



**FINAL REPORT FOR THE PHASE I
REMEDIAL INVESTIGATION AT LOAD
LINE 9 (RVAAP 42)**

**Ravenna Army Ammunition Plant
8451 State Route. 5
Ravenna, Ohio 44266**

Prepared for:



**US Army Joint Munitions Command (JMC)
Rock Island, Illinois 61299-6000**

Prepared By:



**MKM Engineers, Inc.
4153 Bluebonnet Drive
Stafford, TX 77477**

Contract No. DAAA09-02-C-0070

August 2007

**FINAL REPORT FOR THE PHASE I REMEDIAL
INVESTIGATION AT LOAD LINE 9 (RVAAP 42)
AT THE RAVENNA ARMY
AMMUNITION PLANT**



Prepared for

US Army Joint Munitions Command (JMC)
Rock Island, Illinois 61299-6000



Prepared by

MKM ENGINEERS, INC
4153 BLUEBONNET DRIVE
STAFFORD, TEXAS 77477



TABLE OF CONTENTS

LIST OF APPENDICES	vii
LIST OF FIGURES	ix
LIST OF TABLES	xi
LIST OF ACRONYMS AND ABBREVIATIONS	xiv
DOCUMENT DISTRIBUTION	xxi
EXECUTIVE SUMMARY	ES-1
1.0 INTRODUCTION	1-1
1.1 Purpose and Scope	1-1
1.2 Background Information	1-3
1.2.1 General Facility Description	1-3
1.2.2 AOC Description and History	1-4
1.2.3 Previous Investigations	1-6
1.2.4 Regulatory Authorities	1-9
1.3 Report Organization	1-9
2.0 RVAAP ENVIRONMENTAL SETTING	2-1
2.1 Surface Features	2-1
2.2 Meteorology and Climate	2-1
2.3 Surface Water Hydrology	2-3
2.4 Geology	2-4
2.4.1 Glacial Deposits	2-4
2.4.2 Sedimentary Rocks	2-5
2.4.2.1 RVAAP Sedimentary Rocks	2-5
2.4.2.2 LL-9 Sedimentary Bedrock	2-12
2.5 Soils	2-12
2.6 Hydrogeology	2-13
2.6.1 Unconsolidated Sediments	2-13
2.6.2 Bedrock	2-13
2.6.3 Groundwater Utilization	2-14
2.7 Demography and Land Use	2-15
2.8 Ecology	2-15
3.0 LOAD LINE 9 INVESTIGATIONS	3-1
3.1 Field Activities	3-1
3.1.1 MEC Avoidance	3-2
3.1.2 Mowing/Clearing of Sample Locations	3-2
3.1.3 Test Pits	3-2
3.1.4 Work Zones	3-2
3.1.5 Temporary Decontamination Area	3-2
3.1.6 Soil Samples	3-3
3.1.6.1 Azide Screening Soil Samples	3-3
3.1.6.2 RI Soil Samples	3-3
3.1.6.3 Surface Soils	3-5
3.1.6.4 Subsurface Soil Samples	3-7
3.1.7 Groundwater Activities	3-9



1	3.1.7.1	Monitoring Well Installation.....	3-9
2	3.1.7.2	In-Situ Permeability Testing.....	3-11
3	3.1.7.3	Groundwater Sampling.....	3-12
4	3.1.7.4	Water Level Measurements.....	3-12
5	3.1.8	Surface Water.....	3-13
6	3.1.8.1	Azide Screening Surface Water Sample	3-13
7	3.1.8.2	RI Surface Water Samples	3-13
8	3.1.9	Sediment.....	3-15
9	3.1.9.1	Azide Screening Sediment Sample	3-15
10	3.1.9.2	RI Sediment Samples.....	3-15
11	3.1.10	Other RI Samples.....	3-16
12	3.1.10.1	Sewer Manhole Water Sample	3-18
13	3.1.10.2	Sewer Manhole Sediment Sample	3-18
14	3.1.10.3	Subfloor Samples	3-18
15	3.1.10.4	Geotechnical Samples.....	3-19
16	3.1.11	Sampling Location Survey.....	3-19
17	3.1.12	Investigation-Derived Waste	3-20
18	3.1.13	Sample Packaging and Shipping	3-20
19	3.2	Data Analyses and Quality	3-20
20	3.2.1	Field Laboratory Program.....	3-21
21	3.2.2	Laboratory Analysis.....	3-21
22	3.2.3	Data Review, Validation, and Quality Assessment	3-23
23	3.3	Deviations from the Work Plan.....	3-24
24	4.0	NATURE AND EXTENT OF CONTAMINATION	4-1
25	4.1	Soils.....	4-1
26	4.1.1	Azide Screening Surface Soil Results.....	4-2
27	4.1.2	RI Surface Soil Results	4-2
28	4.1.2.1	Explosives and Propellants	4-13
29	4.1.2.2	TAL Metals and Cyanide	4-13
30	4.1.2.3	Pesticides and PCBs	4-15
31	4.1.2.4	Volatiles and Semi-volatiles.....	4-15
32	4.1.3	RI Subsurface Soil Samples.....	4-15
33	4.1.3.1	Explosives and Propellants	4-16
34	4.1.3.2	TAL Metals and Cyanide	4-16
35	4.1.3.3	Pesticides and PCBs	4-23
36	4.1.3.4	Volatiles and Semi-volatiles.....	4-23
37	4.2	Groundwater.....	4-23
38	4.2.1	Hydraulic Conductivity and Water Level Measurements.....	4-23
39	4.2.2	Groundwater Results.....	4-24
40	4.2.2.1	Explosives and Propellants	4-30
41	4.2.2.2	TAL Metals and Cyanide	4-30
42	4.2.2.3	Pesticides and PCBs	4-30
43	4.2.2.4	Volatiles and Semi-volatiles.....	4-31
44	4.3	Surface Water	4-31
45	4.3.1	Azide Screening Results	4-31
46	4.3.2	RI Results.....	4-31
47	4.3.2.1	Explosives and Propellants	4-34
48	4.3.2.2	TAL Metals and Cyanide	4-34
49	4.3.2.3	Pesticides and PCBs	4-36
50	4.3.2.4	Volatiles and Semi-volatiles.....	4-36



1	4.4	Sediment	4-36
2	4.4.1	Azide Screening Results	4-36
3	4.4.2	RI Results.....	4-36
4	4.4.2.1	Explosives and Propellants	4-37
5	4.4.2.2	TAL Metals and Cyanide	4-37
6	4.4.2.3	Pesticides and PCBs	4-42
7	4.4.2.4	Volatiles and Semi-volatiles.....	4-42
8	4.4.2.5	TOC and Grain Size Analyses.....	4-43
9	4.5	Other Analyses.....	4-43
10	4.5.1	Sump and Sewer Manhole Water and Residue	4-43
11	4.5.2	Subfloor.....	4-43
12	4.5.3	Geotechnical Samples	4-43
13	5.0	CONTAMINANT FATE AND TRANSPORT	5-1
14	5.1	Chemical and Physical Properties Affecting Chemical Mobility	5-1
15	5.1.1	Specific Gravity	5-3
16	5.1.2	Vapor Pressure	5-4
17	5.1.3	Water Solubility	5-4
18	5.1.4	Henry's Law Constant	5-5
19	5.1.5	Octanol/Water Partition Coefficient (K_{ow}).....	5-5
20	5.1.6	Organic Carbon-Water Partition Coefficient (K_{oc})	5-5
21	5.1.7	Soil-Water Distribution Coefficient (K_d).....	5-6
22	5.1.8	Chemical Complex Formation	5-7
23	5.1.9	Bioconcentration Factor	5-8
24	5.1.10	Mobility Index	5-8
25	5.1.11	Miscellaneous Characteristics.....	5-9
26	5.2	Chemical Persistence and Degradation Process	5-9
27	5.2.1	Volatile Organic Chemicals	5-13
28	5.2.2	Semi-volatile Organic Chemicals	5-13
29	5.2.3	Pesticides and Polychlorinated Biphenyls.....	5-14
30	5.2.4	Explosive and Propellant Compounds	5-14
31	5.2.5	Metals.....	5-15
32	5.3	Contaminant Transport Pathways.....	5-22
33	5.3.1	Leaching of Soil Contaminants to Groundwater.....	5-23
34	5.3.2	Migration of Groundwater Contaminants.....	5-27
35	5.3.3	Mixing of Groundwater (i.e., Creek Bed Seepage) with Surface Water in Adjacent Stream Channels During Storm Events	5-28
36	5.3.4	Migration of Contaminants from Surface Soil to Surface Water.....	5-28
37	5.3.5	Leaching of Contaminants from Creek Sediment to Surface Water.....	5-28
38	5.3.6	Migration of Contaminants in Surface Water as Dissolved or Sorbed Phases During Storm Events	5-29
39	5.3.7	Volatilization from Soil, Groundwater, or Surface Water	5-29
40	5.4	Conceptual Site Model	5-29
43	6.0	BASELINE HUMAN HEALTH RISK ASSESSMENT	6-1
44	6.1	Introduction.....	6-1
45	6.2	Data Evaluation	6-2
46	6.2.1	Data Collection and Evaluation	6-3
47	6.2.2	Selection of COPCs	6-4
48	6.2.2.1	SRC Screens.....	6-4
49	6.2.2.2	COPC Toxicity Screen.....	6-6



1	6.2.2.3	Screening for Lead	6-7
2	6.2.2.4	COPC Screening Assumptions	6-8
3	6.2.3	COPCs for LL-9	6-8
4	6.2.3.1	Surface Soil	6-8
5	6.2.3.2	Subsurface Soil	6-15
6	6.2.3.3	Deep Surface (0-4 ft bgs) Soils.....	6-15
7	6.2.3.4	Groundwater.....	6-16
8	6.2.3.5	Surface Water.....	6-16
9	6.2.3.6	Sediment	6-17
10	6.2.3.7	Summary of COPC Selection	6-18
11	6.3	Exposure Assessment.....	6-18
12	6.3.1	Exposure Setting	6-18
13	6.3.1.1	Land Use	6-19
14	6.3.1.2	Receptors	6-20
15	6.3.2	Exposure Pathways.....	6-20
16	6.3.2.1	Modified Caretaker – Managed Recreational	6-21
17	6.3.2.2	Security Guard – Maintenance Worker	6-21
18	6.3.2.3	Hunter/Trapper	6-21
19	6.3.2.4	National Guard – Managed Recreational	6-28
20	6.3.2.5	Open Residential	6-28
21	6.3.2.6	Trespasser.....	6-29
22	6.3.3	Exposure Quantification	6-29
23	6.3.3.1	Exposure to Surface Soil	6-29
24	6.3.3.2	Exposure to Subsurface Soil	6-30
25	6.3.3.3	Exposure to Surface Soil and Subsurface Soil	6-30
26	6.3.3.4	Exposure to Groundwater.....	6-30
27	6.3.3.5	Exposure to Surface Water	6-31
28	6.3.3.6	Exposure to Sediment	6-31
29	6.3.3.7	Exposure to Venison	6-32
30	6.3.3.8	Exposure to Beef	6-32
31	6.3.3.9	Exposure to Milk	6-33
32	6.3.3.10	Exposure to Vegetables	6-33
33	6.3.4	Exposure Point Concentrations.....	6-33
34	6.4	Toxicity Assessment.....	6-34
35	6.4.1	Toxicity Information and USEPA Guidance for Non-carcinogens	6-34
36	6.4.2	Toxicity Information and USEPA Guidance for Carcinogens	6-35
37	6.4.3	Estimation of Toxicity Values for Dermal Exposure	6-36
38	6.4.4	Toxicity Criteria Assumptions	6-37
39	6.4.5	Chemicals Lacking USEPA Toxicity Criteria	6-37
40	6.5	Risk Characterization	6-37
41	6.5.1	Methodology.....	6-37
42	6.5.1.1	Methods for Estimation of Carcinogenic Risks	6-37
43	6.5.1.2	Methods for Estimation of Non-carcinogenic Risks	6-38
44	6.5.2	Risk Characterization Results	6-39
45	6.5.2.1	Carcinogenic Risks for LL-9.....	6-46
46	6.5.2.2	Summary of COCs for Each Media/Receptor.....	6-48
47	6.5.2.3	Lead Exposure Analysis	6-49
48	6.5.2.4	Sample Location of Concern	6-49
49	6.6	Uncertainty Analysis.....	6-49
50	6.6.1	Uncertainties Associated with COPC Selection	6-52
51	6.6.1.1	Existing Databases.....	6-52



1	6.6.1.2	Chemicals Potentially Attributable to Background.....	6-52
2	6.6.1.3	COPC Screening Levels.....	6-53
3	6.6.1.4	Absence of COPC Screening Levels.....	6-53
4	6.6.1.5	Frequency of Detection	6-54
5	6.6.1.6	Data Analysis.....	6-54
6	6.6.2	<i>Uncertainties Associated with the Exposure Assessment.....</i>	<i>6-54</i>
7	6.6.2.1	Land Use	6-54
8	6.6.2.2	Exposure Point Concentration.....	6-54
9	6.6.2.3	Exposure Parameters.....	6-54
10	6.6.3	<i>Uncertainties Related to Toxicity Information.....</i>	<i>6-56</i>
11	6.6.3.1	Derivation of Toxicity Criteria.....	6-56
12	6.6.3.2	Chromium Toxicity Criteria	6-57
13	6.6.3.3	Toxicity Criteria for Arsenic.....	6-58
14	6.6.3.4	Toxicity Criteria for Aluminum.....	6-58
15	6.6.4	<i>Uncertainties and Assumptions in the Risk Characterization.....</i>	<i>6-58</i>
16	6.6.4.1	Unavailable or Provisional Toxicity Criteria	6-59
17	6.6.4.2	Foodstuffs Pathway	6-59
18	6.6.4.3	Summation of Risk	6-59
19	6.7	Remedial Goal Options	6-59
20	6.8	Summary and Conclusions	6-61
21	6.8.1	<i>Summary.....</i>	<i>6-61</i>
22	6.8.2	<i>Conclusions.....</i>	<i>6-70</i>
23	7.0	SCREENING-LEVEL ECOLOGICAL RISK ASSESSMENT	7-1
24	7.1	Introduction.....	7-1
25	7.2	Preliminary Problem Formulation	7-2
26	7.2.1	<i>Habitat Types and Ecological Receptors</i>	<i>7-2</i>
27	7.2.2	<i>Major Chemical Sources and Migration Pathways.....</i>	<i>7-2</i>
28	7.2.3	<i>Exposure Routes.....</i>	<i>7-3</i>
29	7.2.3.1	Surface Soil	7-5
30	7.2.3.2	Groundwater.....	7-5
31	7.2.3.3	Surface Water/Sediment	7-6
32	7.2.3.4	Air	7-6
33	7.2.4	<i>Preliminary Assessment and Measurement Endpoints.....</i>	<i>7-6</i>
34	7.2.4.1	Assessment Endpoints	7-6
35	7.2.4.2	Measurement Endpoints.....	7-8
36	7.2.5	<i>Selection of Receptor Species for Food Chain Model.....</i>	<i>7-9</i>
37	7.3	Preliminary Ecological Effects Evaluation	7-9
38	7.4	Preliminary Risk Calculation and Selection of COPC	7-10
39	7.4.1	<i>Selection of Soil COPCs</i>	<i>7-11</i>
40	7.4.2	<i>Selection of Sediment COPCs</i>	<i>7-11</i>
41	7.4.3	<i>Selection of Surface Water COPCs</i>	<i>7-12</i>
42	7.5	Methodology for Refining Conservative Exposure Assumptions.....	7-12
43	7.5.1	<i>Surface Soil.....</i>	<i>7-18</i>
44	7.5.2	<i>Sediment.....</i>	<i>7-21</i>
45	7.5.3	<i>Surface Water</i>	<i>7-23</i>
46	7.5.4	<i>Other Step 3a Factors</i>	<i>7-23</i>
47	7.5.5	<i>Terrestrial Food Chain Modeling.....</i>	<i>7-24</i>
48	7.5.5.1	Characterization of Exposure	7-25
49	7.5.5.2	Ecological Risk Characterization	7-27
50	7.6	Step 3a Refinement.....	7-27



1	7.6.1	<i>Risks to Terrestrial Plants and Invertebrates – Surface Soil</i>	7-27
2	7.6.2	<i>Risks to Benthic Invertebrates – Sediment</i>	7-34
3	7.6.3	<i>Risks to Aquatic Organisms – Surface Water</i>	7-36
4	7.7	<i>Risks to Terrestrial Wildlife</i>	7-40
5	7.7.1	<i>Risks to Insectivorous/Herbivorous Species</i>	7-40
6	7.7.2	<i>Risks to Carnivorous Species</i>	7-45
7	7.8	<i>Ecological Risk Uncertainty Analysis</i>	7-45
8	7.8.1	<i>Measurement and Assessment Endpoints</i>	7-45
9	7.8.2	<i>Exposure Characterization</i>	7-46
10	7.8.3	<i>Ecological Effects Data</i>	7-46
11	7.8.4	<i>Risk Characterization</i>	7-49
12	7.9	<i>Conclusions</i>	7-49
13	7.9.1	<i>Terrestrial Plants and Soil Invertebrates</i>	7-50
14	7.9.2	<i>Benthic Invertebrates</i>	7-50
15	7.9.3	<i>Aquatic Organisms</i>	7-51
16	7.9.4	<i>Terrestrial Wildlife</i>	7-51
17	8.0	SUMMARY AND CONCLUSIONS	8-1
18	8.1	<i>Summary</i>	8-1
19	8.2	<i>Nature and Extent of Contamination</i>	8-1
20	8.3	<i>Contaminant Fate and Transport</i>	8-2
21	8.4	<i>Baseline Human Health Risk Assessment</i>	8-3
22	8.5	<i>Screening Ecological Risk Assessment</i>	8-5
23	9.0	RECOMMENDATIONS	9-1
24	10.0	REFERENCES	10-1
25			



1 LIST OF APPENDICES

- 2 Appendix A Soil Boring Logs
- 3 Appendix B RVAAP List of Rare Species
- 4 Appendix C Photographic Log and Aerial Map
- 5 Appendix D UXO Avoidance Report
- 6 Appendix E Azide Screening
- 7 Appendix F Surface Soil Analytical Results Summary and Laboratory Reports
- 8 Appendix G Surface Soil Field Sampling Forms
- 9 Appendix H Subsurface Soil Field Sampling Forms
- 10 Appendix I Subsurface Soil Analytical Results Summary and Laboratory Reports
- 11 Appendix J Field Technical Change Memorandums
- 12 Appendix K Monitoring Well Data – Boring Logs, Construction Diagrams, Well Development
13 Forms, Purging Forms, Groundwater Sampling Forms, Groundwater Level
14 Elevation Data, and Detailed Core Descriptions
- 15 Appendix L Slug Test Data
- 16 Appendix M Groundwater Analytical Results Summary and Laboratory Reports
- 17 Appendix N Well Survey Data
- 18 Appendix O Surface and Sewer Water Sampling Forms
- 19 Appendix P Surface and Sewer Water Analytical Results and Laboratory Reports
- 20 Appendix Q Sediment Sewer Sediment Sampling Forms
- 21 Appendix R Sediment and Sewer Sediment Analytical Results and Laboratory Reports
- 22 Appendix S Geotechnical Analytical Results and Laboratory Reports
- 23 Appendix T Investigated Derived Waste Sampling Forms, Laboratory Reports & Manifests



1	Appendix U	Quality Control Summary Report and Data Validation Reports
2	Appendix V	Toxicity Profiles and Human Health Risk Assessment Tables
3	Appendix W	Ecological Risk Assessment Tables
4	Appendix X	White Paper for Load Lines 6, 9 & 11
5	Appendix Y	Human Health Risk Assessment Adult and Juvenile Trespasser Receptors
6		Tables
7		



1 LIST OF FIGURES

Figure Number	Title	Page
1-1	RVAAP Location Map	1-2
1-2	Facility Location Map	1-7
1-3	Site Map	1-8
2-1	Glacial Till Map	2-2
2-2	Location of Geologic Cross Sections Map	2-6
2-3	Geologic Cross Section A	2-7
2-4	Geologic Cross Section B	2-8
2-5	Geologic Cross Section C	2-9
2-6	Geologic Cross Section D	2-10
2-7	Bedrock Geology Map	2-11
3-1	Surface Soil Sample Location Map	3-4
3-2	Subsurface Soil Sample Location Map	3-8
3-3	Groundwater Monitoring Well Location Map	3-10
3-4	Surface Water Sample Location Map	3-14
3-5	Sediment Sample Location Map	3-17
4-1	Surface Soil Sample (0-1 ft) Exceedances Explosives/Propellants	4-3
4-2	Surface Soil Sample (0-1 ft) Exceedances TAL Metals/Cyanide	4-4
4-3	Surface Soil Sample (0-1 ft) Exceedances VOCs/SVOCs	4-5
4-4	Surface Soil Sample (0-1 ft) Distribution of Aluminum	4-6
4-5	Surface Soil Sample (0-1 ft) Distribution of Arsenic	4-7



1

Figure Number	Title	Page
4-6	Surface Soil Sample (0-1 ft) Distribution of Chromium	4-8
4-7	Surface Soil Sample (0-1 ft) Distribution of Copper	4-9
4-8	Surface Soil Sample (0-1 ft) Distribution of Lead	4-10
4-9	Surface Soil Sample (0-1 ft) Distribution of Manganese	4-11
4-10	Surface Soil Sample (0-1 ft) Distribution of Mercury	4-12
4-11	Subsurface Soil Sample (>1 ft) Exceedances Explosives/Propellants	4-17
4-12	Subsurface Soil Sample (>1 ft) Exceedances TAL Metals and Cyanide	4-18
4-13	Subsurface Soil Sample (>1 ft) Distribution of Arsenic	4-19
4-14	Subsurface Soil Sample (>1 ft) Distribution of Chromium	4-20
4-15	Distribution of Mercury in Subsurface Soil	4-21
4-16	Potentiometric Surface December 8, 2003	4-25
4-17	Potentiometric Surface December 17, 2003	4-26
4-18	Potentiometric Surface June 25, 2004	4-27
4-19	Groundwater Sample Exceedances Explosives/Propellants	4-28
4-20	Groundwater Sample Exceedances TAL Metals and Cyanide	4-29
4-21	Surface Water Exceedances Explosives/Propellants	4-32
4-22	Surface Water Exceedances TAL Metals and Cyanide	4-33
4-23	Sediment Exceedances Explosives/Propellants	4-38
4-24	Sediment Exceedances TAL Metals and Cyanide	4-39
4-25	Sediment Exceedances VOCs and SVOCs	4-40
7-1	Ecological Conceptual Site Model	7-4



1 LIST OF TABLES

Table Number	Title	Page
1-1	Building Usage And Hazard Classifications For LL-9 at the RVAAP	1-4
3-1	Rationale for Selection of Soil Sampling Intervals	3-5
3-2	Summary of Samples and Analysis	3-6
3-3	LL-9 Groundwater Elevations	3-13
4-1	Summary of Azide Screening Surface Soil Results (0 – 1 ft)	4-44
4-2	Summary of Surface Soil Exceedances (0 – 1 ft bgs)	4-46
4-3	Summary of Surface Soil Results (0 – 1 ft bgs)	4-61
4-4	Summary of Subsurface Soil Results (>1 ft)	4-86
4-5	Summary of Subsurface Soil Exceedances (>1 ft)	4-115
4-6	Hydraulic Conductivity and Water Level Measurements	4-24
4-7	Summary of Groundwater Results	4-126
4-8	Summary of Groundwater Exceedances	4-130
4-9	Azide Screening Surface Water Sample Results Summary	4-131
4-10	Summary of Surface Water Exceedances	4-133
4-11	Summary of Surface Water Results	4-137
4-12	Summary of Sediment Exceedances	4-141
4-13	Summary of Sediment Results	4-147
5-1	Environmental Fate and Transport Properties of Organic Chemicals	5-2
5-2	Relative Mobility of Select Materials under Various Environmental Conditions of Eh and pH	5-3



1

Table Number	Title	Page
5-3	Soil-Water Partition Coefficients for Metals	5-6
5-4	Ranges of Biodegradation Rates for Representative Organic Chemicals	5-11
6-1	Human Health Risk Assessment Dataset for Surface Soils	6-9
6-2	Human Health Risk Assessment Dataset for Subsurface Soils	6-10
6-3	Human Health Risk Assessment Dataset for Deep Surface Soils	6-11
6-4	Human Health Risk Assessment Dataset for Groundwater	6-13
6-5	Human Health Risk Assessment Dataset for Surface Water	6-13
6-6	Human Health Risk Assessment Dataset for Sediments	6-13
6-7	Summary of Quantitative and Qualitative COPCs for Environmental Media	6-14
6-8	Potential Receptors for the LL-9 BHHRA	6-20
6-9	Receptors and Exposure Pathways	6-22
6-10	Exposure Assumptions	6-23
6-11	Summary of Cancer Risks and Hazard Indices	6-40
6-12	Non-Carcinogenic Risks for LL-9	6-43
6-13	Incremental Lifetime Cancer Risk Results	6-46
6-14	Potential Land Use/Receptor/Medium Pathways COCs	6-50
6-15	Surface Soil (0 to 1 ft bgs) RGOs	6-62
6-16	Subsurface Soil RGOs	6-63
6-17	National Guard Trainee Deep Surface Soil (0-4 ft bgs) RGOs	6-64
6-18	Groundwater RGOs	6-65



1

Table Number	Title	Page
6-19	Surface Water RGOs	6-66
7-1	Ecological COPC Selection Table of Shallow Soil (0 – 4 ft bgs)	7-13
7-2	Ecological COPC Selection Table for Sediment	7-16
7-3	Ecological COPC Selection Table for Surface Water	7-17
7-4	Direct Toxicity to Earthworms	7-19
7-5	Direct Toxicity to Plants	7-20
7-6	Step 3a Table for Evaluating Risks to Sediment Invertebrates	7-22
7-7	Terrestrial Food Chain Model– Conservative Scenario (Insectivorous, Herbivorous and Carnivorous Receptors)	7-41
7-8	Terrestrial Food Chain Model– Average Scenario (Insectivorous, Herbivorous and Carnivorous Receptors)	7-43
7-9	GPL Explosives Data Shallow Soil Samples (0–4 ft bgs)	7-48
8-1	Summary of COPCs Selected for the Quantitative Human Health Risk Assessment	8-3
8-2	Human Health Risk Assessment Calculation Summary	8-4
8-3	Ecological Risk Calculations	8-6

2



1 LIST OF ACRONYMS AND ABBREVIATIONS

2	95% UCL-L	log-normal 95% upper confidence level
3	95% UCL-N	normal 95% upper confidence level
4	AET	apparent effects threshold
5	ABS _{GI}	gastrointestinal absorption fraction
6	amsl	above mean sea level
7	AOC	area of concern
8	AST	above-ground storage tank
9	ATSDR	Agent for Toxic Substances and Disease Registry
10	AWQC	ambient water quality criteria
11	BCF	bioconcentration factor
12	BERA	baseline ecological risk assessment
13	bgs	below ground surface
14	BHHRA	baseline human health risk assessment
15	BRACD	Base Realignment and Closure Division
16	CERCLA	Comprehensive Environmental Response, Compensation, and Liability Act
17	C	Celsius
18	CDI	chronic daily intake
19	cm/s	centimeter(s) per second
20	COC	chemical of concern
21	COPC	chemical of potential concern
22	CSF	cancer slope factor



1	CSM	conceptual site model
2	DLF	dust loading factor
3	DoD	Department of Defense
4	DQO	data quality objective
5	EB	explosive building
6	Eco-SSL	ecological soil screening level
7	EDD	electronic data deliverable
8	EDQL	ecological data quality level
9	EEQ	estrogenic equivalent
10	Eh	oxidation–reduction potential
11	EPC	exposure point concentration
12	ESL	ecological screening level
13	ESV	ecological screening value
14	F	Fahrenheit
15	F _{oc}	fraction of organic carbon
16	ft	foot or feet
17	FWSAP	Facility–Wide Sampling and Analysis Plan
18	GAF	gastrointestinal absorption factor
19	GOCO	government–owned, contractor–operated
20	GPD/ft	gallon(s) per day per foot
21	GPM	gallon(s) per minute
22	GPL	General Physics Laboratories



1	HI	hazard index
2	HQ	hazard quotient
3	HSA	hollow-stem auger
4	IDW	investigation-derived waste
5	ILCR	incremental lifetime cancer risk
6	in	inch
7	IOC	Industrial Operations Command
8	IRP	Installation Restoration Program
9	JMC	Joint Munitions Command
10	K _d	soil-water distribution coefficient
11	K _{oc}	organic carbon partition coefficient
12	K _{ow}	octanol/water partition coefficient
13	km	kilometer(s)
14	L	liter(s)
15	LL	load line
16	LOAEL	lowest-observed-adverse-effect level
17	LOEC	lowest-observed-effect concentration
18	LPD/m	liter(s) per day per meter
19	LPM	liter(s) per minute
20	m	meter(s)
21	MCL	maximum contaminant level
22	MEC	munitions and explosives of concern



1	mg/kg	milligrams per kilogram
2	mg/L	milligrams per liter
3	mg/m ³	milligrams per cubic meter
4	MI	mobility index
5	MKM	MKM Engineers, Inc.
6	ml/g	milliliters per gram
7	mm	millimeter(s)
8	mph	miles per hour
9	MS/MSD	matrix spike/matrix spike duplicate
10	MW	monitoring well
11	NAAQS	National Ambient Air Quality Standards
12	NAPL	non-aqueous phase liquids
13	nc	non-carcinogenic
14	NCEA	National Center for Environmental Assessment
15	NEB	non-explosive building
16	NO ₃	nitrate
17	NOAEL	no-observed-adverse-effect level
18	NOEC	no-observed-effect concentration
19	NTU	Nephelometric Turbidity Unit
20	NWS	National Weather Service
21	Ohio EPA	Ohio Environmental Protection Agency
22	OHARNG	Ohio Army National Guard



1	OMZM	outside mixing zone maximum
2	PAH	polycyclic aromatic hydrocarbon
3	PBT	persistent, bioaccumulative, and toxic
4	PCB	polychlorinated biphenyl
5	PEC	probable effects concentration
6	PEF	particulate emission factor
7	pH	measurement used to express the intensity of the acid or alkaline condition of a
8		solution
9	PID	photoionization detector
10	PM10	particulate matter ≤ 10 microns in diameter
11	PPRTV	Peer-Reviewed Toxicity Value
12	PRG	preliminary remediation goal
13	PVC	polyvinyl chloride
14	QA	quality assurance
15	QC	quality control
16	QCSR	quality control summary report
17	RAGS	Risk Assessment Guidance for Superfund
18	RBSC	risk-based screening concentration
19	RCRA	Resource Conservation and Recovery Act
20	RDA	recommended daily allowance
21	RDI	recommended daily intake
22	RDF	regional disposal facility
23	RDX	hexahydro-1,3,5-trinitro-1,3,5-triazine



1	RfC	reference concentration
2	RfD	reference dose
3	RGO	remedial goal option
4	RI	remedial investigation
5	RME	reasonable maximum exposure
6	RRSE	relative risk site evaluation
7	RTLS	Ravenna Training and Logistics Site
8	RVAAP	Ravenna Army Ammunition Plant
9	SAIC	Science Applications International Corporation
10	SdSL	sediment screening level
11	SDWA	Safe Drinking Water Act
12	SERA	Screening Ecological Risk Assessment
13	SQG	soil quality guideline
14	SRC	site related chemical
15	SRV	sediment reference value
16	SSL	soil screening levels
17	SSSL	shallow soil screening level
18	STL	Severn Trent Laboratories, Inc.
19	STSC	Superfund Health Risk Technical Support Center
20	SVOC	semi-volatile organic compound
21	SWSL	surface water screening level
22	TAL	target analyte list



1	TCE	trichloroethene
2	TEC	threshold effect concentration
3	TNT	2,4,6-trinitrotoluene
4	TOC	total organic carbon
5	TRV	toxicity reference value
6	TSCA	Toxic Substances Control Act
7	UCL	upper confidence level
8	µg/kg	micrograms per kilogram
9	µg/L	micrograms per liter
10	USACE	U.S. Army Corps of Engineers
11	USAEC	U.S. Army Environmental Command
12	USACHPPM	U.S. Army Center for Health Promotion and Preventive Medicine
13	USEPA	U.S. Environmental Protection Agency
14	USP&FO	U.S. Property and Fiscal Officer
15	UTL	upper tolerance limit
16	UXO	unexploded ordnance
17	VF	volitization factor
18	VOC	volatile organic compound
19	WBG	Winklepeck Burning Grounds
20	WQC	water quality criteria



1 DOCUMENT DISTRIBUTION

Name/Organization	No. of Hard Copies	No. of Electronic Copies
Eileen Mohr/Ohio Environmental Protection Agency (Ohio EPA)	2	2
Bonnie Buthker/Ohio EPA	1	1
Irv Venger/Ravenna Army Ammunition Plant (RVAAP)	2	2
Major Meade/Ohio Army National Guard (OHARNG)	1	1
John Jent/United States Army Corps of Engineers (USACE) Louisville District	2	2
Mark Krivansky/United States Army Environmental Command (USAEC)	0	1
Adam Deck/United States Army Center for Health Promotion and Preventive Medicine (USACHPPM)	0	1



EXECUTIVE SUMMARY

This Phase I Remedial Investigation (RI) Report characterizes the nature and extent of contamination, evaluates the fate and transport of contaminants, and assesses the potential risk to human health and the environment resulting from past operations at Load Line 9 (LL-9), also known as RVAAP-42, at the Ravenna Army Ammunition Plant (RVAAP), within Portage and Trumbull counties, Ravenna, Ohio. LL-9 is located in the south-central region of the RVAAP along Fuze and Booster Road.

The RI was conducted to fulfill the requirements of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) and to prioritize future environmental restoration efforts at RVAAP's areas of concern (AOCs). The results of a United States Army Center for Health Promotion and Preventive Medicine (USACHPPM) investigation and an azide screening operation, as well as past activities at LL-9, were used to develop the objectives and sampling design of the RI. The objectives of the RI were to:

- Define the vertical and horizontal extent of contamination at LL-9;
- Gather sufficient data to conduct a Baseline Human Health Risk Assessment (BHHRA) and a Screening Ecological Risk Assessment (SERA);
- Define potential transport pathways and receptor populations; and
- Provide sufficient data for the selection of remedial action alternatives.

To fulfill these objectives, field investigations were conducted in March 2002 and October to December 2003. During the field investigations, samples were collected from surface soils, subsurface soil borings, surface water, sediments, and groundwater. Samples were analyzed for possible contamination resulting from the production of artillery fuzes such as explosives, propellants, target analyte list (TAL) metals, nitrates, cyanide, sulfides, semi-volatile organic compounds (SVOCs), and volatile organic compounds (VOCs). The surface and subsurface soil, groundwater, surface water, and sediment results are summarized for each matrix.

Surface Soils (0 to 1 ft below ground surface [bgs])

Propellants (nitrocellulose and nitroguanidine), SVOCs (benzo[a]pyrene and dibenzo[a,h]anthracene) and metals (aluminum, arsenic, barium, cadmium, chromium, cobalt, copper, iron, lead, manganese, mercury, nickel, selenium, vanadium, and zinc) were detected in surface soils at concentrations greater than RVAAP installation background (USACE, 2001b) concentrations, Region 9 residential soil preliminary remediation goals (PRGs), or both.



1 **Subsurface Soils (1 – 3 ft bgs)**

2 Aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, iron, lead, manganese,
3 mercury, vanadium, and zinc were detected at concentrations greater than the RVAAP
4 installation background concentrations, Region 9 residential soil PRGs, or both. In addition, one
5 propellant (nitrocellulose) and one explosive (hexahydro-1,3,5-trinitro-1,3,5-triazine [RDX])
6 were detected in subsurface soils. However, only 10% of the subsurface samples collected
7 were analyzed for explosives and propellants.

8 **Groundwater**

9 Nitrocellulose and five metals (antimony, cobalt, copper, iron, and manganese) were detected in
10 seven groundwater samples (and 1 duplicate) at concentrations exceeding the RVAAP-specific
11 consolidated bedrock background criteria, the Region 9 tap water PRG or both. However, the
12 nitrocellulose concentrations results were qualified as estimated (J), or (B) the concentration of
13 nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J).

14 **Surface Water**

15 Metals including aluminum, arsenic, barium, beryllium, chromium, cobalt, copper, lead,
16 manganese, mercury, nickel, vanadium, and zinc were detected in surface water at
17 concentration exceeding the RVAAP-specific surface water installation background values
18 established by USACE (2000), the Region 9 tap water PRGs, or both. Nitrocellulose was
19 detected in two samples at concentrations above the method detection limit but below the
20 reporting limit. However, the nitrocellulose concentrations results were qualified as estimated
21 (J), or (B) the concentration of nitrocellulose in the method blank was detected above the
22 reporting limit, or both (B) (J).

23 **Sediments**

24 Propellants (nitrocellulose and nitroguanidine), SVOCs (benzo(a)pyrene, benzo(b)fluoranthene,
25 dibenzo(a,h)anthracene, indeno(1,2,3-cd)pyrene) and metals (aluminum, antimony, arsenic,
26 barium, beryllium, cadmium, chromium, cobalt, copper, lead, manganese, mercury, nickel,
27 selenium and vanadium) were detected in sediment samples at concentrations exceeding the
28 RVAAP-specific sediment background values established by USACE (2000), Region 9
29 residential soil PRGs, or both.

30 **Fate and Transport**

31 The primary contaminant migration pathways for the contaminants identified during the RI
32 conducted at LL-9 are:

- 33 • Leaching from soils to groundwater,
- 34 • Leaching from sediments to surface water,



- Leaching from sewer lines to groundwater, and
- Transport in surface drainage channels.

Baseline Human Health Risk Assessment

A BHHRA was performed to assess the potential current and future risks associated with human exposure to site-related contaminants found in LL-9. Risks were evaluated for a National Guard trainee (the primary receptor of concern), a National Guard resident/trainer, a hunter/trapper, security guard/maintenance worker, and a resident farmer (adult and child). The risk assessment demonstrated that some potential risks exist for the following receptors:

- National Guard resident trainer – $HI > 1$; the total cancer risk estimate is within United States Environmental Protection Agency (USEPA) target incremental cancer risk range of $1E-04$ to $1E-06$, and exceeds Ohio Environmental Protection Agency's (Ohio EPA) target risk value of $1E-05$.
- National guard trainee – Total $HI > 1$; the total cancer risk estimate is within USEPA target incremental cancer risk range of $1E-04$ to $1E-06$, and exceeds Ohio EPA's target risk value of $1E-05$.
- Adult resident farmer – $HI > 1$; the total cancer risk estimate exceeds the USEPA target incremental cancer risk range of $1E-04$ to $1E-06$, and exceeds Ohio EPA's target risk value of $1E-05$.
- Child resident farmer – $HI > 1$; the total cancer risk estimate exceeds the USEPA target incremental cancer risk range of $1E-04$ to $1E-06$, and exceeds Ohio EPA's target risk value of $1E-05$.
- Lifelong resident farmer – The total cancer risk estimate is within USEPA target incremental cancer risk range of $1E-04$ to $1E-06$, and exceeds Ohio EPA's target risk value of $1E-05$.

Several significant uncertainties associated with the risk assessment were identified and must be considered carefully when making any risk management decisions. A significant uncertainty is the fact that the nature and extent of the contamination in soil, sediment, surface water, and groundwater at LL-9 have not been completely defined. Therefore, the results of the human health risk assessment are preliminary and subject to change based on the data collected in the Phase II RI.

Screening Ecological Risk Assessment

A SERA was performed to assess whether adverse ecological impacts are present as a result of site-related contaminants found in LL-9. Ecological impact was evaluated for plants, soil and sediment invertebrates, aquatic organisms, and terrestrial wildlife. Three types of mammals and



birds were evaluated: insectivores, herbivores, and carnivores. The ecological risk calculations showed that the only significant potential risks for terrestrial plants and soil invertebrates are from metals (copper, lead, mercury, and zinc) in the area surrounding sample location LL9SS-011-0001-SO. However, the nature and extent of the contamination in soil, sediment, surface water, and groundwater at LL-9 have not been determined. Therefore, the results of the ecological risk assessment are preliminary and subject to change based on the data collected in the Phase II RI.

Conclusions

Contaminants in surface soil, subsurface soil and sediments were detected at concentrations greater than the RVAAP installation background concentrations, Region 9 residential soil PRGs, or both. The lateral and vertical extent of contamination including explosives, propellants, metals, and SVOCs in surface soils, subsurface soil, and sediments were not fully delineated during the Phase I RI. In addition, the presence of metals in groundwater and surface water at concentrations greater than RVAAP-specific background criteria (USACE, 2001b) indicate that contamination present in soils and sediment may be leaching into surface water and groundwater.

The Phase I risk assessment demonstrated that potential risks exist for National Guard resident/trainer and a resident farmer (adult and child). The total cancer risk estimate summarized for each media (2E-05) for the primary receptor of concern, the National Guard trainee, is within the USEPA target cancer risk range (1E-04 to 1E-06) but exceeds the Ohio EPA target risk criterion of 1E-05. The total non-cancer risk (HI) estimate summarized for each media for the primary receptor of concern equals 2. Manganese (inhalation of air particulates being the route of exposure) was the significant contributor to the non-carcinogenic risk. However, significant uncertainties were identified, including the risk estimates for arsenic, manganese, and chromium in soils, and for the inhalation pathway and should be considered by the risk management team for RVAAP when making further remedial decisions for LL-9.

Copper, lead, mercury, and zinc were retained as chemicals of potential concern (COPCs) for terrestrial plants and soil invertebrates in the area surrounding sample location LL9SS-011-0001-SO. No COPCs were retained for direct toxicity to benthic invertebrates, aquatic organism, and terrestrial wildlife.

Recommendations

Concentrations of target analytes (primarily metals, SVOCs, and propellants) were detected in surface soil, subsurface soil, and sediment samples at concentrations exceeding RVAAP installation background (USACE, 2001b) concentrations, Region 9 residential soil PRGs, or both. Most of the contaminants (except for metals) were detected in very few samples and there are insufficient occurrences to provide statistically valid analysis for contaminant distribution. Therefore, the nature and extent of the contamination in soil, sediment, surface



1 water and groundwater at LL-9 have not been fully determined and additional sampling is
2 necessary to fully identify the lateral and vertical extent of the contamination at LL-9. A Phase
3 II RI should be conducted to fill data gaps identified in this Phase I RI.

4 The conclusions of the risk assessments are based on available data. If data collected in the
5 Phase II RI alters the conceptual site model presented in this Phase I RI, then the risk
6 assessment findings presented in this report will require revision. Although the BHHRA
7 indicated potential risk to a National Guard trainee due to exposure to arsenic, chromium, and
8 manganese, conservative toxicity and exposure values (such as PEF) were used to calculate
9 the potential risk. Thus, the risk estimates presented for those chemicals likely overestimate the
10 potential for non-carcinogenic risk. It is recommended that the risk management team consider
11 the need for any further remedial action based on the risk assessment results presented for this
12 receptor. Copper, lead, mercury, and zinc in surface soil require further evaluation to determine
13 the extent of contamination because these chemicals may pose a risk to plants and
14 invertebrates.



1.0 INTRODUCTION

This report documents the results of the Phase I Remedial Investigation (RI) conducted at Load Line 9 (LL-9) by MKM Engineers, Inc. (MKM) under Contract Number DAAA09-98-G-0001/DAAA09-02-C-0070. LL-9 is located at the U. S. Army Base Realignment and Closure Division (BRACD) Ravenna Army Ammunition Plant (RVAAP), Ravenna, Ohio (Figure 1-1). The RI was conducted under the U.S. Department of Defense (DoD) Installation Restoration Program (IRP) and performed in accordance with the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA). The RI was conducted in accordance with work plans reviewed and approved by the Ohio Environmental Protection Agency (Ohio EPA) (MKM, 2003a).

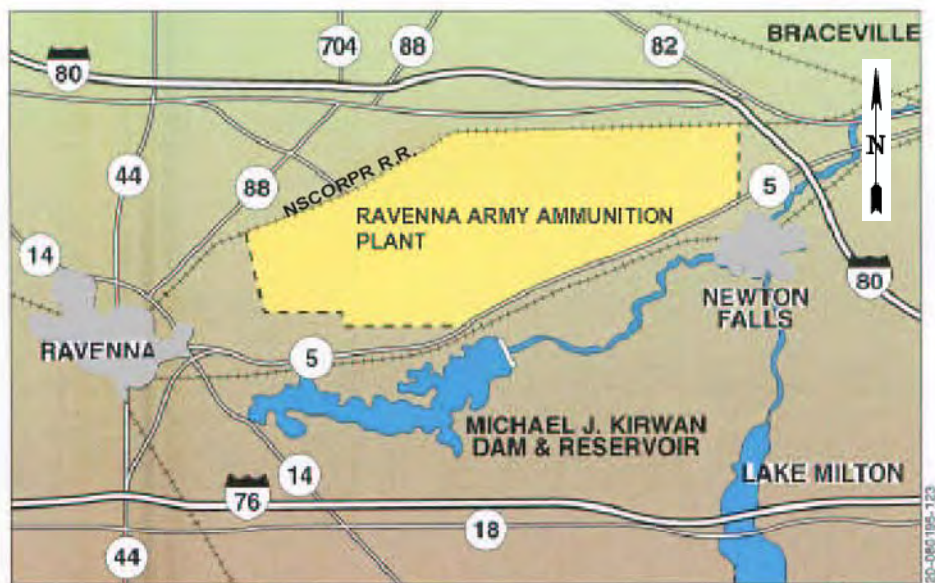
This document summarizes the results of the field activities conducted during March 2002 (azide screening) and the RI conducted from October 2003 to December 2003 at LL-9. The environmental setting, fieldwork, nature, and extent of contamination are discussed. Human health and ecological risk assessments were performed as part of the RI. Results of the data analysis and risk assessments were used to develop a revised conceptual model for LL-9. The conclusions and recommendations of the RI form the framework for decisions regarding future remedial actions at LL-9.

1.1 Purpose and Scope

The purposes of this RI Report are to (1) describe the investigations conducted at RVAAP LL-9; (2) evaluate and identify the environmental impact posed by past DoD industrial operations; (3) determine if there is contamination that poses a risk; or exceeds a preliminary remediation goal (PRG); (cleanup criteria established by the United States Environmental Protection Agency [USEPA]). The specific objectives of the RI are to:

- Describe the physical environment of LL-9 and its surroundings to the extent necessary to define potential transport pathways and receptor populations and provide sufficient data for preliminary screening of remedial action alternatives.
- Characterize the nature and extent of contamination at LL-9 such that a baseline risk assessment can be conducted to evaluate the potential risks to human health and the environment.
- Identify the sources of contamination at LL-9 to evaluate selection of remedial actions. Source locations, types and amounts of contaminants, potential releases, physical and chemical properties of wastes present and engineering characteristics have been evaluated.

Investigation-specific objectives were developed using the data quality objective (DQO) approach presented in the *Final Facility-Wide Sampling and Analysis Plan (FWSAP)* for Ravenna Army Ammunition Plant (USACE, 2001a).



LOCATION MAP



NOT TO SCALE

Figure 1-1. Ravenna Army Ammunition Plant Location Map



The investigative approach to the RI at LL-9 involved a combination of field and laboratory activities to characterize the area of concern (AOC). Field investigation techniques included shallow soil sampling, subsurface soil sampling, surface water sampling, sediment sampling, monitoring well (MW) installation, groundwater sampling, and aquifer testing. The field program was conducted in accordance with the FWSAP (USACE, 2001a) and the *FWSAP Addendum for the Remedial Investigation at Load Line 9 (RVAAP 42)* (MKM, 2003a).

1.2 Background Information

This section briefly describes the RVAAP facility and LL-9, previous investigations, and the regulatory guidance followed when conducting the RI.

1.2.1 General Facility Description

Until 1999, the RVAAP was defined as a 21,419 acre installation. The Ohio Army National Guard (OHARNG) surveyed the property boundary and the total acreage was found to be 21,683.289 acres. As of February 2006, a total of 20,403 acres of the former 21,683 acre RVAAP has been transferred to the United States Property and Fiscal Officer (USP&FO) for Ohio for use as an OHARNG training site. Currently, RVAAP consists of 1,280 acres in several distinct parcels scattered throughout the confines of the OHARNG's Ravenna Training and Logistics Site (RTLS). RVAAP's remaining parcels of land are located completely within the RTLS, and are completely enclosed by the RTLS perimeter fence.

The RTLS is located in northeastern Ohio within Portage and Trumbull counties, approximately 4.8 kilometers (km) (3 miles) east/northeast of the City of Ravenna and approximately 1.6 km (1 mile) northwest of the City of Newton Falls. The RVAAP portions of the property are completely located within Portage County. The RTLS (inclusive of RVAAP) is a parcel of property approximately 17.7 km (11 miles) long and 5.6 km (3.5 miles) wide. The facility is bounded by State Route 5, the Michael J. Kirwan Reservoir, and the CSX System Railroad on the south; Garrett, McCormick, and Berry Roads on the west; the Norfolk Southern Corporation railroad on the north; and State Route 534 on the east.

The RTLS is surrounded by several communities: Windham on the north, Garrettsville 9.6 km (6 miles) to the northwest; City of Newton Falls 1.6 km (1 mile) to the southeast; Charlestown to the southwest; and Wayland 4.8 km (3 miles) to the south. RTLS did not exist when the RVAAP was operational, and the entire 21,683-acre parcel was a government-owned, contractor-operated (GOCO) industrial facility. The RVAAP Installation Restoration Program (IRP) encompasses investigation and cleanup of past activities over the entire 21,683 acres of the former RVAAP, so references to the RVAAP in this document consider the historical extent of the RVAAP, inclusive of the combined acreages of the current RTLS and RVAAP, unless otherwise specifically stated.



DoD activities at RVAAP began in 1940 and have included manufacturing, loading, handling, storage, and disposition of military ammunition and explosives. The industrial operations at RVAAP consisted of 12 munitions assembly facilities, referred to as load lines (Figure 1-2). Load Lines 1-4 and 12 were used as melt pour facilities to load high explosives into projectiles and bombs. Load Lines 5, 6, 7, 8, and 10 were used to manufacture fuzes, primers, and boosters. LL-9 was used to produce detonators. Potential contaminants in these load lines may include lead azide, lead styphnate, lead thiocyanate, antimony sulfide, potassium chlorate, mercury fulminate, black powder, 2,4,6-trinitrotoluene (TNT), composition B, octol, nitrocellulose, nitroguanidine, and nitroglycerin.

1.2.2 AOC Description and History

LL-9, which is approximately 69 acres in size, is located in the south-central portion of RVAAP at the intersection of Fuze and Booster Road and George Road (Figure 1-3). From 1941 to 1945, LL-9 produced detonators. In 1945, the load line was deactivated, and the equipment was removed. There are no documented activities at LL-9 since closure in 1945.

Infrastructure at LL-9 consists mainly of a gravel road that follows the perimeter of main production area. The buildings at LL-9 were thermally decontaminated and demolished in 2003 (MKM, 2003b). Demolition was completed to 2 ft below ground surface (bgs) and the foundations were subsequently removed by Lakeshore Engineering. The concrete and brick were crushed for reuse at the RVAAP to maintain the roads. An unused water tower is the only structure remaining at LL-9. Table 1-1 lists the building usage and hazard classifications for LL-9 at the RVAAP.

Table 1-1

Building Usage and Hazard Classifications for LL-9 at the RVAAP

BLDG. #	Function	Explosive	Reference	Hazard Class
DT-1	Napkin Prep Fulminate	Mercury Fulminate	Property Cards and Maps	EB
DT-2	Mix House Fulminate	Mercury Fulminate	Property Cards and Maps	EB
DT-3	Dry House Fulminate	Mercury Fulminate	Property Cards and Maps	EB
DT-4	Receptacle House Azide	Lead Azide	Property Cards and Maps	EB
DT-5	Mix House Azide	Lead Azide	Property Cards and Maps	EB
DT-6	Dry House Azide	Lead Azide	Property Cards and Maps	EB
DT-7	Screen House Azide	Lead Azide	Property Cards and Maps	EB
DT-8	Dry House Azide	Lead Azide	Property Cards and Maps	EB
DT-9	Dry House Azide	Lead Azide	Property Cards and Maps	EB



1

Table 1-1 (Continued)

2

Building Usage and Hazard Classifications for LL-9 at the RVAAP

BLDG. #	Function	Explosive	Reference	Hazard Class
DT-10	Vacuum Pump House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-11	Napkin Prep Azide	Lead Azide	Property Cards and Maps	EB
DT-12	Saw Dust Storage	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-13	Final Insp, Pack, & Ship	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-14	Detonator Rumbling	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-15	Detonator Rest House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-16	M22 Booster Det Assembly	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-17	M22 Charge House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-18	Detonator Test Bldg	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-18A	Detonator Test Bldg	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-19	Prep Pimer Mix Azide Mag	Lead Azide	Property Cards and Maps	EB
DT-20	Loading Bldg	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-21	Loading Bldg	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-22	Loading Bldg	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-23	Tetryl Pellet Mag	Tetryl	Property Cards and Maps	EB
DT-24	Tetryl Pellet Bldg	Tetryl	Property Cards and Maps	EB
DT-25	Service Mag Fulminate	Mercury Fulminate	Property Cards and Maps	EB
DT-26	Tetryl Screening & Blending	Tetryl	Property Cards and Maps	EB
DT-27	Tetryl Service Mag	Tetryl	Property Cards and Maps	EB
DT-28	Change House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-29	Change House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-30*	Inert Storage	n/a	Property Cards and Maps	NEB
DT-31*	Shipping Bldg	n/a	Property Cards and Maps	NEB
DT-33	Solvent Storage	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-34	Detonator Destroying House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB



Table 1-1 (Continued)

Building Usage and Hazard Classifications for LL-9 at the RVAAP

BLDG. #	Function	Explosive	Reference	Hazard Class
DT-35	Control House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-52	Dining Facility	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-54	Dry House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-55	Dry House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
DT-56	Dry House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB
9-51	Gate House	Lead Azide/Mercury Fulminate	Property Cards and Maps	EB

Notes:

EB – Explosive building (Shaded)

NEB – Non-explosive building (Un-shaded)

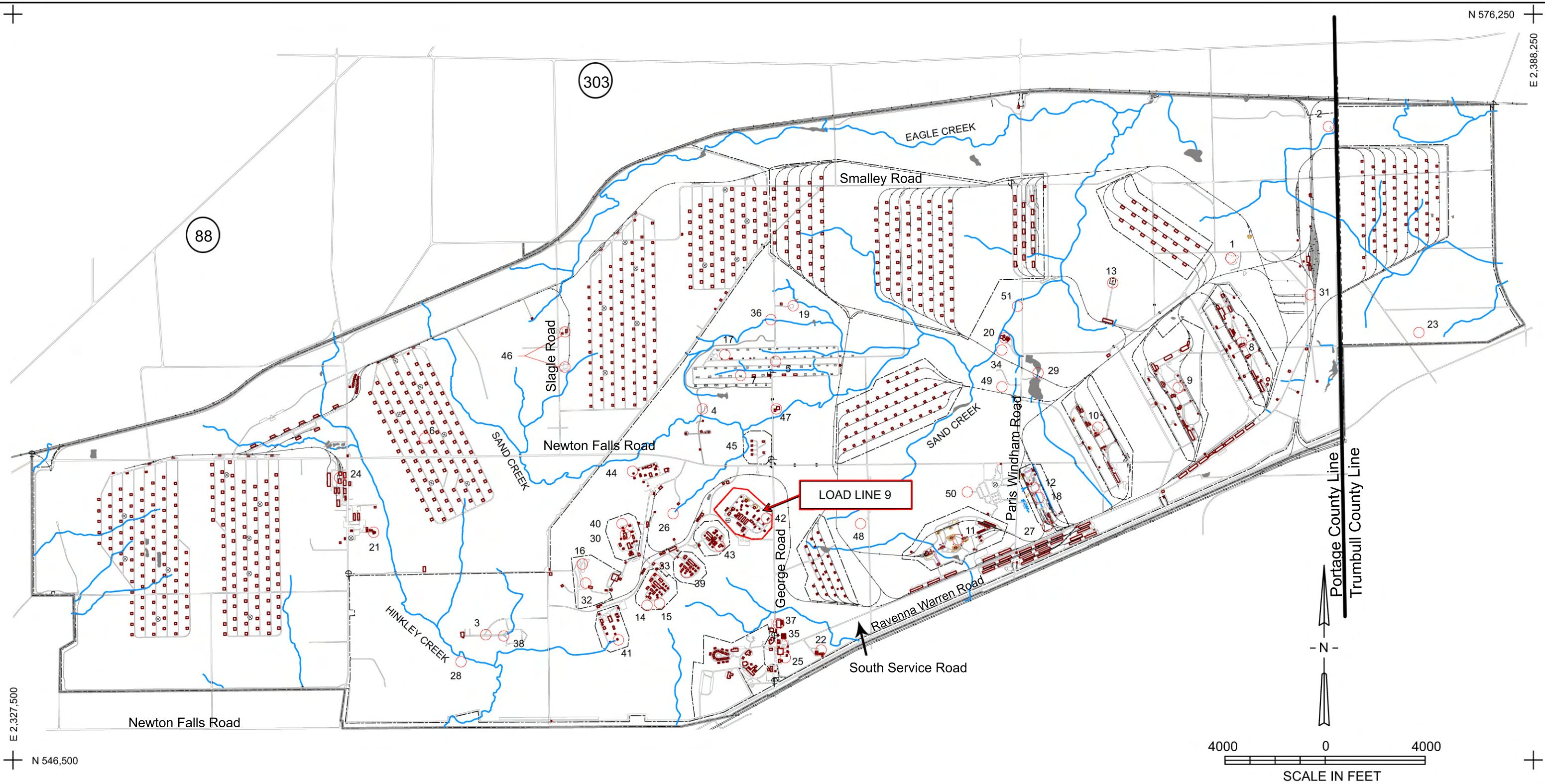
* Un-shaded building numbers indicate a NEB.

1.2.3 Previous Investigations

LL-9 has undergone several investigations since its closure as a production facility. The U.S. Army Center for Health Promotion and Preventive Medicine (USACHPPM) conducted a Relative Risk Site Evaluation (RRSE) for newly added sites at the RVAAP in 1998. The samples were analyzed for explosives and metals. Based on sampling results, the 1998 USACHPPM report identified surface/subsurface soils and groundwater as potential media for contaminant migration. Hunters and recyclers (contractors removing recyclable materials) were identified as potential receptors. The Relative Risk Site Evaluation for the AOC was scored Medium.

In March 2002, the Operational Support Command-RVAAP, in conjunction with USACE, conducted an azide screening operation at LL-9 for the purposes of evaluating the health and safety of future operations as well as to provide additional information for the selection of analytes for the RI. Samples were collected from strategic locations within the load line to screen for primary (azide) and secondary (TNT and hexahydro-1,3,5-trinitro-1,3,5-triazine [RDX]) explosives. Results from the March 2002 sample screening indicate that there is no detectable safety concern related to azide contamination at LL-9. The March 2002 screening results also indicated that there is minimal contamination of secondary explosives at LL-9.

Six shallow soil, three surface water, and three sediment samples were collected during the March 2002 azide screening operation and submitted for laboratory analysis of the TAL metals list. Metals were reported in concentrations that exceeded the RVAAP installation background criteria (USACE, 2001b) values, Region 9 PRGs for residential soil, or both. Since these data were confirmed by laboratory analyses, the results are included in the RI and subsequent risk assessment and are discussed in various portions of this document.




LEGEND OF SITES:

1 RAMSDELL QUARRY LANDFILL	13 BLDG 1200 AND DILUTION/SETTLING POND	25 BLDG 1034 MOTOR POOL WASTE OIL TANK	37 PESTICIDE BUILDING S-44S2	49 CENTRAL BURN PITS
2 ERIE BURNING GROUNDS	14 LOAD LINE 6, EVAPORATION UNIT	26 FUZE BOOSTER AREA SETTLING TANKS	38 NACA TEST AREA	50 ATLAS SCRAP YARD
3 DEMOLITIONS AREA #1	15 LOAD LINE 6, TREATMENT PLANT	27 BLDG 854-PCB STORAGE	39 LOAD LINE 5/FUZE LINE 1	51 DUMP ALONG PARIS - WINDHAM ROAD
4 DEMOLITIONS AREA #2	16 QUARRY LANDFILL/FORMER FUZE & BOOSTER BURNING PITS	28 MUSTARD AGENT BURIAL SITE	40 LOAD LINE 7/BOOSTER LINE 1	+++++ RAILROAD TRACKS
5 WINKLEPECK BURNING GROUNDS	17 DEACTIVATION FURNACE	29 UPPER AND LOWER COBBS POND COMPLEX	41 LOAD LINE 8/BOOSTER LINE 1 FENCELINE
6 C BLOCK QUARRY	18 LOAD LINE 12 PINK WASTE WATER TREATMENT	30 TREATMENT PLANT LOAD LINE 7 PINK WASTEWATER	42 LOAD LINE 9/DETONATOR LINE PROPERTY BOUNDARY
7 BLDG 1601 HAZARDOUS WASTE STORAGE	19 LANDFILL NORTH OF WINKLEPECK BURNING GROUND	31 ORE PILE RETENTION POND	43 LOAD LINE 10/PERCUSSION ELEMENT STREAM OR CREEK
8 LOAD LINE 1 AND DILUTION/SETTLING POND	20 SAND CREEK SEWAGE TREATMENT PLANT	32 40 AND 60 MM FIRING RANGE	44 LOAD LINE 11/ARTILLERY PRIMER	○ 20 AREAS OF CONCERN
9 LOAD LINE 2 AND DILUTION/SETTLING POND	21 DEPOT SEWAGE TREATMENT PLANT	33 FIRESTONE TEST FACILITY	45 WET STORAGE AREA	□ IGLOO
10 LOAD LINE 3 AND DILUTION/SETTLING POND	22 GEORGE ROAD SEWAGE TREATMENT PLANT	34 SAND CREEK DISPOSAL ROAD LANDFILL	46 BUILDINGS F-15 AND F-16	□ BUILDING
11 LOAD LINE 4 AND DILUTION/SETTLING POND	23 UNIT TRAINING SITE WASTE OIL TANK	35 1037 BUILDING-LAUNDRY WASTEWATER SUMP	47 BUILDING T-5301 DECONTAMINATION	
12 LOAD LINE 12 AND DILUTION/SETTLING POND	24 RESERVE UNIT MAINTENANCE AREA WASTE OIL TANK	36 PISTOL RANGE	48 ANCHOR TEXT AREA	

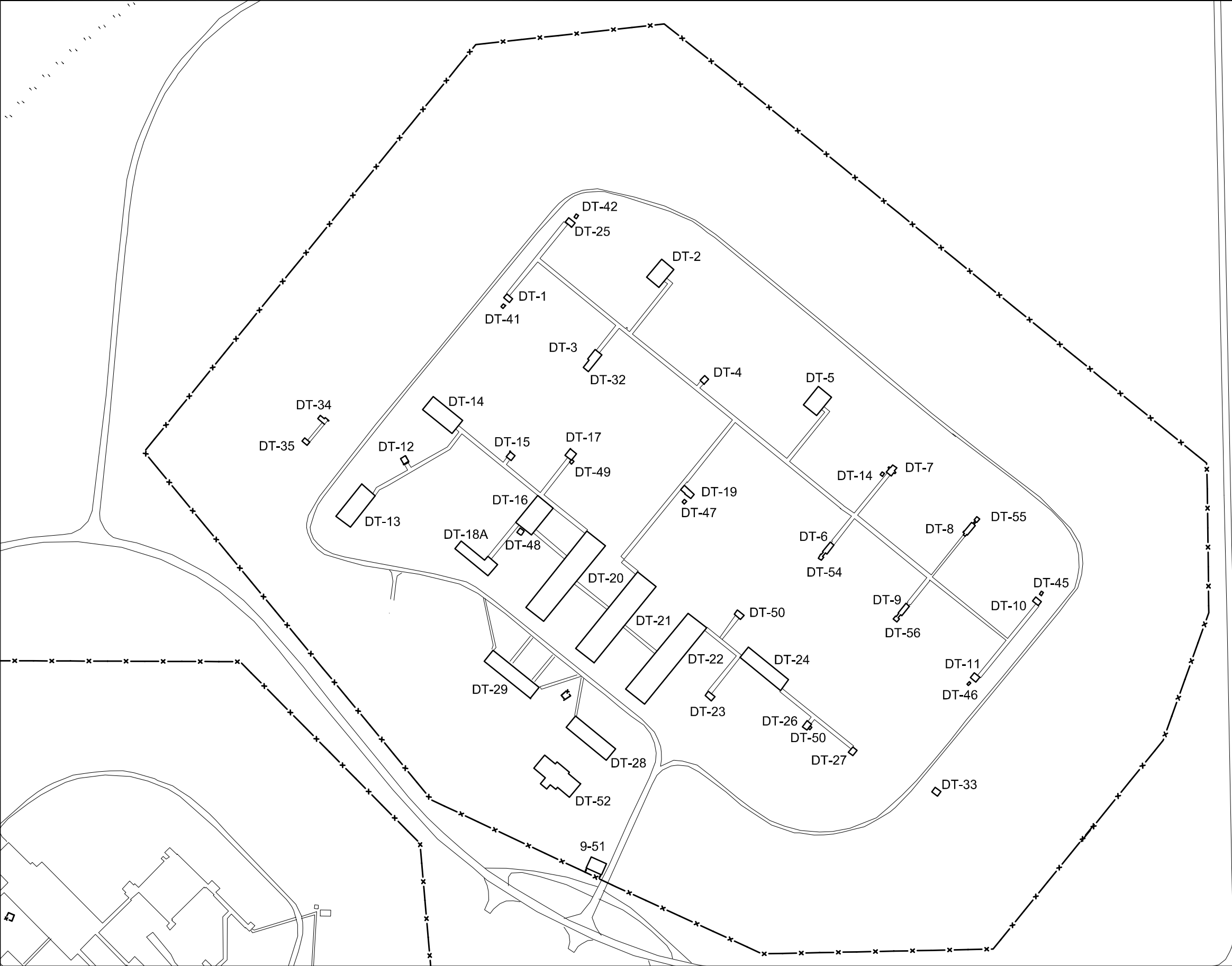
FIGURE 1-2 Revised
FACILITY LOCATION MAP
LOAD LINE 9 - REMEDIAL INVESTIGATION
Ravenna Army Ammunition Plant
Ravenna, OH

Drawn By: ST


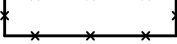


Date Drawn: 10/17/07




MKM Engineers, Inc.
4153 Bluebonnet Road
Stafford, TX 77477

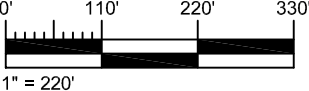



LEGEND:



ROAD
BUILDING
FENCE
DT
DETONATOR LINE







MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO
FIGURE 1-3
LOAD LINE 9 SITE MAP
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: AUG 2007	REV 3
DRAWN BY	KP	APPR. BY	MT



1.2.4 Regulatory Authorities

The approach to addressing environmental conditions at RVAAP is regulatory-based following the frameworks established by the following regulatory drivers: CERCLA, Resource Conservation and Recovery Act (RCRA), Toxic Substances Control Act (TSCA), and applicable Ohio environmental regulations including the State of Ohio EPA Director's Final Findings and Orders (DFFO).

1.3 Report Organization

This RI Report is organized to meet Ohio EPA requirements in accordance with guidance from the USEPA, CERCLA Superfund process, and USACE. The remainder of the report covers the following:

- Section 2.0 – Physical setting of the study area;
- Section 3.0 – Data collection methods, data management, and laboratory programs;
- Section 4.0 – Summarizes LL-9 RI data and the nature and extent of contamination;
- Section 5.0 – Fate and transport of the contaminants of concern;
- Section 6.0 – Baseline human health risk assessment;
- Section 7.0 – Screening ecological risk assessment;
- Section 8.0 – Summary of results and conclusions;
- Section 9.0 – Recommendations; and
- Section 10.0 – References.

The appendices contain RI data and information including boring logs, well construction diagrams, analytical data tables, data quality assessments, a quality assurance summary, risk assessment data tables, and other detailed information used to interpret existing LL-9 conditions.



2.0 RVAAP ENVIRONMENTAL SETTING

This section describes the physical characteristics of the RVAAP facility, including LL-9 (RVAAP-42) and its adjacent environment. These characteristics are factors in interpreting the potential contaminant transport pathways, receptor populations, and exposure scenarios with respect to the evaluation of human health and ecological risks.

2.1 Surface Features

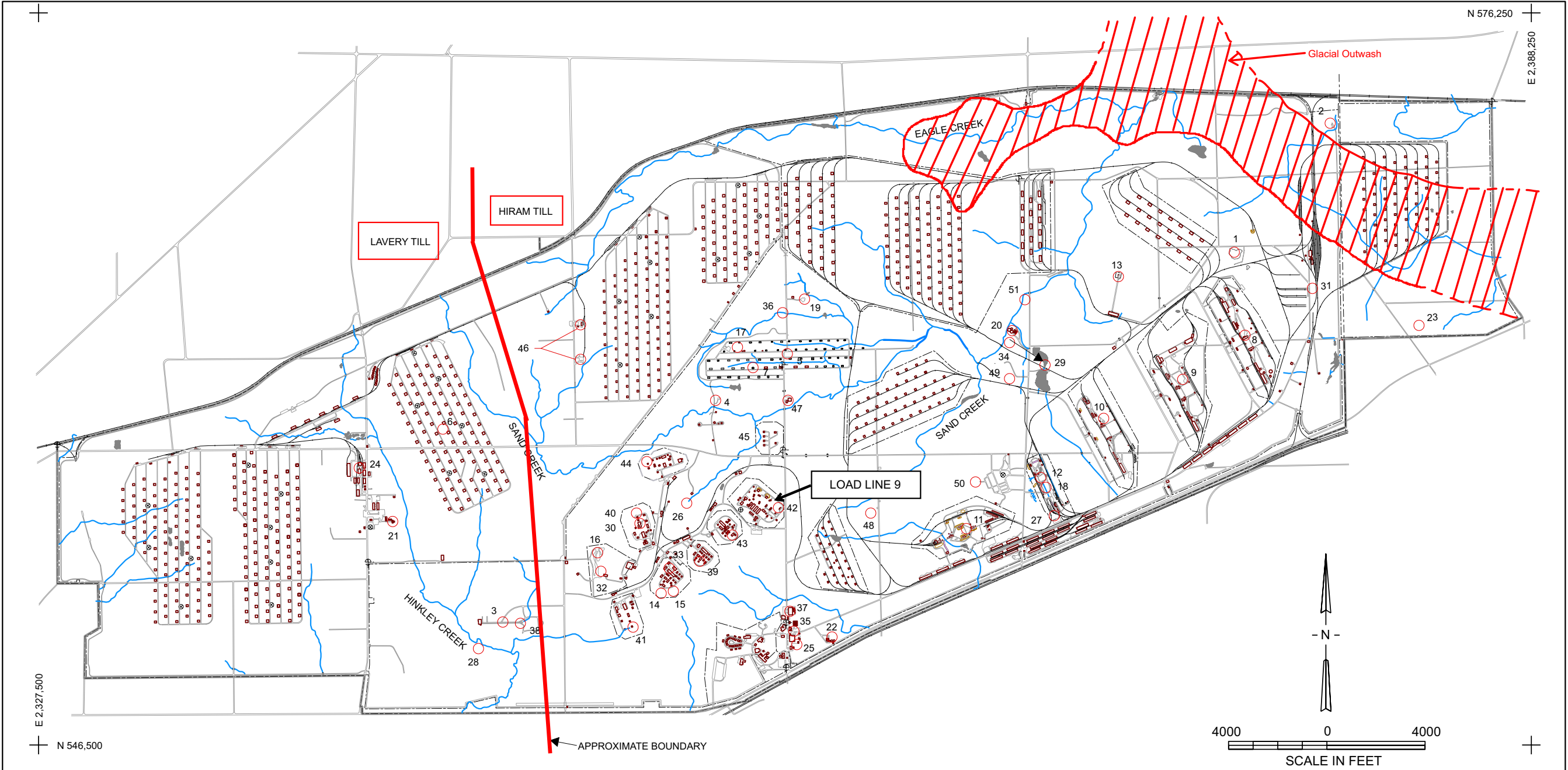
The topography at RVAAP is characterized by gently undulating contours that decrease in elevation from a topographic high in the far western portion of the facility of approximately 1,220 ft above mean sea level (amsl) to lows in far eastern portion of the facility of approximately 930 ft amsl (Figure 2-1). The topography of LL-9 consists of gently sloping (to the north) land where the LL-9 buildings were located. There is a sharp drop in elevation north and east of the LL-9 perimeter road. Elevations range from 1,135 ft in the production area (southwest) to 1,090 ft to the north along the perimeter road.

USACE mapped the installation topography in February 1998 using a 0.61-meter (m) (2 ft) contour interval with an accuracy of 0.204 m (0.67 ft). USACE based the topographic information on aerial photographs taken during spring 1997. The USACE survey is the basis for the topographical information illustrated in figures included in this RI report.

2.2 Meteorology and Climate

The general climate of the RVAAP area is continental, characterized by moderately warm and humid summers, reasonably cold and cloudy winters, and wide variations in precipitation from year to year. The following climatological data were obtained from the National Weather Service Office (NWS, 1995) at the Youngstown-Warren Regional Airport in Trumbull County and are based on a 30-year average.

Total annual rainfall in the RVAAP area is approximately 37.3 inches (in), with the highest monthly average occurring in July (4.07 in) and the lowest monthly average occurring in February (2.03 in). Average annual snowfall totals approximately 56.2 in with the highest monthly average occurring in January (12.9 in). It should be noted that snowfall totals vary widely throughout northeastern Ohio, given the influence of lake-effect snowfall events associated with Lake Erie (located approximately 35 miles to the northwest of RVAAP). The average annual daily temperature in the RVAAP area is 48.3 degrees Fahrenheit (°F), with an average daily high temperature of 57.7 °F and an average daily low temperature of 38.7 °F.




LEGEND OF SITES:

1	RAMSDELL QUARRY LANDFILL	13	BLDG 1200 AND DILUTION/SETTLING POND	25	BLDG 1034 MOTOR POOL WASTE OIL TANK	37	PESTICIDE BUILDING S-44S2
2	ERIE BURNING GROUNDS	14	LOAD LINE 6, EVAPORATION UNIT	26	FUZE BOOSTER AREA SETTLING TANKS	38	NACA TEST AREA
3	DEMOLITIONS AREA #1	15	LOAD LINE 6, TREATMENT PLANT	27	BLDG 854-PCB STORAGE	39	LOAD LINE 5/FUZE LINE 1
4	DEMOLITIONS AREA #2	16	QUARRY LANDFILL/FORMER FUZE & BOOSTER BURNING PITS	28	MUSTARD AGENT BURIAL SITE	40	LOAD LINE 7/BOOSTER LINE 1
5	WINKLEPECK BURNING GROUNDS	17	DEACTIVATION FURNACE	29	UPPER AND LOWER COBBS POND COMPLEX	41	LOAD LINE 8/BOOSTER LINE 1
6	C BLOCK QUARRY	18	LOAD LINE 12 PINK WASTE WATER TREATMENT	30	TREATMENT PLANT LOAD LINE 7 PINK WASTEWATER	42	LOAD LINE 9/DETONATOR LINE
7	BLDG 1601 HAZARDOUS WASTE STORAGE	19	LANDFILL NORTH OF WINKLEPECK BURNING GROUND	31	ORE PILE RETENTION POND	43	LOAD LINE 10/PERCUSSION ELEMENT
8	LOAD LINE 1 AND DILUTION/SETTLING POND	20	SAND CREEK SEWAGE TREATMENT PLANT	32	40 AND 60 MM FIRING RANGE	44	LOAD LINE 11/ARTILLERY PRIMER
9	LOAD LINE 2 AND DILUTION/SETTLING POND	21	DEPOT SEWAGE TREATMENT PLANT	33	LOAD LINE 6/FIRESTONE TEST FACILITY	45	WET STORAGE AREA
10	LOAD LINE 3 AND DILUTION/SETTLING POND	22	GEORGE ROAD SEWAGE TREATMENT PLANT	34	SAND CREEK DISPOSAL ROAD LANDFILL	46	BUILDINGS F-15 AND F-16
11	LOAD LINE 4 AND DILUTION/SETTLING POND	23	UNIT TRAINING SITE WASTE OIL TANK	35	1037 BUILDING-LAUNDRY WASTEWATER SUMP	47	BUILDING T-5301 DECONTAMINATION
12	LOAD LINE 12 AND DILUTION/SETTLING POND	24	RESERVE UNIT MAINTENANCE AREA WASTE OIL TANK	36	PISTOL RANGE	48	ANCHOR TEXT AREA

49	CENTRAL BURN PITS
50	ATLAS SCRAP YARD
51	DUMP ALONG PARIS - WINDHAM ROAD
+++++	RAILROAD TRACKS
-----	FENCELINE
-----	PROPERTY BOUNDARY
-----	STREAM OR CREEK
○ 20	AREAS OF CONCERN
■	IGLOO
□	BUILDING

Figure 2-1
Load Line 9 Remedial Investigation Glacial Till Map
Ravenna Army Ammunition Plant, Ravenna, Ohio

Drawn By: STDate Drawn: 06/17/2005



MKM Engineers, Inc.
4153 Bluebonnet Road
Stafford, TX 77477

Glacial data from - White, George, Glacial Geology of Northeast Ohio. Bulletin 68, 1982



The record high temperature of 100 °F occurred in July 1988, and the record low temperature of -22 °F occurred in January 1994. The prevailing wind direction at RVAAP is from the southwest, with the highest average wind speed occurring in January (11.6 miles per hour [mph]) and the lowest average wind speed occurring in August (7.4 mph). Thunderstorms occur on approximately 35 days per year and are most abundant from April through August. The RVAAP area is susceptible to tornadoes; minor structural damage to several buildings on the facility occurred as the result of a tornado in 1985.

2.3 Surface Water Hydrology

The entire RVAAP facility is situated within the Mahoning River Basin, with the West Branch of the Mahoning River representing the major surface stream in the area. The West Branch generally flows in a north-to-south direction adjacent to the west end of the facility. It eventually flows into the Michael J. Kirwan Reservoir, which is located to the south of State Route 5. The West Branch flows out of the reservoir along the southern facility boundary before joining the Mahoning River east of RVAAP.

The western and northern portions of the RVAAP facility have low hills and a dendritic surface drainage pattern. The eastern and southern portions are characterized by an undulating to moderately level surface, with less dissection of the surface drainage. The facility is marked with marshy areas and flowing and intermittent streams whose headwaters are located in the facility's hills. Three primary water courses drain RVAAP: (1) the South Fork of Eagle Creek, (2) Sand Creek, and (3) Hinckley Creek. Each of these water courses has many associated tributaries. Sand Creek, with a drainage area of 13.9 square mile (sq mi), flows generally in a northeast direction to its confluence with the South Fork of Eagle Creek. In turn, the South Fork of Eagle Creek then continues in a northerly direction for 2.7 miles to its confluence with Eagle Creek. The drainage area of the South Fork of Eagle Creek is 26.2 sq mi, including the area drained by Sand Creek. Hinckley Creek originates just southeast of the intersection between State Routes 88 and 303 to the north of the facility. Hinckley Creek, with a drainage area of 11.0 sq mi, flows in a southerly direction through the facility to its confluence with the West Branch of the Mahoning River south of the facility.

Approximately 50 ponds are scattered throughout the facility. Many were built within natural drainage ways to function as settling ponds or basins for process effluent and runoff. Others are natural in origin, resulting from glacial action or beaver (*Castor canadensis*) activity. Most water bodies at RVAAP support an abundance of aquatic vegetation and contain large fish populations. The ponds within the installation are not used as a water supply source. Stormwater runoff is controlled primarily by natural drainage, except in facility operations areas where an extensive storm sewer network helps to direct runoff to drainage ditches and settling ponds. In addition, the storm sewer system was one of the primary drainage mechanisms for process effluent during the period that production facilities (i.e., LL-9) were in operation.



Surface water follows the LL-9 surface contour gradient and drainage ditch system. The general gradient of the site is to the northeast. Surface runoff flows to and follows constructed drainage ditches, which feed into a natural drainage channel located in the north-central portion of the site. This natural drainage channel flows to an unnamed tributary of the Michael J. Kirwan Reservoir.

2.4 Geology

Lithologic logs from seven soil borings, which were advanced during the RI and completed as monitoring wells, were used to characterize the LL-9 surface and subsurface geology. The soils described during the advancement of the soil borings included medium to fine-grained sand and sandy silt. Grain size analysis from the Shelby tube samples included 19.8% to 40% clay, 38% to 40% silt, and 18% to 30% sand. Bedrock was encountered at a depth ranging from 2.0 to 16.0 ft bgs. The bedrock was cored and included brown, orange, and gray sandstone. Depositional and structural features are detailed in Section 2.4.2. The boring logs, which detail the vertical lithologic sequences, are found in Appendix A.

2.4.1 Glacial Deposits

RVAAP is located within the Glaciated Allegheny Plateau physiographic region of Ohio. Pennsylvanian bedrock is overlain by Wisconsinian Age glacial deposits. Two glacial advances during the Wisconsin Age of the Pleistocene Epoch resulted in the deposition of glacial till over the entire RVAAP installation. The Lavery Till is exposed in the western part of the installation, and the younger Hiram Till is exposed in the remaining portion of RVAAP, including LL-6, which is located in the south-central portion of RVAAP (White, 1982). Glacial outwash deposits are found in the north eastern part of RVAAP in the Eagle creek drainage area and the area described as Erie Burning Grounds (White, 1982).

The first glacial advance deposited the Lavery Till over the facility. The Lavery Till consists mostly of clay and silt with a few cobbles and sporadic boulders. In general the Lavery Till consists of 28% sand and 30% clay. The second glacial advance deposited the Hiram Till over the eastern two-thirds of the facility. The Hiram Till has the highest clay content of tills in northeastern Ohio (White, 1982) (Figure 2-1). Soil associated with the Hiram Till at RVAAP includes the Mahoning silt loam complex. The specific soils developed at LL-9 are detailed in Section 2.5. The typical Hiram Till consists of 12% sand, 41% silt, and 47% illite and chlorite clay minerals, and ranges in depth from 1.5 to 4.6 m (5 to 15 ft) bgs. Unweathered Hiram Till is dark gray and turns a dark brown when exposed to the atmosphere. The Hiram Till overlies thin beds of sandy outwash material in the far northeastern corner of the facility. Soil associated with the Hiram Till at RVAAP typically includes the Mahoning silt loam complex. The thickness of the glacial deposits varies across the installation.



Subsurface lithology at LL-9 consists mostly of thin clay to sand-rich silt tills/soils with interbedded sands scattered throughout. The soil profile at LL-9 has developed over sandstone bedrock high and is relatively thin (2.0 to 16.0 ft range with most in the 2.0 to 6.0 ft range). Based upon the subsurface sampling at the RVAAP, much of the till deposited (if any) was incorporated into the soil profile. Deposits with higher concentrations of sand generally control the elevation of the shallow water table zone (unconsolidated) if enough thickness is present over the bedrock. Cross-sections of the subsurface at LL-9 illustrate the lateral distribution and variation of these thin discontinuous glaciated sediments (Figures 2-2 to 2-6).

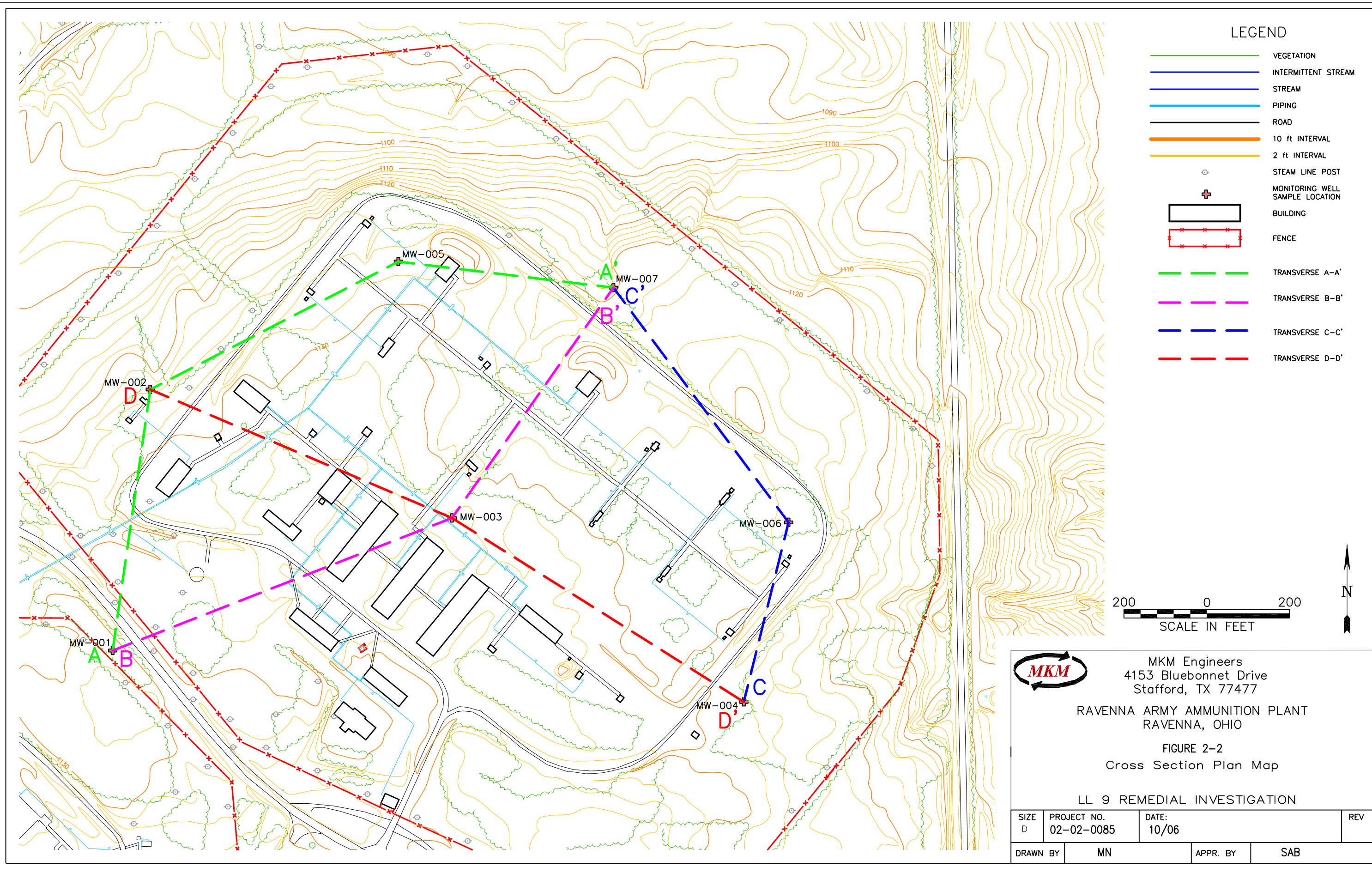
2.4.2 Sedimentary Rocks

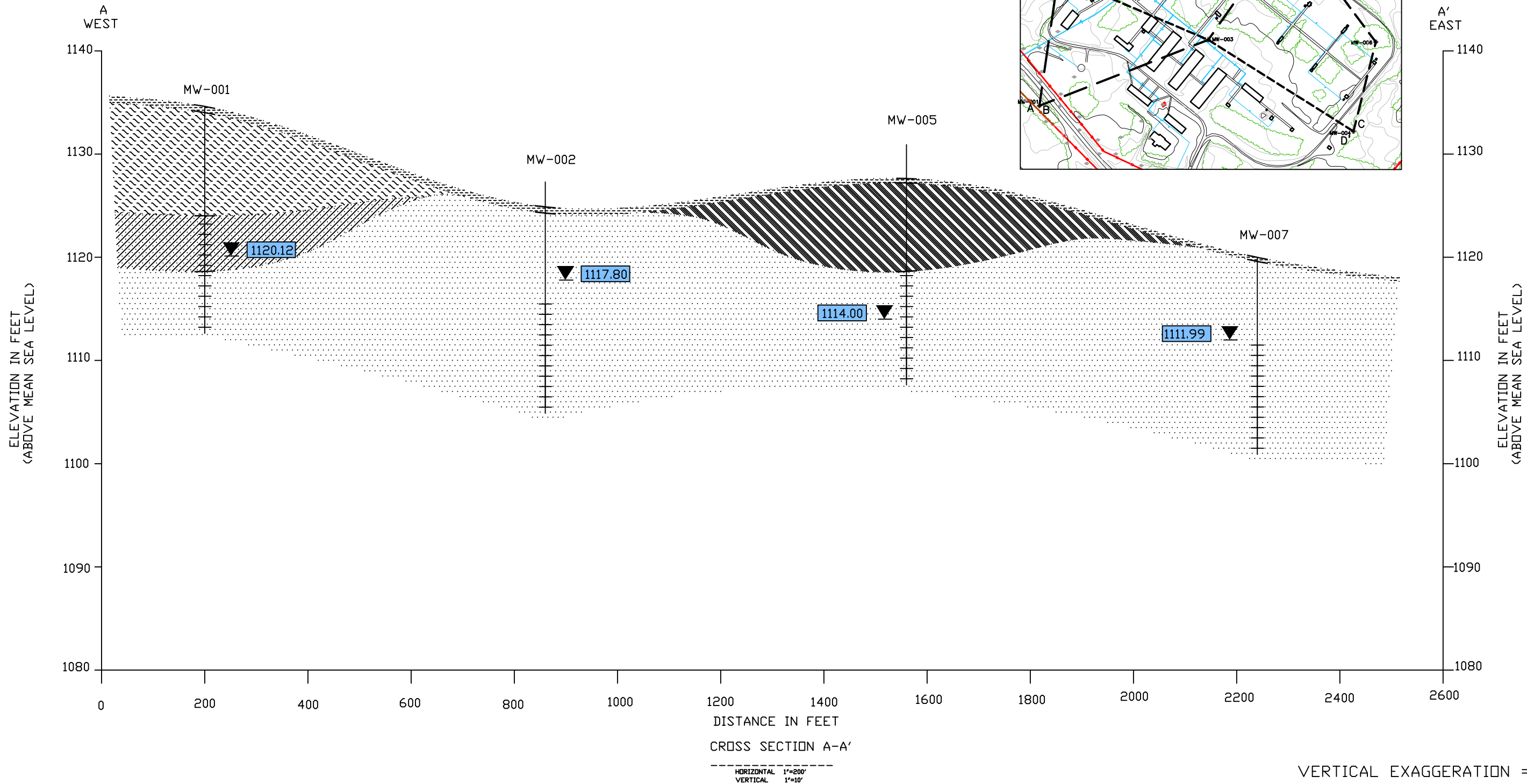
Bedrock outcrops at several areas across RVAAP. According to general geological documents published by the State of Ohio, both Mississippian and Pennsylvanian Age bedrock is believed to underlay RVAAP (USACE, 2001a). The bedrock geology of Portage County is shown on Figure 2-7.

2.4.2.1 RVAAP Sedimentary Rocks

The bedrock geology of RVAAP consists of Carboniferous Age sedimentary rocks that lie stratigraphically beneath the glacial deposits of the Lavery and Hiram Tills. The oldest outcropping of bedrock at the surface within the facility is the Cuyahoga Group of the Mississippian Age. Three members comprise this formation: (1) the Orangeville Shale, (2) the Sharpsville Sandstone, and (3) the Meadville Shale. The Cuyahoga Group outcrops in the far northeastern corner of the facility and generally consists of blue-gray silty shale with interbedded sandstone. The regional dip of the Cuyahoga Group strata is between 1.5 and 3.0 m (5 and 11.5 ft) per mile to the south. The remainder of the facility is underlain by bedrock associated with the Pottsville Formation of Pennsylvanian Age. The Pottsville Formation, which lies unconformably on an erosional surface of the Cuyahoga Group, is divided into four members: (1) the Sharon, (2) the Connoquennessing, (3) the Mercer, and (4) the Homewood Sandstone. The Sharon Member consists of two individual units: the Sharon Conglomerate and the Sharon Shale.

The Sharon Conglomerate is porous, coarse-grained, gray-white sandstone that often exhibits thin layers of milky white quartzite pebbles. The Sharon Conglomerate also has locally occurring thin shale lenses in the upper portion of the unit. Owing to the differences in lithology between the Sharon Conglomerate and the underlying shales of the Cuyahoga Group, the contact between the Pottsville Formation and the Cuyahoga Group usually is quite distinct. The Sharon Shale overlies the Sharon Conglomerate and consists of sandy, gray-black, fissile shale with some plant fragments and thin flagstone beds. Sharon sandstones are exposed on the ground surface at Load Line 1 and the former Ramsdell Quarry.





LEGEND

	TOP SOIL		SANDY SILT		GROUNDWATER ELEVATION (ft) - JUNE 2004		KNOWN SUBSURFACE SOIL CONDITIONS
	SILTY SAND		SILTY CLAY		DEPTH GROUNDWATER ENCOUNTERED		EXPECTED SUBSURFACE SOIL CONDITIONS
	SANDSTONE HOMEWOOD MEMBER		SCREEN INTERVAL				

REVISIONS				
ZONE	REV	DESCRIPTION	DATE	APPROVED
	1		10/15/06	SAB
	2		04/31/07	SL

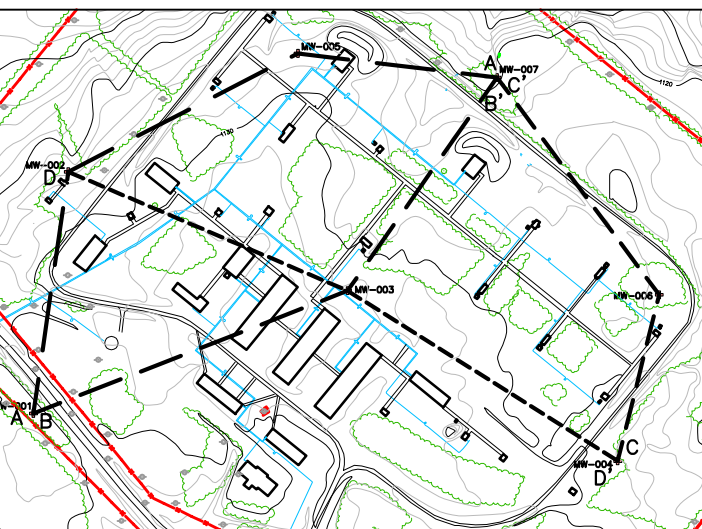
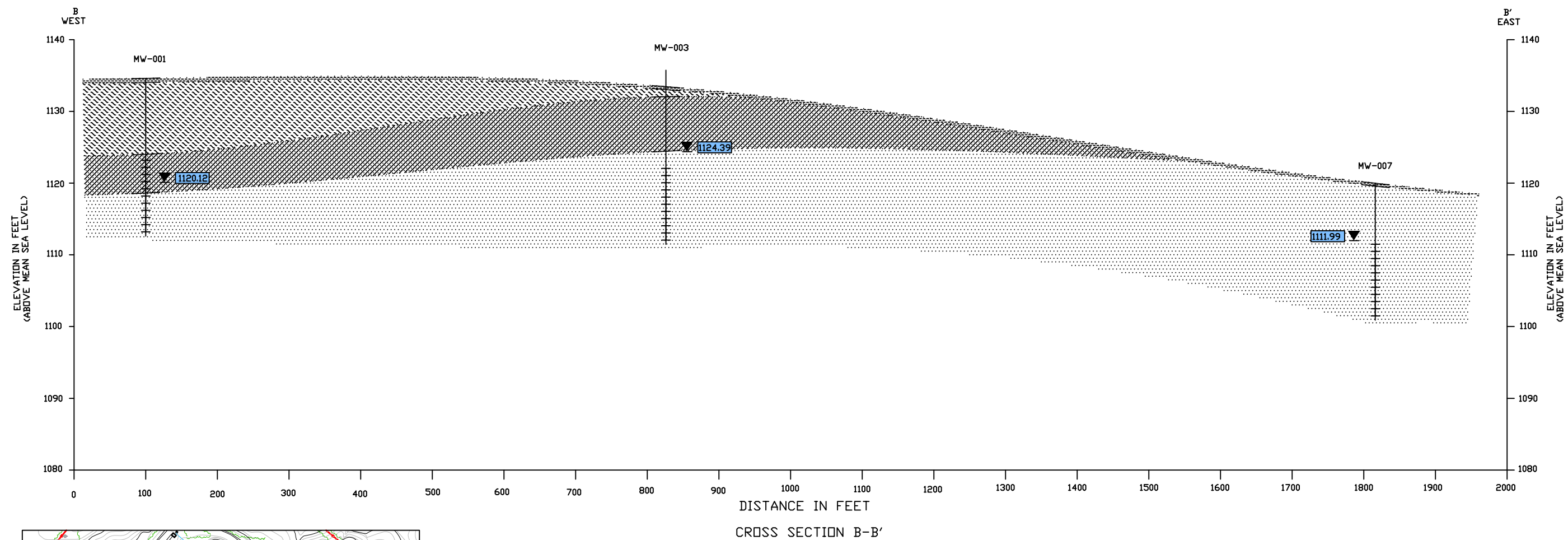


MKM ENGINEERS, INC.

DATE DRAWN 11/18/05

FIGURE 2-3
LOAD LINE 9 REMEDIAL INVESTIGATION
GEOLOGIC CROSS SECTION A-A'
RAVENNA ARMY AMMUNITION PLANT, RAVENNA OHIO

SIZE D	PROJECT NO. 02-02-0085	DWG NO.	REV 2
DRAWN BY	ST/KP	APPR. BY	SRL



VERTICAL EXAGGERATION = 10X

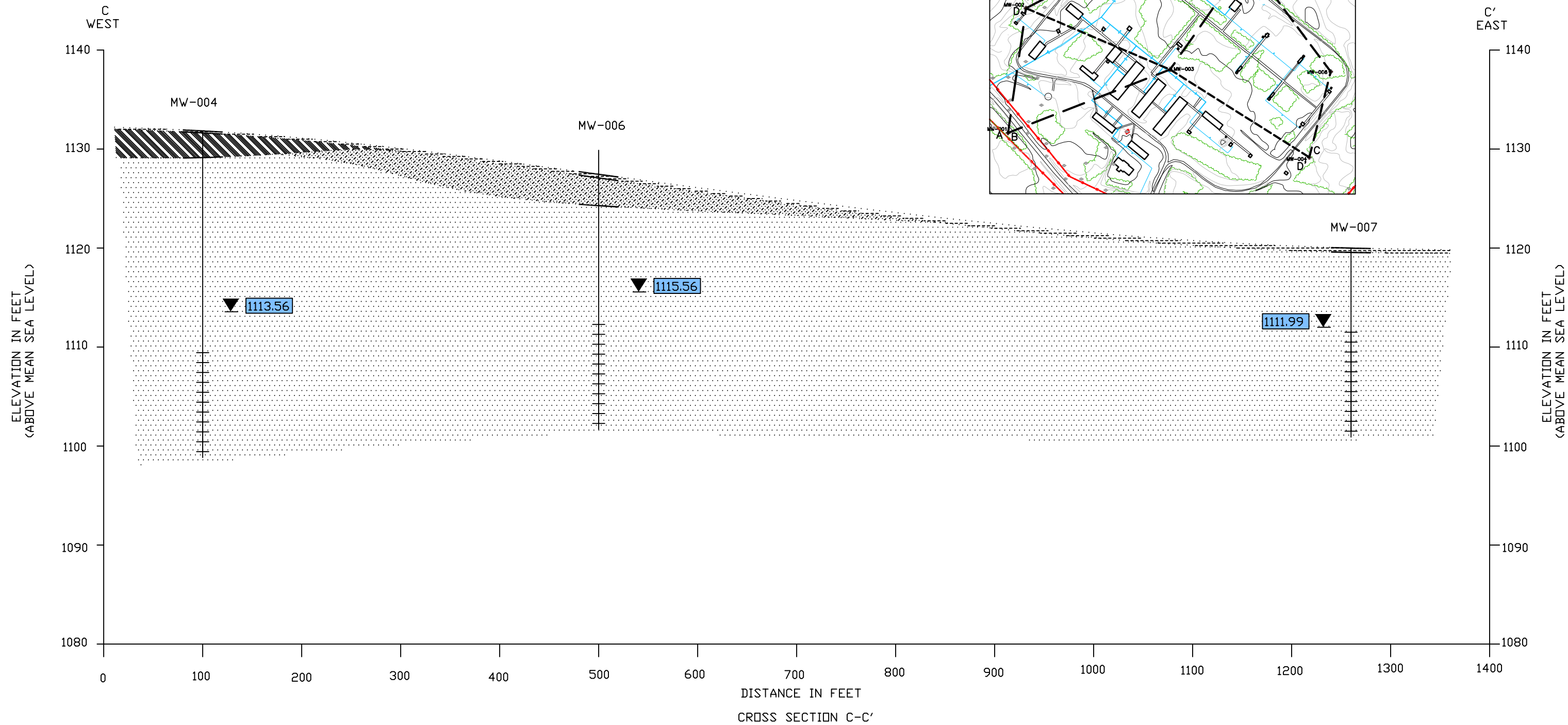
LEGEND

	TOP SOIL		SANDY SILT		GROUNDWATER ELEVATION (ft) - JUNE 2004		KNOWN SUBSURFACE SOIL CONDITIONS
	SILTY SAND		SCREEN INTERVAL		DEPTH GROUNDWATER ENCOUNTERED		EXPECTED SUBSURFACE SOIL CONDITIONS
	SANDSTONE HOMEWOOD MEMBER						

REVISIONS				
ZONE	REV	DESCRIPTION	DATE	APPROVED
	1		10/15/06	SAB
	2		04/31/07	SL

MKM ENGINEERS, INC.	
DATE DRAWN	11/18/05

FIGURE 2-4 LOAD LINE 9 REMEDIAL INVESTIGATION GEOLOGIC CROSS SECTION B-B' RAVENNA ARMY AMMUNITION PLANT, RAVENNA OHIO			
SIZE D	PROJECT NO. 02-02-0085	DWG NO.	REV 2
DRAWN BY	ST	APPR. BY	SRL



HORIZONTAL 1"=100'
VERTICAL 1"=10'

VERTICAL EXAGGERATION = 10X

LEGEND

	TOP SOIL		SILTY CLAY		GROUNDWATER ELEVATION (ft) - JUNE 2004		KNOWN SUBSURFACE SOIL CONDITIONS
	CLAYEY SILT		SCREEN INTERVAL		DEPTH GROUNDWATER ENCOUNTERED		EXPECTED SUBSURFACE SOIL CONDITIONS
	SANDSTONE HOMEWOOD MEMBER						

REVISIONS				
ZONE	REV	DESCRIPTION	DATE	APPROVED
	1		10/15/06	SAB
	2		04/31/07	SL


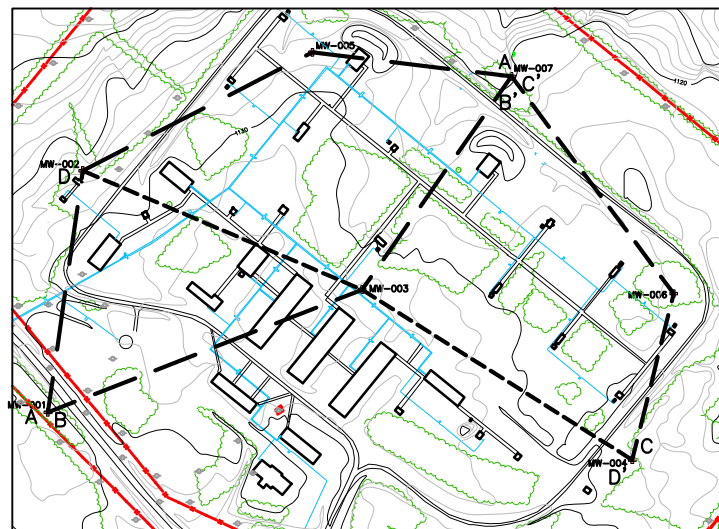
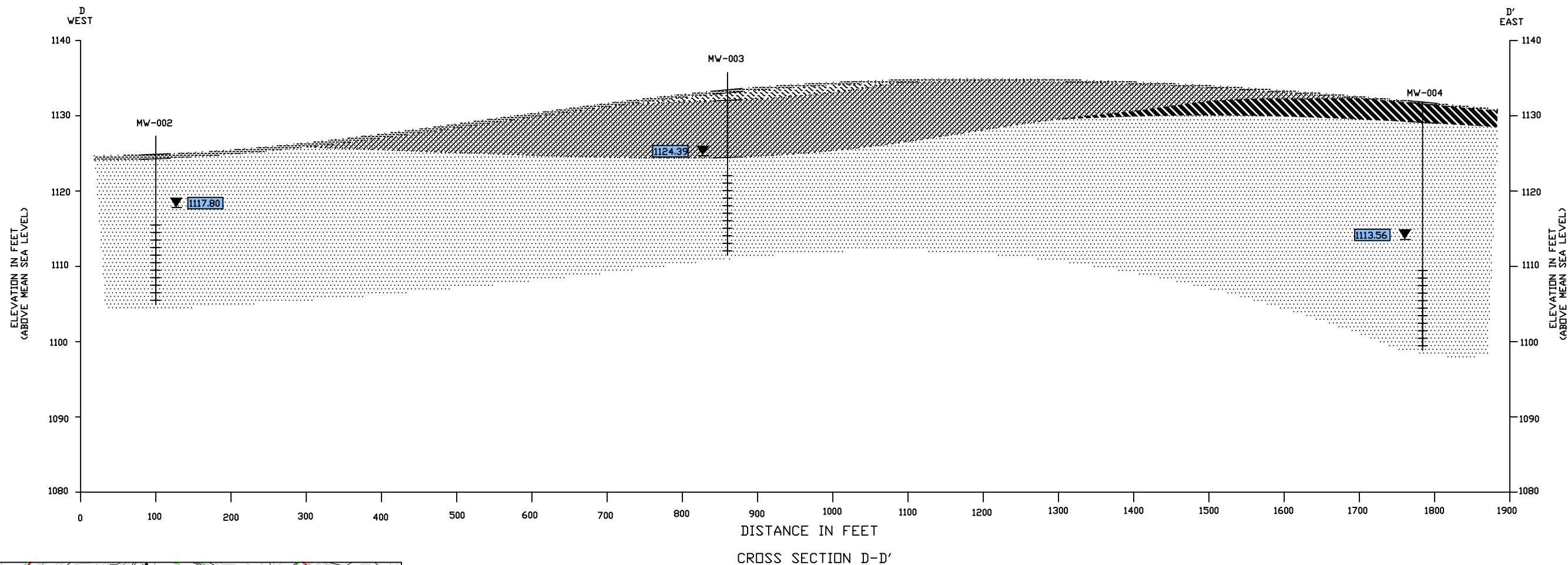

MKM ENGINEERS, INC.
DATE DRAWN 11/17/05

FIGURE 2-5 LOAD LINE 9 REMEDIAL INVESTIGATION GEOLOGIC CROSS SECTION C-C' RAVENNA ARMY AMMUNITION PLANT, RAVENNA OHIO				
SIZE D	PROJECT NO. 02-02-0085	DWG NO.		REV 2
DRAWN BY	ST	APPR. BY	SRL	



VERTICAL EXAGGERATION = 10X

LEGEND

	TOP SOIL		SILTY SILT		SILTY CLAY		KNOWN SUBSURFACE SOIL CONDITIONS
	SILTY SAND		CLAYEY SILT		EXPECTED SUBSURFACE SOIL CONDITIONS		DEPTH GROUNDWATER ENCOUNTERED
	SANDSTONE HOMEWOOD MEMBER		GROUNDWATER ELEVATION (ft) - JUNE 2004		SCREEN INTERVAL		

REVISIONS				
ZONE	REV	DESCRIPTION	DATE	APPROVED
	1		10/16/06	SAB
	2		04/31/07	SL

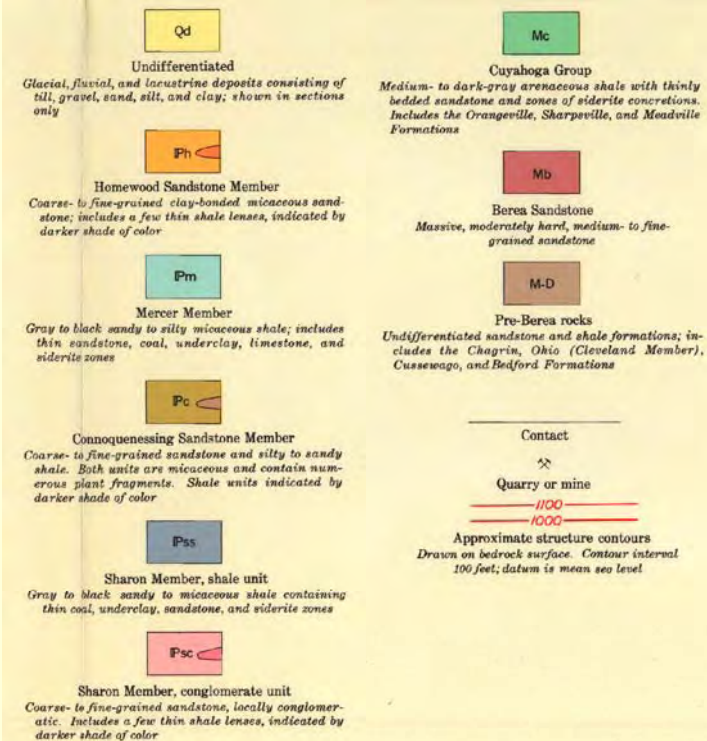
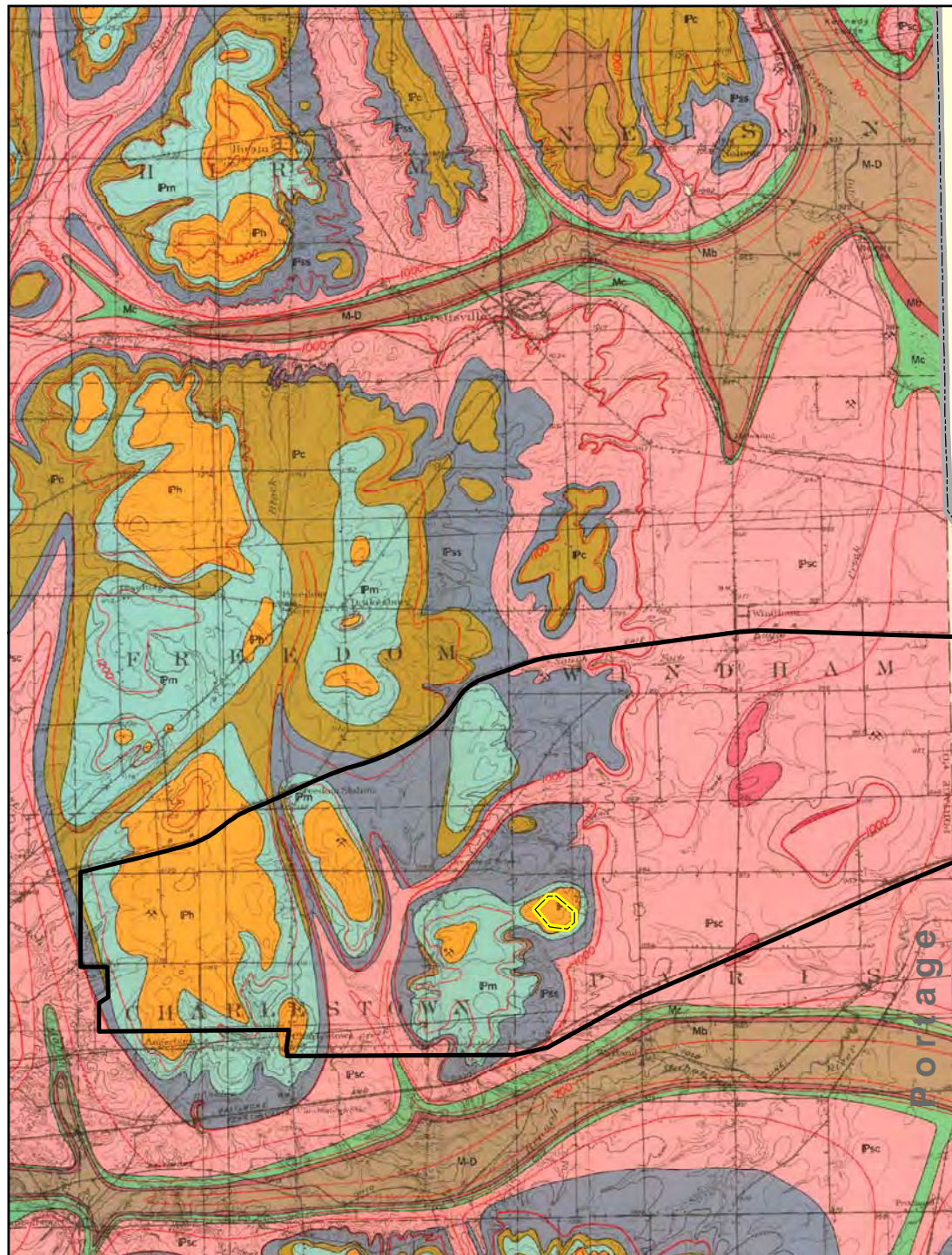


MKM ENGINEERS, INC.

DATE DRAWN 11/17/05

FIGURE 2-6
LOAD LINE 9 REMEDIAL INVESTIGATION
GEOLOGIC CROSS SECTION D-D'
RAVENNA ARMY AMMUNITION PLANT, RAVENNA OHIO

SIZE	PROJECT NO.	DWG NO.	REV
D	02-02-0085		2
DRAWN BY	ST	APPR. BY	SRL



LL-9 Site Boundary
Ravenna Army Ammunition Plant
County Boundary

0 5,000 10,000
Feet

Geology Map Source: Winslow, J.D. and White, G.W., 1966, Geology and ground-water resources of Portage County, Ohio: U.S. Geological Survey, Professional Paper 511, scale 1:62500



MKM Engineers, Inc.
4153 Blubonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

FIGURE 2-7
BEDROCK GEOLOGY MAP

LL-9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: AUG 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	KA



The Connoquenessing Sandstone Member of the Pottsville Formation unconformably overlies the Sharon Member and is a medium to coarse-grained, gray-white sandstone with more feldspar and clay than the Sharon Conglomerate. Thin interbeds and partings of sandy shale also are common in the Connoquenessing. The Mercer Member of the Pottsville Formation overlies the Connoquenessing and consists of silty to carbonaceous shale with abundant thin, discontinuous sandstone lenses in the upper portion. Regionally, the Mercer also has been noted to contain interbeds of coal. The Homewood Member of the Pottsville Formation unconformably overlies the Mercer Member and consists of coarse-grained cross-bedded sandstones that contain discontinuous shale lenses. The Connoquenessing, Mercer, and Homewood Members are present only in the western half of the RVAAP facility. The Sharon Conglomerate unit is the upper bedrock surface in most of the eastern half of the facility. The regional dip of the Pottsville Formation strata is between 1.5 and 3.5 m per 1.6 km (5 and 10 ft per mi.) to the south.

2.4.2.2 LL-9 Sedimentary Bedrock

The Homewood Member of the Pottsville Formation is the upper bedrock surface present beneath the glacial/soils cover at LL 9. At LL-9 weathered bedrock (Homewood Member Sandstone) was encountered in seven locations, at depths from 2 to 16 ft, during the advancement of soil borings for monitoring well construction at LL-9. In addition, weathered bedrock was found in 21 additional locations at depths from 2 to 13 ft during the advancement of soil borings (not for monitoring well construction) at LL-9. The cored bedrock was described as a generally weathered coarse to fine grained well cemented sandstone with some interbedded siltstone. Sandstone colors included brown, orange, and gray. Depositional features included bedding plains, laminations, rip-up clasts, and weathered contacts. Structural features included horizontal and vertical fractures and clay-filled fractures. Structural features included horizontal and vertical fractures and clay-filled fractures.

2.5 Soils

According to the *Soil Survey of Portage County, Ohio* (USDASCS, 1978) and *Soil Survey of Trumbull County, Ohio* (USDASCS, 1982), RVAAP soils, which formed in glacial tills, are described as being nearly level to gently sloping and range from poorly drained to moderately well drained. Three soil types are found at LL-9: Dekalb channery loam, 2–6% slopes and 6–12% slopes; Loudonville silt loam, 2–6% slopes; and Mahoning silt loam, 2–6% slopes. Dekalb channery loam soils formed in residuum weathered from the underlying sandstone bedrock. Loudonville silt loam soils formed from a combination of residuum weathered from the underlying sandstone bedrock and glacial till. Mahoning silt loam soils formed in silty clay loam or clay loam glacial till.



Dekalb channery loam (2–6%) is found in the north and east portion of LL–9. Dekalb channery loam (2–6%) is gently sloping, well drained soil with medium runoff and is usually found on the upper part of hillsides. Dekalb channery loam (6–12%) is found adjacent to the Dekalb channery loam (2–6%) further to the north and east of LL–9. Dekalb channery loam (6–12%) is sloping, well drained soil with medium to rapid runoff and is found in elongated areas on the upper part of hillsides. Loudonville silt loam (2–6%) is found in the central and southern portion of LL–9 and consist of gently sloping soil commonly found on the upper part of hillsides adjacent to Dekalb series soils. Mahoning silt loam (2–6%) is found in the northernmost and westernmost portions of LL–9. Mahoning silt loam is gently sloping, poorly drained soil with medium to rapid runoff, severe seasonal wetness, and slow permeability.

2.6 Hydrogeology

This section describes the unconsolidated sediments, bedrock, and groundwater utilization found at RVAAP and LL–9.

2.6.1 Unconsolidated Sediments

Limited subsurface information indicates that a buried bedrock valley may be present to the west and north of the area known as “Fuze and Booster Hill”. However, the glacial deposits that fill this valley have not been utilized for groundwater production at RVAAP. The water-bearing characteristics for the sand and gravel aquifers in the vicinity of the RVAAP installation are poorly documented. Wells that penetrate these aquifers can yield up to 6,080 liters per minute (LPM) [1,600 gallons per minute (GPM)]. However, yields from wells penetrating silty or clay till materials are significantly lower. In general, the Lavery and Hiram Tills are too thin and impermeable to produce useful quantities of water.

2.6.2 Bedrock

The Sharon Conglomerate and Connoquenessing Sandstone bedrock were the primary sources of potable groundwater at RVAAP during its active phase. Many facility production wells were completed in either the Sharon Conglomerate or the Connoquenessing Sandstone. Some wells were completed so that water was produced from both the Sharon Conglomerate and the Connoquenessing Sandstone, as well as the intervening Sharon Shale (to the extent that the Sharon Shale could produce water). The highest yields were determined to come from the quartzite-pebble conglomerate facies and from jointed and fractured zones. Weathered bedrock (Homewood Member Sandstone of the Pottsville Formation) was encountered in seven locations, at depths from 5 to 16 ft, during the advancement of soil borings for monitoring well construction at LL–9. In addition, weathered bedrock was found in 21 locations at depths from 2 to 13 ft during the advancement of soil borings (not for monitoring well construction) at LL–9.



The most important bedrock sources of groundwater in the vicinity of the RVAAP facility are the sandstone/conglomerate members of the Pottsville Formation. These aquifers, together with two other deeper Mississippian/Devonian sandstone aquifers, represent the most important bedrock sources of groundwater in Northeastern Ohio. The Sharon Conglomerate is the primary source of groundwater at RVAAP and maintains the most significant well yields of the Pottsville Formation Members with hydraulic conductivity values of 19 to 7,600 liters per day per meter (LPD/m) [from 5 to 2,000 gallons per day per foot (GPD/ft)]. Past studies of the Sharon Conglomerate indicate that the highest yields are associated with the true conglomerate phase (coarse-grained sandstone with abundant quartzite pebbles) and with joints and fractures in the bedrock; however, there is no facility-specific information available regarding variations in aquifer properties resulting from these factors. Where present, the overlying Sharon Shale acts as a relatively impermeable confining layer for the Sharon Conglomerate.

Several flowing artesian production wells have been noted at the facility. The Connoquenessing Sandstone and the Homewood Sandstone are the remaining aquifers of the Pottsville Formation and exhibit hydraulic conductivities of from 19 to 1,140 LPD/m (from 5 to 300 GPD/ft) and from 19 to 760 LPD/m (from 5 to 200 GPD/ft), respectively. Well yields in the Connoquenessing and Homewood sandstones, although lower than the Sharon Conglomerate, are high enough to provide significant quantities of water. Several wells at the RVAAP facility have penetrated both the Sharon Conglomerate and the Connoquenessing Sandstone and reportedly produced water from both units. In general, hydraulic conductivities in the shales of the Sharon and Mercer Members of the Pottsville Formation are low and result in insignificant groundwater yields. The primary porosity of the shales is likely secondary, owing to joints and fractures in the bedrock; however, there is no facility-specific information available regarding the occurrence of joints and fractures in these units.

There are no records documenting the use of groundwater from LL-9. However, since the underlying Sharon Conglomerate was extensively pumped as a source of groundwater in the vicinity of LL-9 (Kammer, 1982), consideration should be given to the potential for historical contaminant migration from the AOC via groundwater draw down through the Homewood Sandstone. This phenomenon (downward migration between sandstone units) while possible would only be a migratory possibility if the Homewood Sandstone was impacted from LL-9 operations and sufficient downward permeability was present.

2.6.3 Groundwater Utilization

This section describes the unconsolidated sediments, bedrock, and groundwater utilization found at RVAAP and LL-9. Production wells scattered throughout the facility provided necessary sanitary and process water for RVAAP operations. The remaining process production wells, except for the well at former Building T-5301 (not currently in use), were permanently abandoned in 1992. Currently, only two groundwater production wells remain in



operation. These wells, located in the central portion of the facility, provide sanitary water to the remaining site personnel. Residential groundwater use in the surrounding area is similar to that for RVAAP, with the Sharon Sandstone acting as the major producing aquifer in the area. The Connoquenessing Sandstone and the Homewood Sandstone also provide limited groundwater resources, primarily near the western half of the RVAAP facility. Many of the local residential wells surrounding RVAAP are completed in the unconsolidated glacial material.

2.7 Demography and Land Use

As of February 2006, a total of 20,403 acres of the former 21,683-acre RVAAP have been transferred to the United States Property and Fiscal Officer (USP&FO) for Ohio for use as an OHARNG training site. Currently, RVAAP consists of 1,280 acres in several distinct parcels scattered throughout the confines of the OHARNG's Ravenna Training and Logistics Site (RTLS). RVAAP's remaining parcels of land are located completely within the RTLS, and are completely enclosed by the RTLS perimeter fence.

The RTLS is located in northeastern Ohio within Portage and Trumbull counties, approximately 4.8 kilometers (km) (3 miles) east/northeast of the City of Ravenna and approximately 1.6 km (1 mile) northwest of the City of Newton Falls. The RVAAP portions of the property are completely located within Portage County. The RTLS (inclusive of RVAAP) is a parcel of property approximately 17.7 km (11 miles) long and 5.6 km (3.5 Miles) wide. The facility is bounded by State Route 5, the Michael J. Kirwan Reservoir, and the CSX System Railroad on the south; Garrett, McCormick, and Berry Roads on the west; the Norfolk Southern Corporation railroad on the north; and State Route 534 on the east.

The RTLS is surrounded by several communities: Windham on the north, Garrettsville 9.6 km (6 miles) to the northwest; City of Newton Falls 1.6 km (1 mile) to the southeast; Charlestown to the southwest, and Wayland 4.8 km (3 miles) to the south. RTLS did not exist when the RVAAP was operational, and the entire 21,683-acre parcel was a GOCO industrial facility.

The RVAAP facility is located in a rural area, not close to any major industrial or other developed areas. Based on data from the United States Census Bureau (1992) and the *Portage County Soil and Water Conservation District Resources Inventory* (1985), approximately 55% of Portage County, in which a majority of installation acreage is located, consists of either woodland or farmland acreage. The Michael J. Kirwan Reservoir (also known as the West Branch Reservoir) is the closest major recreational area and is located adjacent to the western half of RVAAP south of State Route 5.

2.8 Ecology

Before the government acquired the property in the 1940s, much of the land at RVAAP was cleared for agricultural use. Over 80% of RVAAP is now forest. The limited field cover growth in the remaining 20% of RVAAP is the result of earlier agricultural practices that left these sites



with poor topsoil that still limits forest growth. Several thousand acres of agricultural fields were planted with trees during the 1950s and 1960s, but these plantings did not grow well in areas with poor topsoil. Some fields, leased for cattle grazing during the same period, were subsequently delayed in their reversion to forest. A few fields have undergone periodic brush removal to maintain them as open fields.

Portions of the RVAAP facility satisfy the regulatory definition of a jurisdictional wetland. Wetland areas at RVAAP include seasonally saturated wetlands, wet fields, and forested wetlands. Some wetland areas are associated with anthropogenic settling ponds and drainage areas. Some of the wetland areas are the result of natural drainage or beaver (*Castor canadensis*) activity. Some wetland areas may have been impacted by former operations due to effluent discharged to settling ponds and the natural drainage of the area in the past. However, many wetlands on the facility are of high quality.

The flora and fauna present at RVAAP are varied and widespread. The following 12 plant species listed as Ohio State Potentially Threatened have been identified at RVAAP:

1. Gray birch (*Betula populifolia*),
2. Butternut (*Juglans cinera*),
3. Northern rose azalea (*Rhododendron nudiflorum* var. *roseum*),
4. Hobblebush (*Viburnum alnifolium*),
5. Long beech fern (*Phegopteris connectilius*),
6. Straw sedge (*Carex straminea*),
7. Water avens (*Geum rivale*),
8. Tall St. John's wort (*Hypericum majus*),
9. Swamp oats (*Sphenopholis pensylvanica*),
10. Shining ladies'-tresses (*Spiranthes lucida*),
11. Arbor vitae (*Thuja occidentalis*), and
12. American chestnut (*Castanea dentata*).

In addition to being a State Potentially Threatened Plant species, butternut also is listed as a Federal Candidate (Category 2) species. A complete list of rare species (plant and animal) found at RVAAP is provided in Appendix B.



The following nine plant and animal species listed as Ohio State Threatened have been identified at RVAAP:

1. Barn owl (*Tyto alba*),
2. Dark-eyed junco (*Junco hyemalis*)(migrant),
3. Hermit thrush (*Catharus guttatus*)(migrant),
4. Least bittern (*Ixobrychus exilis*),
5. Least flycatcher (*Empidonax minimus*),
6. *Psilotreta indecisa* (caddis fly),
7. Simple willow-herb (*Epilobium strictum*),
8. Woodland horsetail (*Equisetum sylvaticum*), and
9. Pale sedge (*Carex pallescens*).

Animals and plants listed as Ohio State Endangered include the sandhill crane (*Grus canadensis*), American bittern (*Botaurus lentiginosus*), northern harrier (*Circus cyaneus*), yellow-bellied sapsucker (*Sphyrapicus varius*), golden-winged warbler (*Vermivora chrysoptera*), osprey (*Pandion haliaetus*), trumpeter swan (*Cygnus buccinator*), mountain brook lamprey (*Ichthyomyzon greeleyi*), graceful underwing (*Catocala gracilis*), ovate spikerush (*Eleocharis ovata*), tufted moisture-loving moss (*Philonotis fontana* var. *caespitosa*), narrow-necked Pohl's moss (*Pohlia elongata* var. *elongata*) and bobcat (*Lynx rufus*). Twenty-one animal species present at RVAAP are also listed as Ohio State Species of Concern:

1. Pygmy shrew (*Sorex hovi*),
2. Star-nosed mole (*Condylura cristata*),
3. Woodland jumping mouse (*Napaeozapus insignis*),
4. Sharp-shinned hawk (*Accipiter striatus*),
5. Marsh wren (*Cistothorus palustris*),
6. Henslow's sparrow (*Ammodramus henslowii*),
7. Cerulean warbler (*Dendroica cerulean*),
8. Prothonotary warbler (*Protonotaria citrea*),



9. Bobolink (*Dolichonyx oryzivorus*),
10. Northern bobwhite (*Colinus virginianus*),
11. Common moorhen (*Gallinula chloropus*),
12. Great egret (*Casmerodius albus*),
13. Sora (*Porzana Carolina*),
14. Virginia rail (*Rallus limicola*),
15. Creek heelsplitter (*Lasmigona compressa*),
16. Sedge wren (*Cistothorus platensis*)
17. Eastern box turtle (*Terrapene carolina*),
18. Four-toed salamander (*Hemidactylium scutatum*),
19. Mayfly (*Stenonema ithica*),
20. Moth (*Apamea mixta*), and
21. Moth (*Brachylomia algens*).

The OHARNG currently manages the Integrated Natural Resource Management Plan for the majority of the facility except for the property that is still maintained by the U.S. Army (as referenced in Section 1.2.1). Restricted land use and forest management practices have preserved and enabled large forest tracts to mature. Habitat conversion at RVAAP has focused on restoration of the forests that covered the area before it was cleared for agriculture. The reversion of these agricultural fields to mature forest provides a diversity of habitats from old field through several successional stages. Overall, the increasing forest cover enhances the area for use by both plant and animal forest species.

Future IRP activities will require consideration of these species to ensure detrimental effects on Ohio threatened or endangered flora and fauna at RVAAP do not occur. This will be discussed in the Ecological Risk Assessment presented in Section 7.0. There are currently no Federally listed species or critical habitat on the RTLS property. There are a few species currently under federal observation for listing but none listed.



3.0 LOAD LINE 9 INVESTIGATIONS

This section describes field activities, sample collection, and sample analyses conducted during the investigation of LL-9. The field and analytical programs were performed in accordance with the RVAAP FWSAP (USACE, 2001a) and the FWSAP Addendum for the Remedial Investigation at LL 9 (MKM, 2003a). Investigation objectives, rationale for sampling locations, sampling methods, and sample locations are discussed in this section. Deviations to the FWSAP (USACE, 2001a) or FWSAP Addendum for LL-9 (MKM, 2003a) are also documented in this section.

3.1 Field Activities

An initial screening field effort for lead azide was conducted in March 2002. RI-related field activities were conducted from October to December 2003 and included:

- Conducting a munitions and explosives of concern (MEC) avoidance screen before field activities were initiated;
- Mowing/clearing sampling areas and access routes;
- Excavation of test pits;
- Establishing work zones;
- Establishing temporary decontamination areas;
- Collecting surface soil samples;
- Collecting subsurface soil samples;
- Collecting sediment samples;
- Collecting surface water samples;
- Installing and developing monitoring wells;
- Collecting geotechnical samples from the borings (Shelby tubes);
- Collecting groundwater samples from monitoring wells;
- Conducting in-situ permeability testing (slug tests);
- Collecting water level measurements;
- Conducting a sample location and monitoring well survey; and
- Packing and shipping samples.



The following sections summarize the LL-9 RI activities including pre-mobilization, mobilization, and project-related field activities. A photographic log of the characterization activities is provided in Appendix C.

3.1.1 MEC Avoidance

MEC activities were conducted as specified in the MEC Avoidance Plan found in the FWSAP Addendum for Load Line 9 (MKM, 2003a). The UXO Avoidance Report is provided in Appendix D. A MEC survey was conducted to ensure worker safety during the RI investigation before initiating field activities. An unexploded ordnance (UXO)-qualified technician surveyed areas where personnel or equipment might traverse. Anomalies were noted, and safe pathways were established. The surface of the point to be penetrated was cleared, and an anomaly-free area was established before intrusive activities were initiated. Down-hole surveys were conducted at 2 ft intervals to a depth of 4 ft, where MEC avoidance was terminated.

3.1.2 Mowing/Clearing of Sample Locations

Ground-level vegetation was mowed using a hydro axe, brush hog, and/or hand clearing with chainsaw, machete, and weed eater. The vegetation was cut or cleared to allow personnel and equipment to access designated sampling locations.

3.1.3 Test Pits

Two test pits were excavated before installation of the monitoring wells to investigate the depth to bedrock and soil profile. The test pits were completed in accordance with Section 4.4.2.4.2 of the FWSAP (USACE, 2001a) as outlined below. The test pit excavations provided information for drilling and installation of monitoring wells; however, no samples were collected during the trenching operations.

3.1.4 Work Zones

MKM established work zones during drilling and trenching operations. The work zone (exclusion zone) was delineated by yellow DO NOT ENTER caution tape and road cones. To ensure site control during sample collection, the access gate to the AOC was closed and padlocked. A site sign-in log was maintained at the access gate for field personnel and site visitors.

3.1.5 Temporary Decontamination Area

A temporary field decontamination area was constructed to facilitate decontamination of the drilling rig, augers, rods and other associated equipment and personnel. The field



decontamination area was located outside the main gate of LL 6 in the parking area (the RI field program at LL 6 (located close to LL-9) was performed concurrently with the LL-9 RI field program). A lined decontamination pad was constructed to capture decontamination fluids. In addition, several investigation-derived waste (IDW) water and clean water above-ground storage tanks (ASTs) were staged at this location. Sampling and drilling equipment was decontaminated in accordance with the procedures outlined in Sections 4.4.2.8 and 4.3.8 of the FWSAP (USACE, 2001a).

3.1.6 Soil Samples

3.1.6.1 Azide Screening Soil Samples

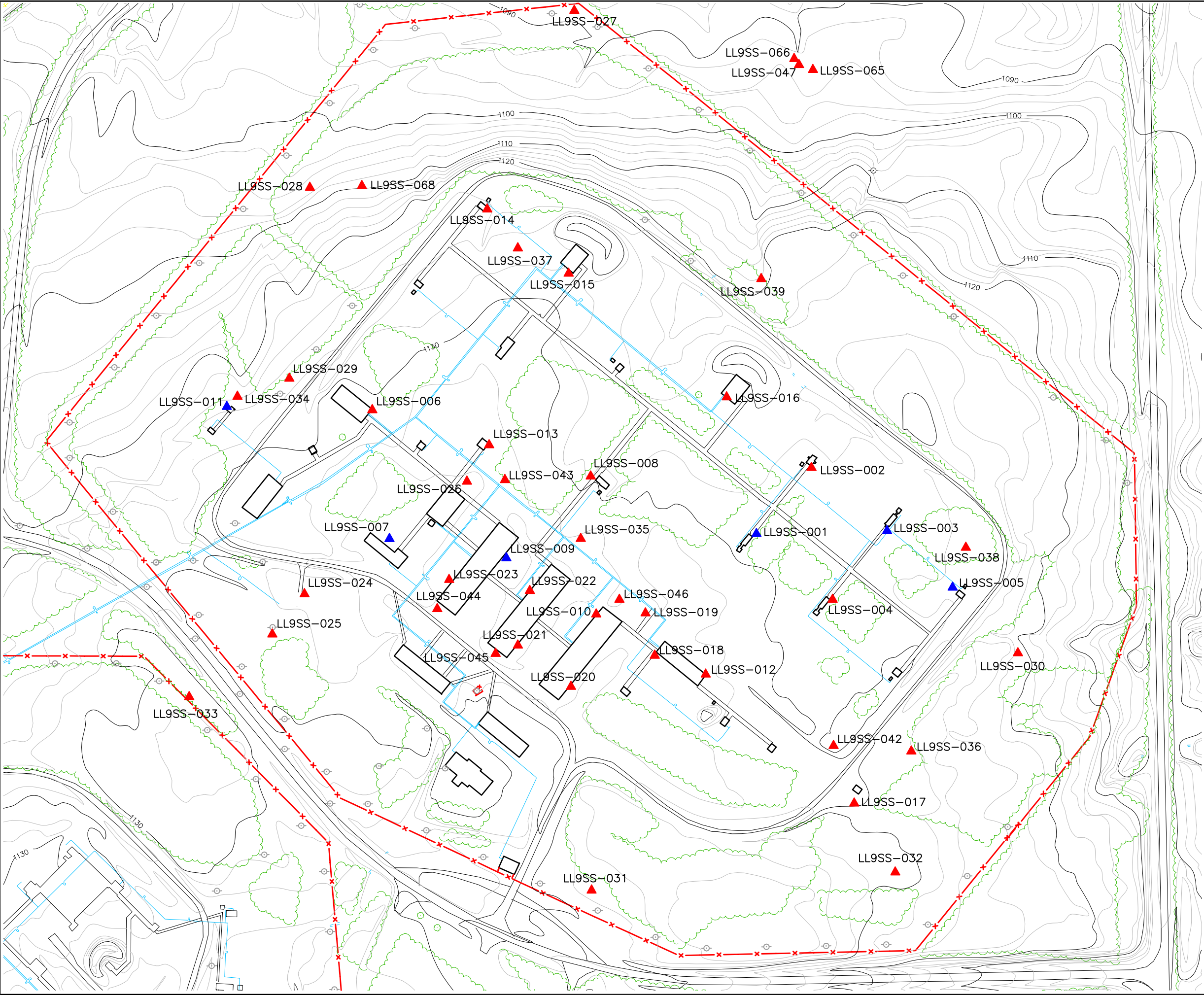
Eleven surface soil samples (LL9SS-001-0001-SO through LL9SS-011-0001-SO) were collected at LL-9 during the March 2002 azide screening event. The 11 soil samples were field analyzed for lead azide, and each was field tested for TNT and RDX using the Jenkins method. Additionally, 6 of the soil samples (LL9SS-001-0001-S), LL9SS-003-0001-SO, LL9SS-005-0001-SO, LL9SS-007-0001-SO, LL9SS-009-0001-SO, and LL9SS-011-0001-SO) were submitted to Severn Trent Laboratories, Inc. (STL) for explosives and metals analysis and split samples were sent to General Physics Laboratories (GPL) to evaluate the accuracy of the new GPL screening method for explosives (modified 8330). In addition, one additional split sample was collected for USACE and submitted for explosives analysis.

Soil samples were collected using a bucket hand auger (stainless steel) during the azide screening. Soil samples were collected from the shallow 0 to 1 ft interval. The surface soil samples were collected in and around bulk handling and process buildings and from locations within the load line. Figure 3-1 shows the location of the surface soil samples. Surface soil samples collected during the azide screening were documented on the field sampling forms included in Appendix E. Results of the surface soil screening and samples analyzed by GPL and STL laboratories are included in Appendix F.

3.1.6.2 RI Soil Samples

Soil sampling during the LL-9 RI was conducted using a combination of hollow-stem auger/split-spoon, direct push (Geoprobe), and hand auger methods. Soil samples were collected from surface (0 to 1 ft interval) and at-depth (1 to 3 ft interval or deeper) soils. The types of soil samples collected, the sampling intervals (if applicable), and the rationale for the depth of sample collection are listed in Table 3-1.

After a soil sample was collected, it was immediately labeled and placed into a plastic bag inside a cooler containing ice. Each cooler shipped to the laboratory was accompanied by a completed chain-of-custody form.



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- SURFACE SOIL (0-1 Ft) SAMPLE LOCATION
- SURFACE SOIL (0-1 Ft) SAMPLING LOCATION TAKEN DURING AZIDE SCREENING
- BUILDING
- FENCE

0' 100' 200' 300'

1" = 200'

N



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO
FIGURE 3-1
SURFACE SOIL SAMPLE
LOCATION MAP
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY R. BOWMAN		APPR. BY	SL



Table 3-1

Rationale for Selection of Soil Sampling Intervals

Sample Type	Sampling Intervals	Rationale for Actual Depth
Surface Soil	0 to 1 ft	Predetermined depths specified in the work plan
Subsurface Soil	1 to 3 ft	Predetermined depths specified in the work plan
Monitoring Well Boreholes	1 st interval: 0 to 1 ft 2 nd interval: None	Visual observation and/or headspace reading No sample collected from rock cores
Subfloor	NA	Floor/soil interface in five LL-9 buildings
Sump/Dry Well	1 st interval: 0 to 1 ft 2 nd interval: Varied	Visual observations Depth of the bottom of the sump/dry well
Sewer	1 st interval: 0 to 1 ft 2 nd interval: Varied	Visual observations Depth of the bottom of the sewer manholes
VOC Screening (Grid)	Varied	Highest headspace reading or strongest VOC odors

3.1.6.3 Surface Soils

Surface soil samples were collected at predetermined depths specified in the work plan to:

- Assess the potential impact of LL-9 activities on surrounding soil,
- Evaluate condition of soil outside the production area,
- Determine the nature and extent of contamination, and
- Determine whether chemical releases had occurred from LL-9 infrastructure.

Figure 3-1 shows the location of the surface soil samples. Forty surface soil samples were collected by hand auger within the LL-9 AOC. These samples included 27 surface soil samples, two adjacent sewers, four adjacent to dry wells, and seven monitoring well locations (LL9SS-033-0001-SO through LL9SS-039-0001-SO). In addition, seven split samples were collected for USACE and submitted for analysis.

Table 3-2 lists the sample numbers and the method used to collect each sample. The soil/floor interface samples were collected after the buildings were demolished. The samples were collected using a hand auger from 0 to 1 ft bgs. One discrete soil sample was collected from each location using a clean, decontaminated, stainless steel hand auger.



**Table 3-2
Summary of Samples and Analysis**

Sample Type	Collection Method	Sample Numbers ¹	Sample Depth	Activity	Date of Field Work	Laboratory Parameters
Surface Soil (40)						
Surface Soil	Hand Auger	Azide Field Screening SS001 - SS010 Lab Analysis SS001, SS003, SS005, SS007, and SS009	0-1 ft	Azide Screening	March 2002	Azide Field Screening, TAL Metals and Explosives (STL) and 10 % Explosives Screening Modified Method 8330 (USACE Comparative Study)
Surface Soil	Hand Auger	SS002, SS004, SS006, SS008, SS010 SS012 - SS032	0-1 ft	R.I	Nov - Dec 2003	TAL Metals and 10% Full Suite
Monitoring Well	Hand Auger	SS033-SS039	0-1 ft	R.I	Nov - Dec 2003	TAL Metals and Explosives with 10% Full Suite Analysis
Sewer	Geoprobe	SS042 and SS043	0-1 ft	R.I	Nov - Dec 2003	TAL Metals
Sump / Dry Wells	Hand Auger	SS044, SS045, SS046, and SS047	0-1 ft	R.I	Nov - Dec 2003	TAL Metals
Sub-floor	Hand Auger	SS011	1-3 ft	Azide Screening	March 2002	Explosives and TAL Metals
Subsurface Soil						
Subsurface Soil	Hand Auger	SB001-SB010, SB012-SB032	1-3 ft	R.I	Nov - Dec 2003	TAL Metals and 10% Full Suite
Monitoring Well	Drill Rig / Split Spoon	Lithologic Samples only	NA			
Sewer	Geoprobe	SB042-SB043	8-11 ft	R.I	Nov - Dec 2003	TAL Metals (SB042) and Full Suite (SB043)
Sump	Hand Auger	SB040-SB041	~ 7 ft	R.I	Nov - Dec 2003	TAL Metals, Explosives, VOCs, SVOCs, PCBs, Pesticides, and Cyanide
Dry Wells		SB044-SB047	> 3 ft	R.I	Nov - Dec 2003	TAL Metals
Sub-floor	Hand Auger	SB048 - SB053	1-3 ft	R.I	Nov - Dec 2003	TAL Metals and 10% Full Suite
VOC Grid	Geoprobe	SB055, SS056, SS061, SS065, SS066	> 1 ft	R.I	Nov - Dec 2003	VOCs, SVOCs, TPH (GRO & DRO)
Groundwater						
Groundwater	Bailer / Low Flow Micropurge	MW001-MW007	Varied	R.I	Nov - Dec 2003	TAL Metals, Explosives, Propellants, VOCs, SVOCs, PCBs, Pesticides and Cyanide
Surface Water						
Surface Water	Submersion	SW001, SW005, SW007 and SW008	Surface	Azide Screening	March 2002	TAL Metals and Explosives
Surface Water	Submersion	SW002, SW003, SW004, and SW012	Surface	R.I	Nov - Dec 2003	TAL Metals and 10% (1 sample - SW012) Full Suite
Sump / Sewer Water	Scoop	SW009 and SW010	> 1 ft	R.I	Nov - Dec 2003	TAL Metals
Sediment						
Sediment (6 Ditch and 2 Sump samples)	Hand Core	SD001 - SD008	> 1 ft	Azide Screening	March 2002	TAL Metals and Explosives
Ditch Sediment	Scoop	SD002, SD004, SD006 and SD011 - SD017	0-0.5 ft	R.I	Nov - Dec 2003	TAL Metals and 10% Full Suite
Sump / Sewer Sediment	Scoop	SD009 and SD010	> 1 ft	R.I	Nov - Dec 2003	TAL Metals and 10% Full Suite
Contingency						
Surface Soil	Hand Auger	SS068	0-1 ft	R.I	Nov - Dec 2003	TAL Metals, Explosives, Propellants, VOCs, SVOCs, PCBs, Pesticides and Cyanide
Subsurface Soil (Next to Sumps)	Hand Auger	SB069, SS070	~ 7 ft	R.I	Nov - Dec 2003	TAL Metals

1 - Full Suite = TAL Metals, Explosives, Propellants, VOCs, SVOCs, PCBs, Pesticides and Cyanide

2 - Sample nomenclature was shortened for inclusion in this table. The full sample numbers are used throughout the remainder of this document



Surface soil sample information was documented on the field sampling forms included in Appendix G. Surface soil sampling results are included in Appendix F.

3.1.6.4 Subsurface Soil Samples

Subsurface soil samples were collected at 53 locations, as shown on Figure 3-2. These samples included 31 subsurface soil sampling locations; 6 subfloor locations (see subsection 3.1.10.3); 2 locations at depths equivalent to the bottom of the sewer manholes; 8 locations at depths equivalent to the bottom of the sump/dry wells; and at 6 locations in the VOC screening grid established east of the solvent building at LL-9 (DT-33). Soil samples were collected from predetermined depths to:

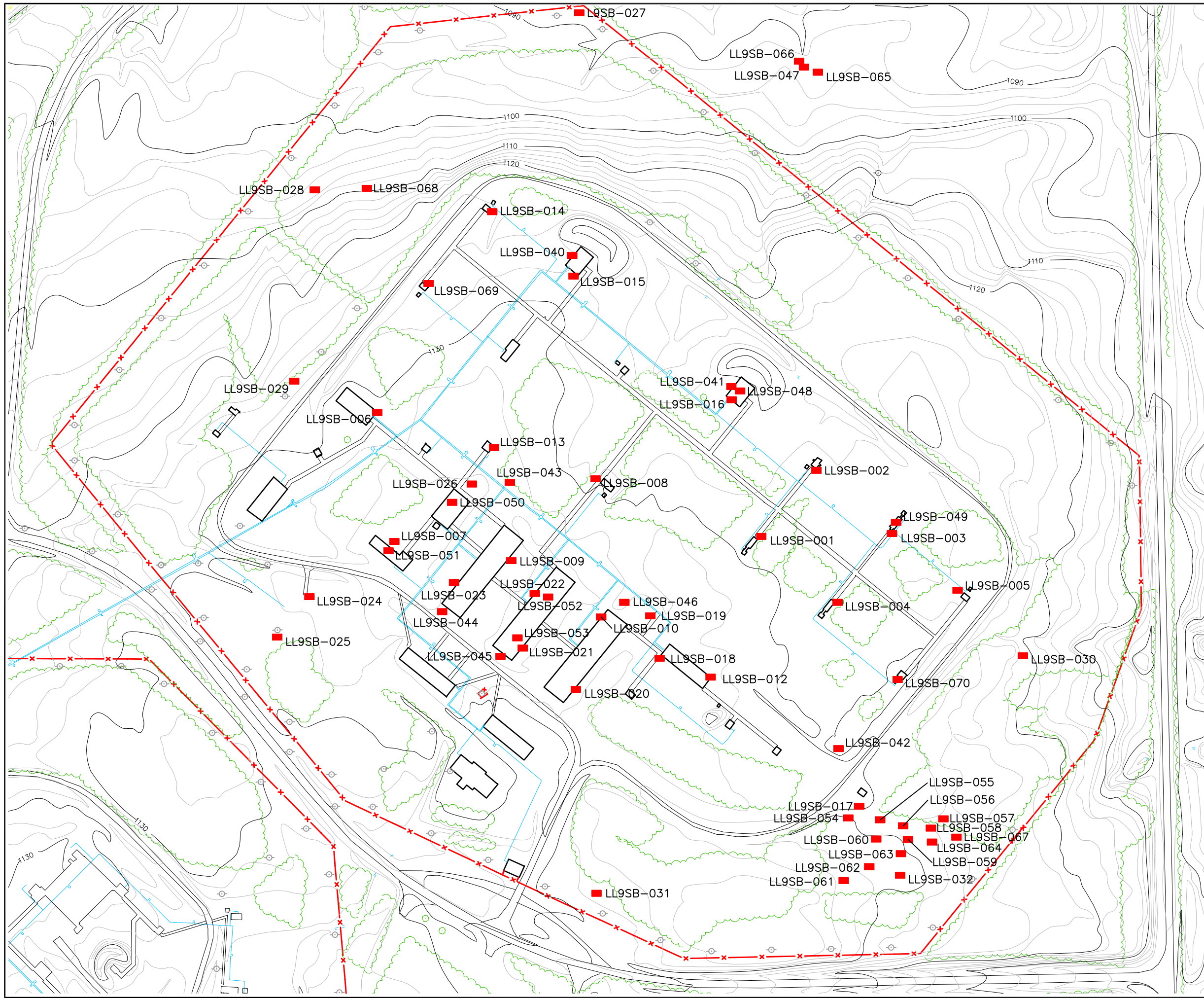
- Assess the potential impact of LL-9 activities on the surrounding soil,
- Determine the nature and horizontal and vertical extent of potential contamination,
- Evaluate the geotechnical characteristics of the LL-9 soils, and
- Determine whether releases had occurred from LL-9 infrastructure.

Subsurface Soil Samples Adjacent to Sewers

Two samples (LL9SB-042-0001-SO and LL9SB-043-0001-SO) were collected at LL-9 sewer manholes. The samples were located at manholes and stress points where sewer integrity was potentially suspect (see Figure 3-2). From each boring, two discrete samples were collected from the 0 to 1 ft bgs interval (surface soil sample), and at a second interval starting at the elevation equal to the total manhole depth and extending to 2 ft below the total depth (subsurface soil sample). The samples were field tested for VOCs using a hand-held photoionization detector (PID) before laboratory analysis.

Subsurface Soil Samples Adjacent to Sumps

Two discrete soil samples (LL9SB-040-0001-SO and LL9SB-041-0001-SO) were collected from each boring at locations adjacent to the LL-9 sumps. The sumps were excavated during the demolition operations conducted before the RI activities began. The samples (LL9SB-040-0001-SO and LL9SB-041-0001-SO) were collected from approximately 7 ft depth, the bottom elevation of the sumps (LL9SB-040-0001-SO [6.5 ft], LL9SB-041-0001-SO [7.0 ft]). Visual observations were used to determine the final depth.



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- SUBSURFACE SOIL SAMPLING LOCATION
- BUILDING
- FENCE

0' 100' 200' 300'

1" = 200'

N



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

**FIGURE 3-2
SUBSURFACE SOIL SAMPLE
LOCATION MAP**

LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY R. BOWMAN		APPR. BY	SL



1 Direct-Push Subsurface Soil Samples (VOC Screening Grid)

2 Of the 14 direct-push soil borings (LL9SB-054-0001-SO through LL9SB-067-0001-SO), six
3 subsurface soil samples (LL9SB-055-0001-SO, LL9SB-056-0001-SO, LL9SB-059-0001-
4 SO, LL9SB-061-0001-SO, LL9SB-064-0001-SO, and LL9SB-067-0001-SO) were collected
5 at a VOC screening grid established east of the solvent building at LL-9 (Bldg DT-33). The
6 VOC screening grid was established and sampled to determine whether releases from the
7 solvent building had occurred. Samples from the soil borings exhibiting the highest headspace
8 readings were selected for analysis.

9 Table 3-2 lists the sample numbers and the method used to collect each sample. In addition,
10 three split samples (LL9SB-005-0001-DUP, LL9SB-043-0001-DUP, and LL9SB-053-0001-
11 DUP) were collected for the USACE and submitted for analysis. After collection, the samples
12 were placed into a cooler containing ice and submitted to the laboratory with completed chain-
13 of-custody forms. The soil boring logs for the direct-push soil samples are included in Appendix
14 A. The subsurface soil samples were documented on the field sampling forms included in
15 Appendix H and the analytical results are included in Appendix I.

16 3.1.7 Groundwater Activities

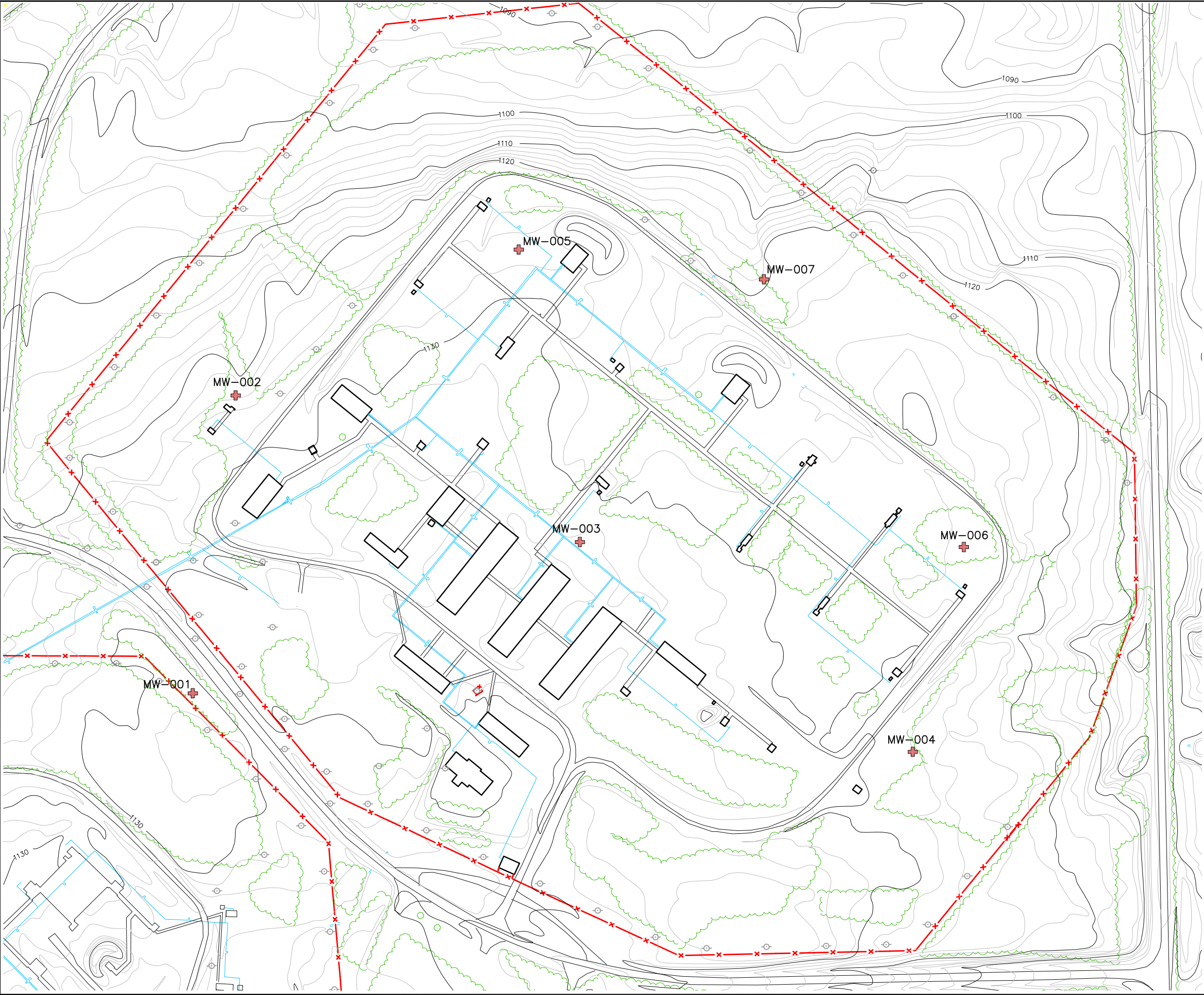
17 During the RI, seven monitoring wells were installed in the shallow consolidated water table
18 zone at LL-9 (see Figure 3-3), slug tests were conducted, and groundwater samples were
19 collected. These groundwater activities were conducted to:

- 20 • Determine whether the production activities had adversely affected groundwater quality
21 downgradient from the process buildings,
- 22 • Evaluate the quality of groundwater upgradient of LL-9, and
- 23 • Collect data about the LL-9 groundwater flow regime.

24 Monitoring wells were sited to evaluate potential subsurface contamination. Six wells were
25 located downgradient from the areas most likely affected by production activities.

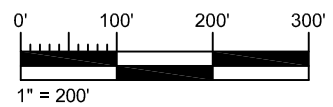
26 3.1.7.1 Monitoring Well Installation

27 The installation, development, and sampling of monitoring wells was conducted in accordance
28 with Section 4.3.2 of the FWSAP (USACE, 2001a) and the FWSAP Addendum for the Remedial
29 Investigation at LL 9 (RVAAP 42) (MKM, 2003a).



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- MONITORING WELL SAMPLE LOCATION
- BUILDING
- FENCE



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 3-3
GROUNDWATER MONITORING WELL
LOCATION MAP

LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	R. BOWMAN	APPR. BY	SL



1 The monitoring wells were installed using hollow stem auger (HSA) drilling methods under the
2 direct supervision of a qualified geologist. A 15.9 cm (6.25 inch) inside-diameter, hollow-stem
3 auger was used to advance the borehole through unconsolidated material to an average depth
4 of 21.04 ft bgs.

5 Monitoring wells were constructed in each borehole, following termination of drilling at the
6 appropriate depth. A 3.05 m (10 ft) section of new, precleaned 5.0 cm (2.0 inch) schedule 40
7 polyvinyl chloride (PVC) with a 0.010 slot screen was set to straddle the static water level
8 determined during drilling activities. Schedule 40 PVC riser was used to complete the well. The
9 screen and riser were placed into the borehole through the drill stem augers during well
10 construction. A clean sand filter pack was tremied in place from the bottom of the boring to
11 approximately 0.6 m (2 ft) above the top of the well screen. The filter pack was sealed with 0.6
12 m (2 ft) of bentonite pellets. Type 1 Portland cement with 7% bentonite grout was tremied to fill
13 the remainder of annular space to the surface. Each well was finished at the surface with
14 protective steel surface casing or surface flush mount. At most locations (where depth to water
15 was 12 ft or greater), the wells were finished at the surface with a protective steel surface casing
16 (pro-casing). Surface (flush) mounts were used where depth to water was less than 12 ft and
17 pro-casings would not fit (refer to Technical Change Letter 12 Nov 03 located in Appendix J).
18 Three steel posts were installed around each well. Monitoring well installation procedures are
19 provided in Section 4.3.2 of the FWSAP for RVAAP (USACE, 2001a). Well construction
20 diagrams are included in Appendix K.

21 Each monitoring well was developed at least 48 hours (and no longer than 7 days) after
22 completion. Static water level and total well depth were recorded. At least five borehole
23 volumes were removed from each well using a submersible pump. Wells were surged with
24 surge blocks, tubing, pumps, check valves, and bailers to remove sediment from screens. For
25 several wells a turbidity of ≤ 5 Nephelometric Turbidity Unit (NTU) could not be reached given
26 the nature of the unconsolidated water bearing zone. Low-yield wells with turbidity > 5 NTU
27 were stopped after 7 borehole volumes were removed or after 48 hours of development. Each
28 well was allowed 48 hours to recover before resampling. All of the issues/procedures were
29 discussed with Ohio EPA during the field investigations. Sediment volume in some of the
30 monitoring wells necessitated the use of alternative development methods to remove the
31 sediment. These methods included tubing and check valve surging. Well development and
32 purging records are included in Appendix K.

33 3.1.7.2 *In-Situ Permeability Testing*

34 Slug tests were performed at the seven LL-9 wells to estimate the hydraulic conductivity of the
35 consolidated bedrock material surrounding each well screen. A transducer was used to collect
36 the falling and rising head permeability data. First, the rising head test was conducted by



inserting a stainless steel slug into the well and recording water levels until the groundwater returned to static levels. After it was determined that the groundwater elevations had stabilized, the falling head test was conducted by removing the slug and collecting data until static conditions were achieved. The slug testing of monitoring wells was conducted in accordance with the FWSAP (USACE, 2001a) and the FWSAP Addendum for LL-9 (MKM, 2003a). Slug test data records are provided in Appendix L. Results of the slug tests are discussed in Section 4.2.1.

3.1.7.3 Groundwater Sampling

Low-flow micropurge sampling technology was used to collect groundwater samples at LL-9. Before samples were collected, the well condition was evaluated and documented on the field form. The casing headspace was analyzed using a hand-held PID and the depth to water and depth to the bottom of the well casing were measured and recorded. Each well was purged using micropurge technology. Purging continued until water quality indicators such as pH, temperature, dissolved oxygen, and conductivity were stabilized (three consecutive readings within 10% of each other). Samples were collected within 24 hours of purging and placed into pre-cleaned bottles provided by STL. All parameters, except VOCs, were collected using low-flow sampling technology. VOC samples were collected with a 2 inch bailer so that potential VOCs were not volatilized or bypassed if light non-aqueous phase liquids were present.

Groundwater sampling was conducted in December 2003 in accordance with the FWSAP (USACE, 2001a). Characteristics of the materials in which the wells are set include abundant mobile fines. Historically, turbidity readings are > 5 NTU have been prevalent at the conclusion of purging. Only bedrock wells have achieved turbidity < 5 NTU. Groundwater samples that were analyzed for TAL dissolved metals were field-filtered immediately after sample collection. Samples were placed into a cooler containing ice and submitted to the lab with completed chain-of-custody forms. Table 3-2 lists the analyses completed for the groundwater samples. Figure 3-3 shows the locations of the monitoring wells. Well purging and sampling forms are provided in Appendix K. Groundwater analytical results are included in Appendix M.

3.1.7.4 Water Level Measurements

Static groundwater levels and total depth measurements were taken and recorded at each monitoring well on three separate occasions to provide LL-9 groundwater flow data. The groundwater elevation data were collected during December 2003 (twice) and June 2004 and are shown in Table 3-3. Additional groundwater elevation data are included in Appendix K. Well survey information is included in Appendix N.



Table 3-3

LL-9 Groundwater Elevations

Easting	Northing	Well ID	TOC	Depth To Water (ft)			GW Elevation (ft MSL)		
				Dec-03	Dec-03	Jun-04	Dec-03	Dec-03	Jun-04
2355817.04	556125.81	LL9MW-001	1134.62	14.22	13.75	14.50	1120.40	1120.87	1120.12
2355907.76	556755.11	LL9MW-002	1127.30	10.10	9.75	9.50	1117.20	1117.55	1117.80
2356635.21	556445.31	LL9MW-003	1135.76	11.43	11.82	11.37	1124.33	1123.94	1124.39
2357338.76	556002.00	LL9MW-004	1131.83	18.69	18.23	18.27	1113.14	1113.60	1113.56
2356505.95	557063.36	LL9MW-005	1130.93	14.42	14.30	16.93	1116.51	1116.63	1114.00
2357446.67	556434.79	LL9MW-006	1129.88	17.04	17.87	14.32	1112.84	1112.01	1115.56
2357024.34	557000.56	LL9MW-007	1119.99	7.97	7.85	8.00	1112.02	1112.14	1111.99

Notes:

ft Feet
GW groundwater
mean sea
MSL level
top of
TOC casing

3.1.8 Surface Water

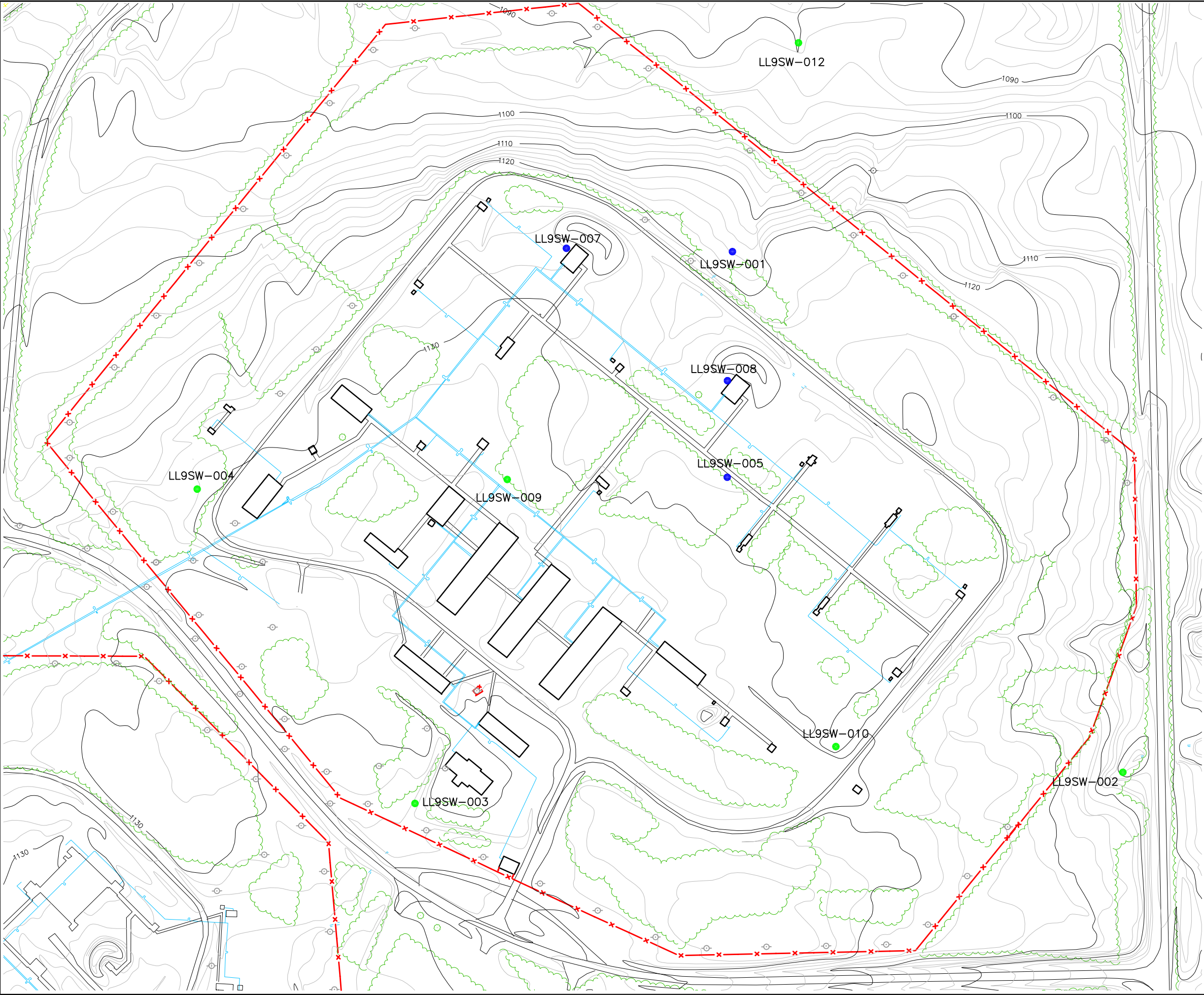
3.1.8.1 Azide Screening Surface Water Sample

Four surface water samples (LL9SW-001-0001-SW, LL9SW-005-0001-SW, LL9SW-007-0001-SW, and LL9SW-008-0001-SW) were collected from LL-9 during the March 2002 azide screening event. In addition, one split sample was collected for the USACE and submitted for analysis. Three locations (LL9SW-002-0001-SW, LL9SW-004-0001-SW, and LL9SW-006-0001-SW) had no surface water present for sample collection. The surface water samples were collected using direct fill of hand-held bottles. Four surface water samples (LL9SW-001-0001-SW, LL9SW-005-0001-SW, LL9SW-007-0001-SW, and LL9SW-008-0001-SW) were submitted to STL for explosives and metals analysis. Figure 3-4 shows the locations of the surface water samples. Surface water sampling forms are included in Appendix O and analytical results from the samples are included in Appendix P.

3.1.8.2 RI Surface Water Samples

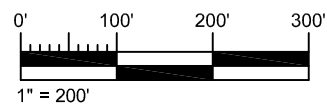
Surface water samples were collected from the main drainage ways at the LL-9 to:

- Evaluate whether surface water is being affected by runoff from the former production area



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- SURFACE WATER SAMPLING LOCATION
- SURFACE WATER SAMPLING LOCATION TAKEN DURING AZIDE SCREENING
- BUILDING
- FENCE



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 3-4
SURFACE WATER SAMPLE
LOCATION MAP
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	R. BOWMAN	APPR. BY	SL



- Identify the migration pathways for potentially contaminated runoff (if any) from the process area, and
- Evaluate whether contaminants migrate beyond the site boundary via LL-9 drainage ditches.

Four locations (LL9SW-002-0001-SW to LL9SW-004-0001SW, and LL9SW-012-0001-SW) were selected to evaluate whether contaminants could be affecting surface water within the AOC boundary (Figure 3-4). Samples LL9SW-009-0001-SW and LL9SW-010-0001-SW were collected from sewer manholes (see subsection 3.1.10.1). In addition, one split sample (LL9SW-012-0001-DUP) was collected for the USACE and submitted for analysis.

At each location, surface water sample collection began at the furthest downstream point and moved upstream, thus minimizing disturbance of the sediment and the turbidity of the collected sample. The surface water was collected as described in Section 4.6.2.1.1 of the FWSAP (USACE, 2001a) and the FWSAP Addendum for LL-9 (MKM, 2003a). Direct fill of hand-held bottles was used to sample water in the drainage ditches. Each container was submerged into the water, with the cap in place. The cap was removed, and the container was allowed to fill slowly and continuously. Water quality measurements (pH, conductivity, dissolved oxygen content, and temperature) were recorded just before to sample collection. Samples were immediately placed into a cooler containing ice, and the cooler was submitted to the laboratory with completed chain-of-custody forms. Table 3-1 lists the surface water samples and analyses. Field sampling forms are provided in Appendix O, and analytical results are included in Appendix P.

3.1.9 Sediment

3.1.9.1 Azide Screening Sediment Sample

Eight sediment samples (LL9SD-001-0001-SD thru LL9SD-008-0001-SD) were collected from LL-9 during the March 2002 azide screening event. Six were ditch sediment samples, and two were sump sediment samples. The sediment samples were co-located with the surface water sample locations. The sediment samples were collected using a 0.61-m (2-ft) hand-core sampler. Four samples (LL9SD-001-0001-SD, LL9SD-003-0001-SD, W-24S, and LL9SD-005-0001-SD) were submitted to STL for explosives and metals analysis. Figure 3-5 shows the locations of the sediment samples. Field sampling forms are included in Appendix Q, and analytical results from the samples are included in Appendix R.

3.1.9.2 RI Sediment Samples

Sediment samples were collected from the main drainage ways at the LL-9 to:



- Evaluate whether sediments are being affected via surface water runoff at the LL-9,
- Identify the migration pathway for potential contaminants that may have been suspended in surface water runoff, and
- Evaluate whether contaminants may have migrated beyond the AOC boundaries.

Ten locations (LL9SD-002-0001-SD, LL9SD-004-0001-SD, LL9SD-006-0001-SDS, and LL9SD-011-0001-SD through LL9SD-017-0001-SD) were selected to evaluate whether the drainage system at LL-9 allowed contaminants to migrate beyond the site boundary (Figure 3-5). Two samples (LL9SD-009-0001-SD and LL9SD-010-0001-SD) were collected from sewer manholes. In addition, two split samples (LL9SD-015-0001-DUP and LL9SD-016-0001-DUP) were collected and submitted for analysis. Three of the sediment samples (LL9SD-002-0001-SD, LL9SD-004-0001-SD, and LL9SD-012-0001-SD) were co-located with the surface water sample locations.

The sediment samples were collected from 0 to 0.15 m (0 to 0.5 ft) below the sediment-water interface along the drainage ditches. Sediment samples were collected using a scoop or trowel. Sampling procedures were in accordance with the FWSAP (USACE, 2001a), Section 4.5.2.1, and the FWSAP Addendum for LL-9 (MKM, 2003a). Sediment samples from LL-9 were homogenized and placed into sample containers. Samples were immediately placed into a cooler containing ice and submitted to the laboratory with completed chain-of-custody forms.

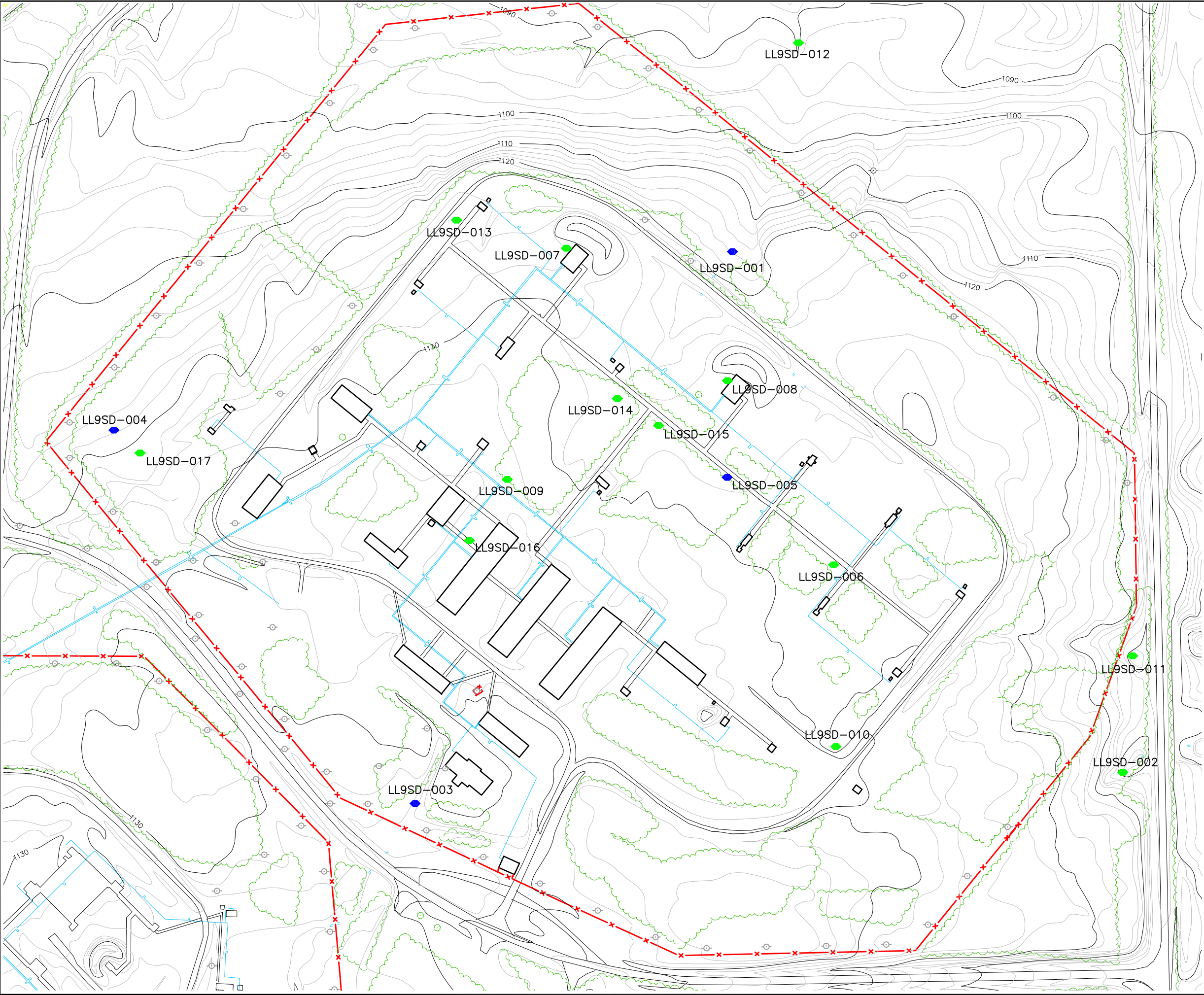
Table 3-1 lists the analyses completed on the sediment samples. Field sampling forms are included in Appendix Q, and analytical results from the samples are included in Appendix R.

3.1.10 Other RI Samples

Environmental samples were collected from sewers and from the soil/sub-base interface beneath the buildings located at LL 9. In addition, 28 samples were collected from two monitoring well borehole locations and from the sediment sample locations. These samples are discussed in Subsections 3.1.10.1 through 3.1.10.4.

No water or sediment sample was collected from the sumps at LL-9 during the RI because the sumps sampled during the azide sediment screening effort were excavated during the demolition operations, which were conducted before the RI activities began. If a sample was not obtained, it was noted on a field sampling report.

Environmental samples were also collected from the soil/sub-base interface beneath the buildings located at LL-9. Geotechnical samples were collected from two of the monitoring well borehole locations and from sediment sample locations.



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- SEDIMENT SAMPLING LOCATION
- SEDIMENT SAMPLING LOCATION TAKEN DURING AZIDE SCREENING
- BUILDING
- FENCE

0' 100' 200' 300'

1" = 200'

N



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

**FIGURE 3-5
SEDIMENT SAMPLE
LOCATION MAP**

LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	R. BOWMAN	APPR. BY	SL



3.1.10.1 Sewer Manhole Water Sample

Water samples were collected from sewer manholes, near the center of the site (LL9SW-009-0001-SW) and the eastern boundary (LL9SW-010-0001-SW) of the LL-9 site. The samples were collected to evaluate whether process operations had affected the sewer manholes. When sewer manhole samples were collected, no personnel entered the structures. A disposable Teflon® bailer was lowered into the manhole to collect the standing water sample. Water samples were immediately placed into a cooler containing ice and submitted to the laboratory with completed chain-of-custody forms. Water samples were not collected from the sewers and sumps located elsewhere at LL-9 because they contained insufficient water for sample collection. Table 3-1 lists the sewer water samples and analyses.

The field sampling forms for the sewer water samples are provided in Appendix O. The analytical results for the sewer water are included in Appendix P.

3.1.10.2 Sewer Manhole Sediment Sample

Sediment samples were collected from sewer manholes near the center of the site (LL9SD-009-0001-SD) and the eastern boundary (LL9SD-010-0001-SD) of the LL-9 site. Sediment samples were not collected from the sewers and sumps located elsewhere at LL-9 because they contained insufficient sediment for sample collection. Table 3-1 lists the sewer sediment samples and analyses.

The field sampling forms for the sewer sediment samples are provided in Appendix Q. The analytical results for the sewer sediment samples are included in Appendix R.

3.1.10.3 Subfloor Samples

Subfloor samples (LL9SB-048-0001-SO through LL9SB-053-0001-SO) at Buildings DT-6, DT-16, DT-18A, DT-21, and DT-55 were collected at the soil/sub-base interface to assess potential impact on the underlying soils from historic activities at LL-9. The location of the subfloor samples are shown in the approved FWSAP Addendum for LL-9 (MKM, 2003a). Although these samples were originally scoped to be collected by boring through the concrete slabs into the undisturbed soil in the 0-1 ft interval, demolition activities were conducted at LL-9 before the RI sampling. The heavy equipment (excavators, bulldozers, and loaders) tracked within the footprint of the buildings, and the subfloor soil was disturbed to depths sometimes greater than 1 ft below the subfloor base. Therefore, subfloor soil in the 0-1 ft interval was not representative of the pre-demolition conditions and samples were collected from the 1-3 ft interval below the disturbed soil. Two samples were collected from Building DT-21 and one each from Buildings DT-6, DT-16, DT-18A, and DT-55 (Figure 3-2).



The soil borings were hand augered from 1 to 3 ft bgs. One discrete soil sample was then collected from each boring location using a clean, decontaminated, stainless steel hand auger. Samples were immediately placed into a cooler containing ice and submitted to the laboratory with completed chain-of-custody forms. Field sampling forms are provided in Appendix H, and analytical results from the samples are included in Appendix I.

3.1.10.4 Geotechnical Samples

Geotechnical samples were collected from two of the seven monitoring well borehole locations in accordance with the scoping meeting held before field operations began. The sample borings were selected in the field by the geologist based upon site knowledge and field conditions encountered during the installation of the borings. The selection of the depth interval was limited because of the shallow occurrence of bedrock (unconsolidated between 0–6 ft bgs). Geotechnical analysis included grain size distribution, moisture content, TOC, hydraulic conductivity, and Atterberg limits. The analytical results match the typical distributions and qualities anticipated for the till/soils types at LLC9. The samples were collected using direct-push Shelby tubes and submitted to a geotechnical laboratory for analysis. Samples were collected from:

- LL9SB-033-0001-SO (MW-001) from 3-ft depth, and
- LL9SB-035-0001-SO (MW-003) from 4-ft depth.

These samples were sent to Prime Engineering & Architecture, Inc., for grain size, hydraulic conductivity, and other geotechnical analyses.

Geotechnical samples for grain size were also obtained from sediment sample locations: LL9SD-002-0001-SD, LL9SD-004-0001-SD, LL9SD-006-0001-SD, LL9SD-009-0001-SD, LL9SD-010-0001-SD, LL9SD-011-0001-SD, LL9SD-012-0001-SDS, LL9SD-013-0001SD, LL9SD-014-0001-SD, LL9SD-015-0001-SD (and duplicate), LL9SD-016-0001-SD (and duplicate), and LL9SD-017-0001-SD. These samples were sent to STL for grain size analysis.

Geotechnical results are included in Appendix S.

3.1.11 Sampling Location Survey

Each newly installed monitoring well, soil, surface water, and sediment sampling location was surveyed as specified in subsection 4.3.2.3.12 of the FWSAP (USACE, 2001a). The soil, surface water, sediment, and monitoring well location survey data can be found in Appendix N.



3.1.12 Investigation-Derived Waste

The IDW (soil and water) generated during the field activities was characterized and disposed. Representative composite samples of the waste soil and water were collected and analyzed per the requirements of the disposal facility. The LL-9 waste management disposition operations are summarized as follows:

- Excess soil cuttings generated from shallow soil sampling, drilling operations, and sediment sampling were containerized in covered roll off boxes and stored until disposition. Based on the composite sampling and analyses, the soil was determined to be non-hazardous. These materials were disposed at the Countywide R & D Landfill regional disposal facility (RDF) landfill in East Sparta, Ohio.
- Monitoring well development/purging and equipment decontamination generated wastewater. Based on laboratory analytical results, the wastewater was determined to be non-hazardous and transported to McCutcheon Enterprises Inc., a bio-solids treatment facility in Cleveland, Ohio, for disposal.

Representative composite samples from each waste stream were collected and analyzed per requirements of the disposal facility. The wastes generated during the LL-9 RI were managed and disposed in accordance with the applicable federal, state and local rules, laws, and regulations. Appendix T contains copies of the composite sample field sampling forms, chain-of-custody forms for composite samples, analytical reports, and manifests used to track the IDW.

3.1.13 Sample Packaging and Shipping

The samples were packaged and shipped per Section 6.0 of the FWSAP (USACE, 2001a) and Section 7.0 of the FWSAP Addendum (MKM, 2003a). Samples were properly packaged for shipment and dispatched to STL and GPL for analysis with completed chain-of-custody forms. When transferring samples, the individual relinquishing custody and the individual receiving the samples signed their names and noted the date and time of transfer on the record. The shipments complied with applicable Department of Transportation regulations for environmental samples.

3.2 Data Analyses and Quality

This section briefly describes the data quality procedures that were followed, and then discusses the quality of the data collected. Supporting quality documentation and relevant records, reports, logs, field notebooks, pictures, subcontractor reports, correspondence, analytical data, and chain-of-custody forms will be maintained in the MKM project files. These



files will remain in the custody of the MKM Program Manager until they are transferred to JMC and OHARNG RTLS.

3.2.1 Field Laboratory Program

Surface soil samples collected during the azide screening were analyzed for lead azide and TNT and RDX using the Jenkins field screening method. QA procedures were completed in accordance with manufacturer requirements, including calibration of the instrument to achieve proper wave length readings and analysis of a control sample.

3.2.2 Laboratory Analysis

Analytical laboratory procedures were completed in accordance with applicable professional standards, USEPA requirements, government regulations and guidelines, and specific project goals and requirements. Samples collected during the investigation were analyzed by STL's facility in Chicago, Illinois, a USACE Center of Excellence-validated laboratory under contract for the investigation. The RI analyses were performed at the STL Chicago site with the exception of the propellant analyses (nitroglycerin, nitrocellulose, and nitroguanidine). STL completed these analyses at their West Sacramento, California, facility. Quality assurance (QA) samples (soil, sediment, surface water, and groundwater) were submitted to GPL in Gaithersburg, Maryland, for analysis.

Samples were analyzed in accordance with the *FWSAP* (USACE, 2001a) and the *FWSAP* Addendum for LL-9 (MKM, 2003a). The data quality objectives (DQOs) established for the LL-9 RI comply with USEPA Region V guidance. The LL-9 requirements for sample collection, handling, analysis criteria, target analytes, laboratory criteria, and data validation criteria are consistent with USEPA requirements for National Priority List sites. DQOs for this project included analytical precision, accuracy, representativeness, completeness, comparability, and sensitivity for the measurement data.

The analytical laboratory was required to strictly adhere to the *FWSAP* (USACE, 2001a) to ensure the quality and usability of the analytical data. The laboratory was required to perform analyses in compliance with USEPA SW-846 (USEPA, 1990), *Test Methods for Evaluating Solid Waste, Physical/Chemical Methods* analytical protocols. The laboratory used USEPA SW-846 analytical methods to analyze the samples for metals, VOCs, SVOCs, pesticides, PCBs, explosives, and cyanide. Laboratories were required to comply with the methods as written; recommended procedures suggested in the methods were considered to be requirements.

Quality control (QC) samples were obtained during the LL-9 RI following the requirements of Section 3.2 of the Facility-Wide Quality Assurance Project Plan (SAIC, 2001). QC samples for



this project included field blanks, trip blanks, field duplicates, laboratory method blanks, laboratory control samples, laboratory duplicates, and matrix spike/matrix spike duplicate (MS/MSD) samples.

- Field duplicates were collected at a frequency of 10%.
- Field equipment rinseates were collected at a frequency of 10% for samples collected with non-dedicated equipment.
- Trip blanks were included with each shipment containing aqueous VOC samples.
- A field blank was collected at the beginning of RI sampling activities.
- Split samples (soil, sediment, surface water, and groundwater) were collected at a frequency of 10% of the environmental samples, and were submitted to GPL in Gaithersburg, Maryland, for analysis.
- MS/MSDs were collected at a frequency of 5%.
- Laboratory method blanks and laboratory control samples were employed to determine the accuracy and precision of the analytical method as implemented by the laboratory.
- Matrix spike sample results provided information about the effect of the sample matrix on the measurement methodology.
- Laboratory sample duplicates and MSD results were used to determine the analytical reproducibility and precision of the analyses for the samples of interest.

Field blanks, consisting of potable water used in the decontamination process, equipment rinseate blanks, and trip blanks, were submitted for analysis along with field duplicate (co-located) samples to provide a means to assess the quality of the data resulting from the field sampling program.

- Field blank samples were analyzed to determine whether field procedures contributed to sample contamination.
- Equipment rinseate blanks were used to assess the adequacy of equipment decontamination processes for groundwater sample collection.
- Trip blanks were used to assess the potential for VOC contamination of samples during sample shipment and storage.
- Field duplicate samples were analyzed to determine sample heterogeneity and sampling methodology reproducibility.

Analytical data reports from STL were forwarded to Purves Environmental, an independent third party validator, for QA review, comparison, and validation. The QC results were evaluated and



1 summarized in the LL-9 Quality Control Summary Report (QCSR), which is included in
2 Appendix U.

3 **3.2.3 Data Review, Validation, and Quality Assessment**

4 Data were produced, reviewed, and reported by the laboratory in accordance with specifications
5 outlined in the FWSAP Addendum for LL-9 (MKM, 2003a) and the laboratory's QA manual.
6 Laboratory reports included documentation verifying compliance with sample log-in procedures,
7 analytical holding times, and QC procedures for analyses. The laboratory reports also provided
8 information about the percent of recovery attained in laboratory spike samples, calibration
9 curves (initial and continuing), dilutions, and detection limits. The laboratory flagged data if
10 results warranted.

11 STL performed in-house analytical data reduction under the direction of STL's Project Manager
12 and QA Officer. These individuals were responsible for assessing data quality and informing
13 MKM about any data that were considered to be unacceptable or that required caution on the
14 part of the data user in terms of its reliability. This notification allowed MKM to determine
15 whether recollection or reanalysis was required to achieve the project data quality objectives.
16 Data reduction, review, and reporting by the laboratory were conducted as follows:

- 17 • Raw data produced by the analyst were given to the analyst's area supervisor.
- 18 • The area supervisor reviewed the data to determine whether they attained the QC
19 criteria as outlined in the established methods and for overall reasonableness.
- 20 • Upon acceptance of the raw data by the area supervisor, a report was generated and
21 sent to STL's Laboratory Project Manager.
- 22 • STL's Project Manager reviewed the reports and, based on that review, generated final
23 reports.
- 24 • The final data were delivered to MKM, who forwarded the packages to Purves
25 Environmental for data validation.

26 STL prepared and retained full analytical and QC documentation for the project in both hard
27 (paper) copy and electronic storage media (e.g., magnetic tape) as directed by the analytical
28 methodologies employed. An electronic data deliverable (EDD) was prepared in the format
29 specified in the FWSAP (USACE, 2001a). STL provided the following information to MKM in
30 each analytical data package submitted:

- 31 • Cover sheets listing the samples included in the report and narrative comments
32 describing problems encountered (if any) during analysis;
- 33 • Tabulated results of inorganic and organic compounds identified and quantified; and



- Analytical results for QC sample spikes, sample duplicates, and initial and continuing calibration verifications of standards and blanks, method blanks, and laboratory control sample information.

MKM compared the data packages to the chain-of-custody forms to ensure that the requested analyses had been conducted and results provided. Purves Environmental systematically verified and validated the data. This process was conducted to ensure that the precision and accuracy of the analytical data were adequate for their intended use. The validation process minimized the potential of using false or negative results in the decision-making process and ensured that detected and non-detected compounds were accurately identified. This approach was consistent with the DQOs for the project and with the analytical methods, and appropriate for determining contaminants of concern and calculating risk.

The data validation determined that the data were 100% complete and usable and satisfied the DQOs for this project. Data validation reports are included in Appendix U.

3.3 Deviations from the Work Plan

Every effort was made to complete the field activities in accordance with the approved work plan. However, in some instances, circumstances or field conditions necessitated a modification. Those changes are noted below.

- Minor adjustments to surface soil sampling locations were made as a consequence of refusal.
- No surface water samples (LL9SW-002-0001-SW, LL9SW-004-0001-SW, and LL9SW-006-0001-SW) were collected from the sumps at LL-9 during the azide sampling because of insufficient sample media.
- Turbidity measured during well development, purging, and sampling did not achieve <5 NTU owing to the highly abundant and mobile fines in the formation.
- The MS/MSD frequency required by the SAP was achieved for Method 8330 analyses; however, the laboratory did not select project samples for MS/MSD analyses for all other methods. Therefore, the overall MS/MSD frequency did not meet the SAP requirements.

Although these deviations reflected unanticipated field conditions, they were discussed with Ohio EPA and the objectives of the RI were still achieved. The technical field change memos are located in Appendix J.



4.0 NATURE AND EXTENT OF CONTAMINATION

This section summarizes the soil (surface and subsurface), groundwater, surface water, sediment, and other analytical results. The results are organized by media or type of sample: soil (surface and subsurface), groundwater, surface water, and sediment, including samples such as standing water and residue found in the sumps and sewer manholes, and geotechnical data results. A table is provided at the end of this section that lists the analytical results that exceeded either the RVAAP installation background criteria (USACE, 2001b) criteria or Region 9 residential soil or tap water PRGs which were used to screen the initial results.

The RVAAP installation background concentrations were calculated from data obtained in the Winklepeck Burning Ground (WBG) Phase II RI (USACE, 2001b). The background criteria are based on concentrations of selected naturally occurring metals in soil, sediment, surface water, and groundwater. Background values were not established for explosives, propellants, VOCs, SVOCs, and TOC; therefore, the background value assigned to these compounds, for the purpose of this RI, is zero. For metals that were “non-detect” in the background study, the background value was established as zero (USACE, 2001b). Some metals (including cyanide and cadmium) are reported as greater than installation background criteria (USACE, 2001b) exceedances even though the reported concentrations are extremely low, compared to their respective PRGs. The Region 9 residential PRGs were obtained from the USEPA website. The Region 9 PRGs for non-carcinogenic (nc) concentrations are reported as one-tenth of the published Region 9 PRG value for soil and sediment media. This procedure takes into account the additive issues with multiple COPCs. The data tables found at the end of Section 4.0 include a column that lists the installation background criteria (USACE, 2001b) concentrations that were established for each analyte/matrix. No background criteria were established for RVAAP soils that exceed 3 ft in depth. Therefore, results for samples collected at depths greater than 3 ft were compared to the background values established for metals found in the 1–3 ft interval.

The significance of the contaminants detected in LL-9 media is discussed in the Human Health and Ecological Risk Assessments (Sections 5 and 6, respectively) of this document. QC sample results (equipment rinsewater samples, trip blanks, and field duplicates) are included in Appendix U.

4.1 Soils

This section summarizes the physical and chemical data from the analysis of surface and subsurface soil samples collected during the LL-9 RI.



4.1.1 Azide Screening Surface Soil Results

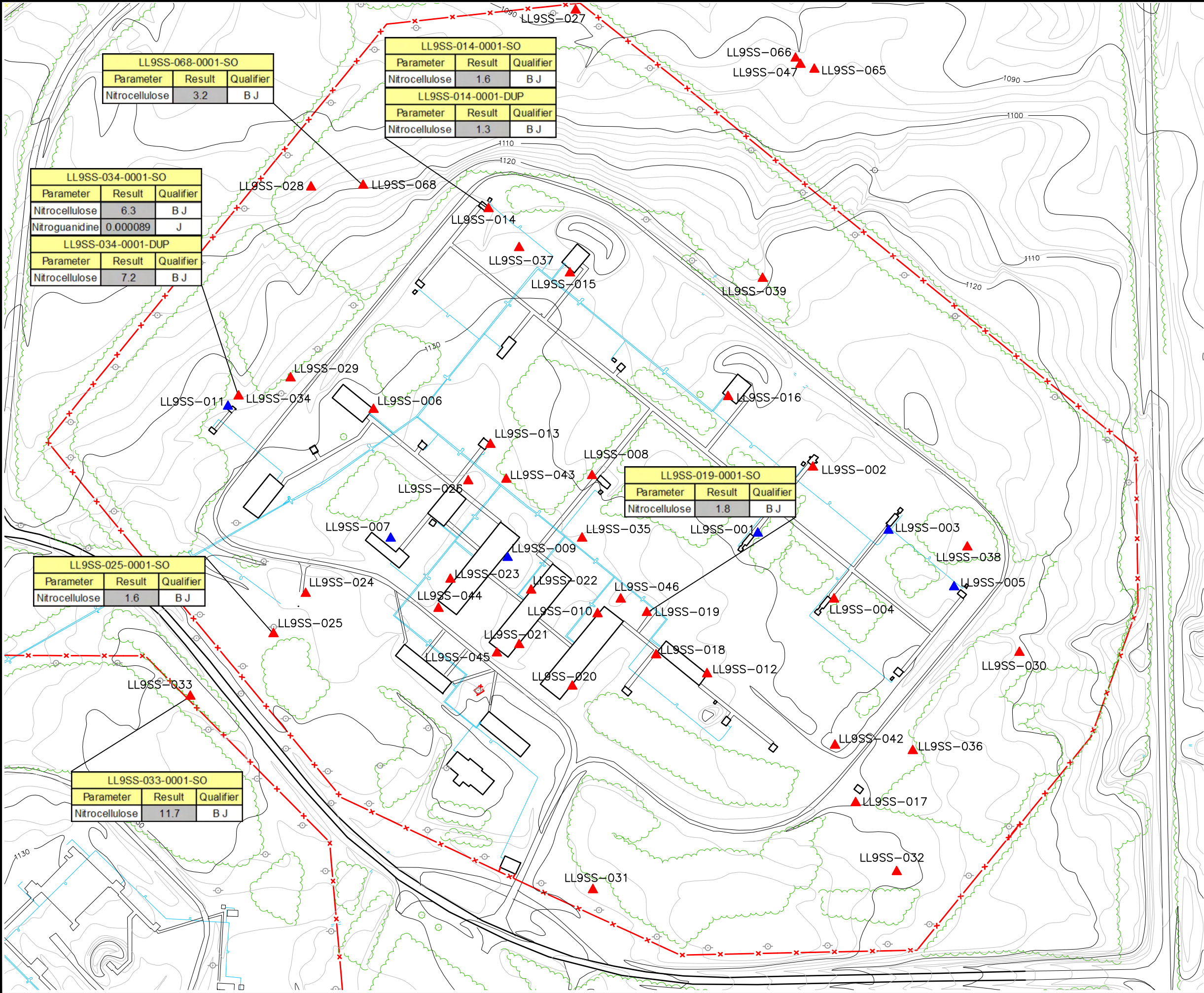
Five surface soil samples were collected and analyzed for explosives and metals during the March 2002 azide screening event; (LL9SS-001-0001-SO, LL9SS-003-0001-SO, LL9SS-005-0001-SO, LL9SS-007-0001-SO, and LL9SS-009-0001-SO). Analytical results are included in Appendix E and summarized in Table 4-1, Summary of Azide Screening Surface Soil Results (0-1 ft) (included at the end of Section 4.0). There were no explosives detected above the reporting limit. Metals exceeded the RVAAP-specific surface soil background values established by USACE (USACE, 2001b) and/or Region 9 residential soil PRG in five samples: LL9SS-001-0001-SO, LL9SS-003-0001-SO, LL9SS-005-0001-SO, LL9SS-007-0001-SO, and LL9SS-009-0001-SO. These exceedances are summarized in Table 4-2 Summary of Surface Soil Exceedences (0-1 ft bgs) (included at the end of Section 4.0).

4.1.2 RI Surface Soil Results

Forty surface soil samples and seven duplicates were collected at various locations at LL-9 during the RI. Each sample was collected from the surface to a depth of 1 ft (0-1ft) bgs. Soils collected at depths greater than 1 ft were considered subsurface soil samples and are discussed in Subsection 4.1.4. The surface soil samples were analyzed for TAL metals. In addition to TAL metals, 10% of the samples collected were analyzed for explosives, cyanides, VOCs, SVOCs, PCBs, pesticides, and propellants. Surface sample analytical results are included in Appendix F and summarized in Table 4-3, Summary of Surface Soil Results (0-1 ft) (included at the end of Section 4.0).

Propellants (nitrocellulose and nitroguanidine), SVOCs (benzo[a]pyrene and dibenzo[a,h]anthracene) and metals (aluminum, arsenic, barium, cadmium, chromium, cobalt, copper, iron, lead, manganese, mercury, nickel, selenium, vanadium, and zinc) were detected in surface soils at concentrations greater than RVAAP installation background (USACE, 2001b) concentrations, Region 9 residential soil preliminary remediation goals (PRGs), or both. Therefore, the lateral and vertical extent of explosives, propellants, SVOCs, and metals were not fully delineated during this Phase I RI. The lateral extent of surface soil contamination at LL-9 (as determined by this Phase I RI) is shown on Figures 4-1 through 4-3. The distribution of selected site related contaminants (aluminum, arsenic, chromium, copper, lead, manganese, and mercury) identified as COPCs in Section 6.2.3 of this document, are shown in Figures 4-4 through 4-11. The distribution and exposure pathways for SRCs are further discussed in the conceptual site model included in Section 5.4.

Exceedances for the 47 surface soil samples (including duplicates) and 5 surface soil samples collected during the azide sampling event are summarized in Table 4-2. Specific exceedences for each analytical parameter are discussed in the following subsections.



LL9SS-068-0001-SO		
Parameter	Result	Qualifier
Nitrocellulose	3.2	B J

LL9SS-014-0001-SO		
Parameter	Result	Qualifier
Nitrocellulose	1.6	B J
LL9SS-014-0001-DUP		
Parameter	Result	Qualifier
Nitrocellulose	1.3	B J

LL9SS-034-0001-SO		
Parameter	Result	Qualifier
Nitrocellulose	6.3	B J
Nitroguanidine	0.000089	J
LL9SS-034-0001-DUP		
Parameter	Result	Qualifier
Nitrocellulose	7.2	B J

LL9SS-019-0001-SO		
Parameter	Result	Qualifier
Nitrocellulose	1.8	B J

LL9SS-025-0001-SO		
Parameter	Result	Qualifier
Nitrocellulose	1.6	B J

LL9SS-033-0001-SO		
Parameter	Result	Qualifier
Nitrocellulose	11.7	B J

LEGEND:

VEGETATION

INTERMITTENT STREAM

STREAM

PIPING

ROAD

10 ft INTERVAL

2 ft INTERVAL

STEAM LINE POST

SURFACE SOIL
SAMPLE LOCATION

SURFACE SOIL
SAMPLING LOCATION
TAKEN DURING AZIDE SCREENING

BUILDING

FENCE

NOTES:

UNITS = mg/kg

IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED

IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT

IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED

IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

ORGANIC QUALIFIERS

U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT

J - RESULT IS AN ESTIMATED VALUE BELOW THE RL

ORGANIC FLAGS

B - METHOD BLANK: BATCH QC IS GREATER THAN RL

* - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT

^ - BATCH QC IS GREATER THAN RL

A - CONCENTRATION IS BELOW THE METHOD REPORTING LIMIT (RL)

H - ALTERNATE PEAK SELECTION UPON ANALYTICAL REVIEW

1" = 200'

MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

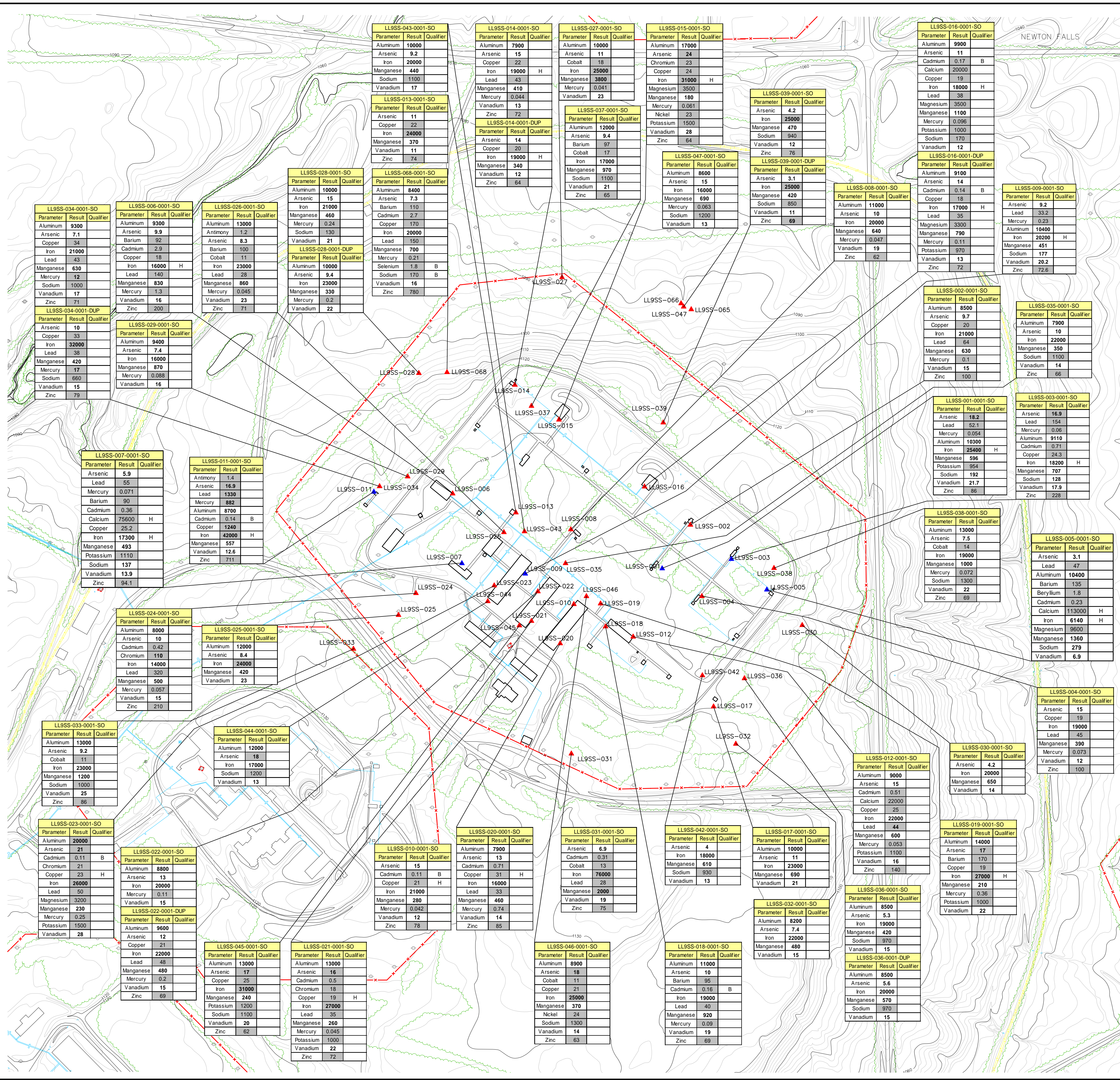
RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-1

SURFACE SOIL SAMPLE (0-1ft)
EXCEEDANCES EXPLOSIVE/PROPELLANTS

LL 9 REMEDIAL INVESTIGATION

SIZE	PROJECT NO.	DATE:	REV
B	02-02-0085	MAY 2007	2
DRAWN BY	R. BOWMAN	APPR. BY	SL



- LEGEND:
- VEGETATION
 - INTERMITTENT STREAM
 - STREAM
 - PIPING
 - ROAD
 - 10 FT. INTERVAL
 - 2 FT. INTERVAL
 - STREAM LINE POST
 - SAMPLE LOCATION
 - SAMPLE LOCATION
 - TAKEN DURING AZIDE SCREENING
 - BUILDING
 - FENCE

NOTES:

RESULTS SHOWN IN mg/kg

IF RESULT > BACKGROUND, THEN THE VALUE IS SHOWN IN BOLD

IF RESULT > THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT

IF RESULT > BACKGROUND AND THE PRG THE VALUE IS BOLD + SHOWN

IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

INORGANIC QUALIFIERS

U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT

J - RESULT IS LESS THAN THE RL, BUT GREATER THAN OR EQUAL TO THE MDL

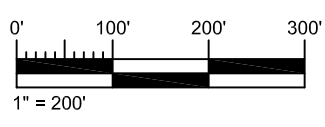
B - RESULT IS LESS THAN THE CRO/RL, BUT GREATER THAN OR EQUAL TO THE IDL/MDL

INORGANIC FLAGS

* - INSTRUMENT RELATED QC EXCEED THE UPPER OR LOWER CONTROL LIMITS

U - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT

H - BATCH QC IS GREATER THAN RL

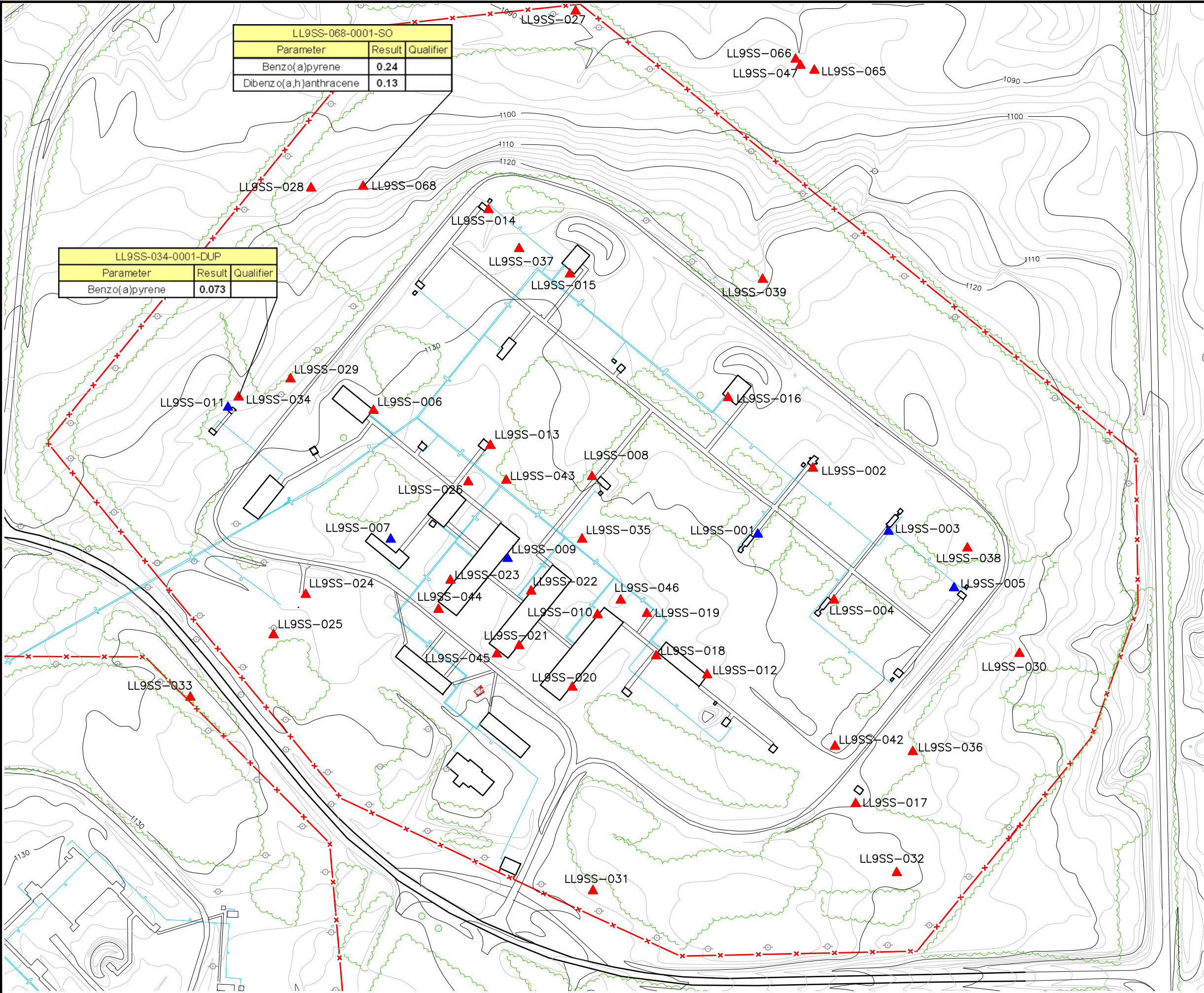


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-2
SURFACE SOIL SAMPLE (0-1ft)
EXCEEDANCES TAL METALS/CYANIDE
LL 9 REMEDIAL INVESTIGATION

SIZE	PROJECT NO.	DATE:	REV
D	02-02-0085	MAY 2007	2
DRAWN BY	R. BOWMAN	APPR. BY	SL



LEGEND:

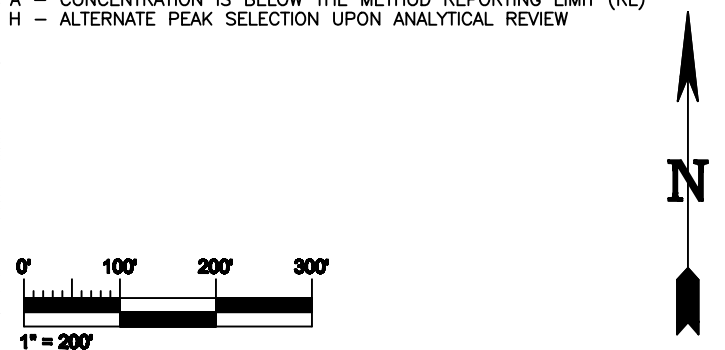
- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKE DURING AZIDE SCREENING
- BUILDING
- FENCE


NOTES:

RESULTS SHOWN IN mg/kg
IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED
IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT
IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED
IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

ORGANIC QUALIFIERS
U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT
J - RESULT IS AN ESTIMATED VALUE BELOW THE RL

ORGANIC FLAGS
B - METHOD BLANK: BATCH QC IS GREATER THAN RL
* - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT
^ - BATCH QC IS GREATER THAN RL
A - CONCENTRATION IS BELOW THE METHOD REPORTING LIMIT (RL)
H - ALTERNATE PEAK SELECTION UPON ANALYTICAL REVIEW



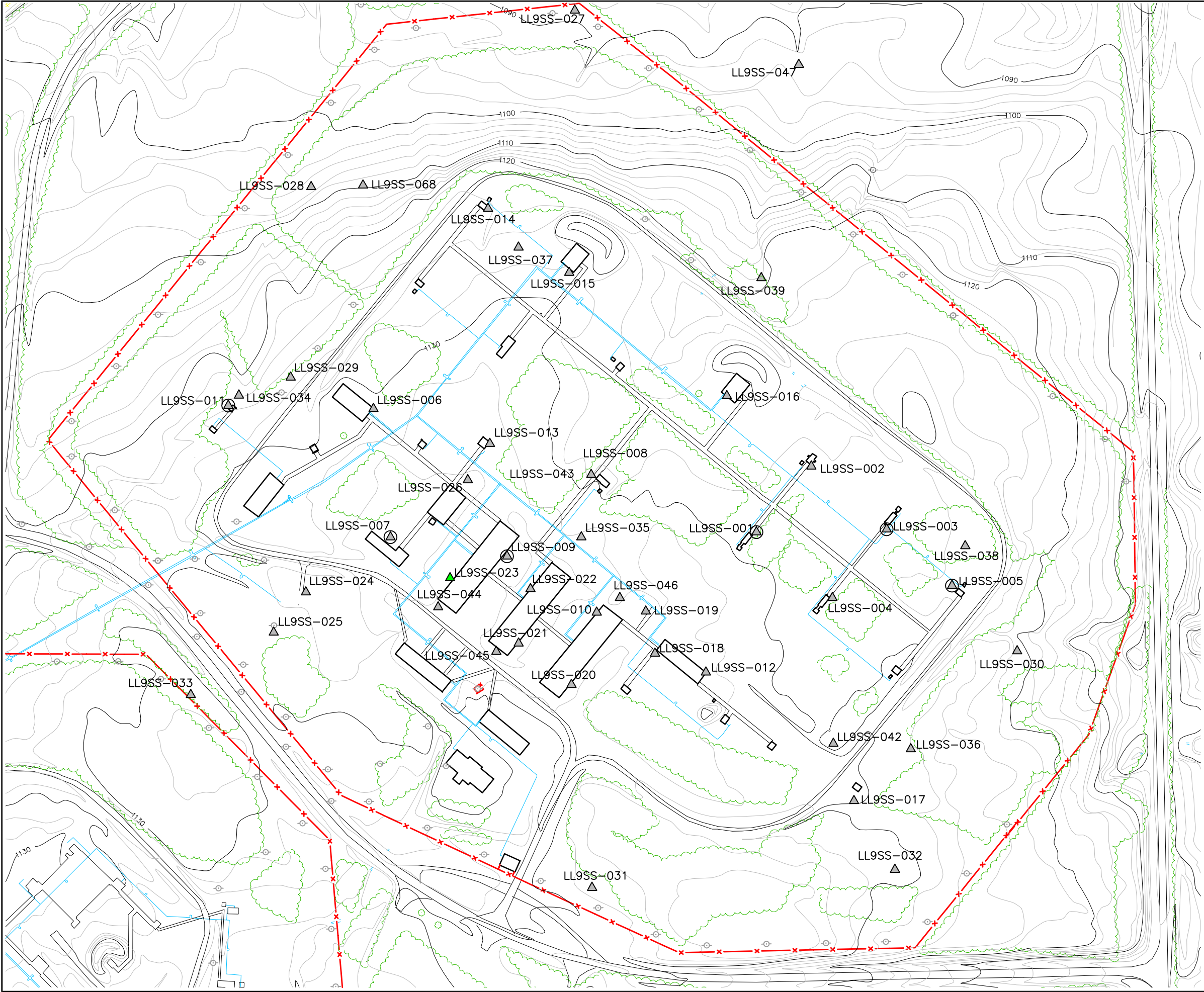


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

FIGURE 4-3
SURFACE SOIL SAMPLE (0-1ft)
EXCEEDANCES VOCs/SVOCs
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY K. PERRY	APPR. BY	SL	



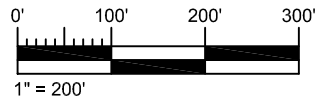
LEGEND:


- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

SURFACE BACKGROUND VALUE FOR ALUMINUM = 17700 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND



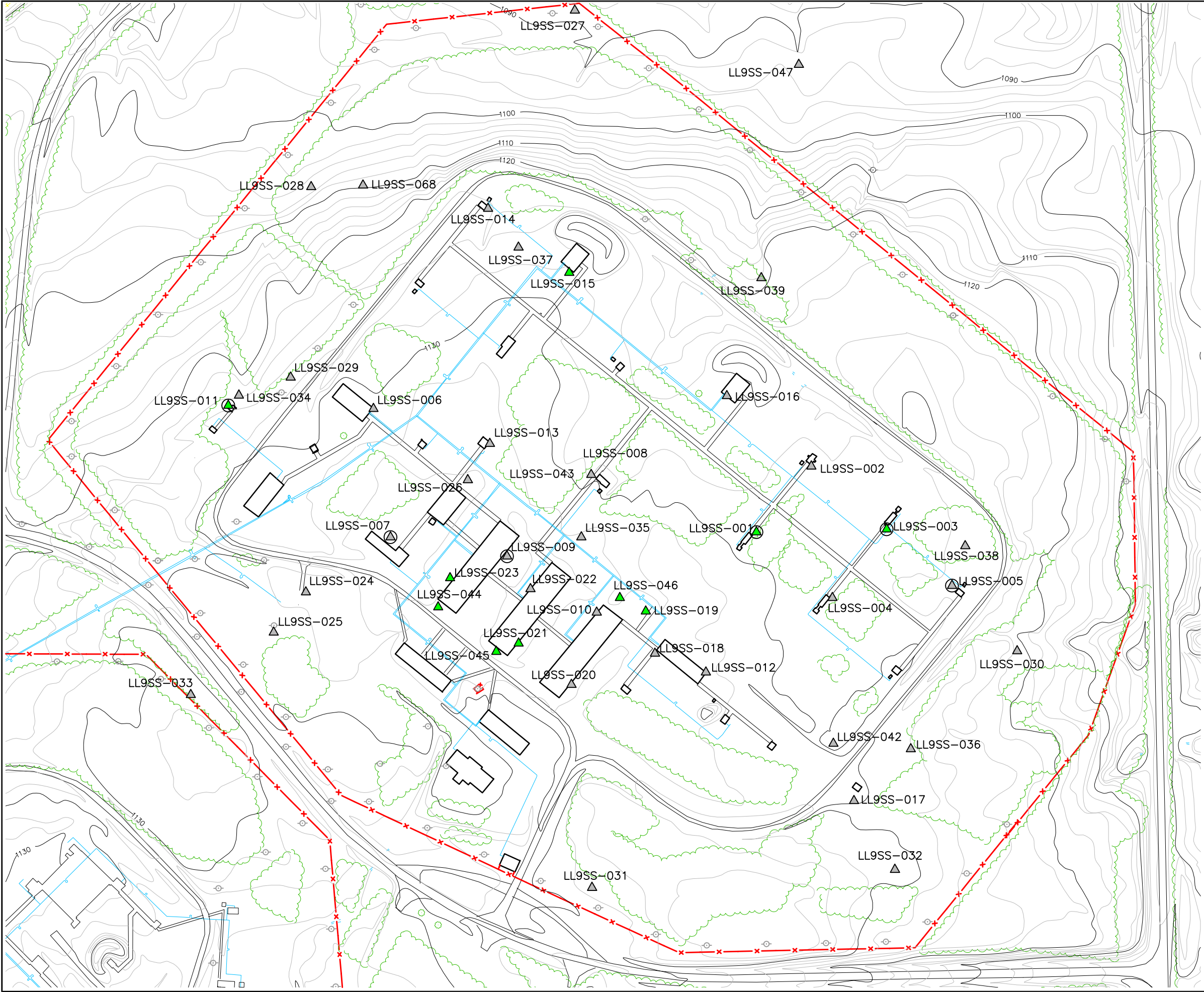


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-4
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF ALUMINUM
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



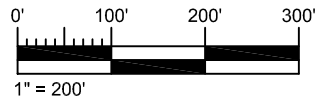
LEGEND:


- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

SURFACE BACKGROUND VALUE FOR ARSENIC = 15.4 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND



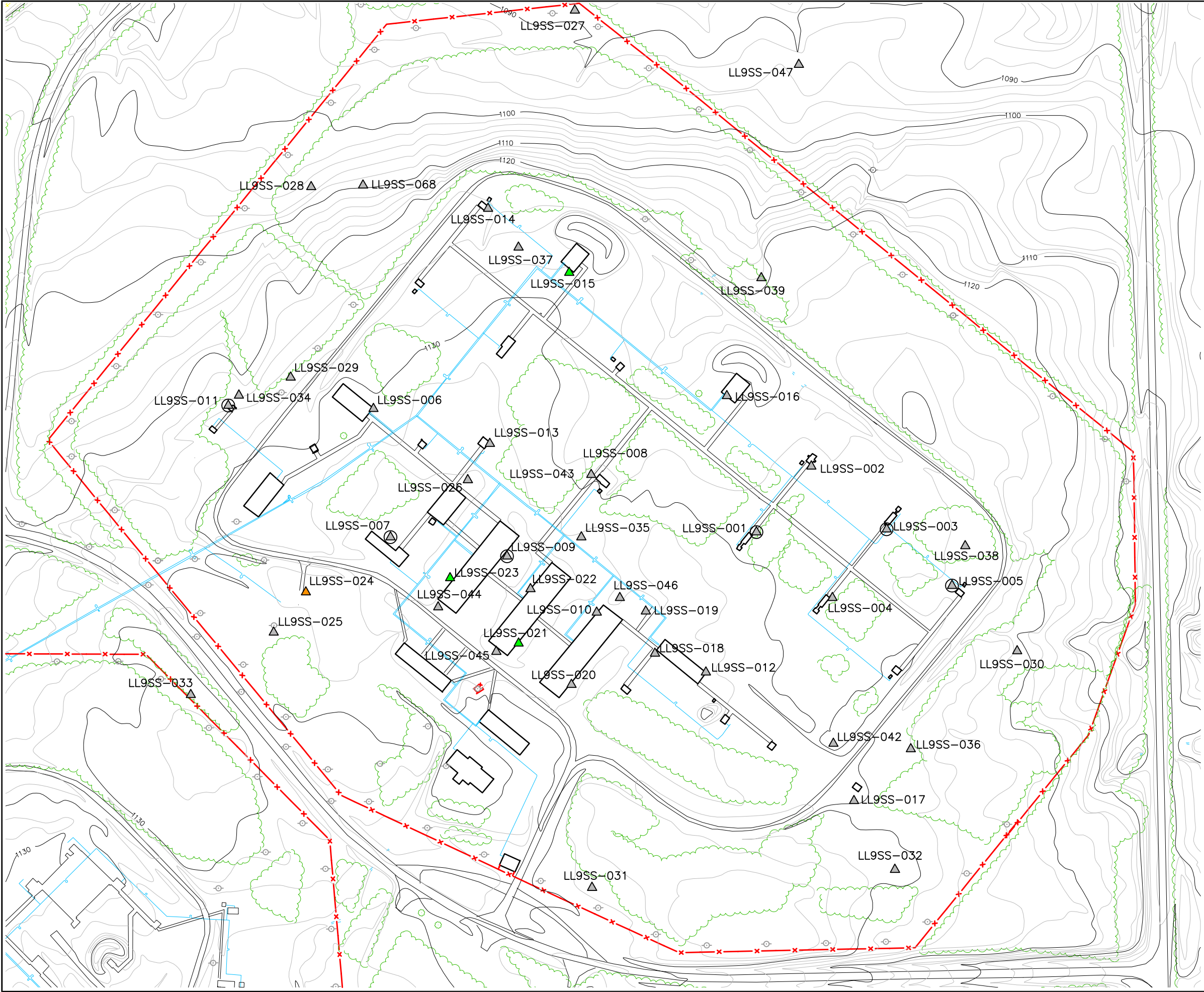


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-5
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF ARSENIC
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING


SURFACE BACKGROUND VALUE FOR CHROMIUM = 17.4 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND

0' 100' 200' 300'
1" = 200'

N

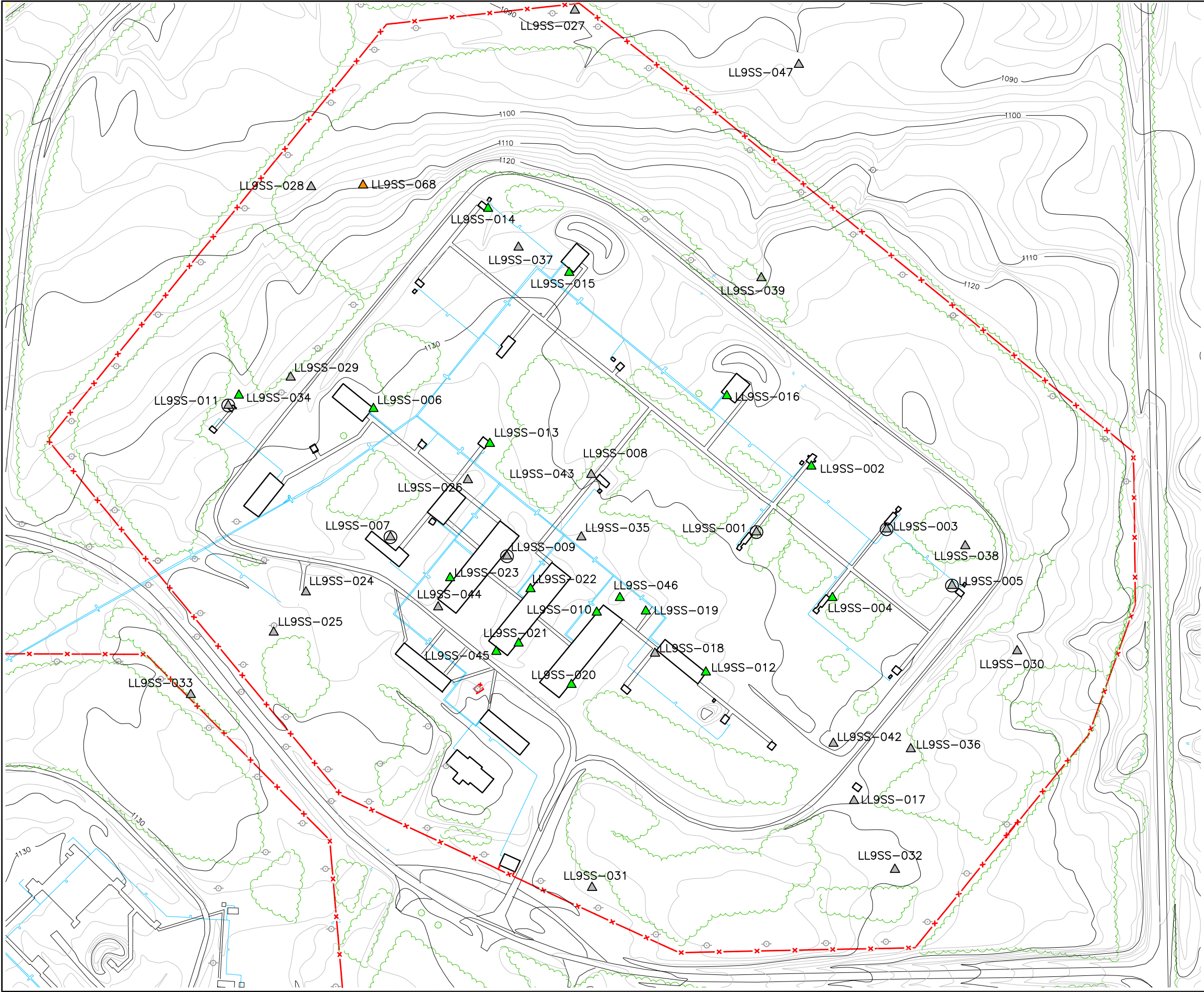


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-6
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF CHROMIUM
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



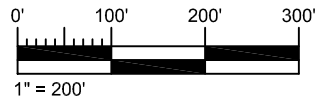
LEGEND:


- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

SURFACE BACKGROUND VALUE FOR COPPER = 17.7 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND



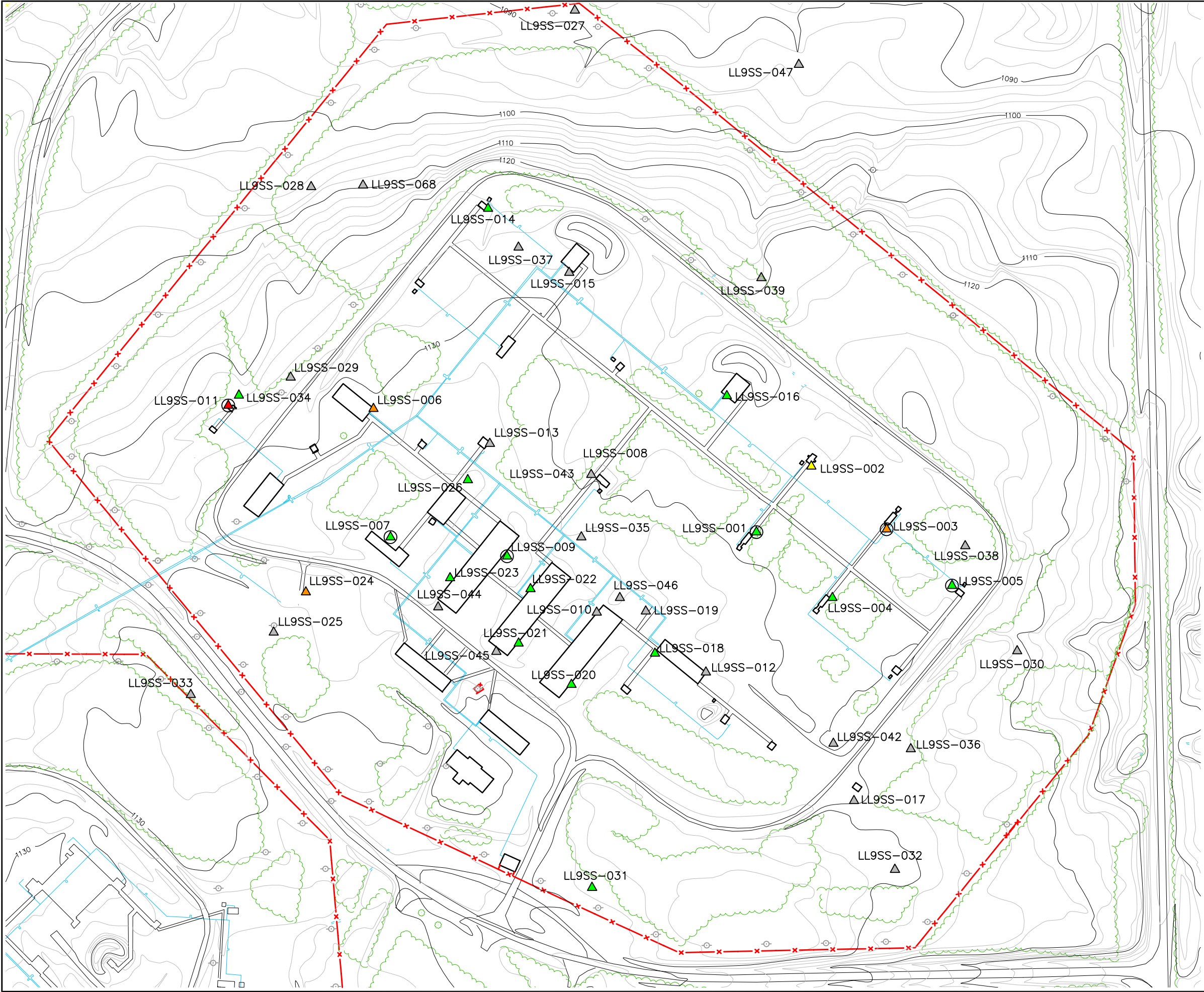


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-7
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF COPPER
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



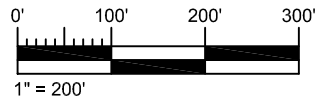
LEGEND:


- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

SURFACE BACKGROUND VALUE FOR LEAD = 26.1 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND



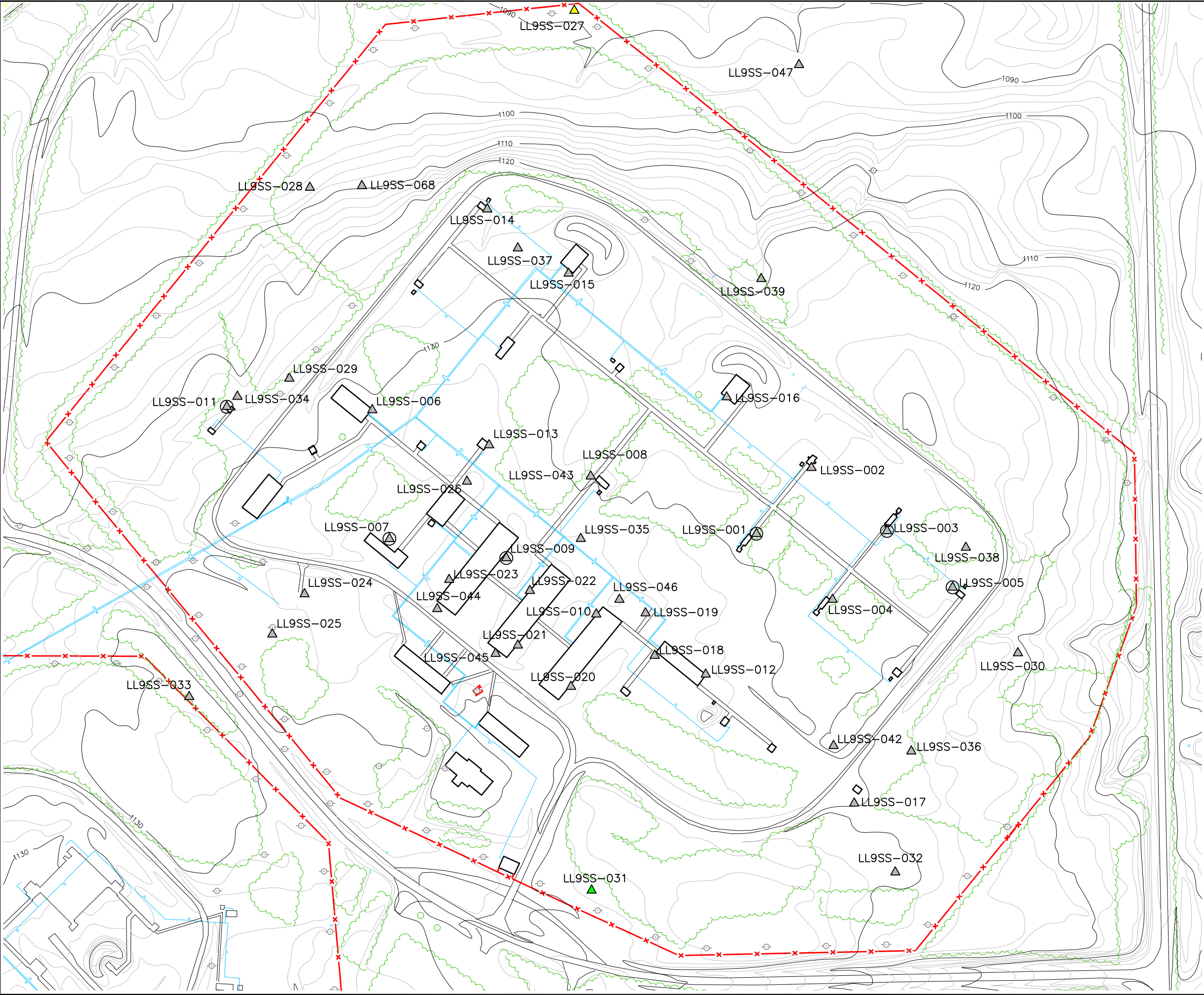


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-8
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF LEAD
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



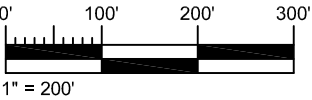
LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

SURFACE BACKGROUND VALUE FOR MANGANESE = 1450 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND



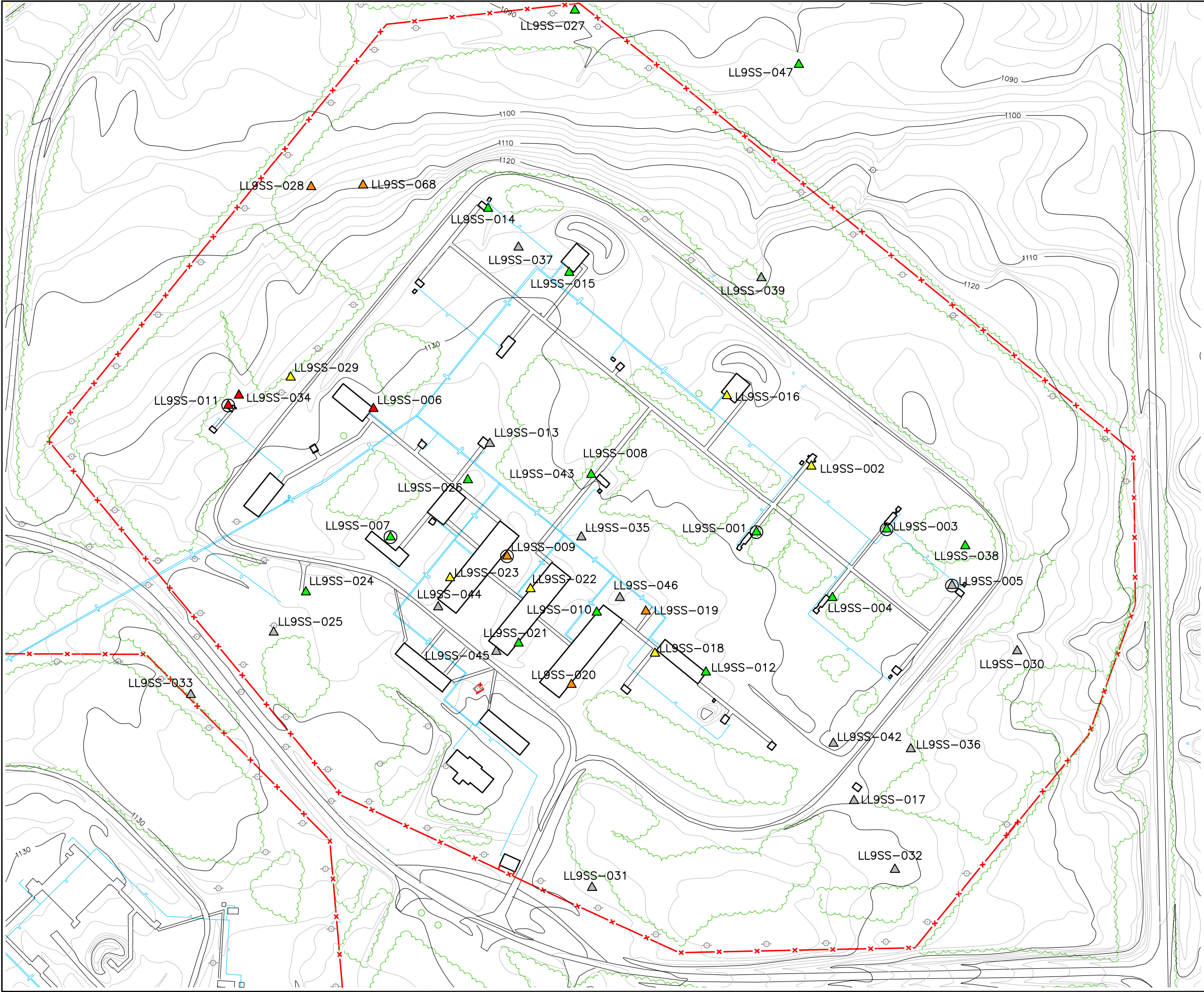


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-9
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF MANGANESE
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION
- SURFACE SOIL SAMPLING LOCATION TAKEN DURING AZIDE SCREENING


SURFACE BACKGROUND VALUE FOR MERCURY = 0.04 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND

0' 100' 200' 300'
1" = 200'

N



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

FIGURE 4-10
SURFACE SOIL SAMPLE (0-1ft)
DISTRIBUTION OF MERCURY
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



4.1.2.1 Explosives and Propellants

Explosives were not detected in the surface soil samples collected during the RI. Nitrocellulose was detected above the RVAAP installation background criteria (USACE, 2001b) in 6 samples and one duplicate. The maximum concentration of 3.2 (B)(J) mg/kg was detected in sample LL9SS-068-0001-SO. However, the maximum result was qualified as estimated (J) because the concentration of nitrocellulose in the method blank was detected above the reporting limit (B). A second propellant, nitroguanidine, was detected in sample LL9SS-034-0001-SO with an estimated concentration of 0.089 (J) mg/kg. Figure 4-1 shows surface soil exceedances for explosives and propellants at LL-9.

4.1.2.2 TAL Metals and Cyanide

Metals were detected in surface samples collected throughout the site. Figure 4-2 shows surface soil exceedances for TAL metals and cyanide at LL-9. Cyanide was not detected in any of the surface soil samples. The following metals were detected above RVAAP installation background criteria (USACE, 2001b) or Region 9 residential soil PRGs for surface soil (0-1 ft bgs):

- Aluminum was detected above the Region 9 residential soil PRG (7,614 mg/kg) in 36 samples, 4 of which were duplicates. Aluminum was also detected above the RVAAP installation background criterion (17,700 mg/kg) and Region 9 residential soil PRG in 1 sample. The maximum aluminum concentration of 20,000 mg/kg was detected in sample LL9SS-023-0001-SO.
- Arsenic was detected above the Region 9 residential soil PRG (0.39 mg/kg) in 40 sample locations, 7 of which were duplicates. Arsenic was also detected above the RVAAP installation background (15.4 mg/kg) and Region 9 residential soil PRG in 7 samples. The maximum arsenic concentration of 24 mg/kg was detected in sample LL9SS-015-0001-SO.
- Barium was detected above the RVAAP installation background criterion (88.4 mg/kg) in 6 samples with the maximum concentration of 170 mg/kg detected in sample LL9SS-019-0001-SO.
- Cadmium was detected above the RVAAP installation background criterion (0.0 mg/kg) in 11 samples, 1 of which was a duplicate. The maximum concentration of 2.9 mg/kg was detected in sample LL9SS-006-0001-SO.
- Calcium was detected above the RVAAP installation background criterion (15,800 mg/kg) in 2 samples. The maximum concentration of 22,000 mg/kg was detected in sample LL9SS-012-0001-SO.
- Chromium was detected above the RVAAP installation background criterion (17.4 mg/kg) at 3 sample locations. Chromium was also detected above the RVAAP installation background and Region 9 residential soil PRG (30 mg/kg) in 1 sample.



The maximum concentration of 110 mg/kg was detected in sample LL9SS-024-0001-SO.

- Cobalt was detected above the RVAAP installation background criterion (10.4 mg/kg) in 7 samples and the maximum concentration of 18 mg/kg was detected in sample LL9SS-027-0001-SO.
- Copper was detected above the RVAAP installation background criterion (17.7 mg/kg) at 21 sample locations, 4 of which were duplicates. The maximum concentration of 170 mg/kg was detected in sample LL9SS-068-0001-SO.
- Iron was detected above the Region 9 residential soil PRG (2,346 mg/kg) in 34 samples, 5 of which were duplicates. Iron was also detected above the RVAAP installation background criterion (23,100 mg/kg) and Region 9 residential soil PRG in 13 samples, 2 of which were duplicates. The maximum concentration of 76,000 mg/kg was detected in sample LL9SS-031-0001-SO.
- Lead was detected above the RVAAP installation background criterion (26.1 mg/kg) in 18 samples, 3 of which were duplicates. The maximum concentration of 320 mg/kg was detected in sample LL9SS-024-0001-SO.
- Magnesium was detected above the RVAAP installation background criterion (3,030 mg/kg) in 4 samples, 1 of which was a duplicate. The maximum concentration of 3,500 mg/kg was detected in samples LL9SS-015-0001-SO and LL9SS-016-0001-SO.
- Manganese was detected above the Region 9 residential soil PRG (176 mg/kg) in 39 samples, 7 of which were duplicates. Manganese was also detected above the RVAAP installation background criterion (1,450 mg/kg) and Region 9 residential soil PRG in 2 samples. The maximum concentration of 3,800 mg/kg was detected in sample LL9SS-027-0001-SO.
- Mercury was detected above the RVAAP installation background criterion (0.04 mg/kg) in 26 samples, 3 of which were duplicates. Mercury was also detected above the RVAAP installation background criterion and Region 9 residential soil PRG (2.3 mg/kg) in 2 samples, 1 of which was a duplicate. The maximum concentration of 17 mg/kg was detected in sample LL9SS-034-0001-DUP.
- Nickel was detected above the RVAAP installation background criterion (21.1 mg/kg) in 2 samples. The maximum concentration of 24 mg/kg was detected in sample LL9SS-046-0001-SO.
- Potassium was detected above the RVAAP installation background criterion (927 mg/kg) in 8 samples, one of which was a duplicate. The maximum concentration of 1,500 mg/kg was detected in samples LL9SS-015-0001-SO and LL9SS-023-0001-SO.
- Selenium was detected above the RVAAP installation background criterion (1.4 mg/kg) in 1 sample with a concentration of 1.8 mg/kg in sample LL9SS-068-0001-SO.



- Sodium was detected above RVAAP installation background criterion (123 mg/kg) in 19 samples, 3 of which were duplicates. The maximum concentration of 1,300 mg/kg was detected in samples LL9SS-038-0001-SO and LL 9SS-046-0001-SO.
- Vanadium was detected above the Region 9 residential soil PRG (7.8 mg/kg) in 47 samples, 7 of which were duplicates. The maximum concentration of 28 mg/kg was detected in samples LL9SS-015-001-SO and LL9SS-023-0001-SO.
- Zinc was detected above the RVAAP installation background criterion (61.8 mg/kg) in 30 samples, 5 of which were duplicates. The maximum concentration of 780 mg/kg was detected in sample LL9SS-068-0001-SO.

4.1.2.3 Pesticides and PCBs

No pesticides or PCBs were reported above the laboratory detection limits in the surface soil samples.

4.1.2.4 Volatiles and Semi-volatiles

Figure 4-3 shows surface soil exceedances for SVOCs and VOCs at LL-9. No VOCs were detected above Region 9 residential soil PRGs. Two SVOCs, benzo(a)pyrene and dibenzo(a,h)anthracene, were detected above the Region 9 residential soil PRG. Benzo(a)pyrene, was detected above the Region 9 residential soil PRG (0.062 mg/kg) in 2 samples, 1 of which was a duplicate not associated with the other exceedance. Benzo(a)pyrene was detected in sample LL9SS-034-0001-DUP at a concentration of 0.073 mg/kg and in sample LL9SS-068-0001-SO at a concentration of 0.240 mg/kg. Dibenzo(a,h)anthracene was detected above the Region 9 residential soil PRG (0.062 mg/kg) in 1 sample at a concentration of 0.130 mg/kg in sample LL9SS-068-0001-SO.

4.1.3 RI Subsurface Soil Samples

Fifty-three (53) subsurface soil samples and 3 duplicates were collected at various locations at LL-9 during the RI. Six of the 53 subsurface soil samples were subfloor subsurface soil samples (initially designated as a subfloor surface soil samples). The samples were re-classified in the field because the subfloor soils in the 0-1 ft interval were not representative of the surface soil conditions because of demolition activities. Therefore, the samples were collected as a subfloor subsurface soil samples from the 1-3 ft interval. Most samples were collected from a depth of 1-3 ft bgs. The depths of subsurface soil samples are listed in Table 4-4 (included at the end of Section 4.0). The samples were submitted for analysis of TAL metals. In addition, 10% of the samples collected were analyzed for explosives, cyanides, VOCs, SVOCs, PCBs, pesticides, and propellants. Subsurface soil sample results are summarized in Table 4-4 (included at the end of Section 4.0) and included in Appendix I.

Aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, iron, lead, manganese, mercury, vanadium, and zinc were detected at concentrations greater than the RVAAP



installation background concentrations, Region 9 residential soil PRGs, or both. In addition, one propellant (nitrocellulose) and one explosive (RDX) were detected in subsurface soils. However, only 10% of the subsurface samples collected were analyzed for explosives and propellants. Therefore, the lateral and vertical extent of explosives, propellants and TAL metals has not been fully delineated during this Phase I RI. The lateral extent of subsurface soil contamination at LL-9 (as determined by this Phase I RI) is shown on Figures 4-12 and 4-13. The distribution of selected site related contaminants (arsenic, chromium, copper, and mercury) identified as subsurface COPCs in Section 6.2.3 of this document, are shown in Figures 4-14 through 4-16. The distribution and exposure pathways for SRCs are further discussed in the conceptual site model included in Section 5.4.

Exceedances for the fifty-three subsurface soil samples, one subfloor subsurface sample and 3 duplicate samples are summarized in Table 4-5 (included at the end of Section 4.0). Specific exceedances for each analytical parameter are discussed in the following subsections.

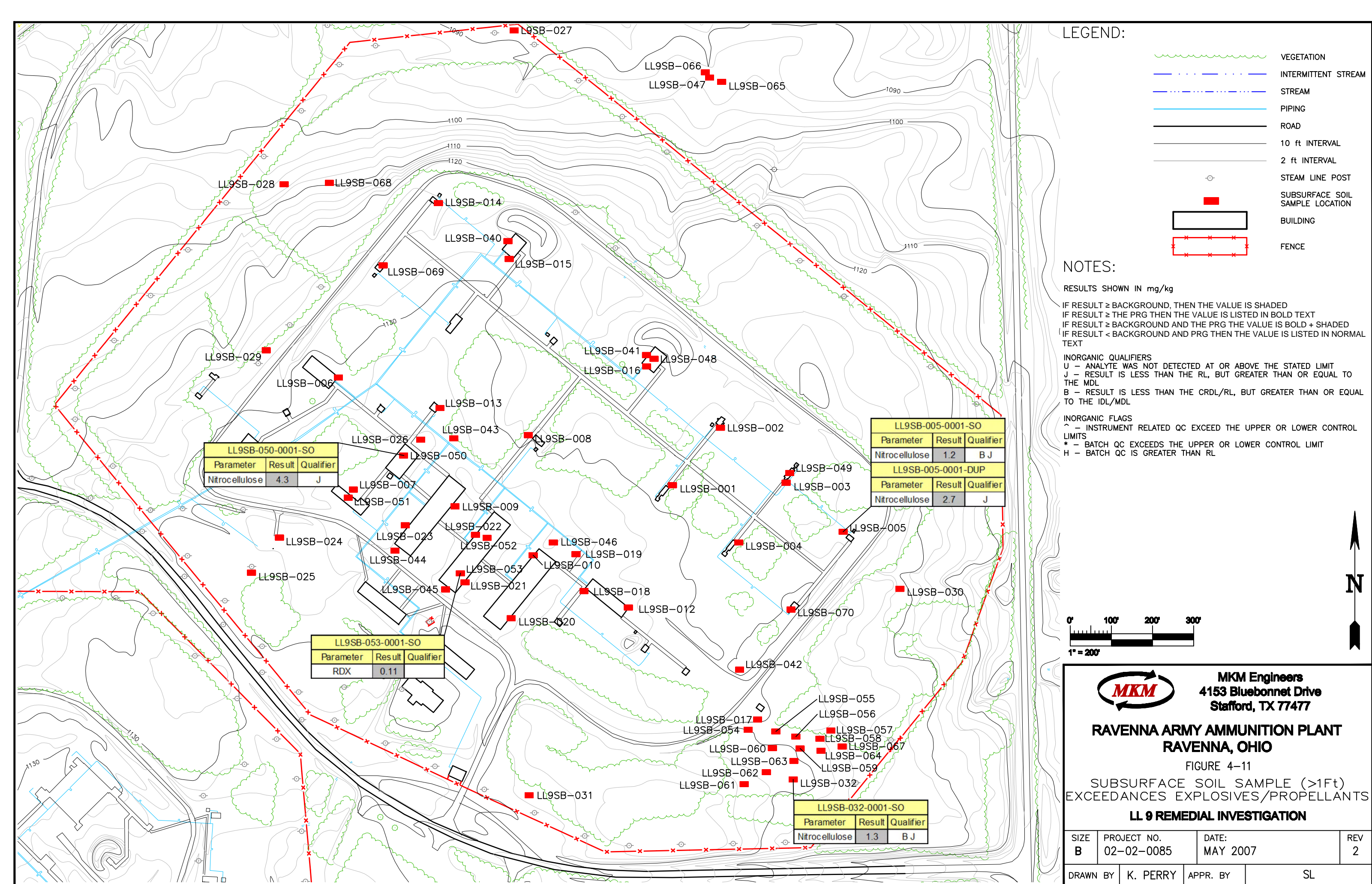
4.1.3.1 Explosives and Propellants

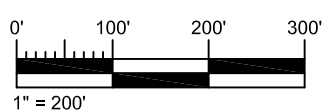
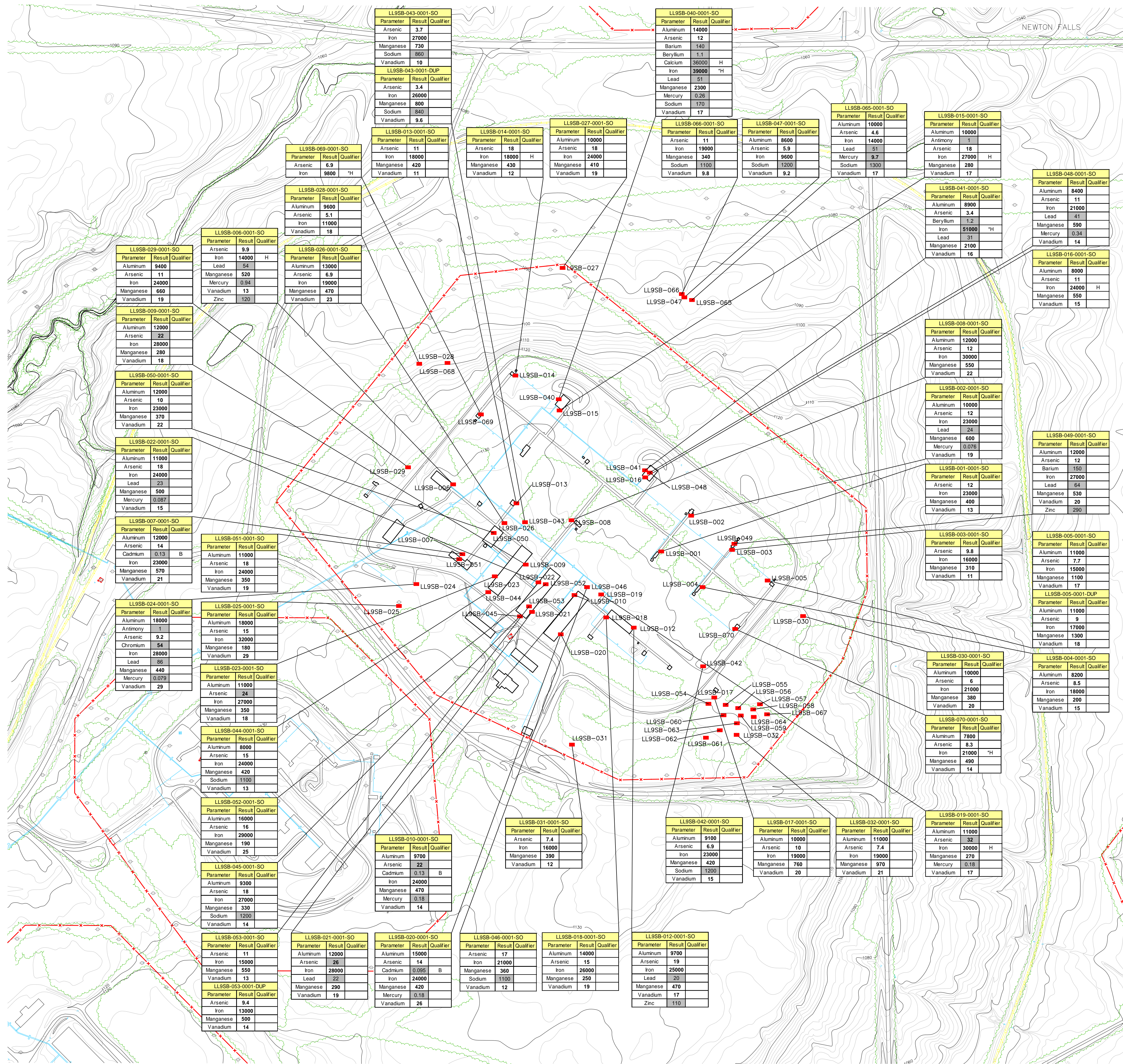
RDX was detected above the RVAAP background criterion (0.0 mg/kg) at a concentration of 0.11 mg/kg in sample, LL9SB-053-0001-SO. Nitrocellulose was detected at concentrations above the background criterion (0.0 mg/kg) in sample LL9SB-005-0001-SO at a concentration of 1.2 (B) (J) mg/kg; sample LL9SB-005-0001-DUP at a concentration of 2.7 (J) mg/kg; sample, LL9SB-032-0001-SO at a concentration of 1.3 (B) (J) mg/kg; and sample LL9SB-050-0001-SO at a concentration of 4.3 (J) mg/kg. However, the results were qualified as (J), or (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J). In addition, one of the detections was a duplicate sample. Based on the qualified analytical results, a determination of vertical extent of explosive and propellant contamination cannot be made. Figure 4-4 shows subsurface soil exceedances for explosives and propellants at LL-9.

4.1.3.2 TAL Metals and Cyanide

Figure 4-5 shows subsurface soil exceedances for TAL metals and cyanide at LL-9. The following metals were detected above comparable RVAAP installation background criteria (USACE, 2001b), Region 9 residential soil PRGs for subsurface soil (>1 ft bgs), or both:

- Aluminum was detected above the Region 9 residential soil PRG (7,614mg/kg) in 39 samples, 1 of which was a duplicate. The maximum concentration of 18,000 mg/kg was detected in sample LL9SB-024-0001-SO and LL9SB-025-0001-SO.
- Antimony was detected above the RVAAP installation background criterion (0.96 mg/kg) in 2 samples. The maximum concentration of 1 mg/kg was detected in samples LL9SB-015-0001-SO and LL9SB-024-0001-SO.

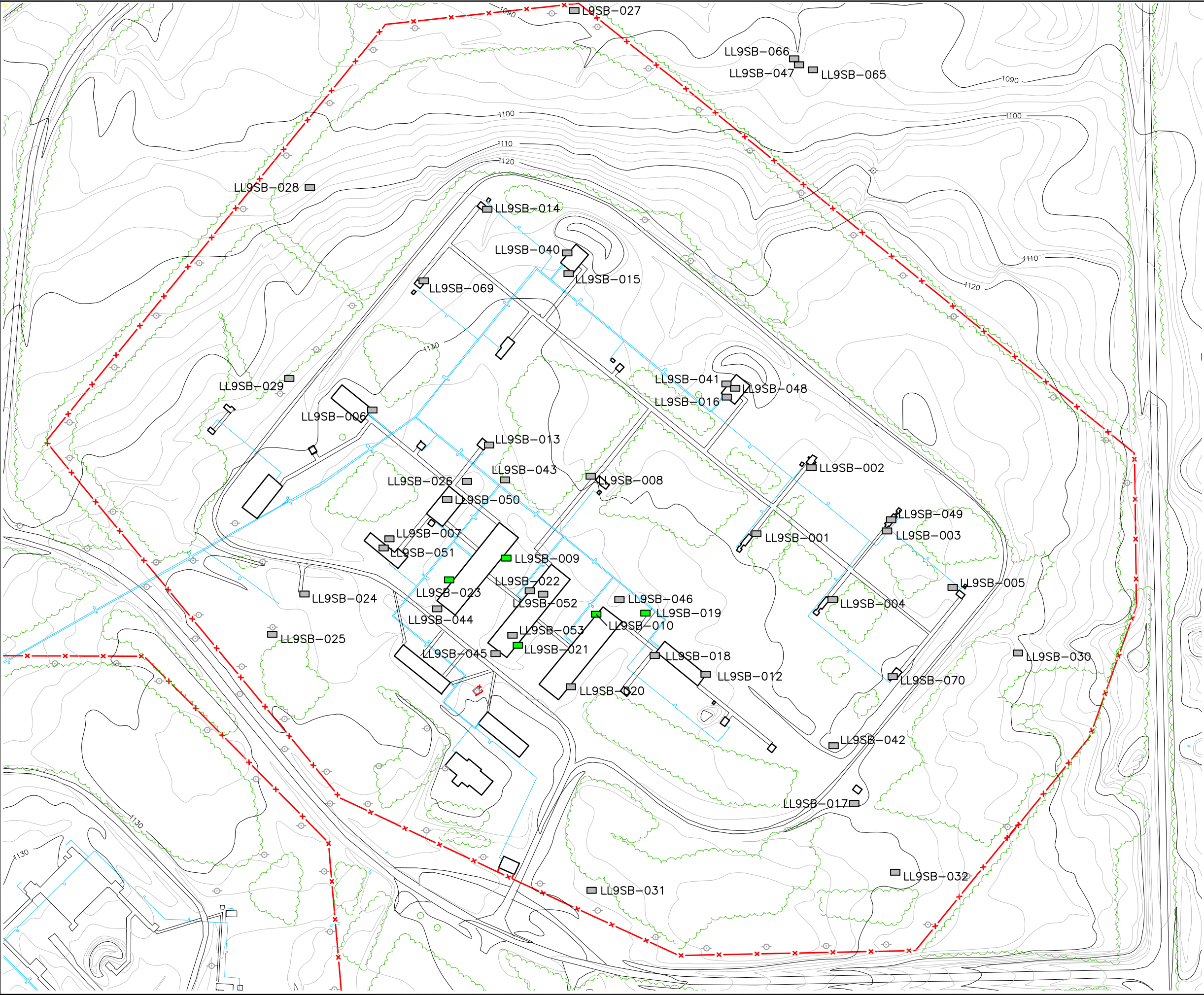




MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO
FIGURE 4-12
SUBSURFACE SOIL SAMPLE (>1ft)
EXCEEDANCES TAL METALS AND CYANIDE
LL 9 REMEDIAL INVESTIGATION

SIZE D	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY R. BOWMAN	APPR. BY	SL	



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION

SURFACE BACKGROUND VALUE FOR ARSENIC = 19.8 mg/Kg


CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND

0' 100' 200' 300'

1" = 200'

N

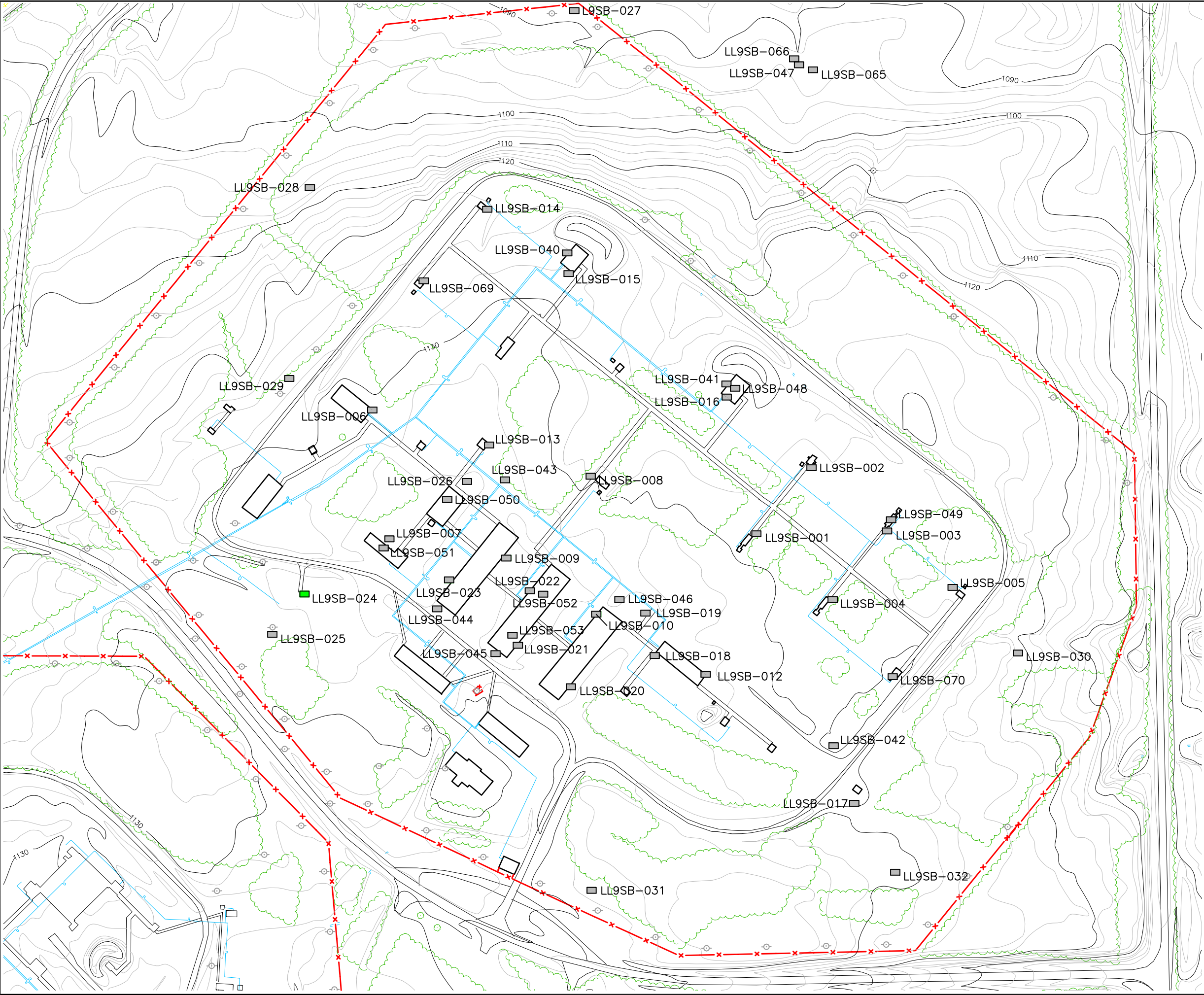


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

**FIGURE 4-13
SUBSURFACE SOIL SAMPLE (>1ft)
DISTRIBUTION OF ARSENIC
LL 9 REMEDIAL INVESTIGATION**

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY K. PERRY	APPR. BY	SL	



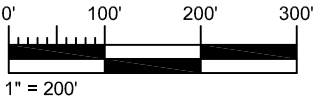
LEGEND:


- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION

SURFACE BACKGROUND VALUE FOR CHROMIUM = 27.2 mg/Kg

CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND



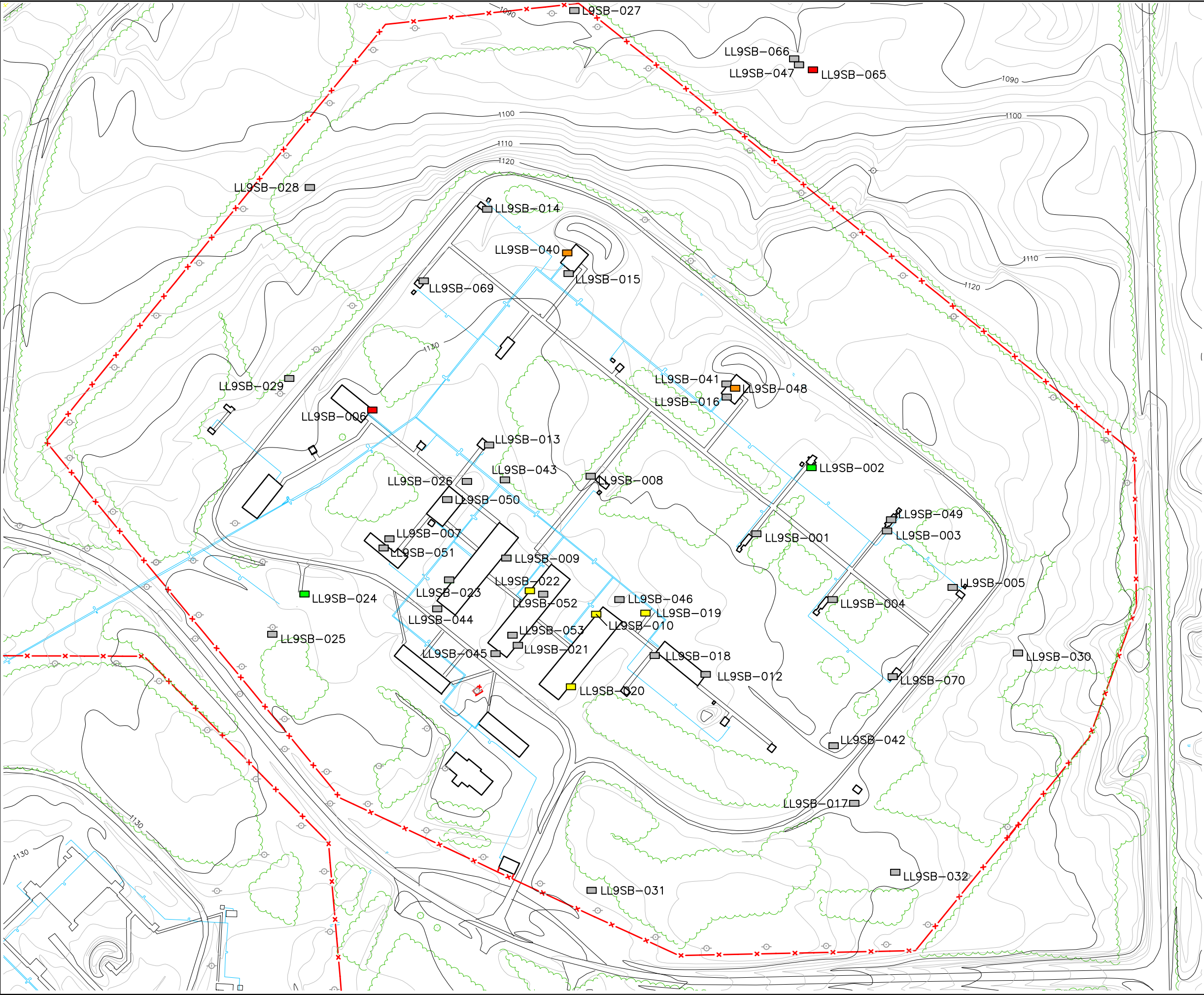


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-14
SUBSURFACE SOIL SAMPLE (>1Ft)
DISTRIBUTION OF CHROMIUM
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- BUILDING
- FENCE
- SURFACE SOIL SAMPLE LOCATION

SURFACE BACKGROUND VALUE FOR MERCURY = 0.04 mg/Kg


CONCENTRATION

- NON-DETECT OR \leq BACKGROUND
- > 1 TO 2 TIMES BACKGROUND
- > 2 TO 5 TIMES BACKGROUND
- > 5 TO 20 TIMES BACKGROUND
- > 20 TIMES BACKGROUND

0' 100' 200' 300'

1" = 200'

N



MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-15
SUBSURFACE SOIL SAMPLE (>1Ft)
DISTRIBUTION OF MERCURY
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



- 1 • Arsenic was detected above the Region 9 residential soil PRG (0.39 mg/kg) in 47
2 samples, 3 of which were duplicates. Arsenic was also detected above the RVAAP
3 installation background criterion (19.8 mg/kg) and Region 9 residential soil PRG in 5
4 sample locations. The maximum concentration of 26 mg/kg was detected in sample
5 LL9SB-021-0001-SO.

- 6 • Barium was detected above the RVAAP installation background criterion (124 mg/kg)
7 in 2 samples. The maximum concentration of 150 mg/kg was detected in sample
8 LL9SB-049-0001-SO.

- 9 • Beryllium was detected above the RVAAP installation background criterion (0.88
10 mg/kg) in 2 samples. The maximum concentration of 1.2 mg/kg was detected in
11 sample LL9SB-041-0001-SO.

- 12 • Cadmium was detected above the RVAAP installation background criterion (0.00
13 mg/kg) in 3 samples. The maximum concentration of 0.13 mg/kg in samples LL9SB-
14 007-0001-SO and LL9SB-010-001-SO.

- 15 • Calcium was detected above the RVAAP installation background criterion (35,500
16 mg/kg) in 1 sample. The concentration of 36,000 mg/kg was detected in sample
17 LL9SB-040-0001-SO.

- 18 • Chromium was detected above the RVAAP installation background criterion (27.2
19 mg/kg) and Region 9 residential soil PRG (30 mg/kg) in 1 sample. The maximum
20 concentration of 54 mg/kg was detected in sample LL9SB-024-0001-SO.

- 21 • Iron was detected above the Region 9 residential soil PRG (2,346 mg/kg) in 48
22 samples, 3 of which were duplicates. Iron was also detected above the RVAAP
23 installation background criterion (35,200 mg/k) and Region 9 residential soil PRG in 2
24 samples. The maximum concentration of 51,000 mg/kg was detected in sample
25 LL9SB-041-0001-SO.

- 26 • Lead was detected above the RVAAP installation background criterion (19.1 mg/kg) in
27 11 samples. The maximum concentration of 86 mg/kg was detected in sample
28 LL9SB-024-0001-SO.

- 29 • Manganese was detected above the Region 9 residential soil PRG (0.39 mg/kg) in 48
30 samples, 3 of which were duplicates. The maximum concentration of 2,300 mg/kg was
31 detected in sample LL9SB-040-0001-SO.

- 32 • Mercury was detected above the RVAAP installation background criterion (0.04 mg/kg)
33 in 9 samples and above the Region 9 residential soil PRG (2.3 mg/kg) and the RVAAP
34 installation background in 1 sample. The maximum concentration of 9.7 mg/kg was
35 detected in sample LL9SB-065-0001-SO.

- 36 • Sodium was detected above the RVAAP installation background criterion (145 mg/kg)
37 in 10 samples, 1 of which was a duplicate. The maximum concentration of 1,300
38 mg/kg was detected in sample LL9SB-065-0001-SO.



- Vanadium was detected above the Region 9 residential soil PRG (7.8 mg/kg) in 51 samples, 3 of which were duplicates. The maximum concentration of 29 mg/kg was detected in samples LL9SB-024-0001-SO and LL9SB-025-0001-SO.
- Zinc was detected above the RVAAP installation background criterion (93.9 mg/kg) in 3 samples. The maximum concentration of 290 mg/kg was detected in sample LL9SB-049-0001-SO.

4.1.3.3 Pesticides and PCBs

No pesticides or PCBs were reported above laboratory detection limits in the subsurface soil samples.

4.1.3.4 Volatiles and Semi-volatiles

No VOCs were reported above laboratory detection limits in subsurface soil samples. Although some SVOCs were detected, no SVOCs exceeded the Region 9 residential soil PRGs in the subsurface soil samples.

4.2 Groundwater

This section summarizes the physical and chemical data collected from the LL-9 groundwater monitoring wells during the RI field effort. Groundwater samples were not collected during the azide screening in March 2002.

4.2.1 Hydraulic Conductivity and Water Level Measurements

In June 2004, slug tests were performed on the 7 newly installed bedrock wells at LL-9. Slug test data are found in Appendix L. As shown in Table 4-6, hydraulic conductivities ranged from 2.11×10^{-4} to 7.92×10^{-4} centimeters/second (cm/s) in the screened materials underlying LL-9. The average hydraulic conductivity at LL-9 is 3.99×10^{-4} cm/s (see Appendix L). Slug tests performed on wells located at other RVAAP areas indicate average hydraulic conductivities that range between $1.45\text{E-}03$ cm/s to $6.09\text{E-}06$ cm/s for unconsolidated soils and bedrock. The hydraulic conductivities measured at LL-9 are consistent with bedrock conductivities measured at other locations in the RVAAP. Groundwater levels were measured on three occasions: December 8, 2003; December 17, 2003; and June 25, 2004 (see Appendix K). Data from those three rounds of well gauging were used to produce potentiometric surface maps for LL-9. These maps are shown on Figures 4-16, 4-17, and 4-18.



Table 4-6

Hydraulic Conductivity and Water Level Measurements

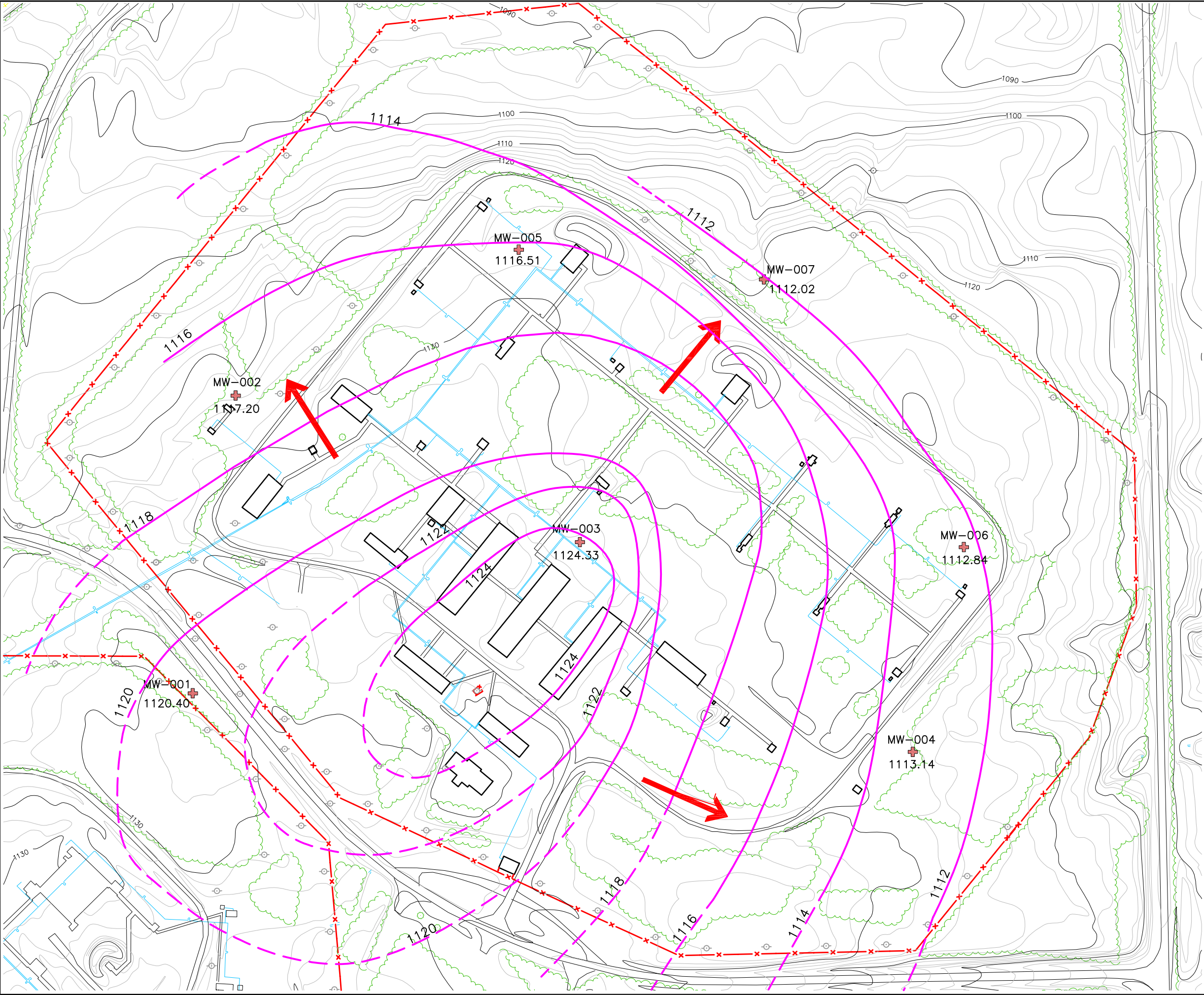
Monitoring Well	Screened Interval Depth (bgs)	Total Depth (ft)	Geologic Material Adjacent to Screen	Hydraulic Conductivity (cm/s)
MW-001	11-21	23.15	Brown sand/sandstone	2.57×10^{-4}
MW-002	10-20	22.40	Reddish brown sandstone	3.21×10^{-4}
MW-003	11.5-21.5	24.03	Weathered red sandstone/brown sandstone	2.90×10^{-4}
MW-004	22-32	34.49	Grey sandstone/siltstone	7.92×10^{-4}
MW-005	10-20	23.32	Brown sandstone	2.11×10^{-4}
MW-006	16-26	28.63	Grey sandstone	2.31×10^{-4}
MW-007	8.5-18.5	17.95	Sandstone	4.61×10^{-4}

4.2.2 Groundwater Results

Groundwater samples were collected from each of the 7 newly installed monitoring wells (MW-001 through MW-007) and analyzed for TAL metals, cyanide, explosives, propellants, VOCs, SVOCs, pesticides and PCBs. The results of the groundwater are summarized in Table 4-7 (included at the end of Section 4.0) and included in Appendix M.

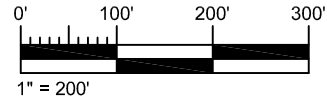
Nitrocellulose and five metals (antimony, cobalt, copper, iron, and manganese) were detected in seven groundwater samples (and 1 duplicate) at concentrations exceeding the RVAAP-specific consolidated bedrock background criteria, the Region 9 tap water PRG or both. However, the nitrocellulose concentrations results were qualified as estimated (J), or (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J).

Exceedances for groundwater at LL-9 are summarized in Table 4-8 (included at the end of Section 4.0). Figures 4-19 and 4-20 show the analytical exceedances above installation consolidated aquifer background criteria (USACE, 2001b) levels (if available), Region 9 tap water PRGs (if available) for explosives/propellants, TAL metals/cyanide, pesticides/PCBs, VOCs, and SVOCs, or both. The results were compared to Region 9 tap water PRGs and consolidated aquifer background values for screening purposes. Purging and field sampling forms are provided in Appendix K; tabulated analytical results and laboratory reports are provided in Appendix M.



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- POTENTIOMETRIC LINE
- POTENTIOMETRIC LINE (inferred)
- STEAM LINE POST
- MONITORING WELL SAMPLE LOCATION
- BUILDING
- FENCE
- GROUNDWATER FLOW DIRECTION

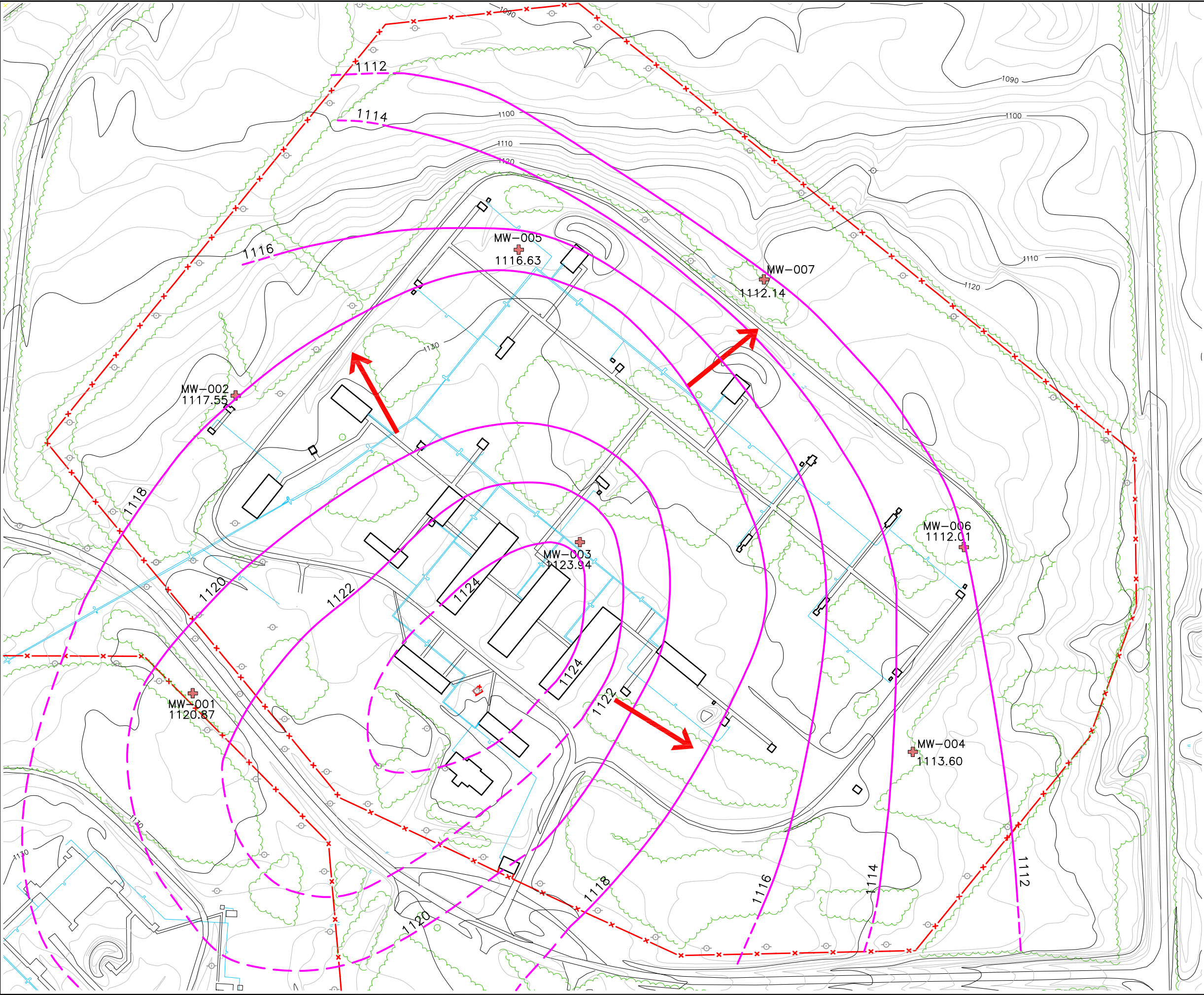


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

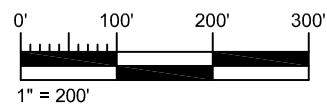
FIGURE 4-16
POTENTIOMETRIC SURFACE MAP
DECEMBER 8, 2003
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	R. BOWMAN	APPR. BY	SL



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- POTENTIOMETRIC LINE
- POTENTIOMETRIC LINE (inferred)
- STEAM LINE POST
- MONITORING WELL SAMPLE LOCATION
- BUILDING
- FENCE
- GROUNDWATER FLOW DIRECTION

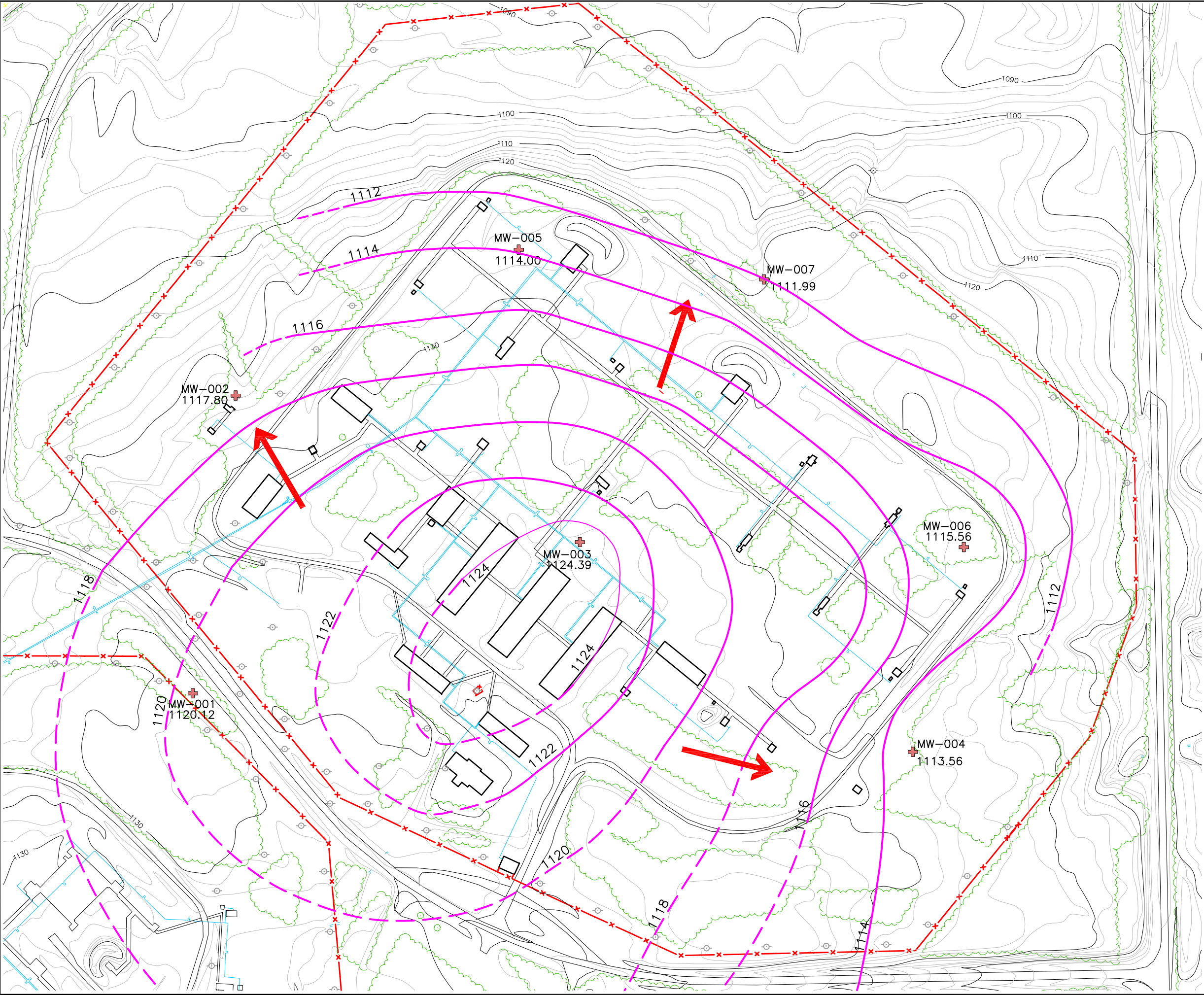


MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

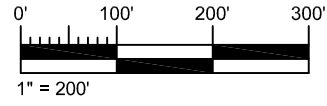
FIGURE 4-17
POTENTIOMETRIC SURFACE MAP
DECEMBER 17, 2003
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY R. BOWMAN	APPR. BY	SL	



LEGEND:

- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- POTENTIOMETRIC LINE
- POTENTIOMETRIC LINE (inferred)
- STEAM LINE POST
- MONITORING WELL SAMPLE LOCATION
- BUILDING
- FENCE
- GROUNDWATER FLOW DIRECTION

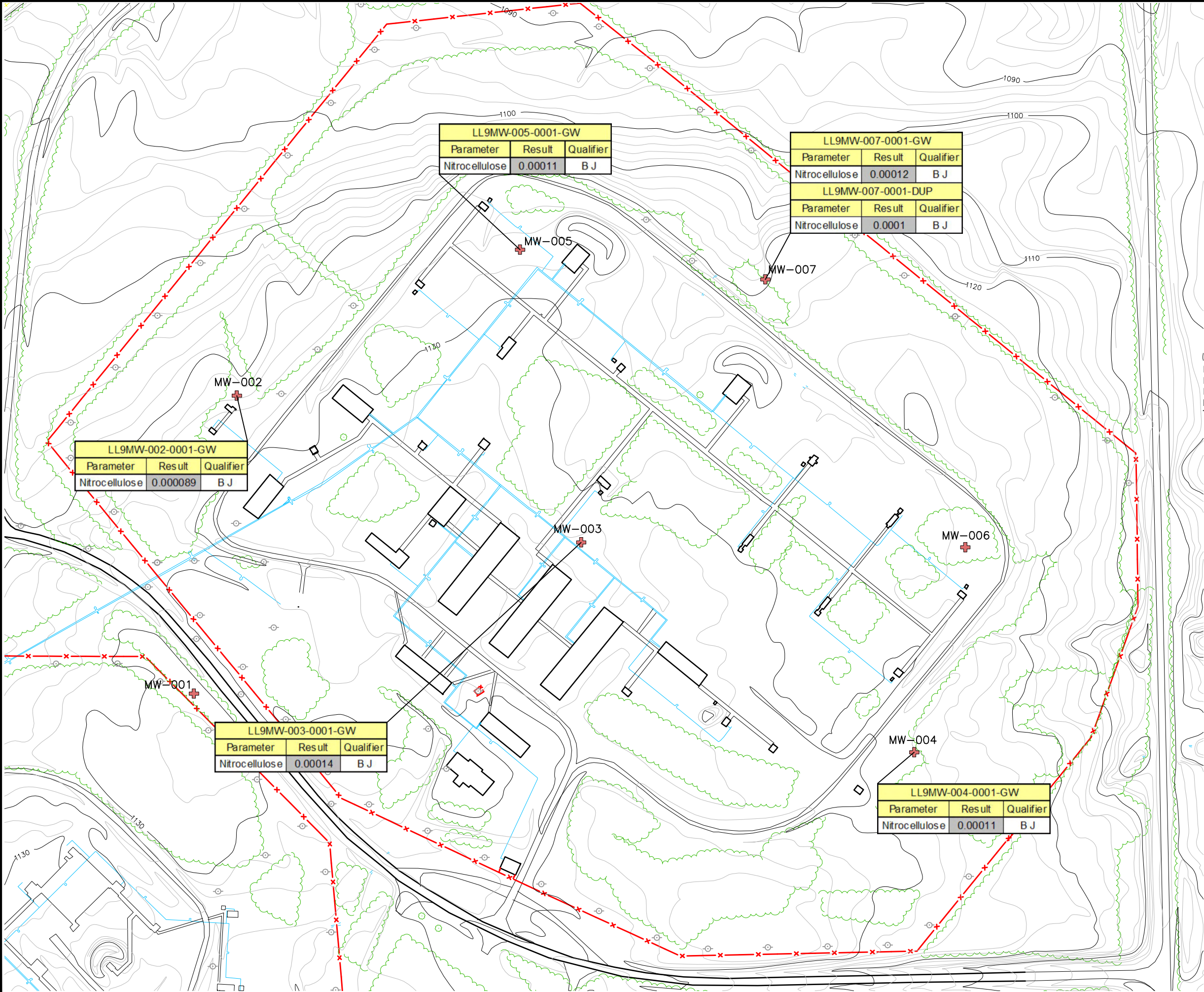




MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO
FIGURE 4-18
POTENTIOMETRIC SURFACE MAP
JUNE 25, 2004
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY R. BOWMAN		APPR. BY	SL



LEGEND:

VEGETATION

INTERMITTENT STREAM

STREAM

PIPING

ROAD

10 ft INTERVAL

2 ft INTERVAL

STEAM LINE POST

MONITORING WELL
SAMPLE LOCATION

BUILDING

FENCE

NOTES:

RESULTS SHOWN IN ug/L

IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED

IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT

IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED

IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

ORGANIC QUALIFIERS

U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT

J - RESULT IS AN ESTIMATED VALUE BELOW THE RL

ORGANIC FLAGS

B - METHOD BLANK: BATCH QC IS GREATER THAN RL

* - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT

^ - BATCH QC IS GREATER THAN RL

A - CONCENTRATION IS BELOW THE METHOD REPORTING LIMIT (RL)

H - ALTERNATE PEAK SELECTION UPON ANALYTICAL REVIEW

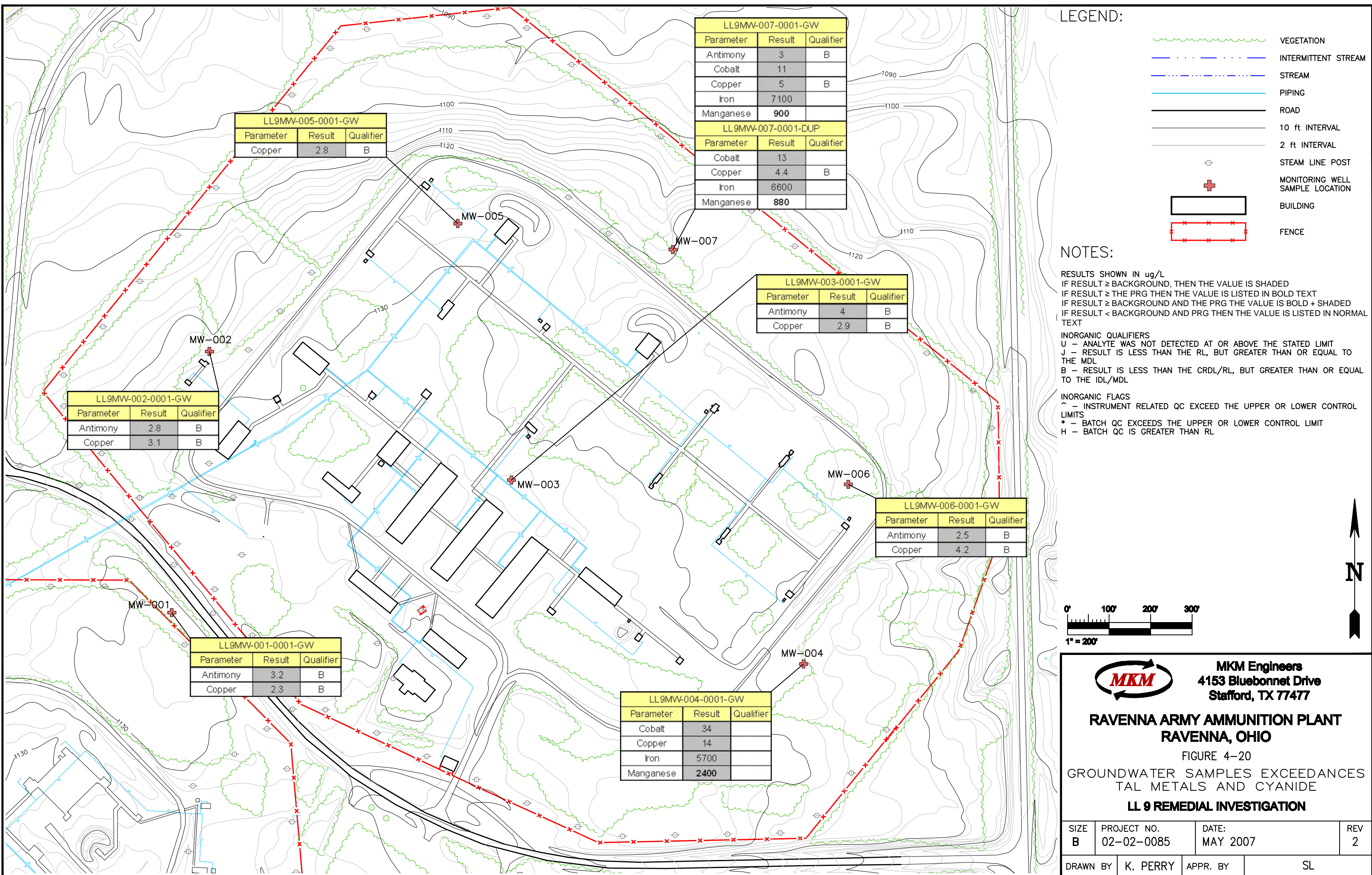
MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-19
GROUNDWATER SAMPLE EXCEEDANCES
EXPLOSIVES/PROPELLANTS

LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: AUG 2007	REV 3
DRAWN BY K. PERRY		APPR. BY	MT





4.2.2.1 Explosives and Propellants

Nitrocellulose was detected above the RVAAP-specific consolidated bedrock background criterion (0.0 mg/kg) in 6 samples, 1 of which was a duplicate. The maximum concentration of 0.00014 (B)(J) $\mu\text{g/L}$ was detected in sample LL9GW-003-0001-GW. Detections refer to individual samples (six samples total), one of which was a duplicate. Figure 4-19 shows groundwater exceedances for explosives and propellants at LL-9.

4.2.2.2 TAL Metals and Cyanide

When the RVAAP-specific consolidated bedrock background criteria for groundwater were established, no background criteria were established for antimony, cobalt or copper. Therefore, the background criteria assigned to these metals, for the purpose of this RI, was zero. Figure 4-20 shows groundwater exceedances for TAL metals and cyanide at LL-9. Thirteen inorganic compounds were detected in the 7 groundwater samples (and 1 duplicate) collected as part of the LL-9 RI. Only 5 of the 13 metals detected exceeded the RVAAP-specific consolidated bedrock background criteria, the Region 9 tap water PRG or both. The exceedences are summarized as follows:

- Antimony was detected above the RVAAP-specific consolidated bedrock background criterion (0.0 $\mu\text{g/L}$) in 5 samples. The maximum concentration of 4 (B) $\mu\text{g/L}$ was detected in sample LL9GW-003-0001-GW.
- Cobalt was detected above the RVAAP-specific consolidated bedrock background criterion (0.0 $\mu\text{g/L}$) in 3 samples, 1 of which was a duplicate. The maximum concentration of 34 $\mu\text{g/L}$ was detected in sample LL9GW-004-0001-GW.
- Copper was detected above the RVAAP-specific consolidated bedrock background criterion (0.0 $\mu\text{g/L}$) in 8 samples, 1 of which was a duplicate. The maximum concentration of 14 $\mu\text{g/L}$ was detected in sample LL9GW-004-0001-GW.
- Iron was detected above the RVAAP-specific consolidated bedrock background criterion (1,430 $\mu\text{g/L}$) in 3 samples, 1 of which was a duplicate. The maximum concentration of 7,100 $\mu\text{g/L}$ was detected in sample LL9GW-007-0001-GW.
- Manganese was detected above the Region 9 tap water PRG (876 $\mu\text{g/L}$) in 2 samples, 1 of which was a duplicate. Manganese was also detected above the RVAAP-specific consolidated bedrock background criterion (1,340 $\mu\text{g/L}$) and the Region 9 tap water PRG in 1 sample. The maximum concentration of 2,400 $\mu\text{g/L}$ was detected in sample LL9GW-004-0001-GW.

4.2.2.3 Pesticides and PCBs

No pesticides or PCBs were reported above laboratory detection limits in the groundwater samples.



4.2.2.4 *Volatiles and Semi-volatiles*

No VOCs or SVOCs were reported above laboratory detection limits in the groundwater samples

4.3 **Surface Water**

This section summarizes the physical and chemical data from the LL-9 surface water collected during the RI field program.

4.3.1 **Azide Screening Results**

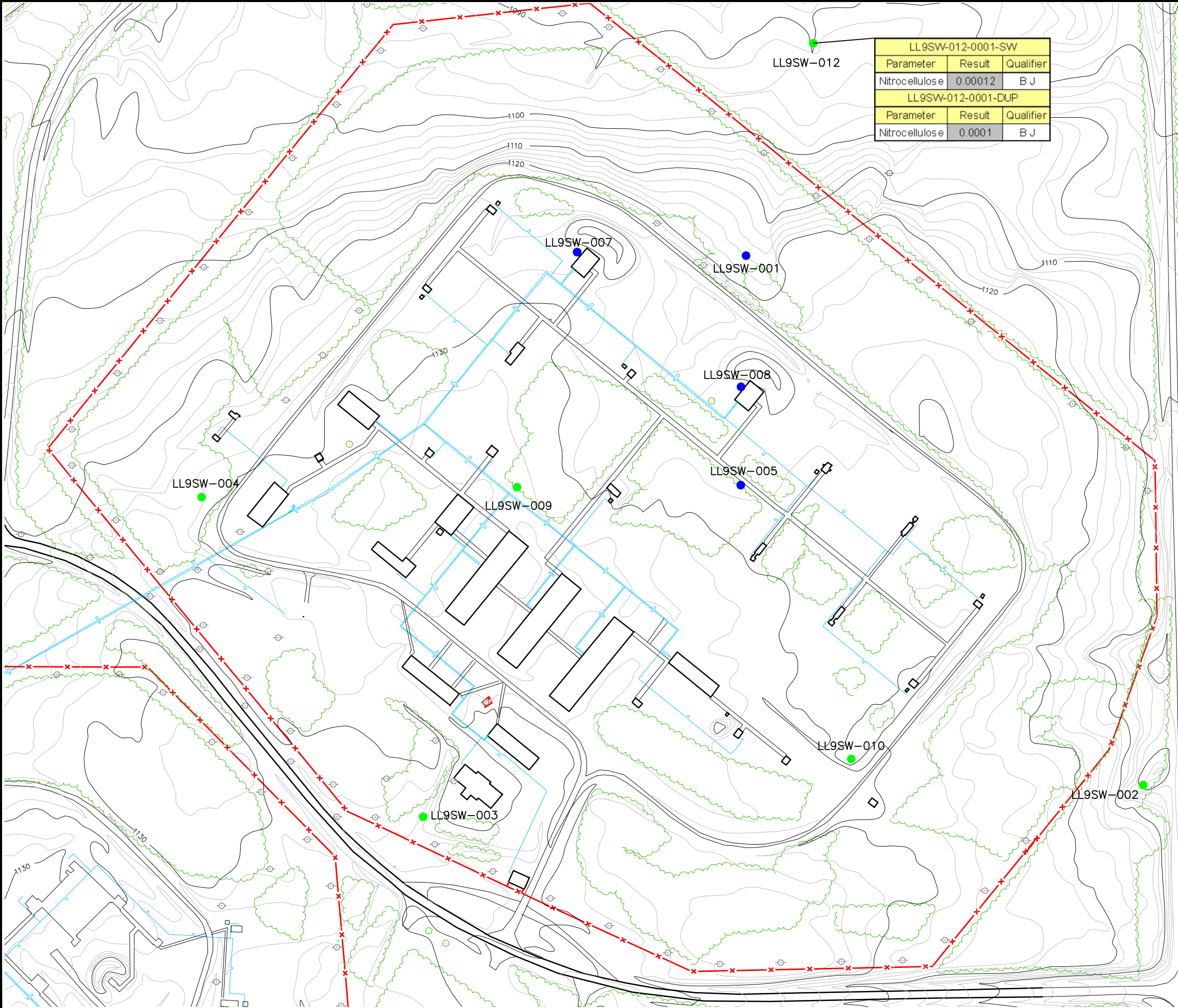
Four surface water samples and one duplicate were collected and analyzed for explosives in accordance with SW846 Method 8330 and TAL metals during the azide screening field program in March 2002. Analytical results are included in Appendix P and summarized in Table 4-9. There were no explosives detected above the reporting limit. Metals exceeded the RVAAP-specific surface water installation background values established by USACE (2000), the Region 9 tap water PRGs, or both in three of the samples and the duplicate; LL9SW-005-0001-SW, LL9SW-007-0001-SW, LL9SW-008-0001-SW, and LL9SW-008-0001-SW (FD). These exceedances are included in the surface water exceedances tabulation at the end of this subsection.

4.3.2 **RI Results**

Four ditch surface water samples, two sewer manhole samples, and one duplicate were collected and analyzed for TAL metals, explosives and cyanide. Figure 4-21 and 4-22 shows the analytical exceedances above RVAAP-specific surface water background values established by USACE (2000) (if available) and/or Region 9 tap water PRGs (if available) for explosives/propellants, TAL metals/cyanide, pesticides/PCBs, VOCs, and SVOCs.

Metals including aluminum, arsenic, barium, beryllium, chromium, cobalt, copper, lead, manganese, mercury, nickel, vanadium, and zinc were detected in surface water at concentration exceeding the RVAAP-specific surface water installation background values established by USACE (2000), the Region 9 tap water PRGs, or both. Nitrocellulose was detected in two samples at concentrations above the method detection limit but below the reporting limit. However, the nitrocellulose concentrations results were qualified as estimated (J), or (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J).

The surface water sampling locations are shown on Figure 3-4. Table 4-10 (included at the end of Section 4.0) lists each surface water sample that exceeded an installation background concentration or Region 9 tap water PRG and the associated analytical result.



LL9SW-012-0001-SW		
Parameter	Result	Qualifier
Nitrocellulose	0.00012	B J
LL9SW-012-0001-DUP		
Parameter	Result	Qualifier
Nitrocellulose	0.0001	B J

LEGEND:

VEGETATION

INTERMITTENT STREAM

STREAM

PIPING

ROAD

10 ft INTERVAL

2 ft INTERVAL

STEAM LINE POST

SURFACE WATER SAMPLING LOCATION

SURFACE WATER SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

BUILDING

FENCE

NOTES:

RESULTS SHOWN IN ug/L

IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED

IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT

IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED

IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

ORGANIC QUALIFIERS

U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT

J - RESULT IS AN ESTIMATED VALUE BELOW THE RL

ORGANIC FLAGS

B - METHOD BLANK: BATCH QC IS GREATER THAN RL

* - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT

^ - BATCH QC IS GREATER THAN RL

A - CONCENTRATION IS BELOW THE METHOD REPORTING LIMIT (RL)

H - ALTERNATE PEAK SELECTION UPON ANALYTICAL REVIEW

1" = 200'

MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

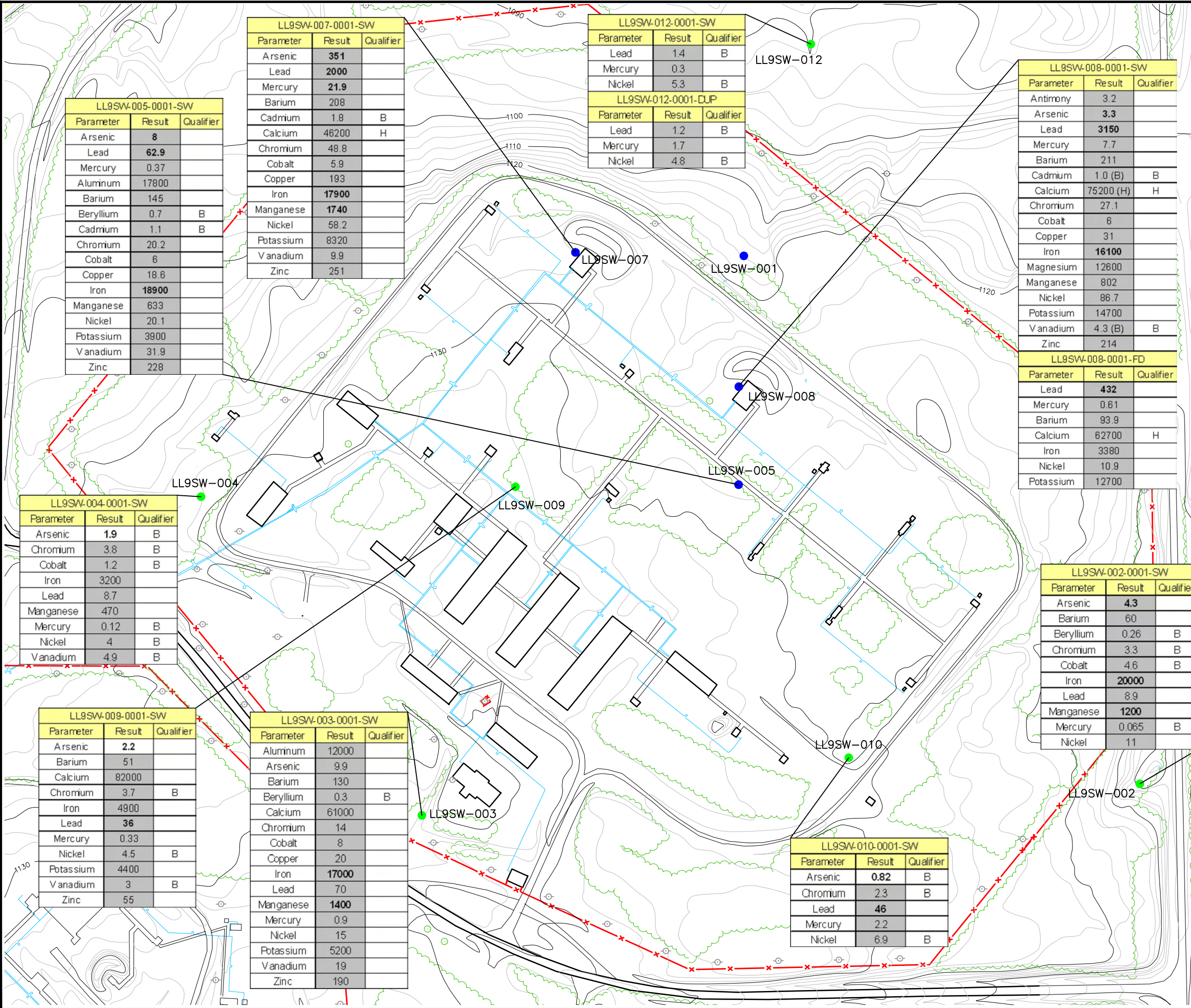
RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-21

SURFACE WATER EXCEEDANCES
EXPLOSIVES/PROPELLANTS

LL 9 REMEDIAL INVESTIGATION

SIZE	PROJECT NO.	DATE:	REV
B	02-02-0085	MAY 2007	2
DRAWN BY	K. PERRY	APPR. BY	SL



LEGEND:

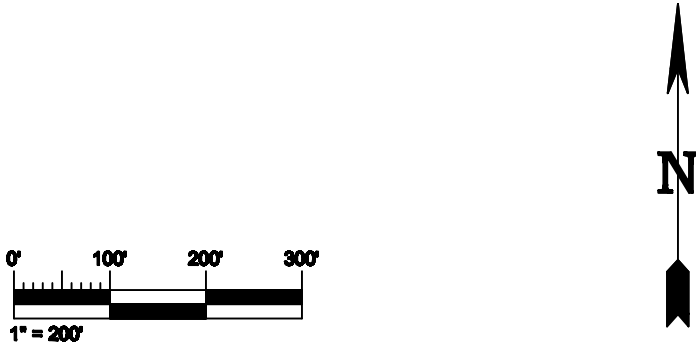
- VEGETATION
- INTERMITTENT STREAM
- STREAM
- PIPING
- ROAD
- 10 ft INTERVAL
- 2 ft INTERVAL
- STEAM LINE POST
- SURFACE WATER SAMPLING LOCATION
- SURFACE WATER SAMPLING LOCATION TAKEN DURING AZIDE SCREENING
- BUILDING
- FENCE


NOTES:

RESULTS SHOWN IN ug/L
IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED
IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT
IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED
IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

INORGANIC QUALIFIERS
U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT
J - RESULT IS LESS THAN THE RL, BUT GREATER THAN OR EQUAL TO THE MDL
B - RESULT IS LESS THAN THE CRDL/RL, BUT GREATER THAN OR EQUAL TO THE IDL/MDL

INORGANIC FLAGS
^ - INSTRUMENT RELATED QC EXCEED THE UPPER OR LOWER CONTROL LIMITS
* - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT
H - BATCH QC IS GREATER THAN RL





MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

**RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO**

FIGURE 4-22
SURFACE WATER EXCEEDANCES
TAL METALS AND CYANIDE
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY K. PERRY		APPR. BY	SL



Surface water sampling forms are found in Appendix O. Analytical results are tabulated and laboratory reports are provided in Appendix P and summarized in Table 4-11 (included at the end of Section 4.0).

4.3.2.1 Explosives and Propellants

No explosives were detected above the method detection limit. Nitrocellulose was detected above the RVAAP installation background criterion (0.0 mg/kg) in 2 samples, 1 of which was a duplicate. Nitrocellulose was detected above the method detection limit but below the reporting limit in sample LL9SW-012-0001-SW at a concentration of 0.00012 (B) (J) µg/L and LL9SW-012-0001-DUP at a concentration of 0.0001 (B) (J) µg/L. However, the nitrocellulose concentrations results were qualified as estimated (J), or (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J). Figure 4-21 shows surface water exceedances for explosives and propellants at LL-9.

4.3.2.2 TAL Metals and Cyanide

When the RVAAP installation background criteria for surface water were established, no background criteria were established for beryllium, chromium, cobalt, lead, mercury, nickel or vanadium; therefore, the background criteria assigned to these metals, for the purpose of this RI, was zero. Figure 4-22 shows groundwater exceedances for TAL metals and cyanide at LL-9. The surface water metals and cyanide samples collected from LL-9 ditches were not filtered, so the results represent total concentrations. The following metals were detected above comparable RVAAP installation background criteria (USACE, 2001b), Region 9 tap water PRGs, or both:

- Aluminum was detected above the RVAAP installation background criterion (3,370 µg/L) in 1 sample. The concentration of 12000 µg/L was detected in sample LL9SW-003-0001-SW.
- Arsenic was detected above the Region 9 tap water PRG (0.045 µg/L) in 3 samples and was detected above RVAAP installation background criterion (3.2 µg/L) and the Region 9 tap water PRG in 2 samples. The maximum concentration of 9.9 µg/L was detected in sample LL9SW-003-0001-SW.
- Barium was detected above the RVAAP installation background criterion (47.5 µg/L) in 3 samples. The maximum concentration of 130 µg/L was detected in sample LL9SW-003-0001-SW.
- Beryllium was detected above the RVAAP installation background criterion (0.0 µg/L) in 2 samples. The maximum concentration of 0.3 µg/L was detected in sample LL9SW-003-0001-SW.
- Calcium was detected above the RVAAP installation background criterion (41,400 µg/L) in 2 samples. The maximum concentration of 82,000µg/L was detected in sample LL9SW-009-0001-SW.



- 1 • Chromium was detected above the RVAAP installation background criterion (0.0 µg/L)
2 in 5 samples. The maximum concentration of 14 µg/L was detected in sample LL9SW-
3 003-0001-SW.
- 4 • Cobalt was detected above the RVAAP installation background criterion (0.0 µg/L) in 3
5 samples. The maximum concentration of 8 µg/L in sample LL9SW-003-0001-SW.
- 6 • Copper was detected above the RVAAP installation background criterion (7.9 µg/L) in
7 1 sample with a concentration of 20 µg/L was detected in sample LL9SW-003-0001-
8 SW.
- 9 • Iron was detected above the RVAAP installation background criterion (2,560 µg/L) in 2
10 samples and was detected above the Region 9 tap water PRG (10,950 µg/L) and
11 RVAAP installation background criterion in 2 samples. The maximum concentration of
12 20,000 µg/L was detected in sample LL9SW-002-0001-SW.
- 13 • Lead was detected above RVAAP installation background criterion (0.0 µg/L) in 4
14 samples, 1 of which was a duplicate. Lead also was detected above the Region 9 tap
15 water PRG (15 µg/L) and the RVAAP installation background criterion in 3 samples.
16 The maximum concentration of 70 µg/L was detected in sample LL9SW-003-0001-
17 SW.
- 18 • Manganese was detected above the RVAAP installation background criterion (391
19 µg/L) in 1 sample and was detected above the Region 9 tap water PRG (876 µg/L) and
20 the RVAAP installation background criterion in 2 samples. The maximum
21 concentration of 1,400 µg/L was detected in sample LL9SW-003-0001-SW.
- 22 • Mercury was detected above the RVAAP installation background criterion (0.0 µg/L) in
23 7 samples, 1 of which was a duplicate. The maximum concentration of 2.2 µg/L was
24 detected in sample LL9SW-003-0001-SW.
- 25 • Nickel was detected above the RVAAP installation background criterion (0.0 µg/L) in 7
26 samples, 1 of which was a duplicate. The maximum concentration of 15 µg/L was
27 detected in sample LL9SW-003-0001-SW.
- 28 • Potassium was detected above the RVAAP installation background criterion (3,170
29 µg/L) in 2 samples. The maximum concentration of 5,200 µg/L was detected in
30 sample LL9SW-003-0001-SW.
- 31 • Vanadium was detected above the RVAAP installation background criterion (0.0 µg/L)
32 in 3 samples. The maximum concentration of 19 µg/L was detected in sample
33 LL9SW-003-0001-SW.
- 34 • Zinc was detected above the RVAAP installation background criterion (42 µg/L) in 2
35 samples. The maximum concentration of 190 µg/L was detected in sample LL9SW-
36 003-0001-SW.



4.3.2.3 Pesticides and PCBs

No pesticides or PCBs were reported above laboratory detection limits in the surface water samples.

4.3.2.4 Volatiles and Semi-volatiles

No VOCs or SVOCs were reported above laboratory detection limits in the surface water samples.

4.4 Sediment

This section summarizes the physical and chemical data from the LL-9 sediment collected during the RI field program.

4.4.1 Azide Screening Results

Five sediment samples were collected and analyzed for explosives in accordance with SW846 Method 8330, TAL metals and lead azide during the azide screening field program in March 2002. Analytical results are included in Appendix R. There no explosives were detected above the method detection limits in any of the samples. Metals exceeded the RVAAP-specific sediment background values established by USACE (2000), Region 9 residential soil PRGs, or both, in the 4 samples analyzed for metals: LL9SD001-001-SD, LL9SD-003-0001-SD, LL9SD-005-0001-SD, and W-24S. These exceedances are included in the sediment exceedances tabulation at the end of this subsection.

4.4.2 RI Results

Twelve sediment samples (and 2 duplicates) were collected from LL-9; 11 from the drainage ways and 1 sewer sediment sample. The samples were analyzed for TAL metals, TOC and grain size. Ten percent of the sediment samples were analyzed for PCBs, pesticides, VOCs and SVOCs. No sump sediment samples were collected during the RI sampling effort. Section 3.3, Deviations from the Work Plan, lists the locations where samples were not collected. The analytical results for the sediment samples that exceeded RVAAP installation background or Region 9 residential soil PRG values are summarized in Table 4-12 (included at the end of Section 4.0). These results include the 5 sediment samples collected and analyzed for explosives, TAL metals and lead azide during the azide screening field program in March 2002.

Propellants (nitrocellulose and nitroguanidine), SVOCs (benzo(a)pyrene, benzo(b)fluoranthene, dibenzo(a,h)anthracene, indeno(1,2,3-cd)pyrene) and metals (aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, cobalt, copper, lead, manganese, mercury, nickel, selenium and vanadium) were detected in sediment samples at concentrations exceeding the RVAAP-specific sediment background values established by



USACE (2000), Region 9 residential soil PRGs, or both. Therefore, the lateral and vertical extent of propellants, SVOCs, and metals were not fully delineated during this Phase I RI. Figures 4-23 through 4-25 show the analytical exceedances above installation background criteria (USACE, 2001b) levels (if available) and/or Region 9 residential soil PRGs (if available) for explosives/propellants, TAL metals/cyanide, pesticides/PCBs, VOCs, and SVOCs. Figures 4-23 through 4-25 also show the lateral extent, as determined by this Phase I RI, of sediment contamination at LL-9. Summary analytical reports are provided in Appendix R and summarized in Table 4-13 (included at the end of Section 4.0). The distribution and exposure pathways for SRCs are further discussed in the conceptual site model included in Section 5.4.

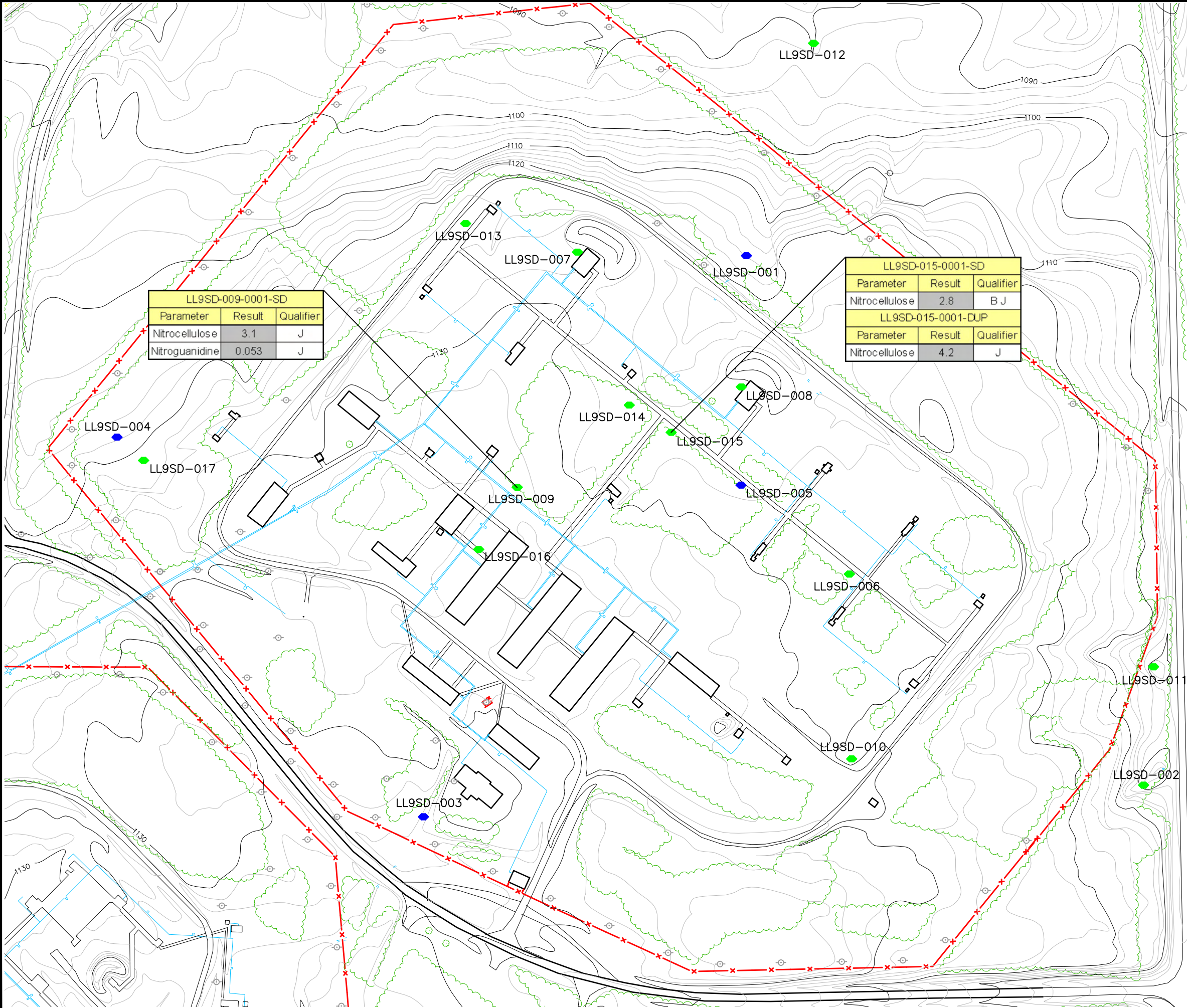
4.4.2.1 Explosives and Propellants

Nitrocellulose was detected above the RVAAP installation background criterion in 3 samples, 1 of which was a duplicate. Nitrocellulose was detected in sample LL9SD-009-0001-SD at a concentration of 3.1 (J) mg/kg; LL9SD-015-0001-SD at a concentration of 2.8 (B) (J) mg/kg; and LL9SD-015-0001-DUP at a concentration of 4.2 (J) mg/kg. Nitroguanidine was detected above the RVAAP installation background criterion in 1 sample at a concentration of 0.053 (J) mg/kg in sample LL9SD-009-0001-SD. Based on the qualified analytical results, a determination of vertical extent of propellants cannot be made. Figure 4-23 shows sediment exceedances for explosives and propellants at LL-9.

4.4.2.2 TAL Metals and Cyanide

When the RVAAP installation background criteria for surface water were established, no background criteria were established for antimony or cadmium; therefore, the background criteria assigned to these metals, for the purpose of this RI, was zero. Figure 4-24 shows sediment exceedances for TAL metals and cyanide at LL-9. The following metals were detected above comparable RVAAP installation background criteria (USACE, 2001b) and/or Region 9 residential soil PRGs:

- Aluminum was detected above the Region 9 residential soil PRG (7,614 mg/kg) in 11 samples, 2 of which were duplicates. Aluminum was also detected above the RVAAP installation background criterion (13,900 mg/kg) and the Region 9 residential soil PRG in 1 sample. The maximum concentration of 14,000 mg/kg was detected in sample LL9SD-004-0001-SD.
- Antimony was detected above the installation background criterion (0.0 mg/kg) in 1 sample. The concentration of 0.81 mg/kg was detected in sample LL9SD-002-0001-SD.



LL9SD-009-0001-SD		
Parameter	Result	Qualifier
Nitrocellulose	3.1	J
Nitroguanidine	0.053	J

LL9SD-015-0001-SD		
Parameter	Result	Qualifier
Nitrocellulose	2.8	B J
LL9SD-015-0001-DUP		
Parameter	Result	Qualifier
Nitrocellulose	4.2	J

LEGEND:

VEGETATION

INTERMITTENT STREAM

STREAM

PIPING

ROAD

10 ft INTERVAL

2 ft INTERVAL

STEAM LINE POST

SEDIMENT SAMPLING LOCATION

SEDIMENT SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

BUILDING

FENCE

NOTES:

RESULTS SHOWN IN mg/kg

IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED

IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT

IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED

IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

ORGANIC QUALIFIERS

U - ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT

J - RESULT IS AN ESTIMATED VALUE BELOW THE RL

ORGANIC FLAGS

B - METHOD BLANK: BATCH QC IS GREATER THAN RL

* - BATCH QC EXCEEDS THE UPPER OR LOWER CONTROL LIMIT

^ - BATCH QC IS GREATER THAN RL

A - CONCENTRATION IS BELOW THE METHOD REPORTING LIMIT (RL)

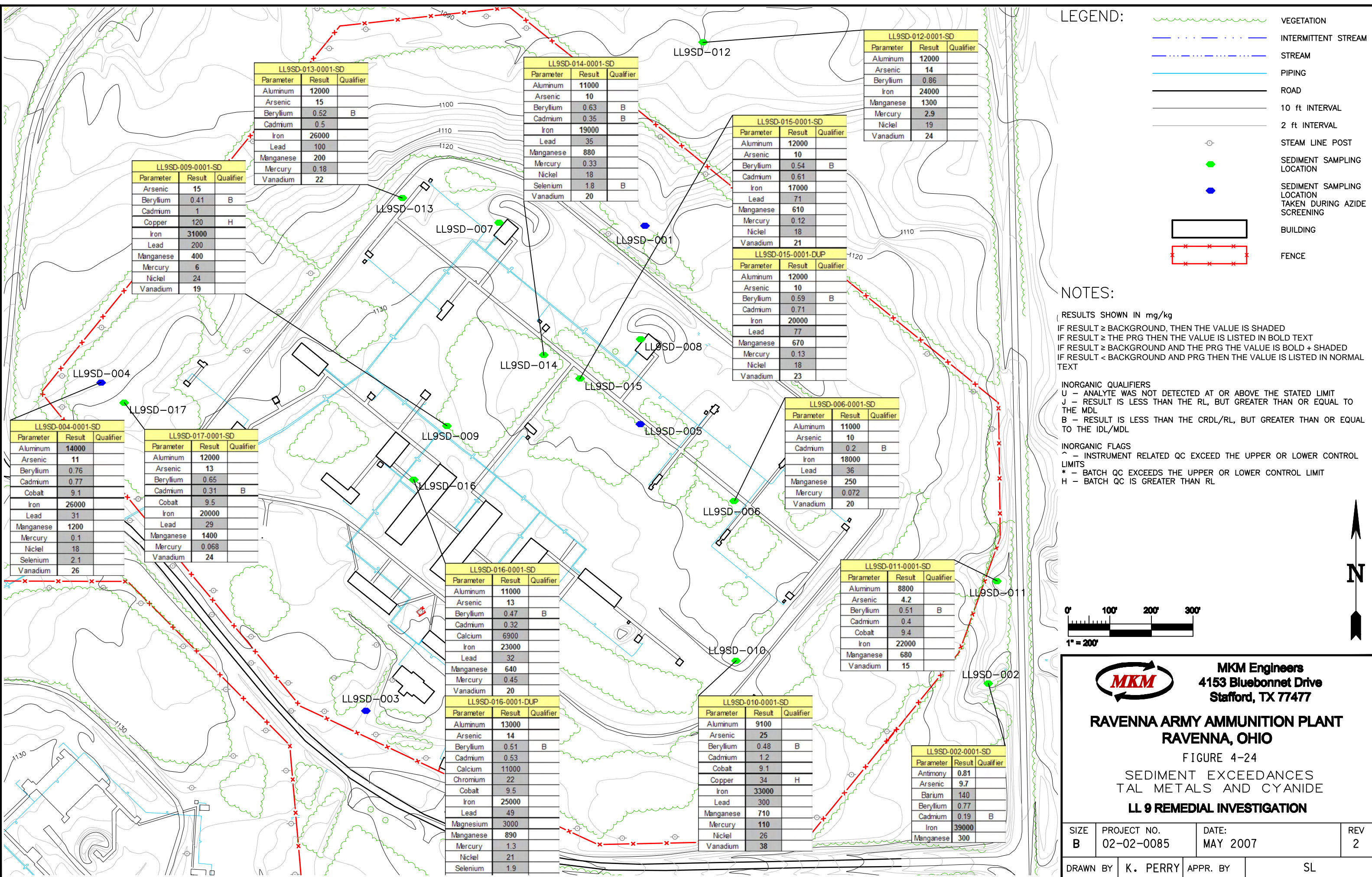
H - ALTERNATE PEAK SELECTION UPON ANALYTICAL REVIEW

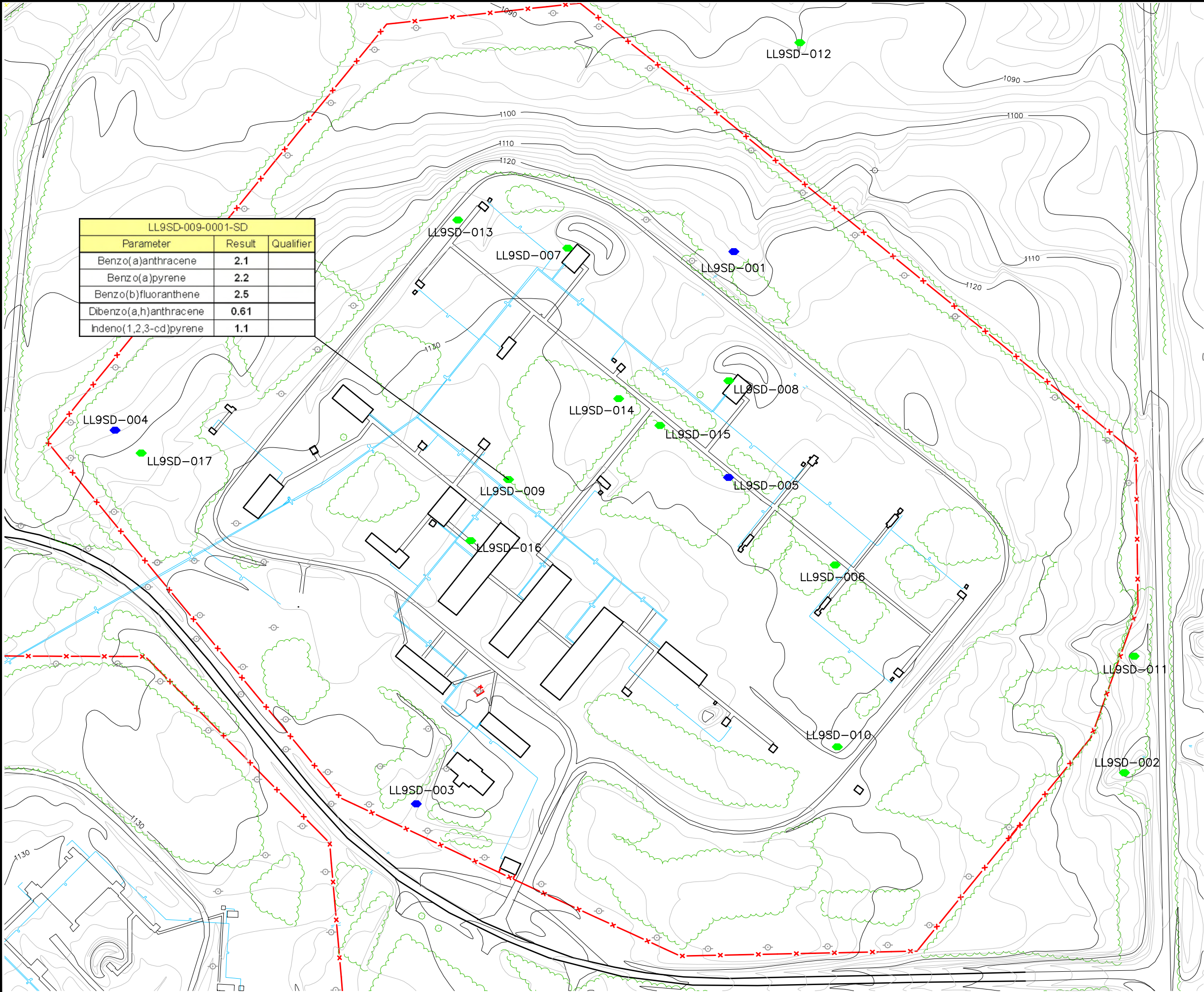
MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-23
SEDIMENT EXCEEDANCES
EXPLOSIVES/PROPELLANTS
LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL





LL9SD-009-0001-SD		
Parameter	Result	Qualifier
Benzo(a)anthracene	2.1	
Benzo(a)pyrene	2.2	
Benzo(b)fluoranthene	2.5	
Dibenzo(a,h)anthracene	0.61	
Indeno(1,2,3-cd)pyrene	1.1	

LEGEND:

VEGETATION

INTERMITTENT STREAM

STREAM

PIPING

ROAD

10 ft INTERVAL

2 ft INTERVAL

STEAM LINE POST

SEDIMENT SAMPLING LOCATION

SEDIMENT SAMPLING LOCATION TAKEN DURING AZIDE SCREENING

BUILDING

FENCE

NOTES:

RESULTS SHOWN IN mg/kg

IF RESULT ≥ BACKGROUND, THEN THE VALUE IS SHADED

IF RESULT ≥ THE PRG THEN THE VALUE IS LISTED IN BOLD TEXT

IF RESULT ≥ BACKGROUND AND THE PRG THE VALUE IS BOLD + SHADED

IF RESULT < BACKGROUND AND PRG THEN THE VALUE IS LISTED IN NORMAL TEXT

INORGANIC QUALIFIERS

U – ANALYTE WAS NOT DETECTED AT OR ABOVE THE STATED LIMIT

J – RESULT IS LESS THAN THE RL, BUT GREATER THAN OR EQUAL TO THE MDL

B – RESULT IS LESS THAN THE CRDL/RL, BUT GREATER THAN OR EQUAL TO THE IDL/MDL

MKM Engineers
4153 Bluebonnet Drive
Stafford, TX 77477

RAVENNA ARMY AMMUNITION PLANT
RAVENNA, OHIO

FIGURE 4-25
SEDIMENT EXCEEDANCES
VOCs AND SVOCs

LL 9 REMEDIAL INVESTIGATION

SIZE B	PROJECT NO. 02-02-0085	DATE: MAY 2007	REV 2
DRAWN BY	K. PERRY	APPR. BY	SL



- Arsenic was detected above the Region 9 residential soil PRG (0.39 mg/kg) in 13 samples, 2 of which were duplicates. Arsenic was also detected above the RVAAP installation background criterion (19.5 mg/kg) and the Region 9 residential soil PRG in 1 sample. The maximum concentration of 25 mg/kg was detected in sample LL9SD-010-0001-SD.
- Barium was detected above the installation background criterion (123 mg/kg) in 1 sample with a concentration of 140 mg/kg in sample LL9SD-002-0001-SD.
- Beryllium was detected above the RVAAP installation background criterion (0.38 mg/kg) in 13 samples, 2 of which were duplicates. The maximum concentration of 0.86 mg/kg was detected in sample LL9SD-012-0001-SD.
- Cadmium was detected above the RVAAP installation background criterion (0.0 mg/kg) in 13 samples, 2 of which were duplicates. The maximum concentration of 0.77 mg/kg was detected in sample LL9SD-004-0001-SD.
- Calcium was detected above the RVAAP installation background criterion (5,510 mg/kg) in 2 samples, 1 of which was a duplicate. The maximum concentration of 11,000 mg/kg was detected in sample LL9SD-016-0001-DUP.
- Chromium was detected above the RVAAP installation background criterion (18.1 mg/kg) in 1 sample which was a duplicate. The maximum concentration of 22 mg/kg was detected in sample LL9SD-016-0001-DUP. Chromium was detected in sample LL9SD-016-0001-SD but the concentration was below the RVAAP installation background criterion with a concentration of 16 mg/kg.
- Cobalt was detected above the RVAAP installation background criterion (9.1 mg/kg) in 5 samples, 1 of which was a duplicate. The maximum concentration of 9.5 mg/kg was detected in sample LL9SD-016-0001-DUP and LL9SD-017-0001-SD. Sample LL9SD-016-0001-SD was detected but was below the RVAAP installation background criterion with a concentration of 8.2 mg/kg.
- Copper was detected above the RVAAP installation background criterion (27.6 mg/kg) in 2 samples. The maximum concentration of 120 mg/kg was detected in sample LL9SD-009-0001-SD.
- Iron was detected above the Region 9 residential soil PRG (2,346 mg/kg) in 11 samples, 2 of which were duplicates. Iron was also detected above the RVAAP installation background criterion (28,200 mg/kg) and the Region 9 residential soil PRG in 3 samples. The maximum concentration of 39,000 mg/kg was detected in sample LL9SD-002-0001-SD.
- Lead was detected above the RVAAP installation background criterion (27.4 mg/kg) in 11 samples, 2 of which were duplicates. The maximum concentration of 0.86 mg/kg was detected in sample LL9SD-012-0001-SD.
- Magnesium was detected above the RVAAP installation background criterion (2,760 mg/kg) in 1 sample which was a duplicate. The maximum concentration of 3,000 mg/kg was detected in sample LL9SD-016-0001-DUP. Sample LL9SD-016-0001-SD was detected but was below the RVAAP installation background criterion with a concentration of 2,700 mg/kg.



- Manganese was detected above the Region 9 residential soil PRG (176 mg/kg) in 14 samples, 2 of which were duplicates. The maximum concentration of 1400 mg/kg was detected in sample LL9SD-017-0001-SD.
- Mercury was detected above the RVAAP installation background criterion (0.06 mg/kg) in 9 samples, 2 of which were duplicates. Mercury was also detected above the Region 9 residential soil PRG (2.3 mg/kg) and the RVAAP installation background criterion in 3 samples. The maximum concentration of 110 mg/kg was detected in sample LL9SD-010-0001-SD.
- Nickel was detected above the RVAAP installation background criterion (17.7 mg/kg) in 8 samples, 2 of which were duplicates. The maximum concentration of 26 mg/kg was detected in sample LL9SD-010-0001-SD.
- Selenium was detected above the RVAAP installation background criterion (1.7 mg/kg) in 3 samples, 1 of which was a duplicate. The maximum concentration of 2.1 mg/kg was detected in sample LL9SD-004-0001-SD.
- Vanadium was detected above the Region 9 residential soil PRG (7.8 mg/kg) in 12 samples, 2 of which were duplicates. Vanadium was also detected above the RVAAP installation background criterion (26.1 mg/kg) in 1 sample. The maximum concentration of 38 mg/kg was detected in sample LL9SD-010-0001-SD.

4.4.2.3 Pesticides and PCBs

No pesticides or PCBs were reported above method detection limits in the subsurface soil samples.

4.4.2.4 Volatiles and Semi-volatiles

Figure 4-25 shows sediment exceedances for SVOCs at LL-9. No VOCs were reported above method detection limits in subsurface soil samples. Twenty-two SVOCs were reported above method detection limits in the sediment samples, however only 5 of the 22 SVOCs exceeded the Region 9 residential soil PRG. The following SVOCs were detected above the Region 9 residential soil PRGs:

- Benzo(a)anthracene was detected above the Region 9 PRG (0.62 mg/kg) in 1 sample. The concentration of 2.1 mg/kg was detected in sample LL9SD-009-0001-SD.
- Benzo(a)pyrene was detected above the Region 9 PRG (0.62 mg/kg) in 1 sample. The concentration of 2.5 mg/kg was detected in sample LL9SD-009-0001-SD.
- Benzo(b)fluoranthene was detected above the Region 9 PRG (0.062 mg/kg) in 1 sample. The concentration of 2.2 mg/kg was detected in sample LL9SD-009-0001-SD.
- Dibenzo(a,h)anthracene was detected above the Region 9 PRG (0.062 mg/kg) in 1 sample. The concentration of 0.61 mg/kg was detected in sample LL9SD-009-0001-SD.



- Indeno(1,2,3-cd)pyrene was detected above the Region 9 PRG (0.62 mg/kg) in 1 sample. The concentration of 1.1 mg/kg was detected in sample LL9SD-009-0001-SD.

4.4.2.5 TOC and Grain Size Analyses

TOC was measured in the sediment samples, with a maximum concentration of 24,000 mg/kg (LL9SD-013-0001-SD) and a minimum concentration of 3,500 mg/kg (LL9SD-012-0001SD).

Samples were also analyzed for grain size in accordance with ASTM D-422-63. Results for grain size analyses are included in Appendix S.

4.5 Other Analyses

This section summarizes the physical and chemical data from the LL-9 sumps, sewers, subfloor and geotechnical samples collected during the RI field program.

4.5.1 Sump and Sewer Manhole Water and Residue

LL-9 sump and sewer manholes sampling results are included and discussed with surface water (Section 4.3) and sediment (Section 4.4) sampling results.

4.5.2 Subfloor

Subsurface soil samples LL9SB-048-0001-SO, LL9SB-049-0001-SO, LL9SB-050-0001-SO, LL9SB-051-0001-SO, LL9SB-052-0001-SO, and LL9SB-053-0001-SO were collected from the subfloor locations of selected buildings at LL-9. LL-9 subfloor sampling results are included and discussed with subsurface soil (Section 4.1.3) sampling results.

4.5.3 Geotechnical Samples

Geotechnical samples were obtained from some subsurface and sediment sampling locations. Two geotechnical samples were obtained from the borings that were advanced to install monitoring wells (MW-001 [sample LL9SB-033-0001-SO], and MW-003 [sample LL9SB-035-0001-SO]). Geotechnical samples were also obtained from 12 sampling and 2 duplicate sampling locations. Results obtained included grain size analysis (from all samples) and hydraulic conductivity (from the boring samples). Geotechnical sampling results are included in Appendix S.

Grain size analysis from the Shelby tube samples included 19.8% to 40% clay, 38% to 40% silt, and 18% to 30% sand. Average hydraulic conductivities ranged from $3.4\text{E-}08$ to $1.1\text{E-}06$. The conductivity results for LL9SB-033-0001-SO/MW-001 ($3.4\text{E-}08$) match the visual description of mostly lean clay. The conductivity results for LL9SB-035-0001-SO/MW-003 match the anticipated values based upon a visual description of sandy silty clay. Since these samples represent the range of visually described soil samples at the AOC, the hydraulic conductivity results can be used as a reference to similarly described soils.



Table 4-1
Summary of Azide Screening Surface Soil Results (0-1 ft)

ANALYTE, UNITS, METHOD NO.	Ravenna Soil Background Criteria (0-1 ft) mg/kg	Region 9 PRG Data (Residential Soil)	D-14S	LL9SS-001-0001-SO	LL9SS-003-0001-SO	LL9SS-005-0001-SO	LL9SS-007-0001-SO	LL9SS-009-0001-SO	LL9SS-011-0001-SO
Sample Date		ppm	3/15/2002	3/11/2002	3/11/2002	3/11/2002	3/11/2002	3/11/2002	3/11/2002
Explosives 8330 mg/kg									
HMX	0.0	306 nc	200 (J) (a)	0.250 U	0.250 U	0.250 U	0.250 U	0.250 U	0.250 U
RDX	0.0	4.4 ca	BRL	0.100 U	0.100 U	0.100 U	0.100 U	0.099 U	0.098 U
1,3,5-Trinitrotoluene	0.0	183 nc	BRL	0.100 U	0.100 U	0.100 U	0.100 U	0.099 U	0.098 U
1,3-Dinitrobenzene	0.0	0.61 nc	BRL	0.100 U	0.100 U	0.100 U	0.100 U	0.099 U	0.098 U
Nitrobenzene	0.0	2 nc	BRL	0.100 U	0.100 U	0.100 U	0.100 U	0.099 U	0.098 U
2,4,6-Trinitrotoluene	0.0	16 ca	BRL	0.100 U	0.100 U	0.100 U	0.100 U	0.099 U	0.098 U
Tetryl	0.0	61 nc	BRL	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U
2,4-Dinitrotoluene	0.0	12 nc	BRL	0.100 U	0.100 U	0.100 U	0.100 U	0.099 U	0.098 U
2,6-Dinitrotoluene	0.0	6.1 nc	BRL	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U
2-Nitrotoluene	0.0	0.88 ca	BRL	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U
4-Nitrotoluene	0.0	12 ca	BRL	0.500 U	0.500 U	0.500 U	0.500 U	0.500 U	0.490 U
3-Nitrotoluene	0.0	73 nc	BRL	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U	0.200 U
Metals 6010B mg/kg (ppm)									
Antimony	0.96	3.1 nc	NT	0.20 (B)	0.46	0.19 (B)	0.33 U	0.33	1.4
Arsenic	15.4	0.39 ca	NT	18.2	16.9	3.1	5.9	9.2	16.9
Lead	26.1	400 pbk	NT	52.1	154	47.0	55.0	33.2	1330
Thallium	0.0	0.52 nc	NT	0.19 U	0.25 U	0.19 U	0.22 U	0.20 U	0.19 U
Mercury	0.04	2.3 nc	NT	0.054	0.060	0.017 (B)	0.071	0.23	882
Aluminum	17700	7614 nc	NT	10300	9110	10400	7200	10400	8700
Barium	88.4	538 nc	NT	64.9	87.3	135	90.0	58.6	40.2
Beryllium	0.88	15 nc	NT	0.61	0.50	1.8	0.48	0.49	0.34 (B)
Cadmium	0.0	3.7 nc	NT	0.24 U	0.71	0.23	0.36	0.22 U	0.14 (B)
Calcium	15800	-- [n]	NT	1380	1230	113000 (H)	75600 (H)	1780	5090
Chromium	17.4	30 ca	NT	15.0	12.6	12.0	11.8	14.2	16.1
Cobalt	10.4	902 ca	NT	8.7	8.3	1.7	6.2	7.7	9.8
Copper	17.7	313 nc	NT	15.9	24.3	9.6	25.2	14.9	1240
Iron	23100	2346 nc	NT	25400 (H)	18200 (H)	6140 (H)	17300 (H)	20200 (H)	42000 (H)
Magnesium	3030	-- [n]	NT	2040	1630	9600	2470	1990	2230
Manganese	1450	176 nc	NT	596	707	1360	493	451	557



Table 4-1
Summary of Azide Screening Surface Soil Results (0-1 ft)

ANALYTE, UNITS, METHOD NO.	Ravenna Soil Background Criteria (0-1 ft) mg/kg	Region 9 PRG Data (Residential Soil)	D-14S	LL9SS-001-0001-SO	LL9SS-003-0001-SO	LL9SS-005-0001-SO	LL9SS-007-0001-SO	LL9SS-009-0001-SO	LL9SS-011-0001-SO
Sample Date		ppm	3/15/2002	3/11/2002	3/11/2002	3/11/2002	3/11/2002	3/11/2002	3/11/2002
Nickel	21.1	156 nc	NT	15.6	13.7	5.7	16.6	13.9	17.3
Potassium	927	-- [n]	NT	954	756.0	903	1110	771	815
Selenium	1.4	39 nc	NT	1.2 U	1.1 U	1.1 U	1.0 U	1.1 U	0.91 U
Silver	0.0	39 nc	NT	0.59 U	0.55 U	0.53 U	0.52 U	0.54 U	0.46 U
Sodium	123	-- [n]	NT	192	128	279	137	177	94.3
Vanadium	31.1	7.8 nc	NT	21.7	17.9	6.9	13.9	20.2	12.6
Zinc	61.8	2346 nc	NT	86.0	228	46.6	94.1	72.6	711

PRGs - Preliminary Remediation Goals

nc - non-cancer basis, value is 1/10 the published PRG

ca - cancer basis

pbk - based on PBK modeling

[n] - nutrient

U - analyte not detected

NT - not tested

If Result = or > Background, then the value is presented with a shaded/highlighted style

If Result = or > Background & PRG, then the value is presented with a bold + shaded/highlighted style

If Result = or > PRG, then the value is presented with a bold style

If Result < PRG & Background, then the value is presented with a normal style

-- -no background/PRG value available for this analyte

INORGANIC FLAGS/QUALIFIERS

B - result is < CRDL/RL, but ≥ IDL/MDL

H - MB, EB1, EB2, EB3: Batch QC is > RL or had a negative instrument reading lower than the absolute value of the RL

ORGANIC FLAGS/QUALIFIERS

J - Result is an estimated value below the RL or a TIC

a - Concentration is below the method RL



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-001-0001-SO	3/11/02	0-1 ft	Aluminum	10300		17700	7614	nc	NO	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Arsenic	18.2		15.4	0.39	ca	YES	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Calcium	1380		15800		[n]	NO	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Iron	25400	H	23100	2346	nc	YES	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Lead	52.1		26.1	400	pbk	YES	NO
LL9SS-001-0001-SO	3/11/02	0-1 ft	Magnesium	2040		3030		[n]	NO	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Manganese	596		1450	176	nc	NO	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Mercury	0.054		0.04	2.3	nc	YES	NO
LL9SS-001-0001-SO	3/11/02	0-1 ft	Potassium	954		927		[n]	YES	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Sodium	192		123		[n]	YES	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Vanadium	21.7		31.1	7.8	nc	NO	YES
LL9SS-001-0001-SO	3/11/02	0-1 ft	Zinc	86		61.8	2346	nc	YES	NO
LL9SS-002-0001-SO	12/04/03	0-1 ft	Aluminum	8500		17700	7614	nc	NO	YES
LL9SS-002-0001-SO	12/04/03	0-1 ft	Arsenic	9.7		15.4	0.39	ca	NO	YES
LL9SS-002-0001-SO	12/04/03	0-1 ft	Copper	20		17.7	313	nc	YES	NO
LL9SS-002-0001-SO	12/04/03	0-1 ft	Iron	21000		23100	2346	nc	NO	YES
LL9SS-002-0001-SO	12/04/03	0-1 ft	Lead	64		26.1	400	pbk	YES	NO
LL9SS-002-0001-SO	12/04/03	0-1 ft	Manganese	630		1450	176	nc	NO	YES
LL9SS-002-0001-SO	12/04/03	0-1 ft	Mercury	0.1		0.04	2.3	nc	YES	NO
LL9SS-002-0001-SO	12/04/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-002-0001-SO	12/04/03	0-1 ft	Zinc	100		61.8	2346	nc	YES	NO
LL9SS-003-0001-SO	3/11/02	0-1 ft	Aluminum	9110		17700	7614	nc	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Arsenic	16.9		15.4	0.39	ca	YES	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Cadmium	0.71		0	3.7	nc	YES	NO
LL9SS-003-0001-SO	3/11/02	0-1 ft	Calcium	1230		15800		[n]	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Copper	24.3		17.7	313	nc	YES	NO
LL9SS-003-0001-SO	3/11/02	0-1 ft	Iron	18200	H	23100	2346	nc	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Lead	154		26.1	400	pbk	YES	NO
LL9SS-003-0001-SO	3/11/02	0-1 ft	Magnesium	1630		3030		[n]	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Manganese	707		1450	176	nc	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Mercury	0.06		0.04	2.3	nc	YES	NO
LL9SS-003-0001-SO	3/11/02	0-1 ft	Potassium	756		927		[n]	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Sodium	128		123		[n]	YES	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Vanadium	17.9		31.1	7.8	nc	NO	YES
LL9SS-003-0001-SO	3/11/02	0-1 ft	Zinc	228		61.8	2346	nc	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-004-0001-SO	12/04/03	0-1 ft	Arsenic	15		15.4	0.39	ca	NO	YES
LL9SS-004-0001-SO	12/04/03	0-1 ft	Copper	19		17.7	313	nc	YES	NO
LL9SS-004-0001-SO	12/04/03	0-1 ft	Iron	19000		23100	2346	nc	NO	YES
LL9SS-004-0001-SO	12/04/03	0-1 ft	Lead	45		26.1	400	pbk	YES	NO
LL9SS-004-0001-SO	12/04/03	0-1 ft	Manganese	390		1450	176	nc	NO	YES
LL9SS-004-0001-SO	12/04/03	0-1 ft	Mercury	0.073		0.04	2.3	nc	YES	NO
LL9SS-004-0001-SO	12/04/03	0-1 ft	Vanadium	12		31.1	7.8	nc	NO	YES
LL9SS-004-0001-SO	12/04/03	0-1 ft	Zinc	100		61.8	2346	nc	YES	NO
LL9SS-005-0001-SO	3/11/02	0-1 ft	Aluminum	10400		17700	7614	nc	NO	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Arsenic	3.1		15.4	0.39	ca	NO	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Barium	135		88.4	538	nc	YES	NO
LL9SS-005-0001-SO	3/11/02	0-1 ft	Beryllium	1.8		0.88	15	nc	YES	NO
LL9SS-005-0001-SO	3/11/02	0-1 ft	Cadmium	0.23		0	3.7	nc	YES	NO
LL9SS-005-0001-SO	3/11/02	0-1 ft	Calcium	113000	H	15800		[n]	YES	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Iron	6140	H	23100	2346	nc	NO	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Lead	47		26.1	400	pbk	YES	NO
LL9SS-005-0001-SO	3/11/02	0-1 ft	Magnesium	9600		3030		[n]	YES	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Manganese	1360		1450	176	nc	NO	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Potassium	903		927		[n]	NO	YES
LL9SS-005-0001-SO	3/11/02	0-1 ft	Sodium	279		123		[n]	YES	YES
LL9SS-006-0001-SO	12/11/03	0-1 ft	Aluminum	9300		17700	7614	nc	NO	YES
LL9SS-006-0001-SO	12/11/03	0-1 ft	Arsenic	9.9		15.4	0.39	ca	NO	YES
LL9SS-006-0001-SO	12/11/03	0-1 ft	Barium	92		88.4	538	nc	YES	NO
LL9SS-006-0001-SO	12/11/03	0-1 ft	Cadmium	2.9		0	3.7	nc	YES	NO
LL9SS-006-0001-SO	12/11/03	0-1 ft	Copper	18		17.7	313	nc	YES	NO
LL9SS-006-0001-SO	12/11/03	0-1 ft	Iron	16000	H	23100	2346	nc	NO	YES
LL9SS-006-0001-SO	12/11/03	0-1 ft	Lead	140		26.1	400	pbk	YES	NO
LL9SS-006-0001-SO	12/11/03	0-1 ft	Manganese	830		1450	176	nc	NO	YES
LL9SS-006-0001-SO	12/11/03	0-1 ft	Mercury	1.3		0.04	2.3	nc	YES	NO
LL9SS-006-0001-SO	12/11/03	0-1 ft	Vanadium	16		31.1	7.8	nc	NO	YES
LL9SS-006-0001-SO	12/11/03	0-1 ft	Zinc	200		61.8	2346	nc	YES	NO
LL9SS-007-0001-SO	3/11/02	0-1 ft	Arsenic	5.9		15.4	0.39	ca	NO	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Barium	90		88.4	538	nc	YES	NO
LL9SS-007-0001-SO	3/11/02	0-1 ft	Cadmium	0.36		0	3.7	nc	YES	NO
LL9SS-007-0001-SO	3/11/02	0-1 ft	Calcium	75600	H	15800		[n]	YES	YES



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-007-0001-SO	3/11/02	0-1 ft	Copper	25.2		17.7	313	nc	YES	NO
LL9SS-007-0001-SO	3/11/02	0-1 ft	Iron	17300	H	23100	2346	nc	NO	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Lead	55		26.1	400	pbk	YES	NO
LL9SS-007-0001-SO	3/11/02	0-1 ft	Magnesium	2470		3030		[n]	NO	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Manganese	493		1450	176	nc	NO	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Mercury	0.071		0.04	2.3	nc	YES	NO
LL9SS-007-0001-SO	3/11/02	0-1 ft	Potassium	1110		927		[n]	YES	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Sodium	137		123		[n]	YES	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Vanadium	13.9		31.1	7.8	nc	NO	YES
LL9SS-007-0001-SO	3/11/02	0-1 ft	Zinc	94.1		61.8	2346	nc	YES	NO
LL9SS-008-0001-SO	12/10/03	0-1 ft	Aluminum	11000		17700	7614	nc	NO	YES
LL9SS-008-0001-SO	12/10/03	0-1 ft	Arsenic	10		15.4	0.39	ca	NO	YES
LL9SS-008-0001-SO	12/10/03	0-1 ft	Iron	20000		23100	2346	nc	NO	YES
LL9SS-008-0001-SO	12/10/03	0-1 ft	Manganese	640		1450	176	nc	NO	YES
LL9SS-008-0001-SO	12/10/03	0-1 ft	Mercury	0.047		0.04	2.3	nc	YES	NO
LL9SS-008-0001-SO	12/10/03	0-1 ft	Vanadium	19		31.1	7.8	nc	NO	YES
LL9SS-008-0001-SO	12/10/03	0-1 ft	Zinc	62		61.8	2346	nc	YES	NO
LL9SS-009-0001-SO	3/11/02	0-1 ft	Aluminum	10400		17700	7614	nc	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Arsenic	9.2		15.4	0.39	ca	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Calcium	1780		15800		[n]	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Iron	20200	H	23100	2346	nc	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Lead	33.2		26.1	400	pbk	YES	NO
LL9SS-009-0001-SO	3/11/02	0-1 ft	Magnesium	1990		3030		[n]	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Manganese	451		1450	176	nc	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Mercury	0.23		0.04	2.3	nc	YES	NO
LL9SS-009-0001-SO	3/11/02	0-1 ft	Potassium	771		927		[n]	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Sodium	177		123		[n]	YES	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Vanadium	20.2		31.1	7.8	nc	NO	YES
LL9SS-009-0001-SO	3/11/02	0-1 ft	Zinc	72.6		61.8	2346	nc	YES	NO
LL9SS-010-0001-SO	12/08/03	0-1 ft	Arsenic	15		15.4	0.39	ca	NO	YES
LL9SS-010-0001-SO	12/08/03	0-1 ft	Cadmium	0.11	B	0	3.7	nc	YES	NO
LL9SS-010-0001-SO	12/08/03	0-1 ft	Copper	21	H	17.7	313	nc	YES	NO
LL9SS-010-0001-SO	12/08/03	0-1 ft	Iron	21000		23100	2346	nc	NO	YES
LL9SS-010-0001-SO	12/08/03	0-1 ft	Manganese	280		1450	176	nc	NO	YES
LL9SS-010-0001-SO	12/08/03	0-1 ft	Mercury	0.042		0.04	2.3	nc	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-010-0001-SO	12/08/03	0-1 ft	Vanadium	12		31.1	7.8	nc	NO	YES
LL9SS-010-0001-SO	12/08/03	0-1 ft	Zinc	78		61.8	2346	nc	YES	NO
LL9SS-011-0001-SO	3/11/02	0-1 ft	Aluminum	8700		17700	7614	nc	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Antimony	1.4		0.96	3.1	nc	YES	NO
LL9SS-011-0001-SO	3/11/02	0-1 ft	Arsenic	16.9		15.4	0.39	ca	YES	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Cadmium	0.14	B	0	3.7	nc	YES	NO
LL9SS-011-0001-SO	3/11/02	0-1 ft	Calcium	5090		15800		[n]	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Copper	1240		17.7	313	nc	YES	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Iron	42000	H	23100	2346	nc	YES	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Lead	1330		26.1	400	pbk	YES	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Magnesium	2230		3030		[n]	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Manganese	557		1450	176	nc	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Mercury	882		0.04	2.3	nc	YES	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Potassium	815		927		[n]	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Sodium	94.3		123		[n]	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Vanadium	12.6		31.1	7.8	nc	NO	YES
LL9SS-011-0001-SO	3/11/02	0-1 ft	Zinc	711		61.8	2346	nc	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Aluminum	9000		17700	7614	nc	NO	YES
LL9SS-012-0001-SO	12/10/03	0-1 ft	Arsenic	15		15.4	0.39	ca	NO	YES
LL9SS-012-0001-SO	12/10/03	0-1 ft	Cadmium	0.51		0	3.7	nc	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Calcium	22000		15800	--[n]	0	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Copper	25		17.7	313	nc	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Iron	22000		23100	2346	nc	NO	YES
LL9SS-012-0001-SO	12/10/03	0-1 ft	Lead	44		26.1	400	pbk	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Manganese	600		1450	176	nc	NO	YES
LL9SS-012-0001-SO	12/10/03	0-1 ft	Mercury	0.053		0.04	2.3	nc	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Potassium	1100		927	--[n]	0	YES	NO
LL9SS-012-0001-SO	12/10/03	0-1 ft	Vanadium	16		31.1	7.8	nc	NO	YES
LL9SS-012-0001-SO	12/10/03	0-1 ft	Zinc	140		61.8	2346	nc	YES	NO
LL9SS-013-0001-SO	12/10/03	0-1 ft	Arsenic	11		15.4	0.39	ca	NO	YES
LL9SS-013-0001-SO	12/10/03	0-1 ft	Copper	22		17.7	313	nc	YES	NO
LL9SS-013-0001-SO	12/10/03	0-1 ft	Iron	24000		23100	2346	nc	YES	YES
LL9SS-013-0001-SO	12/10/03	0-1 ft	Manganese	370		1450	176	nc	NO	YES
LL9SS-013-0001-SO	12/10/03	0-1 ft	Vanadium	11		31.1	7.8	nc	NO	YES
LL9SS-013-0001-SO	12/10/03	0-1 ft	Zinc	74		61.8	2346	nc	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Arsenic	14		15.4	0.39	ca	NO	YES
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Copper	20		17.7	313	nc	YES	NO
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Iron	19000	H	23100	2346	nc	NO	YES
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Manganese	340		1450	176	nc	NO	YES
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Nitrocellulose	1.3	B J	0	--	0	YES	NO
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Vanadium	12		31.1	7.8	nc	NO	YES
LL9SS-014-0001-DUP	12/11/03	0-1 ft	Zinc	64		61.8	2346	nc	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Aluminum	7900		17700	7614	nc	NO	YES
LL9SS-014-0001-SO	12/11/03	0-1 ft	Arsenic	15		15.4	0.39	ca	NO	YES
LL9SS-014-0001-SO	12/11/03	0-1 ft	Copper	22		17.7	313	nc	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Iron	19000	H	23100	2346	nc	NO	YES
LL9SS-014-0001-SO	12/11/03	0-1 ft	Lead	43		26.1	400	pbk	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Manganese	410		1450	176	nc	NO	YES
LL9SS-014-0001-SO	12/11/03	0-1 ft	Mercury	0.044		0.04	2.3	nc	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Nitrocellulose	1.6	B J	0	--	0	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Nitroguanidine	0.965		0	611	nc	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Nitroguanidine	1		0	611	nc	YES	NO
LL9SS-014-0001-SO	12/11/03	0-1 ft	Vanadium	13		31.1	7.8	nc	NO	YES
LL9SS-014-0001-SO	12/11/03	0-1 ft	Zinc	72		61.8	2346	nc	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Aluminum	17000		17700	7614	nc	NO	YES
LL9SS-015-0001-SO	12/11/03	0-1 ft	Arsenic	24		15.4	0.39	ca	YES	YES
LL9SS-015-0001-SO	12/11/03	0-1 ft	Chromium	23		17.4	30	ca	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Copper	24		17.7	313	nc	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Iron	31000	H	23100	2346	nc	YES	YES
LL9SS-015-0001-SO	12/11/03	0-1 ft	Magnesium	3500		3030	--[n]	0	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Manganese	180		1450	176	nc	NO	YES
LL9SS-015-0001-SO	12/11/03	0-1 ft	Mercury	0.061		0.04	2.3	nc	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Nickel	23		21.1	156	nc	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Potassium	1500		927	--[n]	0	YES	NO
LL9SS-015-0001-SO	12/11/03	0-1 ft	Vanadium	28		31.1	7.8	nc	NO	YES
LL9SS-015-0001-SO	12/11/03	0-1 ft	Zinc	64		61.8	2346	nc	YES	NO
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Aluminum	9100		17700	7614	nc	NO	YES
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Arsenic	14		15.4	0.39	ca	NO	YES
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Cadmium	0.14	B	0	3.7	nc	YES	NO
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Copper	18		17.7	313	nc	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Iron	17000	H	23100	2346	nc	NO	YES
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Lead	35		26.1	400	pbk	YES	NO
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Magnesium	3300		3030	--[n]	0	YES	NO
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Manganese	790		1450	176	nc	NO	YES
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Mercury	0.11		0.04	2.3	nc	YES	NO
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Potassium	970		927	--[n]	0	YES	NO
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Vanadium	13		31.1	7.8	nc	NO	YES
LL9SS-016-0001-DUP	12/11/03	0-1 ft	Zinc	72		61.8	2346	nc	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Aluminum	9900		17700	7614	nc	NO	YES
LL9SS-016-0001-SO	12/11/03	0-1 ft	Arsenic	11		15.4	0.39	ca	NO	YES
LL9SS-016-0001-SO	12/11/03	0-1 ft	Cadmium	0.17	B	0	3.7	nc	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Calcium	20000		15800	--[n]	0	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Chromium	19.33	*	17.4	30	ca	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Copper	19		17.7	313	nc	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Iron	18000	H	23100	2346	nc	NO	YES
LL9SS-016-0001-SO	12/11/03	0-1 ft	Lead	38		26.1	400	pbk	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Magnesium	3500		3030	--[n]	0	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Manganese	577.68	*	1450	176	nc	NO	YES
LL9SS-016-0001-SO	12/11/03	0-1 ft	Manganese	1100		1450	176	nc	NO	YES
LL9SS-016-0001-SO	12/11/03	0-1 ft	Mercury	0.096		0.04	2.3	nc	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Potassium	1000		927	--[n]	0	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Sodium	170		123	--[n]	0	YES	NO
LL9SS-016-0001-SO	12/11/03	0-1 ft	Vanadium	12		31.1	7.8	nc	NO	YES
LL9SS-016-0001-SO	12/11/03	0-1 ft	Zinc	70		61.8	2346	nc	YES	NO
LL9SS-017-0001-SO	12/04/03	0-1 ft	Aluminum	10000		17700	7614	nc	NO	YES
LL9SS-017-0001-SO	12/04/03	0-1 ft	Arsenic	11		15.4	0.39	ca	NO	YES
LL9SS-017-0001-SO	12/04/03	0-1 ft	Iron	23000		23100	2346	nc	NO	YES
LL9SS-017-0001-SO	12/04/03	0-1 ft	Manganese	690		1450	176	nc	NO	YES
LL9SS-017-0001-SO	12/04/03	0-1 ft	Vanadium	21		31.1	7.8	nc	NO	YES
LL9SS-018-0001-SO	12/10/03	0-1 ft	Aluminum	11000		17700	7614	nc	NO	YES
LL9SS-018-0001-SO	12/10/03	0-1 ft	Arsenic	10		15.4	0.39	ca	NO	YES
LL9SS-018-0001-SO	12/10/03	0-1 ft	Barium	95		88.4	538	nc	YES	NO
LL9SS-018-0001-SO	12/10/03	0-1 ft	Cadmium	0.16	B	0	3.7	nc	YES	NO
LL9SS-018-0001-SO	12/10/03	0-1 ft	Iron	19000		23100	2346	nc	NO	YES
LL9SS-018-0001-SO	12/10/03	0-1 ft	Lead	40		26.1	400	pbk	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-018-0001-SO	12/10/03	0-1 ft	Manganese	920		1450	176	nc	NO	YES
LL9SS-018-0001-SO	12/10/03	0-1 ft	Mercury	0.09		0.04	2.3	nc	YES	NO
LL9SS-018-0001-SO	12/10/03	0-1 ft	Vanadium	19		31.1	7.8	nc	NO	YES
LL9SS-018-0001-SO	12/10/03	0-1 ft	Zinc	69		61.8	2346	nc	YES	NO
LL9SS-019-0001-SO	12/11/03	0-1 ft	Aluminum	14000		17700	7614	nc	NO	YES
LL9SS-019-0001-SO	12/11/03	0-1 ft	Arsenic	17		15.4	0.39	ca	YES	YES
LL9SS-019-0001-SO	12/11/03	0-1 ft	Barium	170		88.4	538	nc	YES	NO
LL9SS-019-0001-SO	12/11/03	0-1 ft	Copper	19		17.7	313	nc	YES	NO
LL9SS-019-0001-SO	12/11/03	0-1 ft	Iron	27000	H	23100	2346	nc	YES	YES
LL9SS-019-0001-SO	12/11/03	0-1 ft	Manganese	210		1450	176	nc	NO	YES
LL9SS-019-0001-SO	12/11/03	0-1 ft	Mercury	0.36		0.04	2.3	nc	YES	NO
LL9SS-019-0001-SO	12/11/03	0-1 ft	Nitrocellulose	1.8	B J	0	--	0	YES	NO
LL9SS-019-0001-SO	12/11/03	0-1 ft	Potassium	1000		927	--[n]	0	YES	NO
LL9SS-019-0001-SO	12/11/03	0-1 ft	Vanadium	22		31.1	7.8	nc	NO	YES
LL9SS-020-0001-SO	12/08/03	0-1 ft	Aluminum	7900		17700	7614	nc	NO	YES
LL9SS-020-0001-SO	12/08/03	0-1 ft	Arsenic	13		15.4	0.39	ca	NO	YES
LL9SS-020-0001-SO	12/08/03	0-1 ft	Cadmium	0.71		0	3.7	nc	YES	NO
LL9SS-020-0001-SO	12/08/03	0-1 ft	Copper	31	H	17.7	313	nc	YES	NO
LL9SS-020-0001-SO	12/08/03	0-1 ft	Iron	16000		23100	2346	nc	NO	YES
LL9SS-020-0001-SO	12/08/03	0-1 ft	Lead	33		26.1	400	pbk	YES	NO
LL9SS-020-0001-SO	12/08/03	0-1 ft	Manganese	460		1450	176	nc	NO	YES
LL9SS-020-0001-SO	12/08/03	0-1 ft	Mercury	0.74		0.04	2.3	nc	YES	NO
LL9SS-020-0001-SO	12/08/03	0-1 ft	Vanadium	14		31.1	7.8	nc	NO	YES
LL9SS-020-0001-SO	12/08/03	0-1 ft	Zinc	85		61.8	2346	nc	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Aluminum	13000		17700	7614	nc	NO	YES
LL9SS-021-0001-SO	12/08/03	0-1 ft	Arsenic	16		15.4	0.39	ca	YES	YES
LL9SS-021-0001-SO	12/08/03	0-1 ft	Cadmium	0.5		0	3.7	nc	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Chromium	18		17.4	30	ca	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Copper	19	H	17.7	313	nc	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Iron	27000		23100	2346	nc	YES	YES
LL9SS-021-0001-SO	12/08/03	0-1 ft	Lead	35		26.1	400	pbk	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Manganese	260		1450	176	nc	NO	YES
LL9SS-021-0001-SO	12/08/03	0-1 ft	Mercury	0.045		0.04	2.3	nc	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Potassium	1000		927	--[n]	0	YES	NO
LL9SS-021-0001-SO	12/08/03	0-1 ft	Vanadium	22		31.1	7.8	nc	NO	YES



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-021-0001-SO	12/08/03	0-1 ft	Zinc	72		61.8	2346	nc	YES	NO
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Aluminum	9600		17700	7614	nc	NO	YES
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Arsenic	12		15.4	0.39	ca	NO	YES
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Copper	21		17.7	313	nc	YES	NO
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Iron	22000		23100	2346	nc	NO	YES
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Lead	48		26.1	400	pbk	YES	NO
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Manganese	480		1450	176	nc	NO	YES
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Mercury	0.2		0.04	2.3	nc	YES	NO
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-022-0001-DUP	12/10/03	0-1 ft	Zinc	69		61.8	2346	nc	YES	NO
LL9SS-022-0001-SO	12/10/03	0-1 ft	Aluminum	8800		17700	7614	nc	NO	YES
LL9SS-022-0001-SO	12/10/03	0-1 ft	Arsenic	13		15.4	0.39	ca	NO	YES
LL9SS-022-0001-SO	12/10/03	0-1 ft	Iron	20000		23100	2346	nc	NO	YES
LL9SS-022-0001-SO	12/10/03	0-1 ft	Mercury	0.11		0.04	2.3	nc	YES	NO
LL9SS-022-0001-SO	12/10/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-023-0001-SO	12/08/03	0-1 ft	Aluminum	20000		17700	7614	nc	YES	YES
LL9SS-023-0001-SO	12/08/03	0-1 ft	Arsenic	21		15.4	0.39	ca	YES	YES
LL9SS-023-0001-SO	12/08/03	0-1 ft	Cadmium	0.11	B	0	3.7	nc	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Chromium	21		17.4	30	ca	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Copper	23	H	17.7	313	nc	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Iron	26000		23100	2346	nc	YES	YES
LL9SS-023-0001-SO	12/08/03	0-1 ft	Lead	50		26.1	400	pbk	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Magnesium	3200		3030	--[n]	0	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Manganese	230		1450	176	nc	NO	YES
LL9SS-023-0001-SO	12/08/03	0-1 ft	Mercury	0.25		0.04	2.3	nc	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Potassium	1500		927	--[n]	0	YES	NO
LL9SS-023-0001-SO	12/08/03	0-1 ft	Vanadium	28		31.1	7.8	nc	NO	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Aluminum	8000		17700	7614	nc	NO	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Arsenic	10		15.4	0.39	ca	NO	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Cadmium	0.42		0	3.7	nc	YES	NO
LL9SS-024-0001-SO	12/10/03	0-1 ft	Chromium	110		17.4	30	ca	YES	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Iron	14000		23100	2346	nc	NO	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Lead	320		26.1	400	pbk	YES	NO
LL9SS-024-0001-SO	12/10/03	0-1 ft	Manganese	500		1450	176	nc	NO	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Mercury	0.057		0.04	2.3	nc	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-024-0001-SO	12/10/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-024-0001-SO	12/10/03	0-1 ft	Zinc	210		61.8	2346	nc	YES	NO
LL9SS-025-0001-SO	12/10/03	0-1 ft	Aluminum	12000		17700	7614	nc	NO	YES
LL9SS-025-0001-SO	12/10/03	0-1 ft	Arsenic	8.4		15.4	0.39	ca	NO	YES
LL9SS-025-0001-SO	12/10/03	0-1 ft	Iron	24000		23100	2346	nc	YES	YES
LL9SS-025-0001-SO	12/10/03	0-1 ft	Manganese	420		1450	176	nc	NO	YES
LL9SS-025-0001-SO	12/10/03	0-1 ft	Nitrocellulose	1.6	B J	0	--	0	YES	NO
LL9SS-025-0001-SO	12/10/03	0-1 ft	Vanadium	23		31.1	7.8	nc	NO	YES
LL9SS-026-0001-SO	12/10/03	0-1 ft	Aluminum	13000		17700	7614	nc	NO	YES
LL9SS-026-0001-SO	12/10/03	0-1 ft	Antimony	1.2		0.96	3.1	nc	YES	NO
LL9SS-026-0001-SO	12/10/03	0-1 ft	Arsenic	8.3		15.4	0.39	ca	NO	YES
LL9SS-026-0001-SO	12/10/03	0-1 ft	Barium	100		88.4	538	nc	YES	NO
LL9SS-026-0001-SO	12/10/03	0-1 ft	Cobalt	11		10.4	30	ca	YES	NO
LL9SS-026-0001-SO	12/10/03	0-1 ft	Iron	23000		23100	2346	nc	NO	YES
LL9SS-026-0001-SO	12/10/03	0-1 ft	Lead	28		26.1	400	pbk	YES	NO
LL9SS-026-0001-SO	12/10/03	0-1 ft	Manganese	860		1450	176	nc	NO	YES
LL9SS-026-0001-SO	12/10/03	0-1 ft	Mercury	0.045		0.04	2.3	nc	YES	NO
LL9SS-026-0001-SO	12/10/03	0-1 ft	Thallium	0.26		0	0.52	nc	YES	NO
LL9SS-026-0001-SO	12/10/03	0-1 ft	Vanadium	23		31.1	7.8	nc	NO	YES
LL9SS-026-0001-SO	12/10/03	0-1 ft	Zinc	71		61.8	2346	nc	YES	NO
LL9SS-027-0001-SO	12/04/03	0-1 ft	Aluminum	10000		17700	7614	nc	NO	YES
LL9SS-027-0001-SO	12/04/03	0-1 ft	Arsenic	11		15.4	0.39	ca	NO	YES
LL9SS-027-0001-SO	12/04/03	0-1 ft	Cobalt	18		10.4	30	ca	YES	NO
LL9SS-027-0001-SO	12/04/03	0-1 ft	Iron	25000		23100	2346	nc	YES	YES
LL9SS-027-0001-SO	12/04/03	0-1 ft	Manganese	3800		1450	176	nc	YES	YES
LL9SS-027-0001-SO	12/04/03	0-1 ft	Mercury	0.041		0.04	2.3	nc	YES	NO
LL9SS-027-0001-SO	12/04/03	0-1 ft	Vanadium	23		31.1	7.8	nc	NO	YES
LL9SS-028-0001-DUP	12/04/03	0-1 ft	Aluminum	10000		17700	7614	nc	NO	YES
LL9SS-028-0001-DUP	12/04/03	0-1 ft	Arsenic	9.4		15.4	0.39	ca	NO	YES
LL9SS-028-0001-DUP	12/04/03	0-1 ft	Iron	23000		23100	2346	nc	NO	YES
LL9SS-028-0001-DUP	12/04/03	0-1 ft	Manganese	330		1450	176	nc	NO	YES
LL9SS-028-0001-DUP	12/04/03	0-1 ft	Mercury	0.2		0.04	2.3	nc	YES	NO
LL9SS-028-0001-DUP	12/04/03	0-1 ft	Vanadium	22		31.1	7.8	nc	NO	YES
LL9SS-028-0001-SO	12/04/03	0-1 ft	Aluminum	10000		17700	7614	nc	NO	YES
LL9SS-028-0001-SO	12/04/03	0-1 ft	Arsenic	15		15.4	0.39	ca	NO	YES



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-028-0001-SO	12/04/03	0-1 ft	Iron	21000		23100	2346	nc	NO	YES
LL9SS-028-0001-SO	12/04/03	0-1 ft	Manganese	460		1450	176	nc	NO	YES
LL9SS-028-0001-SO	12/04/03	0-1 ft	Mercury	0.24		0.04	2.3	nc	YES	NO
LL9SS-028-0001-SO	12/04/03	0-1 ft	Vanadium	21		31.1	7.8	nc	NO	YES
LL9SS-029-0001-SO	12/04/03	0-1 ft	Aluminum	9400		17700	7614	nc	NO	YES
LL9SS-029-0001-SO	12/04/03	0-1 ft	Arsenic	7.4		15.4	0.39	ca	NO	YES
LL9SS-029-0001-SO	12/04/03	0-1 ft	Iron	16000		23100	2346	nc	NO	YES
LL9SS-029-0001-SO	12/04/03	0-1 ft	Manganese	870		1450	176	nc	NO	YES
LL9SS-029-0001-SO	12/04/03	0-1 ft	Mercury	0.088		0.04	2.3	nc	YES	NO
LL9SS-029-0001-SO	12/04/03	0-1 ft	Vanadium	16		31.1	7.8	nc	NO	YES
LL9SS-030-0001-SO	12/04/03	0-1 ft	Arsenic	4.2		15.4	0.39	ca	NO	YES
LL9SS-030-0001-SO	12/04/03	0-1 ft	Iron	20000		23100	2346	nc	NO	YES
LL9SS-030-0001-SO	12/04/03	0-1 ft	Manganese	650		1450	176	nc	NO	YES
LL9SS-030-0001-SO	12/04/03	0-1 ft	Vanadium	14		31.1	7.8	nc	NO	YES
LL9SS-031-0001-SO	12/04/03	0-1 ft	Arsenic	6.9		15.4	0.39	ca	NO	YES
LL9SS-031-0001-SO	12/04/03	0-1 ft	Cadmium	0.31		0	3.7	nc	YES	NO
LL9SS-031-0001-SO	12/04/03	0-1 ft	Cobalt	13		10.4	30	ca	YES	NO
LL9SS-031-0001-SO	12/04/03	0-1 ft	Iron	76000		23100	2346	nc	YES	YES
LL9SS-031-0001-SO	12/04/03	0-1 ft	Lead	28		26.1	400	pbk	YES	NO
LL9SS-031-0001-SO	12/04/03	0-1 ft	Manganese	2000		1450	176	nc	YES	YES
LL9SS-031-0001-SO	12/04/03	0-1 ft	Vanadium	19		31.1	7.8	nc	NO	YES
LL9SS-031-0001-SO	12/04/03	0-1 ft	Zinc	75		61.8	2346	nc	YES	NO
LL9SS-032-0001-SO	12/04/03	0-1 ft	Aluminum	8200		17700	7614	nc	NO	YES
LL9SS-032-0001-SO	12/04/03	0-1 ft	Arsenic	7.4		15.4	0.39	ca	NO	YES
LL9SS-032-0001-SO	12/04/03	0-1 ft	Iron	22000		23100	2346	nc	NO	YES
LL9SS-032-0001-SO	12/04/03	0-1 ft	Manganese	480		1450	176	nc	NO	YES
LL9SS-032-0001-SO	12/04/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-033-0001-SO	11/11/03	0-1 ft	Aluminum	13000		17700	7614	nc	NO	YES
LL9SS-033-0001-SO	11/11/03	0-1 ft	Arsenic	9.2		15.4	0.39	ca	NO	YES
LL9SS-033-0001-SO	11/11/03	0-1 ft	Cobalt	11		10.4	30	ca	YES	NO
LL9SS-033-0001-SO	11/11/03	0-1 ft	Iron	23000		23100	2346	nc	NO	YES
LL9SS-033-0001-SO	11/11/03	0-1 ft	Manganese	1200		1450	176	nc	NO	YES
LL9SS-033-0001-SO	11/11/03	0-1 ft	Nitrocellulose	1.4	B	0	--	0	YES	NO
LL9SS-033-0001-SO	11/11/03	0-1 ft	Sodium	1000		123	--[n]	0	YES	NO
LL9SS-033-0001-SO	11/11/03	0-1 ft	Vanadium	25		31.1	7.8	nc	NO	YES



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-033-0001-SO	11/11/03	0-1 ft	Zinc	86		61.8	2346	nc	YES	NO
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Arsenic	10		15.4	0.39	ca	NO	YES
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Benzo(a)pyrene	0.073		--	0.062	ca	NO	YES
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Copper	33		17.7	313	nc	YES	NO
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Iron	32000		23100	2346	nc	YES	YES
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Lead	38		26.1	400	pbk	YES	NO
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Manganese	420		1450	176	nc	NO	YES
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Mercury	17		0.04	2.3	nc	YES	YES
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Sodium	660		123	--[n]	0	YES	NO
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-034-0001-DUP	11/11/03	0-1 ft	Zinc	79		61.8	2346	nc	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Aluminum	9300		17700	7614	nc	NO	YES
LL9SS-034-0001-SO	11/11/03	0-1 ft	Arsenic	7.61		15.4	0.39	ca	NO	YES
LL9SS-034-0001-SO	11/11/03	0-1 ft	Copper	34		17.7	313	nc	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Iron	21000		23100	2346	nc	NO	YES
LL9SS-034-0001-SO	11/11/03	0-1 ft	Lead	43		26.1	400	pbk	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Manganese	630		1450	176	nc	NO	YES
LL9SS-034-0001-SO	11/11/03	0-1 ft	Mercury	12		0.04	2.3	nc	YES	YES
LL9SS-034-0001-SO	11/11/03	0-1 ft	Nitrocellulose	1.1	B	0	--	0	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Nitroglycerin	4.84		0	--	0	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Nitroguanidine	0.089	J	0	611	nc	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Sodium	1000		123	--[n]	0	YES	NO
LL9SS-034-0001-SO	11/11/03	0-1 ft	Vanadium	17		31.1	7.8	nc	NO	YES
LL9SS-034-0001-SO	11/11/03	0-1 ft	Zinc	71		61.8	2346	nc	YES	NO
LL9SS-035-0001-SO	11/11/03	0-1 ft	Aluminum	7900		17700	7614	nc	NO	YES
LL9SS-035-0001-SO	11/11/03	0-1 ft	Arsenic	10		15.4	0.39	ca	NO	YES
LL9SS-035-0001-SO	11/11/03	0-1 ft	Iron	22000		23100	2346	nc	NO	YES
LL9SS-035-0001-SO	11/11/03	0-1 ft	Manganese	350		1450	176	nc	NO	YES
LL9SS-035-0001-SO	11/11/03	0-1 ft	Sodium	1100		123	--[n]	0	YES	NO
LL9SS-035-0001-SO	11/11/03	0-1 ft	Vanadium	14		31.1	7.8	nc	NO	YES
LL9SS-035-0001-SO	11/11/03	0-1 ft	Zinc	66		61.8	2346	nc	YES	NO
LL9SS-036-0001-DUP	11/11/03	0-1 ft	Aluminum	8500		17700	7614	nc	NO	YES
LL9SS-036-0001-DUP	11/11/03	0-1 ft	Arsenic	5.6		15.4	0.39	ca	NO	YES
LL9SS-036-0001-DUP	11/11/03	0-1 ft	Iron	20000		23100	2346	nc	NO	YES
LL9SS-036-0001-DUP	11/11/03	0-1 ft	Manganese	570		1450	176	nc	NO	YES



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-036-0001-DUP	11/11/03	0-1 ft	Sodium	970		123	--[n]	0	YES	NO
LL9SS-036-0001-DUP	11/11/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-036-0001-SO	11/11/03	0-1 ft	Aluminum	8500		17700	7614	nc	NO	YES
LL9SS-036-0001-SO	11/11/03	0-1 ft	Arsenic	5.3		15.4	0.39	ca	NO	YES
LL9SS-036-0001-SO	11/11/03	0-1 ft	Iron	19000		23100	2346	nc	NO	YES
LL9SS-036-0001-SO	11/11/03	0-1 ft	Manganese	420		1450	176	nc	NO	YES
LL9SS-036-0001-SO	11/11/03	0-1 ft	Sodium	970		123	--[n]	0	YES	NO
LL9SS-036-0001-SO	11/11/03	0-1 ft	Vanadium	15		31.1	7.8	nc	NO	YES
LL9SS-037-0001-SO	11/11/03	0-1 ft	Aluminum	12000		17700	7614	nc	NO	YES
LL9SS-037-0001-SO	11/11/03	0-1 ft	Arsenic	9.4		15.4	0.39	ca	NO	YES
LL9SS-037-0001-SO	11/11/03	0-1 ft	Barium	97		88.4	538	nc	YES	NO
LL9SS-037-0001-SO	11/11/03	0-1 ft	Cobalt	17		10.4	30	ca	YES	NO
LL9SS-037-0001-SO	11/11/03	0-1 ft	Iron	17000		23100	2346	nc	NO	YES
LL9SS-037-0001-SO	11/11/03	0-1 ft	Manganese	970		1450	176	nc	NO	YES
LL9SS-037-0001-SO	11/11/03	0-1 ft	Sodium	1100		123	--[n]	0	YES	NO
LL9SS-037-0001-SO	11/11/03	0-1 ft	Vanadium	21		31.1	7.8	nc	NO	YES
LL9SS-037-0001-SO	11/11/03	0-1 ft	Zinc	65		61.8	2346	nc	YES	NO
LL9SS-038-0001-SO	11/11/03	0-1 ft	Aluminum	13000		17700	7614	nc	NO	YES
LL9SS-038-0001-SO	11/11/03	0-1 ft	Arsenic	7.5		15.4	0.39	ca	NO	YES
LL9SS-038-0001-SO	11/11/03	0-1 ft	Cobalt	14		10.4	30	ca	YES	NO
LL9SS-038-0001-SO	11/11/03	0-1 ft	Iron	19000		23100	2346	nc	NO	YES
LL9SS-038-0001-SO	11/11/03	0-1 ft	Manganese	1000		1450	176	nc	NO	YES
LL9SS-038-0001-SO	11/11/03	0-1 ft	Mercury	0.072		0.04	2.3	nc	YES	NO
LL9SS-038-0001-SO	11/11/03	0-1 ft	Sodium	1300		123	--[n]	0	YES	NO
LL9SS-038-0001-SO	11/11/03	0-1 ft	Vanadium	22		31.1	7.8	nc	NO	YES
LL9SS-038-0001-SO	11/11/03	0-1 ft	Zinc	69		61.8	2346	nc	YES	NO
LL9SS-039-0001-DUP	11/11/03	0-1 ft	Arsenic	3.1		15.4	0.39	ca	NO	YES
LL9SS-039-0001-DUP	11/11/03	0-1 ft	Iron	25000		23100	2346	nc	YES	YES
LL9SS-039-0001-DUP	11/11/03	0-1 ft	Manganese	420		1450	176	nc	NO	YES
LL9SS-039-0001-DUP	11/11/03	0-1 ft	Sodium	850		123	--[n]	0	YES	NO
LL9SS-039-0001-DUP	11/11/03	0-1 ft	Vanadium	11		31.1	7.8	nc	NO	YES
LL9SS-039-0001-DUP	11/11/03	0-1 ft	Zinc	69		61.8	2346	nc	YES	NO
LL9SS-039-0001-SO	11/11/03	0-1 ft	Arsenic	4.2		15.4	0.39	ca	NO	YES
LL9SS-039-0001-SO	11/11/03	0-1 ft	Iron	25000		23100	2346	nc	YES	YES
LL9SS-039-0001-SO	11/11/03	0-1 ft	Manganese	470		1450	176	nc	NO	YES



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-039-0001-SO	11/11/03	0-1 ft	Mercury	0.05		0.04	2.3	nc	YES	NO
LL9SS-039-0001-SO	11/11/03	0-1 ft	Sodium	940		123	--[n]	0	YES	NO
LL9SS-039-0001-SO	11/11/03	0-1 ft	Vanadium	12		31.1	7.8	nc	NO	YES
LL9SS-039-0001-SO	11/11/03	0-1 ft	Zinc	76		61.8	2346	nc	YES	NO
LL9SS-042-0001-SO	11/06/03	0-1 ft	Arsenic	4		15.4	0.39	ca	NO	YES
LL9SS-042-0001-SO	11/06/03	0-1 ft	Iron	18000		23100	2346	nc	NO	YES
LL9SS-042-0001-SO	11/06/03	0-1 ft	Manganese	610		1450	176	nc	NO	YES
LL9SS-042-0001-SO	11/06/03	0-1 ft	Sodium	930		123	--[n]	0	YES	NO
LL9SS-042-0001-SO	11/06/03	0-1 ft	Vanadium	13		31.1	7.8	nc	NO	YES
LL9SS-043-0001-SO	11/06/03	0-1 ft	Aluminum	10000		17700	7614	nc	NO	YES
LL9SS-043-0001-SO	11/06/03	0-1 ft	Arsenic	9.2		15.4	0.39	ca	NO	YES
LL9SS-043-0001-SO	11/06/03	0-1 ft	Iron	20000		23100	2346	nc	NO	YES
LL9SS-043-0001-SO	11/06/03	0-1 ft	Manganese	440		1450	176	nc	NO	YES
LL9SS-043-0001-SO	11/06/03	0-1 ft	Sodium	1100		123	--[n]	0	YES	NO
LL9SS-043-0001-SO	11/06/03	0-1 ft	Vanadium	17		31.1	7.8	nc	NO	YES
LL9SS-044-0001-SO	11/06/03	0-1 ft	Aluminum	12000		17700	7614	nc	NO	YES
LL9SS-044-0001-SO	11/06/03	0-1 ft	Arsenic	18		15.4	0.39	ca	YES	YES
LL9SS-044-0001-SO	11/06/03	0-1 ft	Barium	92.39		88.4	538	nc	YES	NO
LL9SS-044-0001-SO	11/06/03	0-1 ft	Iron	17000		23100	2346	nc	NO	YES
LL9SS-044-0001-SO	11/06/03	0-1 ft	Sodium	1200		123	--[n]	0	YES	NO
LL9SS-044-0001-SO	11/06/03	0-1 ft	Vanadium	13		31.1	7.8	nc	NO	YES
LL9SS-045-0001-SO	11/06/03	0-1 ft	Aluminum	13000		17700	7614	nc	NO	YES
LL9SS-045-0001-SO	11/06/03	0-1 ft	Arsenic	17		15.4	0.39	ca	YES	YES
LL9SS-045-0001-SO	11/06/03	0-1 ft	Copper	25		17.7	313	nc	YES	NO
LL9SS-045-0001-SO	11/06/03	0-1 ft	Iron	31000		23100	2346	nc	YES	YES
LL9SS-045-0001-SO	11/06/03	0-1 ft	Manganese	240		1450	176	nc	NO	YES
LL9SS-045-0001-SO	11/06/03	0-1 ft	Potassium	1200		927	--[n]	0	YES	NO
LL9SS-045-0001-SO	11/06/03	0-1 ft	Sodium	1100		123	--[n]	0	YES	NO
LL9SS-045-0001-SO	11/06/03	0-1 ft	Vanadium	20		31.1	7.8	nc	NO	YES
LL9SS-045-0001-SO	11/06/03	0-1 ft	Zinc	62		61.8	2346	nc	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-046-0001-SO	11/06/03	0-1 ft	Aluminum	8900		17700	7614	nc	NO	YES
LL9SS-046-0001-SO	11/06/03	0-1 ft	Arsenic	18		15.4	0.39	ca	YES	YES
LL9SS-046-0001-SO	11/06/03	0-1 ft	Cobalt	11		10.4	30	ca	YES	NO
LL9SS-046-0001-SO	11/06/03	0-1 ft	Copper	21		17.7	313	nc	YES	NO
LL9SS-046-0001-SO	11/06/03	0-1 ft	Iron	25000		23100	2346	nc	YES	YES
LL9SS-046-0001-SO	11/06/03	0-1 ft	Manganese	370		1450	176	nc	NO	YES
LL9SS-046-0001-SO	11/06/03	0-1 ft	Nickel	24		21.1	156	nc	YES	NO
LL9SS-046-0001-SO	11/06/03	0-1 ft	Sodium	1300		123	--[n]	0	YES	NO
LL9SS-046-0001-SO	11/06/03	0-1 ft	Vanadium	14		31.1	7.8	nc	NO	YES
LL9SS-046-0001-SO	11/06/03	0-1 ft	Zinc	63		61.8	2346	nc	YES	NO
LL9SS-047-0001-SO	11/06/03	0-1 ft	Aluminum	8600		17700	7614	nc	NO	YES
LL9SS-047-0001-SO	11/06/03	0-1 ft	Arsenic	15		15.4	0.39	ca	NO	YES
LL9SS-047-0001-SO	11/06/03	0-1 ft	Iron	16000		23100	2346	nc	NO	YES
LL9SS-047-0001-SO	11/06/03	0-1 ft	Manganese	690		1450	176	nc	NO	YES
LL9SS-047-0001-SO	11/06/03	0-1 ft	Mercury	0.063		0.04	2.3	nc	YES	NO
LL9SS-047-0001-SO	11/06/03	0-1 ft	Sodium	1200		123	--[n]	0	YES	NO
LL9SS-047-0001-SO	11/06/03	0-1 ft	Vanadium	13		31.1	7.8	nc	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Aluminum	8400		17700	7614	nc	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Arsenic	7.3		15.4	0.39	ca	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Barium	110		88.4	538	nc	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Benzo(a)pyrene	0.24		--	0.062	ca	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Cadmium	2.7		0	3.7	nc	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Copper	170		17.7	313	nc	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Dibenzo(a,h)anthracene	0.13		--	0.062	ca	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Iron	20000		23100	2346	nc	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Lead	150		26.1	400	pbk	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Manganese	700		1450	176	nc	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Mercury	0.21		0.04	2.3	nc	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Nitrocellulose	3.2	B J	0	--	0	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Selenium	1.8	B	1.4	39	nc	YES	NO
LL9SS-068-0001-SO	12/12/03	0-1 ft	Sodium	170	B	123	--[n]	0	YES	NO



Table 4-2
Summary of Surface Soil Exceedances (0-1 ft bgs)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Surface Soil Background Criteria	Surface Soil Region 9 Criteria	SS PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SS-068-0001-SO	12/12/03	0-1 ft	Vanadium	16		31.1	7.8	nc	NO	YES
LL9SS-068-0001-SO	12/12/03	0-1 ft	Zinc	780		61.8	2346	nc	YES	NO

Notes:

-- - no background/PRG value is available for this analyte

blank cell indicates that the analyte was a non-detect (with a "U" qualifier) or analysis was not performed

PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL))

nc - non-cancer basis

ca - cancer basis

pbk - based on PBK modeling

mcl - based on CWA maximum contaminant level

max - ceiling limit

sat - soil saturation

[n] - nutrient

UC - unconsolidated

Inorganic Qualifiers

U - analyte was not detected at or above the stated limit

J - result is less than the RL, but greater than or equal to the MDL

B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL

InOrganic Flags

^ - instrument related QC exceed the upper or lower control limits

* - batch QC exceeds the upper or lower control limit

H - batch QC is greater than RL

Organic Qualifiers

U - analyte was not detected at or above the stated limit

J - result is an estimated value below the RL

Organic Flags

B - Method Blank: batch QC is greater than RL

* - batch QC exceeds the upper or lower control limit

^ - batch QC is greater than RL

a - concentration is below the method Reporting Limit (RL)

H - alternate peak selection upon analytical review



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-002-0001-SO		LL9SS-004-0001-SO		LL9SS-006-0001-SO		LL9SS-008-0001-SO		LL9SS-010-0001-SO		LL9SS-012-0001-SO		LL9SS-013-0001-SO		LL9SS-014-0001-DU	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc															0.13	U
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc															0.017	U
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc															0.017	U
		2,4,6-TNT	mg/Kg	0	16	ca															0.033	U
		2,4-Dinitrotoluene	mg/Kg	0	12	nc															0.035	U
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc															0.046	U
		2-Nitrotoluene	mg/Kg	0	0.88	ca															0.032	U
		3-Nitrotoluene	mg/Kg	0	73	nc															0.049	U
		4-Nitrotoluene	mg/Kg	0	12	ca															0.045	U
		HMX	mg/Kg	0	306	nc															0.11	U
		Nitrobenzene	mg/Kg	0	2	nc															0.022	U
		RDX	mg/Kg	0	4.4	ca															0.057	U
		Tetryl	mg/Kg	0	61	nc															0.042	U
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc															0.0021	U
		Aroclor 1221	mg/Kg	--	0.22	ca															0.0022	U
		Aroclor 1232	mg/Kg	--	0.22	ca															0.0022	U
		Aroclor 1242	mg/Kg	--	0.22	ca															0.0022	U
		Aroclor 1248	mg/Kg	--	0.22	ca															0.0022	U
		Aroclor 1254	mg/Kg	--	0.22	ca															0.0022	U
		Aroclor 1260	mg/Kg	--	0.22	ca															0.0015	U
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca															0.0004	U
		4,4'-DDE	mg/Kg	--	1.7	ca															0.00072	U
		4,4'-DDT	mg/Kg	--	1.7	ca															0.00041	U
		Aldrin	mg/Kg	--	0.029	ca															0.00014	U
		alpha-BHC	mg/Kg	--	0.09	sat															0.00018	U
		alpha-Chlordane	mg/Kg	--	1.6	ca															0.00013	U
		beta-BHC	mg/Kg	--	0.32	ca															0.00017	U
		delta-BHC	mg/Kg	--	--	0															0.00012	U
		Dieldrin	mg/Kg	--	0.030	ca															0.00038	U
		Endosulfan I	mg/Kg	--	37	nc															0.0003	U
		Endosulfan II	mg/Kg	--	37	nc															0.00031	U
		Endosulfan sulfate	mg/Kg	--	37	nc															0.00032	U
		Endrin	mg/Kg	--	1.8	nc															0.00048	U
		Endrin aldehyde	mg/Kg	--	--	0															0.00036	U
		Endrin ketone	mg/Kg	--	--	0															0.00032	U
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca															0.00025	U
		gamma-Chlordane	mg/Kg	--	1.6	ca															0.00017	U
		Heptachlor	mg/Kg	--	0.11	ca															0.0002	U
		Heptachlor epoxide	mg/Kg	--	0.053	ca															0.00015	U
		Methoxychlor	mg/Kg	--	31	nc															0.0025	U
		Toxaphene	mg/Kg	--	0	0															0.0051	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-002-0001-SO		LL9SS-004-0001-SO		LL9SS-006-0001-SO		LL9SS-008-0001-SO		LL9SS-010-0001-SO		LL9SS-012-0001-SO		LL9SS-013-0001-SO		LL9SS-014-0001-DU			
							Sample Date		12/04/03		12/04/03		12/11/03		12/10/03		12/08/03		12/10/03		12/10/03		12/11/03	
							Sample Depth		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft			
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier		
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0															1.3	B J		
	8330	Nitroglycerin	mg/Kg	0	--	0															0.5	U		
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc															0.25	U		
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc															0.065	U		
		1,2-Dichlorobenzene	mg/Kg	--	600	sat															0.088	U		
		1,3-Dichlorobenzene	mg/Kg	--	53	nc															0.088	U		
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca															0.079	U		
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca															0.083	U		
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc															0.042	U		
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc															0.052	U		
		2,4-Dichlorophenol	mg/Kg	--	18	nc															0.053	U		
		2,4-Dimethylphenol	mg/Kg	--	122	nc															0.067	U		
		2,4-Dinitrophenol	mg/Kg	--	12	nc															0.13	U		
		2-Chloronaphthalene	mg/Kg	--	494	nc															0.053	U		
		2-Chlorophenol	mg/Kg	--	6.3	nc															0.065	U		
		2-Methylnaphthalene	mg/Kg	--	--	0															0.0017	U		
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc															0.0093	U		
		2-Nitroaniline	mg/Kg	--	18.3	nc															0.038	U		
		2-Nitrophenol	mg/Kg	--	--	0															0.07	U		
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca															0.02	U		
		3-Nitroaniline	mg/Kg	--	1.8	nc															0.12	U		
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc															0.11	U		
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0															0.0034	U		
		4-Chloro-3-methylphenol	mg/Kg	--	--	0															0.042	U		
		4-Chloroaniline	mg/Kg	--	24	nc															0.11	U		
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0															0.004	U		
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc															0.0064	U		
		4-Nitroaniline	mg/Kg	--	23	ca															0.043	U		
		4-Nitrophenol	mg/Kg	--	--	0															0.091	U		
		Acenaphthene	mg/Kg	--	368	nc															0.0016	U		
		Acenaphthylene	mg/Kg	--	--	0															0.001	U		
		Anthracene	mg/Kg	--	2189	nc															0.00095	U		
		Benzo(a)anthracene	mg/Kg	--	0.62	ca															0.0012	U		
		Benzo(a)pyrene	mg/Kg	--	0.062	ca															0.0024	U		
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca															0.0023	U		
		Benzo(ghi)perylene	mg/Kg	--	--	0															0.0021	U		
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca															0.0031	U		
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0															0.0032	U		
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca															0.0022	U		
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca															0.047	J		
		Butyl benzyl phthalate	mg/Kg	--	1222	nc															0.0046	U		
		Carbazole	mg/Kg	--	24	ca															0.039	U		
		Chrysene	mg/Kg	--	62	ca															0.002	U		
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca															0.0024	U		



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-002-0001-SO		LL9SS-004-0001-SO		LL9SS-006-0001-SO		LL9SS-008-0001-SO		LL9SS-010-0001-SO		LL9SS-012-0001-SO		LL9SS-013-0001-SO		LL9SS-014-0001-DU			
							Sample Date		12/04/03		12/04/03		12/11/03		12/10/03		12/08/03		12/10/03		12/10/03		12/11/03	
							Sample Depth		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier		
		Dibenzofuran	mg/Kg	--	15	nc															0.003	U		
		Diethyl phthalate	mg/Kg	--	4888	nc															0.0041	U		
		Dimethyl phthalate	mg/Kg	--	100000	max															0.004	U		
		Di-n-butyl phthalate	mg/Kg	--	611	nc															0.022	U		
		Di-n-octyl phthalate	mg/Kg	--	244	nc															0.0097	U		
		Fluoranthene	mg/Kg	--	229	nc															0.0012	U		
		Fluorene	mg/Kg	--	275	nc															0.0018	U		
		Hexachlorobenzene	mg/Kg	--	0.30	ca															0.002	U		
		Hexachlorobutadiene	mg/Kg	--	6.2	ca															0.0037	U		
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc															0.06	U		
		Hexachloroethane	mg/Kg	--	35	ca															0.0037	U		
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca															0.0023	U		
		Isophorone	mg/Kg	--	512	ca															0.0027	U		
		Naphthalene	mg/Kg	--	5.6	nc															0.0019	U		
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca															0.0026	U		
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca															0.0032	U		
		Pentachlorophenol	mg/Kg	--	3.0	ca															0.11	U		
		Phenanthrene	mg/Kg	--	--	0															0.0011	U		
Phenol	mg/Kg	--	1833	nc															0.0018	U				
Pyrene	mg/Kg	--	232	nc															0.0022	U				
TAL Metals 6010B	6010B	Aluminum	mg/Kg	17700	7614	nc	8500		7100		9300		11000		7200		9000		6000		6800			
		Barium	mg/Kg	88.4	538	nc	65		44		92		60		27		71		29		35			
		Beryllium	mg/Kg	0.88	15	nc	0.62		0.41	B	0.49		0.46		0.27	B	0.5		0.31	B	0.23	B		
		Cadmium	mg/Kg	0.00	3.7	nc	0.084	U	0.091	U	2.9		0.091	U	0.11	B	0.51		0.08	U	0.08	U		
		Calcium	mg/Kg	15800	--	0	2100		1200		4900		1200		1100		22000		2500		1200			
		Chromium	mg/Kg	17.4	30	ca	13		10		14		14		10		13		8.4		8.9			
		Cobalt	mg/Kg	10.4	902	ca	8.7		6.2		9.2		8.5		6		7.1		7.9		5.7			
		Copper	mg/Kg	17.7	313	nc	20		19		18		13		21	H	25		22		20			
		Iron	mg/Kg	23100	2346	nc	21000		19000		16000	H	20000		21000		22000		24000		19000	H		
		Magnesium	mg/Kg	3030	--	0	1900		1800		2000		2000		1800		3000		2400		1700			
		Manganese	mg/Kg	1450	176	nc	630		390		830		640		280		600		370		340			
		Nickel	mg/Kg	21.1	156	nc	16		15		15		16		15		17		16		13			
		Potassium	mg/Kg	927	--	0	670		620		700		790		790		1100		610		560			
		Selenium	mg/Kg	1.4	39	nc	0.54	B	0.63	B	0.45	B	0.46	U	0.47	B	0.45	U	0.4	U	0.31	B		
		Silver	mg/Kg	0.00	39	nc	0.33	U	0.35	U	0.37	U	0.35	U	0.33	U	0.35	U	0.31	U	0.31	U		
		Sodium	mg/Kg	123	--	0	92	U	99	U	100	U	99	U	92	U	97	U	87	U	87	U		
		Vanadium	mg/Kg	31.1	7.8	nc	15		12		16		19		12		16		11		12			
		Zinc	mg/Kg	61.8	2346	nc	100		100		200		62		78		140		74		64			
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.49	U	0.41	U	0.51	U	0.5	U	0.46	U	0.95		0.47	U	0.45	U		
	7060A	Arsenic	mg/Kg	15.4	0.39	ca	9.7		15		9.9		10		15		15		11		14			
	7421	Lead	mg/Kg	26.1	400	pbk	64		45		140		18		11		44		15		15			
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.1		0.073		1.3		0.047		0.042		0.053		0.024		0.024			
	7841	Thallium	mg/Kg	0.00	0.52	nc	0.21	U	0.36	U	0.22	U	0.22	U	0.2	U	0.21	U	0.2	U	0.2	U		



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-002-0001-SO		LL9SS-004-0001-SO		LL9SS-006-0001-SO		LL9SS-008-0001-SO		LL9SS-010-0001-SO		LL9SS-012-0001-SO		LL9SS-013-0001-SO		LL9SS-014-0001-DU	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat															0.0012	U
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca															0.0011	U
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca															0.0012	U
		1,1-Dichloroethane	mg/Kg	--	51	nc															0.0011	U
		1,1-Dichloroethene	mg/Kg	--	12	nc															0.0015	U
		1,2-Dichloroethane	mg/Kg	--	0.28	ca															0.0011	U
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc															0.0024	U
		1,2-Dichloropropane	mg/Kg	--	0.34	ca															0.0011	U
		2-Butanone (MEK)	mg/Kg	--	2231	nc															0.0044	U
		2-Hexanone	mg/Kg	--	530	nc															0.0012	U
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc															0.0011	U
		Acetone	mg/Kg	--	1412	nc															0.0052	U
		Benzene	mg/Kg	--	0.64	ca															0.0012	U
		Bromodichloromethane	mg/Kg	--	0.82	ca															0.0011	U
		Bromoform	mg/Kg	--	62	ca															0.00084	U
		Bromomethane	mg/Kg	--	0.39	nc															0.0015	U
		Carbon disulfide	mg/Kg	--	36	nc															0.0013	U
		Carbon tetrachloride	mg/Kg	--	0.25	ca															0.0012	U
		Chlorobenzene	mg/Kg	--	15	nc															0.0012	U
		Chloroethane	mg/Kg	--	3.0	ca															0.0011	U
		Chloroform	mg/Kg	--	0.22	ca															0.0012	U
		Chloromethane	mg/Kg	--	4.7	nc															0.0012	U
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca															0.001	U
		Dibromochloromethane	mg/Kg	--	1.1	ca															0.00089	U
		Ethylbenzene	mg/Kg	--	395	sat															0.0012	U
		Methylene chloride	mg/Kg	--	9.1	ca															0.0033	U
		Styrene	mg/Kg	--	1700	sat															0.0012	U
		Tetrachloroethene	mg/Kg	--	0.48	ca															0.0013	U
		Toluene	mg/Kg	--	520	sat															0.0012	U
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca															0.00089	U
		Trichloroethene	mg/Kg	--	0.053	ca															0.0012	U
		Vinyl chloride	mg/Kg	--	0.079	ca															0.0012	U
		Xylenes (total)	mg/Kg	--	0.079	ca															0.0038	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-014-0001-SO		LL9SS-015-0001-SO		LL9SS-016-0001-DU		LL9SS-016-0001-SO		LL9SS-017-0001-SO		LL9SS-018-0001-SO		LL9SS-019-0001-SO		LL9SS-020-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc	0.18	U											0.18	U		
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc	0.017	U											0.017	U		
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc	0.018	U											0.018	U		
		2,4,6-TNT	mg/Kg	0	16	ca	0.033	U											0.033	U		
		2,4-Dinitrotoluene	mg/Kg	0	12	nc	0.035	U											0.035	U		
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc	0.047	U											0.047	U		
		2-Nitrotoluene	mg/Kg	0	0.88	ca	0.033	U											0.033	U		
		3-Nitrotoluene	mg/Kg	0	73	nc	0.049	U											0.05	U		
		4-Nitrotoluene	mg/Kg	0	12	ca	0.046	U											0.046	U		
		HMX	mg/Kg	0	306	nc	0.11	U											0.11	U		
		Nitrobenzene	mg/Kg	0	2	nc	0.022	U											0.022	U		
		RDX	mg/Kg	0	4.4	ca	0.058	U											0.058	U		
		Tetryl	mg/Kg	0	61	nc	0.043	U											0.043	U		
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc	0.002	U											0.0023	U		
		Aroclor 1221	mg/Kg	--	0.22	ca	0.0021	U											0.0025	U		
		Aroclor 1232	mg/Kg	--	0.22	ca	0.0021	U											0.0025	U		
		Aroclor 1242	mg/Kg	--	0.22	ca	0.0021	U											0.0025	U		
		Aroclor 1248	mg/Kg	--	0.22	ca	0.0021	U											0.0025	U		
		Aroclor 1254	mg/Kg	--	0.22	ca	0.0021	U											0.0025	U		
		Aroclor 1260	mg/Kg	--	0.22	ca	0.0015	U											0.0017	U		
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca	0.00038	U											0.00044	U		
		4,4'-DDE	mg/Kg	--	1.7	ca	0.00069	U											0.0008	U		
		4,4'-DDT	mg/Kg	--	1.7	ca	0.00039	U											0.00046	U		
		Aldrin	mg/Kg	--	0.029	ca	0.00014	U											0.00016	U		
		alpha-BHC	mg/Kg	--	0.09	sat	0.00017	U											0.0002	U		
		alpha-Chlordane	mg/Kg	--	1.6	ca	0.00013	U											0.00015	U		
		beta-BHC	mg/Kg	--	0.32	ca	0.00016	U											0.00018	U		
		delta-BHC	mg/Kg	--	--	0	0.00012	U											0.00014	U		
		Dieldrin	mg/Kg	--	0.030	ca	0.00036	U											0.00042	U		
		Endosulfan I	mg/Kg	--	37	nc	0.00029	U											0.00033	U		
		Endosulfan II	mg/Kg	--	37	nc	0.0003	U											0.00034	U		
		Endosulfan sulfate	mg/Kg	--	37	nc	0.00031	U											0.00036	U		
		Endrin	mg/Kg	--	1.8	nc	0.00046	U											0.00053	U		
		Endrin aldehyde	mg/Kg	--	--	0	0.00035	U											0.00041	U		
		Endrin ketone	mg/Kg	--	--	0	0.00031	U											0.00036	U		
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca	0.00024	U											0.00028	U		
		gamma-Chlordane	mg/Kg	--	1.6	ca	0.00016	U											0.00018	U		
		Heptachlor	mg/Kg	--	0.11	ca	0.00019	U											0.00022	U		
		Heptachlor epoxide	mg/Kg	--	0.053	ca	0.00015	U											0.00017	U		
		Methoxychlor	mg/Kg	--	31	nc	0.0024	U											0.0028	U		
		Toxaphene	mg/Kg	--	0	0	0.0049	U											0.0057	U		



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-014-0001-SO		LL9SS-015-0001-SO		LL9SS-016-0001-DU		LL9SS-016-0001-SO		LL9SS-017-0001-SO		LL9SS-018-0001-SO		LL9SS-019-0001-SO		LL9SS-020-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0	1.6	B J											1.8	B J		
	8330	Nitroglycerin	mg/Kg	0	--	0	0.5	U											0.5	U		
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc	0.25	U											0.25	U		
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc	0.065	U											0.073	U		
		1,2-Dichlorobenzene	mg/Kg	--	600	sat	0.087	U											0.098	U		
		1,3-Dichlorobenzene	mg/Kg	--	53	nc	0.087	U											0.098	U		
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca	0.078	U											0.088	U		
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca	0.083	U											0.093	U		
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc	0.042	U											0.047	U		
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc	0.052	U											0.058	U		
		2,4-Dichlorophenol	mg/Kg	--	18	nc	0.053	U											0.059	U		
		2,4-Dimethylphenol	mg/Kg	--	122	nc	0.066	U											0.074	U		
		2,4-Dinitrophenol	mg/Kg	--	12	nc	0.13	U											0.14	U		
		2-Chloronaphthalene	mg/Kg	--	494	nc	0.053	U											0.059	U		
		2-Chlorophenol	mg/Kg	--	6.3	nc	0.065	U											0.073	U		
		2-Methylnaphthalene	mg/Kg	--	--	0	0.0017	U											0.0019	U		
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc	0.0093	U											0.01	U		
		2-Nitroaniline	mg/Kg	--	18.3	nc	0.038	U											0.042	U		
		2-Nitrophenol	mg/Kg	--	--	0	0.07	U											0.078	U		
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca	0.02	U											0.022	U		
		3-Nitroaniline	mg/Kg	--	1.8	nc	0.12	U											0.14	U		
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc	0.1	U											0.12	U		
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0	0.0034	U											0.0038	U		
		4-Chloro-3-methylphenol	mg/Kg	--	--	0	0.042	U											0.047	U		
		4-Chloroaniline	mg/Kg	--	24	nc	0.11	U											0.12	U		
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0	0.004	U											0.0045	U		
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc	0.0064	U											0.0072	U		
		4-Nitroaniline	mg/Kg	--	23	ca	0.043	U											0.048	U		
		4-Nitrophenol	mg/Kg	--	--	0	0.091	U											0.1	U		
		Acenaphthene	mg/Kg	--	368	nc	0.0015	U											0.0017	U		
		Acenaphthylene	mg/Kg	--	--	0	0.001	U											0.0011	U		
		Anthracene	mg/Kg	--	2189	nc	0.00095	U											0.0011	U		
		Benzo(a)anthracene	mg/Kg	--	0.62	ca	0.0012	U											0.0072	J		
		Benzo(a)pyrene	mg/Kg	--	0.062	ca	0.0024	U											0.0068	J		
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca	0.0023	U											0.0073	J		
		Benzo(ghi)perylene	mg/Kg	--	--	0	0.0021	U											0.0053	J		
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca	0.0031	U											0.0049	J		
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0	0.0032	U											0.0036	U		
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca	0.0022	U											0.0025	U		
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca	0.15	J											0.061	J		
		Butyl benzyl phthalate	mg/Kg	--	1222	nc	0.0045	U											0.0051	U		
		Carbazole	mg/Kg	--	24	ca	0.039	U											0.043	U		
		Chrysene	mg/Kg	--	62	ca	0.002	U											0.0052	J		
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca	0.0024	U											0.0027	U		



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-014-0001-SO		LL9SS-015-0001-SO		LL9SS-016-0001-DU		LL9SS-016-0001-SO		LL9SS-017-0001-SO		LL9SS-018-0001-SO		LL9SS-019-0001-SO		LL9SS-020-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzofuran	mg/Kg	--	15	nc	0.003	U											0.0033	U		
		Diethyl phthalate	mg/Kg	--	4888	nc	0.0041	U											0.0046	U		
		Dimethyl phthalate	mg/Kg	--	100000	max	0.004	U											0.0045	U		
		Di-n-butyl phthalate	mg/Kg	--	611	nc	0.022	U											0.025	U		
		Di-n-octyl phthalate	mg/Kg	--	244	nc	0.0096	U											0.011	U		
		Fluoranthene	mg/Kg	--	229	nc	0.0012	U											0.01	J		
		Fluorene	mg/Kg	--	275	nc	0.0018	U											0.002	U		
		Hexachlorobenzene	mg/Kg	--	0.30	ca	0.002	U											0.0022	U		
		Hexachlorobutadiene	mg/Kg	--	6.2	ca	0.0036	U											0.0041	U		
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc	0.06	U											0.067	U		
		Hexachloroethane	mg/Kg	--	35	ca	0.0036	U											0.0041	U		
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca	0.0023	U											0.0026	U		
		Isophorone	mg/Kg	--	512	ca	0.0027	U											0.003	U		
		Naphthalene	mg/Kg	--	5.6	nc	0.0019	U											0.0021	U		
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca	0.0025	U											0.0028	U		
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca	0.0032	U											0.0036	U		
		Pentachlorophenol	mg/Kg	--	3.0	ca	0.11	U											0.12	U		
		Phenanthrene	mg/Kg	--	--	0	0.0011	U											0.0051	J		
		Phenol	mg/Kg	--	1833	nc	0.0018	U											0.002	U		
		Pyrene	mg/Kg	--	232	nc	0.0022	U											0.01	J		
TAL Metals 6010B	6010B	Aluminum	mg/Kg	17700	7614	nc	7900		17000		9100		9900		10000		11000		14000		7900	
		Barium	mg/Kg	88.4	538	nc	39		62		67		82		65		95		170		47	
		Beryllium	mg/Kg	0.88	15	nc	0.22	B	0.51		0.54		0.72		0.69		0.49		0.41	B	0.32	B
		Cadmium	mg/Kg	0.00	3.7	nc	0.083	U	0.093	U	0.14	B	0.17	B	0.09	U	0.16	B	0.085	U	0.71	
		Calcium	mg/Kg	15800	--	0	1100		2200		12000		20000		650		1200		900		1600	
		Chromium	mg/Kg	17.4	30	ca	11		23		12		10		14		14		17		11	
		Cobalt	mg/Kg	10.4	902	ca	7.2		8.3		6.1		7		8		10		8		7	
		Copper	mg/Kg	17.7	313	nc	22		24		18		19		12		12		19		31	H
		Iron	mg/Kg	23100	2346	nc	19000	H	31000	H	17000	H	18000	H	23000		19000		27000	H	16000	
		Magnesium	mg/Kg	3030	--	0	2000		3500		3300		3500		1800		2000		2600		1800	
		Manganese	mg/Kg	1450	176	nc	410		180		790		1100		690		920		210		460	
		Nickel	mg/Kg	21.1	156	nc	16		23		15		17		14		16		17		14	
		Potassium	mg/Kg	927	--	0	620		1500		970		1000		660		700		1000		660	
		Selenium	mg/Kg	1.4	39	nc	0.42	B	0.56	B	0.67	B	0.66	B	0.45	U	0.45	U	0.43	B	0.52	B
		Silver	mg/Kg	0.00	39	nc	0.32	U	0.36	U	0.32	U	0.31	U	0.35	U	0.35	U	0.33	U	0.33	U
		Sodium	mg/Kg	123	--	0	90	U	100	U	100		170		97	U	98	U	93	U	92	U
		Vanadium	mg/Kg	31.1	7.8	nc	13		28		13		12		21		19		22		14	
		Zinc	mg/Kg	61.8	2346	nc	72		64		72		70		52		69		51		85	
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.46	U	0.7	B	0.45	U	0.49	U	0.46	U	0.54	B	0.52	U	0.48	U
	7060A	Arsenic	mg/Kg	15.4	0.39	ca	15		24		14		11		11		10		17		13	
	7421	Lead	mg/Kg	26.1	400	pbk	43		18		35		38		16		40		20		33	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.044		0.061		0.11		0.096		0.037		0.09		0.36		0.74	
	7841	Thallium	mg/Kg	0.00	0.52	nc	0.2	U	0.23	U	0.2	U	0.21	U	1	U	0.22	U	0.23	U	0.21	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-014-0001-SO		LL9SS-015-0001-SO		LL9SS-016-0001-DU		LL9SS-016-0001-SO		LL9SS-017-0001-SO		LL9SS-018-0001-SO		LL9SS-019-0001-SO		LL9SS-020-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat	0.0012	U											0.0014	U		
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca	0.0011	U											0.0012	U		
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca	0.0012	U											0.0014	U		
		1,1-Dichloroethane	mg/Kg	--	51	nc	0.0011	U											0.0012	U		
		1,1-Dichloroethene	mg/Kg	--	12	nc	0.0015	U											0.0016	U		
		1,2-Dichloroethane	mg/Kg	--	0.28	ca	0.0011	U											0.0012	U		
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc	0.0023	U											0.0026	U		
		1,2-Dichloropropane	mg/Kg	--	0.34	ca	0.0011	U											0.0012	U		
		2-Butanone (MEK)	mg/Kg	--	2231	nc	0.0044	U											0.0049	U		
		2-Hexanone	mg/Kg	--	530	nc	0.0012	U											0.0014	U		
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc	0.0011	U											0.0012	U		
		Acetone	mg/Kg	--	1412	nc	0.0051	U											0.0058	U		
		Benzene	mg/Kg	--	0.64	ca	0.0012	U											0.0014	U		
		Bromodichloromethane	mg/Kg	--	0.82	ca	0.0011	U											0.0012	U		
		Bromoform	mg/Kg	--	62	ca	0.00084	U											0.00094	U		
		Bromomethane	mg/Kg	--	0.39	nc	0.0015	U*											0.0016	U*		
		Carbon disulfide	mg/Kg	--	36	nc	0.0013	U											0.0015	U		
		Carbon tetrachloride	mg/Kg	--	0.25	ca	0.0012	UM											0.0014	U		
		Chlorobenzene	mg/Kg	--	15	nc	0.0012	U											0.0014	U		
		Chloroethane	mg/Kg	--	3.0	ca	0.0011	U											0.0012	U		
		Chloroform	mg/Kg	--	0.22	ca	0.0012	U											0.0014	U		
		Chloromethane	mg/Kg	--	4.7	nc	0.0012	U											0.0014	U		
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.001	U											0.0012	U		
		Dibromochloromethane	mg/Kg	--	1.1	ca	0.00088	U											0.00099	U		
		Ethylbenzene	mg/Kg	--	395	sat	0.0012	U											0.0014	U		
		Methylene chloride	mg/Kg	--	9.1	ca	0.0032	U											0.0036	U		
		Styrene	mg/Kg	--	1700	sat	0.0012	U											0.0014	U		
		Tetrachloroethene	mg/Kg	--	0.48	ca	0.0013	U											0.0015	U		
		Toluene	mg/Kg	--	520	sat	0.0012	U											0.0014	U		
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.00088	U											0.00099	U		
		Trichloroethene	mg/Kg	--	0.053	ca	0.0012	U											0.0014	U		
		Vinyl chloride	mg/Kg	--	0.079	ca	0.0012	U											0.0014	U		
		Xylenes (total)	mg/Kg	--	0.079	ca	0.0038	U											0.0042	U		



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-021-0001-SO		LL9SS-022-0001-DU		LL9SS-022-0001-SO		LL9SS-023-0001-SO		LL9SS-024-0001-SO		LL9SS-025-0001-SO		LL9SS-026-0001-SO		LL9SS-027-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc											0.24	U				
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc											0.017	U				
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc											0.018	U				
		2,4,6-TNT	mg/Kg	0	16	ca											0.033	U				
		2,4-Dinitrotoluene	mg/Kg	0	12	nc											0.035	U				
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc											0.047	U				
		2-Nitrotoluene	mg/Kg	0	0.88	ca											0.033	U				
		3-Nitrotoluene	mg/Kg	0	73	nc											0.049	U				
		4-Nitrotoluene	mg/Kg	0	12	ca											0.046	U				
		HMX	mg/Kg	0	306	nc											0.11	U				
		Nitrobenzene	mg/Kg	0	2	nc											0.022	U				
		RDX	mg/Kg	0	4.4	ca											0.058	U				
		Tetryl	mg/Kg	0	61	nc											0.043	U				
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc											0.0024	U				
		Aroclor 1221	mg/Kg	--	0.22	ca											0.0025	U				
		Aroclor 1232	mg/Kg	--	0.22	ca											0.0025	U				
		Aroclor 1242	mg/Kg	--	0.22	ca											0.0025	U				
		Aroclor 1248	mg/Kg	--	0.22	ca											0.0025	U				
		Aroclor 1254	mg/Kg	--	0.22	ca											0.0025	U				
		Aroclor 1260	mg/Kg	--	0.22	ca											0.0017	U				
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca											0.0022	U				
		4,4'-DDE	mg/Kg	--	1.7	ca											0.0041	U				
		4,4'-DDT	mg/Kg	--	1.7	ca											0.0023	U				
		Aldrin	mg/Kg	--	0.029	ca											0.00081	U				
		alpha-BHC	mg/Kg	--	0.09	sat											0.001	U				
		alpha-Chlordane	mg/Kg	--	1.6	ca											0.00075	U				
		beta-BHC	mg/Kg	--	0.32	ca											0.00094	U				
		delta-BHC	mg/Kg	--	--	0											0.00069	U				
		Dieldrin	mg/Kg	--	0.030	ca											0.0021	U				
		Endosulfan I	mg/Kg	--	37	nc											0.0017	U				
		Endosulfan II	mg/Kg	--	37	nc											0.0017	U				
		Endosulfan sulfate	mg/Kg	--	37	nc											0.0018	U				
		Endrin	mg/Kg	--	1.8	nc											0.0027	U				
		Endrin aldehyde	mg/Kg	--	--	0											0.0021	U				
		Endrin ketone	mg/Kg	--	--	0											0.0018	U				
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca											0.0014	U				
		gamma-Chlordane	mg/Kg	--	1.6	ca											0.00094	U				
		Heptachlor	mg/Kg	--	0.11	ca											0.0011	U				
		Heptachlor epoxide	mg/Kg	--	0.053	ca											0.00087	U				
		Methoxychlor	mg/Kg	--	31	nc											0.014	U				
		Toxaphene	mg/Kg	--	0	0											0.029	U				



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-021-0001-SO		LL9SS-022-0001-DU		LL9SS-022-0001-SO		LL9SS-023-0001-SO		LL9SS-024-0001-SO		LL9SS-025-0001-SO		LL9SS-026-0001-SO		LL9SS-027-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0											1.6	B J				
	8330	Nitroglycerin	mg/Kg	0	--	0											0.5	U				
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc											0.25	U				
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc											0.073	U				
		1,2-Dichlorobenzene	mg/Kg	--	600	sat											0.097	U				
		1,3-Dichlorobenzene	mg/Kg	--	53	nc											0.097	U				
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca											0.087	U				
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca											0.092	U				
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc											0.047	U				
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc											0.058	U				
		2,4-Dichlorophenol	mg/Kg	--	18	nc											0.059	U				
		2,4-Dimethylphenol	mg/Kg	--	122	nc											0.074	U				
		2,4-Dinitrophenol	mg/Kg	--	12	nc											0.14	U				
		2-Chloronaphthalene	mg/Kg	--	494	nc											0.059	U				
		2-Chlorophenol	mg/Kg	--	6.3	nc											0.073	U				
		2-Methylnaphthalene	mg/Kg	--	--	0											0.0018	U				
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc											0.01	U				
		2-Nitroaniline	mg/Kg	--	18.3	nc											0.042	U				
		2-Nitrophenol	mg/Kg	--	--	0											0.077	U				
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca											0.022	U				
		3-Nitroaniline	mg/Kg	--	1.8	nc											0.14	U				
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc											0.12	U				
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0											0.0038	U				
		4-Chloro-3-methylphenol	mg/Kg	--	--	0											0.047	U				
		4-Chloroaniline	mg/Kg	--	24	nc											0.12	U				
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0											0.0044	U				
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc											0.0071	U				
		4-Nitroaniline	mg/Kg	--	23	ca											0.048	U				
		4-Nitrophenol	mg/Kg	--	--	0											0.1	U				
		Acenaphthene	mg/Kg	--	368	nc											0.0017	U				
		Acenaphthylene	mg/Kg	--	--	0											0.0011	U				
		Anthracene	mg/Kg	--	2189	nc											0.0011	U				
		Benzo(a)anthracene	mg/Kg	--	0.62	ca											0.012	Ja				
		Benzo(a)pyrene	mg/Kg	--	0.062	ca											0.012	Ja				
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca											0.011	Ja				
		Benzo(ghi)perylene	mg/Kg	--	--	0											0.011	Ja				
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca											0.01	Ja				
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0											0.0036	U				
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca											0.0025	U				
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca											0.012	U				
		Butyl benzyl phthalate	mg/Kg	--	1222	nc											0.005	U				
		Carbazole	mg/Kg	--	24	ca											0.043	U				
		Chrysene	mg/Kg	--	62	ca											0.012	Ja				
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca											0.03	Ja				



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-021-0001-SO		LL9SS-022-0001-DU		LL9SS-022-0001-SO		LL9SS-023-0001-SO		LL9SS-024-0001-SO		LL9SS-025-0001-SO		LL9SS-026-0001-SO		LL9SS-027-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzofuran	mg/Kg	--	15	nc											0.0033	U				
		Diethyl phthalate	mg/Kg	--	4888	nc											0.0046	U				
		Dimethyl phthalate	mg/Kg	--	100000	max											0.0044	U				
		Di-n-butyl phthalate	mg/Kg	--	611	nc											0.025	U				
		Di-n-octyl phthalate	mg/Kg	--	244	nc											0.011	U				
		Fluoranthene	mg/Kg	--	229	nc											0.019	Ja				
		Fluorene	mg/Kg	--	275	nc											0.002	U				
		Hexachlorobenzene	mg/Kg	--	0.30	ca											0.0022	U				
		Hexachlorobutadiene	mg/Kg	--	6.2	ca											0.0041	U				
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc											0.066	U				
		Hexachloroethane	mg/Kg	--	35	ca											0.0041	U				
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca											0.024	Ja				
		Isophorone	mg/Kg	--	512	ca											0.003	U				
		Naphthalene	mg/Kg	--	5.6	nc											0.0021	U				
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca											0.0028	U				
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca											0.0036	U				
		Pentachlorophenol	mg/Kg	--	3.0	ca											0.12	U				
		Phenanthrene	mg/Kg	--	--	0											0.01	Ja				
		Phenol	mg/Kg	--	1833	nc											0.002	U				
		Pyrene	mg/Kg	--	232	nc											0.019	Ja				
TAL Metals 6010B	6010B	Aluminum	mg/Kg	17700	7614	nc	13000		9600		8800		20000		8000		12000		13000		10000	
		Barium	mg/Kg	88.4	538	nc	55		55		38		66		49		57		100		81	
		Beryllium	mg/Kg	0.88	15	nc	0.37	B	0.49		0.36	B	0.43	B	0.31	B	0.35	B	0.6		0.84	
		Cadmium	mg/Kg	0.00	3.7	nc	0.5		0.09	U	0.086	U	0.11	B	0.42		0.096	U	0.095	U	0.1	U
		Calcium	mg/Kg	15800	--	0	2700		4300		1300		5300		1100		1100		1200		430	
		Chromium	mg/Kg	17.4	30	ca	18		12		12		21		110		16		17		11	
		Cobalt	mg/Kg	10.4	902	ca	6		6.9		8.6		6.5		6.6		8		11		18	
		Copper	mg/Kg	17.7	313	nc	19	H	21		17		23	H	13		9.9		12		7.1	
		Iron	mg/Kg	23100	2346	nc	27000		22000		20000		26000		14000		24000		23000		25000	
		Magnesium	mg/Kg	3030	--	0	2200		2300		1900		3200		1500		2300		2300		1300	
		Manganese	mg/Kg	1450	176	nc	260		480		150		230		500		420		860		3800	
		Nickel	mg/Kg	21.1	156	nc	13		15		15		16		9.9		12		16		12	
		Potassium	mg/Kg	927	--	0	1000		750		750		1500		560		790		860		540	
		Selenium	mg/Kg	1.4	39	nc	0.44	U	0.45	U	0.43	U	0.89	B	0.5	U	0.48	U	0.48	U	0.71	B
		Silver	mg/Kg	0.00	39	nc	0.34	U	0.35	U	0.33	U	0.38	U	0.39	U	0.37	U	0.37	U	0.39	U
		Sodium	mg/Kg	123	--	0	94	U	98	U	93	U	110	U	110	U	100	U	100	U	110	U
		Vanadium	mg/Kg	31.1	7.8	nc	22		15		15		28		15		23		23		23	
		Zinc	mg/Kg	61.8	2346	nc	72		69		48		57		210		46		71		47	
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.5	U	0.52	U	0.49	U	0.55	U	0.85		0.54	U	1.2		0.55	U
	7060A	Arsenic	mg/Kg	15.4	0.39	ca	16		12		13		21		10		8.4		8.3		11	
	7421	Lead	mg/Kg	26.1	400	pbk	35		48		20		50		320		25		28		16	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.045		0.2		0.11		0.25		0.057		0.032		0.045		0.041	
	7841	Thallium	mg/Kg	0.00	0.52	nc	0.22	U	0.23	U	0.21	U	0.24	U	0.23	U	0.23	U	0.23	U	0.24	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-021-0001-SO		LL9SS-022-0001-DU		LL9SS-022-0001-SO		LL9SS-023-0001-SO		LL9SS-024-0001-SO		LL9SS-025-0001-SO		LL9SS-026-0001-SO		LL9SS-027-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat											0.0014	U				
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca											0.0012	U				
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca											0.0014	U				
		1,1-Dichloroethane	mg/Kg	--	51	nc											0.0013	U				
		1,1-Dichloroethene	mg/Kg	--	12	nc											0.0016	U				
		1,2-Dichloroethane	mg/Kg	--	0.28	ca											0.0012	U				
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc											0.0026	U				
		1,2-Dichloropropane	mg/Kg	--	0.34	ca											0.0013	U				
		2-Butanone (MEK)	mg/Kg	--	2231	nc											0.0049	U				
		2-Hexanone	mg/Kg	--	530	nc											0.0014	U				
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc											0.0013	U				
		Acetone	mg/Kg	--	1412	nc											0.0058	U				
		Benzene	mg/Kg	--	0.64	ca											0.0014	U				
		Bromodichloromethane	mg/Kg	--	0.82	ca											0.0012	U				
		Bromoform	mg/Kg	--	62	ca											0.00094	U				
		Bromomethane	mg/Kg	--	0.39	nc											0.0016	U				
		Carbon disulfide	mg/Kg	--	36	nc											0.0015	U				
		Carbon tetrachloride	mg/Kg	--	0.25	ca											0.0014	U				
		Chlorobenzene	mg/Kg	--	15	nc											0.0014	U				
		Chloroethane	mg/Kg	--	3.0	ca											0.0013	U				
		Chloroform	mg/Kg	--	0.22	ca											0.0014	U				
		Chloromethane	mg/Kg	--	4.7	nc											0.0014	U				
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca											0.0012	U				
		Dibromochloromethane	mg/Kg	--	1.1	ca											0.00099	U				
		Ethylbenzene	mg/Kg	--	395	sat											0.0014	U				
		Methylene chloride	mg/Kg	--	9.1	ca											0.0036	U				
		Styrene	mg/Kg	--	1700	sat											0.0014	U				
		Tetrachloroethene	mg/Kg	--	0.48	ca											0.0015	U				
		Toluene	mg/Kg	--	520	sat											0.0014	U				
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca											0.00099	U				
		Trichloroethene	mg/Kg	--	0.053	ca											0.0014	U				
		Vinyl chloride	mg/Kg	--	0.079	ca											0.0014	U				
		Xylenes (total)	mg/Kg	--	0.079	ca											0.0043	U				



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-028-0001-DU		LL9SS-028-0001-SO		LL9SS-029-0001-SO		LL9SS-030-0001-SO		LL9SS-031-0001-SO		LL9SS-032-0001-SO		LL9SS-033-0001-SO		LL9SS-034-0001-DU	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc													0.16	U	0.17	U
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc													0.017	U	0.017	U
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc													0.018	U	0.018	U
		2,4,6-TNT	mg/Kg	0	16	ca													0.034	U	0.034	U
		2,4-Dinitrotoluene	mg/Kg	0	12	nc													0.035	U	0.035	U
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc													0.047	U*	0.047	U*
		2-Nitrotoluene	mg/Kg	0	0.88	ca													0.033	U	0.033	U
		3-Nitrotoluene	mg/Kg	0	73	nc													0.05	U	0.05	U
		4-Nitrotoluene	mg/Kg	0	12	ca													0.046	U	0.046	U
		HMX	mg/Kg	0	306	nc													0.11	U	0.11	U
		Nitrobenzene	mg/Kg	0	2	nc													0.022	U	0.022	U
		RDX	mg/Kg	0	4.4	ca													0.058	U	0.058	U
		Tetryl	mg/Kg	0	61	nc													0.043	U	0.043	U
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc													0.0026	U	0.0023	U
		Aroclor 1221	mg/Kg	--	0.22	ca													0.0027	U	0.0024	U
		Aroclor 1232	mg/Kg	--	0.22	ca													0.0027	U	0.0024	U
		Aroclor 1242	mg/Kg	--	0.22	ca													0.0027	U	0.0024	U
		Aroclor 1248	mg/Kg	--	0.22	ca													0.0027	U	0.0024	U
		Aroclor 1254	mg/Kg	--	0.22	ca													0.0027	U	0.0024	U
		Aroclor 1260	mg/Kg	--	0.22	ca													0.0019	U	0.0017	U
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca													0.00049	U	0.00043	U
		4,4'-DDE	mg/Kg	--	1.7	ca													0.00089	U	0.00077	U
		4,4'-DDT	mg/Kg	--	1.7	ca													0.00051	U	0.00044	U
		Aldrin	mg/Kg	--	0.029	ca													0.00018	U	0.00015	U
		alpha-BHC	mg/Kg	--	0.09	sat													0.00022	U	0.00019	U
		alpha-Chlordane	mg/Kg	--	1.6	ca													0.00016	U	0.00014	U
		beta-BHC	mg/Kg	--	0.32	ca													0.00021	U	0.00018	U
		delta-BHC	mg/Kg	--	--	0													0.00015	U	0.00013	U
		Dieldrin	mg/Kg	--	0.030	ca													0.00047	U	0.0004	U
		Endosulfan I	mg/Kg	--	37	nc													0.00037	U	0.00032	U
		Endosulfan II	mg/Kg	--	37	nc													0.00038	U	0.00033	U
		Endosulfan sulfate	mg/Kg	--	37	nc													0.0004	U	0.00035	U
		Endrin	mg/Kg	--	1.8	nc													0.00059	U	0.00051	U
		Endrin aldehyde	mg/Kg	--	--	0													0.00045	U	0.00039	U
		Endrin ketone	mg/Kg	--	--	0													0.0004	U	0.00035	U
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca													0.00032	U	0.00027	U
		gamma-Chlordane	mg/Kg	--	1.6	ca													0.00021	U	0.00018	U
		Heptachlor	mg/Kg	--	0.11	ca													0.00025	U	0.00021	U
		Heptachlor epoxide	mg/Kg	--	0.053	ca													0.00019	U	0.00017	U
		Methoxychlor	mg/Kg	--	31	nc													0.0032	U	0.0027	U
		Toxaphene	mg/Kg	--	0	0													0.0063	U	0.0055	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-028-0001-DU		LL9SS-028-0001-SO		LL9SS-029-0001-SO		LL9SS-030-0001-SO		LL9SS-031-0001-SO		LL9SS-032-0001-SO		LL9SS-033-0001-SO		LL9SS-034-0001-DU	
							Sample Date		12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		11/11/03		11/11/03	
							Sample Depth		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0													1.4	B J	2.5	U
	8330	Nitroglycerin	mg/Kg	0	--	0													0.5	U	0.5	U
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc													0.25	U	0.25	U
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc													0.08	U	0.071	U
		1,2-Dichlorobenzene	mg/Kg	--	600	sat													0.11	U	0.095	U
		1,3-Dichlorobenzene	mg/Kg	--	53	nc													0.11	U	0.095	U
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca													0.096	U	0.085	U
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca													0.1	U	0.09	U
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc													0.051	U	0.046	U
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc													0.064	U	0.057	U
		2,4-Dichlorophenol	mg/Kg	--	18	nc													0.065	U	0.058	U
		2,4-Dimethylphenol	mg/Kg	--	122	nc													0.081	U	0.072	U
		2,4-Dinitrophenol	mg/Kg	--	12	nc													0.15	U	0.14	U
		2-Chloronaphthalene	mg/Kg	--	494	nc													0.065	U	0.058	U
		2-Chlorophenol	mg/Kg	--	6.3	nc													0.08	U	0.071	U
		2-Methylnaphthalene	mg/Kg	--	--	0													0.002	U	0.0018	U
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc													0.011	U	0.01	U
		2-Nitroaniline	mg/Kg	--	18.3	nc													0.046	U	0.041	U
		2-Nitrophenol	mg/Kg	--	--	0													0.085	U	0.076	U
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca													0.024	U	0.022	U
		3-Nitroaniline	mg/Kg	--	1.8	nc													0.15	U	0.13	U
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc													0.13	U*	0.11	U*
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0													0.0042	U	0.0037	U
		4-Chloro-3-methylphenol	mg/Kg	--	--	0													0.051	U	0.046	U
		4-Chloroaniline	mg/Kg	--	24	nc													0.14	U	0.12	U
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0													0.0049	U	0.0043	U
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc													0.0078	U	0.007	U
		4-Nitroaniline	mg/Kg	--	23	ca													0.053	U	0.047	U
		4-Nitrophenol	mg/Kg	--	--	0													0.11	U	0.099	U
		Acenaphthene	mg/Kg	--	368	nc													0.0019	U	0.011	Ja
		Acenaphthylene	mg/Kg	--	--	0													0.0012	U	0.0011	U
		Anthracene	mg/Kg	--	2189	nc													0.0012	U	0.022	Ja
		Benzo(a)anthracene	mg/Kg	--	0.62	ca													0.017	Ja	0.079	
		Benzo(a)pyrene	mg/Kg	--	0.062	ca													0.025	Ja	0.073	
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca													0.031	Ja	0.098	
		Benzo(ghi)perylene	mg/Kg	--	--	0													0.02	Ja	0.056	
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca													0.018	Ja	0.058	M
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0													0.0039	U	0.0035	U
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca													0.0027	U	0.0024	U
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca													0.013	U	0.014	Ja
		Butyl benzyl phthalate	mg/Kg	--	1222	nc													0.0055	U	0.0049	U
		Carbazole	mg/Kg	--	24	ca													0.047	U	0.042	U
		Chrysene	mg/Kg	--	62	ca													0.025	Ja	0.12	
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca													0.01	Ja	0.027	Ja



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-028-0001-DU		LL9SS-028-0001-SO		LL9SS-029-0001-SO		LL9SS-030-0001-SO		LL9SS-031-0001-SO		LL9SS-032-0001-SO		LL9SS-033-0001-SO		LL9SS-034-0001-DU	
							Sample Date		12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		11/11/03	
							Sample Depth		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzofuran	mg/Kg	--	15	nc													0.0036	U	0.0032	U
		Diethyl phthalate	mg/Kg	--	4888	nc													0.005	U	0.0044	U
		Dimethyl phthalate	mg/Kg	--	100000	max													0.0049	U	0.0043	U
		Di-n-butyl phthalate	mg/Kg	--	611	nc													0.027	U	0.024	U
		Di-n-octyl phthalate	mg/Kg	--	244	nc													0.012	U	0.01	U
		Fluoranthene	mg/Kg	--	229	nc													0.035	Ja	0.2	
		Fluorene	mg/Kg	--	275	nc													0.0022	U	0.0085	Ja
		Hexachlorobenzene	mg/Kg	--	0.30	ca													0.0024	U	0.0022	U
		Hexachlorobutadiene	mg/Kg	--	6.2	ca													0.0045	U	0.004	U
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc													0.073	U	0.065	U
		Hexachloroethane	mg/Kg	--	35	ca													0.0045	U	0.004	U
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca													0.018	Ja	0.056	
		Isophorone	mg/Kg	--	512	ca													0.0032	U	0.0029	U
		Naphthalene	mg/Kg	--	5.6	nc													0.0023	U	0.002	U
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca													0.0031	U	0.0028	U
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca													0.0039	U*	0.0035	U*
		Pentachlorophenol	mg/Kg	--	3.0	ca													0.14	U	0.12	U
		Phenanthrene	mg/Kg	--	--	0													0.02	Ja	0.1	
		Phenol	mg/Kg	--	1833	nc													0.0022	U	0.0019	U
		Pyrene	mg/Kg	--	232	nc													0.039	Ja	0.16	
TAL Metals 6010B	6010B	Aluminum	mg/Kg	17700	7614	nc	10000		10000		9400		6800		6400		8200		13000		7000	
		Barium	mg/Kg	88.4	538	nc	52		64		80		52		72		45		77		41	
		Beryllium	mg/Kg	0.88	15	nc	0.51		0.57	B	0.75		0.61		0.74		0.58		0.55		0.46	
		Cadmium	mg/Kg	0.00	3.7	nc	0.09	U	0.12	U	0.1	U	0.096	U	0.31		0.088	U	0.11	U	0.086	U
		Calcium	mg/Kg	15800	--	0	730		1100		530		530		2600		600		990		900	
		Chromium	mg/Kg	17.4	30	ca	12		12		11		9.2		11		12		15		12	
		Cobalt	mg/Kg	10.4	902	ca	5.1		5.3		8.1		5.6		13		7.5		11		6.1	
		Copper	mg/Kg	17.7	313	nc	6.4		7.2		8.9		8.1		8.5		13		9.3		33	
		Iron	mg/Kg	23100	2346	nc	23000		21000		16000		20000		76000		22000		23000		32000	
		Magnesium	mg/Kg	3030	--	0	1300		1400		1600		1200		1300		1700		1900		1500	
		Manganese	mg/Kg	1450	176	nc	330		460		870		650		2000		480		1200		420	
		Nickel	mg/Kg	21.1	156	nc	7.9		8.9		13		10		18		13		12		16	
		Potassium	mg/Kg	927	--	0	570		580		600		450		540		660		770		550	
		Selenium	mg/Kg	1.4	39	nc	0.51	B	0.92	B	0.79	B	0.48	U	0.45	U	0.52	B	1	B	0.61	B
		Silver	mg/Kg	0.00	39	nc	0.35	U	0.45	U	0.4	U	0.37	U	0.35	U	0.34	U	0.41	U	0.33	U
		Sodium	mg/Kg	123	--	0	98	U	130	U	110	U	100	U	97	U	95	U	1000		660	
		Vanadium	mg/Kg	31.1	7.8	nc	22		21		16		14		19		15		25		15	
		Zinc	mg/Kg	61.8	2346	nc	36		43		56		48		75		47		86		79	
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.53	U	0.62	U	0.54	U	0.52	U	0.48	U	0.49	U	0.54	U	0.49	U
	7060A	Arsenic	mg/Kg	15.4	0.39	ca	9.4		15		7.4		4.2		6.9		7.4		9.2		10	
	7421	Lead	mg/Kg	26.1	400	pbk	13		13		17		13		28		12		25		38	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.2		0.24		0.088		0.034		0.032		0.028		0.033		17	
	7841	Thallium	mg/Kg	0.00	0.52	nc	0.23	U	0.27	U	0.23	U	0.22	U	1	U	0.21	U	1.2	U	0.21	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-028-0001-DU		LL9SS-028-0001-SO		LL9SS-029-0001-SO		LL9SS-030-0001-SO		LL9SS-031-0001-SO		LL9SS-032-0001-SO		LL9SS-033-0001-SO		LL9SS-034-0001-DU	
							Sample Date		12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		11/11/03		11/11/03	
							Sample Depth		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat													0.00084	U*	0.00073	U*
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca													0.00088	U	0.00077	U
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca													0.00098	U	0.00086	U
		1,1-Dichloroethane	mg/Kg	--	51	nc													0.0012	U	0.0011	U
		1,1-Dichloroethene	mg/Kg	--	12	nc													0.0014	U	0.0012	U
		1,2-Dichloroethane	mg/Kg	--	0.28	ca													0.0008	U	0.0007	U
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc													0.0026	U	0.0023	U
		1,2-Dichloropropane	mg/Kg	--	0.34	ca													0.0013	U	0.0012	U
		2-Butanone (MEK)	mg/Kg	--	2231	nc													0.0058	U	0.0051	U
		2-Hexanone	mg/Kg	--	530	nc													0.0023	U	0.002	U
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc													0.0041	U	0.0036	U
		Acetone	mg/Kg	--	1412	nc													0.0056	U*	0.0049	U*
		Benzene	mg/Kg	--	0.64	ca													0.00091	U	0.0008	U
		Bromodichloromethane	mg/Kg	--	0.82	ca													0.00093	U	0.00082	U
		Bromoform	mg/Kg	--	62	ca													0.0012	U	0.0011	U
		Bromomethane	mg/Kg	--	0.39	nc													0.004	U*	0.0035	U*
		Carbon disulfide	mg/Kg	--	36	nc													0.0027	U	0.0024	U
		Carbon tetrachloride	mg/Kg	--	0.25	ca													0.0011	U	0.001	U
		Chlorobenzene	mg/Kg	--	15	nc													0.0012	U	0.0011	U
		Chloroethane	mg/Kg	--	3.0	ca													0.0022	U	0.0019	U
		Chloroform	mg/Kg	--	0.22	ca													0.00085	U	0.00075	U
		Chloromethane	mg/Kg	--	4.7	nc													0.0013	U	0.0011	U
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca													0.0011	U	0.00095	U
		Dibromochloromethane	mg/Kg	--	1.1	ca													0.00095	U	0.00083	U
		Ethylbenzene	mg/Kg	--	395	sat													0.0015	U	0.0013	U
		Methylene chloride	mg/Kg	--	9.1	ca													0.0025	U	0.0022	U
		Styrene	mg/Kg	--	1700	sat													0.0014	U	0.0012	U
		Tetrachloroethene	mg/Kg	--	0.48	ca													0.00092	U	0.00081	U
		Toluene	mg/Kg	--	520	sat													0.0014	U	0.0012	U
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca													0.0012	U	0.001	U
		Trichloroethene	mg/Kg	--	0.053	ca													0.00081	U	0.00071	U
		Vinyl chloride	mg/Kg	--	0.079	ca													0.001	U	0.00089	U
		Xylenes (total)	mg/Kg	--	0.079	ca													0.004	U	0.0035	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-034-0001-SO		LL9SS-035-0001-SO		LL9SS-036-0001-DU		LL9SS-036-0001-SO		LL9SS-037-0001-SO		LL9SS-038-0001-SO		LL9SS-039-0001-DU		LL9SS-039-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc	0.1	U														
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc	0.017	U	0.017	U					0.018	U			0.018	U	0.018	U
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc	0.018	U	0.018	U					0.018	U			0.018	U	0.018	U
		2,4,6-TNT	mg/Kg	0	16	ca	0.033	U	0.034	U					0.034	U			0.034	U	0.034	U
		2,4-Dinitrotoluene	mg/Kg	0	12	nc	0.035	U	0.035	U					0.036	U			0.036	U	0.036	U
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc	0.047	U*	0.047	U*					0.048	U*			0.048	U*	0.048	U*
		2-Nitrotoluene	mg/Kg	0	0.88	ca	0.033	U	0.033	U					0.033	U			0.033	U	0.033	U
		3-Nitrotoluene	mg/Kg	0	73	nc	0.05	U	0.05	U					0.05	U			0.05	U	0.05	U
		4-Nitrotoluene	mg/Kg	0	12	ca	0.046	U	0.046	U					0.047	U			0.047	U	0.047	U
		HMX	mg/Kg	0	306	nc	0.11	U	0.11	U					0.11	U			0.11	U	0.11	U
		Nitrobenzene	mg/Kg	0	2	nc	0.022	U	0.022	U					0.022	U			0.022	U	0.022	U
		RDX	mg/Kg	0	4.4	ca	0.058	U	0.058	U					0.059	U			0.059	U	0.059	U
		Tetryl	mg/Kg	0	61	nc	0.043	U	0.043	U					0.043	U			0.043	U	0.043	U
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc	0.0023	U														
		Aroclor 1221	mg/Kg	--	0.22	ca	0.0024	U														
		Aroclor 1232	mg/Kg	--	0.22	ca	0.0024	U														
		Aroclor 1242	mg/Kg	--	0.22	ca	0.0024	U														
		Aroclor 1248	mg/Kg	--	0.22	ca	0.0024	U														
		Aroclor 1254	mg/Kg	--	0.22	ca	0.0024	U														
		Aroclor 1260	mg/Kg	--	0.22	ca	0.0017	U														
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca	0.00043	U														
		4,4'-DDE	mg/Kg	--	1.7	ca	0.00077	U														
		4,4'-DDT	mg/Kg	--	1.7	ca	0.00044	U														
		Aldrin	mg/Kg	--	0.029	ca	0.00015	U														
		alpha-BHC	mg/Kg	--	0.09	sat	0.00019	U														
		alpha-Chlordane	mg/Kg	--	1.6	ca	0.00014	U														
		beta-BHC	mg/Kg	--	0.32	ca	0.00018	U														
		delta-BHC	mg/Kg	--	--	0	0.00013	U														
		Dieldrin	mg/Kg	--	0.030	ca	0.0004	U														
		Endosulfan I	mg/Kg	--	37	nc	0.00032	U														
		Endosulfan II	mg/Kg	--	37	nc	0.00033	U														
		Endosulfan sulfate	mg/Kg	--	37	nc	0.00034	U														
		Endrin	mg/Kg	--	1.8	nc	0.00051	U														
		Endrin aldehyde	mg/Kg	--	--	0	0.00039	U														
		Endrin ketone	mg/Kg	--	--	0	0.00034	U														
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca	0.00027	U														
		gamma-Chlordane	mg/Kg	--	1.6	ca	0.00018	U														
		Heptachlor	mg/Kg	--	0.11	ca	0.00021	U														
		Heptachlor epoxide	mg/Kg	--	0.053	ca	0.00017	U														
		Methoxychlor	mg/Kg	--	31	nc	0.0027	U														
		Toxaphene	mg/Kg	--	0	0	0.0055	U														



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-034-0001-SO		LL9SS-035-0001-SO		LL9SS-036-0001-DU		LL9SS-036-0001-SO		LL9SS-037-0001-SO		LL9SS-038-0001-SO		LL9SS-039-0001-DU		LL9SS-039-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0	1.1	B														
	8330	Nitroglycerin	mg/Kg	0	--	0	0.5	U														
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc	0.089	J														
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc	0.071	U														
		1,2-Dichlorobenzene	mg/Kg	--	600	sat	0.094	U														
		1,3-Dichlorobenzene	mg/Kg	--	53	nc	0.094	U														
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca	0.085	U														
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca	0.09	U														
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc	0.045	U														
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc	0.056	U														
		2,4-Dichlorophenol	mg/Kg	--	18	nc	0.057	U														
		2,4-Dimethylphenol	mg/Kg	--	122	nc	0.072	U														
		2,4-Dinitrophenol	mg/Kg	--	12	nc	0.14	U														
		2-Chloronaphthalene	mg/Kg	--	494	nc	0.057	U														
		2-Chlorophenol	mg/Kg	--	6.3	nc	0.071	U														
		2-Methylnaphthalene	mg/Kg	--	--	0	0.0018	U														
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc	0.01	U														
		2-Nitroaniline	mg/Kg	--	18.3	nc	0.041	U														
		2-Nitrophenol	mg/Kg	--	--	0	0.075	U														
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca	0.022	U														
		3-Nitroaniline	mg/Kg	--	1.8	nc	0.13	U														
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc	0.11	U*														
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0	0.0037	U														
		4-Chloro-3-methylphenol	mg/Kg	--	--	0	0.045	U														
		4-Chloroaniline	mg/Kg	--	24	nc	0.12	U														
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0	0.0043	U														
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc	0.0069	U														
		4-Nitroaniline	mg/Kg	--	23	ca	0.047	U														
		4-Nitrophenol	mg/Kg	--	--	0	0.098	U														
		Acenaphthene	mg/Kg	--	368	nc	0.0017	U														
		Acenaphthylene	mg/Kg	--	--	0	0.0011	U														
		Anthracene	mg/Kg	--	2189	nc	0.001	U														
		Benzo(a)anthracene	mg/Kg	--	0.62	ca	0.026	Ja														
		Benzo(a)pyrene	mg/Kg	--	0.062	ca	0.023	Ja														
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca	0.032	Ja														
		Benzo(ghi)perylene	mg/Kg	--	--	0	0.017	Ja														
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca	0.025	Ja														
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0	0.0035	U														
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca	0.0024	U														
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca	0.011	U														
		Butyl benzyl phthalate	mg/Kg	--	1222	nc	0.0049	U														
		Carbazole	mg/Kg	--	24	ca	0.042	U														
		Chrysene	mg/Kg	--	62	ca	0.03	Ja														
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca	0.0026	U														



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-034-0001-SO		LL9SS-035-0001-SO		LL9SS-036-0001-DU		LL9SS-036-0001-SO		LL9SS-037-0001-SO		LL9SS-038-0001-SO		LL9SS-039-0001-DU		LL9SS-039-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzofuran	mg/Kg	--	15	nc	0.0032	U														
		Diethyl phthalate	mg/Kg	--	4888	nc	0.0044	U														
		Dimethyl phthalate	mg/Kg	--	100000	max	0.0043	U														
		Di-n-butyl phthalate	mg/Kg	--	611	nc	0.024	U														
		Di-n-octyl phthalate	mg/Kg	--	244	nc	0.01	U														
		Fluoranthene	mg/Kg	--	229	nc	0.058															
		Fluorene	mg/Kg	--	275	nc	0.0019	U														
		Hexachlorobenzene	mg/Kg	--	0.30	ca	0.0022	U														
		Hexachlorobutadiene	mg/Kg	--	6.2	ca	0.0039	U														
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc	0.065	U														
		Hexachloroethane	mg/Kg	--	35	ca	0.0039	U														
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca	0.0025	U														
		Isophorone	mg/Kg	--	512	ca	0.0029	U														
		Naphthalene	mg/Kg	--	5.6	nc	0.002	U														
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca	0.0028	U														
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca	0.0035	U*														
		Pentachlorophenol	mg/Kg	--	3.0	ca	0.12	U														
		Phenanthrene	mg/Kg	--	--	0	0.021	Ja														
		Phenol	mg/Kg	--	1833	nc	0.0019	U														
		Pyrene	mg/Kg	--	232	nc	0.045															
TAL Metals 6010B	6010B	Aluminum	mg/Kg	17700	7614	nc	9300		7900		8500		8500		12000		13000		5700		6000	
		Barium	mg/Kg	88.4	538	nc	57		38		47		44		97		81		34		37	
		Beryllium	mg/Kg	0.88	15	nc	0.42	B	0.41	B	0.39	B	0.36	B	0.58		0.59		0.49		0.51	
		Cadmium	mg/Kg	0.00	3.7	nc	0.087	U	0.092	U	0.092	U	0.09	U	0.092	U	0.11	U	0.091	U	0.099	U
		Calcium	mg/Kg	15800	--	0	1300		4100		370		320		610		210		2000		1600	
		Chromium	mg/Kg	17.4	30	ca	14		13		14		14		16		15		9.8		9.7	
		Cobalt	mg/Kg	10.4	902	ca	8.7		7.5		8.1		7		17		14		4.5		4.8	
		Copper	mg/Kg	17.7	313	nc	34		14		9.1		8.8		9.2		11		7.1		7.8	
		Iron	mg/Kg	23100	2346	nc	21000		22000		20000		19000		17000		19000		25000		25000	
		Magnesium	mg/Kg	3030	--	0	1900		2100		1500		1600		1800		1700		1100		1100	
		Manganese	mg/Kg	1450	176	nc	630		350		570		420		970		1000		420		470	
		Nickel	mg/Kg	21.1	156	nc	15		14		12		12		15		14		9.4		10	
		Potassium	mg/Kg	927	--	0	760		820		660		650		680		720		520		550	
		Selenium	mg/Kg	1.4	39	nc	0.66	B	0.84	B	1.1	B	0.56	B	0.99	B	1.1	B	1.1	B	1.2	B
		Silver	mg/Kg	0.00	39	nc	0.34	U	0.35	U	0.36	U	0.35	U	0.36	U	0.44	U	0.35	U	0.38	U
		Sodium	mg/Kg	123	--	0	1000		1100		970		970		1100		1300		850		940	
		Vanadium	mg/Kg	31.1	7.8	nc	17		14		15		15		21		22		11		12	
		Zinc	mg/Kg	61.8	2346	nc	71		66		49		48		65		69		69		76	
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.5	U	0.47	U	0.52	U	0.52	U	0.5	U	0.54	U	0.48	U	0.55	U
	7060A	Arsenic	mg/Kg	15.4	0.39	ca	7.1		10		5.6		5.3		9.4		7.5		3.1		4.2	
	7421	Lead	mg/Kg	26.1	400	pbk	43		21		13		13		21		21		19		22	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	12		0.033		0.031		0.026		0.03		0.072		0.032		0.039	
	7841	Thallium	mg/Kg	0.00	0.52	nc	0.22	U	0.2	U	1.1	U	1.1	U	0.22	U	0.24	U	1	U	0.24	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-034-0001-SO		LL9SS-035-0001-SO		LL9SS-036-0001-DU		LL9SS-036-0001-SO		LL9SS-037-0001-SO		LL9SS-038-0001-SO		LL9SS-039-0001-DU		LL9SS-039-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat	0.00074	U*														
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca	0.00077	U														
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca	0.00086	U														
		1,1-Dichloroethane	mg/Kg	--	51	nc	0.0011	U														
		1,1-Dichloroethene	mg/Kg	--	12	nc	0.0012	U														
		1,2-Dichloroethane	mg/Kg	--	0.28	ca	0.0007	U														
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc	0.0023	U														
		1,2-Dichloropropane	mg/Kg	--	0.34	ca	0.0012	U														
		2-Butanone (MEK)	mg/Kg	--	2231	nc	0.0051	U														
		2-Hexanone	mg/Kg	--	530	nc	0.0021	U														
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc	0.0036	U														
		Acetone	mg/Kg	--	1412	nc	0.005	U*														
		Benzene	mg/Kg	--	0.64	ca	0.0008	U														
		Bromodichloromethane	mg/Kg	--	0.82	ca	0.00082	U														
		Bromoform	mg/Kg	--	62	ca	0.0011	U														
		Bromomethane	mg/Kg	--	0.39	nc	0.0035	U*														
		Carbon disulfide	mg/Kg	--	36	nc	0.0024	U														
		Carbon tetrachloride	mg/Kg	--	0.25	ca	0.001	U														
		Chlorobenzene	mg/Kg	--	15	nc	0.0011	U														
		Chloroethane	mg/Kg	--	3.0	ca	0.0019	U														
		Chloroform	mg/Kg	--	0.22	ca	0.00075	U														
		Chloromethane	mg/Kg	--	4.7	nc	0.0011	U														
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.00096	U														
		Dibromochloromethane	mg/Kg	--	1.1	ca	0.00083	U														
		Ethylbenzene	mg/Kg	--	395	sat	0.0013	U														
		Methylene chloride	mg/Kg	--	9.1	ca	0.0022	U														
		Styrene	mg/Kg	--	1700	sat	0.0012	U														
		Tetrachloroethene	mg/Kg	--	0.48	ca	0.00081	U														
		Toluene	mg/Kg	--	520	sat	0.0012	U														
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.001	U														
		Trichloroethene	mg/Kg	--	0.053	ca	0.00071	U														
		Vinyl chloride	mg/Kg	--	0.079	ca	0.00089	U														
		Xylenes (total)	mg/Kg	--	0.079	ca	0.0035	U														



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-042-0001-SO		LL9SS-043-0001-SO		LL9SS-044-0001-SO		LL9SS-045-0001-SO		LL9SS-046-0001-SO		LL9SS-047-0001-SO		LL9SS-068-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc													0.26	U
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc													0.018	U
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc													0.018	U
		2,4,6-TNT	mg/Kg	0	16	ca													0.034	U
		2,4-Dinitrotoluene	mg/Kg	0	12	nc													0.036	U
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc													0.048	U
		2-Nitrotoluene	mg/Kg	0	0.88	ca													0.033	U
		3-Nitrotoluene	mg/Kg	0	73	nc													0.05	U
		4-Nitrotoluene	mg/Kg	0	12	ca													0.047	U
		HMX	mg/Kg	0	306	nc													0.11	U
		Nitrobenzene	mg/Kg	0	2	nc													0.022	U
		RDX	mg/Kg	0	4.4	ca													0.059	U
		Tetryl	mg/Kg	0	61	nc													0.043	U
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc													0.0036	U
		Aroclor 1221	mg/Kg	--	0.22	ca													0.0038	U
		Aroclor 1232	mg/Kg	--	0.22	ca													0.0038	U
		Aroclor 1242	mg/Kg	--	0.22	ca													0.0038	U
		Aroclor 1248	mg/Kg	--	0.22	ca													0.0038	U
		Aroclor 1254	mg/Kg	--	0.22	ca													0.0038	U
		Aroclor 1260	mg/Kg	--	0.22	ca													0.0026	U
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca													0.00068	U
		4,4'-DDE	mg/Kg	--	1.7	ca													0.0012	U
		4,4'-DDT	mg/Kg	--	1.7	ca													0.0007	U
		Aldrin	mg/Kg	--	0.029	ca													0.00025	U
		alpha-BHC	mg/Kg	--	0.09	sat													0.0003	U
		alpha-Chlordane	mg/Kg	--	1.6	ca													0.00023	U
		beta-BHC	mg/Kg	--	0.32	ca													0.00028	U
		delta-BHC	mg/Kg	--	--	0													0.00021	U
		Dieldrin	mg/Kg	--	0.030	ca													0.00064	U
		Endosulfan I	mg/Kg	--	37	nc													0.00051	U
		Endosulfan II	mg/Kg	--	37	nc													0.00053	U
		Endosulfan sulfate	mg/Kg	--	37	nc													0.00055	U
		Endrin	mg/Kg	--	1.8	nc													0.00081	U
		Endrin aldehyde	mg/Kg	--	--	0													0.00062	U
		Endrin ketone	mg/Kg	--	--	0													0.00055	U
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca													0.00043	U
		gamma-Chlordane	mg/Kg	--	1.6	ca													0.00028	U
		Heptachlor	mg/Kg	--	0.11	ca													0.00034	U
		Heptachlor epoxide	mg/Kg	--	0.053	ca													0.00026	U
		Methoxychlor	mg/Kg	--	31	nc													0.0043	U
		Toxaphene	mg/Kg	--	0	0													0.0087	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-042-0001-SO		LL9SS-043-0001-SO		LL9SS-044-0001-SO		LL9SS-045-0001-SO		LL9SS-046-0001-SO		LL9SS-047-0001-SO		LL9SS-068-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0													3.2	B J
	8330	Nitroglycerin	mg/Kg	0	--	0													0.5	U
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc													0.25	U
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc													0.11	U
		1,2-Dichlorobenzene	mg/Kg	--	600	sat													0.15	U
		1,3-Dichlorobenzene	mg/Kg	--	53	nc													0.15	U
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca													0.13	U
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca													0.14	U
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc													0.07	U
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc													0.087	U
		2,4-Dichlorophenol	mg/Kg	--	18	nc													0.089	U
		2,4-Dimethylphenol	mg/Kg	--	122	nc													0.11	U
		2,4-Dinitrophenol	mg/Kg	--	12	nc													0.21	U
		2-Chloronaphthalene	mg/Kg	--	494	nc													0.089	U
		2-Chlorophenol	mg/Kg	--	6.3	nc													0.11	U
		2-Methylnaphthalene	mg/Kg	--	--	0													0.015	J
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc													0.016	U
		2-Nitroaniline	mg/Kg	--	18.3	nc													0.063	U
		2-Nitrophenol	mg/Kg	--	--	0													0.12	U
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca													0.033	U
		3-Nitroaniline	mg/Kg	--	1.8	nc													0.21	U
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc													0.18	U
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0													0.0057	U
		4-Chloro-3-methylphenol	mg/Kg	--	--	0													0.07	U
		4-Chloroaniline	mg/Kg	--	24	nc													0.18	U
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0													0.0067	U
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc													0.011	U
		4-Nitroaniline	mg/Kg	--	23	ca													0.072	U
		4-Nitrophenol	mg/Kg	--	--	0													0.15	U
		Acenaphthene	mg/Kg	--	368	nc													0.011	J
		Acenaphthylene	mg/Kg	--	--	0													0.05	J
		Anthracene	mg/Kg	--	2189	nc													0.048	J
		Benzo(a)anthracene	mg/Kg	--	0.62	ca													0.23	
		Benzo(a)pyrene	mg/Kg	--	0.062	ca													0.24	
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca													0.24	M
		Benzo(ghi)perylene	mg/Kg	--	--	0													0.17	
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca													0.2	M
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0													0.0054	U
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca													0.0037	U
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca													0.03	J
		Butyl benzyl phthalate	mg/Kg	--	1222	nc													0.0076	U
		Carbazole	mg/Kg	--	24	ca													0.065	U
		Chrysene	mg/Kg	--	62	ca													0.25	
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca													0.13	



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-042-0001-SO		LL9SS-043-0001-SO		LL9SS-044-0001-SO		LL9SS-045-0001-SO		LL9SS-046-0001-SO		LL9SS-047-0001-SO		LL9SS-068-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzofuran	mg/Kg	--	15	nc													0.013	J
		Diethyl phthalate	mg/Kg	--	4888	nc													0.0068	U
		Dimethyl phthalate	mg/Kg	--	100000	max													0.0067	U
		Di-n-butyl phthalate	mg/Kg	--	611	nc													0.037	U
		Di-n-octyl phthalate	mg/Kg	--	244	nc													0.016	U
		Fluoranthene	mg/Kg	--	229	nc													0.36	
		Fluorene	mg/Kg	--	275	nc													0.025	J
		Hexachlorobenzene	mg/Kg	--	0.30	ca													0.0033	U
		Hexachlorobutadiene	mg/Kg	--	6.2	ca													0.0061	U
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc													0.1	U
		Hexachloroethane	mg/Kg	--	35	ca													0.0061	U
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca													0.17	
		Isophorone	mg/Kg	--	512	ca													0.0044	U
		Naphthalene	mg/Kg	--	5.6	nc													0.019	J
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca													0.0043	U
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca													0.0054	U
		Pentachlorophenol	mg/Kg	--	3.0	ca													0.18	U
		Phenanthrene	mg/Kg	--	--	0													0.28	
		Phenol	mg/Kg	--	1833	nc													0.003	U
		Pyrene	mg/Kg	--	232	nc													0.4	
TAL Metals 6010B	6010B	Aluminum	mg/Kg	17700	7614	nc	7200		10000		12000		13000		8900		8600		8400	
		Barium	mg/Kg	88.4	538	nc	41		57		85		40		53		61		110	
		Beryllium	mg/Kg	0.88	15	nc	0.47		0.47		0.27	B	0.48		0.42	B	0.26	B	0.54	B
		Cadmium	mg/Kg	0.00	3.7	nc	0.078	U	0.086	U	0.09	U	0.082	U	0.09	U	0.088	U	2.7	
		Calcium	mg/Kg	15800	--	0	470		630		2800		750		890		770		1900	
		Chromium	mg/Kg	17.4	30	ca	10		13		16		17		13		12		15	
		Cobalt	mg/Kg	10.4	902	ca	9.2		9.3		4.2		8.6		11		6.4		6.8	
		Copper	mg/Kg	17.7	313	nc	11		13		9.7		25		21		15		170	
		Iron	mg/Kg	23100	2346	nc	18000		20000		17000		31000		25000		16000		20000	
		Magnesium	mg/Kg	3030	--	0	1700		2400		2200		2900		3000		2000		1200	
		Manganese	mg/Kg	1450	176	nc	610		440		150		240		370		690		700	
		Nickel	mg/Kg	21.1	156	nc	13		17		19		21		24		13		15	
		Potassium	mg/Kg	927	--	0	710		860		770		1200		900		890		680	
		Selenium	mg/Kg	1.4	39	nc	0.39	B	0.62	B	0.45	U	0.41	U	0.45	U	0.45	B	1.8	B
		Silver	mg/Kg	0.00	39	nc	0.3	U	0.33	U	0.35	U	0.32	U	0.35	U	0.34	U	0.57	U
		Sodium	mg/Kg	123	--	0	930		1100		1200		1100		1300		1200		170	B
		Vanadium	mg/Kg	31.1	7.8	nc	13		17		13		20		14		13		16	
		Zinc	mg/Kg	61.8	2346	nc	46		50		40		62		63		45		780	
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.41	U	0.49	U	0.47	U	0.49	U	0.54	B	0.52	U	0.79	U
	7060A	Arsenic	mg/Kg	15.4	0.39	ca	4		9.2		18		17		18		15		7.3	
	7421	Lead	mg/Kg	26.1	400	pbk	9.2		14		10		16		13		12		150	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.0076	B	0.026		0.0057	B	0.0081	B	0.016	B	0.063		0.21	
	7841	Thallium	mg/Kg	0.00	0.52	nc	0.89	U	0.21	U	1	U	0.21	U	1	U	1.1	U	0.34	U



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

							LL9SS-042-0001-SO		LL9SS-043-0001-SO		LL9SS-044-0001-SO		LL9SS-045-0001-SO		LL9SS-046-0001-SO		LL9SS-047-0001-SO		LL9SS-068-0001-SO			
							Sample Date		11/06/03		11/06/03		11/06/03		11/06/03		11/06/03		11/06/03		12/12/03	
							Sample Depth		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 1 ft		0 - 0.3 ft	
Group	Method	Analyte	Units	RVAAP Surface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SS PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier		
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat													0.0021	U		
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca													0.0018	U		
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca													0.0021	U		
		1,1-Dichloroethane	mg/Kg	--	51	nc														0.0019	U	
		1,1-Dichloroethene	mg/Kg	--	12	nc														0.0025	U	
		1,2-Dichloroethane	mg/Kg	--	0.28	ca														0.0018	U	
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc														0.004	U	
		1,2-Dichloropropane	mg/Kg	--	0.34	ca														0.0019	U	
		2-Butanone (MEK)	mg/Kg	--	2231	nc														0.0074	U	
		2-Hexanone	mg/Kg	--	530	nc														0.0021	U	
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc														0.0019	U*	
		Acetone	mg/Kg	--	1412	nc														0.0087	U	
		Benzene	mg/Kg	--	0.64	ca														0.0021	U	
		Bromodichloromethane	mg/Kg	--	0.82	ca														0.0018	U	
		Bromoform	mg/Kg	--	62	ca														0.0014	U	
		Bromomethane	mg/Kg	--	0.39	nc														0.0025	U	
		Carbon disulfide	mg/Kg	--	36	nc														0.0023	U	
		Carbon tetrachloride	mg/Kg	--	0.25	ca														0.0021	U	
		Chlorobenzene	mg/Kg	--	15	nc														0.0021	U	
		Chloroethane	mg/Kg	--	3.0	ca														0.0019	U	
		Chloroform	mg/Kg	--	0.22	ca														0.0021	U	
		Chloromethane	mg/Kg	--	4.7	nc														0.0021	U	
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca														0.0018	U	
		Dibromochloromethane	mg/Kg	--	1.1	ca														0.0015	U	
		Ethylbenzene	mg/Kg	--	395	sat														0.0021	U	
		Methylene chloride	mg/Kg	--	9.1	ca														0.0055	U	
		Styrene	mg/Kg	--	1700	sat														0.0021	U	
		Tetrachloroethene	mg/Kg	--	0.48	ca														0.0023	U	
		Toluene	mg/Kg	--	520	sat														0.0021	U	
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca														0.0015	U	
		Trichloroethene	mg/Kg	--	0.053	ca														0.0021	U	
		Vinyl chloride	mg/Kg	--	0.079	ca														0.0021	U	
		Xylenes (total)	mg/Kg	--	0.079	ca														0.0064	U	



Table 4-3
Summary of Surface Soil Results (0-1 ft bgs)

Notes:
-- - no background/PRG value is available for this analyte
blank cell indicates that the analysis was not performed
PRG - preliminary remediation goals
(The PRG for lead is the Maximum Contaminant Level (MCL))
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background and PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG and Background, then the value is presented with a normal style

Inorganic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL
InOrganic Flags
^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL
Organic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL
Organic Flags
B - Method Blank: batch QC is greater than R
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-001-0001-SO		LL9SB-002-0001-SO		LL9SB-003-0001-SO		LL9SB-004-0001-SO		LL9SB-005-0001-DUP		LL9SB-005-0001-SO		LL9SB-006-0001-SO		LL9SB-007-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc									0.17	U	0.25	U				
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc	0.017	U							0.017	U	0.018	U				
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc	0.017	U							0.018	U	0.018	U				
		2,4,6-TNT	mg/Kg	0	16	ca	0.033	U							0.034	U	0.034	U				
		2,4-Dinitrotoluene	mg/Kg	0	12	nc	0.035	U							0.035	U	0.036	U				
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc	0.047	U*							0.047	U*	0.048	U*				
		2-Nitrotoluene	mg/Kg	0	0.88	ca	0.033	U							0.033	U	0.033	U				
		3-Nitrotoluene	mg/Kg	0	73	nc	0.049	U							0.05	U	0.05	U				
		4-Nitrotoluene	mg/Kg	0	12	ca	0.046	U							0.046	U	0.047	U				
		HMX	mg/Kg	0	306	nc	0.11	U							0.11	U	0.11	U				
		Nitrobenzene	mg/Kg	0	2	nc	0.022	U							0.022	U	0.022	U				
		RDX	mg/Kg	0	4.4	ca	0.057	U							0.058	U	0.059	U				
		Tetryl	mg/Kg	0	61	nc	0.043	U							0.043	U	0.043	U				
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--																
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--																
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc									0.0023	U	0.0022	U				
		Aroclor 1221	mg/Kg	--	0.22	ca									0.0024	U	0.0023	U				
		Aroclor 1232	mg/Kg	--	0.22	ca									0.0024	U	0.0023	U				
		Aroclor 1242	mg/Kg	--	0.22	ca									0.0024	U	0.0023	U				
		Aroclor 1248	mg/Kg	--	0.22	ca									0.0024	U	0.0023	U				
		Aroclor 1254	mg/Kg	--	0.22	ca									0.0024	U	0.0023	U				
		Aroclor 1260	mg/Kg	--	0.22	ca									0.0017	U	0.0016	U				
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca									0.00043	U	0.00041	U				
		4,4'-DDE	mg/Kg	--	1.7	ca									0.00078	U	0.00075	U				
		4,4'-DDT	mg/Kg	--	1.7	ca									0.00044	U	0.00043	U				
		Aldrin	mg/Kg	--	0.029	ca									0.00016	U	0.00015	U				
		alpha-BHC	mg/Kg	--	0.09	sat									0.00019	U	0.00018	U				
		alpha-Chlordane	mg/Kg	--	1.6	ca									0.00014	U	0.00014	U				
		beta-BHC	mg/Kg	--	0.32	ca									0.00018	U	0.00017	U				
		delta-BHC	mg/Kg	--	--	0									0.00013	U	0.00013	U				
		Dieldrin	mg/Kg	--	0.030	ca									0.00041	U	0.00039	U				
		Endosulfan I	mg/Kg	--	37	nc									0.00032	U	0.00031	U				
		Endosulfan II	mg/Kg	--	37	nc									0.00034	U	0.00032	U				
		Endosulfan sulfate	mg/Kg	--	37	nc									0.00035	U	0.00033	U				
		Endrin	mg/Kg	--	1.8	nc									0.00052	U	0.0005	U				
		Endrin aldehyde	mg/Kg	--	--	0									0.0004	U	0.00038	U				
		Endrin ketone	mg/Kg	--	--	0									0.00035	U	0.00033	U				
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca									0.00028	U	0.00027	U				
		gamma-Chlordane	mg/Kg	--	1.6	ca									0.00018	U	0.00017	U				
		Heptachlor	mg/Kg	--	0.11	ca									0.00022	U	0.00021	U				
		Heptachlor epoxide	mg/Kg	--	0.053	ca									0.00017	U	0.00016	U				
		Methoxychlor	mg/Kg	--	31	nc									0.0028	U	0.0027	U				
		Toxaphene	mg/Kg	--	0	0									0.0055	U	0.0053	U				



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-001-0001-SO		LL9SB-002-0001-SO		LL9SB-003-0001-SO		LL9SB-004-0001-SO		LL9SB-005-0001-DUP		LL9SB-005-0001-SO		LL9SB-006-0001-SO		LL9SB-007-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0									2.7	J	1.2	B J				
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0									0.5	U	0.5	U				
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc									0.25	U	0.25	U				
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc									0.073	U	0.07	U				
		1,2-Dichlorobenzene	mg/Kg	--	600	sat									0.097	U	0.094	U				
		1,3-Dichlorobenzene	mg/Kg	--	53	nc									0.097	U	0.094	U				
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca									0.087	U	0.085	U				
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca									0.092	U	0.089	U				
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc									0.047	U	0.045	U				
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc									0.058	U	0.056	U				
		2,4-Dichlorophenol	mg/Kg	--	18	nc									0.059	U	0.057	U				
		2,4-Dimethylphenol	mg/Kg	--	122	nc									0.074	U	0.071	U				
		2,4-Dinitrophenol	mg/Kg	--	12	nc									0.14	U	0.14	U				
		2-Chloronaphthalene	mg/Kg	--	494	nc									0.059	U	0.057	U				
		2-Chlorophenol	mg/Kg	--	6.3	nc									0.073	U	0.07	U				
		2-Methylnaphthalene	mg/Kg	--	--	0									0.0018	U	0.0018	U				
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc									0.01	U	0.01	U				
		2-Nitroaniline	mg/Kg	--	18.3	nc									0.042	U	0.04	U				
		2-Nitrophenol	mg/Kg	--	--	0									0.078	U	0.075	U				
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca									0.022	U	0.021	U				
		3-Nitroaniline	mg/Kg	--	1.8	nc									0.14	U	0.13	U				
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc									0.12	U*	0.11	U*				
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0									0.0038	U	0.0037	U				
		4-Chloro-3-methylphenol	mg/Kg	--	--	0									0.047	U	0.045	U				
		4-Chloroaniline	mg/Kg	--	24	nc									0.12	U	0.12	U				
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0									0.0044	U	0.0043	U				
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc									0.0071	U	0.0069	U				
		4-Nitroaniline	mg/Kg	--	23	ca									0.048	U	0.046	U				
		4-Nitrophenol	mg/Kg	--	--	0									0.1	U	0.098	U				
		Acenaphthene	mg/Kg	--	368	nc									0.0017	U	0.0017	U				
		Acenaphthylene	mg/Kg	--	--	0									0.0011	U	0.0011	U				
		Anthracene	mg/Kg	--	2189	nc									0.0011	U	0.001	U				
		Benzo(a)anthracene	mg/Kg	--	0.62	ca									0.0014	U	0.0013	U				
		Benzo(a)pyrene	mg/Kg	--	0.062	ca									0.0027	U	0.0026	U				
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca									0.0026	U	0.0025	U				
		Benzo(ghi)perylene	mg/Kg	--	--	0									0.0038	J	0.0023	U				
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca									0.0034	U	0.0033	U				
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0									0.0036	U	0.0035	U				
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca									0.0025	U	0.0024	U				
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca									0.012	U	0.011	U				



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-001-0001-SO		LL9SB-002-0001-SO		LL9SB-003-0001-SO		LL9SB-004-0001-SO		LL9SB-005-0001-DUP		LL9SB-005-0001-SO		LL9SB-006-0001-SO		LL9SB-007-0001-SO	
							12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		12/11/03		12/08/03	
							1 - 3 ft		1 - 3 ft		1 - 2 ft		1 - 3 ft		1 - 3 ft		1 - 3 ft		1 - 2 ft		1 - 3 ft	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc									0.005	U	0.0049	U				
		Carbazole	mg/Kg	--	24	ca									0.043	U	0.042	U				
		Chrysene	mg/Kg	--	62	ca									0.0022	U	0.0021	U				
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca									0.0027	U	0.0026	U				
		Dibenzofuran	mg/Kg	--	15	nc									0.0033	U	0.0032	U				
		Diethyl phthalate	mg/Kg	--	4888	nc									0.0046	U	0.0044	U				
		Dimethyl phthalate	mg/Kg	--	100000	max									0.0044	U	0.0043	U				
		Di-n-butyl phthalate	mg/Kg	--	611	nc									0.025	U	0.024	U				
		Di-n-octyl phthalate	mg/Kg	--	244	nc									0.011	U	0.01	U				
		Fluoranthene	mg/Kg	--	229	nc									0.0014	U	0.0013	U				
		Fluorene	mg/Kg	--	275	nc									0.002	U	0.0019	U				
		Hexachlorobenzene	mg/Kg	--	0.30	ca									0.0022	U	0.0021	U				
		Hexachlorobutadiene	mg/Kg	--	6.2	ca									0.0041	U	0.0039	U				
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc									0.066	U	0.064	U				
		Hexachloroethane	mg/Kg	--	35	ca									0.0041	U	0.0039	U				
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca									0.018	J	0.017	J				
		Isophorone	mg/Kg	--	512	ca									0.003	U	0.0029	U				
		Naphthalene	mg/Kg	--	5.6	nc									0.0021	U	0.002	U				
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca									0.0028	U	0.0027	U				
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca									0.0036	U	0.0035	U				
		Pentachlorophenol	mg/Kg	--	3.0	ca									0.12	U	0.12	U				
		Phenanthrene	mg/Kg	--	--	0									0.0012	U	0.0012	U				
		Phenol	mg/Kg	--	1833	nc									0.002	U	0.0019	U				
		Pyrene	mg/Kg	--	232	nc									0.0025	U	0.0024	U				
TAL Metals 6010B	6010B	Aluminum	mg/Kg	19500	7614	nc	7500		10000		6600		8200		11000		11000		7000		12000	
		Barium	mg/Kg	124	538	nc	30		64		26		50		80		70		66		75	
		Beryllium	mg/Kg	0.88	15	nc	0.33	B	0.44	B	0.38	B	0.43		0.8		0.63		0.31	B	0.55	
		Cadmium	mg/Kg	0.00	3.7	nc	0.091	U	0.092	U	0.085	U	0.084	U	0.091	U	0.09	U	2		0.13	B
		Calcium	mg/Kg	35500	--	0	800		4300		530		1100		360		330		2700		1900	
		Chromium	mg/Kg	27.2	30	ca	10		16		8.4		10		13		13		11		16	
		Cobalt	mg/Kg	23.2	902	ca	6.8		8		5.5		3.8		11		11		6.2		10	
		Copper	mg/Kg	32.3	313	nc	27		19		17		6.5		9.3		9.1		15		21	H
		Iron	mg/Kg	35200	2346	nc	23000		23000		16000		18000		17000		15000		14000	H	23000	
		Magnesium	mg/Kg	8790	--	0	2100		2200		1600		1200		1700		1600		1500		2700	
		Manganese	mg/Kg	3030	176	nc	400		600		310		200		1300		1100		520		570	
		Nickel	mg/Kg	60.7	156	nc	18		17		13		7.1		16		14		12		21	
		Potassium	mg/Kg	3350	--	0	630		660		480		490		560		550		550		1100	
		Selenium	mg/Kg	1.5	39	nc	0.45	U	0.77	B	0.43	U	0.42	U	0.91	B	0.69	B	0.38	B	0.46	U
		Silver	mg/Kg	0.00	39	nc	0.35	U	0.36	U	0.33	U	0.32	U	0.35	U	0.35	U	0.34	U	0.36	U
		Sodium	mg/Kg	145	--	0	98	U	99	U	93	U	91	U	99	U	97	U	95	U	100	U
		Vanadium	mg/Kg	37.6	7.8	nc	13		19		11		15		18		17		13		21	
		Zinc	mg/Kg	93.3	2346	nc	60		68		48		37		49		45		120		62	



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-001-0001-SO		LL9SB-002-0001-SO		LL9SB-003-0001-SO		LL9SB-004-0001-SO		LL9SB-005-0001-DUP		LL9SB-005-0001-SO		LL9SB-006-0001-SO		LL9SB-007-0001-SO	
Sample Date							12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		12/04/03		12/11/03		12/08/03	
Sample Depth							1 - 3 ft		1 - 3 ft		1 - 2 ft		1 - 3 ft		1 - 3 ft		1 - 3 ft		1 - 2 ft		1 - 3 ft	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.52	U	0.48	U	0.48	U	0.5	U	0.53	U	0.48	U	0.48	U	0.5	U
	7060A	Arsenic	mg/Kg	19.8	0.39	ca	12		12		9.8		8.5		9		7.7		9.9		14	
	7421	Lead	mg/Kg	19.1	400	pbk	9.2		24		10		9.9		12		11		54		19	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.021		0.076		0.022		0.019	B	0.039		0.034		0.94		0.025	
	7841	Thallium	mg/Kg	0.91	0.52	nc	0.23	U	0.21	U	0.21	U	0.22	U	0.23	U	0.21	U	0.21	U	0.22	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat									0.0014	U	0.0013	U				
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca									0.0012	U	0.0012	U				
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca									0.0014	U	0.0013	U				
		1,1-Dichloroethane	mg/Kg	--	51	nc									0.0012	U	0.0012	U				
		1,1-Dichloroethene	mg/Kg	--	12	nc									0.0016	U	0.0016	U				
		1,2-Dichloroethane	mg/Kg	--	0.28	ca									0.0012	U	0.0011	U				
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc									0.0026	U	0.0025	U				
		1,2-Dichloropropane	mg/Kg	--	0.34	ca									0.0012	U	0.0012	U				
		2-Butanone (MEK)	mg/Kg	--	2231	nc									0.0048	U	0.0047	U				
		2-Hexanone	mg/Kg	--	530	nc									0.0014	U	0.0013	U				
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc									0.0012	U	0.0012	U				
		Acetone	mg/Kg	--	1412	nc									0.0057	U	0.0055	U				
		Benzene	mg/Kg	--	0.64	ca									0.0014	U	0.0013	U				
		Bromodichloromethane	mg/Kg	--	0.82	ca									0.0012	U	0.0012	U				
		Bromoform	mg/Kg	--	62	ca									0.00093	U	0.0009	U				
		Bromomethane	mg/Kg	--	0.39	nc									0.0016	U*	0.0016	U*				
		Carbon disulfide	mg/Kg	--	36	nc									0.0015	U	0.0014	U				
		Carbon tetrachloride	mg/Kg	--	0.25	ca									0.0014	U	0.0013	U				
		Chlorobenzene	mg/Kg	--	15	nc									0.0014	U	0.0013	U				
		Chloroethane	mg/Kg	--	3.0	ca									0.0012	U	0.0012	U				
		Chloroform	mg/Kg	--	0.22	ca									0.0014	U	0.0013	U				
		Chloromethane	mg/Kg	--	4.7	nc									0.0014	U	0.0013	U				
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca									0.0012	U	0.0011	U				
		Dibromochloromethane	mg/Kg	--	1.1	ca									0.00098	U	0.00095	U				
		Ethylbenzene	mg/Kg	--	395	sat									0.0014	U	0.0013	U				
		Methylene chloride	mg/Kg	--	9.1	ca									0.0036	U	0.0035	U				
		Styrene	mg/Kg	--	1700	sat									0.0014	U	0.0013	U				
		Tetrachloroethene	mg/Kg	--	0.48	ca									0.0015	U	0.0014	U				
		Toluene	mg/Kg	--	520	sat									0.0014	U	0.0013	U				
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca									0.00098	U	0.00095	U				
		Trichloroethene	mg/Kg	--	0.053	ca									0.0014	U	0.0013	U				
		Vinyl chloride	mg/Kg	--	0.079	ca									0.0014	U	0.0013	U				
		Xylenes (total)	mg/Kg	--	0.079	ca									0.0042	U	0.0041	U				



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-008-0001-SO		LL9SB-009-0001-SO		LL9SB-010-0001-SO		LL9SB-012-0001-SO		LL9SB-013-0001-SO		LL9SB-014-0001-SO		LL9SB-015-0001-SO		LL9SB-016-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc																
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc																
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc																
		2,4,6-TNT	mg/Kg	0	16	ca																
		2,4-Dinitrotoluene	mg/Kg	0	12	nc																
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc																
		2-Nitrotoluene	mg/Kg	0	0.88	ca																
		3-Nitrotoluene	mg/Kg	0	73	nc																
		4-Nitrotoluene	mg/Kg	0	12	ca																
		HMX	mg/Kg	0	306	nc																
		Nitrobenzene	mg/Kg	0	2	nc																
		RDX	mg/Kg	0	4.4	ca																
		Tetryl	mg/Kg	0	61	nc																
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--																
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--																
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc																
		Aroclor 1221	mg/Kg	--	0.22	ca																
		Aroclor 1232	mg/Kg	--	0.22	ca																
		Aroclor 1242	mg/Kg	--	0.22	ca																
		Aroclor 1248	mg/Kg	--	0.22	ca																
		Aroclor 1254	mg/Kg	--	0.22	ca																
		Aroclor 1260	mg/Kg	--	0.22	ca																
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca																
		4,4'-DDE	mg/Kg	--	1.7	ca																
		4,4'-DDT	mg/Kg	--	1.7	ca																
		Aldrin	mg/Kg	--	0.029	ca																
		alpha-BHC	mg/Kg	--	0.09	sat																
		alpha-Chlordane	mg/Kg	--	1.6	ca																
		beta-BHC	mg/Kg	--	0.32	ca																
		delta-BHC	mg/Kg	--	--	0																
		Dieldrin	mg/Kg	--	0.030	ca																
		Endosulfan I	mg/Kg	--	37	nc																
		Endosulfan II	mg/Kg	--	37	nc																
		Endosulfan sulfate	mg/Kg	--	37	nc																
		Endrin	mg/Kg	--	1.8	nc																
		Endrin aldehyde	mg/Kg	--	--	0																
		Endrin ketone	mg/Kg	--	--	0																
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca																
		gamma-Chlordane	mg/Kg	--	1.6	ca																
		Heptachlor	mg/Kg	--	0.11	ca																
		Heptachlor epoxide	mg/Kg	--	0.053	ca																
		Methoxychlor	mg/Kg	--	31	nc																
		Toxaphene	mg/Kg	--	0	0																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-008-0001-SO		LL9SB-009-0001-SO		LL9SB-010-0001-SO		LL9SB-012-0001-SO		LL9SB-013-0001-SO		LL9SB-014-0001-SO		LL9SB-015-0001-SO		LL9SB-016-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0																
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0																
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc																
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc																
		1,2-Dichlorobenzene	mg/Kg	--	600	sat																
		1,3-Dichlorobenzene	mg/Kg	--	53	nc																
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca																
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca																
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc																
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc																
		2,4-Dichlorophenol	mg/Kg	--	18	nc																
		2,4-Dimethylphenol	mg/Kg	--	122	nc																
		2,4-Dinitrophenol	mg/Kg	--	12	nc																
		2-Chloronaphthalene	mg/Kg	--	494	nc																
		2-Chlorophenol	mg/Kg	--	6.3	nc																
		2-Methylnaphthalene	mg/Kg	--	--	0																
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc																
		2-Nitroaniline	mg/Kg	--	18.3	nc																
		2-Nitrophenol	mg/Kg	--	--	0																
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca																
		3-Nitroaniline	mg/Kg	--	1.8	nc																
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc																
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0																
		4-Chloro-3-methylphenol	mg/Kg	--	--	0																
		4-Chloroaniline	mg/Kg	--	24	nc																
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0																
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc																
		4-Nitroaniline	mg/Kg	--	23	ca																
		4-Nitrophenol	mg/Kg	--	--	0																
		Acenaphthene	mg/Kg	--	368	nc																
		Acenaphthylene	mg/Kg	--	--	0																
		Anthracene	mg/Kg	--	2189	nc																
		Benzo(a)anthracene	mg/Kg	--	0.62	ca																
		Benzo(a)pyrene	mg/Kg	--	0.062	ca																
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca																
		Benzo(ghi)perylene	mg/Kg	--	--	0																
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca																
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0																
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca																
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-008-0001-SO		LL9SB-009-0001-SO		LL9SB-010-0001-SO		LL9SB-012-0001-SO		LL9SB-013-0001-SO		LL9SB-014-0001-SO		LL9SB-015-0001-SO		LL9SB-016-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc																
		Carbazole	mg/Kg	--	24	ca																
		Chrysene	mg/Kg	--	62	ca																
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca																
		Dibenzofuran	mg/Kg	--	15	nc																
		Diethyl phthalate	mg/Kg	--	4888	nc																
		Dimethyl phthalate	mg/Kg	--	100000	max																
		Di-n-butyl phthalate	mg/Kg	--	611	nc																
		Di-n-octyl phthalate	mg/Kg	--	244	nc																
		Fluoranthene	mg/Kg	--	229	nc																
		Fluorene	mg/Kg	--	275	nc																
		Hexachlorobenzene	mg/Kg	--	0.30	ca																
		Hexachlorobutadiene	mg/Kg	--	6.2	ca																
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc																
		Hexachloroethane	mg/Kg	--	35	ca																
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca																
		Isophorone	mg/Kg	--	512	ca																
		Naphthalene	mg/Kg	--	5.6	nc																
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca																
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca																
		Pentachlorophenol	mg/Kg	--	3.0	ca																
		Phenanthrene	mg/Kg	--	--	0																
		Phenol	mg/Kg	--	1833	nc																
TAL Metals 6010B	6010B	Pyrene	mg/Kg	--	232	nc																
		Aluminum	mg/Kg	19500	7614	nc	12000		12000		9700		9700		6700		7100		10000		8000	
		Barium	mg/Kg	124	538	nc	71		51		57		55		29		34		60		47	
		Beryllium	mg/Kg	0.88	15	nc	0.58		0.52		0.55		0.34	B	0.25	B	0.23	B	0.54		0.43	
		Cadmium	mg/Kg	0.00	3.7	nc	0.095	U	0.091	U	0.13	B	0.091	U	0.084	U	0.081	U	0.079	U	0.077	U
		Calcium	mg/Kg	35500	--	0	1100		420		4900		2600		4400		900		750		2200	
		Chromium	mg/Kg	27.2	30	ca	15		17		14		13		9		12		15		12	
		Cobalt	mg/Kg	23.2	902	ca	10		11		9.3		8.8		6		7.2		11		7.1	
		Copper	mg/Kg	32.3	313	nc	19		21	H	21	H	26		24		20		22		19	
		Iron	mg/Kg	35200	2346	nc	30000		28000		24000		25000		18000		18000	H	27000	H	24000	H
		Magnesium	mg/Kg	8790	--	0	2100		3000		3200		2500		2600		1800		3200		2100	
		Manganese	mg/Kg	3030	176	nc	550		280		470		470		420		430		280		550	
		Nickel	mg/Kg	60.7	156	nc	17		24		25		19		15		16		32		17	
		Potassium	mg/Kg	3350	--	0	780		1300		1200		900		640		620		1300		760	
		Selenium	mg/Kg	1.5	39	nc	0.47	U	0.46	U	0.44	U	0.46	U	0.42	U	0.32	B	0.15	B	0.3	B
		Silver	mg/Kg	0.00	39	nc	0.37	U	0.35	U	0.34	U	0.35	U	0.32	U	0.31	U	0.31	U	0.3	U
		Sodium	mg/Kg	145	--	0	100	U	99	U	96	U	99	U	91	U	87	U	85	U	83	U
		Vanadium	mg/Kg	37.6	7.8	nc	22		18		14		17		11		12		17		15	
		Zinc	mg/Kg	93.3	2346	nc	64		62		72		110		64		65		62		82	



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-008-0001-SO		LL9SB-009-0001-SO		LL9SB-010-0001-SO		LL9SB-012-0001-SO		LL9SB-013-0001-SO		LL9SB-014-0001-SO		LL9SB-015-0001-SO		LL9SB-016-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.53	U	0.47	U	0.49	U	0.8		0.46	U	0.48	U	1		0.47	U
	7060A	Arsenic	mg/Kg	19.8	0.39	ca	12		22		22		19		11		18		18		11	
	7421	Lead	mg/Kg	19.1	400	pbk	14		13		19		20		11		15		14		17	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.037		0.014	B	0.18		0.033		0.016	B	0.02		0.015	B	0.029	
	7841	Thallium	mg/Kg	0.91	0.52	nc	0.23	U	0.2	U	0.21	U	0.2	U	0.016	U	0.021	U	0.015	U	0.029	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat																
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca																
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca																
		1,1-Dichloroethane	mg/Kg	--	51	nc																
		1,1-Dichloroethene	mg/Kg	--	12	nc																
		1,2-Dichloroethane	mg/Kg	--	0.28	ca																
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc																
		1,2-Dichloropropane	mg/Kg	--	0.34	ca																
		2-Butanone (MEK)	mg/Kg	--	2231	nc																
		2-Hexanone	mg/Kg	--	530	nc																
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc																
		Acetone	mg/Kg	--	1412	nc																
		Benzene	mg/Kg	--	0.64	ca																
		Bromodichloromethane	mg/Kg	--	0.82	ca																
		Bromoform	mg/Kg	--	62	ca																
		Bromomethane	mg/Kg	--	0.39	nc																
		Carbon disulfide	mg/Kg	--	36	nc																
		Carbon tetrachloride	mg/Kg	--	0.25	ca																
		Chlorobenzene	mg/Kg	--	15	nc																
		Chloroethane	mg/Kg	--	3.0	ca																
		Chloroform	mg/Kg	--	0.22	ca																
		Chloromethane	mg/Kg	--	4.7	nc																
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca																
		Dibromochloromethane	mg/Kg	--	1.1	ca																
		Ethylbenzene	mg/Kg	--	395	sat																
		Methylene chloride	mg/Kg	--	9.1	ca																
		Styrene	mg/Kg	--	1700	sat																
		Tetrachloroethene	mg/Kg	--	0.48	ca																
		Toluene	mg/Kg	--	520	sat																
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca																
		Trichloroethene	mg/Kg	--	0.053	ca																
		Vinyl chloride	mg/Kg	--	0.079	ca																
		Xylenes (total)	mg/Kg	--	0.079	ca																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-017-0001-SO		LL9SB-018-0001-SO		LL9SB-019-0001-SO		LL9SB-020-0001-SO		LL9SB-021-0001-SO		LL9SB-022-0001-SO		LL9SB-023-0001-SO		LL9SB-024-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc																
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc																
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc																
		2,4,6-TNT	mg/Kg	0	16	ca																
		2,4-Dinitrotoluene	mg/Kg	0	12	nc																
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc																
		2-Nitrotoluene	mg/Kg	0	0.88	ca																
		3-Nitrotoluene	mg/Kg	0	73	nc																
		4-Nitrotoluene	mg/Kg	0	12	ca																
		HMX	mg/Kg	0	306	nc																
		Nitrobenzene	mg/Kg	0	2	nc																
		RDX	mg/Kg	0	4.4	ca																
		Tetryl	mg/Kg	0	61	nc																
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--																
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--																
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc																
		Aroclor 1221	mg/Kg	--	0.22	ca																
		Aroclor 1232	mg/Kg	--	0.22	ca																
		Aroclor 1242	mg/Kg	--	0.22	ca																
		Aroclor 1248	mg/Kg	--	0.22	ca																
		Aroclor 1254	mg/Kg	--	0.22	ca																
		Aroclor 1260	mg/Kg	--	0.22	ca																
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca																
		4,4'-DDE	mg/Kg	--	1.7	ca																
		4,4'-DDT	mg/Kg	--	1.7	ca																
		Aldrin	mg/Kg	--	0.029	ca																
		alpha-BHC	mg/Kg	--	0.09	sat																
		alpha-Chlordane	mg/Kg	--	1.6	ca																
		beta-BHC	mg/Kg	--	0.32	ca																
		delta-BHC	mg/Kg	--	--	0																
		Dieldrin	mg/Kg	--	0.030	ca																
		Endosulfan I	mg/Kg	--	37	nc																
		Endosulfan II	mg/Kg	--	37	nc																
		Endosulfan sulfate	mg/Kg	--	37	nc																
		Endrin	mg/Kg	--	1.8	nc																
		Endrin aldehyde	mg/Kg	--	--	0																
		Endrin ketone	mg/Kg	--	--	0																
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca																
		gamma-Chlordane	mg/Kg	--	1.6	ca																
		Heptachlor	mg/Kg	--	0.11	ca																
		Heptachlor epoxide	mg/Kg	--	0.053	ca																
		Methoxychlor	mg/Kg	--	31	nc																
		Toxaphene	mg/Kg	--	0	0																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-017-0001-SO		LL9SB-018-0001-SO		LL9SB-019-0001-SO		LL9SB-020-0001-SO		LL9SB-021-0001-SO		LL9SB-022-0001-SO		LL9SB-023-0001-SO		LL9SB-024-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0																
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0																
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc																
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc																
		1,2-Dichlorobenzene	mg/Kg	--	600	sat																
		1,3-Dichlorobenzene	mg/Kg	--	53	nc																
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca																
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca																
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc																
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc																
		2,4-Dichlorophenol	mg/Kg	--	18	nc																
		2,4-Dimethylphenol	mg/Kg	--	122	nc																
		2,4-Dinitrophenol	mg/Kg	--	12	nc																
		2-Chloronaphthalene	mg/Kg	--	494	nc																
		2-Chlorophenol	mg/Kg	--	6.3	nc																
		2-Methylnaphthalene	mg/Kg	--	--	0																
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc																
		2-Nitroaniline	mg/Kg	--	18.3	nc																
		2-Nitrophenol	mg/Kg	--	--	0																
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca																
		3-Nitroaniline	mg/Kg	--	1.8	nc																
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc																
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0																
		4-Chloro-3-methylphenol	mg/Kg	--	--	0																
		4-Chloroaniline	mg/Kg	--	24	nc																
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0																
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc																
		4-Nitroaniline	mg/Kg	--	23	ca																
		4-Nitrophenol	mg/Kg	--	--	0																
		Acenaphthene	mg/Kg	--	368	nc																
		Acenaphthylene	mg/Kg	--	--	0																
		Anthracene	mg/Kg	--	2189	nc																
		Benzo(a)anthracene	mg/Kg	--	0.62	ca																
		Benzo(a)pyrene	mg/Kg	--	0.062	ca																
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca																
		Benzo(ghi)perylene	mg/Kg	--	--	0																
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca																
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0																
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca																
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-017-0001-SO		LL9SB-018-0001-SO		LL9SB-019-0001-SO		LL9SB-020-0001-SO		LL9SB-021-0001-SO		LL9SB-022-0001-SO		LL9SB-023-0001-SO		LL9SB-024-0001-SO	
							12/04/03		12/10/03		12/11/03		12/08/03		12/08/03		12/10/03		12/08/03		12/10/03	
							1 - 3 ft		1 - 3 ft		1 - 3 ft		1 - 3 ft		2.5 ft		1 - 3 ft		1 - 3 ft		1 - 3 ft	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc																
		Carbazole	mg/Kg	--	24	ca																
		Chrysene	mg/Kg	--	62	ca																
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca																
		Dibenzofuran	mg/Kg	--	15	nc																
		Diethyl phthalate	mg/Kg	--	4888	nc																
		Dimethyl phthalate	mg/Kg	--	100000	max																
		Di-n-butyl phthalate	mg/Kg	--	611	nc																
		Di-n-octyl phthalate	mg/Kg	--	244	nc																
		Fluoranthene	mg/Kg	--	229	nc																
		Fluorene	mg/Kg	--	275	nc																
		Hexachlorobenzene	mg/Kg	--	0.30	ca																
		Hexachlorobutadiene	mg/Kg	--	6.2	ca																
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc																
		Hexachloroethane	mg/Kg	--	35	ca																
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca																
		Isophorone	mg/Kg	--	512	ca																
		Naphthalene	mg/Kg	--	5.6	nc																
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca																
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca																
		Pentachlorophenol	mg/Kg	--	3.0	ca																
		Phenanthrene	mg/Kg	--	--	0																
		Phenol	mg/Kg	--	1833	nc																
TAL Metals 6010B	6010B	Pyrene	mg/Kg	--	232	nc																
		Aluminum	mg/Kg	19500	7614	nc	10000		14000		11000		15000		12000		11000		11000		18000	
		Barium	mg/Kg	124	538	nc	71		58		95		65		43		73		69		78	
		Beryllium	mg/Kg	0.88	15	nc	0.6		0.53		0.65		0.36	B	0.56		0.69		0.63		0.52	
		Cadmium	mg/Kg	0.00	3.7	nc	0.083	U	0.09	U	0.092	U	0.095	B	0.085	U	0.094	U	0.089	U	0.099	U
		Calcium	mg/Kg	35500	--	0	410		580		870		1700		970		6700		1200		650	
		Chromium	mg/Kg	27.2	30	ca	14		18		16		19		17		15		16		54	
		Cobalt	mg/Kg	23.2	902	ca	6.7		9.1		12		6.4		11		10		13		14	
		Copper	mg/Kg	32.3	313	nc	9.4		20		25		20	H	22	H	25		21	H	24	
		Iron	mg/Kg	35200	2346	nc	19000		26000		30000	H	24000		28000		24000		27000		28000	
		Magnesium	mg/Kg	8790	--	0	1600		3000		3200		2500		2700		3600		3300		4700	
		Manganese	mg/Kg	3030	176	nc	760		250		270		420		290		500		350		440	
		Nickel	mg/Kg	60.7	156	nc	12		24		29		15		23		25		29		31	
		Potassium	mg/Kg	3350	--	0	570		1200		1100		1200		1200		950		1300		1600	
		Selenium	mg/Kg	1.5	39	nc	0.67	B	0.45	U	0.3	B	0.45	U	0.63	B	0.47	U	0.45	U	0.5	U
		Silver	mg/Kg	0.00	39	nc	0.32	U	0.35	U	0.36	U	0.35	U	0.33	U	0.36	U	0.35	U	0.39	U
		Sodium	mg/Kg	145	--	0	90	U	97	U	100	U	98	U	93	U	100	U	97	U	110	U
		Vanadium	mg/Kg	37.6	7.8	nc	20		19		17		26		19		15		18		29	
		Zinc	mg/Kg	93.3	2346	nc	36		63		75		53		78		85		58		83	



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-017-0001-SO		LL9SB-018-0001-SO		LL9SB-019-0001-SO		LL9SB-020-0001-SO		LL9SB-021-0001-SO		LL9SB-022-0001-SO		LL9SB-023-0001-SO		LL9SB-024-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.85		0.82		0.49	U	0.85		0.52	B	0.53	U	0.49	U	1	
	7060A	Arsenic	mg/Kg	19.8	0.39	ca	10		15		32		14		26		18		24		9.2	
	7421	Lead	mg/Kg	19.1	400	pbk	10		12		19		12		22		23		12		86	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.029		0.035		0.18		0.18		0.021		0.087		0.018	B	0.079	
	7841	Thallium	mg/Kg	0.91	0.52	nc	0.2	U	0.21	U	0.21	U	0.21	U	0.33		0.23	U	0.21	U	0.24	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat																
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca																
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca																
		1,1-Dichloroethane	mg/Kg	--	51	nc																
		1,1-Dichloroethene	mg/Kg	--	12	nc																
		1,2-Dichloroethane	mg/Kg	--	0.28	ca																
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc																
		1,2-Dichloropropane	mg/Kg	--	0.34	ca																
		2-Butanone (MEK)	mg/Kg	--	2231	nc																
		2-Hexanone	mg/Kg	--	530	nc																
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc																
		Acetone	mg/Kg	--	1412	nc																
		Benzene	mg/Kg	--	0.64	ca																
		Bromodichloromethane	mg/Kg	--	0.82	ca																
		Bromoform	mg/Kg	--	62	ca																
		Bromomethane	mg/Kg	--	0.39	nc																
		Carbon disulfide	mg/Kg	--	36	nc																
		Carbon tetrachloride	mg/Kg	--	0.25	ca																
		Chlorobenzene	mg/Kg	--	15	nc																
		Chloroethane	mg/Kg	--	3.0	ca																
		Chloroform	mg/Kg	--	0.22	ca																
		Chloromethane	mg/Kg	--	4.7	nc																
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca																
		Dibromochloromethane	mg/Kg	--	1.1	ca																
		Ethylbenzene	mg/Kg	--	395	sat																
		Methylene chloride	mg/Kg	--	9.1	ca																
		Styrene	mg/Kg	--	1700	sat																
		Tetrachloroethene	mg/Kg	--	0.48	ca																
		Toluene	mg/Kg	--	520	sat																
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca																
		Trichloroethene	mg/Kg	--	0.053	ca																
		Vinyl chloride	mg/Kg	--	0.079	ca																
		Xylenes (total)	mg/Kg	--	0.079	ca																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-025-0001-SO		LL9SB-026-0001-SO		LL9SB-027-0001-SO		LL9SB-028-0001-SO		LL9SB-029-0001-SO		LL9SB-030-0001-SO		LL9SB-031-0001-SO		LL9SB-032-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc															0.21	U
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc													0.018	U	0.017	U
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc													0.018	U	0.018	U
		2,4,6-TNT	mg/Kg	0	16	ca													0.034	U	0.034	U
		2,4-Dinitrotoluene	mg/Kg	0	12	nc													0.036	U	0.035	U
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc													0.048	U*	0.047	U*
		2-Nitrotoluene	mg/Kg	0	0.88	ca													0.033	U	0.033	U
		3-Nitrotoluene	mg/Kg	0	73	nc													0.05	U	0.05	U
		4-Nitrotoluene	mg/Kg	0	12	ca													0.047	U	0.046	U
		HMX	mg/Kg	0	306	nc													0.11	U	0.11	U
		Nitrobenzene	mg/Kg	0	2	nc													0.022	U	0.022	U
		RDX	mg/Kg	0	4.4	ca													0.059	U	0.058	U
		Tetryl	mg/Kg	0	61	nc													0.043	U	0.043	U
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--																
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--																
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc															0.0022	U
		Aroclor 1221	mg/Kg	--	0.22	ca															0.0024	U
		Aroclor 1232	mg/Kg	--	0.22	ca															0.0024	U
		Aroclor 1242	mg/Kg	--	0.22	ca															0.0024	U
		Aroclor 1248	mg/Kg	--	0.22	ca															0.0024	U
		Aroclor 1254	mg/Kg	--	0.22	ca															0.0024	U
		Aroclor 1260	mg/Kg	--	0.22	ca															0.0016	U
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca															0.00042	U
		4,4'-DDE	mg/Kg	--	1.7	ca															0.00076	U
		4,4'-DDT	mg/Kg	--	1.7	ca															0.00044	U
		Aldrin	mg/Kg	--	0.029	ca															0.00015	U
		alpha-BHC	mg/Kg	--	0.09	sat															0.00019	U
		alpha-Chlordane	mg/Kg	--	1.6	ca															0.00014	U
		beta-BHC	mg/Kg	--	0.32	ca															0.00018	U
		delta-BHC	mg/Kg	--	--	0															0.00013	U
		Dieldrin	mg/Kg	--	0.030	ca															0.0004	U
		Endosulfan I	mg/Kg	--	37	nc															0.00032	U
		Endosulfan II	mg/Kg	--	37	nc															0.00033	U
		Endosulfan sulfate	mg/Kg	--	37	nc															0.00034	U
		Endrin	mg/Kg	--	1.8	nc															0.00051	U
		Endrin aldehyde	mg/Kg	--	--	0															0.00039	U
		Endrin ketone	mg/Kg	--	--	0															0.00034	U
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca															0.00027	U
		gamma-Chlordane	mg/Kg	--	1.6	ca															0.00018	U
		Heptachlor	mg/Kg	--	0.11	ca															0.00021	U
		Heptachlor epoxide	mg/Kg	--	0.053	ca															0.00016	U
		Methoxychlor	mg/Kg	--	31	nc															0.0027	U
		Toxaphene	mg/Kg	--	0	0															0.0054	U



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-025-0001-SO		LL9SB-026-0001-SO		LL9SB-027-0001-SO		LL9SB-028-0001-SO		LL9SB-029-0001-SO		LL9SB-030-0001-SO		LL9SB-031-0001-SO		LL9SB-032-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0															1.3	B J
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0															0.5	U
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc															0.25	U
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc															0.071	U
		1,2-Dichlorobenzene	mg/Kg	--	600	sat															0.095	U
		1,3-Dichlorobenzene	mg/Kg	--	53	nc															0.095	U
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca															0.086	U
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca															0.091	U
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc															0.046	U
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc															0.057	U
		2,4-Dichlorophenol	mg/Kg	--	18	nc															0.058	U
		2,4-Dimethylphenol	mg/Kg	--	122	nc															0.072	U
		2,4-Dinitrophenol	mg/Kg	--	12	nc															0.14	U
		2-Chloronaphthalene	mg/Kg	--	494	nc															0.058	U
		2-Chlorophenol	mg/Kg	--	6.3	nc															0.071	U
		2-Methylnaphthalene	mg/Kg	--	--	0															0.0018	U
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc															0.01	U
		2-Nitroaniline	mg/Kg	--	18.3	nc															0.041	U
		2-Nitrophenol	mg/Kg	--	--	0															0.076	U
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca															0.022	U
		3-Nitroaniline	mg/Kg	--	1.8	nc															0.13	U
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc															0.11	U*
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0															0.0037	U
		4-Chloro-3-methylphenol	mg/Kg	--	--	0															0.046	U
		4-Chloroaniline	mg/Kg	--	24	nc															0.12	U
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0															0.0043	U
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc															0.007	U
		4-Nitroaniline	mg/Kg	--	23	ca															0.047	U
		4-Nitrophenol	mg/Kg	--	--	0															0.099	U
		Acenaphthene	mg/Kg	--	368	nc															0.0017	U
		Acenaphthylene	mg/Kg	--	--	0															0.0011	U
		Anthracene	mg/Kg	--	2189	nc															0.001	U
		Benzo(a)anthracene	mg/Kg	--	0.62	ca															0.0031	J
		Benzo(a)pyrene	mg/Kg	--	0.062	ca															0.0027	U
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca															0.0045	J
		Benzo(ghi)perylene	mg/Kg	--	--	0															0.0033	J
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca															0.0034	U
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0															0.0035	U
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca															0.0024	U
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca															0.011	U



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-025-0001-SO		LL9SB-026-0001-SO		LL9SB-027-0001-SO		LL9SB-028-0001-SO		LL9SB-029-0001-SO		LL9SB-030-0001-SO		LL9SB-031-0001-SO		LL9SB-032-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc															0.0049	U
		Carbazole	mg/Kg	--	24	ca															0.042	U
		Chrysene	mg/Kg	--	62	ca															0.0031	J
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca															0.026	J
		Dibenzofuran	mg/Kg	--	15	nc															0.0033	U
		Diethyl phthalate	mg/Kg	--	4888	nc															0.0045	U
		Dimethyl phthalate	mg/Kg	--	100000	max															0.0043	U
		Di-n-butyl phthalate	mg/Kg	--	611	nc															0.024	U
		Di-n-octyl phthalate	mg/Kg	--	244	nc															0.01	U
		Fluoranthene	mg/Kg	--	229	nc															0.0064	J
		Fluorene	mg/Kg	--	275	nc															0.0019	U
		Hexachlorobenzene	mg/Kg	--	0.30	ca															0.0022	U
		Hexachlorobutadiene	mg/Kg	--	6.2	ca															0.004	U
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc															0.065	U
		Hexachloroethane	mg/Kg	--	35	ca															0.004	U
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca															0.018	J
		Isophorone	mg/Kg	--	512	ca															0.0029	U
		Naphthalene	mg/Kg	--	5.6	nc															0.0021	U
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca															0.0028	U
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca															0.0035	U
		Pentachlorophenol	mg/Kg	--	3.0	ca															0.12	U
		Phenanthrene	mg/Kg	--	--	0															0.0012	U
		Phenol	mg/Kg	--	1833	nc															0.0019	U
		Pyrene	mg/Kg	--	232	nc															0.004	J
TAL Metals 6010B	6010B	Aluminum	mg/Kg	19500	7614	nc	18000		13000		10000		9600		9400		10000		6400		11000	
		Barium	mg/Kg	124	538	nc	45		70		54		37		62		43		28		100	
		Beryllium	mg/Kg	0.88	15	nc	0.5		0.36	B	0.61		0.33	B	0.58		0.55		0.4	B	0.84	
		Cadmium	mg/Kg	0.00	3.7	nc	0.094	U	0.093	U	0.087	U	0.092	U	0.08	U	0.094	U	0.087	U	0.086	U
		Calcium	mg/Kg	35500	--	0	890		600		630		600		320		500		470		760	
		Chromium	mg/Kg	27.2	30	ca	23		15		13		11		14		13		9.9		13	
		Cobalt	mg/Kg	23.2	902	ca	8.5		8.1		6.7		2.4		8.7		5.9		5.5		11	
		Copper	mg/Kg	32.3	313	nc	23		8.1		14		5.2		9.1		11		6.3		12	
		Iron	mg/Kg	35200	2346	nc	32000		19000		24000		11000		24000		21000		16000		19000	
		Magnesium	mg/Kg	8790	--	0	4000		2200		1700		1200		1700		1700		1400		1600	
		Manganese	mg/Kg	3030	176	nc	180		470		410		50		660		380		390		970	
		Nickel	mg/Kg	60.7	156	nc	25		14		14		6.6		12		13		10		13	
		Potassium	mg/Kg	3350	--	0	1600		860		670		610		530		560		440		510	
		Selenium	mg/Kg	1.5	39	nc	0.47	U	0.47	U	0.43	U	0.46	U	0.65	B	0.58	B	0.43	U	0.43	U
		Silver	mg/Kg	0.00	39	nc	0.37	U	0.36	U	0.34	U	0.36	U	0.31	U	0.36	U	0.34	U	0.33	U
		Sodium	mg/Kg	145	--	0	100	U	100	U	94	U	100	U	87	U	100	U	94	U	93	U
		Vanadium	mg/Kg	37.6	7.8	nc	29		23		19		18		19		20		12		21	
		Zinc	mg/Kg	93.3	2346	nc	61		45		47		27		42		45		38		44	



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-025-0001-SO		LL9SB-026-0001-SO		LL9SB-027-0001-SO		LL9SB-028-0001-SO		LL9SB-029-0001-SO		LL9SB-030-0001-SO		LL9SB-031-0001-SO		LL9SB-032-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.7	B	0.56	B	0.52	U	0.47	U	0.46	U	0.46	U	0.47	U	0.46	U
	7060A	Arsenic	mg/Kg	19.8	0.39	ca	15		6.9		18		5.1		11		6		7.4		7.4	
	7421	Lead	mg/Kg	19.1	400	pbk	11		14		14		5.5		11		8.8		6.5		15	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.0095	B	0.024		0.035		0.032		0.038		0.038		0.016	B	0.015	B
	7841	Thallium	mg/Kg	0.91	0.52	nc	0.22	U	0.22	U	1.1	U	0.2	U	0.99	U	0.2	U	1	U	0.2	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat															0.0013	U
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca															0.0012	U
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca															0.0013	U
		1,1-Dichloroethane	mg/Kg	--	51	nc															0.0012	U
		1,1-Dichloroethene	mg/Kg	--	12	nc															0.0016	U
		1,2-Dichloroethane	mg/Kg	--	0.28	ca															0.0011	U
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc															0.0026	U
		1,2-Dichloropropane	mg/Kg	--	0.34	ca															0.0012	U
		2-Butanone (MEK)	mg/Kg	--	2231	nc															0.0047	U
		2-Hexanone	mg/Kg	--	530	nc															0.0013	U
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc															0.0012	U
		Acetone	mg/Kg	--	1412	nc															0.0056	U
		Benzene	mg/Kg	--	0.64	ca															0.0013	U
		Bromodichloromethane	mg/Kg	--	0.82	ca															0.0012	U
		Bromoform	mg/Kg	--	62	ca															0.00091	U
		Bromomethane	mg/Kg	--	0.39	nc															0.0016	U*
		Carbon disulfide	mg/Kg	--	36	nc															0.0015	U
		Carbon tetrachloride	mg/Kg	--	0.25	ca															0.0013	U
		Chlorobenzene	mg/Kg	--	15	nc															0.0013	U
		Chloroethane	mg/Kg	--	3.0	ca															0.0012	U
		Chloroform	mg/Kg	--	0.22	ca															0.0013	U
		Chloromethane	mg/Kg	--	4.7	nc															0.0013	U
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca															0.0011	U
		Dibromochloromethane	mg/Kg	--	1.1	ca															0.00096	U
		Ethylbenzene	mg/Kg	--	395	sat															0.0013	U
		Methylene chloride	mg/Kg	--	9.1	ca															0.0035	U
		Styrene	mg/Kg	--	1700	sat															0.0013	U
		Tetrachloroethene	mg/Kg	--	0.48	ca															0.0015	U
		Toluene	mg/Kg	--	520	sat															0.0013	U
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca															0.00096	U
		Trichloroethene	mg/Kg	--	0.053	ca															0.0013	U
		Vinyl chloride	mg/Kg	--	0.079	ca															0.0013	U
		Xylenes (total)	mg/Kg	--	0.079	ca															0.0041	U



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-040-0001-SO		LL9SB-041-0001-SO		LL9SB-042-0001-SO		LL9SB-043-0001-DUP		LL9SB-043-0001-SO		LL9SB-044-0001-SO		LL9SB-045-0001-SO		LL9SB-046-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc	0.17	U	0.17	U			0.077	U^	0.13	U						
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc	0.018	U	0.017	U			0.017	U	0.017	U						
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc	0.018	U	0.018	U			0.018	U	0.018	U						
		2,4,6-TNT	mg/Kg	0	16	ca	0.034	U	0.034	U			0.034	U	0.033	U						
		2,4-Dinitrotoluene	mg/Kg	0	12	nc	0.036	U	0.035	U			0.035	U	0.035	U						
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc	0.048	U	0.047	U			0.047	U	0.047	U						
		2-Nitrotoluene	mg/Kg	0	0.88	ca	0.033	U	0.033	U			0.033	U	0.033	U						
		3-Nitrotoluene	mg/Kg	0	73	nc	0.05	U	0.05	U			0.05	U	0.05	U						
		4-Nitrotoluene	mg/Kg	0	12	ca	0.047	U	0.046	U			0.046	U	0.046	U						
		HMX	mg/Kg	0	306	nc	0.11	U	0.11	U			0.11	U	0.11	U						
		Nitrobenzene	mg/Kg	0	2	nc	0.022	U	0.022	U			0.022	U	0.022	U						
		RDX	mg/Kg	0	4.4	ca	0.059	U	0.058	U			0.058	U	0.058	U						
		Tetryl	mg/Kg	0	61	nc	0.043	U	0.043	U			0.043	U	0.043	U						
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--																
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--																
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc	0.0022	U	0.0022	U			0.018	U	0.019	U						
		Aroclor 1221	mg/Kg	--	0.22	ca	0.0023	U	0.0023	U			0.018	U	0.019	U						
		Aroclor 1232	mg/Kg	--	0.22	ca	0.0023	U	0.0023	U			0.018	U	0.019	U						
		Aroclor 1242	mg/Kg	--	0.22	ca	0.0023	U	0.0023	U			0.018	U	0.019	U						
		Aroclor 1248	mg/Kg	--	0.22	ca	0.0023	U	0.0023	U			0.018	U	0.019	U						
		Aroclor 1254	mg/Kg	--	0.22	ca	0.0023	U	0.0023	U			0.018	U	0.019	U						
		Aroclor 1260	mg/Kg	--	0.22	ca	0.0016	U	0.0016	U			0.018	U	0.019	U						
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca	0.0021	U	0.00042	U			0.0004	U	0.0004	U						
		4,4'-DDE	mg/Kg	--	1.7	ca	0.0038	U	0.00075	U			0.00072	U	0.00073	U						
		4,4'-DDT	mg/Kg	--	1.7	ca	0.0021	U	0.00043	U			0.00041	U	0.00041	U						
		Aldrin	mg/Kg	--	0.029	ca	0.00075	U	0.00015	U			0.00014	U	0.00015	U						
		alpha-BHC	mg/Kg	--	0.09	sat	0.00093	U	0.00018	U			0.00018	U	0.00018	U						
		alpha-Chlordane	mg/Kg	--	1.6	ca	0.0007	U	0.00014	U			0.00013	U	0.00013	U						
		beta-BHC	mg/Kg	--	0.32	ca	0.00087	U	0.00017	U			0.00017	U	0.00017	U						
		delta-BHC	mg/Kg	--	--	0	0.00064	U	0.00013	U			0.00012	U	0.00012	U						
		Dieldrin	mg/Kg	--	0.030	ca	0.002	U	0.00039	U			0.00037	U	0.00038	U						
		Endosulfan I	mg/Kg	--	37	nc	0.0016	U	0.00031	U			0.0003	U	0.0003	U						
		Endosulfan II	mg/Kg	--	37	nc	0.0016	U	0.00032	U			0.00031	U	0.00031	U						
		Endosulfan sulfate	mg/Kg	--	37	nc	0.0017	U	0.00034	U			0.00032	U	0.00032	U						
		Endrin	mg/Kg	--	1.8	nc	0.0025	U	0.0005	U			0.00047	U	0.00048	U						
		Endrin aldehyde	mg/Kg	--	--	0	0.0019	U	0.00038	U			0.00036	U	0.00037	U						
		Endrin ketone	mg/Kg	--	--	0	0.0017	U	0.00034	U			0.00032	U	0.00032	U						
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca	0.0013	U	0.00027	U			0.00025	U	0.00026	U						
		gamma-Chlordane	mg/Kg	--	1.6	ca	0.00087	U	0.00017	U			0.00017	U	0.00017	U						
		Heptachlor	mg/Kg	--	0.11	ca	0.001	U	0.00021	U			0.0002	U	0.0002	U						
		Heptachlor epoxide	mg/Kg	--	0.053	ca	0.00081	U	0.00016	U			0.00015	U	0.00016	U						
		Methoxychlor	mg/Kg	--	31	nc	0.013	U	0.0027	U			0.0025	U	0.0026	U						
				Toxaphene	mg/Kg	--	0	0	0.027	U	0.0053	U			0.0051	U	0.0051	U				



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-040-0001-SO		LL9SB-041-0001-SO		LL9SB-042-0001-SO		LL9SB-043-0001-DUP		LL9SB-043-0001-SO		LL9SB-044-0001-SO		LL9SB-045-0001-SO		LL9SB-046-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0							2.3	U	2.3	U						
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0							0.5	U	0.5	U						
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc							0.25	U	0.25	U						
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc	0.068	U	0.068	U			0.065	U	0.065	U						
		1,2-Dichlorobenzene	mg/Kg	--	600	sat	0.091	U	0.091	U			0.087	U	0.087	U						
		1,3-Dichlorobenzene	mg/Kg	--	53	nc	0.091	U	0.091	U			0.087	U	0.087	U						
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca	0.082	U	0.082	U			0.078	U	0.078	U						
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca	0.086	U	0.086	U			0.083	U	0.083	U						
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc	0.044	U	0.044	U			0.042	U	0.042	U						
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc	0.054	U	0.054	U			0.052	U	0.052	U						
		2,4-Dichlorophenol	mg/Kg	--	18	nc	0.055	U	0.055	U			0.053	U	0.053	U						
		2,4-Dimethylphenol	mg/Kg	--	122	nc	0.069	U	0.069	U			0.066	U	0.066	U						
		2,4-Dinitrophenol	mg/Kg	--	12	nc	0.13	U	0.13	U			0.13	U	0.13	U						
		2-Chloronaphthalene	mg/Kg	--	494	nc	0.055	U	0.055	U			0.053	U	0.053	U						
		2-Chlorophenol	mg/Kg	--	6.3	nc	0.068	U	0.068	U			0.065	U	0.065	U						
		2-Methylnaphthalene	mg/Kg	--	--	0	0.0017	U	0.0017	U			0.0017	U	0.0017	U						
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc	0.0096	U	0.0097	U			0.0092	U	0.0093	U						
		2-Nitroaniline	mg/Kg	--	18.3	nc	0.039	U	0.039	U			0.037	U*	0.038	U*						
		2-Nitrophenol	mg/Kg	--	--	0	0.072	U	0.072	U			0.069	U	0.07	U						
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca	0.021	U	0.021	U			0.02	U	0.02	U						
		3-Nitroaniline	mg/Kg	--	1.8	nc	0.13	U	0.13	U			0.12	U	0.12	U						
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc	0.11	U	0.11	U			0.1	U*	0.1	U*						
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0	0.0036	U	0.0036	U			0.0034	U*	0.0034	U*						
		4-Chloro-3-methylphenol	mg/Kg	--	--	0	0.044	U	0.044	U			0.042	U	0.042	U						
		4-Chloroaniline	mg/Kg	--	24	nc	0.11	U	0.12	U			0.11	U	0.11	U						
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0	0.0041	U	0.0041	U			0.004	U*	0.004	U*						
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc	0.0067	U	0.0067	U			0.0064	U	0.0064	U						
		4-Nitroaniline	mg/Kg	--	23	ca	0.045	U	0.045	U			0.043	U	0.043	U						
		4-Nitrophenol	mg/Kg	--	--	0	0.094	U	0.094	U			0.09	U	0.091	U						
		Acenaphthene	mg/Kg	--	368	nc	0.0016	U	0.0016	U			0.0015	U	0.0015	U						
		Acenaphthylene	mg/Kg	--	--	0	0.001	U	0.001	U			0.001	U*	0.001	U*						
		Anthracene	mg/Kg	--	2189	nc	0.00099	U	0.00099	U			0.00095	U	0.00095	U						
		Benzo(a)anthracene	mg/Kg	--	0.62	ca	0.053		0.0013	U			0.0012	U	0.0012	U						
		Benzo(a)pyrene	mg/Kg	--	0.062	ca	0.052		0.0025	U			0.0024	U	0.0024	U						
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca	0.062		0.0024	U			0.0023	U	0.0023	U						
		Benzo(ghi)perylene	mg/Kg	--	--	0	0.034	Ja	0.0022	U			0.0021	U	0.0021	U						
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca	0.059	M	0.0032	U			0.0031	U	0.0031	U						
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0	0.0033	U	0.0033	U			0.0032	U	0.0032	U						
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca	0.0023	U	0.0023	U			0.0022	U	0.0022	U						
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca	0.011	U	0.011	U			0.01	U	0.01	U						



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-040-0001-SO		LL9SB-041-0001-SO		LL9SB-042-0001-SO		LL9SB-043-0001-DUP		LL9SB-043-0001-SO		LL9SB-044-0001-SO		LL9SB-045-0001-SO		LL9SB-046-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc	0.0047	U	0.0047	U			0.0045	U	0.0045	U						
		Carbazole	mg/Kg	--	24	ca	0.04	U	0.04	U			0.039	U	0.039	U						
		Chrysene	mg/Kg	--	62	ca	0.058		0.0021	U			0.002	U	0.002	U						
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca	0.018	Ja	0.0025	U			0.0024	U	0.0024	U						
		Dibenzofuran	mg/Kg	--	15	nc	0.0031	U	0.0031	U			0.003	U	0.003	U						
		Diethyl phthalate	mg/Kg	--	4888	nc	0.0043	U	0.0043	U			0.0041	U*	0.0041	U*						
		Dimethyl phthalate	mg/Kg	--	100000	max	0.0041	U	0.0041	U			0.004	U*	0.004	U*						
		Di-n-butyl phthalate	mg/Kg	--	611	nc	0.14	Ja	0.087	Ja			0.022	U	0.022	U						
		Di-n-octyl phthalate	mg/Kg	--	244	nc	0.01	U	0.01	U			0.0096	U	0.0096	U						
		Fluoranthene	mg/Kg	--	229	nc	0.17		0.0013	U			0.0012	U	0.0012	U						
		Fluorene	mg/Kg	--	275	nc	0.0018	U	0.0018	U			0.0018	U	0.0018	U						
		Hexachlorobenzene	mg/Kg	--	0.30	ca	0.0021	U	0.0021	U			0.002	U*	0.002	U*						
		Hexachlorobutadiene	mg/Kg	--	6.2	ca	0.0038	U	0.0038	U			0.0036	U	0.0036	U						
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc	0.062	U	0.062	U			0.059	U	0.06	U						
		Hexachloroethane	mg/Kg	--	35	ca	0.0038	U	0.0038	U			0.0036	U	0.0036	U						
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca	0.034	Ja	0.0024	U			0.0023	U	0.0023	U						
		Isophorone	mg/Kg	--	512	ca	0.0028	U	0.0028	U			0.0026	U	0.0027	U						
		Naphthalene	mg/Kg	--	5.6	nc	0.002	U	0.002	U			0.0019	U	0.0019	U						
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca	0.0026	U	0.0026	U			0.0025	U	0.0025	U						
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca	0.0033	U	0.0033	U			0.0032	U	0.0032	U						
		Pentachlorophenol	mg/Kg	--	3.0	ca	0.11	U	0.12	U			0.11	U	0.11	U						
		Phenanthrene	mg/Kg	--	--	0	0.061		0.0012	U			0.0011	U	0.0011	U						
		Phenol	mg/Kg	--	1833	nc	0.0018	U	0.0018	U			0.0018	U	0.0018	U						
		Pyrene	mg/Kg	--	232	nc	0.12		0.0023	U			0.0022	U	0.0022	U						
TAL Metals 6010B	6010B	Aluminum	mg/Kg	19500	7614	nc	14000		8900		9100		4300		4800		8000		9300		6700	
		Barium	mg/Kg	124	538	nc	140		57		57		24		26		32		34		26	
		Beryllium	mg/Kg	0.88	15	nc	1.1		1.2		0.43	B	0.4		0.42		0.35	B	0.43		0.29	B
		Cadmium	mg/Kg	0.00	3.7	nc	0.083	U	0.086	U	0.087	U	0.078	U	0.084	U	0.08	U	0.085	U	0.092	U
		Calcium	mg/Kg	35500	--	0	36000	H	6900	H	480		420		420		1500		700		760	
		Chromium	mg/Kg	27.2	30	ca	17		18		14		8.8		9.7		13		14		10	
		Cobalt	mg/Kg	23.2	902	ca	6.1		19		8.7		6.5		6.8		10		11		7.7	
		Copper	mg/Kg	32.3	313	nc	15		20		13		6.7		7.4		19		20		18	
		Iron	mg/Kg	35200	2346	nc	39000	*H	51000	*H	23000		26000		27000		24000		27000		21000	
		Magnesium	mg/Kg	8790	--	0	5000		3000		2200		1400		1500		2800		3100		2200	
		Manganese	mg/Kg	3030	176	nc	2300		2100		420		800		730		420		330		360	
		Nickel	mg/Kg	60.7	156	nc	11		37		17		12		12		25		23		17	
		Potassium	mg/Kg	3350	--	0	1200		1700		810		520		560		1100		900		910	
		Selenium	mg/Kg	1.5	39	nc	0.42	U	0.43	U	0.43	U	0.39	U	0.42	U	0.4	U	0.5	B	0.46	U
		Silver	mg/Kg	0.00	39	nc	0.32	U	0.33	U	0.34	U	0.3	U	0.33	U	0.31	U	0.33	U	0.36	U
		Sodium	mg/Kg	145	--	0	170		93	U	1200		840		860		1100		1200		1100	
		Vanadium	mg/Kg	37.6	7.8	nc	17		16		15		9.6		10		13		14		12	
		Zinc	mg/Kg	93.3	2346	nc	68	H	64	H	52		51		56		58		60		55	



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-040-0001-SO		LL9SB-041-0001-SO		LL9SB-042-0001-SO		LL9SB-043-0001-DUP		LL9SB-043-0001-SO		LL9SB-044-0001-SO		LL9SB-045-0001-SO		LL9SB-046-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.47	U	0.49	U	0.44	U	0.44	U	0.48	U	0.5	B	0.5	U	0.54	B
	7060A	Arsenic	mg/Kg	19.8	0.39	ca	12		3.4		6.9		3.4		3.7		15		18		17	
	7421	Lead	mg/Kg	19.1	400	pbk	51		31		11		5.6		6		9.3		11		9	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.26		0.017	B	0.02		0.012	B	0.018	B	0.014	B	0.014	B	0.013	B
	7841	Thallium	mg/Kg	0.91	0.52	nc	0.2	U	0.21	U	0.95	U	0.96	U	1	U	1	U	1.1	U	0.97	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat	0.00071	U	0.00072	U			0.02	U	0.0007	U						
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca	0.00075	U	0.00075	U			0.022	U	0.00073	U						
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca	0.00083	U	0.00084	U			0.038	U	0.00081	U						
		1,1-Dichloroethane	mg/Kg	--	51	nc	0.001	U	0.001	U			0.016	U	0.001	U						
		1,1-Dichloroethene	mg/Kg	--	12	nc	0.0012	U	0.0012	U			0.017	U	0.0011	U						
		1,2-Dichloroethane	mg/Kg	--	0.28	ca	0.00068	U	0.00068	U			0.026	U	0.00066	U						
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc	0.0022	U	0.0022	U			0.035	U	0.0022	U						
		1,2-Dichloropropane	mg/Kg	--	0.34	ca	0.0011	U	0.0011	U			0.021	U	0.0011	U						
		2-Butanone (MEK)	mg/Kg	--	2231	nc	0.0049	U	0.0049	U			0.062	U	0.0048	U						
		2-Hexanone	mg/Kg	--	530	nc	0.002	U	0.002	U			0.063	U	0.0019	U						
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc	0.0035	U	0.0035	U			0.045	U	0.0034	U						
		Acetone	mg/Kg	--	1412	nc	0.0048	U	0.0048	U			0.035	U	0.0047	U						
		Benzene	mg/Kg	--	0.64	ca	0.00077	U	0.00078	U			0.017	U	0.00075	U						
		Bromodichloromethane	mg/Kg	--	0.82	ca	0.0008	U	0.0008	U			0.023	U	0.00078	U						
		Bromoform	mg/Kg	--	62	ca	0.0011	U	0.0011	U			0.022	U	0.001	U						
		Bromomethane	mg/Kg	--	0.39	nc	0.0034	U	0.0034	U			0.013	U	0.0033	U						
		Carbon disulfide	mg/Kg	--	36	nc	0.0023	U	0.0024	U			0.025	U	0.0023	U						
		Carbon tetrachloride	mg/Kg	--	0.25	ca	0.00097	U	0.00098	U			0.02	U	0.00095	U						
		Chlorobenzene	mg/Kg	--	15	nc	0.0011	U	0.0011	U			0.027	U	0.001	U						
		Chloroethane	mg/Kg	--	3.0	ca	0.0019	U	0.0019	U			0.024	U	0.0018	U						
		Chloroform	mg/Kg	--	0.22	ca	0.00073	U	0.00073	U			0.022	U	0.00071	U						
		Chloromethane	mg/Kg	--	4.7	nc	0.0011	U	0.0011	U			0.028	U	0.0011	U						
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.00093	U	0.00093	U			0.027	U	0.0009	U						
		Dibromochloromethane	mg/Kg	--	1.1	ca	0.00081	U	0.00081	U			0.023	U	0.00079	U						
		Ethylbenzene	mg/Kg	--	395	sat	0.0013	U	0.0013	U			0.027	U	0.0013	U						
		Methylene chloride	mg/Kg	--	9.1	ca	0.0021	U	0.0021	U			0.024	U	0.0021	U						
		Styrene	mg/Kg	--	1700	sat	0.0012	U	0.0012	U			0.034	U	0.0011	U						
		Tetrachloroethene	mg/Kg	--	0.48	ca	0.00078	U	0.00079	U			0.028	U	0.00076	U						
		Toluene	mg/Kg	--	520	sat	0.0012	U	0.0012	U			0.022	U	0.0011	U						
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.00098	U	0.00099	U			0.024	U	0.00096	U						
		Trichloroethene	mg/Kg	--	0.053	ca	0.00069	U	0.00069	U			0.026	U	0.00067	U						
		Vinyl chloride	mg/Kg	--	0.079	ca	0.00087	U	0.00087	U			0.022	U	0.00084	U						
		Xylenes (total)	mg/Kg	--	0.079	ca	0.0034	U	0.0034	U			0.06	U	0.0033	U						



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-047-0001-SO		LL9SB-048-0001-SO		LL9SB-049-0001-SO		LL9SB-050-0001-SO		LL9SB-051-0001-SO		LL9SB-052-0001-SO		LL9SB-053-0001-DUP		LL9SB-053-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc							0.19	U								
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc							0.018	U					0.017	U	0.017	U
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc							0.018	U					0.018	U	0.017	U
		2,4,6-TNT	mg/Kg	0	16	ca							0.034	U					0.033	U	0.033	U
		2,4-Dinitrotoluene	mg/Kg	0	12	nc							0.036	U					0.035	U	0.035	U
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc							0.048	U					0.047	U	0.046	U
		2-Nitrotoluene	mg/Kg	0	0.88	ca							0.033	U					0.033	U	0.032	U
		3-Nitrotoluene	mg/Kg	0	73	nc							0.05	U					0.049	U	0.049	U
		4-Nitrotoluene	mg/Kg	0	12	ca							0.047	U					0.046	U	0.045	U
		HMX	mg/Kg	0	306	nc							0.11	U					0.11	U	0.11	U
		Nitrobenzene	mg/Kg	0	2	nc							0.022	U					0.022	U	0.022	U
		RDX	mg/Kg	0	4.4	ca							0.059	U					0.058	U	0.11	
		Tetryl	mg/Kg	0	61	nc							0.043	U					0.043	U	0.042	U
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--																
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--																
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc							0.0022	U								
		Aroclor 1221	mg/Kg	--	0.22	ca							0.0023	U								
		Aroclor 1232	mg/Kg	--	0.22	ca							0.0023	U								
		Aroclor 1242	mg/Kg	--	0.22	ca							0.0023	U								
		Aroclor 1248	mg/Kg	--	0.22	ca							0.0023	U								
		Aroclor 1254	mg/Kg	--	0.22	ca							0.0023	U								
		Aroclor 1260	mg/Kg	--	0.22	ca							0.0016	U								
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca							0.00042	U								
		4,4'-DDE	mg/Kg	--	1.7	ca							0.00075	U								
		4,4'-DDT	mg/Kg	--	1.7	ca							0.00043	U								
		Aldrin	mg/Kg	--	0.029	ca							0.00015	U								
		alpha-BHC	mg/Kg	--	0.09	sat							0.00019	U								
		alpha-Chlordane	mg/Kg	--	1.6	ca							0.00014	U								
		beta-BHC	mg/Kg	--	0.32	ca							0.00017	U								
		delta-BHC	mg/Kg	--	--	0							0.00013	U								
		Dieldrin	mg/Kg	--	0.030	ca							0.00039	U								
		Endosulfan I	mg/Kg	--	37	nc							0.00031	U								
		Endosulfan II	mg/Kg	--	37	nc							0.00033	U								
		Endosulfan sulfate	mg/Kg	--	37	nc							0.00034	U								
		Endrin	mg/Kg	--	1.8	nc							0.0005	U								
		Endrin aldehyde	mg/Kg	--	--	0							0.00038	U								
		Endrin ketone	mg/Kg	--	--	0							0.00034	U								
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca							0.00027	U								
		gamma-Chlordane	mg/Kg	--	1.6	ca							0.00017	U								
		Heptachlor	mg/Kg	--	0.11	ca							0.00021	U								
		Heptachlor epoxide	mg/Kg	--	0.053	ca							0.00016	U								
		Methoxychlor	mg/Kg	--	31	nc							0.0027	U								
		Toxaphene	mg/Kg	--	0	0							0.0053	U								



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-047-0001-SO		LL9SB-048-0001-SO		LL9SB-049-0001-SO		LL9SB-050-0001-SO		LL9SB-051-0001-SO		LL9SB-052-0001-SO		LL9SB-053-0001-DUP		LL9SB-053-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0							4.3	J								
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0							0.5	U								
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc							0.25	U								
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc							0.07	U								
		1,2-Dichlorobenzene	mg/Kg	--	600	sat							0.094	U								
		1,3-Dichlorobenzene	mg/Kg	--	53	nc							0.094	U								
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca							0.084	U								
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca							0.089	U								
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc							0.045	U								
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc							0.056	U								
		2,4-Dichlorophenol	mg/Kg	--	18	nc							0.057	U								
		2,4-Dimethylphenol	mg/Kg	--	122	nc							0.071	U								
		2,4-Dinitrophenol	mg/Kg	--	12	nc							0.14	U								
		2-Chloronaphthalene	mg/Kg	--	494	nc							0.057	U								
		2-Chlorophenol	mg/Kg	--	6.3	nc							0.07	U								
		2-Methylnaphthalene	mg/Kg	--	--	0							0.0018	U								
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc							0.01	U								
		2-Nitroaniline	mg/Kg	--	18.3	nc							0.04	U								
		2-Nitrophenol	mg/Kg	--	--	0							0.075	U								
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca							0.021	U								
		3-Nitroaniline	mg/Kg	--	1.8	nc							0.13	U								
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc							0.11	U								
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0							0.0037	U								
		4-Chloro-3-methylphenol	mg/Kg	--	--	0							0.045	U								
		4-Chloroaniline	mg/Kg	--	24	nc							0.12	U								
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0							0.0043	U								
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc							0.0069	U								
		4-Nitroaniline	mg/Kg	--	23	ca							0.046	U								
		4-Nitrophenol	mg/Kg	--	--	0							0.097	U								
		Acenaphthene	mg/Kg	--	368	nc							0.019	Ja								
		Acenaphthylene	mg/Kg	--	--	0							0.0011	U								
		Anthracene	mg/Kg	--	2189	nc							0.03	Ja								
		Benzo(a)anthracene	mg/Kg	--	0.62	ca							0.041									
		Benzo(a)pyrene	mg/Kg	--	0.062	ca							0.042									
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca							0.038	Ja								
		Benzo(ghi)perylene	mg/Kg	--	--	0							0.033	Ja								
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca							0.032	Ja								
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0							0.0034	U								
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca							0.0024	U								
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca							0.011	U								



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-047-0001-SO		LL9SB-048-0001-SO		LL9SB-049-0001-SO		LL9SB-050-0001-SO		LL9SB-051-0001-SO		LL9SB-052-0001-SO		LL9SB-053-0001-DUP		LL9SB-053-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc							0.0049	U								
		Carbazole	mg/Kg	--	24	ca							0.042	U								
		Chrysene	mg/Kg	--	62	ca							0.042									
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca							0.043									
		Dibenzofuran	mg/Kg	--	15	nc							0.0079	Ja								
		Diethyl phthalate	mg/Kg	--	4888	nc							0.0044	U								
		Dimethyl phthalate	mg/Kg	--	100000	max							0.0043	U								
		Di-n-butyl phthalate	mg/Kg	--	611	nc							0.024	U								
		Di-n-octyl phthalate	mg/Kg	--	244	nc							0.01	U								
		Fluoranthene	mg/Kg	--	229	nc							0.13									
		Fluorene	mg/Kg	--	275	nc							0.015	Ja								
		Hexachlorobenzene	mg/Kg	--	0.30	ca							0.0021	U								
		Hexachlorobutadiene	mg/Kg	--	6.2	ca							0.0039	U								
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc							0.064	U								
		Hexachloroethane	mg/Kg	--	35	ca							0.0039	U								
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca							0.044									
		Isophorone	mg/Kg	--	512	ca							0.0028	U								
		Naphthalene	mg/Kg	--	5.6	nc							0.002	U								
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca							0.0027	U								
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca							0.0034	U								
		Pentachlorophenol	mg/Kg	--	3.0	ca							0.12	U								
		Phenanthrene	mg/Kg	--	--	0							0.1									
		Phenol	mg/Kg	--	1833	nc							0.0019	U								
		Pyrene	mg/Kg	--	232	nc							0.1									
TAL Metals 6010B	6010B	Aluminum	mg/Kg	19500	7614	nc	8600		8400		12000		12000		11000		16000		6900		7400	
		Barium	mg/Kg	124	538	nc	61		51		150		56		64		66		54		57	
		Beryllium	mg/Kg	0.88	15	nc	0.16	B	0.4	B	0.47		0.33	B	0.55		0.55		0.19	B	0.24	B
		Cadmium	mg/Kg	0.00	3.7	nc	0.082	U	0.085	U	0.089	U	0.092	U	0.11	B	0.089	U	0.08	U	0.09	U
		Calcium	mg/Kg	35500	--	0	1000		5100		990		970		6800		850		1300		790	
		Chromium	mg/Kg	27.2	30	ca	11		12		24		16		15		21		8.1		8.8	
		Cobalt	mg/Kg	23.2	902	ca	1.6		9		9.2		6.8		9.6		7.4		5.4		6.2	
		Copper	mg/Kg	32.3	313	nc	8.7		20		21		11		21	H	22	H	11		12	
		Iron	mg/Kg	35200	2346	nc	9600		21000		27000		23000		24000		29000		13000	H	15000	H
		Magnesium	mg/Kg	8790	--	0	1300		2500		2500		2200		3100		3600		1400		1500	
		Manganese	mg/Kg	3030	176	nc	92		590		530		370		350		190		500		550	
		Nickel	mg/Kg	60.7	156	nc	9.6		18		21		13		22		20		11		13	
		Potassium	mg/Kg	3350	--	0	740		850		1100		1000		1400		1600		670		700	
		Selenium	mg/Kg	1.5	39	nc	0.41	U	0.43	U	0.44	U	0.46	U	0.44	U	0.44	U	0.46	B	0.41	B
		Silver	mg/Kg	0.00	39	nc	0.32	U	0.33	U	0.34	U	0.36	U	0.34	U	0.34	U	0.31	U	0.35	U
		Sodium	mg/Kg	145	--	0	1200		92	U	96	U	100	U	95	U	120		86	U	97	U
		Vanadium	mg/Kg	37.6	7.8	nc	9.2		14		20		22		19		25		14		13	
		Zinc	mg/Kg	93.3	2346	nc	50		82		290		40		59		53		47		45	



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-047-0001-SO		LL9SB-048-0001-SO		LL9SB-049-0001-SO		LL9SB-050-0001-SO		LL9SB-051-0001-SO		LL9SB-052-0001-SO		LL9SB-053-0001-DUP		LL9SB-053-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc	0.49	U	0.47	U	0.49	U	0.51	U	0.51	U	0.52	U	0.45	U	0.46	U
	7060A	Arsenic	mg/Kg	19.8	0.39	ca	5.9		11		12		10		18		16		9.4		11	
	7421	Lead	mg/Kg	19.1	400	pbk	5.2		41		64		15		13		9.3		10		12	
	7471A	Mercury	mg/Kg	0.04	2.3	nc	0.03		0.34		0.036		0.027		0.027		0.017	B	0.023		0.024	
	7841	Thallium	mg/Kg	0.91	0.52	nc	1	U	0.2	U	0.21	U	0.22	U	0.22	U	0.22	U	0.19	U	0.2	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat							0.0013	U								
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca							0.0012	U								
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca							0.0013	U								
		1,1-Dichloroethane	mg/Kg	--	51	nc							0.0012	U								
		1,1-Dichloroethene	mg/Kg	--	12	nc							0.0016	U								
		1,2-Dichloroethane	mg/Kg	--	0.28	ca							0.0011	U								
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc							0.0025	U								
		1,2-Dichloropropane	mg/Kg	--	0.34	ca							0.0012	U								
		2-Butanone (MEK)	mg/Kg	--	2231	nc							0.0047	U								
		2-Hexanone	mg/Kg	--	530	nc							0.0013	U								
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc							0.0012	U								
		Acetone	mg/Kg	--	1412	nc							0.0055	U								
		Benzene	mg/Kg	--	0.64	ca							0.0013	U								
		Bromodichloromethane	mg/Kg	--	0.82	ca							0.0012	U								
		Bromoform	mg/Kg	--	62	ca							0.0009	U								
		Bromomethane	mg/Kg	--	0.39	nc							0.0016	U								
		Carbon disulfide	mg/Kg	--	36	nc							0.0014	U								
		Carbon tetrachloride	mg/Kg	--	0.25	ca							0.0013	U								
		Chlorobenzene	mg/Kg	--	15	nc							0.0013	U								
		Chloroethane	mg/Kg	--	3.0	ca							0.0012	U								
		Chloroform	mg/Kg	--	0.22	ca							0.0013	U								
		Chloromethane	mg/Kg	--	4.7	nc							0.0013	U								
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca							0.0011	U								
		Dibromochloromethane	mg/Kg	--	1.1	ca							0.00095	U								
		Ethylbenzene	mg/Kg	--	395	sat							0.0013	U								
		Methylene chloride	mg/Kg	--	9.1	ca							0.0035	U								
		Styrene	mg/Kg	--	1700	sat							0.0013	U								
		Tetrachloroethene	mg/Kg	--	0.48	ca							0.0014	U								
		Toluene	mg/Kg	--	520	sat							0.0013	U								
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca							0.00095	U								
		Trichloroethene	mg/Kg	--	0.053	ca							0.0013	U								
		Vinyl chloride	mg/Kg	--	0.079	ca							0.0013	U								
		Xylenes (total)	mg/Kg	--	0.079	ca							0.0041	U								



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-055-0001-SO		LL9SB-056-0001-SO		LL9SB-059-0001-SO		LL9SB-061-0001-SO		LL9SB-065-0001-SO		LL9SB-066-0001-SO		LL9SB-069-0001-SO		LL9SB-070-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc																
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc																
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc																
		2,4,6-TNT	mg/Kg	0	16	ca																
		2,4-Dinitrotoluene	mg/Kg	0	12	nc																
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc																
		2-Nitrotoluene	mg/Kg	0	0.88	ca																
		3-Nitrotoluene	mg/Kg	0	73	nc																
		4-Nitrotoluene	mg/Kg	0	12	ca																
		HMX	mg/Kg	0	306	nc																
		Nitrobenzene	mg/Kg	0	2	nc																
		RDX	mg/Kg	0	4.4	ca																
		Tetryl	mg/Kg	0	61	nc																
Others	8015B MDRO	Diesel Range Organics (DRO)	mg/Kg	--	--	--	3	U	2.9	U	3	U	2.9	U								
	8015B MGRO	Gasoline Range Organics (GRO)	mg/Kg	--	--	--	0.012	U	0.011	U	0.011	U	0.011	U								
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc																
		Aroclor 1221	mg/Kg	--	0.22	ca																
		Aroclor 1232	mg/Kg	--	0.22	ca																
		Aroclor 1242	mg/Kg	--	0.22	ca																
		Aroclor 1248	mg/Kg	--	0.22	ca																
		Aroclor 1254	mg/Kg	--	0.22	ca																
		Aroclor 1260	mg/Kg	--	0.22	ca																
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca																
		4,4'-DDE	mg/Kg	--	1.7	ca																
		4,4'-DDT	mg/Kg	--	1.7	ca																
		Aldrin	mg/Kg	--	0.029	ca																
		alpha-BHC	mg/Kg	--	0.09	sat																
		alpha-Chlordane	mg/Kg	--	1.6	ca																
		beta-BHC	mg/Kg	--	0.32	ca																
		delta-BHC	mg/Kg	--	--	0																
		Dieldrin	mg/Kg	--	0.030	ca																
		Endosulfan I	mg/Kg	--	37	nc																
		Endosulfan II	mg/Kg	--	37	nc																
		Endosulfan sulfate	mg/Kg	--	37	nc																
		Endrin	mg/Kg	--	1.8	nc																
		Endrin aldehyde	mg/Kg	--	--	0																
		Endrin ketone	mg/Kg	--	--	0																
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca																
		gamma-Chlordane	mg/Kg	--	1.6	ca																
		Heptachlor	mg/Kg	--	0.11	ca																
		Heptachlor epoxide	mg/Kg	--	0.053	ca																
		Methoxychlor	mg/Kg	--	31	nc																
		Toxaphene	mg/Kg	--	0	0																



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-055-0001-SO		LL9SB-056-0001-SO		LL9SB-059-0001-SO		LL9SB-061-0001-SO		LL9SB-065-0001-SO		LL9SB-066-0001-SO		LL9SB-069-0001-SO		LL9SB-070-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0																
SVOCs TCL 8270 C	8330	Nitroglycerin	mg/Kg	0	--	0																
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc																
	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc	0.072	U	0.068	U	0.069	U	0.068	U								
		1,2-Dichlorobenzene	mg/Kg	--	600	sat	0.097	U	0.091	U	0.092	U	0.091	U								
		1,3-Dichlorobenzene	mg/Kg	--	53	nc	0.097	U	0.091	U	0.092	U	0.091	U								
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca	0.087	U	0.081	U	0.083	U	0.082	U								
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca	0.092	U	0.086	U	0.087	U	0.086	U								
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc	0.047	U	0.044	U	0.044	U	0.044	U								
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc	0.058	U	0.054	U	0.055	U	0.054	U								
		2,4-Dichlorophenol	mg/Kg	--	18	nc	0.059	U	0.055	U	0.056	U	0.055	U								
		2,4-Dimethylphenol	mg/Kg	--	122	nc	0.073	U	0.069	U	0.07	U	0.069	U								
		2,4-Dinitrophenol	mg/Kg	--	12	nc	0.14	U	0.13	U	0.13	U	0.13	U								
		2-Chloronaphthalene	mg/Kg	--	494	nc	0.059	U	0.055	U	0.056	U	0.055	U								
		2-Chlorophenol	mg/Kg	--	6.3	nc	0.072	U	0.068	U	0.069	U	0.068	U								
		2-Methylnaphthalene	mg/Kg	--	--	0	0.0018	U	0.0017	U	0.0017	U	0.0017	U								
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc	0.01	U	0.0096	U	0.0098	U	0.0096	U								
		2-Nitroaniline	mg/Kg	--	18.3	nc	0.042	U*	0.039	U*	0.04	U*	0.039	U*								
		2-Nitrophenol	mg/Kg	--	--	0	0.077	U	0.072	U	0.073	U	0.072	U								
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca	0.022	U	0.021	U	0.021	U	0.021	U								
		3-Nitroaniline	mg/Kg	--	1.8	nc	0.14	U	0.13	U	0.13	U	0.13	U								
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc	0.12	U*	0.11	U*	0.11	U*	0.11	U*								
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0	0.0038	U*	0.0036	U*	0.0036	U*	0.0036	U*								
		4-Chloro-3-methylphenol	mg/Kg	--	--	0	0.047	U	0.044	U	0.044	U	0.044	U								
		4-Chloroaniline	mg/Kg	--	24	nc	0.12	U	0.11	U	0.12	U	0.11	U								
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0	0.0044	U*	0.0041	U*	0.0042	U*	0.0041	U*								
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc	0.0071	U	0.0066	U	0.0068	U	0.0067	U								
		4-Nitroaniline	mg/Kg	--	23	ca	0.048	U	0.045	U	0.045	U	0.045	U								
		4-Nitrophenol	mg/Kg	--	--	0	0.1	U	0.094	U	0.096	U	0.094	U								
		Acenaphthene	mg/Kg	--	368	nc	0.0017	U	0.0016	U	0.006	J	0.0016	U								
		Acenaphthylene	mg/Kg	--	--	0	0.0011	U*	0.001	U*	0.0011	U*	0.001	U*								
		Anthracene	mg/Kg	--	2189	nc	0.0011	U	0.00099	U	0.0063	J	0.00099	U								
		Benzo(a)anthracene	mg/Kg	--	0.62	ca	0.0013	U	0.0066	J	0.029	J	0.0066	J								
		Benzo(a)pyrene	mg/Kg	--	0.062	ca	0.0027	U	0.0061	J	0.027	J	0.0071	J								
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca	0.0026	U	0.0072	J	0.032	JH	0.01	J								
		Benzo(ghi)perylene	mg/Kg	--	--	0	0.0023	U	0.0054	J	0.012	J	0.0022	U								
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca	0.0034	U	0.0078	JM	0.03	J	0.0065	JM								
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0	0.0036	U	0.0033	U	0.0034	U	0.0033	U								
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca	0.0024	U	0.0023	U	0.0023	U	0.0023	U								
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca	0.012	U	0.011	U	0.011	U	0.011	U								



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-055-0001-SO		LL9SB-056-0001-SO		LL9SB-059-0001-SO		LL9SB-061-0001-SO		LL9SB-065-0001-SO		LL9SB-066-0001-SO		LL9SB-069-0001-SO		LL9SB-070-0001-SO	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Butyl benzyl phthalate	mg/Kg	--	1222	nc	0.005	U	0.0047	U	0.0048	U	0.0047	U								
		Carbazole	mg/Kg	--	24	ca	0.043	U	0.04	U	0.041	U	0.04	U								
		Chrysene	mg/Kg	--	62	ca	0.0022	U	0.007	J	0.026	J	0.0071	J								
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca	0.0027	U	0.0025	U	0.0064	J	0.0025	U								
		Dibenzofuran	mg/Kg	--	15	nc	0.0033	U	0.0031	U	0.0031	U	0.0031	U								
		Diethyl phthalate	mg/Kg	--	4888	nc	0.0045	U*	0.0042	U*	0.0043	U*	0.0042	U*								
		Dimethyl phthalate	mg/Kg	--	100000	max	0.0044	U*	0.0041	U*	0.0042	U*	0.0041	U*								
		Di-n-butyl phthalate	mg/Kg	--	611	nc	0.024	U	0.023	U	0.023	U	0.023	U								
		Di-n-octyl phthalate	mg/Kg	--	244	nc	0.011	U	0.01	U	0.01	U	0.01	U								
		Fluoranthene	mg/Kg	--	229	nc	0.0013	U	0.015	J	0.071		0.013	J								
		Fluorene	mg/Kg	--	275	nc	0.002	U	0.0018	U	0.0019	U	0.0018	U								
		Hexachlorobenzene	mg/Kg	--	0.30	ca	0.0022	U*	0.0021	U*	0.0021	U*	0.0021	U*								
		Hexachlorobutadiene	mg/Kg	--	6.2	ca	0.004	U	0.0038	U	0.0038	U	0.0038	U								
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc	0.066	U	0.062	U	0.063	U	0.062	U								
		Hexachloroethane	mg/Kg	--	35	ca	0.004	U	0.0038	U	0.0038	U	0.0038	U								
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca	0.0026	U	0.0039	J	0.014	J	0.0041	J								
		Isophorone	mg/Kg	--	512	ca	0.0029	U	0.0028	U	0.0028	U	0.0028	U								
		Naphthalene	mg/Kg	--	5.6	nc	0.0021	U	0.0019	U	0.002	U	0.002	U								
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca	0.0028	U	0.0026	U	0.0027	U	0.0026	U								
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca	0.0036	U	0.0033	U	0.0034	U	0.0033	U								
		Pentachlorophenol	mg/Kg	--	3.0	ca	0.12	U	0.11	U	0.12	U	0.11	U								
		Phenanthrene	mg/Kg	--	--	0	0.0012	U	0.0065	J	0.036	J	0.0049	J								
		Phenol	mg/Kg	--	1833	nc	0.002	U	0.0018	U	0.0019	U	0.0018	U								
		Pyrene	mg/Kg	--	232	nc	0.0024	U	0.0096	J	0.044		0.0079	J								
TAL Metals 6010B	6010B	Aluminum	mg/Kg	19500	7614	nc									10000		5300		2900		7800	
		Barium	mg/Kg	124	538	nc									83		38		13		45	
		Beryllium	mg/Kg	0.88	15	nc									0.43	B	0.44		0.16	B	0.27	B
		Cadmium	mg/Kg	0.00	3.7	nc									0.11	U	0.086	U	0.086	U	0.087	U
		Calcium	mg/Kg	35500	--	0									1100		1200		370	H	1600	H
		Chromium	mg/Kg	27.2	30	ca									12		9.8		4.7		11	
		Cobalt	mg/Kg	23.2	902	ca									5.8		10		2.2		6.6	
		Copper	mg/Kg	32.3	313	nc									8.7		18		5.1		18	
		Iron	mg/Kg	35200	2346	nc									14000		19000		9800	*H	21000	*H
		Magnesium	mg/Kg	8790	--	0									2100		1700		540		2000	
		Manganese	mg/Kg	3030	176	nc									140		340		42		490	
		Nickel	mg/Kg	60.7	156	nc									15		22		4.3		16	
		Potassium	mg/Kg	3350	--	0									790		1000		370		740	
		Selenium	mg/Kg	1.5	39	nc									0.58	B	0.43	U	0.43	U	0.43	U
		Silver	mg/Kg	0.00	39	nc									0.42	U	0.33	U	0.33	U	0.34	U
		Sodium	mg/Kg	145	--	0									1300		1100		93	U	94	U
		Vanadium	mg/Kg	37.6	7.8	nc									17		9.8		6.1		14	
		Zinc	mg/Kg	93.3	2346	nc									72		54		49	H	69	H



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

							LL9SB-055-0001-SO		LL9SB-056-0001-SO		LL9SB-059-0001-SO		LL9SB-061-0001-SO		LL9SB-065-0001-SO		LL9SB-066-0001-SO		LL9SB-069-0001-SO		LL9SB-070-0001-SO	
Sample Date							11/05/03		11/05/03		11/05/03		11/05/03		11/06/03		11/06/03		10/27/03		10/27/03	
Sample Depth							1.5 - 3.5 ft		2 - 4 ft		3 - 5 ft		1 - 3 ft		3 - 5 ft		5 - 7 ft		7 ft		6.5 ft	
Group	Method	Analyte	Units	RVAAP Subsurface Soil Background Criteria	Region 9 (Residential Soil) PRGs	SB PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
	7041	Antimony	mg/Kg	0.96	3.1	nc									0.72	B	0.45	U	0.43	U	0.44	U
	7060A	Arsenic	mg/Kg	19.8	0.39	ca									4.6		11		6.9		8.3	
	7421	Lead	mg/Kg	19.1	400	pbk									51		10		9.2		11	
	7471A	Mercury	mg/Kg	0.04	2.3	nc									9.7		0.016	B	0.025		0.013	B
	7841	Thallium	mg/Kg	0.91	0.52	nc									1.1	U	0.97	U	0.19	U	0.19	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat	0.00075	U	0.00071	U	0.019	U	0.019	U								
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca	0.00079	U	0.00074	U	0.022	U	0.022	U								
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca	0.00087	U	0.00082	U	0.037	U	0.037	U								
		1,1-Dichloroethane	mg/Kg	--	51	nc	0.0011	U	0.001	U	0.016	U	0.016	U								
		1,1-Dichloroethene	mg/Kg	--	12	nc	0.0012	U	0.0012	U	0.017	U	0.016	U								
		1,2-Dichloroethane	mg/Kg	--	0.28	ca	0.00071	U	0.00067	U	0.025	U	0.025	U								
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc	0.0023	U	0.0022	U	0.034	U	0.034	U								
		1,2-Dichloropropane	mg/Kg	--	0.34	ca	0.0012	U	0.0011	U	0.021	U	0.021	U								
		2-Butanone (MEK)	mg/Kg	--	2231	nc	0.0052	U	0.0049	U	0.06	U	0.06	U								
		2-Hexanone	mg/Kg	--	530	nc	0.0021	U	0.002	U	0.061	U	0.061	U								
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc	0.0037	U	0.0035	U	0.044	U	0.044	U								
		Acetone	mg/Kg	--	1412	nc	0.005	U	0.0048	U	0.034	U	0.034	U								
		Benzene	mg/Kg	--	0.64	ca	0.00081	U	0.00077	U	0.017	U	0.016	U								
		Bromodichloromethane	mg/Kg	--	0.82	ca	0.00083	U	0.00079	U	0.022	U	0.022	U								
		Bromoform	mg/Kg	--	62	ca	0.0011	U	0.0011	U	0.021	U	0.021	U								
		Bromomethane	mg/Kg	--	0.39	nc	0.0036	U	0.0034	U	0.012	U	0.012	U								
		Carbon disulfide	mg/Kg	--	36	nc	0.0025	U	0.0023	U	0.024	U	0.024	U								
		Carbon tetrachloride	mg/Kg	--	0.25	ca	0.001	U	0.00096	U	0.019	U	0.019	U								
		Chlorobenzene	mg/Kg	--	15	nc	0.0011	U	0.0011	U	0.026	U	0.026	U								
		Chloroethane	mg/Kg	--	3.0	ca	0.002	U	0.0019	U	0.024	U	0.024	U								
		Chloroform	mg/Kg	--	0.22	ca	0.00076	U	0.00072	U	0.021	U	0.021	U								
		Chloromethane	mg/Kg	--	4.7	nc	0.0012	U	0.0011	U	0.028	U	0.028	U								
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.00097	U	0.00092	U	0.027	U	0.026	U								
		Dibromochloromethane	mg/Kg	--	1.1	ca	0.00085	U	0.0008	U	0.022	U	0.022	U								
		Ethylbenzene	mg/Kg	--	395	sat	0.0013	U	0.0013	U	0.027	U	0.026	U								
		Methylene chloride	mg/Kg	--	9.1	ca	0.0022	U	0.0021	U	0.024	U	0.024	U								
		Styrene	mg/Kg	--	1700	sat	0.0012	U	0.0012	U	0.034	U	0.033	U								
		Tetrachloroethene	mg/Kg	--	0.48	ca	0.00082	U	0.00078	U	0.027	U	0.027	U								
		Toluene	mg/Kg	--	520	sat	0.0012	U	0.0012	U	0.021	U	0.021	U								
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca	0.001	U	0.00097	U	0.023	U	0.023	U								
		Trichloroethene	mg/Kg	--	0.053	ca	0.00072	U	0.00068	U	0.025	U	0.025	U								
		Vinyl chloride	mg/Kg	--	0.079	ca	0.00091	U	0.00086	U	0.021	U	0.021	U								
		Xylenes (total)	mg/Kg	--	0.079	ca	0.0036	U	0.0034	U	0.059	U	0.059	U								



Table 4-4
Summary of Subsurface Soil Results (>1 ft bgs)

Notes:
-- - no background/PRG value is available for this analyte
blank cell indicates that the analysis was not performed
PRG - preliminary remediation goals
(The PRG for lead is the Maximum Contaminant Level (MCL))
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background and PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG and Background, then the value is presented with a normal style

Inorganic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL
InOrganic Flags
^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL
Organic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL
Organic Flags
B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-001-0001-SO	12/04/03	1-3 ft	Arsenic	12		19.8	0.39	ca	NO	YES
LL9SB-001-0001-SO	12/04/03	1-3 ft	Iron	23000		35200	2346	nc	NO	YES
LL9SB-001-0001-SO	12/04/03	1-3 ft	Manganese	400		3030	176	nc	NO	YES
LL9SB-001-0001-SO	12/04/03	1-3 ft	Vanadium	13		37.6	7.8	nc	NO	YES
LL9SB-002-0001-SO	12/04/03	1-3 ft	Aluminum	10000		19500	7614	nc	NO	YES
LL9SB-002-0001-SO	12/04/03	1-3 ft	Arsenic	12		19.8	0.39	ca	NO	YES
LL9SB-002-0001-SO	12/04/03	1-3 ft	Iron	23000		35200	2346	nc	NO	YES
LL9SB-002-0001-SO	12/04/03	1-3 ft	Lead	24		19.1	400	pbk	YES	NO
LL9SB-002-0001-SO	12/04/03	1-3 ft	Manganese	600		3030	176	nc	NO	YES
LL9SB-002-0001-SO	12/04/03	1-3 ft	Mercury	0.076		0.04	2.3	nc	YES	NO
LL9SB-002-0001-SO	12/04/03	1-3 ft	Vanadium	19		37.6	7.8	nc	NO	YES
LL9SB-003-0001-SO	12/04/03	1-2 ft	Arsenic	9.8		19.8	0.39	ca	NO	YES
LL9SB-003-0001-SO	12/04/03	1-2 ft	Iron	16000		35200	2346	nc	NO	YES
LL9SB-003-0001-SO	12/04/03	1-2 ft	Manganese	310		3030	176	nc	NO	YES
LL9SB-003-0001-SO	12/04/03	1-2 ft	Vanadium	11		37.6	7.8	nc	NO	YES
LL9SB-004-0001-SO	12/04/03	1-3 ft	Aluminum	8200		19500	7614	nc	NO	YES
LL9SB-004-0001-SO	12/04/03	1-3 ft	Arsenic	8.5		19.8	0.39	ca	NO	YES
LL9SB-004-0001-SO	12/04/03	1-3 ft	Iron	18000		35200	2346	nc	NO	YES
LL9SB-004-0001-SO	12/04/03	1-3 ft	Manganese	200		3030	176	nc	NO	YES
LL9SB-004-0001-SO	12/04/03	1-3 ft	Vanadium	15		37.6	7.8	nc	NO	YES
LL9SB-005-0001-DUP	12/04/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-005-0001-DUP	12/04/03	1-3 ft	Arsenic	9		19.8	0.39	ca	NO	YES
LL9SB-005-0001-DUP	12/04/03	1-3 ft	Iron	17000		35200	2346	nc	NO	YES
LL9SB-005-0001-DUP	12/04/03	1-3 ft	Manganese	1300		3030	176	nc	NO	YES
LL9SB-005-0001-DUP	12/04/03	1-3 ft	Nitrocellulose	2.7	J	0	--	0	YES	NO
LL9SB-005-0001-DUP	12/04/03	1-3 ft	Vanadium	18		37.6	7.8	nc	NO	YES
LL9SB-005-0001-SO	12/04/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-005-0001-SO	12/04/03	1-3 ft	Arsenic	7.7		19.8	0.39	ca	NO	YES
LL9SB-005-0001-SO	12/04/03	1-3 ft	Iron	15000		35200	2346	nc	NO	YES
LL9SB-005-0001-SO	12/04/03	1-3 ft	Manganese	1100		3030	176	nc	NO	YES
LL9SB-005-0001-SO	12/04/03	1-3 ft	Nitrocellulose	1.2	B J	0	--	0	YES	NO
LL9SB-005-0001-SO	12/04/03	1-3 ft	Nitroglycerin	4.82		0	--	0	YES	NO



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-005-0001-SO	12/04/03	1-3 ft	Nitroglycerin	4.9		0	--	0	YES	NO
LL9SB-005-0001-SO	12/04/03	1-3 ft	Nitroguanidine	0.945		0	611	nc	YES	NO
LL9SB-005-0001-SO	12/04/03	1-3 ft	Nitroguanidine	0.912		0	611	nc	YES	NO
LL9SB-005-0001-SO	12/04/03	1-3 ft	Vanadium	17		37.6	7.8	nc	NO	YES
LL9SB-006-0001-SO	12/11/03	1-2 ft	Arsenic	9.9		19.8	0.39	ca	NO	YES
LL9SB-006-0001-SO	12/11/03	1-2 ft	Cadmium	2		0	3.7	nc	YES	NO
LL9SB-006-0001-SO	12/11/03	1-2 ft	Iron	14000	H	35200	2346	nc	NO	YES
LL9SB-006-0001-SO	12/11/03	1-2 ft	Lead	54		19.1	400	pbk	YES	NO
LL9SB-006-0001-SO	12/11/03	1-2 ft	Manganese	520		3030	176	nc	NO	YES
LL9SB-006-0001-SO	12/11/03	1-2 ft	Mercury	0.94		0.04	2.3	nc	YES	NO
LL9SB-006-0001-SO	12/11/03	1-2 ft	Vanadium	13		37.6	7.8	nc	NO	YES
LL9SB-006-0001-SO	12/11/03	1-2 ft	Zinc	120		93.3	2346	nc	YES	NO
LL9SB-007-0001-SO	12/08/03	1-3 ft	Aluminum	12000		19500	7614	nc	NO	YES
LL9SB-007-0001-SO	12/08/03	1-3 ft	Arsenic	14		19.8	0.39	ca	NO	YES
LL9SB-007-0001-SO	12/08/03	1-3 ft	Cadmium	0.13	B	0	3.7	nc	YES	NO
LL9SB-007-0001-SO	12/08/03	1-3 ft	Iron	23000		35200	2346	nc	NO	YES
LL9SB-007-0001-SO	12/08/03	1-3 ft	Manganese	570		3030	176	nc	NO	YES
LL9SB-007-0001-SO	12/08/03	1-3 ft	Vanadium	21		37.6	7.8	nc	NO	YES
LL9SB-008-0001-SO	12/10/03	1-3 ft	Aluminum	12000		19500	7614	nc	NO	YES
LL9SB-008-0001-SO	12/10/03	1-3 ft	Arsenic	12		19.8	0.39	ca	NO	YES
LL9SB-008-0001-SO	12/10/03	1-3 ft	Iron	30000		35200	2346	nc	NO	YES
LL9SB-008-0001-SO	12/10/03	1-3 ft	Manganese	550		3030	176	nc	NO	YES
LL9SB-008-0001-SO	12/10/03	1-3 ft	Vanadium	22		37.6	7.8	nc	NO	YES
LL9SB-009-0001-SO	12/08/03	1-3 ft	Aluminum	12000		19500	7614	nc	NO	YES
LL9SB-009-0001-SO	12/08/03	1-3 ft	Arsenic	22		19.8	0.39	ca	YES	YES
LL9SB-009-0001-SO	12/08/03	1-3 ft	Iron	28000		35200	2346	nc	NO	YES
LL9SB-009-0001-SO	12/08/03	1-3 ft	Manganese	280		3030	176	nc	NO	YES
LL9SB-009-0001-SO	12/08/03	1-3 ft	Vanadium	18		37.6	7.8	nc	NO	YES
LL9SB-010-0001-SO	12/08/03	2.5 ft	Aluminum	9700		19500	7614	nc	NO	YES
LL9SB-010-0001-SO	12/08/03	2.5 ft	Arsenic	22		19.8	0.39	ca	YES	YES
LL9SB-010-0001-SO	12/08/03	2.5 ft	Cadmium	0.13	B	0	3.7	nc	YES	NO
LL9SB-010-0001-SO	12/08/03	2.5 ft	Iron	24000		35200	2346	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-010-0001-SO	12/08/03	2.5 ft	Manganese	470		3030	176	nc	NO	YES
LL9SB-010-0001-SO	12/08/03	2.5 ft	Mercury	0.18		0.04	2.3	nc	YES	NO
LL9SB-010-0001-SO	12/08/03	2.5 ft	Vanadium	14		37.6	7.8	nc	NO	YES
LL9SB-012-0001-SO	12/10/03	1-3 ft	Aluminum	9700		19500	7614	nc	NO	YES
LL9SB-012-0001-SO	12/10/03	1-3 ft	Arsenic	19		19.8	0.39	ca	NO	YES
LL9SB-012-0001-SO	12/10/03	1-3 ft	Iron	25000		35200	2346	nc	NO	YES
LL9SB-012-0001-SO	12/10/03	1-3 ft	Lead	20		19.1	400	pbk	YES	NO
LL9SB-012-0001-SO	12/10/03	1-3 ft	Manganese	470		3030	176	nc	NO	YES
LL9SB-012-0001-SO	12/10/03	1-3 ft	Vanadium	17		37.6	7.8	nc	NO	YES
LL9SB-012-0001-SO	12/10/03	1-3 ft	Zinc	110		93.3	2346	nc	YES	NO
LL9SB-013-0001-SO	12/10/03	1-3 ft	Arsenic	11		19.8	0.39	ca	NO	YES
LL9SB-013-0001-SO	12/10/03	1-3 ft	Iron	18000		35200	2346	nc	NO	YES
LL9SB-013-0001-SO	12/10/03	1-3 ft	Manganese	420		3030	176	nc	NO	YES
LL9SB-013-0001-SO	12/10/03	1-3 ft	Vanadium	11		37.6	7.8	nc	NO	YES
LL9SB-014-0001-SO	12/11/03	1-3 ft	Arsenic	18		19.8	0.39	ca	NO	YES
LL9SB-014-0001-SO	12/11/03	1-3 ft	Iron	18000	H	35200	2346	nc	NO	YES
LL9SB-014-0001-SO	12/11/03	1-3 ft	Manganese	430		3030	176	nc	NO	YES
LL9SB-014-0001-SO	12/11/03	1-3 ft	Vanadium	12		37.6	7.8	nc	NO	YES
LL9SB-015-0001-SO	12/11/03	1-3 ft	Aluminum	10000		19500	7614	nc	NO	YES
LL9SB-015-0001-SO	12/11/03	1-3 ft	Antimony	1		0.96	3.1	nc	YES	NO
LL9SB-015-0001-SO	12/11/03	1-3 ft	Arsenic	18		19.8	0.39	ca	NO	YES
LL9SB-015-0001-SO	12/11/03	1-3 ft	Iron	27000	H	35200	2346	nc	NO	YES
LL9SB-015-0001-SO	12/11/03	1-3 ft	Manganese	280		3030	176	nc	NO	YES
LL9SB-015-0001-SO	12/11/03	1-3 ft	Vanadium	17		37.6	7.8	nc	NO	YES
LL9SB-016-0001-SO	12/11/03	1-3 ft	Aluminum	8000		19500	7614	nc	NO	YES
LL9SB-016-0001-SO	12/11/03	1-3 ft	Arsenic	11		19.8	0.39	ca	NO	YES
LL9SB-016-0001-SO	12/11/03	1-3 ft	Iron	24000	H	35200	2346	nc	NO	YES
LL9SB-016-0001-SO	12/11/03	1-3 ft	Manganese	550		3030	176	nc	NO	YES
LL9SB-016-0001-SO	12/11/03	1-3 ft	Vanadium	15		37.6	7.8	nc	NO	YES
LL9SB-017-0001-SO	12/04/03	1-3 ft	Aluminum	10000		19500	7614	nc	NO	YES
LL9SB-017-0001-SO	12/04/03	1-3 ft	Arsenic	10		19.8	0.39	ca	NO	YES
LL9SB-017-0001-SO	12/04/03	1-3 ft	Iron	19000		35200	2346	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-017-0001-SO	12/04/03	1-3 ft	Manganese	760		3030	176	nc	NO	YES
LL9SB-017-0001-SO	12/04/03	1-3 ft	Vanadium	20		37.6	7.8	nc	NO	YES
LL9SB-018-0001-SO	12/10/03	1-3 ft	Aluminum	14000		19500	7614	nc	NO	YES
LL9SB-018-0001-SO	12/10/03	1-3 ft	Arsenic	15		19.8	0.39	ca	NO	YES
LL9SB-018-0001-SO	12/10/03	1-3 ft	Iron	26000		35200	2346	nc	NO	YES
LL9SB-018-0001-SO	12/10/03	1-3 ft	Manganese	250		3030	176	nc	NO	YES
LL9SB-018-0001-SO	12/10/03	1-3 ft	Vanadium	19		37.6	7.8	nc	NO	YES
LL9SB-019-0001-SO	12/11/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-019-0001-SO	12/11/03	1-3 ft	Arsenic	32		19.8	0.39	ca	YES	YES
LL9SB-019-0001-SO	12/11/03	1-3 ft	Iron	30000	H	35200	2346	nc	NO	YES
LL9SB-019-0001-SO	12/11/03	1-3 ft	Manganese	270		3030	176	nc	NO	YES
LL9SB-019-0001-SO	12/11/03	1-3 ft	Mercury	0.18		0.04	2.3	nc	YES	NO
LL9SB-019-0001-SO	12/11/03	1-3 ft	Vanadium	17		37.6	7.8	nc	NO	YES
LL9SB-020-0001-SO	12/08/03	1-3 ft	Aluminum	15000		19500	7614	nc	NO	YES
LL9SB-020-0001-SO	12/08/03	1-3 ft	Arsenic	14		19.8	0.39	ca	NO	YES
LL9SB-020-0001-SO	12/08/03	1-3 ft	Cadmium	0.095	B	0	3.7	nc	YES	NO
LL9SB-020-0001-SO	12/08/03	1-3 ft	Iron	24000		35200	2346	nc	NO	YES
LL9SB-020-0001-SO	12/08/03	1-3 ft	Manganese	420		3030	176	nc	NO	YES
LL9SB-020-0001-SO	12/08/03	1-3 ft	Mercury	0.18		0.04	2.3	nc	YES	NO
LL9SB-020-0001-SO	12/08/03	1-3 ft	Vanadium	26		37.6	7.8	nc	NO	YES
LL9SB-021-0001-SO	12/08/03	2.5 ft	Aluminum	12000		19500	7614	nc	NO	YES
LL9SB-021-0001-SO	12/08/03	2.5 ft	Arsenic	26		19.8	0.39	ca	YES	YES
LL9SB-021-0001-SO	12/08/03	2.5 ft	Iron	28000		35200	2346	nc	NO	YES
LL9SB-021-0001-SO	12/08/03	2.5 ft	Lead	22		19.1	400	pbk	YES	NO
LL9SB-021-0001-SO	12/08/03	2.5 ft	Manganese	290		3030	176	nc	NO	YES
LL9SB-021-0001-SO	12/08/03	2.5 ft	Vanadium	19		37.6	7.8	nc	NO	YES
LL9SB-022-0001-SO	12/10/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-022-0001-SO	12/10/03	1-3 ft	Arsenic	18		19.8	0.39	ca	NO	YES
LL9SB-022-0001-SO	12/10/03	1-3 ft	Iron	24000		35200	2346	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-022-0001-SO	12/10/03	1-3 ft	Lead	23		19.1	400	pbk	YES	NO
LL9SB-022-0001-SO	12/10/03	1-3 ft	Manganese	500		3030	176	nc	NO	YES
LL9SB-022-0001-SO	12/10/03	1-3 ft	Mercury	0.087		0.04	2.3	nc	YES	NO
LL9SB-022-0001-SO	12/10/03	1-3 ft	Vanadium	15		37.6	7.8	nc	NO	YES
LL9SB-023-0001-SO	12/08/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-023-0001-SO	12/08/03	1-3 ft	Arsenic	24		19.8	0.39	ca	YES	YES
LL9SB-023-0001-SO	12/08/03	1-3 ft	Iron	27000		35200	2346	nc	NO	YES
LL9SB-023-0001-SO	12/08/03	1-3 ft	Manganese	350		3030	176	nc	NO	YES
LL9SB-023-0001-SO	12/08/03	1-3 ft	Vanadium	18		37.6	7.8	nc	NO	YES
LL9SB-024-0001-SO	12/10/03	1-3 ft	Aluminum	18000		19500	7614	nc	NO	YES
LL9SB-024-0001-SO	12/10/03	1-3 ft	Antimony	1		0.96	3.1	nc	YES	NO
LL9SB-024-0001-SO	12/10/03	1-3 ft	Arsenic	9.2		19.8	0.39	ca	NO	YES
LL9SB-024-0001-SO	12/10/03	1-3 ft	Chromium	54		27.2	30	ca	YES	YES
LL9SB-024-0001-SO	12/10/03	1-3 ft	Iron	28000		35200	2346	nc	NO	YES
LL9SB-024-0001-SO	12/10/03	1-3 ft	Lead	86		19.1	400	pbk	YES	NO
LL9SB-024-0001-SO	12/10/03	1-3 ft	Manganese	440		3030	176	nc	NO	YES
LL9SB-024-0001-SO	12/10/03	1-3 ft	Mercury	0.079		0.04	2.3	nc	YES	NO
LL9SB-024-0001-SO	12/10/03	1-3 ft	Vanadium	29		37.6	7.8	nc	NO	YES
LL9SB-025-0001-SO	12/10/03	1-3 ft	Aluminum	18000		19500	7614	nc	NO	YES
LL9SB-025-0001-SO	12/10/03	1-3 ft	Arsenic	15		19.8	0.39	ca	NO	YES
LL9SB-025-0001-SO	12/10/03	1-3 ft	Iron	32000		35200	2346	nc	NO	YES
LL9SB-025-0001-SO	12/10/03	1-3 ft	Manganese	180		3030	176	nc	NO	YES
LL9SB-025-0001-SO	12/10/03	1-3 ft	Vanadium	29		37.6	7.8	nc	NO	YES
LL9SB-026-0001-SO	12/10/03	1-3 ft	Aluminum	13000		19500	7614	nc	NO	YES
LL9SB-026-0001-SO	12/10/03	1-3 ft	Arsenic	6.9		19.8	0.39	ca	NO	YES
LL9SB-026-0001-SO	12/10/03	1-3 ft	Iron	19000		35200	2346	nc	NO	YES
LL9SB-026-0001-SO	12/10/03	1-3 ft	Manganese	470		3030	176	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-026-0001-SO	12/10/03	1-3 ft	Vanadium	23		37.6	7.8	nc	NO	YES
LL9SB-027-0001-SO	12/04/03	1-3 ft	Aluminum	10000		19500	7614	nc	NO	YES
LL9SB-027-0001-SO	12/04/03	1-3 ft	Arsenic	18		19.8	0.39	ca	NO	YES
LL9SB-027-0001-SO	12/04/03	1-3 ft	Iron	24000		35200	2346	nc	NO	YES
LL9SB-027-0001-SO	12/04/03	1-3 ft	Manganese	410		3030	176	nc	NO	YES
LL9SB-027-0001-SO	12/04/03	1-3 ft	Vanadium	19		37.6	7.8	nc	NO	YES
LL9SB-028-0001-SO	12/04/03	1-3 ft	Aluminum	9600		19500	7614	nc	NO	YES
LL9SB-028-0001-SO	12/04/03	1-3 ft	Arsenic	5.1		19.8	0.39	ca	NO	YES
LL9SB-028-0001-SO	12/04/03	1-3 ft	Iron	11000		35200	2346	nc	NO	YES
LL9SB-028-0001-SO	12/04/03	1-3 ft	Vanadium	18		37.6	7.8	nc	NO	YES
LL9SB-029-0001-SO	12/04/03	1-3 ft	Aluminum	9400		19500	7614	nc	NO	YES
LL9SB-029-0001-SO	12/04/03	1-3 ft	Arsenic	11		19.8	0.39	ca	NO	YES
LL9SB-029-0001-SO	12/04/03	1-3 ft	Iron	24000		35200	2346	nc	NO	YES
LL9SB-029-0001-SO	12/04/03	1-3 ft	Manganese	660		3030	176	nc	NO	YES
LL9SB-029-0001-SO	12/04/03	1-3 ft	Vanadium	19		37.6	7.8	nc	NO	YES
LL9SB-030-0001-SO	12/04/03	1-3 ft	Aluminum	10000		19500	7614	nc	NO	YES
LL9SB-030-0001-SO	12/04/03	1-3 ft	Arsenic	6		19.8	0.39	ca	NO	YES
LL9SB-030-0001-SO	12/04/03	1-3 ft	Iron	21000		35200	2346	nc	NO	YES
LL9SB-030-0001-SO	12/04/03	1-3 ft	Manganese	380		3030	176	nc	NO	YES
LL9SB-030-0001-SO	12/04/03	1-3 ft	Vanadium	20		37.6	7.8	nc	NO	YES
LL9SB-031-0001-SO	12/04/03	1-3 ft	Arsenic	7.4		19.8	0.39	ca	NO	YES
LL9SB-031-0001-SO	12/04/03	1-3 ft	Iron	16000		35200	2346	nc	NO	YES
LL9SB-031-0001-SO	12/04/03	1-3 ft	Manganese	390		3030	176	nc	NO	YES
LL9SB-031-0001-SO	12/04/03	1-3 ft	Vanadium	12		37.6	7.8	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-032-0001-SO	12/04/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-032-0001-SO	12/04/03	1-3 ft	Arsenic	7.4		19.8	0.39	ca	NO	YES
LL9SB-032-0001-SO	12/04/03	1-3 ft	Iron	19000		35200	2346	nc	NO	YES
LL9SB-032-0001-SO	12/04/03	1-3 ft	Manganese	970		3030	176	nc	NO	YES
LL9SB-032-0001-SO	12/04/03	1-3 ft	Nitrocellulose	1.3	B J	0	--	0	YES	NO
LL9SB-032-0001-SO	12/04/03	1-3 ft	Vanadium	21		37.6	7.8	nc	NO	YES
LL9SB-040-0001-SO	10/27/03	6.5 ft	Aluminum	14000		19500	7614	nc	NO	YES
LL9SB-040-0001-SO	10/27/03	6.5 ft	Arsenic	12		19.8	0.39	ca	NO	YES
LL9SB-040-0001-SO	10/27/03	6.5 ft	Barium	140		124	538	nc	YES	NO
LL9SB-040-0001-SO	10/27/03	6.5 ft	Beryllium	1.1		0.88	15	nc	YES	NO
LL9SB-040-0001-SO	10/27/03	6.5 ft	Calcium	36000	H	35500	--[n]	0	YES	NO
LL9SB-040-0001-SO	10/27/03	6.5 ft	Iron	39000	*H	35200	2346	nc	YES	YES
LL9SB-040-0001-SO	10/27/03	6.5 ft	Lead	51		19.1	400	pbk	YES	NO
LL9SB-040-0001-SO	10/27/03	6.5 ft	Manganese	2300		3030	176	nc	NO	YES
LL9SB-040-0001-SO	10/27/03	6.5 ft	Mercury	0.26		0.04	2.3	nc	YES	NO
LL9SB-040-0001-SO	10/27/03	6.5 ft	Sodium	170		145	--[n]	0	YES	NO
LL9SB-040-0001-SO	10/27/03	6.5 ft	Vanadium	17		37.6	7.8	nc	NO	YES
LL9SB-041-0001-SO	10/27/03	7.0 ft	Aluminum	8900		19500	7614	nc	NO	YES
LL9SB-041-0001-SO	10/27/03	7.0 ft	Arsenic	3.4		19.8	0.39	ca	NO	YES
LL9SB-041-0001-SO	10/27/03	7.0 ft	Beryllium	1.2		0.88	15	nc	YES	NO
LL9SB-041-0001-SO	10/27/03	7.0 ft	Iron	51000	*H	35200	2346	nc	YES	YES
LL9SB-041-0001-SO	10/27/03	7.0 ft	Lead	31		19.1	400	pbk	YES	NO
LL9SB-041-0001-SO	10/27/03	7.0 ft	Manganese	2100		3030	176	nc	NO	YES
LL9SB-041-0001-SO	10/27/03	7.0 ft	Vanadium	16		37.6	7.8	nc	NO	YES
LL9SB-042-0001-SO	11/06/03	8-10 ft	Aluminum	9100		19500	7614	nc	NO	YES
LL9SB-042-0001-SO	11/06/03	8-10 ft	Arsenic	6.9		19.8	0.39	ca	NO	YES
LL9SB-042-0001-SO	11/06/03	8-10 ft	Iron	23000		35200	2346	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-042-0001-SO	11/06/03	8-10 ft	Manganese	420		3030	176	nc	NO	YES
LL9SB-042-0001-SO	11/06/03	8-10 ft	Sodium	1200		145	--[n]	0	YES	NO
LL9SB-042-0001-SO	11/06/03	8-10 ft	Vanadium	15		37.6	7.8	nc	NO	YES
LL9SB-043-0001-DUP	11/06/03	9-11 ft	Arsenic	3.4		19.8	0.39	ca	NO	YES
LL9SB-043-0001-DUP	11/06/03	9-11 ft	Iron	26000		35200	2346	nc	NO	YES
LL9SB-043-0001-DUP	11/06/03	9-11 ft	Manganese	800		3030	176	nc	NO	YES
LL9SB-043-0001-DUP	11/06/03	9-11 ft	Sodium	840		145	--[n]	0	YES	NO
LL9SB-043-0001-DUP	11/06/03	9-11 ft	Vanadium	9.6		37.6	7.8	nc	NO	YES
LL9SB-043-0001-SO	11/06/03	9-11 ft	Arsenic	3.7		19.8	0.39	ca	NO	YES
LL9SB-043-0001-SO	11/06/03	9-11 ft	Iron	27000		35200	2346	nc	NO	YES
LL9SB-043-0001-SO	11/06/03	9-11 ft	Manganese	730		3030	176	nc	NO	YES
LL9SB-043-0001-SO	11/06/03	9-11 ft	Sodium	860		145	--[n]	0	YES	NO
LL9SB-043-0001-SO	11/06/03	9-11 ft	Vanadium	10		37.6	7.8	nc	NO	YES
LL9SB-044-0001-SO	11/06/03	8-10 ft	Aluminum	8000		19500	7614	nc	NO	YES
LL9SB-044-0001-SO	11/06/03	8-10 ft	Arsenic	15		19.8	0.39	ca	NO	YES
LL9SB-044-0001-SO	11/06/03	8-10 ft	Iron	24000		35200	2346	nc	NO	YES
LL9SB-044-0001-SO	11/06/03	8-10 ft	Manganese	420		3030	176	nc	NO	YES
LL9SB-044-0001-SO	11/06/03	8-10 ft	Sodium	1100		145	--[n]	0	YES	NO
LL9SB-044-0001-SO	11/06/03	8-10 ft	Vanadium	13		37.6	7.8	nc	NO	YES
LL9SB-045-0001-SO	11/06/03	8-10 ft	Aluminum	9300		19500	7614	nc	NO	YES
LL9SB-045-0001-SO	11/06/03	8-10 ft	Arsenic	18		19.8	0.39	ca	NO	YES
LL9SB-045-0001-SO	11/06/03	8-10 ft	Iron	27000		35200	2346	nc	NO	YES
LL9SB-045-0001-SO	11/06/03	8-10 ft	Manganese	330		3030	176	nc	NO	YES
LL9SB-045-0001-SO	11/06/03	8-10 ft	Sodium	1200		145	--[n]	0	YES	NO
LL9SB-045-0001-SO	11/06/03	8-10 ft	Vanadium	14		37.6	7.8	nc	NO	YES
LL9SB-046-0001-SO	11/06/03	4-6 ft	Arsenic	17		19.8	0.39	ca	NO	YES
LL9SB-046-0001-SO	11/06/03	4-6 ft	Iron	21000		35200	2346	nc	NO	YES
LL9SB-046-0001-SO	11/06/03	4-6 ft	Manganese	360		3030	176	nc	NO	YES
LL9SB-046-0001-SO	11/06/03	4-6 ft	Sodium	1100		145	--[n]	0	YES	NO
LL9SB-046-0001-SO	11/06/03	4-6 ft	Vanadium	12		37.6	7.8	nc	NO	YES
LL9SB-047-0001-SO	11/06/03	3-5 ft	Aluminum	8600		19500	7614	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-047-0001-SO	11/06/03	3-5 ft	Arsenic	5.9		19.8	0.39	ca	NO	YES
LL9SB-047-0001-SO	11/06/03	3-5 ft	Iron	9600		35200	2346	nc	NO	YES
LL9SB-047-0001-SO	11/06/03	3-5 ft	Sodium	1200		145	--[n]	0	YES	NO
LL9SB-047-0001-SO	11/06/03	3-5 ft	Vanadium	9.2		37.6	7.8	nc	NO	YES
LL9SB-048-0001-SO	12/10/03	1-3 ft	Aluminum	8400		19500	7614	nc	NO	YES
LL9SB-048-0001-SO	12/10/03	1-3 ft	Arsenic	11		19.8	0.39	ca	NO	YES
LL9SB-048-0001-SO	12/10/03	1-3 ft	Iron	21000		35200	2346	nc	NO	YES
LL9SB-048-0001-SO	12/10/03	1-3 ft	Lead	41		19.1	400	pbk	YES	NO
LL9SB-048-0001-SO	12/10/03	1-3 ft	Manganese	590		3030	176	nc	NO	YES
LL9SB-048-0001-SO	12/10/03	1-3 ft	Mercury	0.34		0.04	2.3	nc	YES	NO
LL9SB-048-0001-SO	12/10/03	1-3 ft	Vanadium	14		37.6	7.8	nc	NO	YES
LL9SB-049-0001-SO	12/10/03	1-3 ft	Aluminum	12000		19500	7614	nc	NO	YES
LL9SB-049-0001-SO	12/10/03	1-3 ft	Arsenic	12		19.8	0.39	ca	NO	YES
LL9SB-049-0001-SO	12/10/03	1-3 ft	Barium	150		124	538	nc	YES	NO
LL9SB-049-0001-SO	12/10/03	1-3 ft	Iron	27000		35200	2346	nc	NO	YES
LL9SB-049-0001-SO	12/10/03	1-3 ft	Lead	64		19.1	400	pbk	YES	NO
LL9SB-049-0001-SO	12/10/03	1-3 ft	Manganese	530		3030	176	nc	NO	YES
LL9SB-049-0001-SO	12/10/03	1-3 ft	Vanadium	20		37.6	7.8	nc	NO	YES
LL9SB-049-0001-SO	12/10/03	1-3 ft	Zinc	290		93.3	2346	nc	YES	NO
LL9SB-050-0001-SO	12/10/03	1-3 ft	Aluminum	12000		19500	7614	nc	NO	YES
LL9SB-050-0001-SO	12/10/03	1-3 ft	Arsenic	10		19.8	0.39	ca	NO	YES
LL9SB-050-0001-SO	12/10/03	1-3 ft	Iron	23000		35200	2346	nc	NO	YES
LL9SB-050-0001-SO	12/10/03	1-3 ft	Manganese	370		3030	176	nc	NO	YES
LL9SB-050-0001-SO	12/10/03	1-3 ft	Nitrocellulose	4.3	J	0	--	0	YES	NO
LL9SB-050-0001-SO	12/10/03	1-3 ft	Vanadium	22		37.6	7.8	nc	NO	YES
LL9SB-051-0001-SO	12/08/03	1-3 ft	Aluminum	11000		19500	7614	nc	NO	YES
LL9SB-051-0001-SO	12/08/03	1-3 ft	Arsenic	18		19.8	0.39	ca	NO	YES
LL9SB-051-0001-SO	12/08/03	1-3 ft	Cadmium	0.11	B	0	3.7	nc	YES	NO
LL9SB-051-0001-SO	12/08/03	1-3 ft	Iron	24000		35200	2346	nc	NO	YES
LL9SB-051-0001-SO	12/08/03	1-3 ft	Manganese	350		3030	176	nc	NO	YES
LL9SB-051-0001-SO	12/08/03	1-3 ft	Vanadium	19		37.6	7.8	nc	NO	YES
LL9SB-052-0001-SO	12/08/03	1-3 ft	Aluminum	16000		19500	7614	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-052-0001-SO	12/08/03	1-3 ft	Arsenic	16		19.8	0.39	ca	NO	YES
LL9SB-052-0001-SO	12/08/03	1-3 ft	Iron	29000		35200	2346	nc	NO	YES
LL9SB-052-0001-SO	12/08/03	1-3 ft	Manganese	190		3030	176	nc	NO	YES
LL9SB-052-0001-SO	12/08/03	1-3 ft	Vanadium	25		37.6	7.8	nc	NO	YES
LL9SB-053-0001-DUP	12/11/03	1-3 ft	Arsenic	9.4		19.8	0.39	ca	NO	YES
LL9SB-053-0001-DUP	12/11/03	1-3 ft	Iron	13000	H	35200	2346	nc	NO	YES
LL9SB-053-0001-DUP	12/11/03	1-3 ft	Manganese	500		3030	176	nc	NO	YES
LL9SB-053-0001-DUP	12/11/03	1-3 ft	Vanadium	14		37.6	7.8	nc	NO	YES
LL9SB-053-0001-SO	12/11/03	1-3 ft	Arsenic	11		19.8	0.39	ca	NO	YES
LL9SB-053-0001-SO	12/11/03	1-3 ft	Iron	15000	H	35200	2346	nc	NO	YES
LL9SB-053-0001-SO	12/11/03	1-3 ft	Manganese	550		3030	176	nc	NO	YES
LL9SB-053-0001-SO	12/11/03	1-3 ft	RDX	0.11		0	4.4	ca	YES	NO
LL9SB-053-0001-SO	12/11/03	1-3 ft	Vanadium	13		37.6	7.8	nc	NO	YES
LL9SB-065-0001-SO	11/06/03	3-5 ft	Aluminum	10000		19500	7614	nc	NO	YES
LL9SB-065-0001-SO	11/06/03	3-5 ft	Arsenic	4.6		19.8	0.39	ca	NO	YES
LL9SB-065-0001-SO	11/06/03	3-5 ft	Iron	14000		35200	2346	nc	NO	YES
LL9SB-065-0001-SO	11/06/03	3-5 ft	Lead	51		19.1	400	pbk	YES	NO
LL9SB-065-0001-SO	11/06/03	3-5 ft	Mercury	9.7		0.04	2.3	nc	YES	YES
LL9SB-065-0001-SO	11/06/03	3-5 ft	Sodium	1300		145	--[n]	0	YES	NO
LL9SB-065-0001-SO	11/06/03	3-5 ft	Vanadium	17		37.6	7.8	nc	NO	YES
LL9SB-066-0001-SO	11/06/03	5-7 ft	Arsenic	11		19.8	0.39	ca	NO	YES
LL9SB-066-0001-SO	11/06/03	5-7 ft	Iron	19000		35200	2346	nc	NO	YES
LL9SB-066-0001-SO	11/06/03	5-7 ft	Manganese	340		3030	176	nc	NO	YES
LL9SB-066-0001-SO	11/06/03	5-7 ft	Sodium	1100		145	--[n]	0	YES	NO
LL9SB-066-0001-SO	11/06/03	5-7 ft	Vanadium	9.8		37.6	7.8	nc	NO	YES
LL9SB-069-0001-SO	10/27/03	7.0 ft	Arsenic	6.9		19.8	0.39	ca	NO	YES
LL9SB-069-0001-SO	10/27/03	7.0 ft	Iron	9800	*H	35200	2346	nc	NO	YES
LL9SB-070-0001-SO	10/27/03	6.5 ft	Aluminum	7800		19500	7614	nc	NO	YES
LL9SB-070-0001-SO	10/27/03	6.5 ft	Arsenic	8.3		19.8	0.39	ca	NO	YES
LL9SB-070-0001-SO	10/27/03	6.5 ft	Iron	21000	*H	35200	2346	nc	NO	YES



Table 4-5
Summary of Subsurface Soil Exceedances (>1 ft)

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Subsurface Soil Background Criteria	Subsurface Soil Region 9 Criteria	SB PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SB-070-0001-SO	10/27/03	6.5 ft	Manganese	490		3030	176	nc	NO	YES
LL9SB-070-0001-SO	10/27/03	6.5 ft	Vanadium	14		37.6	7.8	nc	NO	YES

Notes:

-- - no background/PRG value is available for this analyte

PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL)

nc - non-cancer basis

ca - cancer basis

pbk - based on PBK modeling

mcl - based on CWA maximum contaminant level

max - ceiling limit

sat - soil saturation

[n] - nutrient

UC - unconsolidated

Inorganic Qualifiers

U - analyte was not detected at or above the stated limit

J - result is less than the RL, but greater than or equal to the MDL

B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL

InOrganic Flags

^ - instrument related QC exceed the upper or lower control limits

* - batch QC exceeds the upper or lower control limit

H - batch QC is greater than RL

Organic Qualifiers

U - analyte was not detected at or above the stated limit

J - result is an estimated value below the RL

Organic Flags

B - Method Blank: batch QC is greater than RL

* - batch QC exceeds the upper or lower control limit

^ - batch QC is greater than RL

a - concentration is below the method Reporting Limit (RL)

H - alternate peak selection upon analytical review



Table 4-7
Summary of Groundwater Results

							LL9MW-001-0001 -GW		LL9MW-002-0001 -GW		LL9MW-003-0001 -GW		LL9MW-004-0001 -GW		LL9MW-005-0001 -GW		LL9MW-006-0001 -GW		LL9MW-007-0001 -DUP		LL9MW-007-0001 -GW	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval	
Group	Method	Analyte	Units	RVAAP Consolidated Wells Background Criteria	Consolidated Wells Region 9 (Tap Water) PRGs	C-MW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	µg/L	0.00	730	nc	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U
Explosives 8330	8330	1,3,5-Trinitrobenzene	µg/L	0	1095	nc	0.11	U	0.1	U	0.08	U	0.11	U	0.096	U	0.08	U	0.084	U	0.12	U
		1,3-Dinitrobenzene	µg/L	0	3.6	nc	0.071	U	0.066	U	0.053	U	0.07	U	0.064	U	0.053	U	0.056	U	0.08	U
		2,4,6-TNT	µg/L	0	2.2	ca	0.091	U	0.085	U	0.068	U	0.09	U	0.082	U	0.068	U	0.071	U	0.1	U
		2,4-Dinitrotoluene	µg/L	0	73	nc	0.056	U	0.052	U	0.042	U	0.055	U	0.05	U	0.042	U	0.044	U	0.063	U
		2,6-Dinitrotoluene	µg/L	0	36	nc	0.28	U	0.26	U	0.21	U	0.27	U	0.25	U	0.21	U	0.22	U	0.31	U
		2-Nitrotoluene	µg/L	0	0.049	ca	0.22	U	0.2	U	0.16	U	0.21	U	0.2	U	0.16	U	0.17	U	0.24	U
		3-Nitrotoluene	µg/L	0	122	nc	0.14	U	0.13	U	0.1	U	0.13	U	0.12	U	0.1	U	0.11	U	0.15	U
		4-Nitrotoluene	µg/L	0	0.66	ca	0.45	U	0.42	U	0.34	U	0.44	U	0.4	U	0.34	U	0.35	U	0.51	U
		HMX	µg/L	0	1825	nc	0.3	U	0.28	U	0.22	U	0.3	U	0.27	U	0.22	U	0.24	U	0.34	U
		Nitrobenzene	µg/L	0	3.4	nc	0.12	U	0.12	U	0.092	U	0.12	U	0.11	U	0.092	U	0.097	U	0.14	U
		RDX	µg/L	0	0.61	ca	0.18	U	0.17	U	0.13	U	0.18	U	0.16	U	0.13	U	0.14	U	0.2	U
		Tetryl	µg/L	0	365	nc	0.29	U	0.27	U	0.22	U	0.29	U	0.26	U	0.22	U	0.23	U	0.33	U
PCB TCL 8082	8082	Aroclor 1016	µg/L	--	0.96	ca	0.16	U	0.17	U	0.17	U	0.16	U	0.17	U	0.16	U	0.17	U	0.17	U
		Aroclor 1221	µg/L	--	0.034	ca	0.44	U	0.46	U	0.45	U	0.44	U	0.46	U	0.43	U	0.47	U	0.46	U
		Aroclor 1232	µg/L	--	0.034	ca	0.21	U	0.22	U	0.21	U	0.21	U	0.22	U	0.21	U	0.22	U	0.22	U
		Aroclor 1242	µg/L	--	0.034	ca	0.18	U	0.19	U	0.18	U	0.18	U	0.19	U	0.18	U	0.19	U	0.19	U
		Aroclor 1248	µg/L	--	0.034	ca	0.2	U	0.21	U	0.2	U	0.2	U	0.21	U	0.2	U	0.21	U	0.21	U
		Aroclor 1254	µg/L	--	0.034	ca	0.12	U	0.13	U	0.13	U	0.12	U	0.13	U	0.12	U	0.13	U	0.13	U
		Aroclor 1260	µg/L	--	0.034	ca	0.14	U	0.15	U	0.15	U	0.14	U	0.15	U	0.14	U	0.15	U	0.15	U
Pesticides 8081A	8081A	4,4'-DDD	µg/L	--	0.28	ca	0.034	U	0.036	U	0.035	U	0.034	U	0.036	U	0.034	U	0.037	U	0.036	U
		4,4'-DDE	µg/L	--	0.20	ca	0.022	U	0.023	U	0.022	U	0.022	U	0.023	U	0.021	U	0.023	U	0.023	U
		4,4'-DDT	µg/L	--	0.20	ca	0.047	U	0.049	U	0.048	U	0.047	U	0.049	U	0.046	U	0.05	U	0.049	U
		Aldrin	µg/L	--	0.0040	ca	0.027	U	0.028	U	0.027	U	0.027	U	0.028	U	0.026	U	0.029	U	0.028	U
		alpha-BHC	µg/L	--	0.011	nc	0.045	U	0.047	U	0.046	U	0.045	U	0.047	U	0.044	U	0.048	U	0.047	U
		alpha-Chlordane	µg/L	--	0.19	ca	0.015	U	0.016	U	0.016	U	0.015	U	0.016	U	0.015	U	0.016	U	0.016	U
		beta-BHC	µg/L	--	0.037	ca	0.027	U	0.028	U	0.027	U	0.027	U	0.028	U	0.026	U	0.029	U	0.028	U
		delta-BHC	µg/L	--	--	0	0.024	U	0.025	U	0.024	U	0.024	U	0.025	U	0.023	U	0.026	U	0.025	U
		Dieldrin	µg/L	--	0.0042	ca	0.017	U	0.018	U	0.017	U	0.017	U	0.018	U	0.017	U	0.018	U	0.018	U
		Endosulfan I	µg/L	--	220	nc	0.02	U	0.021	U	0.02	U	0.02	U	0.021	U	0.02	U	0.021	U	0.021	U
		Endosulfan II	µg/L	--	220	nc	0.04	U	0.042	U	0.041	U	0.04	U	0.042	U	0.039	U	0.043	U	0.042	U
		Endosulfan sulfate	µg/L	--	220	nc	0.042	U	0.044	U	0.043	U	0.042	U	0.044	U	0.041	U	0.045	U	0.044	U
		Endrin	µg/L	--	11	nc	0.016	U	0.017	U	0.017	U	0.016	U	0.017	U	0.016	U	0.017	U	0.017	U
		Endrin aldehyde	µg/L	--	--	0	0.033	U	0.035	U*	0.034	U*	0.033	U	0.035	U	0.033	U*	0.036	U	0.035	U
		Endrin ketone	µg/L	--	--	0	0.028	U	0.029	U*	0.028	U*	0.028	U	0.029	U	0.027	U*	0.03	U	0.029	U
		gamma-BHC (Lindane)	µg/L	--	0.052	ca	0.04	U	0.042	U	0.041	U	0.04	U	0.042	U	0.039	U	0.043	U	0.042	U
		gamma-Chlordane	µg/L	--	0.19	ca	0.016	U	0.017	U	0.017	U	0.016	U	0.017	U	0.016	U	0.017	U	0.017	U
		Heptachlor	µg/L	--	0.015	ca	0.039	U	0.041	U	0.04	U	0.039	U	0.041	U	0.038	U	0.042	U	0.041	U
		Heptachlor epoxide	µg/L	--	0.0074	ca	0.034	U	0.036	U	0.035	U	0.034	U	0.036	U	0.034	U	0.037	U	0.036	U
		Methoxychlor	µg/L	--	182	nc	0.16	U	0.17	U	0.17	U	0.16	U	0.17	U	0.16	U	0.17	U	0.17	U
		Toxaphene	µg/L	--	0	0	0.13	U	0.14	U	0.14	U	0.13	U	0.14	U	0.13	U	0.14	U	0.14	U



Table 4-7
Summary of Groundwater Results

							LL9MW-001 -0001 -GW		LL9MW-002 -0001 -GW		LL9MW-003 -0001 -GW		LL9MW-004 -0001 -GW		LL9MW-005 -0001 -GW		LL9MW-006 -0001 -GW		LL9MW-007 -0001 -DUP		LL9MW-007 -0001 -GW	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval		Screened Interval	
Group	Method	Analyte	Units	RVAAP Consolidated Wells Background Criteria	Consolidated Wells Region 9 (Tap Water) PRGs	C-MW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	µg/L	0	--	0	0.0005	U	0.000089	B J	0.00014	B J	0.00011	B J	0.00011	B J	0.0005	U	0.0001	B J	0.00012	B J
	8330	Nitroglycerin	µg/L	0	--	0	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U
	UV/HPLC per SOP	Nitroguanidine	µg/L	0	3650	nc	20	U	20	U	20	U	20	U	20	U	20	U	20	U	20	U
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	µg/L	--	7.2	nc	0.33	U	0.35	U	0.33	U	0.33	U	0.35	U	0.34	U	0.33	U	0.32	U
		1,2-Dichlorobenzene	µg/L	--	370	nc	0.34	U	0.36	U	0.34	U	0.34	U	0.36	U	0.35	U	0.34	U	0.33	U
		1,3-Dichlorobenzene	µg/L	--	182	nc	0.41	U	0.44	U	0.41	U	0.41	U	0.44	U	0.43	U	0.42	U	0.41	U
		1,4-Dichlorobenzene	µg/L	--	0.50	ca	0.32	U	0.34	U	0.32	U	0.32	U	0.34	U	0.33	U	0.32	U	0.31	U
		2,2-oxybis (1-chloropropane)	µg/L	--	0.27	ca	0.27	U	0.29	U	0.27	U	0.27	U	0.29	U	0.28	U	0.27	U	0.26	U
		2,4,5-Trichlorophenol	µg/L	--	3650	nc	1.3	U	1.4	U	1.3	U	1.3	U	1.4	U	1.4	U	1.4	U	1.3	U
		2,4,6-Trichlorophenol	µg/L	--	3.6	nc	0.2	U	0.21	U	0.2	U	0.2	U	0.21	U	0.21	U	0.2	U	0.2	U
		2,4-Dichlorophenol	µg/L	--	109	nc	0.87	U	0.93	U	0.87	U	0.87	U	0.93	U	0.9	U	0.88	U	0.86	U
		2,4-Dimethylphenol	µg/L	--	730	nc	1.2	U	1.3	U	1.2	U	1.2	U	1.3	U	1.3	U	1.3	U	1.2	U
		2,4-Dinitrophenol	µg/L	--	73	nc	3.2	U	3.4	U	3.2	U	3.2	U	3.4	U	3.3	U	3.2	U	3.1	U
		2-Chloronaphthalene	µg/L	--	487	nc	0.25	U	0.27	U	0.25	U	0.25	U	0.27	U	0.26	U	0.25	U	0.25	U
		2-Chlorophenol	µg/L	--	30	nc	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.11	U
		2-Methylnaphthalene	µg/L	--	--	0	0.12	U	0.13	U	0.12	U	0.12	U	0.13	U	0.13	U	0.13	U	0.12	U
		2-Methylphenol (o-cresol)	µg/L	--	1825	nc	0.25	U	0.27	U	0.25	U	0.25	U	0.27	U	0.26	U	0.25	U	0.25	U
		2-Nitroaniline	µg/L	--	109	nc	0.21	U	0.22	U	0.21	U	0.21	U	0.22	U	0.22	U	0.21	U	0.21	U
		2-Nitrophenol	µg/L	--	--	0	0.79	U	0.84	U	0.79	U	0.79	U	0.84	U	0.81	U	0.8	U	0.77	U
		3,3-Dichlorobenzidine	µg/L	--	0.15	ca	0.69	U	0.73	U	0.69	U	0.69	U	0.73	U	0.71	U	0.7	U	0.68	U
		3-Nitroaniline	µg/L	--	3.2	ca	2	U	2.1	U	2	U	2	U	2.1	U	2.1	U	2	U	2	U
		4,6-Dinitro-2-methylphenol	µg/L	--	3.6	nc	2.3	U	2.4	U	2.3	U	2.3	U	2.4	U	2.4	U	2.3	U	2.3	U
		4-Bromophenyl phenyl ether	µg/L	--	--	0	0.18	U	0.19	U	0.18	U	0.18	U	0.19	U	0.19	U	0.18	U	0.18	U
		4-Chloro-3-methylphenol	µg/L	--	--	0	2.3	U	2.4	U	2.3	U	2.3	U	2.4	U	2.4	U	2.3	U	2.3	U
		4-Chloroaniline	µg/L	--	146	nc	2.7	U	2.9	U	2.7	U	2.7	U	2.9	U	2.8	U	2.7	U	2.6	U
		4-Chlorophenyl phenyl ether	µg/L	--	--	0	0.72	U	0.77	U	0.72	U	0.72	U	0.77	U	0.74	U	0.73	U	0.71	U
		4-Methylphenol (m/p-cresol)	µg/L	--	182	nc	0.096	U	0.1	U	0.096	U	0.096	U	0.1	U	0.099	U	0.097	U	0.094	U
		4-Nitroaniline	µg/L	--	3.2	ca	2.2	U	2.3	U	2.2	U	2.2	U	2.3	U	2.3	U	2.2	U	2.2	U
		4-Nitrophenol	µg/L	--	--	0	3.6	U	3.8	U	3.6	U	3.6	U	3.8	U	3.7	U	3.6	U	3.5	U
		Acenaphthene	µg/L	--	365	nc	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.11	U
		Acenaphthylene	µg/L	--	--	0	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.11	U
		Anthracene	µg/L	--	1825	nc	0.14	U	0.15	U	0.14	U	0.14	U	0.15	U	0.15	U	0.15	U	0.14	U
		Benzo(a)anthracene	µg/L	--	0.092	ca	0.047	U	0.05	U	0.047	U	0.047	U	0.05	U	0.049	U	0.048	U	0.046	U
		Benzo(a)pyrene	µg/L	--	0.0092	ca	0.081	U	0.086	U	0.081	U	0.081	U	0.086	U	0.083	U	0.082	U	0.079	U
		Benzo(b)fluoranthene	µg/L	--	0.092	ca	0.064	U	0.068	U	0.064	U	0.064	U	0.068	U	0.066	U	0.065	U	0.063	U
		Benzo(ghi)perylene	µg/L	--	--	0	0.18	U	0.19	U	0.18	U	0.18	U	0.19	U	0.19	U	0.18	U	0.18	U
		Benzo(k)fluoranthene	µg/L	--	0.92	ca	0.069	U	0.073	U	0.069	U	0.069	U	0.073	U	0.071	U	0.07	U	0.068	U
		Bis(2-chloroethoxy)methane	µg/L	--	--	0	0.3	U	0.32	U	0.3	U	0.3	U	0.32	U	0.31	U	0.3	U	0.29	U
		Bis(2-chloroethyl)ether	µg/L	--	0.010	ca	0.29	U	0.31	U	0.29	U	0.29	U	0.31	U	0.3	U	0.29	U	0.28	U
		Bis(2-ethylhexyl)phthalate	µg/L	--	4.8	ca	3.7	U	4	U	3.7	U	3.7	U	4	U	3.9	U	3.8	U	4.3	J
		Butyl benzyl phthalate	µg/L	--	7300	nc	0.37	U	0.4	U	0.37	U	0.37	U	0.4	U	0.39	U	0.38	U	0.37	U
		Carbazole	µg/L	--	3.4	ca	0.28	U	0.3	U	0.28	U	0.28	U	0.3	U	0.29	U	0.28	U	0.27	U
		Chrysene	µg/L	--	9.2	ca	0.043	U	0.046	U	0.043	U	0.043	U	0.046	U	0.045	U	0.044	U	0.042	U



Table 4-7
Summary of Groundwater Results

							LL9MW-001-0001-GW		LL9MW-002-0001-GW		LL9MW-003-0001-GW		LL9MW-004-0001-GW		LL9MW-005-0001-GW		LL9MW-006-0001-GW		LL9MW-007-0001-DUP		LL9MW-007-0001-GW	
							Sample Date		12/17/03		12/16/03		12/16/03		12/17/03		12/17/03		12/16/03		12/15/03	
							Screened Interval		10.5-20.5 ft.		10-20 ft.		11.5-21.5 ft.		22-32 ft		10-20 ft		16-26 ft.		8.5-18.5 ft	
Group	Method	Analyte	Units	RVAAP Consolidated Wells Background Criteria	Consolidated Wells Region 9 (Tap Water) PRGs	C-MW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzo(a,h)anthracene	µg/L	--	0.0092	ca	0.12	U	0.13	U	0.12	U	0.12	U	0.13	U	0.13	U	0.13	U	0.12	U
		Dibenzofuran	µg/L	--	12	nc	0.12	U	0.13	U	0.12	U	0.12	U	0.13	U	0.13	U	0.13	U	0.12	U
		Diethyl phthalate	µg/L	--	29199	nc	0.14	U	0.15	U	0.14	U	0.14	U	0.15	U	0.15	U	0.15	U	0.14	U
		Dimethyl phthalate	µg/L	--	364867	nc	0.2	U	0.21	U	0.2	U	0.2	U	0.21	U	0.21	U	0.2	U	0.2	U
		Di-n-butyl phthalate	µg/L	--	3650	nc	0.35	U	0.37	U	0.35	U	0.35	U	0.37	U	0.36	U	0.35	U	0.34	U
		Di-n-octyl phthalate	µg/L	--	1460	nc	2.4	U	2.6	U	2.4	U	2.4	U	2.6	U	2.5	U	2.4	U	2.4	U
		Fluoranthene	µg/L	--	1460	nc	0.13	U	0.14	U	0.13	U	0.13	U	0.14	U	0.14	U	0.14	U	0.13	U
		Fluorene	µg/L	--	243	nc	0.12	U	0.13	U	0.12	U	0.12	U	0.13	U	0.13	U	0.13	U	0.12	U
		Hexachlorobenzene	µg/L	--	0.042	ca	0.093	U	0.099	U	0.093	U	0.093	U	0.099	U	0.096	U	0.094	U	0.092	U
		Hexachlorobutadiene	µg/L	--	0.86	ca	0.62	U	0.65	U	0.62	U	0.62	U	0.65	U	0.63	U	0.62	U	0.6	U
		Hexachlorocyclopentadiene	µg/L	--	219	nc	0.62	U	0.66	U	0.62	U	0.62	U	0.66	U	0.64	U	0.63	U	0.61	U
		Hexachloroethane	µg/L	--	4.8	ca	0.59	U	0.62	U	0.59	U	0.59	U	0.62	U	0.6	U	0.59	U	0.58	U
		Indeno(1,2,3-cd)pyrene	µg/L	--	0.092	ca	0.083	U	0.088	U	0.083	U	0.083	U	0.088	U	0.085	U	0.083	U	0.081	U
		Isophorone	µg/L	--	71	ca	0.25	U	0.27	U	0.25	U	0.25	U	0.27	U	0.26	U	0.25	U	0.25	U
		Naphthalene	µg/L	--	6.2	nc	0.15	U	0.16	U	0.15	U	0.15	U	0.16	U	0.16	U	0.16	U	0.15	U
		n-Nitroso-di-n-propylamine	µg/L	--	0.0096	ca	0.078	U	0.083	U	0.078	U	0.078	U	0.083	U	0.08	U	0.079	U	0.076	U
		n-Nitrosodiphenylamine	µg/L	--	14	ca	0.12	U	0.13	U	0.12	U	0.12	U	0.13	U	0.13	U	0.13	U	0.12	U
		Pentachlorophenol	µg/L	--	0.56	ca	1.6	U	1.7	U	1.6	U	1.6	U	1.7	U	1.7	U	1.7	U	1.6	U
		Phenanthrene	µg/L	--	--	0	0.13	U	0.14	U	0.13	U	0.13	U	0.14	U	0.14	U	0.14	U	0.13	U
		Phenol	µg/L	--	10950	nc	0.34	U	0.36	U	0.34	U	0.34	U	0.36	U	0.35	U	0.34	U	0.33	U
		Pyrene	µg/L	--	182	nc	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.11	U
TAL Metals 6010B	6010B	Aluminum	µg/L	--	36499	nc	24	U	170	B	31	B	24	U	25	B	76	B	24	U	24	U
		Barium	µg/L	256	2555	nc	15		4.7	B	26		40		6	B	69		18		18	
		Beryllium	µg/L	0.00	73	nc	0.17	U	0.17	U	0.17	U	0.17	U	0.17	U	0.17	U	0.17	U	0.17	U
		Cadmium	µg/L	0.00	18	nc	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U
		Calcium	µg/L	53100	--	0	34000	H	17000	H	14000	H	13000	H	8800	H	4400	H	11000	H	12000	H
		Chromium	µg/L	0.00	109	nc	1.5	U	1.5	U	1.5	U	1.5	U	1.5	U	1.5	U	1.5	U	1.5	U
		Cobalt	µg/L	0.00	730	nc	1	U	1	U	1	U	34		1	U	1	U	13		11	
		Copper	µg/L	0.00	1460	nc	2.3	B	3.1	B	2.9	B	14		2.8	B	4.2	B	4.4	B	5	B
		Iron	µg/L	1430	10950	nc	40	U	40	U	62		5700		40	U	40	U	6600		7100	
		Magnesium	µg/L	15000	--	0	9600		6800		5400		10000		3700		5000		6000		6400	
		Manganese	µg/L	1340	876	nc	160		190		140		2400		15		73		880		900	
		Nickel	µg/L	83.4	730	nc	1.9	U	29		5.1	B	45		2.3	B	9.8	B	25		24	
		Potassium	µg/L	5770	--	0	1200		1200		1900		1200		750		1400		1300		1400	
		Selenium	µg/L	0.00	182	nc	5	U	5	U	5	U	5	U	5	U	5	U	5	U	5	U
		Silver	µg/L	0.00	182	nc	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U
		Sodium	µg/L	51400	--	0	4600		2600		3600		5500		3200		2200		3700		3700	
		Vanadium	µg/L	0.00	36	nc	2.1	U	2.1	U	2.1	U	2.1	U	2.1	U	2.1	U	2.1	U	2.1	U
		Zinc	µg/L	52.3	10950	nc	10	U	13		40		31		24		21		38		38	
		Antimony	µg/L	0.00	15	nc	3.2	B	2.8	B	4	B	2.2	U	2.2	U	2.5	B	2.2	U	3	B
		Arsenic	µg/L	0.00	0.045	ca	0.73	U	0.73	U	0.73	U	0.73	U	0.73	U	0.73	U	0.73	U	0.73	U
		Lead	µg/L	0.00	15	mcl	0.78	U	0.78	U	0.78	U	0.78	U	0.78	U	0.78	U	0.78	U	0.78	U
		Mercury	µg/L	0.00	11	nc	0.049	U	0.049	U	0.049	U	0.049	U	0.049	U	0.049	U	0.049	U	0.049	U
		Thallium	µg/L	0.00	2.4	nc	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	µg/L	--	3172	nc	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U
		1,1,2,2-Tetrachloroethane	µg/L	--	0.055	ca	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U
		1,1,2-Trichloroethane	µg/L	--	0.20	ca	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U
		1,1-Dichloroethane	µg/L	--	811	nc	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U
		1,1-Dichloroethene	µg/L	--	339	nc	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U



Table 4-7
Summary of Groundwater Results

							LL9MW-001-0001-GW		LL9MW-002-0001-GW		LL9MW-003-0001-GW		LL9MW-004-0001-GW		LL9MW-005-0001-GW		LL9MW-006-0001-GW		LL9MW-007-0001-DUP		LL9MW-007-0001-GW	
							Sample Date		12/17/03		12/16/03		12/16/03		12/17/03		12/17/03		12/16/03		12/15/03	
							Screened Interval		10.5-20.5 ft.		10-20 ft.		11.5-21.5 ft.		22-32 ft		10-20 ft		16-26 ft.		8.5-18.5 ft	
Group	Method	Analyte	Units	RVAAP Consolidated Wells Background Criteria	Consolidated Wells Region 9 (Tap Water) PRGs	C-MW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		1,2-Dichloroethane	µg/L	--	0.12	ca	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U
		1,2-Dichloroethene (total)	µg/L	--	120	nc	0.23	U	0.23	U	0.23	U	0.23	U	0.23	U	0.23	U	0.23	U	0.23	U
		1,2-Dichloropropane	µg/L	--	0.16	ca	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U
		2-Butanone (MEK)	µg/L	--	6968	nc	1.2	U	1.2	U	1.2	U	1.2	U	1.2	U	1.2	U	1.2	U	1.2	U
		2-Hexanone	µg/L	--	2000	nc	0.53	U	0.53	U	0.53	U	0.53	U	0.53	U	0.53	U	0.53	U	0.53	U
		4-Methyl-2-pentanone (MIBK)	µg/L	--	1993	nc	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U	0.65	U
		Acetone	µg/L	--	5475	nc	1.8	U	1.8	U	1.8	U	1.8	U	1.8	U	1.8	U	1.8	U	1.8	U
		Benzene	µg/L	--	0.35	ca	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U
		Bromodichloromethane	µg/L	--	0.18	ca	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U
		Bromoform	µg/L	--	8.5	ca	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U
		Bromomethane	µg/L	--	8.7	nc	0.1	U	0.1	U*	0.1	U	0.1	U	0.1	U	0.1	U*	0.1	U*	0.1	U*
		Carbon disulfide	µg/L	--	1043	nc	0.2	U	0.2	U	0.2	U	0.2	U	0.2	U	0.2	U	0.2	U	0.2	U
		Carbon tetrachloride	µg/L	--	0.17	ca	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U
		Chlorobenzene	µg/L	--	106	nc	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U
		Chloroethane	µg/L	--	4.6	ca	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U
		Chloroform	µg/L	--	0.17	ca	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U	0.11	U
		Chloromethane	µg/L	--	158	nc	0.08	U	0.08	U*	0.08	U	0.08	U	0.08	U	0.08	U*	0.08	U*	0.08	U*
		cis-1,3-Dichloropropene	µg/L	--	0.40	ca	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U	0.12	U
		Dibromochloromethane	µg/L	--	0.13	ca	0.06	U	0.06	U	0.06	U	0.06	U	0.06	U	0.06	U	0.06	U	0.06	U
		Ethylbenzene	µg/L	--	1340	nc	0.07	U	0.07	U	0.07	U	0.07	U	0.07	U	0.07	U	0.07	U	0.07	U
		Methylene chloride	µg/L	--	4.3	ca	0.35	U	0.35	U	0.35	U	0.35	U	0.35	U	0.35	U	0.35	U	0.35	U
		Styrene	µg/L	--	1641	nc	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U	0.13	U
		Tetrachloroethene	µg/L	--	0.10	ca	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U	0.09	U
		Toluene	µg/L	--	723	nc	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U
		trans-1,3-Dichloropropene	µg/L	--	0.40	ca	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U	0.15	U
		Trichloroethene	µg/L	--	0.028	ca	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U	0.1	U
		Vinyl chloride	µg/L	--	0.020	ca	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U	0.08	U
		Xylenes (total)	µg/L	--	0.020	ca	0.28	U	0.28	U	0.28	U	0.28	U	0.28	U	0.28	U	0.28	U	0.28	U

Notes:
-- - no background/PRG value is available for this analyte
blank cell indicates that the analysis was not performed
PRG - preliminary remediation goals
(The PRG for lead is the Maximum Contaminant Level (MCL)
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background and PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG and Background, then the value is presented with a normal style

Inorganic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL
InOrganic Flags
^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL
Organic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL
Organic Flags
B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-8
Summary of Groundwater Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result	Units	Qualifier	RVAAP Consolidated Wells Background Criteria	Region 9 (Tap Water) PRGs	C-MW PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9MW-001-0001-GW	12/17/2003	10.5-20.5 ft	Copper	2.3	µg/L	H	0.00	1460	nc	YES	NO
LL9MW-001-0001-GW	12/17/2003	10.5-20.5 ft	Antimony	3.2	µg/L		0.00	15	nc	YES	NO
LL9MW-002-0001-GW	12/16/2003	10-20 ft.	Nitrocellulose	0.000089	µg/L	BJ	0	--	0	YES	NO
LL9MW-002-0001-GW	12/16/2003	10-20 ft.	Copper	3.1	µg/L	B	0.00	1460	nc	YES	NO
LL9MW-002-0001-GW	12/16/2003	10-20 ft.	Antimony	2.8	µg/L	B	0.00	15	nc	YES	NO
LL9MW-003-0001-GW	12/16/2003	11.5-21.5 ft.	Nitrocellulose	0.00014	µg/L	BJ	0	--	0	YES	NO
LL9MW-003-0001-GW	12/16/2003	11.5-21.5 ft.	Copper	2.9	µg/L	B	0.00	1460	nc	YES	NO
LL9MW-003-0001-GW	12/16/2003	11.5-21.5 ft.	Antimony	4	µg/L	B	0.00	15	nc	YES	NO
LL9MW-004-0001-GW	12/17/2003	22-32 ft	Nitrocellulose	0.00011	µg/L		0	--	0	YES	NO
LL9MW-004-0001-GW	12/17/2003	22-32 ft	Cobalt	34	µg/L		0.00	730	nc	YES	NO
LL9MW-004-0001-GW	12/17/2003	22-32 ft	Copper	14	µg/L		0.00	1460	nc	YES	NO
LL9MW-004-0001-GW	12/17/2003	22-32 ft	Iron	5700	µg/L		1430	10950	nc	YES	NO
LL9MW-004-0001-GW	12/17/2003	22-32 ft	Manganese	2400	µg/L		1340	876	nc	YES	YES
LL9MW-005-0001-GW	12/17/2003	10-20 ft	Nitrocellulose	0.00011	µg/L	BJ	0	--	0	YES	NO
LL9MW-005-0001-GW	12/17/2003	10-20 ft	Copper	2.8	µg/L	B	0.00	1460	nc	YES	NO
LL9MW-006-0001-GW	12/16/2003	16-26 ft.	Copper	4.2	µg/L	B	0.00	1460	nc	YES	NO
LL9MW-006-0001-GW	12/16/2003	16-26 ft.	Antimony	2.5	µg/L	B	0.00	15	nc	YES	NO
LL9MW-007-0001-DUP	12/15/2003	8.5-18.5 ft	Nitrocellulose	0.0001	µg/L	BJ	0	--	0	YES	NO
LL9MW-007-0001-DUP	12/15/2003	8.5-18.5 ft	Cobalt	13	µg/L		0.00	730	nc	YES	NO
LL9MW-007-0001-DUP	12/15/2003	8.5-18.5 ft	Copper	4.4	µg/L	B	0.00	1460	nc	YES	NO
LL9MW-007-0001-DUP	12/15/2003	8.5-18.5 ft	Iron	6600	µg/L		1430	10950	nc	YES	NO
LL9MW-007-0001-DUP	12/15/2003	8.5-18.5 ft	Manganese	880	µg/L		1340	876	nc	NO	YES
LL9MW-007-0001-GW	12/15/2003	8.5-18.5 ft	Nitrocellulose	0.00012	µg/L	BJ	0	--	0	YES	NO
LL9MW-007-0001-GW	12/15/2003	8.5-18.5 ft	Cobalt	11	µg/L		0.00	730	nc	YES	NO
LL9MW-007-0001-GW	12/15/2003	8.5-18.5 ft	Copper	5	µg/L	B	0.00	1460	nc	YES	NO
LL9MW-007-0001-GW	12/15/2003	8.5-18.5 ft	Iron	7100	µg/L		1430	10950	nc	YES	NO
LL9MW-007-0001-GW	12/15/2003	8.5-18.5 ft	Manganese	900	µg/L		1340	876	nc	NO	YES
LL9MW-007-0001-GW	12/15/2003	8.5-18.5 ft	Antimony	3	µg/L	B	0.00	15	nc	YES	NO

Notes:

-- - no background/PRG value is available for this analyte
blank cell indicates that the analyte was a non-detect (with a "U" qualifier) or analysis was not performed
PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL))
nc - non-cancer basis
ca - cancer basis

Inorganic Qualifiers

U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL

Organic Qualifiers

U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL



Table 4-9
Azide Screening Surface Water Sample Results Summary

ANALYTE**, UNITS, METHOD NO.	RVAAP Surface Water Background Criteria ug/L	Region 9 PRG Data (Tap Water)	LL9SW-001-0001-SW	LL9SW-005-0001-SW	LL9SW-007-0001-SW	LL9SW-008-0001-SW	LL9SW-008-0001-FD
Sample Date			3/12/2002	3/13/2002	3/12/2002	3/13/2002	3/13/2002
Explosives 8330 ug/L							
HMX	0.0	1825 nc	0.55 U	0.65 U	0.44 U	0.70 U	0.57 U
RDX	0.0	0.61 ca	0.23 U	0.27 U	0.18 U	0.29 U	0.23 U
1,3,5-Trinitrotoluene	0.0	1095 nc	0.23 U	0.27 U	0.18 U	0.29 U	0.23 U
1,3-Dinitrobenzene	0.0	3.6 nc	0.23 U	0.27 U	0.18 U	0.29 U	0.23 U
Nitrobenzene	0.0	3.4 nc	0.23 U	0.27 U	0.18 U	0.29 U	0.23 U
2,4,6-Trinitrtoluene	0.0	2.2 ca	0.23 U	0.27 U	0.18 U	0.29 U	0.23 U
Tetryl	0.0	365 nc	0.44 U	0.52 U	0.35 U	0.55 U	0.45 U
2,4-Dinitrotoluene	0.0	73 nc	0.23 U	0.27 U	0.18 U	0.29 U	0.23 U
2,6-Dinitrotoluene	0.0	36 nc	0.44 U	0.52 U	0.35 U	0.55 U	0.45 U
2-Nitrotoluene	0.0	0.049 ca	0.44 U	0.52 U	0.35 U	0.55 U	0.45 U
4-Nitrtoluene	0.0	0.66 ca	1.1 U	1.3 U	0.88 U	1.4 U	1.1 U
3-Nitrotoluene	0.0	122 nc	0.44 U	0.52 U	0.35 U	0.55 U	0.45 U
TAL Metals 6010B ug/L							
Antimony	0.0	15 nc	3.0 U	3.0 U	3.0 U	3.2	3.0 U
Arsenic	3.2	0.045 ca	2.0 U	8.0	351	3.3	2.0 U
Lead	0.0	15 mcl	2.0 U	62.9	2000	3150	432.0
Thallium	0.0	2.4 nc	2.0 U	2.0 U	2.0 U	2.0 U	2.0 U
Mercury	0.0	11 nc	0.20 U	0.37	21.9	7.7	0.61
Aluminum	3370	36499 nc	339	17800	3030	1610	243
Barium	47.5	2555 nc	21.4	145	208.0	211	93.9
Beryllium	0.0	73 nc	4.0 U	0.70 (B)	4.0 U	4.0 U	4.0 U
Cadmium	0.0	18 nc	2.0 U	1.1 (B)	1.8 (B)	1.0 (B)	2.0 U
Calcium	41400	-- [n]	9830 (H)	13900 (H)	46200 (H)	75200 (H)	62700 (H)
Chromium	0.0	109 nc	10.0 U	20.2	48.8	27.1	10.0 U
Cobalt	0.0	730 nc	5.0 U	6.0	5.9	6.0	5.0 U
Copper	7.9	1460 nc	10.0 U	18.6	193	31	4.4 (B)
Iron	2560.0	10950 nc	249	18900	17900	16100	3380
Magnesium	10800.0	-- [n]	2110	5360	2770	12600	3550
Manganese	391	876 nc	10.1	633	1740	802	271
Nickel	0.0	730 nc	10.0 U	20.1	58.2	86.7	10.9
Potassium	3170	-- [n]	798	3900	8320	14700	12700



Table 4-9
Azide Screening Surface Water Sample Results Summary

ANALYTE**, UNITS, METHOD NO.	RVAAP Surface Water Background Criteria ug/L	Region 9 PRG Data (Tap Water)	LL9SW-001-0001-SW	LL9SW-005-0001-SW	LL9SW-007-0001-SW	LL9SW-008-0001-SW	LL9SW-008-0001-FD
Sample Date			3/12/2002	3/13/2002	3/12/2002	3/13/2002	3/13/2002
Selenium	0	182 nc	10. 0 U	10. 0 U	10.0 U	10.0 U	10.0 U
Silver	0	182 nc	5. 0 U	10.0 U	5.0 U	5.0 U	5.0 U
Sodium	21300	-- [n]	966 (B)	1350	1470	2970	2410
Vanadium	0.0	36 nc	5. 0 U	31.9	9.9	4.3 (B)	5.0 U
Zinc	42	10950 nc	32.4	228	251	214	33.0
Total Organic Carbon							
Organic Carbon							

PRGs - Preliminary Remediation Goals
nc - non-cancer basis, value is 1/10 the published PRG
ca - cancer basis
pbk - based on PBK modeling
[n] - nutrient
U - analyte not detected
NT - not tested
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background & PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG & Background, then the value is presented with a normal style
-- - no background/PRG value is available for this analyte

INORGANIC FLAGS/QUALIFIERS
B - result is < CRDL/RL, but ≥ IDL/MDL
H - MB, EB1, EB2, EB3: Batch QC is > RL or had a negative instrument reading lower than the absolute value of the RL

ORGANIC FLAGS/QUALIFIERS
J - Result is an estimated value below the RL or a TIC
a - Concentration is below the method RL



Table 4-10
Summary of Surface Water Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (ug/l)	Units	Qualifier	Surface Water Background Criteria	Surface Water Region 9 Criteria	SW PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SW-002-0001-SW	12/05/03	0	Arsenic	4.3	µg/L		3.2	0.045	ca	YES	YES
LL9SW-002-0001-SW	12/05/03	0	Barium	60	µg/L		47.5	2555	nc	YES	NO
LL9SW-002-0001-SW	12/05/03	0	Beryllium	0.26	µg/L	B	0	73	nc	YES	NO
LL9SW-002-0001-SW	12/05/03	0	Chromium	3.3	µg/L	B	0	109	nc	YES	NO
LL9SW-002-0001-SW	12/05/03	0	Cobalt	4.6	µg/L	B	0	730	nc	YES	NO
LL9SW-002-0001-SW	12/05/03	0	Iron	20000	µg/L		2560	10950	nc	YES	YES
LL9SW-002-0001-SW	12/05/03	0	Lead	8.9	µg/L		0	15	mcl	YES	NO
LL9SW-002-0001-SW	12/05/03	0	Manganese	1200	µg/L		391	876	nc	YES	YES
LL9SW-002-0001-SW	12/05/03	0	Mercury	0.065	µg/L	B	0	11	nc	YES	NO
LL9SW-002-0001-SW	12/05/03	0	Nickel	11	µg/L		0	730	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Aluminum	12000	µg/L		3370	36499	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Arsenic	9.9	µg/L		3.2	0.045	ca	YES	YES
LL9SW-003-0001-SW	12/05/03	0	Barium	130	µg/L		47.5	2555	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Beryllium	0.3	µg/L	B	0	73	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Calcium	61000	µg/L		41400	--[n]	0	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Chromium	14	µg/L		0	109	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Cobalt	8	µg/L		0	730	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Copper	20	µg/L		7.9	1460	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Iron	17000	µg/L		2560	10950	nc	YES	YES
LL9SW-003-0001-SW	12/05/03	0	Lead	70	µg/L		0	15	mcl	YES	YES
LL9SW-003-0001-SW	12/05/03	0	Manganese	1400	µg/L		391	876	nc	YES	YES
LL9SW-003-0001-SW	12/05/03	0	Mercury	0.39	µg/L		0	11	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Nickel	15	µg/L		0	730	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Potassium	5200	µg/L		3170	--[n]	0	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Vanadium	19	µg/L		0	36	nc	YES	NO
LL9SW-003-0001-SW	12/05/03	0	Zinc	190	µg/L		42	10950	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Arsenic	1.9	µg/L	B	3.2	0.045	ca	NO	YES
LL9SW-004-0001-SW	12/05/03	0	Chromium	3.8	µg/L	B	0	109	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Cobalt	1.2	µg/L	B	0	730	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Iron	3200	µg/L		2560	10950	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Lead	8.7	µg/L		0	15	mcl	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Manganese	470	µg/L		391	876	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Mercury	0.12	µg/L	B	0	11	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Nickel	4	µg/L	B	0	730	nc	YES	NO
LL9SW-004-0001-SW	12/05/03	0	Vanadium	4.9	µg/L	B	0	36	nc	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Aluminum	17800	µg/L		3370	36499	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Arsenic	8	µg/L		3.2	0.045	CA	YES	YES
LL9SW-005-0001-SW	3/13/2002	0	Barium	145	µg/L		47.5	2555	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Beryllium	0.7	µg/L	B	0	73	NC	YES	NO



Table 4-10
Summary of Surface Water Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (ug/l)	Units	Qualifier	Surface Water Background Criteria	Surface Water Region 9 Criteria	SW PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SW-005-0001-SW	3/13/2002	0	Cadmium	1.1	µg/L	B	0	18	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Chromium	20.2	µg/L		0	109	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Cobalt	6	µg/L		0	730	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Copper	18.6	µg/L		7.9	1460	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Iron	18900	µg/L		2560	10950	NC	YES	YES
LL9SW-005-0001-SW	3/13/2002	0	Lead	62.9	µg/L		0	15	MCL	YES	YES
LL9SW-005-0001-SW	3/13/2002	0	Manganese	633	µg/L		391	876	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Mercury	0.37	µg/L		0	11	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Nickel	20.1	µg/L		0	730	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Potassium	3900	µg/L		3170	--	[n]	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Vanadium	31.9	µg/L		0	36	NC	YES	NO
LL9SW-005-0001-SW	3/13/2002	0	Zinc	228	µg/L		42	10950	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Arsenic	351	µg/L		3.2	0.045	CA	YES	YES
LL9SW-007-0001-SW	3/12/2002	0	Barium	208	µg/L		47.5	2555	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Cadmium	1.8	µg/L	B	0	18	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Calcium	46200	µg/L	H	41400	--	[n]	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Chromium	48.8	µg/L		0	109	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Cobalt	5.9	µg/L		0	730	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Copper	193	µg/L		7.9	1460	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Iron	17900	µg/L		2560	10950	NC	YES	YES
LL9SW-007-0001-SW	3/12/2002	0	Lead	2000	µg/L		0	15	MCL	YES	YES
LL9SW-007-0001-SW	3/12/2002	0	Manganese	1740	µg/L		391	876	NC	YES	YES
LL9SW-007-0001-SW	3/12/2002	0	Mercury	21.9	µg/L		0	11	NC	YES	YES
LL9SW-007-0001-SW	3/12/2002	0	Nickel	58.2	µg/L		0	730	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Potassium	8320	µg/L		3170	--	[n]	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Vanadium	9.9	µg/L		0	36	NC	YES	NO
LL9SW-007-0001-SW	3/12/2002	0	Zinc	251	µg/L		42	10950	NC	YES	NO
LL9SW-008-0001-FD	3/13/2002	0	4-Nitrtoluene	1.1	µg/L		0	0.66	CA	YES	YES
LL9SW-008-0001-FD	3/13/2002	0	Barium	93.9	µg/L		47.5	2555	NC	YES	NO
LL9SW-008-0001-FD	3/13/2002	0	Calcium	62700	µg/L	H	41400	--	[n]	YES	NO
LL9SW-008-0001-FD	3/13/2002	0	Iron	3380	µg/L		2560	10950	NC	YES	NO
LL9SW-008-0001-FD	3/13/2002	0	Lead	432	µg/L		0	15	MCL	YES	YES
LL9SW-008-0001-FD	3/13/2002	0	Mercury	0.61	µg/L		0	11	NC	YES	NO
LL9SW-008-0001-FD	3/13/2002	0	Nickel	10.9	µg/L		0	730	NC	YES	NO
LL9SW-008-0001-FD	3/13/2002	0	Potassium	12700	µg/L		3170	--	[n]	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Antimony	3.2	µg/L		0	15	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Arsenic	3.3	µg/L		3.2	0.045	CA	YES	YES
LL9SW-008-0001-SW	3/13/2002	0	Barium	211	µg/L		47.5	2555	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Cadmium	1	µg/L	B	0	18	NC	YES	NO



Table 4-10
Summary of Surface Water Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (ug/l)	Units	Qualifier	Surface Water Background Criteria	Surface Water Region 9 Criteria	SW PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SW-008-0001-SW	3/13/2002	0	Calcium	75200	µg/L	H	41400	--	[n]	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Chromium	27.1	µg/L		0	109	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Cobalt	6	µg/L		0	730	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Copper	31	µg/L		7.9	1460	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Iron	16100	µg/L		2560	10950	NC	YES	YES
LL9SW-008-0001-SW	3/13/2002	0	Lead	3150	µg/L		0	15	MCL	YES	YES
LL9SW-008-0001-SW	3/13/2002	0	Magnesium	12600	µg/L		10800	--	[n]	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Manganese	802	µg/L		391	876	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Mercury	7.7	µg/L		0	11	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Nickel	86.7	µg/L		0	730	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Potassium	14700	µg/L		3170	--	[n]	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Vanadium	4.3	µg/L	B	0	36	NC	YES	NO
LL9SW-008-0001-SW	3/13/2002	0	Zinc	214	µg/L		42	10950	NC	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Arsenic	2.2	µg/L		3.2	0.045	ca	NO	YES
LL9SW-009-0001-SW	12/09/03	0	Barium	51	µg/L		47.5	2555	nc	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Calcium	82000	µg/L		41400	--[n]	0	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Chromium	3.7	µg/L	B	0	109	nc	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Iron	4900	µg/L		2560	10950	nc	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Lead	36	µg/L		0	15	mcl	YES	YES
LL9SW-009-0001-SW	12/09/03	0	Mercury	0.33	µg/L		0	11	nc	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Nickel	4.5	µg/L	B	0	730	nc	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Potassium	4400	µg/L		3170	--[n]	0	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Vanadium	3	µg/L	B	0	36	nc	YES	NO
LL9SW-009-0001-SW	12/09/03	0	Zinc	55	µg/L		42	10950	nc	YES	NO
LL9SW-010-0001-SW	12/09/03	0	Arsenic	0.82	µg/L	B	3.2	0.045	ca	NO	YES
LL9SW-010-0001-SW	12/09/03	0	Chromium	2.3	µg/L	B	0	109	nc	YES	NO
LL9SW-010-0001-SW	12/09/03	0	Lead	46	µg/L		0	15	mcl	YES	YES
LL9SW-010-0001-SW	12/09/03	0	Mercury	2.2	µg/L		0	11	nc	YES	NO
LL9SW-010-0001-SW	12/09/03	0	Nickel	6.9	µg/L	B	0	730	nc	YES	NO
LL9SW-012-0001-DUP	12/09/03	0	Lead	1.2	µg/L	B	0	15	mcl	YES	NO
LL9SW-012-0001-DUP	12/09/03	0	Mercury	1.7	µg/L		0	11	nc	YES	NO
LL9SW-012-0001-DUP	12/09/03	0	Nickel	4.8	µg/L	B	0	730	nc	YES	NO
LL9SW-012-0001-DUP	12/09/03	0	Nitrocellulose	0.0001	µg/L	B J	0	--	0	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Lead	1.4	µg/L	B	0	15	mcl	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Mercury	0.3	µg/L		0	11	nc	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Nickel	5.3	µg/L	B	0	730	nc	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Nitrocellulose	0.00012	µg/L	B J	0	--	0	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Nitroglycerin	4.9	µg/L		0	--	0	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Nitroglycerin	5.02	µg/L		0	--	0	YES	NO



Table 4-10
Summary of Surface Water Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (ug/l)	Units	Qualifier	Surface Water Background Criteria	Surface Water Region 9 Criteria	SW PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SW-012-0001-SW	12/09/03	0	Nitroguanidine	249	µg/L		0	3650	nc	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Nitroguanidine	271	µg/L		0	3650	nc	YES	NO
LL9SW-012-0001-SW	12/09/03	0	Thallium	1.97	µg/L	B	0	2.4	nc	YES	NO

Notes:

-- - no background/PRG value is available for this analyte
PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL)
nc - non-cancer basis
ca - cancer basis
mcl - based on CWA maximum contaminant level
max - ceiling limit
[n] - nutrient

Inorganic Qualifiers

J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL

InOrganic Flags

^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL

Organic Qualifiers

J - result is an estimated value below the RL

Organic Flags

B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-11
Summary of Surface Water Results

							LL9SW-002-0001-SW		LL9SW-003-0001-SW		LL9SW-004-0001-SW		LL9SW-009-0001-SW		LL9SW-010-0001-SW		LL9SW-012-0001-DUP		LL9SW-012-0001-SW	
							12/05/03		12/05/03		12/05/03		12/09/03		12/09/03		12/09/03		12/09/03	
							Surface		Surface		Surface		Surface		Surface		Surface		Surface	
Group	Method	Analyte	Units	RVAAP Surface Water Background Criteria	Region 9 (Tap Water) PRGs	SW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	µg/L	0.00	730	nc	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U	4.4	U
Explosives 8330	8330	1,3,5-Trinitrobenzene	µg/L	0	1095	nc	0.11	U	0.087	U	0.087	U	0.08	U	0.08	U	0.11	U	0.12	U
		1,3-Dinitrobenzene	µg/L	0	3.6	nc	0.072	U	0.057	U	0.057	U	0.053	U	0.053	U	0.07	U	0.081	U
		2,4,6-TNT	µg/L	0	2.2	ca	0.092	U	0.074	U	0.074	U	0.068	U	0.068	U	0.09	U	0.1	U
		2,4-Dinitrotoluene	µg/L	0	73	nc	0.057	U	0.046	U	0.046	U	0.042	U	0.042	U	0.055	U	0.064	U
		2,6-Dinitrotoluene	µg/L	0	36	nc	0.28	U	0.22	U	0.22	U	0.21	U	0.21	U	0.27	U	0.32	U
		2-Nitrotoluene	µg/L	0	0.049	ca	0.22	U	0.18	U	0.18	U	0.16	U	0.16	U	0.21	U	0.25	U
		3-Nitrotoluene	µg/L	0	122	nc	0.14	U	0.11	U	0.11	U	0.1	U	0.1	U	0.13	U	0.16	U
		4-Nitrotoluene	µg/L	0	0.66	ca	0.45	U	0.37	U	0.37	U	0.34	U	0.34	U	0.44	U	0.52	U
		HMX	µg/L	0	1825	nc	0.3	U	0.24	U	0.24	U	0.22	U	0.22	U	0.3	U	0.35	U
		Nitrobenzene	µg/L	0	3.4	nc	0.12	U	0.1	U	0.1	U	0.092	U	0.092	U	0.12	U	0.14	U
		RDX	µg/L	0	0.61	ca	0.18	U	0.14	U	0.14	U	0.13	U	0.13	U	0.18	U	0.2	U
		Tetryl	µg/L	0	365	nc	0.29	U	0.24	U	0.24	U	0.22	U	0.22	U	0.29	U	0.33	U
PCB TCL 8082	8082	Aroclor 1016	µg/L	--	0.96	ca											0.17	U	0.17	U
		Aroclor 1221	µg/L	--	0.034	ca											0.45	U	0.45	U
		Aroclor 1232	µg/L	--	0.034	ca											0.21	U	0.22	U
		Aroclor 1242	µg/L	--	0.034	ca											0.18	U	0.19	U
		Aroclor 1248	µg/L	--	0.034	ca											0.2	U	0.21	U
		Aroclor 1254	µg/L	--	0.034	ca											0.13	U	0.13	U
		Aroclor 1260	µg/L	--	0.034	ca											0.15	U	0.15	U
Pesticides 8081A	8081A	4,4'-DDD	µg/L	--	0.28	ca											0.035	U	0.035	U
		4,4'-DDE	µg/L	--	0.20	ca											0.022	U	0.023	U
		4,4'-DDT	µg/L	--	0.20	ca											0.048	U	0.048	U
		Aldrin	µg/L	--	0.0040	ca											0.027	U	0.027	U
		alpha-BHC	µg/L	--	0.011	nc											0.046	U	0.046	U
		alpha-Chlordane	µg/L	--	0.19	ca											0.016	U	0.016	U
		beta-BHC	µg/L	--	0.037	ca											0.027	U	0.027	U
		delta-BHC	µg/L	--	--	0											0.024	U	0.025	U
		Dieldrin	µg/L	--	0.0042	ca											0.017	U	0.018	U
		Endosulfan I	µg/L	--	220	nc											0.02	U	0.021	U
		Endosulfan II	µg/L	--	220	nc											0.041	U	0.041	U
		Endosulfan sulfate	µg/L	--	220	nc											0.043	U	0.043	U
		Endrin	µg/L	--	11	nc											0.017	U	0.017	U
		Endrin aldehyde	µg/L	--	--	0											0.034	U	0.034	U
		Endrin ketone	µg/L	--	--	0											0.028	U	0.028	U
		gamma-BHC (Lindane)	µg/L	--	0.052	ca											0.041	U	0.041	U
		gamma-Chlordane	µg/L	--	0.19	ca											0.017	U	0.017	U
		Heptachlor	µg/L	--	0.015	ca											0.04	U	0.04	U
		Heptachlor epoxide	µg/L	--	0.0074	ca											0.035	U	0.035	U
		Methoxychlor	µg/L	--	182	nc											0.17	U	0.17	U
		Toxaphene	µg/L	--	0	0											0.14	U	0.14	U



Table 4-11
Summary of Surface Water Results

							LL9SW-002-0001-SW		LL9SW-003-0001-SW		LL9SW-004-0001-SW		LL9SW-009-0001-SW		LL9SW-010-0001-SW		LL9SW-012-0001-DUF		LL9SW-012-0001-SW	
							12/05/03		12/05/03		12/05/03		12/09/03		12/09/03		12/09/03		12/09/03	
							Surface		Surface		Surface		Surface		Surface		Surface		Surface	
Group	Method	Analyte	Units	RVAAP Surface Water Background Criteria	Region 9 (Tap Water) PRGs	SW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Propellants 8330	353.2	Nitrocellulose	µg/L	0	--	0											0.0001	B J	0.00012	B J
	8330	Nitroglycerin	µg/L	0	--	0											0.65	U	0.65	U
	UV/HPLC per SOP	Nitroguanidine	µg/L	0	3650	nc											20	U	20	U
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	µg/L	--	7.2	nc											0.33	U	0.33	U
		1,2-Dichlorobenzene	µg/L	--	370	nc											0.34	U	0.34	U
		1,3-Dichlorobenzene	µg/L	--	182	nc											0.41	U	0.42	U
		1,4-Dichlorobenzene	µg/L	--	0.50	ca											0.32	U	0.32	U
		2,2-oxybis (1-chloropropane)	µg/L	--	0.27	ca											0.27	U	0.27	U
		2,4,5-Trichlorophenol	µg/L	--	3650	nc											1.3	U	1.4	U
		2,4,6-Trichlorophenol	µg/L	--	3.6	nc											0.2	U	0.21	U
		2,4-Dichlorophenol	µg/L	--	109	nc											0.87	U	0.89	U
		2,4-Dimethylphenol	µg/L	--	730	nc											1.2	U	1.3	U
		2,4-Dinitrophenol	µg/L	--	73	nc											3.2	U	3.2	U
		2-Chloronaphthalene	µg/L	--	487	nc											0.25	U	0.25	U
		2-Chlorophenol	µg/L	--	30	nc											0.12	U	0.12	U
		2-Methylnaphthalene	µg/L	--	--	0											0.12	U	0.13	U
		2-Methylphenol (o-cresol)	µg/L	--	1825	nc											0.25	U	0.25	U
		2-Nitroaniline	µg/L	--	109	nc											0.21	U	0.22	U
		2-Nitrophenol	µg/L	--	--	0											0.79	U	0.8	U
		3,3-Dichlorobenzidine	µg/L	--	0.15	ca											0.69	U	0.71	U
		3-Nitroaniline	µg/L	--	3.2	ca											2	U	2.1	U
		4,6-Dinitro-2-methylphenol	µg/L	--	3.6	nc											2.3	U	2.4	U
		4-Bromophenyl phenyl ether	µg/L	--	--	0											0.18	U	0.19	U
		4-Chloro-3-methylphenol	µg/L	--	--	0											2.3	U	2.4	U
		4-Chloroaniline	µg/L	--	146	nc											2.7	U	2.7	U
		4-Chlorophenyl phenyl ether	µg/L	--	--	0											0.72	U	0.74	U
		4-Methylphenol (m/p-cresol)	µg/L	--	182	nc											0.096	U	0.098	U
		4-Nitroaniline	µg/L	--	3.2	ca											2.2	U	2.3	U
		4-Nitrophenol	µg/L	--	--	0											3.6	U	3.6	U
		Acenaphthene	µg/L	--	365	nc											0.12	U	0.12	U
		Acenaphthylene	µg/L	--	--	0											0.12	U	0.12	U
		Anthracene	µg/L	--	1825	nc											0.14	U	0.15	U
		Benzo(a)anthracene	µg/L	--	0.092	ca											0.047	U	0.048	U
		Benzo(a)pyrene	µg/L	--	0.0092	ca											0.081	U	0.082	U
		Benzo(b)fluoranthene	µg/L	--	0.092	ca											0.064	U	0.066	U
		Benzo(ghi)perylene	µg/L	--	--	0											0.18	U	0.19	U
		Benzo(k)fluoranthene	µg/L	--	0.92	ca											0.069	U	0.071	U
		Bis(2-chloroethoxy)methane	µg/L	--	--	0											0.3	U	0.3	U
		Bis(2-chloroethyl)ether	µg/L	--	0.010	ca											0.29	U	0.29	U
		Bis(2-ethylhexyl)phthalate	µg/L	--	4.8	ca											3.7	U	3.8	U
		Butyl benzyl phthalate	µg/L	--	7300	nc											0.37	U	0.38	U
		Carbazole	µg/L	--	3.4	ca											0.28	U	0.28	U
		Chrysene	µg/L	--	9.2	ca											0.043	U	0.044	U
		Dibenzo(a,h)anthracene	µg/L	--	0.0092	ca											0.12	U	0.13	U



Table 4-11
Summary of Surface Water Results

							LL9SW-002-0001-SW		LL9SW-003-0001-SW		LL9SW-004-0001-SW		LL9SW-009-0001-SW		LL9SW-010-0001-SW		LL9SW-012-0001-DUF		LL9SW-012-0001-SW	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Surface Water Background Criteria	Region 9 (Tap Water) PRGs	SW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		Dibenzofuran	µg/L	--	12	nc											0.12	U	0.13	U
		Diethyl phthalate	µg/L	--	29199	nc											0.14	U	0.15	U
		Dimethyl phthalate	µg/L	--	364867	nc											0.2	U	0.21	U
		Di-n-butyl phthalate	µg/L	--	3650	nc											0.35	U	0.35	U
		Di-n-octyl phthalate	µg/L	--	1460	nc											2.4	U	2.5	U
		Fluoranthene	µg/L	--	1460	nc											0.13	U	0.14	U
		Fluorene	µg/L	--	243	nc											0.12	U	0.13	U
		Hexachlorobenzene	µg/L	--	0.042	ca											0.093	U	0.095	U
		Hexachlorobutadiene	µg/L	--	0.86	ca											0.62	U	0.63	U
		Hexachlorocyclopentadiene	µg/L	--	219	nc											0.62	U	0.64	U
		Hexachloroethane	µg/L	--	4.8	ca											0.59	U	0.6	U
		Indeno(1,2,3-cd)pyrene	µg/L	--	0.092	ca											0.083	U	0.084	U
		Isophorone	µg/L	--	71	ca											0.25	U	0.25	U
		Naphthalene	µg/L	--	6.2	nc											0.15	U	0.16	U
		n-Nitroso-di-n-propylamine	µg/L	--	0.0096	ca											0.078	U	0.079	U
		n-Nitrosodiphenylamine	µg/L	--	14	ca											0.12	U	0.13	U
		Pentachlorophenol	µg/L	--	0.56	ca											1.6	U	1.7	U
		Phenanthrene	µg/L	--	--	0											0.13	U	0.14	U
		Phenol	µg/L	--	10950	nc											0.34	U	0.34	U
		Pyrene	µg/L	--	182	nc											0.12	U	0.12	U
TAL Metals 6010B	6010B	Aluminum	µg/L	3370	36499	nc	1800		12000		3200		1700		1200		440		430	
		Barium	µg/L	47.5	2555	nc	60		130		43		51		31		28		28	
		Beryllium	µg/L	0.00	73	nc	0.26	B	0.3	B	0.17	U	0.17	U	0.17	U	0.17	U	0.17	U
		Cadmium	µg/L	0.00	18	nc	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U	0.44	U
		Calcium	µg/L	41400	--	0	9900		61000		12000		82000		20000		18000		18000	
		Chromium	µg/L	0.00	109	nc	3.3	B	14		3.8	B	3.7	B	2.3	B	1.5	U	1.5	U
		Cobalt	µg/L	0.00	730	nc	4.6	B	8		1.2	B	1	U	1	U	1	U	1	U
		Copper	µg/L	7.9	1460	nc	5.8	B	20		4.4	B	7.3	B	6.6	B	1.6	U	1.6	U
		Iron	µg/L	2560	10950	nc	20000		17000		3200		4900		1900		440		540	
		Magnesium	µg/L	10800	--	0	3000		9600		3000		9600		5400		4300		4500	
		Manganese	µg/L	391	876	nc	1200		1400		470		330		160		110		120	
		Nickel	µg/L	0.00	730	nc	11		15		4	B	4.5	B	6.9	B	4.8	B	5.3	B
		Potassium	µg/L	3170	--	0	1700		5200		1400		4400		1700		1200		1200	
		Selenium	µg/L	0.00	182	nc	5	U	5	U	5	U	5	U	5	U	5	U	5	U
		Silver	µg/L	0.00	182	nc	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U	3.1	U
		Sodium	µg/L	21300	--	0	2000		4300		2000		6300		11000		2300		2400	
		Vanadium	µg/L	0.00	36	nc	2.1	U	19		4.9	B	3	B	2.1	U	2.1	U	2.1	U
		Zinc	µg/L	42	10950	nc	41		190		39		55		38		19		20	
	7041	Antimony	µg/L	0.00	15	nc	2.2	U	2.2	U	2.2	U	2.2	U	2.2	U	2.2	U	2.2	U
	7060A	Arsenic	µg/L	3.2	0.045	ca	4.3		9.9		1.9	B	2.2		0.82	B	0.73	U	0.73	U
	7421	Lead	µg/L	0.00	15	mcl	8.9		70		8.7		36		46		1.2	B	1.4	B
	7470A	Mercury	µg/L	0.00	11	nc	0.065	B	0.39		0.12	B	0.33		2.2		1.7		0.3	
	7841	Thallium	µg/L	0.00	2.4	nc	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U	1.6	U



Table 4-11
Summary of Surface Water Results

							LL9SW-002-0001-SW		LL9SW-003-0001-SW		LL9SW-004-0001-SW		LL9SW-009-0001-SW		LL9SW-010-0001-SW		LL9SW-012-0001-DUF		LL9SW-012-0001-SW	
							12/05/03		12/05/03		12/05/03		12/09/03		12/09/03		12/09/03		12/09/03	
							Surface		Surface		Surface		Surface		Surface		Surface		Surface	
Group	Method	Analyte	Units	RVAAP Surface Water Background Criteria	Region 9 (Tap Water) PRGs	SW PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	µg/L	--	3172	nc											0.08	U	0.08	U
		1,1,2,2-Tetrachloroethane	µg/L	--	0.055	ca											0.09	U	0.09	U
		1,1,2-Trichloroethane	µg/L	--	0.20	ca											0.15	U	0.15	U
		1,1-Dichloroethane	µg/L	--	811	nc											0.11	U	0.11	U
		1,1-Dichloroethene	µg/L	--	339	nc											0.12	U	0.12	U
		1,2-Dichloroethane	µg/L	--	0.12	ca											0.09	U	0.09	U
		1,2-Dichloroethene (total)	µg/L	--	120	nc											0.23	U	0.23	U
		1,2-Dichloropropane	µg/L	--	0.16	ca											0.12	U	0.12	U
		2-Butanone (MEK)	µg/L	--	6968	nc											1.2	U	1.2	U
		2-Hexanone	µg/L	--	2000	nc											0.53	U	0.53	U
		4-Methyl-2-pentanone (MIBK)	µg/L	--	1993	nc											0.65	U	0.65	U
		Acetone	µg/L	--	5475	nc											1.8	U	1.8	U
		Benzene	µg/L	--	0.35	ca											0.09	U	0.09	U
		Bromodichloromethane	µg/L	--	0.18	ca											0.11	U	0.11	U
		Bromoform	µg/L	--	8.5	ca											0.11	U	0.11	U
		Bromomethane	µg/L	--	8.7	nc											0.1	U	0.1	U
		Carbon disulfide	µg/L	--	1043	nc											0.2	U	0.2	U
		Carbon tetrachloride	µg/L	--	0.17	ca											0.13	U	0.13	U
		Chlorobenzene	µg/L	--	106	nc											0.08	U	0.08	U
		Chloroethane	µg/L	--	4.6	ca											0.08	U	0.08	U
		Chloroform	µg/L	--	0.17	ca											0.11	U	0.11	U
		Chloromethane	µg/L	--	158	nc											0.08	U	0.08	U
		cis-1,3-Dichloropropene	µg/L	--	0.40	ca											0.12	U	0.12	U
		Dibromochloromethane	µg/L	--	0.13	ca											0.06	U	0.06	U
		Ethylbenzene	µg/L	--	1340	nc											0.07	U	0.07	U
		Methylene chloride	µg/L	--	4.3	ca											0.35	U	0.35	U
		Styrene	µg/L	--	1641	nc											0.13	U	0.13	U
		Tetrachloroethene	µg/L	--	0.10	ca											0.09	U	0.09	U
		Toluene	µg/L	--	723	nc											0.1	U	0.1	U
		trans-1,3-Dichloropropene	µg/L	--	0.40	ca											0.15	U	0.15	U
		Trichloroethene	µg/L	--	0.028	ca											0.1	U	0.1	U
		Vinyl chloride	µg/L	--	0.020	ca											0.08	U	0.08	U
		Xylenes (total)	µg/L	--	0.020	ca											0.28	U	0.28	U

Notes:

-- - no background/PRG value is available for this analyte
blank cell indicates that the analysis was not performed
PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL))
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background and PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG and Background, then the value is presented with a normal style

Inorganic Qualifiers

U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL

InOrganic Flags

^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL

Organic Qualifiers

U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL

Organic Flags

B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-12
Summary of Sediment Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Sediment Background Criteria	Sediment Region 9 Criteria	SD PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Aluminum	10800		17700	7614.00	NC	NO	YES
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Arsenic	9.4		19.5	0.39	CA	NO	YES
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Iron	22700	H	28200	2346.00	NC	NO	YES
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Manganese	563		1950	176.00	NC	NO	YES
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Vanadium	22.1		26.1	7.80	NC	NO	YES
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	Arsenic	9.7		19.5	0.39	CA	NO	YES
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	Manganese	300		1950	176	NC	NO	YES
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Aluminum	15600		17700	7614.00	NC	NO	YES
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Arsenic	17.3		19.5	0.39	CA	NO	YES
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Manganese	300		1950	176.00	NC	NO	YES
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Arsenic	11		19.5	0.39	CA	NO	YES
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Iron	26000		28200	2346	NC	NO	YES
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Manganese	1200		1950	176	NC	NO	YES
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Vanadium	26		26.1	7.8	NC	NO	YES
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Aluminum	14500		17700	7614.00	NC	NO	YES
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Arsenic	14.4		19.5	0.39	CA	NO	YES
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Iron	26500	H	28200	2346.00	NC	NO	YES
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Manganese	395		1950	176.00	NC	NO	YES
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Aluminum	11000		13900	7614	NC	NO	YES
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Arsenic	10		19.5	0.39	CA	NO	YES
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Iron	18000		28200	2346	NC	NO	YES
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Manganese	250		1950	176	NC	NO	YES
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Vanadium	20		26.1	7.8	NC	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Arsenic	15		19.5	0.39	CA	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Benzo(a)anthracene	2.1		--	0.62	CA	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Benzo(a)pyrene	2.2		--	0.062	CA	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Benzo(b)fluoranthene	2.5		--	0.62	CA	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Dibenzo(a,h)anthracene	0.61		--	0.062	CA	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Indeno(1,2,3-cd)pyrene	1.1		--	0.62	CA	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Manganese	400		1950	176	NC	NO	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Vanadium	19		26.1	7.8	NC	NO	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Aluminum	9100		13900	7614	NC	NO	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Manganese	710		1950	176	NC	NO	YES
LL9SD-011-0001-SD	12/05/03	0-1 ft	Aluminum	8800		13900	7614	NC	NO	YES
LL9SD-011-0001-SD	12/05/03	0-1 ft	Arsenic	4.2		19.5	0.39	CA	NO	YES
LL9SD-011-0001-SD	12/05/03	0-1 ft	Iron	22000		28200	2346	NC	NO	YES
LL9SD-011-0001-SD	12/05/03	0-1 ft	Manganese	680		1950	176	NC	NO	YES
LL9SD-011-0001-SD	12/05/03	0-1 ft	Vanadium	15		26.1	7.8	NC	NO	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Aluminum	12000		13900	7614	NC	NO	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Arsenic	14		19.5	0.39	CA	NO	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Iron	24000		28200	2346	NC	NO	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Manganese	1300		1950	176	NC	NO	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Vanadium	24		26.1	7.8	NC	NO	YES
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Aluminum	12000		13900	7614	NC	NO	YES
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Arsenic	15		19.5	0.39	CA	NO	YES
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Iron	26000		28200	2346	NC	NO	YES



Table 4-12
Summary of Sediment Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Sediment Background Criteria	Sediment Region 9 Criteria	SD PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Manganese	200		1950	176	NC	NO	YES
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Vanadium	22		26.1	7.8	NC	NO	YES
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Aluminum	11000		13900	7614	NC	NO	YES
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Arsenic	10		19.5	0.39	CA	NO	YES
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Iron	19000		28200	2346	NC	NO	YES
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Manganese	880		1950	176	NC	NO	YES
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Vanadium	20		26.1	7.8	NC	NO	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Aluminum	12000		13900	7614	NC	NO	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Arsenic	10		19.5	0.39	CA	NO	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Iron	20000		28200	2346	NC	NO	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Manganese	670		1950	176	NC	NO	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Vanadium	23		26.1	7.8	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Aluminum	12000		13900	7614	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Aluminum	11506.62		13900	7614	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Arsenic	10		19.5	0.39	CA	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Arsenic	9.41		19.5	0.39	CA	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Iron	17000		28200	2346	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Iron	17116.59		28200	2346	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Manganese	610		1950	176	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Manganese	600.66		1950	176	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Vanadium	21		26.1	7.8	NC	NO	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Vanadium	21		26.1	7.8	NC	NO	YES
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Aluminum	13000		13900	7614	NC	NO	YES
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Arsenic	14		19.5	0.39	CA	NO	YES
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Iron	25000		28200	2346	NC	NO	YES
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Manganese	890		1950	176	NC	NO	YES
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Vanadium	23		26.1	7.8	NC	NO	YES
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Aluminum	11000		13900	7614	NC	NO	YES
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Arsenic	13		19.5	0.39	CA	NO	YES
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Iron	23000		28200	2346	NC	NO	YES
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Manganese	640		1950	176	NC	NO	YES
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Vanadium	20		26.1	7.8	NC	NO	YES
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Aluminum	12000		13900	7614	NC	NO	YES
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Arsenic	13		19.5	0.39	CA	NO	YES
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Iron	20000		28200	2346	NC	NO	YES
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Manganese	1400		1950	176	NC	NO	YES
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Vanadium	24		26.1	7.8	NC	NO	YES
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Antimony	0.49		0.00	3.10	NC	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Beryllium	0.99		0.38	15.00	NC	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Cadmium	0.47		0.0	3.70	NC	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Lead	47.2		27.4	400.00	PBK	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Maximum Particle Size	19			--		YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Mercury	0.15		0.06	2.30	NC	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Nickel	22.4		17.7	156.00	NC	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Sodium	136	B	112	--	[N]	YES	NO
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	Barium	140		123	538	NC	YES	NO



Table 4-12
Summary of Sediment Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Sediment Background Criteria	Sediment Region 9 Criteria	SD PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	Beryllium	0.77		0.38	15	NC	YES	NO
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.19	B	0	3.7	NC	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Beryllium	0.67		0.38	15.00	NC	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Chromium	20.6		18.1	30.00	CA	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Lead	37.8		27.4	400.00	PBK	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Magnesium	3450		2760	--	[N]	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Maximum Particle Size	9.5			--		YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Mercury	0.35		0.06	2.30	NC	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Nickel	21.2		17.7	156.00	NC	YES	NO
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Sodium	213		112	--	[N]	YES	NO
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Beryllium	0.76		0.38	15	NC	YES	NO
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.77		0	3.7	NC	YES	NO
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Lead	31		27.4	400	PBK	YES	NO
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Mercury	0.1		0.06	2.3	NC	YES	NO
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Nickel	18		17.7	156	NC	YES	NO
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Selenium	2.1		1.7	39	NC	YES	NO
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Beryllium	0.62		0.38	15.00	NC	YES	NO
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Chromium	18.9		18.1	30.00	CA	YES	NO
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Lead	34.1		27.4	400.00	PBK	YES	NO
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Maximum Particle Size	19			--		YES	NO
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Mercury	0.13		0.06	2.30	NC	YES	NO
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Sodium	204		112	--	[N]	YES	NO
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.2	B	0	3.7	NC	YES	NO
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Lead	36		27.4	400	PBK	YES	NO
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	Mercury	0.072		0.06	2.3	NC	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Beryllium	0.41	B	0.38	15	NC	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Cadmium	1		0	3.7	NC	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Copper	120	H	27.6	313	NC	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Lead	200		27.4	400	PBK	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Nickel	24		17.7	156	NC	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Nitrocellulose	3.1	J	0	--	0	YES	NO
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Nitroguanidine	0.053	J	0	611	NC	YES	NO
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Beryllium	0.48	B	0.38	15	NC	YES	NO
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Cadmium	1.2		0	3.7	NC	YES	NO
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Copper	34	H	27.6	313	NC	YES	NO
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Lead	300		27.4	400	PBK	YES	NO
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Nickel	26		17.7	156	NC	YES	NO
LL9SD-011-0001-SD	12/05/03	0-1 ft	Beryllium	0.51	B	0.38	15	NC	YES	NO
LL9SD-011-0001-SD	12/05/03	0-1 ft	Cadmium	0.4		0	3.7	NC	YES	NO
LL9SD-011-0001-SD	12/05/03	0-1 ft	Cobalt	9.4		9.1	30	CA	YES	NO
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Beryllium	0.86		0.38	15	NC	YES	NO
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Nickel	19		17.7	156	NC	YES	NO
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Beryllium	0.52	B	0.38	15	NC	YES	NO
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.5		0	3.7	NC	YES	NO
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Lead	100		27.4	400	PBK	YES	NO
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	Mercury	0.18		0.06	2.3	NC	YES	NO



Table 4-12
Summary of Sediment Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Sediment Background Criteria	Sediment Region 9 Criteria	SD PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Beryllium	0.63	B	0.38	15	NC	YES	NO
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.35	B	0	3.7	NC	YES	NO
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Lead	35		27.4	400	PBK	YES	NO
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Mercury	0.33		0.06	2.3	NC	YES	NO
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Nickel	18		17.7	156	NC	YES	NO
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	Selenium	1.8	B	1.7	39	NC	YES	NO
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Beryllium	0.59	B	0.38	15	NC	YES	NO
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Cadmium	0.71		0	3.7	NC	YES	NO
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Lead	77		27.4	400	PBK	YES	NO
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Mercury	0.13		0.06	2.3	NC	YES	NO
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Nickel	18		17.7	156	NC	YES	NO
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Nitrocellulose	4.2	J	0	--	0	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Beryllium	0.54	B	0.38	15	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Beryllium	0.55	B	0.38	15	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Cadmium	0.61		0	3.7	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Cadmium	0.64		0	3.7	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Lead	71		27.4	400	PBK	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Lead	75.88		27.4	400	PBK	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Mercury	0.12		0.06	2.3	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Mercury	0.11		0.06	2.3	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nickel	18		17.7	156	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitrocellulose	76	*	0	--	0	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitrocellulose	2.8	B J	0	--	0	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitrocellulose	59		0	--	0	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitroglycerin	4.86		0	--	0	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitroglycerin	4.74		0	--	0	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitroguanidine	0.844		0	611	NC	YES	NO
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitroguanidine	0.873		0	611	NC	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Beryllium	0.51	B	0.38	15	NC	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Cadmium	0.53		0	3.7	NC	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Calcium	11000		5510	--[n]	0	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Chromium	22		18.1	30	CA	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Cobalt	9.5		9.1	30	CA	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Lead	49		27.4	400	PBK	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Magnesium	3000		2760	--[n]	0	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Mercury	1.3		0.06	2.3	NC	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Nickel	21		17.7	156	NC	YES	NO
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	Selenium	1.9		1.7	39	NC	YES	NO
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Beryllium	0.47	B	0.38	15	NC	YES	NO
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.32		0	3.7	NC	YES	NO
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Calcium	6900		5510	--[n]	0	YES	NO
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Lead	32		27.4	400	PBK	YES	NO
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	Mercury	0.45		0.06	2.3	NC	YES	NO
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Beryllium	0.65		0.38	15	NC	YES	NO
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Cadmium	0.31	B	0	3.7	NC	YES	NO
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Cobalt	9.5		9.1	30	CA	YES	NO



Table 4-12
Summary of Sediment Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Sediment Background Criteria	Sediment Region 9 Criteria	SD PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Lead	29		27.4	400	PBK	YES	NO
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	Mercury	0.068		0.06	2.3	NC	YES	NO
LL9SD-001-0001-SD	3/13/2002	0-0.5 ft	Organic Carbon	23000					YES	YES
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	Iron	39000		28200	2346	NC	YES	YES
LL9SD-002-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	11000		0.89	0.52	NC	YES	YES
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Iron	31900	H	28200	2346.00	NC	YES	YES
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Organic Carbon	3700					YES	YES
LL9SD-003-0001-SD	3/13/2002	0-0.5 ft	Vanadium	26.2		26.1	7.80	NC	YES	YES
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	Aluminum	14000		13900	7614	NC	YES	YES
LL9SD-004-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	15000		0.89	0.52	NC	YES	YES
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Organic Carbon	5300					YES	YES
LL9SD-005-0001-SD	3/13/2002	0-0.5 ft	Vanadium	27.2		26.1	7.80	NC	YES	YES
LL9SD-006-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	24000		0.89	0.52	NC	YES	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Iron	31000		28200	2346	NC	YES	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Mercury	6		0.06	2.3	NC	YES	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	TOC Average Duplicates	4200		0.89	0.52	NC	YES	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Arsenic	25		19.5	0.39	CA	YES	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Iron	33000		28200	2346	NC	YES	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Mercury	110		0.06	2.3	NC	YES	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	TOC Average Duplicates	3700		0.89	0.52	NC	YES	YES
LL9SD-010-0001-SD	12/09/03	0-0.5 ft	Vanadium	38		26.1	7.8	NC	YES	YES
LL9SD-011-0001-SD	12/05/03	0-1 ft	TOC Average Duplicates	8700		0.89	0.52	NC	YES	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	Mercury	2.9		0.06	2.3	NC	YES	YES
LL9SD-012-0001-SD	12/09/03	0-0.5 ft	TOC Average Duplicates	3500		0.89	0.52	NC	YES	YES
LL9SD-013-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	28000		0.89	0.52	NC	YES	YES
LL9SD-014-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	24000		0.89	0.52	NC	YES	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	TOC Average Duplicates	12000		0.89	0.52	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Aluminum	NT	NA	17700	7614.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Antimony	NT	NA	0.00	3.10	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Arsenic	NT	NA	19.5	0.39	CA	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Barium	NT	NA	123.0	538.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Beryllium	NT	NA	0.38	15.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Cadmium	NT	NA	0.0	3.70	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Calcium	NT	NA	5500	--	[N]	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Chromium	NT	NA	18.1	30.00	CA	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Cobalt	NT	NA	9.1	902.00	CA	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Copper	NT	NA	27.6	313.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Iron	NT	NA	28200	2346.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Lead	NT	NA	27.4	400.00	PBK	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Magnesium	NT	NA	2760	--	[N]	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Manganese	NT	NA	1950	176.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Maximum Particle Size	NT	NA		--		YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Mercury	NT	NA	0.06	2.30	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Nickel	NT	NA	17.7	156.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Organic Carbon	NT	NA				YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Potassium	NT	NA	1950	--	[N]	YES	YES



Table 4-12
Summary of Sediment Exceedances

Sample Number	Sample Date	Sample Depth	Analyte	Result (mg/kg)	Qualifier	Sediment Background Criteria	Sediment Region 9 Criteria	SD PRG Basis	Exceed Installation Background?	Exceed Region 9 Criteria?
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Selenium	NT	NA	1.7	39.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Silver	NT	NA	0.0	39.00	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Sodium	NT	NA	112	--	[N]	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Thallium	NT	NA	0.9	0.52	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Vanadium	NT	NA	26.1	7.80	NC	YES	YES
LL9SD-004-0001-SD	3/13/2002	0-0.5 ft	Zinc	NT	NA	532.0	2346.00	NC	YES	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	TOC Average Duplicates	16000		0.89	0.52	NC	YES	YES
LL9SD-016-0001-DUP	12/05/03	0-0.5 ft	TOC Average Duplicates	15000		0.89	0.52	NC	YES	YES
LL9SD-016-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	11000		0.89	0.52	NC	YES	YES
LL9SD-017-0001-SD	12/05/03	0-0.5 ft	TOC Average Duplicates	26000		0.89	0.52	NC	YES	YES
LL9SD-009-0001-SD	12/09/03	0-0.5 ft	Nitroglycerin	ND		0	--	0	YES	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Nitroglycerin	ND		0	--	0	YES	YES
LL9SD-015-0001-DUP	12/09/03	0-0.5 ft	Nitroguanidine	ND		0	611	NC	YES	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitroglycerin	ND		0	--	0	YES	YES
LL9SD-015-0001-SD	12/09/03	0-0.5 ft	Nitroguanidine	ND		0	611	NC	YES	YES

Notes:

-- - no background/PRG value is available for this analyte
PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL)
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated

Inorganic Qualifiers

U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL

InOrganic Flags

^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL

Organic Qualifiers

U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL

Organic Flags

B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-13
Summary of Sediment Results

							LL9SD-002-0001-SD		LL9SD-004-0001-SD		LL9SD-006-0001-SD		LL9SD-009-0001-SD		LL9SD-010-0001-SD		LL9SD-011-0001-SD		LL9SD-012-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc							0.25	U						
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc							0.017	U						
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc							0.018	U						
		2,4,6-TNT	mg/Kg	0	16	ca							0.033	U						
		2,4-Dinitrotoluene	mg/Kg	0	12	nc							0.035	U						
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc							0.047	U						
		2-Nitrotoluene	mg/Kg	0	0.88	ca							0.033	U						
		3-Nitrotoluene	mg/Kg	0	73	nc							0.049	U						
		4-Nitrotoluene	mg/Kg	0	12	ca							0.046	U						
		HMX	mg/Kg	0	306	nc							0.11	U						
		Nitrobenzene	mg/Kg	0	2	nc							0.022	U						
		RDX	mg/Kg	0	4.4	ca							0.058	U						
		Tetryl	mg/Kg	0	61	nc							0.043	U						
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc							0.0024	U						
		Aroclor 1221	mg/Kg	--	0.22	ca							0.0026	U						
		Aroclor 1232	mg/Kg	--	0.22	ca							0.0026	U						
		Aroclor 1242	mg/Kg	--	0.22	ca							0.0026	U						
		Aroclor 1248	mg/Kg	--	0.22	ca							0.0026	U						
		Aroclor 1254	mg/Kg	--	0.22	ca							0.0026	U						
		Aroclor 1260	mg/Kg	--	0.22	ca							0.0018	U						
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca							0.0093	U						
		4,4'-DDE	mg/Kg	--	1.7	ca							0.017	U						
		4,4'-DDT	mg/Kg	--	1.7	ca							0.0095	U						
		Aldrin	mg/Kg	--	0.029	ca							0.0033	U						
		alpha-BHC	mg/Kg	--	0.09	sat							0.0041	U						
		alpha-Chlordane	mg/Kg	--	1.6	ca							0.0031	U						
		beta-BHC	mg/Kg	--	0.32	ca							0.0039	U						
		delta-BHC	mg/Kg	--	--	0							0.0028	U						
		Dieldrin	mg/Kg	--	0.030	ca							0.0087	U						
		Endosulfan I	mg/Kg	--	37	nc							0.0069	U						
		Endosulfan II	mg/Kg	--	37	nc							0.0072	U						
		Endosulfan sulfate	mg/Kg	--	37	nc							0.0075	U						
		Endrin	mg/Kg	--	1.8	nc							0.011	U						
		Endrin aldehyde	mg/Kg	--	--	0							0.0085	U						
		Endrin ketone	mg/Kg	--	--	0							0.0075	U						
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca							0.0059	U						
		gamma-Chlordane	mg/Kg	--	1.6	ca							0.0039	U						
		Heptachlor	mg/Kg	--	0.11	ca							0.0046	U						
		Heptachlor epoxide	mg/Kg	--	0.053	ca							0.0036	U						
		Methoxychlor	mg/Kg	--	31	nc							0.059	U						
		Toxaphene	mg/Kg	--	0	0							0.12	U						
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0							3.1	J						
	8330	Nitroglycerin	mg/Kg	0	--	0							0.5	U						
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc							0.053	J						
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc							0.076	U						
		1,2-Dichlorobenzene	mg/Kg	--	600	sat							0.1	U						
		1,3-Dichlorobenzene	mg/Kg	--	53	nc							0.1	U						
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca							0.091	U						
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca							0.096	U						
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc							0.049	U						



Table 4-13
Summary of Sediment Results

							LL9SD-013-0001-SD		LL9SD-014-0001-SD		LL9SD-015-0001-DU		LL9SD-015-0001-SD		LL9SD-016-0001-DU		LL9SD-016-0001-SD		LL9SD-017-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
Cyanide 9012A	9014/9010B	Cyanide, Total	mg/Kg	0.00	122	nc					0.27	U	0.3	U						
Explosives 8330	8330	1,3,5-Trinitrobenzene	mg/Kg	0	183	nc					0.017	U	0.017	U						
		1,3-Dinitrobenzene	mg/Kg	0	0.61	nc					0.017	U	0.018	U						
		2,4,6-TNT	mg/Kg	0	16	ca					0.033	U	0.033	U						
		2,4-Dinitrotoluene	mg/Kg	0	12	nc					0.035	U	0.035	U						
		2,6-Dinitrotoluene	mg/Kg	0	6.1	nc					0.047	U	0.047	U						
		2-Nitrotoluene	mg/Kg	0	0.88	ca					0.033	U	0.033	U						
		3-Nitrotoluene	mg/Kg	0	73	nc					0.049	U	0.049	U						
		4-Nitrotoluene	mg/Kg	0	12	ca					0.046	U	0.046	U						
		HMX	mg/Kg	0	306	nc					0.11	U	0.11	U						
		Nitrobenzene	mg/Kg	0	2	nc					0.022	U	0.022	U						
		RDX	mg/Kg	0	4.4	ca					0.057	U	0.058	U						
		Tetryl	mg/Kg	0	61	nc					0.043	U	0.043	U						
PCB TCL 8082	8082	Aroclor 1016	mg/Kg	--	0.39	nc					0.0035	U	0.0032	U						
		Aroclor 1221	mg/Kg	--	0.22	ca					0.0037	U	0.0034	U						
		Aroclor 1232	mg/Kg	--	0.22	ca					0.0037	U	0.0034	U						
		Aroclor 1242	mg/Kg	--	0.22	ca					0.0037	U	0.0034	U						
		Aroclor 1248	mg/Kg	--	0.22	ca					0.0037	U	0.0034	U						
		Aroclor 1254	mg/Kg	--	0.22	ca					0.0037	U	0.0034	U						
		Aroclor 1260	mg/Kg	--	0.22	ca					0.0026	U	0.0024	U						
Pesticides 8081A	8081A	4,4'-DDD	mg/Kg	--	2.4	ca					0.00066	U	0.00061	U						
		4,4'-DDE	mg/Kg	--	1.7	ca					0.0012	U	0.0011	U						
		4,4'-DDT	mg/Kg	--	1.7	ca					0.00068	U	0.00063	U						
		Aldrin	mg/Kg	--	0.029	ca					0.00024	U	0.00022	U						
		alpha-BHC	mg/Kg	--	0.09	sat					0.00029	U	0.00027	U						
		alpha-Chlordane	mg/Kg	--	1.6	ca					0.00022	U	0.0002	U						
		beta-BHC	mg/Kg	--	0.32	ca					0.00027	U	0.00026	U						
		delta-BHC	mg/Kg	--	--	0					0.0002	U	0.00019	U						
		Dieldrin	mg/Kg	--	0.030	ca					0.00062	U	0.00058	U						
		Endosulfan I	mg/Kg	--	37	nc					0.00049	U	0.00046	U						
		Endosulfan II	mg/Kg	--	37	nc					0.00051	U	0.00048	U						
		Endosulfan sulfate	mg/Kg	--	37	nc					0.00053	U	0.00049	U						
		Endrin	mg/Kg	--	1.8	nc					0.00079	U	0.00073	U						
		Endrin aldehyde	mg/Kg	--	--	0					0.0006	U	0.00056	U						
		Endrin ketone	mg/Kg	--	--	0					0.00053	U	0.00049	U						
		gamma-BHC (Lindane)	mg/Kg	--	0.44	ca					0.00042	U	0.00039	U						
		gamma-Chlordane	mg/Kg	--	1.6	ca					0.00027	U	0.00026	U						
		Heptachlor	mg/Kg	--	0.11	ca					0.00033	U	0.00031	U						
		Heptachlor epoxide	mg/Kg	--	0.053	ca					0.00026	U	0.00024	U						
		Methoxychlor	mg/Kg	--	31	nc					0.0042	U	0.0039	U						
		Toxaphene	mg/Kg	--	0	0					0.0084	U	0.0078	U						
Propellants 8330	353.2	Nitrocellulose	mg/Kg	0	--	0					4.2	J	2.8	B J						
	8330	Nitroglycerin	mg/Kg	0	--	0					0.5	U	0.5	U						
	UV/HPLC per SOP	Nitroguanidine	mg/Kg	0	611	nc					0.25	U	0.25	U						
SVOCs TCL 8270 C	8270C	1,2,4-Trichlorobenzene	mg/Kg	--	6.2	nc					0.11	U	0.099	U						
		1,2-Dichlorobenzene	mg/Kg	--	600	sat					0.15	U	0.13	U						
		1,3-Dichlorobenzene	mg/Kg	--	53	nc					0.15	U	0.13	U						
		1,4-Dichlorobenzene	mg/Kg	--	3.4	ca					0.13	U	0.12	U						
		2,2-oxybis (1-chloropropane)	mg/Kg	--	2.9	ca					0.14	U	0.13	U						
		2,4,5-Trichlorophenol	mg/Kg	--	611	nc					0.071	U	0.064	U						



Table 4-13
Summary of Sediment Results

							LL9SD-002-0001-SD		LL9SD-004-0001-SD		LL9SD-006-0001-SD		LL9SD-009-0001-SD		LL9SD-010-0001-SD		LL9SD-011-0001-SD		LL9SD-012-0001-SD	
Sample Date							12/05/03		12/05/03		12/05/03		12/09/03		12/09/03		12/05/03		12/09/03	
Sample Depth							0 - 0.5 ft		0 - 0.5 ft		0 - 0.5 ft		0 - 0.5 ft		0 - 0.5 ft		0 - 1 ft		0 - 0.5 ft	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc							0.06	U						
		2,4-Dichlorophenol	mg/Kg	--	18	nc							0.062	U						
		2,4-Dimethylphenol	mg/Kg	--	122	nc							0.077	U						
		2,4-Dinitrophenol	mg/Kg	--	12	nc							0.15	U						
		2-Chloronaphthalene	mg/Kg	--	494	nc							0.062	U						
		2-Chlorophenol	mg/Kg	--	6.3	nc							0.076	U						
		2-Methylnaphthalene	mg/Kg	--	--	0							0.011	Ja						
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc							0.011	U						
		2-Nitroaniline	mg/Kg	--	18.3	nc							0.044	U						
		2-Nitrophenol	mg/Kg	--	--	0							0.081	U						
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca							0.023	U						
		3-Nitroaniline	mg/Kg	--	1.8	nc							0.14	U						
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc							0.12	U						
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0							0.004	U						
		4-Chloro-3-methylphenol	mg/Kg	--	--	0							0.049	U						
		4-Chloroaniline	mg/Kg	--	24	nc							0.13	U						
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0							0.0046	U						
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc							0.0075	U						
		4-Nitroaniline	mg/Kg	--	23	ca							0.05	U						
		4-Nitrophenol	mg/Kg	--	--	0							0.11	U						
		Acenaphthene	mg/Kg	--	368	nc							0.075							
		Acenaphthylene	mg/Kg	--	--	0							0.05							
		Anthracene	mg/Kg	--	2189	nc							0.48							
		Benzo(a)anthracene	mg/Kg	--	0.62	ca							2.1							
		Benzo(a)pyrene	mg/Kg	--	0.062	ca							2.2							
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca							2.5							
		Benzo(ghi)perylene	mg/Kg	--	--	0							1.2							
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca							1.3	M						
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0							0.0037	U						
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca							0.0026	U						
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca							0.39							
		Butyl benzyl phthalate	mg/Kg	--	1222	nc							0.0053	U						
		Carbazole	mg/Kg	--	24	ca							0.36							
		Chrysene	mg/Kg	--	62	ca							2.4							
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca							0.61							
		Dibenzofuran	mg/Kg	--	15	nc							0.097							
		Diethyl phthalate	mg/Kg	--	4888	nc							0.0048	U						
		Dimethyl phthalate	mg/Kg	--	100000	max							0.0046	U						
		Di-n-butyl phthalate	mg/Kg	--	611	nc							0.032	Ja						
		Di-n-octyl phthalate	mg/Kg	--	244	nc							0.017	Ja						
		Fluoranthene	mg/Kg	--	229	nc							4.1							
		Fluorene	mg/Kg	--	275	nc							0.22							
		Hexachlorobenzene	mg/Kg	--	0.30	ca							0.0023	U						
		Hexachlorobutadiene	mg/Kg	--	6.2	ca							0.0042	U						
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc							0.069	U						
		Hexachloroethane	mg/Kg	--	35	ca							0.0042	U						
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca							1.1							
		Isophorone	mg/Kg	--	512	ca							0.0031	U						
		Naphthalene	mg/Kg	--	5.6	nc							0.016	Ja						
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca							0.003	U						



Table 4-13
Summary of Sediment Results

							LL9SD-013-0001-SD		LL9SD-014-0001-SD		LL9SD-015-0001-DU		LL9SD-015-0001-SD		LL9SD-016-0001-DU		LL9SD-016-0001-SD		LL9SD-017-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		2,4,6-Trichlorophenol	mg/Kg	--	0.61	nc					0.088	U	0.079	U						
		2,4-Dichlorophenol	mg/Kg	--	18	nc					0.09	U	0.081	U						
		2,4-Dimethylphenol	mg/Kg	--	122	nc					0.11	U	0.1	U						
		2,4-Dinitrophenol	mg/Kg	--	12	nc					0.21	U	0.19	U						
		2-Chloronaphthalene	mg/Kg	--	494	nc					0.09	U	0.081	U						
		2-Chlorophenol	mg/Kg	--	6.3	nc					0.11	U	0.099	U						
		2-Methylnaphthalene	mg/Kg	--	--	0					0.0028	U	0.004	Ja						
		2-Methylphenol (o-cresol)	mg/Kg	--	306	nc					0.016	U	0.014	U						
		2-Nitroaniline	mg/Kg	--	18.3	nc					0.063	U	0.057	U						
		2-Nitrophenol	mg/Kg	--	--	0					0.12	U	0.11	U						
		3,3-Dichlorobenzidine	mg/Kg	--	1.1	ca					0.034	U	0.03	U						
		3-Nitroaniline	mg/Kg	--	1.8	nc					0.21	U	0.19	U						
		4,6-Dinitro-2-methylphenol	mg/Kg	--	0.61	nc					0.18	U	0.16	U						
		4-Bromophenyl phenyl ether	mg/Kg	--	--	0					0.0058	U	0.0052	U						
		4-Chloro-3-methylphenol	mg/Kg	--	--	0					0.071	U	0.064	U						
		4-Chloroaniline	mg/Kg	--	24	nc					0.19	U	0.17	U						
		4-Chlorophenyl phenyl ether	mg/Kg	--	--	0					0.0067	U	0.0061	U						
		4-Methylphenol (m/p-cresol)	mg/Kg	--	31	nc					0.011	U	0.0098	U						
		4-Nitroaniline	mg/Kg	--	23	ca					0.073	U	0.066	U						
		4-Nitrophenol	mg/Kg	--	--	0					0.15	U	0.14	U						
		Acenaphthene	mg/Kg	--	368	nc					0.0026	U	0.0024	U						
		Acenaphthylene	mg/Kg	--	--	0					0.0017	U	0.0018	Ja						
		Anthracene	mg/Kg	--	2189	nc					0.0016	U	0.0031	Ja						
		Benzo(a)anthracene	mg/Kg	--	0.62	ca					0.016	Ja	0.02	Ja						
		Benzo(a)pyrene	mg/Kg	--	0.062	ca					0.021	Ja	0.025	Ja						
		Benzo(b)fluoranthene	mg/Kg	--	0.62	ca					0.02	Ja	0.024	Ja						
		Benzo(ghi)perylene	mg/Kg	--	--	0					0.013	Ja	0.018	Ja						
		Benzo(k)fluoranthene	mg/Kg	--	6.2	ca					0.019	Ja	0.029	Ja						
		Bis(2-chloroethoxy)methane	mg/Kg	--	--	0					0.0054	U	0.0049	U						
		Bis(2-chloroethyl)ether	mg/Kg	--	0.22	ca					0.0037	U	0.0034	U						
		Bis(2-ethylhexyl)phthalate	mg/Kg	--	35	ca					0.55		0.66							
		Butyl benzyl phthalate	mg/Kg	--	1222	nc					0.0077	U	0.0088	Ja						
		Carbazole	mg/Kg	--	24	ca					0.065	U	0.059	U						
		Chrysene	mg/Kg	--	62	ca					0.023	Ja	0.029	Ja						
		Dibenzo(a,h)anthracene	mg/Kg	--	0.062	ca					0.043	Ja	0.0037	U						
		Dibenzofuran	mg/Kg	--	15	nc					0.005	U	0.0046	U						
		Diethyl phthalate	mg/Kg	--	4888	nc					0.0069	U	0.0062	U						
		Dimethyl phthalate	mg/Kg	--	100000	max					0.0067	U	0.0061	U						
		Di-n-butyl phthalate	mg/Kg	--	611	nc					0.037	U	0.038	Ja						
		Di-n-octyl phthalate	mg/Kg	--	244	nc					0.016	U	0.015	U						
		Fluoranthene	mg/Kg	--	229	nc					0.037	Ja	0.048	Ja						
		Fluorene	mg/Kg	--	275	nc					0.003	U	0.0027	U						
		Hexachlorobenzene	mg/Kg	--	0.30	ca					0.0034	U	0.003	U						
		Hexachlorobutadiene	mg/Kg	--	6.2	ca					0.0062	U	0.0056	U						
		Hexachlorocyclopentadiene	mg/Kg	--	37	nc					0.1	U	0.091	U						
		Hexachloroethane	mg/Kg	--	35	ca					0.0062	U	0.0056	U						
		Indeno(1,2,3-cd)pyrene	mg/Kg	--	0.62	ca					0.037	Ja	0.039	Ja						
		Isophorone	mg/Kg	--	512	ca					0.0045	U	0.004	U						
		Naphthalene	mg/Kg	--	5.6	nc					0.0032	U	0.0029	U						
		n-Nitroso-di-n-propylamine	mg/Kg	--	0.069	ca					0.0043	U	0.0039	U						



Table 4-13
Summary of Sediment Results

							LL9SD-002-0001-SD		LL9SD-004-0001-SD		LL9SD-006-0001-SD		LL9SD-009-0001-SD		LL9SD-010-0001-SD		LL9SD-011-0001-SD		LL9SD-012-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca							0.0037	U						
		Pentachlorophenol	mg/Kg	--	3.0	ca							0.13	U						
		Phenanthrene	mg/Kg	--	--	0							2.1							
		Phenol	mg/Kg	--	1833	nc							0.0021	U						
		Pyrene	mg/Kg	--	232	nc							3.7							
TAL Metals 6010B	6010B	Aluminum	mg/Kg	13900	7614	nc	5500		14000		11000		7200		9100		8800		12000	
		Barium	mg/Kg	123	538	nc	140		89		64		63		98		58		100	
		Beryllium	mg/Kg	0.38	15	nc	0.77		0.76		0.31	B	0.41	B	0.48	B	0.51	B	0.86	
		Cadmium	mg/Kg	0.00	3.7	nc	0.19	B	0.77		0.2	B	1		1.2		0.4		0.1	U
		Calcium	mg/Kg	5510	--	0	870		3000		1700		1500		3100		1100		1200	
		Chromium	mg/Kg	18.1	30	ca	6.3		17		14		14		18		12		14	
		Cobalt	mg/Kg	9.1	30	ca	6.2		9.1		5.2		7.6		9.1		9.4		7.1	
		Copper	mg/Kg	27.6	313	nc	13		16		11		120	H	34	H	14		7.6	
		Iron	mg/Kg	28200	2346	nc	39000		26000		18000		31000		33000		22000		24000	
		Magnesium	mg/Kg	2760	--	0	620		2400		1800		2000		2300		2300		1800	
		Manganese	mg/Kg	1950	176	nc	300		1200		250		400		710		680		1300	
		Nickel	mg/Kg	17.7	156	nc	16		18		11		24		26		16		19	
		Potassium	mg/Kg	1950	--	0	350		1100		860		920		1200		830		1000	
		Selenium	mg/Kg	1.7	39	nc	1.4	B	2.1		0.7	B	0.49	U	0.51	B	0.86	B	0.82	B
		Silver	mg/Kg	0.00	39	nc	0.59	U	0.5	U	0.47	U	0.38	U	0.38	U	0.44	U	0.39	U
		Sodium	mg/Kg	112	--	0	170	U	140	U	130	U	110	U	110	U	120	U	110	U
		Vanadium	mg/Kg	26.1	7.8	nc	7.6		26		20		19		38		15		24	
		Zinc	mg/Kg	532	2346	nc	140		420		70		220		120		210		69	
	7041	Antimony	mg/Kg	0.00	3.1	nc	0.81	U	0.8	U	0.69	U	0.55	U	0.56	U	0.62	U	0.57	U
	7060A	Arsenic	mg/Kg	19.5	0.39	ca	9.7		11		10		15		25		4.2		14	
	7421	Lead	mg/Kg	27.4	400	pbk	24		31		36		200		300		15		17	
	7471A	Mercury	mg/Kg	0.06	2.3	nc	0.044		0.1		0.072		6		110		0.026		2.9	
	7841	Thallium	mg/Kg	0.89	0.52	nc	0.35	U	0.35	U	0.3	U	0.24	U	0.24	U	0.27	U	0.31	
	Lloyd Kahn	TOC Average Duplicates	mg/Kg	0.89	0.52	nc	11000		15000		24000		4200		3700		8700		3500	
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat							0.0014	U						
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca							0.0012	U						
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca							0.0014	U						
		1,1-Dichloroethane	mg/Kg	--	51	nc							0.0013	U						
		1,1-Dichloroethene	mg/Kg	--	12	nc							0.0017	U						
		1,2-Dichloroethane	mg/Kg	--	0.28	ca							0.0012	U						
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc							0.0027	U						
		1,2-Dichloropropane	mg/Kg	--	0.34	ca							0.0013	U						
		2-Butanone (MEK)	mg/Kg	--	2231	nc							0.005	U						
		2-Hexanone	mg/Kg	--	530	nc							0.0014	U						
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc							0.0013	U						
		Acetone	mg/Kg	--	1412	nc							0.0059	U						
		Benzene	mg/Kg	--	0.64	ca							0.0014	U						
		Bromodichloromethane	mg/Kg	--	0.82	ca							0.0012	U						
		Bromoform	mg/Kg	--	62	ca							0.00097	U						
		Bromomethane	mg/Kg	--	0.39	nc							0.0017	U						
		Carbon disulfide	mg/Kg	--	36	nc							0.0016	U						
		Carbon tetrachloride	mg/Kg	--	0.25	ca							0.0014	U						
		Chlorobenzene	mg/Kg	--	15	nc							0.0014	U						
		Chloroethane	mg/Kg	--	3.0	ca							0.0013	U						
		Chloroform	mg/Kg	--	0.22	ca							0.0014	U						
		Chloromethane	mg/Kg	--	4.7	nc							0.0014	U						



Table 4-13
Summary of Sediment Results

							LL9SD-013-0001-SD		LL9SD-014-0001-SD		LL9SD-015-0001-DU		LL9SD-015-0001-SD		LL9SD-016-0001-DU		LL9SD-016-0001-SD		LL9SD-017-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		n-Nitrosodiphenylamine	mg/Kg	--	99	ca					0.0054	U	0.0049	U						
		Pentachlorophenol	mg/Kg	--	3.0	ca					0.19	U	0.17	U						
		Phenanthrene	mg/Kg	--	--	0					0.02	Ja	0.02	Ja						
		Phenol	mg/Kg	--	1833	nc					0.003	U	0.0027	U						
		Pyrene	mg/Kg	--	232	nc					0.028	Ja	0.034	Ja						
TAL Metals 6010B	6010B	Aluminum	mg/Kg	13900	7614	nc	12000		11000		12000		12000		13000		11000		12000	
		Barium	mg/Kg	123	538	nc	65		96		92		89		86		73		100	
		Beryllium	mg/Kg	0.38	15	nc	0.52	B	0.63	B	0.59	B	0.54	B	0.51	B	0.47	B	0.65	
		Cadmium	mg/Kg	0.00	3.7	nc	0.5		0.35	B	0.71		0.61		0.53		0.32		0.31	B
		Calcium	mg/Kg	5510	--	0	2100		2700		1900		1800		11000		6900		1700	
		Chromium	mg/Kg	18.1	30	ca	18		15		16		15		22		16		15	
		Cobalt	mg/Kg	9.1	30	ca	7.2		7		7.9		7.4		9.5		8.2		9.5	
		Copper	mg/Kg	27.6	313	nc	18		16		14		13		22		18		12	
		Iron	mg/Kg	28200	2346	nc	26000		19000		20000		17000		25000		23000		20000	
		Magnesium	mg/Kg	2760	--	0	2300		2200		1900		1800		3000		2700		2000	
		Manganese	mg/Kg	1950	176	nc	200		880		670		610		890		640		1400	
		Nickel	mg/Kg	17.7	156	nc	17		18		18		18		21		17		16	
		Potassium	mg/Kg	1950	--	0	1000		1100		950		900		1300		1100		830	
		Selenium	mg/Kg	1.7	39	nc	1.6		1.8	B	1.2	B	0.82	B	1.9		1.4	B	1.4	B
		Silver	mg/Kg	0.00	39	nc	0.51	U	0.74	U	0.56	U	0.49	U	0.57	U	0.49	U	0.49	U
		Sodium	mg/Kg	112	--	0	140	U	210	U	160	U	140	U	160	U	140	U	140	U
		Vanadium	mg/Kg	26.1	7.8	nc	22		20		23		21		23		20		24	
		Zinc	mg/Kg	532	2346	nc	130		85		160		140		120		97		110	
	7041	Antimony	mg/Kg	0.00	3.1	nc	0.88	U	0.99	U	0.78	U	0.69	U	0.85	U	0.69	U	0.74	U
	7060A	Arsenic	mg/Kg	19.5	0.39	ca	15		10		10		10		14		13		13	
	7421	Lead	mg/Kg	27.4	400	pbk	100		35		77		71		49		32		29	
	7471A	Mercury	mg/Kg	0.06	2.3	nc	0.18		0.33		0.13		0.12		1.3		0.45		0.068	
	7841	Thallium	mg/Kg	0.89	0.52	nc	0.38	U	0.43	U	0.34	U	0.3	U	0.37	U	0.3	U	0.32	U
	Lloyd Kahn	TOC Average Duplicates	mg/Kg	0.89	0.52	nc	28000		24000		12000		16000		15000		11000		26000	
VOCs TCL 8260B	8260B	1,1,1-Trichloroethane	mg/Kg	--	1200	sat					0.0021	U	0.0019	U						
		1,1,2,2-Tetrachloroethane	mg/Kg	--	0.41	ca					0.0018	U	0.0016	U						
		1,1,2-Trichloroethane	mg/Kg	--	0.73	ca					0.0021	U	0.0019	U						
		1,1-Dichloroethane	mg/Kg	--	51	nc					0.0019	U	0.0017	U						
		1,1-Dichloroethene	mg/Kg	--	12	nc					0.0025	U	0.0022	U						
		1,2-Dichloroethane	mg/Kg	--	0.28	ca					0.0018	U	0.0016	U						
		1,2-Dichloroethene (total)	mg/Kg	--	6.9	nc					0.004	U	0.0036	U						
		1,2-Dichloropropane	mg/Kg	--	0.34	ca					0.0019	U	0.0017	U						
		2-Butanone (MEK)	mg/Kg	--	2231	nc					0.0075	U	0.0067	U						
		2-Hexanone	mg/Kg	--	530	nc					0.0021	U	0.0019	U						
		4-Methyl-2-pentanone (MIBK)	mg/Kg	--	528	nc					0.0019	U	0.0017	U						
		Acetone	mg/Kg	--	1412	nc					0.0088	U	0.0078	U						
		Benzene	mg/Kg	--	0.64	ca					0.0021	U	0.0019	U						
		Bromodichloromethane	mg/Kg	--	0.82	ca					0.0018	U	0.0016	U						
		Bromoform	mg/Kg	--	62	ca					0.0014	U	0.0013	U						
		Bromomethane	mg/Kg	--	0.39	nc					0.0025	U	0.0022	U						
		Carbon disulfide	mg/Kg	--	36	nc					0.0023	U	0.002	U						
		Carbon tetrachloride	mg/Kg	--	0.25	ca					0.0021	U	0.0019	U						
		Chlorobenzene	mg/Kg	--	15	nc					0.0021	U	0.0019	U						
		Chloroethane	mg/Kg	--	3.0	ca					0.0019	U	0.0017	U						
		Chloroform	mg/Kg	--	0.22	ca					0.0021	U	0.0019	U						
		Chloromethane	mg/Kg	--	4.7	nc					0.0021	U	0.0019	U						



Table 4-13
Summary of Sediment Results

							LL9SD-002-0001-SD		LL9SD-004-0001-SD		LL9SD-006-0001-SD		LL9SD-009-0001-SD		LL9SD-010-0001-SD		LL9SD-011-0001-SD		LL9SD-012-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca							0.0012	U						
		Dibromochloromethane	mg/Kg	--	1.1	ca							0.001	U						
		Ethylbenzene	mg/Kg	--	395	sat							0.0014	U						
		Methylene chloride	mg/Kg	--	9.1	ca							0.0037	U						
		Styrene	mg/Kg	--	1700	sat							0.0014	U						
		Tetrachloroethene	mg/Kg	--	0.48	ca							0.0016	U						
		Toluene	mg/Kg	--	520	sat							0.0014	U						
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca							0.001	U						
		Trichloroethene	mg/Kg	--	0.053	ca							0.0014	U						
		Vinyl chloride	mg/Kg	--	0.079	ca							0.0014	U						
		Xylenes (total)	mg/Kg	--	0.079	ca							0.0044	U						

Notes:
-- - no background/PRG value is available for this analyte
blank cell indicates that the analysis was not performed
PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL)
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background and PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG and Background, then the value is presented with a normal style

Inorganic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL
InOrganic Flags
^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL
Organic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL
Organic Flags
B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



Table 4-13
Summary of Sediment Results

							LL9SD-013-0001-SD		LL9SD-014-0001-SD		LL9SD-015-0001-DU		LL9SD-015-0001-SD		LL9SD-016-0001-DU		LL9SD-016-0001-SD		LL9SD-017-0001-SD	
							Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date		Sample Date	
							Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth		Sample Depth	
Group	Method	Analyte	Units	RVAAP Sediment Background Criteria	Region 9 (Residential Soil) PRGs	SD PRG Basis	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier	Result	Qualifier
		cis-1,3-Dichloropropene	mg/Kg	--	0.78	ca					0.0018	U	0.0016	U						
		Dibromochloromethane	mg/Kg	--	1.1	ca					0.0015	U	0.0013	U						
		Ethylbenzene	mg/Kg	--	395	sat					0.0021	U	0.0019	U						
		Methylene chloride	mg/Kg	--	9.1	ca					0.0056	U	0.0049	U						
		Styrene	mg/Kg	--	1700	sat					0.0021	U	0.0019	U						
		Tetrachloroethene	mg/Kg	--	0.48	ca					0.0023	U	0.002	U						
		Toluene	mg/Kg	--	520	sat					0.0021	U	0.0019	U						
		trans-1,3-Dichloropropene	mg/Kg	--	0.78	ca					0.0015	U	0.0013	U						
		Trichloroethene	mg/Kg	--	0.053	ca					0.0021	U	0.0019	U						
		Vinyl chloride	mg/Kg	--	0.079	ca					0.0021	U	0.0019	U						
		Xylenes (total)	mg/Kg	--	0.079	ca					0.0065	U	0.0058	U						

Notes:
-- - no background/PRG value is available for this analyte
blank cell indicates that the analysis was not performed
PRG - preliminary remediation goals (The PRG for lead is the Maximum Contaminant Level (MCL)
nc - non-cancer basis
ca - cancer basis
pbk - based on PBK modeling
mcl - based on CWA maximum contaminant level
max - ceiling limit
sat - soil saturation
[n] - nutrient
UC - unconsolidated
If Result = or > Background, then the value is presented with a shaded/highlighted style
If Result = or > Background and PRG, then the value is presented with a bold + shaded/highlighted style
If Result = or > PRG, then the value is presented with a bold style
If Result < PRG and Background, then the value is presented with a normal style

Inorganic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is less than the RL, but greater than or equal to the MDL
B - result is less than the CRDL/RL, but greater than or equal to the IDL/MDL
InOrganic Flags
^ - instrument related QC exceed the upper or lower control limits
* - batch QC exceeds the upper or lower control limit
H - batch QC is greater than RL
Organic Qualifiers
U - analyte was not detected at or above the stated limit
J - result is an estimated value below the RL
Organic Flags
B - Method Blank: batch QC is greater than RL
* - batch QC exceeds the upper or lower control limit
^ - batch QC is greater than RL
a - concentration is below the method Reporting Limit (RL)
H - alternate peak selection upon analytical review



5.0 CONTAMINANT FATE AND TRANSPORT

Knowledge about a contaminant's potential to migrate and persist in an environmental medium is important when evaluating its potential to elicit an adverse human health or ecological effect. This section contains information on properties and degradation potential of the chemicals detected at concentrations in excess of background levels, with an emphasis on chemicals selected as COPCs. The environmental conditions of the site and hydrological considerations that are likely to affect contaminant fate and transport near LL-9 and the surrounding area are also discussed. It is expected that the fate and transport of minor site contaminants would be similar to those of the major contaminants (i.e., COPCs) with similar physical and chemical properties.

Section 5.1 contains a general discussion of the various chemicals and their physical properties. Section 5.2 reviews the potential for chemical compounds to biodegrade or undergo other transformations. Section 5.3 discusses transport pathways where migration and attenuation might be occurring and how spatial and temporal variations in hydrologic conditions might be affecting transport. Section 5.4 presents an overall summary of contaminant migration in and around LL-9.

5.1 Chemical and Physical Properties Affecting Chemical Mobility

Table 5-1 presents physical and chemical properties of representative organic chemicals detected at LL-9. Because organic chemicals are not present in background, this table includes properties for the organic chemicals detected at LL-9 with the exception of minor contaminants. The COPCs are included in Table 5-1. Table 5-2 is a simple representation of the gross mobility of metal ions that is expected under various combinations of oxidation reduction (redox) potential and acidity (pH). These properties can be used to qualitatively estimate the environmental mobility and fate of site contaminants. The properties that are discussed include the following:

- Specific gravity,
- Vapor pressure,
- Water solubility-Henry's Law constant,
- Octanol/water partition coefficient (K_{ow}),
- Organic carbon partition coefficient (K_{oc}),
- Soil-water distribution coefficient (K_d),
- Bioconcentration factor (BCF), and
- Mobility index (MI).



Table 5-1
Environmental Fate and Transport Properties for Organic Chemicals

Chemical	Specific Gravity (@ 20/4°C) ⁽¹⁾	Vapor Pressure (mm Hg @ 20°C) ⁽¹⁾	Solubility (mg/L @ 20°C) ⁽¹⁾	Octanol/Water Partition Coefficient ⁽¹⁾	Organic Carbon Partition Coefficient ⁽²⁾	Henry's Law Constant (atm-m3/mole) ⁽¹⁾	Bioconcentration Factor (mg/L/mg/kg) ⁽²⁾	Mobility Index log((solubility*VP)/Koc)
ENERGETIC COMPOUNDS								
Nirocellulose	1.67 ⁽³⁾	NA	NA	NA	NA	NA	NA	NA
Nitroguanidine	1.71 ⁽⁴⁾	NA	NA	NA	NA	NA	NA	NA
RDX	1.82	NA	5.98E+01	7.41E+00 ⁽⁴⁾	7.89E+01	6.30E-08	1.92E+02 ⁽⁴⁾	NA
MONOCYCLIC AROMATICS								
2,6-Dinitrotoluene	1.2833 (111°C)	5.67E-04	1.82E+02	1.12E+02	2.04E+02	9.26E-08	1.20E+01	-3.30E+00
2-Methylphenol	1.047	2.99E-01	2.60E+04	9.55E+01	9.12E+01	1.20E-06	1.60E+01	1.93E+00
3-Methylphenol	1.034	1.40E-01	2.27E+04	9.33E+00	8.99E+00	7.09E-07	1.90E+01	2.55E+00
4-Methylphenol	1.0178	1.1E-1 (25°C)	2.4E+4 (25°C)	8.32E+01	9.0E-1(8)	3.92E-07	1.7E+1 ⁽⁴⁾	3.47E+00
4-Nitrophenol	1.27	5.00E-03	1.16E+04	8.13E+01	3.90E+01	1.30E-08	1.60E+01	1.71E-01
POLYCUCLIC AROMATIC HYDROCARBONS (PAHs)								
2-Methylnaphthalene	1.0058	1E+1 (105°C)	2.6E+1 (25°C)	7.24E+03	7.27E+2 ⁽⁵⁾	4.99E-4 (25°C)	5.1E+2 ⁽⁴⁾	-4.47E-01
Acenaphthene	1.07	5.00E-03	4.24E+02	8.32E+03	7.08E+03	1.55E-04	1.10E+03	-3.52E+00
Acenaphthylene	1.02	2.30E-02	1.61E+01	1.17E+04	2.00E+03	1.14E-04	3.80E+02	-3.73E+00
Anthracene	1.283 (25/4°C)	1.95E-4 (25°C)	1.29E+0 (25°C)	2.82E+04	2.95E+04 ⁽⁶⁾	8.6E-5 (25°C)	4.70E+03	-8.07E+00
Benzo(a)anthracene	1.274	5.00E-09	1E-2 (24°C)	4.07E+05	3.98E+05 ⁽⁶⁾	6.60E-07	5.30E+04	-1.59E+01
Benzo(a)pyrene	1.351	5.00E-09	3.8E-3 (25°C)	9.55E+05	1.02E+06 ⁽⁶⁾	4.9E-7 (25°C)	1.40E+05	-1.67E+01
Benzo(b)fluoranthene	NA	5.00E-07	1.2E-3 (25°C)	3.72E+06	1.23E+06 ⁽⁶⁾	1.20E-05	1.40E+05	-1.53E+01
Benzo(g,h,i)perylene	1.35	1.00E-10	2.6E-4 (25°C)	1.70E+07	1.60E+06	1.4E-7 (25°C)	3.50E+05	-1.98E+01
Benzo(k)fluoranthene	NA	9.59E-11	5.5E-4 (25°C)	6.92E+06	1.23E+06 ⁽⁶⁾	1.04E-03	1.40E+05	-1.94E+01
Chrysene	1.274 (20°C)	6.3E-9 (25°C)	6E-3 (25°C)	4.07E+05	3.98E+05 ⁽⁶⁾	1.05E-6 (25°C)	5.30E+04	-1.60E+01
Dibenzo(a,h)anthracene	1.282	1.00E-10	5E-4 (25°C)	9.33E+05	3.80E+06 ⁽⁶⁾	7.3E-8 (25°C)	6.90E+05	-1.99E+01
Dibenzofuran	1.0886	4.40E-02	4.22E+00	1.32E+04	8.13E+03	NA	8.00E+02	-4.64E+00
Fluoranthene	1.252	5.0E-6 (25°C)	2.65E-1 (25°C)	2.14E+05	1.07E+05 ⁽⁶⁾	6.5E-6 (25°C)	1.20E+04	-1.09E+01
Fluorene	1.202	1.00E+01	1.98E+00	1.62E+04	1.38E+04	6.36E-05	3.80E+03	-2.84E+00
Indeno(1,2,3-cd)pyrene	NA	1E-10 (25°C)	6.20E-02	4.57E+07	3.47E+06 ⁽⁶⁾	6.95E-8 (25°C)	3.50E+05	-1.77E+01
Naphthalene	1.162	8.2E-2 (25°C)	3E+1 (25°C)	2.34E+03	2.00E+03 ⁽⁶⁾	4.83E-4 (25°C)	4.20E+02	-2.91E+00
Phenanthrene	0.980 (4°C)	1E+0 (118.2°C)	8.16E-1 (21°C)	2.88E+04	1.40E+04	3.93E-5 (25°C)	4.70E+03	-4.23E+00
Pyrene	1.271 (23/4°C)	2.5E+0 (200°C)	1.6E-1 (26°C)	1.51E+05	1.05E+05 ⁽⁶⁾	5.1E-6 (25°C)	1.20E+04	-5.42E+00
PHTHALATE ESTERS								
Bis(2-ethylhexyl)phthalate	0.99 (20/20°C)	1.2E+0 (200°C)	4E-1 (25°C)	2.00E+05	1.51E+07 ⁽⁶⁾	3.00E-07	2.30E+08	-7.50E+00
Butylbenzylphthalate	1.113	8.60E-06	7.10E-01	6.92E+04	5.75E+04	1.26E-06	7.72E+02	-9.97E+00
Di-n-butylphthalate	1.047 (20/20°C)	1E-1 (115°C)	4E+2 (25°C)	1.58E+05	3.39E+04 ⁽⁶⁾	2.8E-7 (25°C)	4.70E+04	-2.93E+00

NA -Not Available

Footnotes:

- 1 EPA, September 1992, Handbook of RCRA Groundwater Monitoring Constituents: Chemical and Physical Properties.
- 2 USEPA, December 1982, Aquatic Fate Process Data for Organic Priority Pollutants.
- 3 Lyman et al., 1990; Equation 5-3, Handbook of Chemical Property Estimation Methods.
- 4 Lyman et al., 1990, Eq. 5-2
- 5 Lyman et al., 1990; Equation 4-5
- 6 EPA, July 1996, Soil Screening Guidance.



Table 5-2

**Relative Mobility of Select Materials Under Various
Environmental Conditions of Eh and pH**

Relative Mobility	Environmental Conditions			
	Oxidizing	Acidic	Neutral/Alkaline	Reducing
Very High			Se	
High	Se, Zn	Se, Zn, Cu, Ni, Hg, Ag		
Medium	Cu, Ni, Hg, Ag, As, Cd	As, Cd	As, Cd	
Low	Pb, Ba, Se	Pb, Ba, Be	Pb, Ba, Be	
Very Low	Fe, Cr	Cr	Cr, Zn, Cu, Ni, Hg, Ag	Cr, Se, Zn, Cu, Ni, Hg, Pb, Ba, Be, Ag

Notes:

As = Arsenic
Ag = Silver
Ba = Barium
Be =
Beryllium
Cd =
Cadmium
Cr =
Chromium
Cu = Copper

Fe = Iron
Hg = Mercury
Ni = Nickel

Pb = Lead

Se = Selenium

Zn = Zinc
Eh = Standard Redox Potential

Source: Swartzbaugh, et al. Remediating Sites Contaminated with Heavy Metals.
Hazardous Materials Control, November/December 1992.

Literature values of these properties are included for each organic chemical, when available, in Table 5-1. Calculated values, which were obtained using approximation methods, are listed when literature values are not available and the values could be computed. A discussion of the environmental significance of each of these parameters follows.

5.1.1 Specific Gravity

Specific gravity is the ratio of the density of a given volume of pure chemical at a specified temperature (usually 20 degrees Celsius [°C]) to the density of the same volume of water at a given temperature (usually 4°C). Its primary use is to determine whether a liquid chemical will have a tendency to float or sink in water if it is present as a pure chemical, i.e., at very high



concentrations. Non-aqueous phase liquids (NAPLs) with a specific gravity greater than 1, including halogenated aliphatic compounds, polycyclic aromatic hydrocarbons (PAHs), and pesticides, will tend to sink if present as a pure liquid or mixture of pure liquids. This is true whether the NAPL is present in a water body or in a matrix such as soil. Chemicals with a specific gravity less than 1 (e.g., light hydrocarbons) will tend to float. This physical characteristic becomes important only when the chemicals are at very high concentrations and are liquid when they are in pure phase. However, no NAPLs were detected LL-9. Therefore, the characteristic of specific gravity is not expected to be important for LL-9.

5.1.2 Vapor Pressure

Vapor pressure indicates the tendency for a chemical to volatilize from both solid and aqueous matrices. It is of primary importance at environmental interfaces such as surface soil/air and surface water/air. In unsaturated soils, volatility also governs for interstitial air pockets to become saturated with chemical vapor. Volatilization from stream sediments could also be significant under low-flow conditions (i.e., during summer months and drought conditions) when the sediments are exposed to the atmosphere in a dry creek bed. Volatilization is not as important when evaluating contaminated groundwater and subsurface soils that are not exposed to the atmosphere. Vapor pressures for ketones and halogenated aliphatics are generally many times greater than vapor pressures for PAHs, pesticides, energetic compounds, and phthalate esters. Generally, the more volatile chemicals are liquid at normal temperatures and pressures whereas the less volatile chemicals are solids under the same conditions. Chemicals with greater vapor pressures are expected to enter the atmosphere much more readily than chemicals with lower vapor pressures. Volatilization from surface water, surface soil, and sediment is a significant loss process for VOCs. Volatilization is not significant for most inorganics unless the inorganic materials have been converted to organometallic compounds through biochemical action.

5.1.3 Water Solubility

Solubility is easily determined under laboratory conditions. Many chemicals of environmental interest are often considered to be insoluble in reference books (e.g., any "Handbook of Chemistry and Physics") because they are insoluble, for most practical situations. However, most chemicals are soluble to some extent in water, and even those that are classified as insoluble may be soluble in the parts per billion ($\mu\text{g/L}$) or parts per million (mg/L) concentration ranges. These ranges may yield toxic effects, depending on the chemical, and are routinely measured during environmental work.

The tendency for a chemical to be leached from soil by infiltrating precipitation is governed by its water solubility. More-soluble chemicals are more readily leached than less-soluble chemicals. The water solubility listed in Table 5-1 indicate that ketones and halogenated aliphatics are usually several orders of magnitude more water soluble than pesticides, PCBs, and some of the



PAHs. PCBs, in particular, are not frequently detected as dissolved compounds in aqueous samples because of their low solubility.

5.1.4 Henry's Law Constant

Both vapor pressure and water solubility are useful for determining volatilization rates from surface water bodies and groundwater. The measured ratio of these two parameters (the Henry's Law constant) under equilibrium conditions is used to calculate the equilibrium chemical concentrations in the vapor (air) phase versus the liquid (water) phase for the dilute solutions commonly encountered in environmental settings. In general, chemicals having a Henry's Law constant less than 1×10^{-5} atm-m³/mole, such as pesticides, PCBs, and PAHs, are expected to volatilize very little and to be present only in minute amounts in the atmosphere or soil vapor. For chemicals with Henry's Law constants greater than 5×10^{-3} atm-m³/mole, such as many of the halogenated aliphatics, volatilization and diffusion in soil vapor could be significant.

5.1.5 Octanol/Water Partition Coefficient (K_{ow})

K_{ow} is a measure of the equilibrium partitioning of chemicals between octanol and water as determined under laboratory conditions. A linear relationship between the K_{ow} and the uptake of chemicals by fatty tissues of animal and human receptors, or the bioconcentration factor (BCF), has been established (Lyman et al., 1990). The K_{ow} is also useful in characterizing the sorption of compounds by organic soils where experimental values for soil are not available. Pesticides and aromatic compounds, lacking functional groups that enhance water solubility, are several orders of magnitude more likely to partition into fatty tissues than the more water-soluble VOCs. K_{ow} values are also used to estimate BCFs in aquatic organisms.

5.1.6 Organic Carbon-Water Partition Coefficient (K_{oc})

K_{oc} indicates the tendency of a chemical to adhere to organic matter contained in soils under laboratory conditions. Many VOCs, for example, have relatively low K_{oc} values and they tend to be fairly mobile in the environment as a result of groundwater or surface water movement. Chemicals with high K_{oc} values generally have low water solubility and vice versa. This parameter may be used to infer the relative rates at which the more mobile chemicals (e.g., ketones and halogenated aliphatics) are transported in groundwater. Chemicals, such as most pesticides, PCBs, and PAHs, are relatively immobile in the soil and are preferentially bound to the soil. These compounds are not subject to groundwater transport to the same extent as compounds with higher water solubility. However, these immobile chemicals can be transported by erosional processes when they occur in surface soils.

Several factors affect the measured value of K_{oc} . Values of K_{oc} usually decrease with increasing temperature. The fine silt and clay fraction of soil and sediments may have a greater tendency



to absorb chemicals because they often have a higher concentration of organic matter and more adsorption sites per unit volume.

5.1.7 Soil-Water Distribution Coefficient (K_d)

K_d is a measure of the equilibrium distribution of a chemical in soil/water systems. The K_d of organic chemicals is a function of both the K_{oc} and the fraction of organic carbon in the soil (f_{oc}):

$$K_d = K_{oc} * f_{oc}$$

The degree to which organic chemicals sorb to soils is very important when assessing migration potential. If a chemical tends to sorb strongly to soil, there is much less probability that the chemical will reach groundwater and affect the groundwater quality. In sediments, a high degree of sorption similarly indicates that the chemical is more likely to be transported in entrained sediments than as a dissolved species in surface water.

Chemicals, such as PAHs, that migrate slowly through soil and the upper unsaturated rock units have a relatively long time period where they are subjected to biodegradation processes before they reach the first water-bearing rock unit. As a consequence of low migration potential, there is a very low probability that chemicals with very high K_d values (i.e., PAHs) will reach surface water bodies via groundwater flow. However, if PAHs or pesticides are present at the ground surface, eroded, and transported in surface runoff with soil particles (as a sorbed phase), then there is potential for these compounds to reach adjacent stream channels.

For metals, the sorption coefficients are, in large part, a representation of the ion exchange tendencies of the metals with the soil. Table 5-3 presents sorption coefficients for select metals, several of which were detected at LL-9. Other factors aside, the more positively charged an ion, the more tightly it is bound to soils. Ions that bind tightly tend to displace ions that are less tightly bound because the ions compete for the same ion exchange sites. In addition, soils with high organic material content tend to bind metals in large chemical complexes. This process is described in the section below.

Table 5-3

Soil-Water Partition Coefficients for Metals

Inorganic Element	K _d at pH = 5.5 ⁽¹⁾ (L/kg)	K _d at pH = 6.2 ⁽²⁾ (L/kg)	K _d ⁽³⁾ (L/kg)
Aluminum	NA	NA	1,500
Antimony	NA	NA	45
Arsenic	26	28	200



Table 5-3 (Continued)

Soil-Water Partition Coefficients for Metals

Inorganic Element	Kd at pH = 5.5 ⁽¹⁾ (L/kg)	Kd at pH = 6.2 ⁽²⁾ (L/kg)	Kd ⁽³⁾ (L/kg)
Barium	21	33	41
Beryllium			790
Cadmium	27	42	75
Chromium (as Cr ⁺³)	2,100	420,000	1,800,000
Chromium (as Cr ⁺⁶)	NA	NA	850
Cobalt			45
Copper	NA	NA	428
Iron	NA	NA	25
Lead			900
Magnesium			5
Manganese	NA	NA	65
Mercury			52
Nickel	28	42	65
Selenium	12	75	5
Silver			8
Strontium	NA	NA	35
Thallium			71
Vanadium	NA	NA	1,000
Zinc	26	42	62

(1) Values from Illinois EPA Tiered Approach to Corrective Action Objectives (TACO) Program; Kd values presented assume an average pH of 5.5.

(2) Values from Illinois EPA TACO Program; Kd values presented assume an average pH of 6.2.

(3) Values from the U.S EPA-sponsored Risk Assessment Information System (RAIS) Internet site:
"http://risk.lsd.ornl.gov/rap_hp.html"

NA = Not applicable.

Kd = Soil-water distribution coefficient.

5.1.8 Chemical Complex Formation

Metals may form chemical complexes or combinations that alter the mobility of the metals. This may also occur for non-metals. Some of the most important environmental chemicals that form metal complexes are the humic substances. These complex mixtures of organic acids and other organic matter are formed naturally in shallow surface soils, in pond bottoms, lake bottoms, through the decay of vegetable matter. These substances, after binding with a metal, can increase its mobility by dissolving into an aqueous phase. If the humic substances are



adsorbed to a solid substrate, such as soil, sediment, or vegetation, they will tend to remove metal ions from solution by binding to the metals and fixing them to the solid substrate. The binding equilibria are affected by pH. At low pH, the bound metals are released and at high pH the metals are preferentially bound. Iron is an exception to this rule (Dragun, 1988). Fulvic acid is a component of humic substances. The pH at which fulvic acid complexes begin to release relatively large proportions of bound metals has been shown to be less than 5 (Dragun, 1988). Thus, most metals are expected to be preferentially bound to the humic substances at pH values greater than 5.

5.1.9 Bioconcentration Factor

BCF represents the ratio of aquatic animal tissue concentration to water concentration. The ratio is both contaminant- and species-specific. When site-specific values are not measured, literature values are used or the BCF is derived from the K_{ow} . Many of the pesticides and PAHs will bioconcentrate at levels three to five orders of magnitude greater than those concentrations found in the water, but VOCs and energetic compounds are not as readily bioconcentrated. Any bioconcentration that occurs generally requires that the chemical of interest be in the dissolved state; otherwise, the chemical is inaccessible to the organism. For plants, this means that the chemical must be dissolved in the surrounding soil. For other organisms that can inhale, aspirate, or ingest solid particles, the chemical may be bound to the particles and released after ingestion.

5.1.10 Mobility Index

The MI is a quantitative assessment of mobility that uses water solubility (S), vapor pressure (VP), and the K_{oc} (Laskowski, 1983). It is defined as

$$MI = \log ((S*VP)/K_{oc})$$

A scale to evaluate MI, as presented by Ford and Gurba (1984), is:

<u>Relative MI</u>	<u>Mobility Description</u>
> 5	extremely mobile
0 to 5	very mobile
-5 to 0	slightly mobile
-10 to -5	immobile
< -10	very immobile

Of the organic chemicals detected at LL-9, chlorinated solvents and ketones generally have MIs greater than 5 and are considered extremely mobile. Lighter molecular weight PAHs, such as naphthalene, have MIs ranging from -5 to 0 and are considered slightly mobile. Heavier molecular weight PAHs [e.g., benzo(a)pyrene] are classified as very immobile, having MIs less than -10.



5.1.11 Miscellaneous Characteristics

The solubility and mobility of an inorganic chemical is strongly influenced by its valence state(s) and mineral forms present in soils (e.g., silicates, hydroxides, oxides, and carbonates). The solubility of a metal also depends largely on the pH and oxidation-reduction potential (Eh) of its environment, the ambient temperature, and other ionic species in solution (the Debye-Huckel theory).

Most metals have increased solubility and mobility at lower water pH values (e.g., < 5.0). Some metals such as aluminum, however, have increased solubility at both low and high pH values (greater than 8). Iron and manganese are metals that have more than one common valence state and are more soluble in the reduced valence states. For example, iron(+3) and manganese(+4) are generally less soluble than iron(+2) and manganese(+2) in the environment. As a result, these metals are more mobile under reducing conditions. Chromium(+6), however, is more mobile than Chromium(+3) under typical environmental conditions. Mobility is highly dependent on the anions that are associated with the metal. For example, silver nitrate is very soluble in water, but most other silver compounds are highly insoluble. The solubility product constants reported in the literature vary with the type of chemical complex formed.

Metals can be rapidly transformed from one oxidation state to another as they encounter different local environments. For example, manganese that may have been dissolved from local geologic materials at one location may precipitate out of solution as the groundwater containing the dissolved manganese migrates to another area. The wide array of potential metal-anion combinations and environmental conditions leads to a wide array of potential metal mobility that must be evaluated on a case-by-case basis for potential environmental impact.

5.2 Chemical Persistence and Degradation Process

Degradation and other transformation processes that affect site contaminants are discussed in this section. Degradation leads to the following possibilities, depending on the contaminant of interest and the contaminant's environment:

- Transfer of the chemically unaltered contaminant from one environmental medium to another (physical changes may occur (e.g., transfer of dissolved contaminant to vapor);
- Transformation of the original contaminant into a less toxic chemical;
- Transformation of the original contaminant into a more toxic or otherwise problematic chemical; and
- Transfer of a chemically or biochemically transformed contaminant from one environmental medium to another.



1 These transformations may occur individually or together, and more than one of these
2 processes may occur for any single chemical. Consequently, the number and complexity of
3 possibilities is partly a function of the number of contaminants and partly a function of which
4 contaminants are present.

5 For organic chemicals, degradation typically involves a gradual reduction in the molecular
6 weight of the chemical as molecular bonds are cleaved. Cleavage may occur in sequential
7 steps with the most susceptible bonds being cleaved first. Because of this, larger, more
8 complex molecules that have more chemical bonds commonly have more molecular
9 degradation pathways than smaller molecules. With the changes in molecular structure come
10 changes in chemical and physical properties. For example, daughter compounds are commonly
11 more volatile than their parent compounds. They may also be more or less toxic than the parent
12 compound. For inorganic chemicals, degradation may be a purely inorganic chemical reaction
13 that results in a change of valence state. Depending on the chemical, it may also involve
14 biochemical reactions that transform the inorganic chemical into an organometallic compound or
15 another inorganic species.

16 Hydrolysis, biodegradation, photolysis, and chemical oxidation/reduction reactions are four
17 processes that are important for many environmental investigations because the potential for
18 their occurrence is high, depending on the contaminants that are present and where those
19 contaminants are located. These four processes are discussed in more detail below.

20 Hydrolysis is a reaction between the chemical of interest and water that results in a cleavage of
21 the water molecules. Many of these reactions can be catalyzed by the presence of acids or
22 bases.

23 Biodegradation is a generic term that includes oxidation or reduction transformations resulting
24 from enzymatic or other biochemical processes that occur in or near living organisms. Many of
25 these reactions take place in naturally occurring microorganisms, such as bacteria, that are
26 present in the Earth's crust, especially at the Earth's surface. A chemical or family of chemicals
27 (e.g., chlorinated solvents or ketones) may be highly susceptible to transformation by one type
28 of microorganism but much less so by other organisms. Consequently, the rates of
29 biodegradation can vary widely from no degradation to very rapid degradation, depending on
30 the presence of organisms that can degrade the chemicals. The presence of other chemicals
31 and the ambient conditions can also greatly influence the potential for, and the rate of,
32 degradation. Rates are difficult to predict, however, because they are sensitive to ambient
33 conditions, including the presence or absence of microorganisms that are capable of degrading
34 particular chemicals biochemically. Some of these difficulties are described further below.

35 Photolysis is the cleavage of chemical bonds induced by light, especially the ultraviolet
36 wavelengths. Hence, environmental photolysis characteristically occurs at the Earth's surface in



surface water, shallow surface soils, shallow sediments exposed to air, and in the atmosphere, where exposure of the chemicals to sunlight can occur routinely.

Degradation of chemical compounds can frequently be modeled as a first-order decrease or decay of the concentrations over time. If the concentration of the chemical is plotted over time, the concentration decrease is rapid early in the degradation process but becomes less rapid as time goes on because the rate of degradation at any point in time is directly proportional to the remaining concentration of the chemical. Such models are useful for predicting the concentration of a chemical at any future time. The true degradation, however, may not follow a first-order decay process even though a first-order decay is used to model the degradation. Furthermore, because many different decay processes can occur in concert, which may have different mechanisms, the first-order decay model will effectively be a composite of the pertinent processes. This often leads to significant uncertainties in estimating degradation rates. Any uncertainties associated with estimating the degradation rates translate into uncertainties in estimates of future concentrations.

A characteristic of first-order decays is a relationship between half-life and degradation rate. A half-life is the time required to reduce a concentration to one-half of its value. Chemicals with long half-lives have a long persistence in the environment, whereas chemicals with short half-lives degrade or disperse quickly. For first-order decays, the half-life, $t_{1/2}$, is equal to the natural logarithm (Ln) of 2 divided by the degradation rate constant, k (i.e., $t_{1/2} = \text{Ln}(2)/k$). In the discussions below, mathematical models are not used to estimate degradation rates or to predict future concentrations. The concept of half-life is, however, occasionally referenced to offer perspective on relative degradation rates of various chemicals. Table 5-4 presents half-lives of select chemicals detected at LL-9. The half-lives presented in Table 5-4 and discussed below are taken from literature studies and may not accurately represent degradation rates at LL-9, but they indicate relative degradation rates to be expected for these compounds.

Table 5-4

Ranges of Biodegradation Rates for Representative Organic Chemicals

Chemical	SOIL		GROUND WATER	
	Short Half-Life	Long Half-Life	Short Half-Life	Long Half-Life
	(days)	(days)	(days)	(days)
Energetics				
RDX	NA	NA	NA	NA
Nitrocellulose	NA ⁽¹⁾	NA ⁽¹⁾	NA ⁽¹⁾	NA ⁽¹⁾



Table 5-4 (Continued)

Ranges of Biodegradation Rates for Representative Organic Chemicals

Chemical	SOIL	GROUND WATER	Chemical	SOIL
	Short Half-Life	Long Half-Life	Short Half-Life	Short Half-Life
	(days)	(days)	(days)	(days)
Polycyclic Aromatic Hydrocarbons				
Acenaphthene	12.3	102	24.6	204
Benzo(a)anthracene	102	680	204	1360
Benzo(a)pyrene	57	530	114	1060
Benzo(b)fluoranthene	360	610	720	1220
Benzo(k)fluoranthene	5	23	10	389
Chrysene				
Fluoranthene	140	440	280	880
Indeno(1,2,3-cd)pyrene	560	730	1200	1460
Phenanthrene	16	200	32	400
Pyrene	210	1900	420	3800
Semi-volatile Organic Compounds				
Bis(2-ethylhexyl)phthalate	5	23	10	389

1 Assume to be greater than 365 days, based on properties of cellulose.

2 Rough estimate based on: Erickson, 2001 – See text references for full citation.
Actual half-lives vary widely.

NA = Not available.

The fate and transport of each contaminant detected at LL-9 is described below. The chemicals are discussed in groups that are indicated by their analytical fractions (e.g., VOCs or metals) because these fractions generally represent chemicals with similar properties. Only those processes that are most applicable to the indicated fractions are described. No attempt was made to model the fate and transport of the chemicals mathematically, in part because simple rules of thumb concerning contaminant half-lives are sufficient to gain an understanding of contaminant fate. In addition, the levels of contaminants are generally low, and migration will tend to reduce the concentrations further through dispersion, dilution, and other transport mechanisms. The contaminants that were identified are generally considered to be immobile or slow-moving contaminants so the migration rates will be slow, except when migrating in sediments and surface water as adsorbed or dissolved species.



5.2.1 Volatile Organic Chemicals

No chemicals in this category were detected at LL-9. Therefore, VOCs will not be discussed further.

5.2.2 Semi-volatile Organic Chemicals

Benzo(a)pyrene, dibenzo(a,h)anthracene, and polycyclic aromatic hydrocarbons (PAHs), were the only chemicals in the SVOC category to be selected as human health COPCs for LL-9. Several other SVOCs, most of which are PAHs, were detected at LL-9, and some of these were selected as ecological risk COPCs. Appendix V, Tables 2.1 through 2.5, and Appendix W, Tables 1 and 2, indicate which SVOCs were detected at LL-9 and, of those, which were selected as COPCs in various media. Benzo(a)pyrene and dibenzo(a,h)anthracene were selected as human health COPCs in soil only. Several PAHs were selected as ecological COPCs in surface soils, and dibenzo(a,h)anthracene and bis(2-ethylhexyl)phthalate were selected as ecological COPCs in sediment (see Tables 7-1 and 7-2).

PAHs, as a group, are much more likely to bind to soil and to be transported via erosion and surface water runoff than to be solubilized in surface water or groundwater. PAHs are subject to slow degradation *via* aerobic bacterial metabolism, but may be relatively persistent in the absence of microbial populations or macronutrients such as phosphorus and nitrogen.

Landspreading applications have indicated that PAHs are highly amenable to microbial degradation in soil. This would apply to sewer sediments as well. Although sewers and manholes were plugged, degradation is still expected to occur in those locations if residual contamination remains. The rates of degradation in these media are influenced by temperature, pH, oxygen concentrations, initial chemical concentrations, and moisture. Photolysis, hydrolysis, and oxidation are not important fate processes for the degradation of PAHs in soil (Agency for Toxic Substances and Disease Registry [ATSDR], 1997).

The most important fates of PAHs in water are photo-oxidation, chemical oxidation, and biodegradation. Metabolism in higher animals is another degradation pathway and becomes important when the PAHs are metabolized to carcinogenic species (UN, 1998). PAHs do not contain functional groups that are susceptible to hydrolytic action, and hydrolysis is considered to be an insignificant degradation mechanism. The rate of photo degradation is influenced by water depth, turbidity, and temperature. Benzo(a)pyrene is reported to be resistant to photo degradation. PAHs may also be metabolized by microbes under oxygenated conditions (ATSDR, 1997).

The longest PAH half-life (of those shown in Table 5-4) is about 4 years in groundwater and 1 year in soil. Thus, PAH concentrations should decrease significantly within a decade and should be effectively degraded in less than 30 years in groundwater and less than 10 years in



soil. The small reservoir of PAHs at LL-9 indicates that long-term contamination will not be a problem.

Bis(2-ethylhexyl)phthalate is more soluble in water than the PAHs. Chemical hydrolysis is essentially non-existent, but photo degradation in the atmosphere is rapid (ToxNet, 2004). Aerobic degradation can be affected by several soil microorganisms; however, the microbial degradation in the environment has been reported to be slow and temperature-dependent (ToxNet, 2004). At temperatures below about 10° C, little degradation takes places. Butyl benzyl phthalate and di-n-butyl phthalate, also were detected at LL-9 but only in sediment. These chemicals, are expected to behave similarly to bis(2-ethylhexyl)phthalate with regard to chemical and biochemical transformations. Bis(2-ethylhexyl)phthalate is highly lipophilic and moderately persistent in the environment. The degree of bioaccumulation depends on the capability of an organism to metabolize this compound. The compound accumulates in a variety of aquatic invertebrates, fish, and amphibians. Plant uptake is low (ToxNet, 2004).

5.2.3 Pesticides and Polychlorinated Biphenyls

No pesticide or polychlorinated biphenyl compounds were detected at LL-9 in any of the investigated media. Therefore, these chemicals are not discussed further.

5.2.4 Explosive and Propellant Compounds

Nitroguanidine, nitrocellulose, and RDX were detected in soil samples collected from LL-9. Nitrocellulose was also detected in surface water, groundwater, and sediment; but the concentrations are very low and appear to be potential false detections. Nitroguanidine was detected in surface soil only and at a concentration (89 µg/kg) within the normal detection limit range; therefore, it may be a false detection. Of these, nitrocellulose (in soil, sediment, and surface water) was selected as an ecological COPC for shallow soil because there are no toxicity data for this compound. Nitroguanidine was selected as an ecological COPC in sediment for the same reason. Findings for these chemicals were within the range of normal detection limits, indicating that the detections could be false.

The low solubility of nitrocellulose suggests that it is not very toxic in soil and sediment. Although nitrocellulose (cellulose nitrate) may be incorporated into propellant mixtures, it has a very low solubility and, consequently, exhibits very limited mobility. It is a fibrous material composed essentially of cellulose fibers that have been modified to contain nitrate (NO₃) functional groups. The chemical composition, however, suggests that any environmental degradation of this chemical could release nitrates or nitrites to the environment. Because of its low solubility, relatively slow degradation rates in the environment are expected. Thus, any nitrates or nitrites detected at the site are presumably related to other nitrates that are much more soluble, such as sodium or potassium nitrate. These metal salts, especially sodium nitrate, are used in large quantities in the manufacture of black powder.



Studies (e.g., ANRCP, 1998; and Roberts and Kotharu, no date) indicate that RDX is likely to break down under highly reducing conditions to mono-, di-, and tri-nitroso derivatives as well as smaller molecules. One known reductive degradation pathway involves a sequential conversion of nitro functional groups to nitroso groups followed by ring cleavage to yield relatively non-toxic compounds. Another pathway involves degradation by microbial consortia to yield nitramines, formaldehyde and, ultimately, carbon dioxide and nitrous oxide. RDX is less commonly reported to be subject to oxidative degradation, usually persistent under aggressive oxidizing and moderately reducing conditions (e.g., Adam et al., *J. Environ. Qual.* 33:2165–2173 [2004]). Roberts et al. (no date) also report that oxidative biodegradation occurs via sequential oxidation of the nitro groups with ultimate formation of 4-nitro-2,4-diazabutanal. This dead-end product may react with other chemicals to form yet smaller degradation products. The oxidative pathway, however is generally thought to be much less prevalent than the reductive pathway under typical environmental conditions for pH values.

Detections of energetic compounds were generally sparse with the exception of nitrocellulose; therefore, only the nitrocellulose is considered to be a potential environmental concern. The potential environmental concern is minor because of its low solubility and, hence, low biochemical availability and migration potential. In addition, the majority of the nitrocellulose results were qualified as estimated (J), or (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J). Additional information on the environmental significance of detected energetic compounds is included in Sections 6 and 7.

5.2.5 Metals

Metals are highly persistent environmental contaminants because they do not biodegrade. However, their valence states can readily change based on pH and Eh conditions, biotic uptake and assimilation into living organisms, and decay and decomposition of dead plant matter. The major fate mechanisms for metals are adsorption to the soil matrix or precipitation as a mineral coating, uptake and bioaccumulation in plants, or transport through the hydrologic system. Table 5–2 presents a qualitative characterization of select metal nobilities in the environment as a function of ambient conditions. Because the chemical properties of the metals can vary significantly from metal to metal, only those metals detected at LL–9 are discussed here. Metals that have similar properties (as presented in Table 5–2 or discussed in the text) will behave similarly with regard to fate and transport.

Metals commonly form carbonates, silicates, oxides, and hydroxides after they have been released to the environment and interact with carbon dioxide, oxygen, and water in their surroundings. The carbonate, oxide, and hydroxide equilibria that dictate whether one species predominates over another, are sensitive to the surrounding pH and oxidation-reduction (redox) potentials. At high pH values, oxides and hydroxides form readily. At pH values less than about 4, these species are essentially non-existent, except under highly oxidizing conditions.



The Earth serves as an almost limitless reservoir of metals. Soils, surface water, groundwater, and sediments are typically well-buffered with respect to pH (i.e., pH is not easily changed). Based on geotechnical data from other RVAAP load line soil samples, the pH of soils measured at RVAAP load lines is typically in the range of 5.5 to 7.4 (USACE, 2003a, 20003b, and 2003c). The same pH range is expected for LL-9. Soil samples occasionally exhibit pH values less than 5.0, but they are rare (USACE, 2003a, 20003b, and 2003c). Therefore, it is assumed that low pH values are limited to small soil volumes. The pH buffering capacity near neutral to slightly acidic pH values tends to limit the ability of most metals to migrate rapidly because the metals tend to precipitate as solids or to bind tightly to soil particles through cation exchange. The positively charged metals or oxidation metal ions adsorb to negatively charged hydroxyl groups at the surface of the soil or sediment particles, which are often silicate minerals. In general, the more highly positive a cation, the more tightly bound it will be to the soil. Hence, monovalent ions (e.g., potassium, sodium) are not tightly bound whereas the trivalent aluminum is.

Each of the metals detected at LL-9 (except for the macronutrients: calcium, magnesium, potassium, and sodium) is described briefly below regarding its most important environmental chemistry characteristics and information concerning plant uptake. These factors affect the mobility and, hence, the fate of metals in the environment.

Aluminum, atomic symbol Al (not listed in Table 5-2), is a COPC in LL-9 sediment, surface soil, subsurface soil, and surface water. This metal is a naturally occurring component of silicate minerals and other minerals. It does not migrate readily because its aqueous ion has a positive charge of "+3". This high charge causes aluminum ions in solution to bind preferentially to negatively charged soil adsorption sites, such as OH groups located at the surfaces of silicate minerals. Aluminum also readily precipitates from solutions as aluminum hydroxides. Aluminum does dissolve in infiltrating precipitation, however, and may be taken up by plants from the resulting subsurface and surface soil solutions of aluminum. Any industrial releases of this metal to soils are not expected to migrate significantly. Releases to surface water will have the greatest mobility because aluminum adsorbed to sediments is transported this way, and aluminum in the dissolved phase can move for long distances before being completely removed from solution by binding to sediments and soils. Aluminum that is taken up by plants will be redeposited to soils when the vegetation dies or releases leaves and branches. Aluminum is not known to be transformed biologically.

Antimony, atomic symbol Sb (not listed in Table 5-2), is similar to arsenic (described below) in chemistry. This metal, commonly classified as a metalloid because its chemical properties lie between those of true metals and non-metals, has a mean concentration in soils around the world of approximately 1 mg/kg (Emsley, 2001). Stibnite (Sb_2S_3), a sulfide, is the predominant mineral of antimony found in native ores (Van Nostrand, 1976). The two prevalent valence states of antimony are Sb (III) and Sb (V). These relatively high oxidation charges tend to limit the mobility of antimony in the environment. This metal was selected as a human health COPC in LL-9 groundwater.



Arsenic, atomic symbol As, is a human health COPC in LL-9 surface and subsurface soil and surface water because its concentrations exceed human health screening levels in those media. It was selected as an ecological COPC in surface soil because its maximum concentration exceeded the ecological screening criteria for soil and it is a bioaccumulative chemical. Arsenic is a naturally occurring metal, commonly classified as a metalloid because it has properties that are transitional between metallic and non-metallic chemical elements. In minerals, this element commonly appears as a sulfide (NIH, 2004), which implies that this form of arsenic is a particularly abundant or persistent form. In water, inorganic arsenic occurs primarily in the "+5" valence state when conditions are oxidizing, such as in surface water. Under reducing conditions, such as may be found in groundwater, the inorganic form of arsenic is predominantly the "+3" valence state. In acidic or neutral waters, As (V) is extensively adsorbed but As (III) is not. At high pH (e.g., pH > 9), neither species of arsenic is tightly adsorbed (NIH, 2004). The mobility of arsenic is low to moderate in clay but higher in loamy or sandy soils. As (V) is adsorbed most strongly at pH 5 in clay minerals. As pH increases, more As (III) is adsorbed. Arsenic adsorbs strongly to sediments. If conditions change sufficiently from oxidizing to reducing, arsenic may be released from sediments.

Bacterial and fungal activity can convert inorganic arsenic to methyl arsines. These volatile organometallic compounds are mobile and can be transferred from sediments to the surrounding water and, ultimately, to the atmosphere. The potential for volatilization from moist surfaces varies greatly with local conditions. Tightly sorbed arsenic or complexed arsenic species will be unlikely to volatilize, whereas methylated arsenic species can volatilize fairly readily (NIH, 2004).

Barium, atomic symbol Ba, is not a COPC in any LL-9 media. This metal is commonly found in nature as sulfate (Emsley, 2001). It forms hydroxides, carbonates, and sulfates in soils. Barium carbonates and sulfates are relatively insoluble (negative logarithm of solubility product, pK_{sp} , constants near 9). By contrast, barium hydroxides (pK_{sp} near 2), and especially nitrates and chlorides of barium, are quite soluble (Lang, 1985). Thus, over time, barium is expected to precipitate out of solution as it migrates to areas containing high carbonate or sulfate concentrations and high concentrations of other anions that form barium precipitates. In the absence of these anions, barium is likely to be moderately mobile. Barium is generally classified as a metal of low mobility (See Table 5-2).

Beryllium, atomic symbol Be, was detected in most LL-9 soil samples with a maximum detection site wide of 1.8 mg/kg. The maximum detection in sediment was 0.99 mg/kg. This metal is reported to have a typical concentration of 6 mg/kg in worldwide soils and was not selected as a human health COPC for LL-9 (Emsley, 2001). It was, however selected as an ecological COPC in sediment. Beryllium can be taken up by plants. Typical concentrations in plants are too low to exhibit any effects on animals that live off of them, even in plants that have the ability to concentrate the beryllium (Emsley, 2001). The divalent Be^{+2} ion dominates environmental beryllium chemistry at pH values less than about 4. Above that pH it forms



insoluble hydroxides (Dragun, 1988). Consequently, this metal generally exhibits low mobility under typical environmental conditions.

Cadmium, atomic symbol Cd (not listed in Table 5-1), is not a human health COPC in any LL-9 media but was retained as an ecological COPC in surface soil. While cadmium concentrations are typically less than 1 mg/kg in worldwide soils, local hot spots have been shown to contain cadmium concentrations as high as 40 mg/kg (Emsley, 2001). This element is widely present in food crops at low levels and can be concentrated by the fungus *Amanita muscaria*, even when grown in soils containing low cadmium levels. Tobacco and other large leaf crops are also known to contain cadmium. Cadmium is present in sewage; and crops grown in sewage land farms can lead to an accumulation of cadmium in animals that eat the crops, including humans. This metal mimics zinc in its chemical behavior, especially in the human body. The lifetime in the human body is reported to be about 30 years (Emsley, 2001). Cadmium is expected to be moderately mobile in soil because it is present under most naturally occurring redox and pH conditions as the "+2" ion. If it is combined with anions such as sulfide, it can be very immobile. At pH values greater than about 8, cadmium tends to form insoluble hydroxides. These conditions are not expected at LL-9; therefore, the cadmium mobility is expected to be moderate.

Chromium, atomic symbol Cr, exists primarily as two oxidation states – Cr (III) and Cr (VI). Cr (IV) is very mobile in the environment under oxidizing conditions, but Cr (III) is less mobile except at pH values less than about 4. The trivalent metal predominates under most conditions except those that are low pH (e.g., less than 4) and high oxidation potential (e.g., greater than 0.4). This element forms insoluble hydroxides under low to moderately reducing conditions. Under highly oxidizing conditions, it forms the soluble chromate ion at pH values greater than about 6 and the bichromate ion at lower pH values (Dragun, 1988). Concentration values in soils worldwide are reported to range from 1 to 450 mg/kg (Emsley, 2001). Chromium is mobile in ground and surface water under oxidizing conditions and can move in sediments when adsorbed to sediment grains. Chromium was selected as a human health COPC for LL-9 in soil and surface water. It was also selected as an ecological COPC for surface soil.

Cobalt, atomic symbol Co (not listed in Table 5-1), is not a COPC in any LL-9 media. This metal is an essential nutrient and is found in relatively low abundance (0.1 to 70 mg/kg) in the Earth's crust (Emsley, 2001). Terrestrial plants absorb this element from soils. Based on its mean K_d equal to 4 (Dragun, 1988), this element has moderate to low mobility in soils and sediments. It forms insoluble oxides and hydroxides in the environment at pH values greater than about 8 but is otherwise generally present as the Co (II) ion at most oxidation-reduction potentials (Dragun, 1988).

Copper, atomic symbol Cu, is a relatively immobile element in soils (Emsley, 2001). Its affinity for organic matter causes it to be bound tightly in surface soils. Where subsurface releases of copper occur, this binding mechanism may not be as important because the most organic-rich



layer of soils is generally the surface layer. Plant uptake varies widely for this element and it is transmitted through the food chain. Most of the copper taken up by plants, however, remains in the roots (Emsley, 2001). At low pH, this metal is prevalent as the divalent ion under most redox conditions. Under reducing conditions, the metal may appear as pure metal. At high pH, the metal tends to form hydroxides and oxides (Dragun, 1988). The concentrations of this metal were less than human health screening levels except in surface soil. The highest surface soil copper concentrations (1,240 mg/kg) was about an order of magnitude greater than the next highest concentration (170 mg/kg in sample LL9SS-068-0001-SO). The highest copper concentration was colocated with highest concentrations of lead, and mercury and the second-highest concentrations of zinc in surface soil. Copper was selected as a human health COPC for LL-9 surface soil. It was also selected as an ecological COPC for surface soil and surface water. Transport in surface water is expected to occur in the suspended solids that are washed into streams then washed downstream. The amount of material transported in this manner, however, is expected to be small because the reservoir of high copper concentrations is spatially limited. It appears that the colocated copper, lead, and zinc represent remnants of one or more shell casings and projectiles, which would be consistent with site operations.

Iron, atomic symbol Fe, is a major component of the Earth's crust, being the fourth most abundant element (Emsley, 2001). It occurs widely as in oxide forms and is rarely found in the pure metal state. Iron in soil is readily transformed by organic matter into various oxides and other compounds. The most common oxidation state is "+3". This valence state can, however, be readily altered to the "+2" valence in groundwater and other aqueous environs under chemically reducing conditions. Fe (II) is more mobile than Fe (III). An abundance of iron in the Fe (II) state can serve as a large electron reservoir to electron acceptors during redox reactions. Iron in soil is not expected to be mobile; but under reducing conditions, iron can be mobilized. In sediments, iron is expected to travel primarily as an adsorbed or particulate species in surface drainage channels because the water is mixed with oxygen in the air to maintain the Fe (III) state. The most prevalent occurrence of Fe (II), then, is groundwater under reducing conditions. Pure metal can be found at very low oxidation potentials and neutral to high pH. Iron was not selected as a human health COPC in any medium at LL-9. It was selected as a COPC for ecological risk in surface soil and surface water.

Lead, atomic symbol Pb, is a very immobile element in the environment. This metal has a relatively high soil-water distribution coefficient (ranging from 4.5 to 7,640 milliliters per gram [ml/g]; Dragun, 1988). This indicates that it preferentially binds to soils rather than dissolving in water. When adsorbed to sediments, this metal will be transported with the sediment as a bound species. This metal appears in the divalent ("2") state under most pH and redox conditions. At high pH values (e.g., greater than 8), it readily forms hydroxides and oxides. The highest concentration of soil lead was detected in sample LL9SS-011-0001-SO (1,330 mg/kg) and was co-located with high concentrations of copper, mercury, and zinc. The lead, copper, and zinc may represent residual contamination from brass shell casings (composed primarily of copper and zinc) and lead projectiles. Concentrations of this lead exceeded human health



screening levels in soil but not in any other media except surface water. Therefore, lead was selected as a COPC for LL-9 soil and surface water. It was also selected as an ecological COPC for surface soil, sediment, and surface water.

Manganese, atomic symbol Mn (not shown in Table 5-2), exhibits chemistry similar to that of iron. This essential nutrient, the twelfth most abundant element in the Earth's crust, occurs primarily in the "+2" valence state. In soil, manganese occurs as hydroxides and oxides, and microbial action plays a role in valence state alteration. The least stable form of manganese, Mn (III), is the most biologically active (Emsley, 2001). Manganese can also exist as Mn (IV). Manganese is most soluble in water in the Mn (II) form; thus, its mobility is greatest under reducing conditions. Under these conditions it is most susceptible to migration in groundwater or uptake by plants (Emsley, 2001). When adsorbed to soil or sediment particles, manganese may migrate in surface water. The concentrations of this metal were greater than human health screening levels in surface and subsurface soils, groundwater, and surface water; and manganese was selected as a human health COPC for LL-9 in these media. Its concentrations in shallow soil and surface water also exceeded ecological screening criteria; and it is selected as an ecological COPC in surface soil and surface water.

Mercury, atomic symbol Hg, has a complex environmental chemistry. This metal, under most redox conditions, persists as an immobile species. It can be converted to organometallic species that are much more volatile, especially in wet sediments. The two common valence states are the "+1" and the "+2" states with Hg (II) predominating under typical environmental conditions. Mercury, when deposited as the pure liquid metal (e.g. from barometers, manometers, and thermometers), will persist for decades in the environment. The liquid mercury coalesces into globules that can slowly release dissolved mercury to the environment. However, the mercury thus released would be expected to precipitate as a chloride, carbonate, hydroxide, or oxide over time. Worldwide, concentrations of mercury in uncontaminated soil are reported to be 0.01 to 0.05 or 0.08 µg/kg (Dragun, 1988; Emsley, 2001). The concentrations of this metal in soil exceeded human health screening levels, especially in sample LL9SS-011-0001-SO (882 mg/kg). Mercury concentrations in soil of this magnitude are unusual. The fact that the high detection was co-located with other metals in a relatively small area suggests that the detection is real. The limited spatial extent of contamination, however, limits the total mass of mercury available for migration to other locations. Mercury was selected as a human health and ecological COPC for LL-9 soils. This metal was also selected as a COPC for sediment and surface water.

Nickel, atomic symbol Ni, predominates under typical environmental conditions as the Ni (II) species. Under high pH conditions (e.g., pH greater than 8), this metal will tend to precipitate as an oxide or hydroxide, except under very reducing conditions where pure metal may exist (Dragun, 1988). Typical soil concentrations are reported to be approximately 50 mg/kg (Emsley, 2001) or even as high as 1,000 mg/kg (Dragun, 1988). The concentrations of this metal were less than human health screening levels, and nickel was not selected as a human



health COPC for LL-9; but it was selected as an ecological COPC for surface soil. The mobility of this metal ranges from low in neutral or alkaline conditions to high in acidic environments (see Table 5-2).

Selenium, atomic symbol Se, has chemistry similar to arsenic. This bioaccumulative essential nutrient is a metalloid, thus exhibiting properties similar to both metals and non-metals. Methyl and dimethyl selenide gases have been determined to be produced by anaerobic soils and sediments. Selenium is also taken up and concentrated by some plants which, in turn, may be ingested with ill effect by herbivorous or omnivorous animal species (Emsley, 2001). Selenium, however, is essential for life and participates in an endless cycle of transfer from soil and water to the atmosphere and back to soil and water. This chemical was detected at concentrations greater than ecological screening values in shallow soil and sediment and was, therefore, selected as an ecological COPC. Detected concentrations did not exceed human health risk screening value; therefore, it was not selected as a human health COPC.

Thallium (atomic symbol Tl) minerals are rare, although thallium (not listed in Table 5-2) is ten times more abundant than silver. This metal commonly appears in two different valence states: Tl (I) and Tl (III). When in the "+1" valence state, it has a behavior similar to potassium and sodium. Thallium salts are used in chemical research, and thallium sulfate has been banned as a pesticide in the Western countries (Emsley, 2001). Thallium is absorbed easily by plants, and the degree of absorption is proportional to the concentration of thallium in the associated soil. Tl (III) is easily reduced to Tl (I) (oxidation-reduction potential = 1.25 V). Thus, thallium is a strong oxidizer and the ease of reduction of Tl (III) explains its rarity in nature. Metallic ions of a single positive charge are characteristically mobile in the environment. Thus, thallium is not expected to be strongly bound to soils or sediments and it should move easily in aqueous media. LL-9 thallium concentrations did not exceed the screening levels for any medium and this metal was not selected as a human health or ecological COPC for LL-9.

Vanadium, atomic symbol V, has a very complex environmental chemistry. It exists as numerous species with an array of oxidation states, depending on ambient conditions. Generally, this element is expected to be immobile at neutral to high pH when redox potentials are negative. At moderately positive redox potentials (e.g., greater than 0.2 V), vanadium is expected to exist as soluble vanadate or other oxoanions at all but the lowest pH values. At pH values less than 4, this metal exists predominantly as positively charged oxocations (Dragun, 1988). LL-9 vanadium concentrations exceeded the human health screening values for surface water and sediments, and this metal was selected as a human health COPC for LL-9 in those media. It was selected as an ecological COPC for surface soil only because it exceeded the ecological risk-based screening value.

Zinc, atomic symbol Zn, is widely detected in worldwide soils at concentrations ranging from 5 to 770 mg/kg (Emsley, 2001). Soil concentrations have also been reported to vary from 10 to 300 mg/kg (Dragun, 1988). Plants can take up zinc from surrounding soils, the amount taken up



depending on the amount in the soil (Emsley, 2001). This essential nutrient is vital to proper functioning of enzymes. Zinc is only found in the environment as the divalent Zn (II) ion. At high pH (greater than 8), zinc precipitates as a hydroxide. This metal is moderately to very mobile in soils under most conditions (Table 5-2). The highest detected zinc concentration (780 mg/kg) was observed in sample LL9SS-068-0001-SO. The second-highest concentration was also observed in a surface soil sample, LL9SS-011-0001-SO at 711 mg/kg. This latter concentration was co-located with high concentrations of copper, mercury, and lead, which suggest that the zinc is related to those metals. Its relationship as a component of brass shell casings, along with copper, is understandable. The high lead may represent remnants of projectiles. The relationship of the zinc to mercury, however, is not understood. Zinc concentrations were less than human health screening criteria in all media; hence, this metal was not selected as a human health COPC for LL-9. Zinc was selected as an ecological COPC in surface soil and surface water because it exceeded the ecological risk-based screening values for those media.

5.3 Contaminant Transport Pathways

A contaminant transport pathway represents the physical path or the mechanism by which a contaminant moves or may move from one location (i.e., the source area) to another. A transport pathway may also involve a phase change for the contaminant (e.g., a contaminant is absorbed to soil, volatilizes to soil gas in the vadose zone, and then migrates into a basement as a gas). In addition, contaminant transport pathways provide mechanisms and conduits for contaminants to migrate to a new location where they may contribute to a human health or ecological risk. The determination of whether a pathway is currently causing a risk or could potentially cause a future risk depends on the combination of chemical characteristics, the existence of a potential pathway, the physical site conditions, and the potential for exposure to occur now or in the future.

This section presents a brief summary of contaminant fate and transport pathways that exist at LL-9. Based on the evaluation of existing conditions, the following potential contaminant transport pathways may exist:

- Leaching of soil contaminants to groundwater;
- Migration of groundwater contaminants within the soil and bedrock strata;
- Mixing of groundwater (i.e., creek bed seepage) with surface water in adjacent stream channels during storm events;
- Erosion and runoff of contaminated particles from soil and deposition in surface water bodies;
- Leaching of contaminants from creek sediment to surface water;



- Migration of contaminants in surface water as dissolved or sorbed phases during storm events; and
- Volatilization from soil, groundwater, or surface water.

An inspection of the integrity of the LL-9 sewer lines was not conducted as a part of this RI. Sumps connected to and upgradient of the sewers were removed. Sewers were sampled for sediment and water at readily accessible locations at the manholes. The integrity of the sewer lines is not completely known. These man-made structures typically represent potential contaminant migration pathways. The potential for migration has been reduced by the sump removal.

5.3.1 Leaching of Soil Contaminants to Groundwater

Contaminants that adhere to soil particles or have accumulated in soil pore spaces at LL-9 can leach and migrate vertically to the groundwater as a result of infiltration of precipitation. The rate and extent of this leaching is influenced by the amount of precipitation, rate of infiltration, the physical and chemical properties of the soil, the physical and chemical properties of the contaminant, and the depth of the water table.

During the selection of COPCs in soil (Section 6.2.3), the maximum detected concentrations of individual soil contaminants were compared against screening levels and installation background criteria (USACE, 2001b) concentrations. The soil screening levels (SSLs) included levels protective of groundwater. The SSLs are derived from assumptions regarding aquifer hydraulic conductivity, hydraulic gradient, infiltration rate, mixing zone height, aquifer thickness, source length, and chemical-specific soil-water K_d . These SSLs for protection of groundwater are used to conservatively estimate whether a chemical could leach from soil, migrate to groundwater, and raise the groundwater concentration above the pertinent risk-based screening level for groundwater.

Based on the results of LL-9 soil analyses, six metals-aluminum, arsenic, chromium, cobalt, manganese, and mercury-exhibited concentrations in excess of the soil-to-groundwater SSLs (Appendix V, Tables 2-1 and 2-2). However, only three of these chemicals (aluminum, cobalt, and manganese) were detected in groundwater samples. Hence, leaching of the other three chemicals (arsenic, chromium, and mercury) from soil to groundwater is not considered to be significant. Aluminum, cobalt, and manganese were present within background levels in subsurface soil and were only detected in surface soil above background levels in sporadic locations. This indicates that there is not a large source area in soil; and migration of aluminum, cobalt, and manganese from soil to groundwater is not significant.

The rates of metal leaching from soils to groundwater will not increase unless geological conditions, such as pH or oxidation-reduction potentials, are changed. This is an unlikely event because it involves changes to soils and bedrock that would require massive influxes of



chemical before a meaningful change could occur. If it were to occur, the reservoir of metals in the native minerals would vastly outweigh the amount of site-related contamination.

Aside from geological conditions, such as pH and oxidation-reduction potentials, a major factor affecting metal leaching rates is the length of time that the site has been non-operational. Metals are generally most mobile in soil soon after they are released. As time progresses, the assimilation of the metals into the mineral matrix renders them less mobile. If an excess of metals is not present beyond what the soil can readily bind, a steady state sets up. This steady state limits the leaching rates of the metals. Over time, the rates of leaching decrease as the excess metal reservoir is depleted and the more tightly bound metals are the only metals available for leaching. LL-9 has not been operational for about 60 years, which means that no new contaminants have been generated in the last 60 years and the previously released contaminants have had some time to bind to the mineral matrix. Thus, leaching rates can not be any greater than at the current time unless geologic conditions change.

A second factor is, with spatially localized exceptions, the relatively low concentrations of metal contaminants in soil compared to background concentrations. Using arsenic as the example, the site concentrations are only generally slightly greater than the background values with most values being less than the arsenic background value. The background concentrations for arsenic range from 15.4 to 19.8 mg/kg, depending on soil depth. The maximum arsenic concentration in any LL-9 soil sample was 32 mg/kg (sample LL9SB-019-0001-SO). This is less than two times the maximum background concentration. This may indicate that the arsenic is not a site contaminant or, if it is, that the total mass of arsenic contamination is not much greater than what is present naturally. However, arsenic is retained as a site-related contaminant evaluated by this report. This will further be discussed in the uncertainty section. The metals that had the greatest concentrations (i.e., copper, lead, mercury, and zinc), although much greater than normal background concentrations at the locations of highest contamination, also are generally comparable to background concentrations when the "hot spots" are discounted.

A third factor is the arsenic spatial distribution. Because many of LL-9 soil concentrations were less than the 19.8 mg/kg background value, the breadth of arsenic contamination is evidently limited to isolated locations. Furthermore, the isolated occurrences of screening value exceedances in the data available provide a very limited reservoir of contamination that could be transferred to groundwater. In addition, when the vertical profile of arsenic concentrations in soil is considered, it is clear that arsenic concentrations generally do not differ appreciably with soil depth (Tables 4-1 through 4-2). Instead, the concentration values, after allowing for natural soil sample heterogeneity, appear in the data available to be randomly distributed with depth over the entire observed concentration range up to 32 mg/kg in LL9SB-019-0001-SO. This kind of spatial distribution is often an indication that an apparent contaminant is not an actual contaminant. A similar situation exists in the lateral directions. It is an indication that the site soils are well leached and will not pose a significant risk to groundwater in the future. Of



course, copper, mercury, lead, and zinc at isolated locations are gross exceptions to this general pattern. Arsenic concentrations as high as 32 mg/kg, however, are not normally observed in soil so this one value and perhaps the six (four subsurface and two surface) others that exceed 19.8 mg/kg up to 26 mg/kg could represent true contamination. This may indicate that the arsenic is not a site contaminant or, if it is, that the total mass of arsenic contamination is not much greater than what is present naturally. However, arsenic is retained as a site-related contaminant evaluated by this report. This will be discussed further in the uncertainty section. Thus, despite one or a few isolated concentration values that may represent true contamination, the evidence is strong that LL-9 soils are in a steady-state condition whereby the rate of leaching of metals into the groundwater will not increase over time. Even the grossly contaminated location LL9SB-011-0001-SO is expected to represent a steady-state condition because of the length of time that has passed since the release.

The final factor is the observed groundwater metal concentrations in comparison to soil concentrations. All wells had the same arsenic concentration (0.73 µg/L), regardless of location. The arsenic detection limit was 0.73 µg/L, which is well below the background screening value for groundwater. Furthermore, the consistent detection of arsenic at this value is likely to be an indication that arsenic is not a site-related contaminant. The low concentrations indicate that arsenic is not a contaminant migration concern and that leaching of contaminants from soil to groundwater has not occurred appreciably. Thus, there does not appear to be a link between elevated soil concentrations and elevated groundwater concentrations to suggest that arsenic is leaching from soil.

This last factor indicates that the site is in a steady state in which there is no significant release of metals contamination except that which is available in the natural soils and bedrock. Variations in groundwater arsenic concentrations are likely to be caused by natural fluctuations such as changes in rainfall. The rates will not increase because contaminants are no longer being released to LL-9.

Although the factors discussed above focus on arsenic, similar arguments can be made for other metals whose concentrations may have exceeded soil screening levels at isolated locations. Even in wells downgradient of the most contaminated surface soil sample (LL9SS-011-0001-SO), the groundwater did not exhibit significantly elevated metal concentrations.

As indicated in Section 5.2.5, many metals detected in LL-9 soils were not selected as COPCs in groundwater. This is additional evidence that the soil-to-groundwater leaching pathway is of limited applicability. Furthermore, the leaching potential of metals is expected to decrease over time as the available metals are leached from soils and not replenished. Because the metal COPC concentrations are not currently much greater than background concentrations, however, it would be difficult or impossible to measure any such rates of decrease. The well that exhibited the highest metal concentrations (MW-004) is on the other side of LL-9 from the most contaminated soil location and is side-gradient to that soil location. Soil samples upgradient of



MW-004 did not exhibit elevated metal concentrations that correspond to the elevated groundwater metal concentrations. These observations are an indication that the detected groundwater metal concentrations are not well correlated to soil concentrations.

Thirteen of the analyzed metals were detected in groundwater. Essential human nutrients (magnesium, potassium, calcium, and sodium) are considered toxic only at very high doses and do not have screening levels referenced in this report. These nutrients were eliminated from consideration as COPCs.

This leaves aluminum, antimony, arsenic, barium, beryllium, cadmium, chromium, cobalt, copper, iron, lead, manganese, mercury, nickel, selenium, silver, thallium, vanadium, and zinc as environmentally significant metals detected in groundwater. The detections, however, are generally comparable to other RVAAP AOC upgradient concentrations or what are otherwise considered to represent concentrations in uncontaminated wells worldwide. For example, zinc, atomic symbol Zn, is an essential nutrient at low concentrations. The concentration of this element in worldwide soils is reported to range from 5 mg/kg to 770 mg/kg. It is taken up by plants in amounts that are proportional to the concentration in the surrounding soil. At levels in excess of 500 mg/kg, zinc can interfere with the uptake of other essential nutrients such as iron and manganese in plants (Emsley, 2001). Zinc only appears in the divalent oxidation state, Zn (II). The metal is mobile under most conditions although it forms hydroxide precipitates in aqueous solutions at pH values greater than 8 (Dragun, 1988). The maximum observed zinc concentration was 40 µg/L in sample LL9MW-003-0001-GW. The three other metals (copper, mercury, and lead) that were detected at high concentrations in surface soil sample LL9SS-011-0001-SO do not appear at concentrations in groundwater that reflect transfer of soil contamination to groundwater. For example, the worldwide zinc concentration is reported to range from less than 10 to 2,000 µg/L, the worldwide copper concentration range is less than 1.0 to 30 µg/L, and the worldwide lead concentration range is less than 15 µg/L. The highest observed concentrations of copper, lead, mercury, and zinc in LL-9 groundwater were 14.0 µg/L (LL9MW-004-0001-GW), non-detect, non-detect, and 40 µg/L (LL9MW-003-0001-GW), respectively.

Several organic chemicals [several different PAHs, nitrocellulose, nitroguanidine, RDX, bis(2-ethylhexyl)phthalate, and di-n-butylphthalate] were detected in soil but, of these, only nitrocellulose and bis(2-ethylhexyl)phthalate were detected in groundwater (Appendix V, Tables 2-1, 2-2, and 2-3). The majority of the nitrocellulose detections were qualified as estimated (J), or (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit, or both (B) (J). Therefore, the results may be false-positive detections, especially when the very low solubility of nitrocellulose is considered. The general lack of correlation between organic chemicals detected in soil and groundwater indicates the levels of these chemicals detected in soil do not present enough total mass to yield detectable groundwater concentrations after dilution, dispersion, and other natural mechanisms reduce their concentrations.



5.3.2 Migration of Groundwater Contaminants

Organic groundwater contaminants will alternately adsorb to and desorb from organic matter in the soil and bedrock at the molecular level as they migrate with groundwater radially away from LL-9. The radial movements are caused by the groundwater high located near the center of LL-9 (MW-003). Because general groundwater flow appears to be radial away from LL-9, contaminants in groundwater will ultimately flow in similar radial directions. Inorganic groundwater contaminants will follow a pathway similar to the organic contaminants. The relative migration rates will depend on individual contaminant adsorption characteristics and rates of degradation that may occur, as described in Sections 5.1 and 5.2.

The reservoir of organic contamination in soil that could feed groundwater is small, based on soil concentrations reported in Appendix V, Tables 2-1 and 2-2. Therefore, the level of importance of groundwater migration is expected to be minor. This is borne out by the low frequencies of organic contaminant detections for groundwater (Appendix V, Table 2-3).

Precipitation of metals as a mineral phase may physically transform contaminants. Contaminants may be chemically transformed through hydrolysis, oxidation/reduction, or biodegradation. Besides biodegradation, dilution from surface recharge is also a factor that causes the concentrations of contaminants to decrease in the downgradient directions.

Bis(2-ethylhexyl)phthalate was detected at low concentration (4.3 µg/L) in one out of seven wells; hence, this contaminant is not viewed to be an important LL-9 contaminant. However, migration would follow the same path as the groundwater with a retardation of the bis(2-ethylhexyl)phthalate as it alternately adsorbs to, and desorbs from, soil and bedrock.

Nitrocellulose was detected in nine soil samples, five groundwater samples, one sediment sample, and one surface water sample scattered across LL-9. Most soil detections occurred in the central portion of the site near the locations of former operational buildings. Nitrocellulose is relatively insoluble, so it is expected to persist in soil and sediment for a long time. Half-life information could not be found for this chemical. Therefore, groundwater dissolution of nitrocellulose followed by groundwater transport does not appear to be a significant transport pathway.

Metals in groundwater will follow groundwater flow patterns just as the organic contaminants do. The metals may migrate in soluble form or as ions or non-ionized organometallic complexes bound to colloidal particles. Migration of soluble metals would be retarded by ion exchange with bedrock, and the rate of colloidal flow through the bedrock would depend on particle size with smaller particles moving more quickly than larger particles. In general, highly charged (+3 or greater) metals and complexes will be retarded to the greatest degree, and weakly charged species (+1) will be the most mobile. Lead, chemical symbol Pb, is a notable exception. This metal binds tightly to soil as the Pb^{+2} ion, and is essentially immobile despite its moderate



charge of "+2." This ion also binds tightly to sediments but is mobile when the sediments, themselves, migrate.

As explained in Section 5.3.1, however, the amount of metals leaching into groundwater and, hence, the migration of groundwater metal contaminants, will remain steady or decrease over time relative to the current concentrations.

5.3.3 Mixing of Groundwater (i.e., Creek Bed Seepage) with Surface Water in Adjacent Stream Channels During Storm Events

Groundwater may recharge nearby drainage channels, but knowledge of site conditions suggests that the primary source of drainage channel water is precipitation. Any contaminants that are in the groundwater would be transported to the creek with the groundwater. Rates of transport cannot be predicted; but because the concentrations of organic and metal contaminants are low in LL-9 groundwater, the rates are not significant. Precipitation tends to further dilute groundwater seepage that enters the drainage channels.

5.3.4 Migration of Contaminants from Surface Soil to Surface Water

Rainfall, snowmelt, and surface water runoff that come into direct contact with surface soils can leach contaminants from the soils and transport them to drainage channels via runoff during storm events. Soil particles containing sorbed contaminants can also be dislodged from the soil surface and be physically transported to the creek via overland runoff. The topographic relief at LL-9 is minor, so the amount of overland runoff from this site is expected to be very small.

Nitrocellulose, was detected at an estimated concentration 0.12 (J)(B) mg/L in sample LL9SW-012-0001-SW and 0.1 (J)(B) mg/L in sample LL9SW-012-0001-DUP, was the only organic chemical detected in surface water. However, the results were qualified as estimated (J) and (B) the concentration of nitrocellulose in the method blank was detected above the reporting limit. These two detections at low concentration indicate that surface runoff is not a significant migration pathway. Nitrocellulose, which is not expected to be soluble to any significant degree in surface water, will be transported as suspended particulates. The rate of transport is expected to be small because the reservoir of nitrocellulose is small. The detection of nitrocellulose cited here may be a false detection, based on its similarity to the analytical detection limit and the known limited solubility of nitrocellulose in water.

5.3.5 Leaching of Contaminants from Creek Sediment to Surface Water

The reservoir of organic and inorganic contamination in drainage channels and creek sediments is low and the source of water that LL-9 drainage channels receive is precipitation. Therefore, the opportunity for transfer from creek sediments to surface water is limited by the frequency of rainfall and snowmelt, and this insignificant migration pathway is not discussed further.



5.3.6 Migration of Contaminants in Surface Water as Dissolved or Sorbed Phases During Storm Events

Once contaminants are dislodged from surface soil particles, some of which then become dissolved, they may flow with surface water and entrained sediments to downgradient areas. The masses and concentrations of organic and inorganic contamination available in surface soils and sediments are generally low, with the notable exception of the high metal concentrations at LL9SS-011-0001-SO in surface soil. This location is near the western side of the site where the terrain becomes relatively steep further to the west. There are no clear drainage channels in the immediate vicinity of that location, however, so the erosion potential is concluded to be relatively low. Erosion is also limited by the grass cover in this area. Therefore, while the drainage channels represent potential migration pathways, it does not appear that these migration pathways are significant as a means of transporting contaminants.

5.3.7 Volatilization from Soil, Groundwater, or Surface Water

All detected compounds in soil were SVOCs or other relatively non-volatile chemicals; therefore, volatilization from soil in an insignificant transport pathway. A similar situation exists for the other media. Therefore volatilization from any of these media is considered to be insignificant. This is further supported by the generally small reservoir of contamination present in any of the investigated media. When small masses of contaminants are coupled with slow rates of release, the migration pathway of interest is insignificant. Where mercury concentrations were very high (i.e., LL9SS-011-0001-SO) transformation of mercury to methyl or ethyl mercury compounds is possible with subsequent evaporation from the soil. However, this is most likely to occur under anaerobic conditions, and the high mercury concentrations were detected in shallow soil where anaerobic conditions are not likely. Therefore, even this transformation pathway followed by volatilization is not expected to be significant. The rates of such transformation are low enough that such transformations would not be expected to result in the release of hazardous quantities of these very toxic organomercury compounds chemicals.

5.4 Conceptual Site Model

The CSM is a description based on known site conditions that explains in a conceptual manner how contaminants were or could have been deposited, how these contaminants can or do move in the environment, and the impact they may have on receptors. The focus of this section is on the physical model. Effects and potential effects on environmental organisms, including humans, are discussed in the risk assessment sections.

From 1941 to 1945, LL-9 operated to produce detonators. In 1945, the load line was deactivated, and the equipment was removed. There are no documented activities at LL-9 since closure in 1945. Primary explosives handled in detonators (Lead Azide/Mercury Fulminate) were present in smaller quantities. Secondary high explosives of interest included



tetryl, and possibly others; propellants of interest include nitroglycerine, nitrocellulose, and nitroguanidine. Some of these contaminants and others were detected in several environmental media, although at low concentrations that represent small total masses of contaminants sparsely scattered across the site:

- There were analytical detections of metals, SVOCs, energetic compounds, and some miscellaneous inorganic chemicals. Metal contaminants were detected throughout the site, but primarily in the immediate vicinity of the production buildings. Organic compounds were also detected throughout the site. Nitrocellulose, which has an expected long half-life in the environment, was detected in several samples; but the detections appear to be false detections.
- In soils, metals, nitrocellulose, nitroguanidine, RDX, and SVOCs were detected throughout the site including the periphery areas. Nitroguanidine was detected at the far western edge of the site in surface soil only. When detected, these organic chemicals were generally detected at the greatest concentrations in surface soil, which is consistent with an aerial deposition mechanism.
- In sediment, metals, nitrocellulose, several PAHs, and three phthalate esters were detected. The distribution is similar to the distribution in soil: concentrations are generally higher in surface soils than subsurface soils
- In surface water sampling locations, only metals were detected at concentrations exceeding either a Region 9 tap water PRG or an RVAAP installation background concentration. No pattern of distribution could be discerned.
- Exceedances of RVAAP installation background and/or Region 9 tap water PRG values were reported in the seven groundwater samples. Analytes included metals and one detection of bis(2-ethylhexyl)phthalate. Monitor well MW-007, located downgradient of the production area, had exceedances on inorganics only. Nitrocellulose was reported as detected, but the reported detections appear to be false detections and there are no screening criteria for this operations-related compound.

The concentrations of contaminants are generally low, with a notable exception being a localized spot of high metal concentrations for copper, mercury, lead, and zinc at LL9SS-011-0001-SO in surface soil.

Sumps might have been a primary contamination source from operations water that was routinely diverted to them; however, sumps were removed during the LL-9 interim remedial action. Sewer lines may provide a contaminant transport pathway.

Contamination at LL-9 is sparsely dispersed across the site with some elevated chemical concentrations occurring near operational activities. Most metals appear to be fairly uniformly distributed across the site, with exception of location LL9SS-011-0001-SO as noted above.



Metals do not degrade in the environment, but they can be assimilated into minerals. Their oxidation states can change over time as they migrate from one location to another. Metals in general, however, are relatively immobile except for those with predominantly single positive charges, such as sodium and potassium. A few other metals, such as calcium and magnesium, are also relatively mobile. These four metals, however, generally pose little to no environmental risks to receptors. Receptor exposure and risk are treated more completely in the risk assessment sections (6.0 and 7.0).

Few organic chemicals that are directly related to site operations were selected as COPCs, indicating that the concentrations of detected organic chemicals are generally low enough not to pose a significant threat to the health of receptors.

The primary contaminant migration pathways for contaminants at LL-9 are:

- Leaching from soils to groundwater,
- Leaching from sediments to surface water, and
- Transport in surface drainage channels.

Leaching from soils would be expected to occur around sumps if contaminated soil remained in place after the sump removals. This is not expected to be a significant concern. The sump removal is complete. Given that the quantities of contamination present at LL-9 are generally low elsewhere, even if contaminated soil is in place from the sump removal, the mass of contaminants transported by this migration pathway is expected to be low. The sumps were holding water and appeared to be structurally intact. No staining was observed. Dilution effects should be large in drainage channels. This is reflected in the low concentrations of contaminants detected in surface waters. Similar effects are observed in groundwater. Nitrocellulose, which has evidently been widely distributed in surface and subsurface soils at low concentrations across LL-9, is notably recalcitrant to dissolution; hence, its persistence is expected to be long.

Any soil or sediment contaminants that leach into groundwater would flow radially away from the AOC based upon groundwater elevation mapping. Each contaminant will be retarded in its movement. The degree of retardation depends largely on the adsorption tendency of each contaminant and the amount of organic material in the soils and bedrock.

Surface water drainage channels can transport surface soil contaminants and sediments to downgradient locations; however, the topographical relief is moderate at LL-9 except at the LL-9 peripheries. Therefore, transport pathways are not expected to move sediments rapidly. An exception to this could be during storm events when flow rates increase significantly. Contaminants detected at LL-9 will tend to adsorb to sediments.



1 In summary, the low concentrations and small total masses of energetic compounds at LL-9 are
2 consistent with good health and safety practices. In particular, primary explosives, such as
3 metal azides (e.g., lead azide) and fulminates (e.g., mercury fulminate), which are very unstable
4 with respect to physical shock, would not be expected to be released indiscriminately to non-
5 operational areas. To do so would result in extreme safety hazards. The small reservoir of
6 contamination in soil provides little total contaminant mass for migration to groundwater or other
7 media. The metals will persist and the organics will degrade over time, albeit some of them
8 (e.g., nitrocellulose) will do so slowly. Given the low concentrations observed at this site,
9 modeling to estimate the degradation rates would not be fruitful. SESOIL and AT123D modeling
10 were not performed for six metals (aluminum, arsenic, chromium, cobalt, manganese, and
11 mercury) because aluminum, chromium, and manganese had concentrations that exceeded
12 background less than 5% of the time, arsenic exceeded 26% and mercury 44%, but neither of
13 these were detected in groundwater, indicating that little or no leaching is taking place. Cobalt
14 exceeded background 18% of the time. Maximum concentrations of these metals were
15 generally less than three times background, and the total volume of any contaminated soil is so
16 small, that impacts on groundwater would not be measurable, either through a modeling
17 exercise or through actual groundwater monitoring.



6.0 BASELINE HUMAN HEALTH RISK ASSESSMENT

This section presents the results of the Baseline Human Health Risk Assessment (BHHRA) conducted to characterize the risks to humans associated with potential current or future exposures to chemicals in surface soil, subsurface soil, groundwater, surface water, and sediment within LL-9 at the RVAAP, Ravenna, Ohio. This section also provides a list of chemicals of concern (COCs) for further evaluation by the risk managers.

6.1 Introduction

As described in previous sections, LL-9 is located at the intersection of Fuze and Booster Roads and George Road in the south-central region of RVAAP and consists of approximately 69 acres. LL-9 was utilized primarily for the production of detonators from 1941 to 1945. Site features include a gravel road around the former main production area and a water tower that is no longer connected to a water distribution system. Buildings formerly located at the site were previously thermally decontaminated and demolished (MKM, 2003a).

The methods used in the LL-9 BHHRA are based primarily on the protocol established in the RVAAP *Facility-Wide Human Health Risk Assessors Manual (FWHHRAM)* (USACE, January 2004.) Additional protocol is summarized in the *White Paper – Human Health Risk Assessment Approach for Load Lines 6, 9, and 11 Remedial Investigations, Ravenna Army Ammunition Plant, Ravenna, Ohio* (TtNUS, April 2004). A copy of the White Paper is included in Appendix X. Technical guidance discussed during March 2004 teleconferences with the USACE and Ohio EPA was also considered during preparation of the BHHRA.

The methods for the BHHRA consist of the following six steps:

- Step 1 – Data evaluation – Select site-related chemicals (SRCs) and chemicals of potential concern (COPCs) for environmental media at LL-9. The COPCs identified in Step 1 are evaluated quantitatively (when possible) and discussed qualitatively otherwise. Data evaluation is further discussed in Section 6.2.
- Step 2 – Exposure assessment – Identify potential current and future human receptors that may be exposed to the COPCs, evaluate the pathways by which the receptors may be exposed, and the estimate chemical intake resulting from exposure.
- Step 3 – Toxicity assessment – Assess toxicity of each COPC including the presentation of the toxicity criteria (i.e., cancer slope factors, reference doses) typically used by USEPA to estimate risk.



- Step 4 – Risk characterization – Estimate cancer and non-cancer risk using the intakes and toxicity criteria listed in the exposure assessment and toxicity assessment, respectively.
- Step 5 – Uncertainty analysis – Specify major uncertainties affecting the interpretation of the BHHRA results.
- Step 6 – Summary and Conclusions – Summarize and present the conclusions of the BHHRA.

The results of the BHHRA are used to (1) document and evaluate risks to human health; (2) determine the need, if any, for remedial action; and (3) identify COCs. COCs are those chemicals determined by the BHHRA to pose (or significantly contribute to) a potential risk to human receptors in excess of target risk benchmarks. The development of chemical-specific remediation levels may be required for these chemicals. Per the protocol established in the White Paper for LL-9 (TtNUS, 2004), potential COCs were identified for an environmental medium when the receptor risk exceeded a cumulative cancer risk benchmark of 1×10^{-5} or a total non-cancer hazard index (HI) benchmark of 1.

Three major conditions of chemical contamination and environmental fate and transport are considered when evaluating potential risks: (1) contaminants with toxic characteristics must be found in environmental media and must be released by either natural processes or by human action; (2) potential exposure points must exist; (3) human receptors must be present at the points of exposure. If any of the conditions listed above is absent, risks are not quantitatively evaluated for that exposure pathway.

The quantitative BHHRA for LL-9 includes tables based on RAGS Part D (USEPA, 2001a) and RVAAP specifications.

6.2 Data Evaluation

Data evaluation involves the gathering, evaluation, and organization of data into a form that is appropriate for a baseline risk assessment. The data gathering step involves the compilation of all available data and the sorting of data by medium. The quality of the data is evaluated with respect to qualifiers and codes applied as a consequence of the data validation process and via other quality measures (e.g., the adequacy of the analytical methods used in the site investigation, the adequacy of the sample quantitation limits, etc). COPCs are selected for evaluation in the BHHRA using protocols that include a comparison of site concentrations to facility background concentrations and to conservative toxicity screening concentrations. Section 6.2.1 describes the data sets compiled for use in the BHHRA and the data validation process used to evaluate the quality of the data. Section 6.2.2 summarizes the protocols for and the results of the COPC selection process.



6.2.1 Data Collection and Evaluation

The data used in the BHHRA consisted of the following:

- Analytical data available for soils, groundwater, surface water, and sediment samples collected at LL-9. The data sets evaluated are discussed in detail in Section 6.2.3. The data are the result of the environmental investigations described in Section 3.
- Surface soil was defined as soils collected from 0–1 ft bgs, and subsurface soil was defined as soils collected from depths greater than 1 ft bgs. The deep surface soil data set was composed of surface soil data and subsurface soil data from samples collected 1–4 ft bgs. For groundwater, filtered samples were collected and analyzed for TAL metals, per State of Ohio's directions. Unfiltered samples were collected and analyzed for volatile and semi-volatile compounds. The analytical laboratory selected the most appropriate analytical result if reanalysis/dilutions were necessary, and that result was evaluated in the determination of the SRCs for this risk assessment.
- Some data originally collected as part of the RI were not used in the BHHRA. For example, buildings and sumps were removed from LL-9; therefore, samples collected from these areas removed during the excavations were not used in the BHHRA. Consequently, the data sets evaluated in the BHHRA represent current conditions at LL-9.

Chemicals were analyzed and assessed in accordance with Louisville Chemistry Guidance. The analytical results were evaluated, using the National Functional Guidelines (USEPA, 1994a, and 1994b) to assess data usability and the laboratory's compliance with the analytical methods. The analytical data were reviewed, validated, and evaluated using the criteria specified in the data quality objectives. Validated data (and qualifiers) are included in Appendices F, G, J, L, and N. Unqualified positive detections and "J" qualified detections (estimated values) were considered as detected concentrations for this BHHRA. All non-detect results (indicated with a "U" qualifier) were retained in the BHHRA data set. None of the analytical data evaluated in the BHHRA were rejected as a consequence of the data validation process. A complete data quality summary is included in Appendix U.

The data evaluation included the calculation of basic descriptive statistics for each data set evaluated in the BHHRA. Basic statistics included frequency of detection, range of positive detections, arithmetic mean, normal 95% upper confidence level (95% UCL-N) on the mean, and log-normal 95% upper confidence level (95% UCL-L) on the mean. Appendix V provides the equations used to determine the 95% UCL-L and the 95% UCL-N on the mean. These descriptive statistics were also used in the determination of exposure point concentrations (EPCs) selected in the exposure assessment.



6.2.2 Selection of COPCs

COPCs for LL-9 were identified in two steps. First, SRCs were identified for each medium, then a toxicity screen was conducted to determine which SRCs would be selected as COPCs. The purpose of the screening process was to eliminate chemicals for which no further risk evaluation is needed. The premise of this screening step is that risk is typically dominated by a few chemicals and that, although dozens may actually be detected, many chemicals may contribute minimally to the total risk. Section 6.2.2.1 describes the SRC selection process, and Section 6.2.2.2 describes the COPC screening process.

6.2.2.1 SRC Screens

The following assumptions were made in the determination of the LL-9 SRCs:

- Physical chemistry data (e.g., alkalinity, pH, etc.) are not considered to be SRCs (and, therefore, are not considered to be COPCs) for LL-9.
- Groundwater samples were filtered in the field. Filtered metals data were used for TAL metals to determine inorganic groundwater SRCs. Filtered data are typically more indicative of the soluble or dissolved, and therefore, more mobile chemical concentrations in groundwater. Unfiltered data were used for volatiles and semi-volatile SRCs.
- Soil data were subdivided into three data sets based on the sampling depths for LL-9 and the receptors to be evaluated at LL-9. In accordance with the FWHHRAM, the surface soil data set is comprised of data for samples collected from the 0–1 ft bgs interval. The deep surface soil data set is composed of soil samples collected within the 0–4 ft bgs interval. Subsurface soil data is composed of sample results for soil samples collected below 1 ft bgs. Consequently, data from the surface soil data set were compared to the surface soil RVAAP background criteria, and data from the subsurface soil data set were compared to the subsurface soil background criteria. The chemical concentrations in the deep surface soil samples were compared to the lower of the RVAAP background benchmarks available for the background surface and subsurface soil data sets.

The LL-9 SRC screening process (a process by which chemicals are eliminated from further consideration in the BHHRA) involved two steps: comparing results to already established, RVAAP-specific background concentrations and weight-of-evidence screening. The background comparison was conducted to determine whether the metals detected in the environmental media were naturally occurring or potentially site-related. Per the protocol established in the White Paper (TtNUS, 2004), the RVAAP-specific background values are those presented in the *Phase II Remedial Investigation Report for the Winklepeck Burning Ground at Ravenna Army Ammunition Plant, Ravenna, Ohio* (USACE, 2001b). These values



are either the 95% upper tolerance limits (UTLs) for the background dataset for each environmental media, or they are the maximum detected concentrations if the calculated UTLs exceed the maximum detected concentrations reported for the background samples. An inorganic was selected as a potential SRC if the maximum detected concentration exceeds the RVAAP background value. This background screen was applied to inorganic chemicals only; organic chemicals were not eliminated as SRCs based on background comparisons.

Two weight-of-evidence screens (a frequency-of-detection screen and a screen for naturally occurring essential elements) were also used as follows to eliminate chemicals from further consideration in the BHHRA:

- If a data subset contained 20 or more data points, a frequency-of-detection screen was applied. Chemicals with a frequency of detection of less than 5% of the samples analyzed in any medium may not be considered to be site-related for that medium if:
 - 1) The chemical was not detected at high concentrations relative to the detection limit or to the established background values;
 - 2) The chemical was not detected in another medium; and
 - 3) Knowledge of LL-9's history or processes did not suggest that the chemical would be present. For example, any explosive or propellant detected in any LL-9 medium was considered to be site-related regardless of frequency of detection.
- Naturally occurring essential elements, including calcium, chloride, iodine, iron, magnesium, potassium, phosphorous, and sodium, are typically considered to be toxic only at high concentrations and are typically not selected as SRCs. As detailed in the FWHHRAM, these metals are an integral part of the human food supply and are often added to food as supplements. USEPA recommends that these chemicals not be evaluated as COPCs as long as they are: (1) present at concentrations only slightly elevated above naturally occurring levels and (2) only toxic at doses much higher than those that could be associated with contact at a site. Recommended daily allowance (RDA) and recommended daily intake (RDI) values are available for seven of these metals. Based on the RDA/RDI values, a receptor ingesting 100 mg of soil per day would receive less than the RDA/RDI of calcium, magnesium, phosphorous, potassium, and sodium, even if the soil consisted of the pure mineral (i.e., soil concentrations greater than 1,000,000 mg/kg). Receptors ingesting 100 mg of soil per day would require soil concentrations of 1,500 mg/kg of iodine and 100,000 to 180,000 mg/kg of iron to meet their RDA/RDI for these metals. Concentrations of these metals in the environmental media at LL-9 do not exceed these levels. Consequently, these essential nutrients were not selected as potential SRCs.



The results of the SRC screen are shown for each medium in Appendix V, Tables 2.1 through 2.6.

6.2.2.2 COPC Toxicity Screen

After the SRCs were identified, they were further evaluated using a toxicity screen to develop a list of human health COPCs. For this risk assessment, risk-based and health-based criteria were used to reduce the number of chemicals and exposure routes considered in the LL-9 BHHRA. More importantly, the screening focused the risk assessment on those chemicals most likely to be significant in terms of risk. COPCs were defined as chemicals that were positively detected in an environmental medium at a maximum concentration exceeding screening values.

Environmental sampling results were compared to risk-based screening concentrations (RBSCs) based on USEPA Region 9 residential PRGs (USEPA, 2004c) and, for groundwater only, federal Safe Drinking Water Act (SDWA) Maximum Contaminant Levels (MCLs). The USEPA Region 9 risk-based residential PRGs represent a risk level of 1×10^{-6} for carcinogenic effects (i.e., a 1 in 1,000,000 excess chance of developing cancer over a lifetime) and an HI of 1.0 for non-carcinogenic effects (i.e., adverse non-carcinogenic health effects are not anticipated at or below this exposure concentration). The RBSCs represent a risk level of 1×10^{-6} for carcinogenic effects and an HI of 0.1 to account for additive non-carcinogenic effects. The Region 9 residential PRGs for soil were calculated for a human receptor hypothetically exposed to chemicals in soil assuming a residential land use scenario. The Region 9 PRGs for tap water were calculated for a human receptor hypothetically using a groundwater or surface water resource as a domestic water supply. Conservatively, RBSCs for non-carcinogens are one-tenth the Region 9 PRGs to further account for potential cumulative non-carcinogenic effects. The RBSCs for carcinogens are the Region 9 residential PRGs and represent the 1×10^{-6} cancer risk level. If RBSCs exist for a chemical for both carcinogenic and non-carcinogenic effects, the lower of the two values was used as the COPC selection criterion.

The maximum detected concentration for each SRC in each medium was compared to the risk based screening concentration as follows:

- Chemicals in surface soil, subsurface soil, and sediment were screened against their respective RBSCs. The RBSCs used were the Region 9 residential soil PRGs that reflect residential land use. Data from these media were also compared to their respective industrial soil PRGs; that comparison is presented for informational purposes only.
- Chemicals in groundwater were screened against the lesser of the RBSCs based on the Region 9 tap water PRGs and Federal SDWA MCLs.
- Chemicals in surface water are screened against the RBSCs based on the Region 9 tap water PRGs.



Region 9 PRGs can be found at <http://www.epa.gov/region09/waste/sfund/prg/index>. Drinking water MCLs were obtained from <http://www.epa.gov/safewater/mcl>.

SRCs that exceeded toxicity screening levels and those lacking screening levels were retained as COPCs. Conversely, chemicals that were not detected at maximum concentrations exceeding toxicity screening levels were not retained as COPCs.

If USEPA-approved toxicity information, such as cancer slope factors or reference doses, is available, the COPCs are then classified as quantitative COPCs. If no toxicity information is available, the COPCs are considered to be qualitative COPCs. The qualitative COPCs identified for LL-9 were further evaluated in the uncertainty section of the BHHRA (Section 6.6.1). The quantitative COPCs are further discussed in the toxicity assessment section (Section 6.4). Toxicity profiles are presented for all COPCs in Section 6.4.

Appendix V, Tables 2.1 through 2.6, present the COPC screens for surface soil, deep surface soil (0- to 4-ft interval), subsurface soil, groundwater, surface water, and sediment, respectively. These tables include the following:

- Summary statistics, including the frequency of detection, range of non-detected values, maximum and minimum detected concentrations, location of maximum concentration, arithmetic average concentration, and upper 95% UCL on the mean concentration;
- Screening values (RVAAP background criteria [USACE, 2001b], RBSCs, and MCLs, as appropriate);
- SRC determination; and
- COPC determination.

The data sets evaluated for each environmental medium are presented in Tables 6-1 through 6-6. Table 6-7 summarizes the COPC list for each medium and indicates which COPCs will be addressed quantitatively and qualitatively for each medium.

6.2.2.3 Screening for Lead

USEPA-approved toxicity criteria (i.e., cancer slope factors and reference doses) have not been published for lead. Consequently, a calculated, toxicity criteria-based PRG is not available for this element. For this risk assessment, lead concentrations in soil and sediment at LL-9 (see Appendix V, Tables 2.1, 2.2, 2.3, and 2.6) were compared to the lead soil screening guidance concentration of 400 mg/kg for residential soil published in OSWER Directive #9355.4-12 (USEPA, 1994c). This value is presented as the residential PRG in the USEPA Region 9 table. Lead was not detected in sediment, subsurface soil, or groundwater at LL-9 at concentrations greater than the lead soil screening guidance concentration of 400 mg/kg for residential soil.



Lead was detected in surface water, and the SDWA action level of 15 µg/L was used as the screening level.

6.2.2.4 COPC Screening Assumptions

The following assumptions have been made regarding the use of PRGs in this BHHRA:

- LL-9 environmental samples were analyzed for total chromium. The maximum detected concentrations for chromium were screened against the Region 9 residential PRGs for hexavalent chromium. This is a conservative screening approach, because hexavalent chromium is more toxic than trivalent chromium and is a less commonly occurring form of the metal.
- As recommended by Ohio EPA, the Region 9 Cal-modified PRG for trichloroethene was used as the basis for the RBSC for trichloroethene.

6.2.3 COPCs for LL-9

Sections 6.2.3.1 through 6.2.3.6 identify LL-9 COPCs in surface soil, deep surface soil (i.e., the 0–4 ft interval), subsurface soil, groundwater, surface water, and sediment, respectively. Arsenic was determined to be a COPC in all media at LL-9. Antimony and manganese were selected as COPCs only for groundwater. Carcinogenic PAHs (benzo[a]pyrene and dibenzo[a,h]anthracene) were determined to be a COPC only in surface soil and deep surface soil.

6.2.3.1 Surface Soil

Analytical data available for 53 surface soil samples were evaluated in the BHHRA. Twenty-two metals, 19 SVOCs, nitroguanidine, and nitrocellulose were detected in the surface soil samples collected at LL-9. The samples evaluated in this BHHRA are listed in Table 6–1. The results of COPC selection for surface soils are summarized in Appendix V, Table 2.1. The following chemicals were identified as surface soil COPCs:

- Metals (aluminum, arsenic, chromium, copper, lead, manganese, and mercury);
- Organics (benzo[a]pyrene, and dibenzo[a,h]anthracene); and
- Nitrocellulose.

One aluminum detection, two manganese detections, and four chromium detections out of 53 exceeded their respective background concentrations; therefore, these metals were selected as COPCs.



Table 6-1
Human Health Risk Assessment Dataset for Surface Soils

Sample ID	Depth bgs*
LL9SS-001	0-1
LL9SS-002	0-1
LL9SS-003	0-1
LL9SS-004	0-1
LL9SS-005	0-1
LL9SS-006	0-1
LL9SS-007	0-1
LL9SS-008	0-1
LL9SS-009	0-1
LL9SS-010	0-1
LL9SS-011	0-1
LL9SS-012	0-1
LL9SS-013	0-1
LL9SS-014	0-1
LL9SS-015	0-1
LL9SS-016	0-1
LL9SS-017	0-1
LL9SS-018	0-1
LL9SS-019	0-1
LL9SS-020	0-1
LL9SS-021	0-1
LL9SS-022	0-1
LL9SS-023	0-1
LL9SS-024	0-1
LL9SS-025	0-1
LL9SS-026	0-1
LL9SS-027	0-1
LL9SS-028	0-1
LL9SS-029	0-1
LL9SS-030	0-1
LL9SS-031	0-1
LL9SS-032	0-1
LL9SS-033	0-1
LL9SS-034	0-1
LL9SS-035	0-1
LL9SS-036	0-1
LL9SS-037	0-1
LL9SS-038	0-1
LL9SS-039	0-1
LL9SS-040	0-1
LL9SS-041	0-1
LL9SS-042	0-1
LL9SS-043	0-1
LL9SS-044	0-1
LL9SS-045	0-1
LL9SS-046	0-1
LL9SS-047	0-1
LL9SS-068	0-1



Table 6-2
Human Health Risk Assessment Dataset For Subsurface Soils

Sample ID	Depth bgs*
LL9SB-001	1 - 3
LL9SB-002	1 - 3
LL9SB-003	1 - 2
LL9SB-004	1 - 3
LL9SB-005	1 - 3
LL9SB-006	1 - 2
LL9SB-007	1 - 3
LL9SB-008	1 - 3
LL9SB-009	1 - 3
LL9SB-010	1 - 2.5
LL9SB-012	1 - 3
LL9SB-013	1 - 3
LL9SB-014	1 - 3
LL9SB-015	1 - 3
LL9SB-016	1 - 3
LL9SB-017	1 - 3
LL9SB-018	1 - 3
LL9SB-019	1 - 3
LL9SB-020	1 - 3
LL9SB-021	1 - 2.5
LL9SB-022	1 - 3
LL9SB-023	1 - 3
LL9SB-024	1 - 3
LL9SB-025	1 - 3
LL9SB-026	1 - 3
LL9SB-027	1 - 3
LL9SB-028	1 - 3
LL9SB-029	1 - 3
LL9SB-030	1 - 3
LL9SB-031	1 - 3
LL9SB-032	1 - 3
LL9SB-040	6.5 - 6.5
LL9SB-041	7 - 7
LL9SB-042	8 - 10
LL9SB-043	9 - 11
LL9SB-044	8 - 10
LL9SB-045	8 - 10
LL9SB-046	4 - 6
LL9SB-047	3 - 5
LL9SB-048	1 - 3
LL9SB-049	1 - 3
LL9SB-050	1 - 3
LL9SB-051	1 - 3
LL9SB-052	1 - 3
LL9SB-053	1 - 3
LL9SB-055	1.5 - 3.5
LL9SB-056	2 - 4
LL9SB-059	3 - 5
LL9SB-061	1 - 3
LL9SB-065	3 - 5
LL9SB-066	5 - 7
LL9SB-069	7 - 7
LL9SB-070	6.5 - 6.5



Table 6-3
Human Health Risk Assessment Dataset For Deep Surface Soils

Sample ID	Depth bgs*
LL9SS-001	0-1
LL9SS-002	0-1
LL9SS-003	0-1
LL9SS-004	0-1
LL9SS-005	0-1
LL9SS-006	0-1
LL9SS-007	0-1
LL9SS-008	0-1
LL9SS-009	0-1
LL9SS-010	0-1
LL9SS-011	0-1
LL9SS-012	0-1
LL9SS-013	0-1
LL9SS-014	0-1
LL9SS-015	0-1
LL9SS-016	0-1
LL9SS-017	0-1
LL9SS-018	0-1
LL9SS-019	0-1
LL9SS-020	0-1
LL9SS-021	0-1
LL9SS-022	0-1
LL9SS-023	0-1
LL9SS-024	0-1
LL9SS-025	0-1
LL9SS-026	0-1
LL9SS-027	0-1
LL9SS-028	0-1
LL9SS-029	0-1
LL9SS-030	0-1
LL9SS-031	0-1
LL9SS-032	0-1
LL9SS-033	0-1
LL9SS-034	0-1
LL9SS-035	0-1
LL9SS-036	0-1
LL9SS-037	0-1
LL9SS-038	0-1
LL9SS-039	0-1
LL9SS-040	0-1
LL9SS-041	0-1
LL9SS-042	0-1
LL9SS-043	0-1
LL9SS-044	0-1
LL9SS-045	0-1
LL9SS-046	0-1
LL9SS-047	0-1
LL9SS-068	0-1
LL9SB-001	1-3



Table 6-3
Human Health Risk Assessment Dataset For Deep Surface Soils

Sample ID	Depth bgs*
LL9SB-002	1-3
LL9SB-003	1-2
LL9SB-004	1-3
LL9SB-005	1-3
LL9SB-006	1-2
LL9SB-007	1-3
LL9SB-008	1-3
LL9SB-009	1-3
LL9SB-010	1-2.5
LL9SB-012	1-3
LL9SB-013	1-3
LL9SB-014	1-3
LL9SB-015	1-3
LL9SB-016	1-3
LL9SB-017	1-3
LL9SB-018	1-3
LL9SB-019	1-3
LL9SB-020	1-3
LL9SB-021	1-2.5
LL9SB-022	1-3
LL9SB-023	1-3
LL9SB-024	1-3
LL9SB-025	1-3
LL9SB-026	1-3
LL9SB-027	1-3
LL9SB-028	1-3
LL9SB-029	1-3
LL9SB-030	1-3
LL9SB-031	1-3
LL9SB-032	1-3
LL9SB-047	3-5
LL9SB-048	1-3
LL9SB-049	1-3
LL9SB-050	1-3
LL9SB-051	1-3
LL9SB-052	1-3
LL9SB-053	1-3
LL9SB-055	1.5-3.5
LL9SB-056	2-4
LL9SB-059	3-5
LL9SB-061	1-3
LL9SB-065	3-5



Table 6-4

Human Health Risk Assessment Data Set for Groundwater

Sample ID	Depth bgs*
LL9MW-001	15
LL9MW-002	21
LL9MW-003	18
LL9MW-004	17
LL9MW-005	15
LL9MW-006	15
LL9MW-007	10

*bgs = below ground surface

Table 6-5

Human Health Risk Assessment Data Set for Surface Water

Sample Numbers
LL9SW-001
LL9SW-002
LL9SW-003
LL9SW-004
LL9SW-005
LL9SW-012

Table 6-6

Human Health Risk Assessment Data Set for Sediments

Sample ID	Depth bgs*
LL9SD-002	0-0.5
LL9SD-003	0-0.5
LL9SD-004	0-0.5
LL9SD-005	0-0.5
LL9SD-010	0-0.5
LL9SD-013	0-0.5
LL9SD-014	0-0.5
LL9SD-015	0-0.5

*bgs = below ground surface



Table 6-7

Summary of Quantitative and Qualitative COPCs for Environmental Media

Parameter	Surface Soil	Subsurface Soil	Deep Surface Soil	Ground-water	Surface Water	Sediment
Quantitative COPCs						
Metals						
Aluminum	X		X		X	X
Antimony				X		
Arsenic	X	X	X		X	
Chromium	X	X	X		X	
Lead					X	
Manganese	X		X	X	X	
Mercury	X	X	X		X	X
Vanadium					X	X
Organics						
Benzo(a)pyrene	X		X			
Dibenzo(a,h)anthracene	X		X			
Qualitative COPCs						
Organics						
Nitrocellulose	X	X	X	X	X	X

COPC = Chemical of potential concern

Quantitative COPCs are those COPCs for which risks and/or hazards have been calculated.

Qualitative COPCs are those COPCs for which risks and/or hazards have not been calculated because of a lack of reliable toxicity data.

The 95% UCL on the mean for aluminum, arsenic, chromium, and manganese are less than their respective background concentrations, as discussed further in the uncertainty section.

The maximum detected concentrations of aluminum and copper exceed the RBSCs (set at an HI equal to 0.1), so they were selected as COPCs; however, they do not exceed the residential Region 9 PRGs for soil.

Arsenic was detected in 10 samples at concentrations exceeding the RVAAP background benchmark for surface soil. The concentrations detected are within the literature background values (less than 0.1 to 97 mg/kg) reported in the PRG guidance document (USEPA, 2004c). Arsenic is a component of herbicides commonly used in the United States in the past and is, consequently, often detected at concentrations exceeding naturally occurring background concentrations. Arsenic is retained as a COPC for surface soil. Uncertainties associated with its selection as a COPC are further discussed in the Section 6.6.



Nitrocellulose was selected as a COPC because PRGs are not available for this chemical. However, according to the USEPA Office of Drinking Water, nitrocellulose is essentially non-toxic (USEPA, 2004d). Nitrocellulose is further discussed in the uncertainty section.

No chemicals were eliminated as COPCs for surface soil on the basis of frequency of detection.

6.2.3.2 Subsurface Soil

Analytical data available for 53 subsurface soil samples were evaluated in the BHRRA. Twenty-two metals, 16 SVOCs, RDX, and nitrocellulose were detected in the subsurface soil samples collected at LL-9. The samples evaluated in this BHRRA are listed in Table 6-2. The results of the COPC selection for subsurface soils are summarized in Appendix V, Table 2.2. The following chemicals were identified as subsurface soil COPCs:

- Metals (arsenic, chromium, and mercury); and
- Nitrocellulose.

One chromium detection exceeds the RVAAP background benchmark for subsurface soil. The maximum detected concentrations of mercury exceed the RBSCs (set at an HI equal to 0.1), however, it does not exceed the Region 9 residential PRGs for soil.

Arsenic was detected in 5 of 49 samples at concentrations exceeding the RVAAP background benchmark for subsurface soil.

Nitrocellulose was selected as a COPC because PRGs are not available for this chemical. However, according to the USEPA Office of Drinking Water, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in the uncertainty section.

Aluminum, vanadium, and manganese were eliminated as COPCs for subsurface soil on the basis of the background screen only. The maximum detected concentration of manganese and vanadium exceeded the RBSC (set at an HI equal to 0.1). No chemicals were eliminated as COPCs for subsurface soil on the basis of frequency of detection.

6.2.3.3 Deep Surface (0-4 ft bgs) Soils

Analytical data available for 91 deep surface soil samples were evaluated in the BHRRA. Twenty-two metals, 19 SVOCs, RDX, nitroguanidine, and nitrocellulose were detected in the deep surface (0-4 ft bgs) soil samples collected at LL-9. The samples evaluated in this BHRRA are listed in Table 6-3. The results of the COPC selection for deep surface soil are summarized in Appendix V, Table 2.3. The following chemicals were identified as deep surface soil COPCs:



- Metals (aluminum, arsenic, chromium, copper, lead, manganese, and mercury);
- Organics (benzo[a]pyrene, and dibenzo[a,h]anthracene); and
- Nitrocellulose.

Three aluminum detections and two manganese detections in 85 samples exceeded their respective background concentrations. The maximum detected concentrations of aluminum and copper exceed the RBSCs (set at an HI equal to 0.1). Arsenic was detected in 22 of 85 samples at concentrations exceeding the RVAAP background benchmark for deep surface soil.

Nitrocellulose was selected as a COPC because PRGs are not available for this chemical. However, according to the USEPA Office of Drinking Water, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in the uncertainty section.

No chemicals were eliminated as COPCs for deep surface soil on the basis of frequency of detection.

6.2.3.4 Groundwater

Analytical data available for seven groundwater samples were evaluated in the BHHRA. Thirteen metals, bis(2-ethylhexyl)phthalate, and nitrocellulose were detected in the seven groundwater samples collected at LL-9. The samples evaluated in this BHHRA are listed in Table 6-4. The results of the COPC selection for groundwater are summarized in Appendix V, Table 2.4. The following chemicals were identified as groundwater COPCs:

- Metals (antimony and manganese); and
- Nitrocellulose.

Only one manganese detection is greater than its background concentration. The maximum detection of antimony (4 µg/L) does not exceed the SDWA primary MCL of 6 µg/L. The maximum detections of antimony and manganese exceed the RBSC, which is set at an HI equal to 0.1.

No chemicals were eliminated as COPCs on the basis of frequency of detection.

6.2.3.5 Surface Water

Analytical data available for six surface water samples were evaluated in the BHHRA. Nineteen metals and nitrocellulose were detected in the six surface water samples collected at LL-9. The samples evaluated in this BHHRA are listed in Table 6-5. The results of the COPC selection for



surface water are summarized in Appendix V, Table 2.5. The following chemicals were identified as surface water COPCs:

- Metals (aluminum, arsenic, chromium, lead, manganese, mercury, and vanadium) and
- Nitrocellulose.

Three chromium detections exceed the RBSC; however, no concentrations exceed the SDWA primary MCL (100 µg/L). The maximum detected concentrations of aluminum, chromium, and vanadium exceed the RBSCs, which are set at an HI equal to 0.1.

Nitrocellulose was selected as a COPC because a PRG is not available for this chemical. However, as noted above, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in the uncertainty section.

No chemicals were eliminated as COPCs for surface water on the basis of the background screen.

6.2.3.6 Sediment

Analytical data available for 13 sediment samples were evaluated in the BHHRA. Twenty-two metals, 17 SVOCs, and nitrocellulose were detected in the sediment samples collected at LL-9. The samples evaluated in this BHHRA are listed in Table 6-6. The results of the COPC selection for sediments are summarized in Appendix V, Table 2.6. The following chemicals were identified as sediment COPCs:

- Metals (aluminum, mercury, and vanadium) and
- Nitrocellulose.

The maximum detected concentrations of the inorganic COPCs exceed the RBSCs (set at an HI equal to 0.1). Two vanadium and three aluminum detections exceed the RVAAP's background benchmarks for sediments.

Nitrocellulose was selected as a COPC because a PRG is not available for this chemical. However, as noted above, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in the uncertainty section.

Arsenic and manganese were eliminated as COPCs for sediment on the basis of the background screen only. No chemicals were eliminated as COPCs on the basis of frequency of detection.



6.2.3.7 Summary of COPC Selection

The COPCs selected for LL-9 are summarized in Table 6-7. Several COPCs were detected at maximum concentrations only slightly greater than RVAAP background and may actually represent background conditions. The average concentrations and the 95% UCL on the mean are less than RVAAP background values for many metals in soil (surface soil, subsurface soil, and sediment).

6.3 Exposure Assessment

The exposure assessment defines and evaluates the current and potential future exposures by receptor populations. More specifically, an exposure assessment identifies the pathways by which humans are potentially exposed to COPCs, the magnitude of the potential human exposure, and the frequency and duration of exposure. This process involves the following steps:

- Characterization of the exposure setting in terms of physical characteristics and the populations that may potentially be exposed to site-related chemicals;
- Identification of potential exposure pathways and receptors; and
- Quantification of exposure for each receptor in terms of the amount of chemical that is ingested, inhaled, or absorbed through the skin from potentially complete exposure pathways.

The output of the exposure assessment (i.e., an estimate of COPC intake) is used in conjunction with toxicity criteria (i.e., cancer slope factors and reference doses) identified in the toxicity assessment (Section 6.4) to quantify risks and hazards to receptors during risk characterization (Section 6.5).

6.3.1 Exposure Setting

The exposure setting was described in the FWHHRAM and White Paper (TtNUS, 2004). The RVAAP installation is located in two counties of northeastern Ohio, Portage and Trumbull, with a majority of the facility in Portage County. According to the 2000 Census, the total populations of Portage and Trumbull counties were 152,061 and 225,116, respectively. The largest population centers in the area are the City of Ravenna (population 11,771) approximately 3.2 km (2 miles) to the west, and Newton Falls (population 5,002) approximately 1.6 km (1 mile) to the southeast (Science Applications International Corporation [SAIC, 2001]).

Approximately 55 percent of Portage County is either woodland or farmland (Portage County, 1985; Census Bureau, 1992). To the south of the facility, across State Route 5, is the Michael J. Kirwan Reservoir, which is used for recreational purposes. The reservoir is fed by the West



1 Branch of the Mahoning River, which flows south along the western edge of the installation.
2 Hinckley Creek flows south across the western portion of the facility and eventually flows into
3 the West Branch of the Mahoning River. The other major surface drainages at RVAAP, Sand
4 Creek and the South Fork of Eagle Creek, exit the facility and eventually flow east to the
5 Mahoning River (SAIC, 2001).

6 6.3.1.1 *Land Use*

7 The land use immediately surrounding the facility is primarily farmland with sparse private
8 residences. Residential groundwater use occurs outside the facility. Residential wells in the
9 vicinity of the RVAAP are constructed in both the unconsolidated unit and bedrock (SAIC,
10 2001). The last of the process production wells on the facility were abandoned in 1992. Two
11 groundwater production wells in the central portion of the facility remain in operation to provide
12 sanitary water to the remaining site personnel. An additional groundwater production well is not
13 in operation but could be activated in the future.

14 Access to the facility is restricted by a perimeter fence. In 1992, the land use changed from
15 "inactive -maintained" status to "modified (not maintained) caretaker" status (U.S. Department of
16 the Army, 1993). This new status indicated that the facility was no longer needed to mobilize
17 for production of ammunition for war efforts. The only remaining federally mandated mission for
18 the facility – ammunition and bulk explosives storage – ended in December 2004 (SAIC, 2001).

19 LL-9 lies within the south-central portion of the RVAAP. Site workers (security guards and
20 maintenance workers) infrequently visit LL-9; the load line is no longer mowed on a routine
21 basis. On-site surface water is limited to intermittent drainage ditches or gullies. There are no
22 groundwater or production wells at LL-9. The ground surface at LL-9 slopes gently to the north
23 in the areas of the former LL-9 buildings, and there is a sharp drop in elevation north of the LL-
24 9 perimeter road.

25 Future recreational goals for RVAAP are hunting (mainly deer), fishing, and trapping anywhere
26 suitable habitat exists. In addition, National Guard training operations are very likely to occur in
27 the LL-9 area in the future. The OHARNG-proposed land use for LL-9 is Mounted Training –
28 No Digging. The National Guard trainee is assumed to come in direct contact with soil, surface
29 water, and sediment for 24 days a year on inactive duty and 15 days a year during training.
30 Digging is prohibited in this area, but mounted vehicles (e.g., tanks) are expected to disturb soil
31 up to 4 ft bgs.

32 As noted above, the most plausible long-term use is a combination of OHARNG training and
33 controlled recreation. Although other uses of the LL-9 area are unlikely, this BHHRA also
34 evaluates additional potential future land uses that reflect a broader use of the land, including
35 residential farm use and a more intensive use of the AOC by the National Guard (e.g., a
36 National Guard resident trainer). The resident trainer would be expected to have similar



exposures as the trainee, but the days-per-year exposure frequency is anticipated to be greater because the trainer would supervise the training of several groups of trainees. Table 6-8 summarizes the land uses and receptors that were evaluated as part of the BHHRA.

Table 6-8

Potential Receptors for the LL-9 BHHRA

Land Use Designation	Description	Potential Receptors
Modified Caretaker – Managed Recreational	light maintenance and managed hunting/trapping	security guard/maintenance worker permitted visitor: hunter/trapper
National Guard Training Area– Managed Recreational	Ohio National Guard training activities and managed hunting/ trapping	National Guard trainee National Guard resident/trainer National Guard fire and dust control personnel permitted visitor: hunter/ trapper
Open Residential	unrestricted residential housing and farming	on-site resident farmer

6.3.1.2 Receptors

The potential receptors for each land use scenario are further defined in Table 6-9. The exposure pathways for each receptor and environmental medium evaluated are listed in Table 6-9 and reflect the protocol established in the RVAAP FWHHRAM and the White Paper developed for Load Lines 6, 9, and 11 (TtNUS, 2004). The adult and juvenile trespasser scenario is located in Appendix Y.

6.3.2 Exposure Pathways

A summary of potential exposure pathways for the receptors evaluated in the LL-9 BHHRA are described in Section 5. Information in this section includes sources, release mechanisms, transport pathways, exposure routes, and receptors. The exposure assumptions used to estimate COPC intake for each receptor are shown in Table 6-10. The current and likely future land use scenarios and receptors identified in Section 6.3.1.1 are discussed in the following subsections.



6.3.2.1 *Modified Caretaker – Managed Recreational*

This scenario describes the current land use at LL-9. Receptor/exposure activities under the current land use scenario are expected to be as described in the FWHHRAM and the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004). Consequently, the exposure assumptions for the potential receptors (security guard/maintenance worker and hunter/trapper) presented in Table 6-10 for the current land use scenario are those suggested in the FWHHRAM and White Paper. The evaluation of the following authorized receptors (i.e. receptors allowed access to LL-9 during its future reuse) provides “reasonable worst-case” risk estimates that may be used to make risk management decisions, assuming that the future land use will not change significantly.

6.3.2.2 *Security Guard – Maintenance Worker*

Current activities at LL-9 include maintenance activities and security patrols. Consequently, a security guard or maintenance worker may contact environmental media (i.e., surface soil) at LL-9. However, the contact is likely to be limited because maintenance activities are not routinely scheduled for LL-9, and an individual performing a security patrol is not likely to intentionally contact environmental media on a regular basis. Use of the shallow aquifer for a potable water supply and contact with subsurface soil, surface water, and sediment are not considered likely. Although routine mowing does not currently take place in LL9, it will occur on a regular basis once the site is turned over to the OHARNG. Exposure parameter values used to assess this receptor in the BHHRA are presented in Table 6-10.

6.3.2.3 *Hunter/Trapper*

Information regarding hunting, trapping, and fishing activities at RVAAP is based on information summarized in the FWHHRAM and based on communications with on-site personnel. Permitted deer hunting occurs at RVAAP. A future goal for the RVAAP is to allow hunting, fishing, and trapping anywhere suitable habitat exists. According to the RVAAP FWHHRAM, a hunter is assumed to be within a particular RVAAP area at a particular frequency (e.g., 6 hours per day for 2 days per year), and the trapper is assumed to be exposed less frequently (e.g., 0.5 hour per day for 6 days per year); therefore, the hunter exposure is used as the more conservative scenario. The hunter/trapper is assumed to hunt/trap as long as he/she resides in the area, so 30 years – the residential default exposure duration – is used. A hunter/trapper operating in the vicinity of LL-9 may be exposed COPCs in surface soil, surface water, and sediment as well as through the ingestion of deer meat. Exposure parameter values used to assess the hunter/trapper receptor are found in Table 6-10.

The intermittent drainage ditches at LL-9 are not capable of sustaining fish population; consequently, fishing does not occur at LL-9.



Table 6-9
Receptors and Exposure Pathways

Exposure Pathways	Exposure Media			
	Groundwater	Surface Water	Sediment	Surface Soil ^a
National Guard - Trainee				
Ingestion				
Incidental	--	X	X	X
Drinking water	X	--	--	--
Dermal	X	X	X	X
Inhalation				
Vapor	No VOCs	--	No VOCs	No VOCs
Dust	--	--	X	X
Ingestion of food	--	--	--	--
National Guard - Trainer/Resident				
Ingestion				
Incidental	--	X	X	X
Drinking water	X	--	--	--
Dermal	X	X	X	X
Inhalation				
Vapor	No VOCs	--	No VOCs	No VOCs
Dust	--	--	X	X
Ingestion of food	--	--	--	--
Future Hypothetical Resident				
Ingestion				
Incidental	--	X	X	X
Drinking water	X	--	--	--
Dermal	X	X	X	X
Inhalation				
Vapor	No VOCs	--	No VOCs	No VOCs
Dust	--	--	X	X
Ingestion of beef, milk, and vegetables	X	--	--	X
Ingestion of venison	--	--	--	X
Recreators - Hunter/Trapper				
Ingestion				
Incidental	--	X	X	X
Drinking water	--	--	--	--
Dermal	--	X	X	X
Inhalation				
Vapor	--	--	No VOCs	No VOCs
Dust	--	--	X	X
Ingestion of venison	--	--	--	X
Security and Maintenance Personnel				
Ingestion				
Incidental	--	--	--	X
Drinking water	--	--	--	--
Dermal	--	--	--	X
Inhalation				
Vapor	--	--	--	No VOCs
Dust	--	--	--	X
Ingestion of food	--	--	--	--

^aSurface soil is defined as 0-1 feet for all receptors except the National Guard Trainee, where it is 0-4 feet bgs (deep surface soil) due to the nature of ground training activities.



Table 6-10
Exposure Assumptions

Parameter	Units	Security Guard/ Maintenance Worker	National Guard Trainee	National Guard Trainer/Resident	Hunter/Trapper	Resident Farmer (child/adult)
Pathway						
Surface Soil is defined as 0-1 for all receptors except the National Guard Trainee, where it is 0 to 4 feet bgs due to nature of ground training activities.						
Incidental ingestion						
Soil ingestion rate (Adult)	kg/day	0.0001 ^a	0.0001 ^a	0.0001 ^a	0.0001 ^a	0.0001 ^a
Soil ingestion rate (Child)	kg/day	NA	NA	NA	NA	0.0002 ^a
Exposure time	hours/day	1 ^b	24 ^b	24 ^e	6	24 ^a
Exposure frequency	days/year	250 ^a	39 ^b	250 ^e	2	350 ^a
Exposure duration (Adult)	years	25 ^a	25 ^b	25 ^b	30 ^a	30 ^a
Exposure duration (Child)	years	NA	NA	NA	NA	6 ^a
Body weight (Adult)	kg	70 ^a	70 ^a	70 ^a	70 ^a	70 ^a
Body weight (Child)	kg	NA	NA	NA	NA	15 ^a
Carcinogen averaging time	days	25550 ^a	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time (Adult)	days	9125 ^a	9125 ^a	9125 ^a	10950 ^a	10950 ^a
Noncarcinogen averaging time (Child)	days	NA	NA	NA	NA	2190 ^a
Fraction ingested	unitless	1 ^b	1 ^b	1 ^b	1 ^b	1 ^a
Conversion factor	days/hour	0.0625 ^y	0.042	0.042	0.0625 ^y	0.042
Dermal contact						
Skin area	m ² /event	0.33 ^d	0.33 ^d	0.33 ^d	0.57 ^d	0.22 ^h /0.57 ^d
Adherence factor	mg/cm ²	0.7 ^c	0.3 ^c	0.3 ^c	0.3 ^c	0.2 ^h /0.4 ^c
Absorption fraction	Unitless	Chem. Spec ^p	Chem. Spec ^p	Chem. Spec ^p	Chem. Spec ^p	Chem. Spec ^p
Exposure frequency	events/year	250 ^a	39 ^b	250 ^e	2	350 ^a
Exposure duration	years	25 ^a	25 ^b	25 ^b	30 ^a	6 ^a /30 ^a
Body weight	kg	70 ^a	70 ^a	70 ^a	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	25550 ^a	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	9125 ^a	9125 ^a	9125 ^a	10950 ^a	2190 ^a /10950 ^a
Conversion factor	(kg-cm ²)/(mg-m ²)	0.01	0.01	0.01	0.01	0.01
Inhalation of VOCs and dust						
Inhalation rate	--	0.83 ^a m ³ /hr	44.4 m ³ /day	44.4 m ³ /day	1.47 ^f m ³ /hr	10 ^q /20 ^a m ³ /day
Exposure time	hours/day	1 ^b	24 ^b	24 ^e	6	24/24 ^a
Exposure frequency	days/year	250 ^a	39 ^b	250 ^e	2	350 ^a
Exposure duration	years	25 ^a	25 ^b	25 ^b	30 ^a	6 ^a /30 ^a
Body weight	kg	70 ^a	70 ^a	70 ^a	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	25550 ^a	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	9125 ^a	9125 ^a	9125 ^a	10950 ^a	2190 ^a /10950 ^a
Conversion factor	days/hour	NA	0.042	0.042	NA	0.042
Particulate Emission Factor	m ³ /kg	9.24E+08 ^v	1.67E+06 ^w	9.24E+08 ^v	9.24E+08 ^v	9.24E+08 ^v
SEDIMENT						
Incidental ingestion						
Soil ingestion rate (Adult)	kg/day	NA	0.0001 ^a	0.0001 ^a	0.0001 ^a	0.0001 ^a
Soil ingestion rate (Child)	kg/day	NA	NA	NA	NA	0.0002 ^a
Exposure time	hours/day	NA	24 ^b	24 ^e	6	24 ^a
Exposure frequency	days/year	NA	39 ^b	250 ^e	2	75 ^b
Exposure duration (Adult)	years	NA	25 ^b	25 ^b	30 ^a	30 ^a
Exposure duration (Child)	years	NA	NA	NA	NA	6 ^a
Body weight (Adult)	kg	NA	70 ^a	70 ^a	70 ^a	70 ^a
Body weight (Child)	kg	NA	NA	NA	NA	15 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time (Adult)	days	NA	9125 ^a	9125 ^a	10950 ^a	10950 ^a
Noncarcinogen averaging time (Child)	days	NA	NA	NA	NA	2190 ^a
Fraction ingested	unitless	NA	1 ^b	1 ^b	1 ^b	1 ^a
Conversion factor	days/hour	NA	0.042	0.042	0.0625 ^y	0.042



Table 6-10
Exposure Assumptions

Parameter	Units	Security Guard/ Maintenance Worker	National Guard Trainee	National Guard Trainer/Resident	Hunter/Trapper	Resident Farmer (child/adult)
Pathway						
Surface Soil is defined as 0-1 for all receptors except the National Guard Trainee, where it is 0 to 4 feet bgs due to nature of ground training activities.						
Dermal contact						
Skin area	m ² /event	NA	0.33 ^d	0.33 ^d	0.52 ^d	0.22 ^h /0.57 ^d
Adherence factor	mg/cm ²	NA	0.3 ^c	0.3 ^c	0.3 ^c	0.2 ^h /0.4 ^c
Absorption fraction	unitless	NA	chem. Spec ^p	chem. Spec ^p	chem. Spec ^p	chem. Spec ^p
Exposure frequency	events/year	NA	39 ^b	250 ^e	2	75 ^a
Exposure duration	years	NA	25 ^b	25 ^b	30 ^a	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	10950 ^a	2190 ^a /10950 ^a
Conversion factor	(kg-cm ²)/(mg-m ²)	NA	0.01	0.01	0.01	0.01
Inhalation of VOCs and dust						
Inhalation rate	--	NA	44.4 m ³ /day	44.4 m ³ /day	0.83 ^a m ³ /hr	10 ^q /20 ^a m ³ /day
Exposure time	hours/day	NA	24 ^b	24 ^e	6	24 ^a
Exposure frequency	days/year	NA	39 ^b	250 ^e	2	75 ^b
Exposure duration	years	NA	25 ^b	25 ^b	30 ^a	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	10950 ^a	2190 ^a /10950 ^a
Conversion factor	days/hour	NA	0.042	0.042	0.0625 ^y	0.042
Particulate Emission Factor	m ³ /kg	NA	1.67E+06 ^w	9.24E+08 ^v	9.24E+08 ^v	9.24E+08 ^v
SURFACE WATER						
Incidental ingestion while wading						
Drinking water ingestion rate	L/hr	NA	NA	NA	NA	NA
Incidental water ingestion rate	L/day	NA	0.1 ^b	0.1 ^b	0.05 ^b	0.1 ^b
Exposure frequency	days/year	NA	39 ^b	250 ^e	2	45 ^b
Exposure duration	years	NA	25 ^b	25 ^b	30 ^a	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	10950 ^a	2190 ^a /10950 ^a
Dermal contact while wading						
Skin area	m ²	NA	0.33 ^d	0.33 ^d	0.52 ^d	0.22 ^h /0.57 ^d
Exposure time	hours/day	NA	2 ^b	2 ^b	2 ^b	2 ^b
Exposure frequency	days/year	NA	39 ^b	250 ^e	2	45 ^b
Exposure duration	years	NA	25 ^b	25 ^b	30 ^a	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	10950 ^a	2190 ^a /10950 ^a
Conversion factor	(m/cm)(L/m ³)	NA	10	10	10	10
GROUNDWATER						
Drinking Water Ingestion						
Drinking water ingestion rate	L/day	NA	2 ^a	2 ^a	NA	1.5 ^h /2 ^a
Exposure frequency	days/year	NA	39 ^b	250 ^e	NA	350 ^a
Exposure duration	years	NA	25 ^b	25 ^b	NA	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	NA	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	NA	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	NA	2190 ^a /10950 ^a



Table 6-10
Exposure Assumptions

Parameter	Units	Security Guard/ Maintenance Worker	National Guard Trainee	National Guard Trainer/Resident	Hunter/Trapper	Resident Farmer (child/adult)
Pathway						
Surface Soil is defined as 0-1 for all receptors except the National Guard Trainee, where it is 0 to 4 feet bgs due to nature of ground training activities.						
Dermal contact while showering						
Skin area	m ²	NA	1.94 ^g	1.94 ^g	NA	0.866 ^s /1.94 ^g
Exposure time	hours/day	NA	0.25 ^a	0.25 ^a	NA	0.25 ^a
Exposure frequency	days/year	NA	39 ^b	250 ^e	NA	350 ^a
Exposure duration	years	NA	25 ^b	25 ^b	NA	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	NA	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	NA	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	NA	2190 ^a /10950 ^a
Conversion factor	(m/cm)(L/m ³)	NA	10	10	NA	10
Inhalation of VOCs during household water use (Only if applicable)						
Inhalation rate	--	NA	0.83 ^a m ³ /hr	0.83 ^a m ³ /hr	NA	10 ^q /20 ^a
Exposure time	hours/day	NA	24	24	NA	24
Exposure frequency	days/year	NA	39 ^b	250 ^e	NA	350 ^a
Exposure duration	years	NA	25 ^b	25 ^b	NA	6 ^a /30 ^a
Body weight	kg	NA	70 ^a	70 ^a	NA	15 ^a /70 ^a
Carcinogen averaging time	days	NA	25550 ^a	25550 ^a	NA	25550 ^a
Noncarcinogen averaging time	days	NA	9125 ^a	9125 ^a	NA	2190 ^a /10950 ^a
Volatilization factor	L/m ³	NA	0.5 ^a	0.5 ^a	NA	0.5 ^a
FOODSTUFFS						
Ingestion of venison						
Conversion factor	unitless	NA	NA	NA	1.25	1.25
Browse ingestion rate	kg dry weight/day	NA	NA	NA	0.87 ^b	0.87 ^b
Fraction browse ingested from site	unitless	NA	NA	NA	0.16	0.16
Fat ratio (venison to beef)	unitless	NA	NA	NA	0.2	0.2
Venison ingestion rate	kg/day	NA	NA	NA	0.03 ^b	0.03 ^b
Fraction ingested	unitless	NA	NA	NA	1 ^b	1 ^b
Exposure frequency	days/year	NA	NA	NA	365 ^b	365 ^b
Exposure duration	years	NA	NA	NA	30 ^a	6 ^a /30 ^a
Body weight	kg	NA	NA	NA	70 ^a	15 ^a /70 ^a
Carcinogen averaging time	days	NA	NA	NA	25550 ^a	25550 ^a
Noncarcinogen averaging time	days	NA	NA	NA	10950 ^a	2190 ^a /10950 ^a
Ingestion of beef						
Resuspension multiplier	unitless	NA	NA	NA	NA	0.25 ^l
Quantity of pasture ingested	kg dry weight/day	NA	NA	NA	NA	7.2 ^l
Fraction of year cow is on-site	unitless	NA	NA	NA	NA	1
Fraction of cow's food from on-site	unitless	NA	NA	NA	NA	0.9 ^b
Quantity of soil ingested by cow	kg/day	NA	NA	NA	NA	1 ^k
Beef ingestion rate	kg/day	NA	NA	NA	NA	0.0094/0.044 ^l
Fraction ingested	unitless	NA	NA	NA	NA	1 ^b
Exposure frequency	days/year	NA	NA	NA	NA	365 ^b
Exposure duration	years	NA	NA	NA	NA	6 ^a /30 ^a
Body weight	kg	NA	NA	NA	NA	15 ^a /70 ^a
Carcinogen averaging time	days	NA	NA	NA	NA	25550 ^a
Noncarcinogen averaging time	days	NA	NA	NA	NA	2190 ^a /10950 ^a



Table 6-10
Exposure Assumptions

Parameter	Units	Security Guard/ Maintenance Worker	National Guard Trainee	National Guard Trainer/Resident	Hunter/Trapper	Resident Farmer (child/adult)
Pathway						
Surface Soil is defined as 0-1 for all receptors except the National Guard Trainee, where it is 0 to 4 feet bgs due to nature of ground training activities.						
Ingestion of milk products						
Resuspension multiplier	unitless	NA	NA	NA	NA	0.25 ⁱ
Quantity of pasture ingested	kg dry weight/day	NA	NA	NA	NA	16.1 ^j
Fraction of year cow is on-site	unitless	NA	NA	NA	NA	1 ^b
Fraction of cow's food from on-site	unitless	NA	NA	NA	NA	0.6 ^b
Quantity of soil ingested by cow	kg/day	NA	NA	NA	NA	1 ^k
Milk ingestion rate (Adult)	kg/day	NA	NA	NA	NA	0.305 ^j
Milk ingestion rate (Child)	kg/day	NA	NA	NA	NA	0.509 ^m
Fraction ingested	unitless	NA	NA	NA	NA	1 ^b
Exposure frequency	days/year	NA	NA	NA	NA	365 ^b
Exposure duration (Adult)	years	NA	NA	NA	NA	30 ^a
Exposure duration (Child)	years	NA	NA	NA	NA	6 ^a
Body weight (Adult)	kg	NA	NA	NA	NA	70 ^a
Body weight (Child)	kg	NA	NA	NA	NA	15 ^a
Carcinogen averaging time	days	NA	NA	NA	NA	25550 ^a
Noncarcinogen averaging time (Adult)	days	NA	NA	NA	NA	10950 ^a
Noncarcinogen averaging time (Child)	days	NA	NA	NA	NA	2190 ^a
Ingestion of vegetables						
Resuspension multiplier	unitless	NA	NA	NA	NA	0.26 ⁿ
Vegetable ingestion rate	kg/day	NA	NA	NA	NA	0.043/0.2 ^j
Fraction ingested	unitless	NA	NA	NA	NA	0.4 ^j
Exposure frequency	days/year	NA	NA	NA	NA	365 ^a
Exposure duration	years	NA	NA	NA	NA	6 ^a /30 ^a
Body weight	kg	NA	NA	NA	NA	15 ^a /70 ^a
Carcinogen averaging time	days	NA	NA	NA	NA	25550 ^a
Noncarcinogen averaging time	days	NA	NA	NA	NA	2190 ^a /10950 ^a
Noncarcinogen averaging time	days	NA	NA	NA	10950 ^a	10950 ^a

Note: Resident farmer is chosen instead of resident because the property would be used for farming in the unlikely event it was released from military use. Further, the resident farmer is expected to result in greater, more conservative exposure intake when compared to the resident.

Shading indicates values that differ than those presented in the Facility Wide Human Health Risk Assessors Manual.

NA = not applicable for this scenario.

a RAGS, Part B (EPA 1991a).

b Site-specific (value assumed for site or value obtained from site personnel). National Guard Trainee is assumed to be on –site 24 hrs/d for 24 d/yr for inactive duty training and 24 hrs/d for 15 d/yr for annual training. Both National Guard receptors are assumed to remain at RVAAP and at the AOC of interest for their entire 25 year enlistment. The Hunter is assumed to be on-site 6 hours/day for 2 days/year. The trapper will be exposed less (i.e., 0.5 hours/day for 6 days/year); therefore, the hunter exposure is used as the more conservative scenario. The hunter is assumed to hunt as long as he/she resides in the area, so the residential default exposure duration is used. The Security Guard/Maintenance Worker is assumed to visit each AOC for 1 hour/day for a standard worker default of 250 days/yr and 25 years.

National Guard Trainee, National Guard Resident, and Resident Farmer are assumed to ingest 0.05 L/hour [per RAGS Part A (EPA 1998)] for approximately 2 hours/day spent in the surface water. Hunter/trappers are assumed to ingest 0.05 L/day due to splashing while setting traps.



Table 6-10
Exposure Assumptions

Parameter	Units	Security Guard/ Maintenance Worker	National Guard Trainee	National Guard Trainer/Resident	Hunter/Trapper	Resident Farmer (child/adult)
Pathway						
Surface Soil is defined as 0-1 for all receptors except the National Guard Trainee, where it is 0 to 4 feet bgs due to nature of ground training activities.						

c Security Guard/Maintenance Worker = Adult Groundskeeper (95th percentile); Hunter/Trapper = Residential Default; National Guard Trainee = Construction Worker (95th percentile); Resident Farmer Adult= Adult Farmer (95th percentile) (RAGS, Vol. 1 Part E, Supplemental Guidance for Dermal Risk Assessment, Interim) EPA/540/R/99/005.

d Security Guard/Maintenance Worker, National Guard Trainee = Industrial Default; Hunter/Trapper and Resident Farmer = Adult Residential Default. Exposure Factors Handbook (EPA 1997a) (Note dermal contact for Hunter/trapper during wading is 0.52 based on head, hands, forearms and lower legs from Exposure Factors Handbook.)

e National Guard resident is assumed to reside on-site during the week (i.e., 24 hours/day, 5 days/week, 50 weeks/year) and go home on weekends.

f RAGS, Part A (EPA 1989a).

g Average total body surface area for an adult (EPA 1992b).

h Per OEPA comment 2002.

i Plant mass loading factor for pasture (Hinton 1992).

j International Atomic Energy agency 1994.

k Soil ingestion by dairy cattle (Darwin 1990).

l Exposure Factors Handbook (EPA 1997). 50th percentile beef ingestion 0.626 g/kg-day = 44 g/day for a 70 kg adult and 9.4 for a 15 kg child. 50th percentile vegetable ingestion rate = 2.86 g/kg/day = 200 g/day for a 70 kg adult and 43 g/day for a 15 kg child. Ingestion rates for the total population and not any age-specific group.

m Pao et al. (1982).

n Plant mass loading factor for vegetables (Pinder 1989).

o Based on EPA/FDA Nation Advisory of 1 meal (8 oz uncooked weight) noncommercial fish per week (EPA-823-F-04-008). Adult or Child

p Risk Assessment Guidance for Superfund, Volume 1: Human Health Evaluation Manual (Part E, Supplemental Guidance for Dermal Risk Assessment, Interim), EPA/540/R/99/005.

q Recommended value for child age 6-8 (EPA 1997a).

r The inhalation rate for hunter/trapper is based on an adult engaged in light activities 4 hours/day, moderate activities 1 hour/day, and heavy activities 1 hour/day.

s 50th percentile value for male child age 6-7 (EPA 1997a).

u Ecological Risk Assessment. Ohio EPA/DERR. February 2003

v Default value for Cleveland, Ohio. EPA Soil Screening Calculation Internet Site (http://risk.lsd.ornl.gov/calc_start.shtml).

w Based on a dust loading factor of 600 ug/m³ (DOE, 1993).

x A site specific value will be developed following methodology in EPA's Supplemental Guidance for Developing Soil Screening Levels for Superfund Sites, December 2002, OSWER 9355.4-24.

y Based on a 16 hour day.



6.3.2.4 *National Guard – Managed Recreational*

Three receptors are expected under the National Guard managed recreational land use scenario, a hunter/trapper, National Guard training participant, and National Guard resident trainer. Although National Guard units typically participate in training 2 weeks a year with some weekend training, instructors and other National Guard personnel managing the training activities could be expected to be on site more frequently. The National Guard trainee is assumed to be on site 24 hours per day for 24 days per year for inactive duty training and 24 hours per day for 15 days per year for annual training, for a total of 39 days per year. The National Guard resident trainer is assumed to reside on site during the week (i.e., 24 hours per day, 5 days per week, 50 weeks per year) and go home on weekends, for a total of 250 days per year. Both National Guard receptors are assumed to remain at RVAAP and at the AOC for their entire 25-year enlistment. The National Guard resident trainer receptor is not included in the RVAAP FWHHRAM; however, based on the anticipated future land use for the LL-9, the National Guard trainee is the key receptor for LL-9. The National Guard resident trainer was included in this BHHRA after discussions with both the USACE and Ohio EPA. Risk estimates for this receptor may assist the risk managers for RVAAP when making certain risk management decisions.

The National Guard receptors may be exposed to COPCs in surface soil, subsurface soil, sediment, and surface water during various training activities such as those detailed in the FWHHRAM. Due to the nature of ground training activities (i.e., use of tracked vehicles and other military equipment), surface soil is defined as 0 to 4 ft bgs for the National Guard trainee. The BHHRA includes risk estimates for National Guard receptors assuming the receptors are using the groundwater as a potable water resource. Exposure parameter values used in this BHHRA to evaluate the National Guard receptors are found in Table 6-10.

6.3.2.5 *Open Residential*

This potential scenario was evaluated to provide risk information assuming a future residential land use of the LL-9 area. Conservatively, a resident farmer adult and child are the receptors of concern. However, based on current and anticipated future land use information, it is unlikely that the LL-9 area will be used for residential purposes. This scenario is included principally for purposes of completeness and to provide information to risk managers regarding the need for institutional land use controls.

Adult and child resident farmers may come into direct contact with COPCs in all media and may also be exposed via indirect exposure pathways (i.e., ingestion of venison, beef, milk, and vegetables). Exposure parameters used to represent receptor activity patterns are listed in Table 6-10 and generally come from the FWHHRAM, with exceptions as outlined in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004).



6.3.2.6 Trespasser

This potential scenario was evaluated in the BHHRA, and the results are presented in Appendix Y. Adult and juvenile trespassers may come into direct contact with surface soil, sediment, and surface water. Exposure parameters used to represent receptor activity patterns are listed in the 2005 addendum to the FWHHRAM.

6.3.3 Exposure Quantification

Intake or dose is defined as the amount of a COPC that could be in contact with the body per unit body weight per unit time. For the LL-9 BHHRA, the surface soil (deep surface soil for the National Guard trainee), subsurface soil, sediment, surface water, and groundwater ingestion, dermal, and inhalation intakes for each receptor as well as for the ingestion of homegrown vegetables, milk, meat, and venison were calculated using standard intake equations from USEPA guidance (e.g., USEPA, 1989a). The standard intake equations are presented in the FWHHRAM with exceptions as outlined in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004).

The standard tables published in RAGS Part D (USEPA, 1989a) are used to present the exposure assumptions, intake equations, dose estimates, toxicity criteria (i.e., cancer slope factors, and reference doses), and cancer and non-cancer risk estimates for receptors evaluated in this BHHRA. These tables also include references for exposure assumptions. The RAGS Part D tables and example calculations for each receptor and exposure pathway are presented in Appendix V.

6.3.3.1 Exposure to Surface Soil

Appendix V (Tables 4.1, 4.2, 4.6, 4.11, 4.15, and 4.24) provides the dose equations used to estimate chemical intake for the security guard/maintenance worker, National Guard trainee, National Guard resident trainer, hunter/trapper, child resident farmer, and adult resident farmer hypothetically exposed to COPCs in surface soil. Where available, the exposure assumptions used to calculate COPC intakes were those suggested by standard USEPA guidance documents and listed in the FWHHRAM. Exposure assumptions for the National Guard trainer/resident were presented in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004). Professional judgment was used to select the exposure parameter values in some cases. For example, it was assumed that the conversion factor for ingestion of soil/sediment should be 0.0625 day per hour (not 0.042 day per hour as originally presented in the risk assessments prepared for Winkelpack remedial investigation) for receptors who are not on site for 24 hours a day (security guard/maintenance worker and hunter/trapper). The change was made to reflect the fact that these receptors are typically not exposed to soil/dust while they are sleeping, and the conversion factor used is more conservative than the value in the FWHHRAM.



Intake values for security guard/maintenance worker, National Guard trainee, National Guard resident trainer, hunter/trapper, child resident farmer, and adult resident farmer exposure to LL-9 surface soil COPCs are provided in Appendix V (Tables 7.1, 7.2, 7.3, 7.4, 7.5, and 7.6, respectively).

6.3.3.2 *Exposure to Subsurface Soil*

It is assumed two receptors – the National Guard resident trainer and the resident farmer – will be exposed to both surface soil and subsurface soil. The National Guard resident trainer may be exposed to shallow subsurface soil during certain training activities. Although unlikely, this BHHRA was prepared assuming that the future resident farmer may also come in contact with subsurface soils excavated and spread across the surface of the site. This scenario is included primarily for completeness and to provide information for informed future land use controls.

Appendix V (Tables 4.9, 4.18, and 4.27) provides the dose equations used to estimate chemical intakes for the National Guard resident trainer, child resident farmer, and adult resident farmer exposed to COPCs in subsurface soil. If available, the exposure assumptions used in the equations were those suggested by standard USEPA guidance documents and outlined in the FWHHRAM. Exposure assumptions for the National Guard resident trainer are not found in the FWHHRAM, but were presented in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004).

Intake values for National Guard resident trainer, child resident farmer, and adult resident farmer exposure to LL-9 subsurface soil COPCs are provided in Appendix V (Tables 7.3, 7.5, and 7.6, respectively).

Intake values for National Guard resident trainer, child resident farmer, and adult resident farmer's exposure to LL-9 subsurface soil COPCs are provided in Appendix V (Tables 7.3, 7.5, and 7.6, respectively).

6.3.3.3 *Exposure to Surface Soil and Subsurface Soil*

When exposure to both surface and subsurface soil is anticipated, total receptor risk estimates were calculated separately to avoid an overestimation of risk (i.e., risk estimates for COPCs in surface soil were not be added to risk estimates developed for COPCs in subsurface soil).

6.3.3.4 *Exposure to Groundwater*

Appendix V (Tables 4.5, 4.10, 4.19, and 4.28) lists the dose equations used to estimate chemical intake for the National Guard trainees, National Guard resident trainer, child resident farmer, and adult resident farmer, respectively, hypothetically exposed to COPCs in groundwater. Where available, the exposure assumptions used in the equations were those suggested by standard USEPA guidance documents and listed in the FWHHRAM. Exposure



assumptions for the National Guard resident trainer are presented in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004).

Intake values for National Guard trainee, National Guard resident trainer, child resident farmer, and adult resident farmer exposure to LL-9 groundwater COPCs are provided in Appendix V (Tables 7.2, 7.3, 7.5, and 7.6, respectively).

6.3.3.5 *Exposure to Surface Water*

Appendix V (Tables 4.4, 4.8, 4.13, 4.17, and 4.26) provides the dose equations used to estimate chemical intake for the National Guard trainee, National Guard resident trainer, hunter/trapper, child resident farmer, and adult resident farmer, respectively, exposed to COPCs in surface water. Where available, the exposure assumptions used in the equations were those suggested by standard USEPA guidance documents and listed in the FWHHRAM. Exposure assumptions for the National Guard resident trainer are presented in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004). However, some exposure factor values were not available from the guidance and were selected using site-specific information provided by RVAAP personnel and using professional judgment. These values are presented and discussed in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004).

Intake values for National Guard trainee, National Guard resident trainer, hunter/trappers, child resident farmer, and adult resident farmer exposure to LL-9 surface water COPCs are provided in Appendix V (Tables 7.2, 7.3, 7.4, 7.5, and 7.6, respectively).

6.3.3.6 *Exposure to Sediment*

Appendix V (Tables 4.3, 4.7, 4.12, 4.16, and 4.25) provides the dose equations used to estimate chemical intake for the National Guard trainee, National Guard resident trainer, hunter/trapper, child resident farmer, and adult resident farmer exposed to COPCs in sediments. Where available, the exposure assumptions used in the equations were those suggested by standard USEPA guidance documents and listed in the FWHHRAM. Exposure assumptions for the National Guard resident trainer are presented in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004). However, professional judgment was used to select exposure parameter values in some cases. For example, it was assumed that the conversion factor for ingestion of soil/sediment should be 0.0625 day per hour (not 0.042 day per hour) for receptors who are not on site for 24 hours a day (security guard/maintenance worker and hunter/trapper). The change was made to reflect the fact that these receptors are typically not exposed to soil/dust while they are sleeping, and the conversion factor used is more conservative than the value in the FWHHRAM.



Intake values for National Guard trainee, National Guard resident trainer, hunter/trapper, child resident farmer, and adult resident farmer exposure to LL-9 sediment COPCs are provided in Appendix V (Tables 7.2, 7.3, 7.4, 7.5, and 7.6 respectively).

6.3.3.7 *Exposure to Venison*

Appendix V (Tables 4.14, 4.20, and 4.29) provides the dose equations used to estimate chemical intake for venison consumption by the hunter/trapper, child resident farmer, and adult resident farmer. Where available, the exposure assumptions used in the equations were those suggested by the FWHHRAM. No exposure assumptions were listed in the FWHHRAM for the hunter/trapper; therefore, values used were those recommended in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004). Site-specific information for LL-9 was used to adjust the fraction of the animals' food from the site to 0.16 to reflect the size of the site (approximately 28 ha) compared to the home range of a white-tailed deer (175 ha) (USEPA, 1993a).

Intake values for hunter/trapper, child resident farmer, and adult resident farmer exposure to LL-9 COPCs in venison are provided in Appendix V (Tables 7.4, 7.5, and 7.6, respectively).

6.3.3.8 *Exposure to Beef*

Appendix V (Tables 4.21 and 4.30) provides the dose equations used to estimate chemical intake for beef consumption by child resident farmer and adult resident farmer, respectively. The exposure assumptions used in the equations were those suggested by the FWHHRAM. Site-specific information for LL-9 was used to determine an exposure parameter as outlined in the FWHHRAM:

- The fraction of the year the animals are on site was presumed to be 1.0 (i.e., the animals are assumed to be on-site year round).
- As explained in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004), the FWHHRAM recommends that COPC concentrations in beef products consider contaminant uptake as a result of the animal ingesting contaminated ground or surface water. The contribution from this pathway is considered minor for LL-9 and was not considered in the BHHRA. Note that the water ingestion contribution is shown in the intake equation given in the FWHHRAM, but no exposure parameter is given in Table 5 of the FWHHRAM.

Intake values for child resident farmer and adult resident farmer exposure to LL-9 COPCs in beef are provided in Appendix V (Tables 7.5 and 7.6).



6.3.3.9 *Exposure to Milk*

Appendix V (Tables 4.22 and 4.31) provides the dose equations used to estimate chemical intake for milk consumption by child resident farmer and adult resident farmer, respectively. The exposure assumptions used in the equations were those suggested by the FWHHRAM. However, as explained in the White Paper for Load Lines 6, 9, and 11 (TtNUS, 2004), the FWHHRAM recommends that COPC concentrations in milk products consider contaminant uptake as a result of the animal ingesting contaminated ground or surface water. The contribution from this pathway is considered minor and was not considered in the BHHRA. Note the water ingestion contribution is shown in the intake equation given in the FWHHRAM, but no exposure parameter is given in Table 5 of the FWHHRAM.

Intake values for child resident s and adult resident farmer exposure to LL-9 COPCs in milk are provided in Appendix V (Tables 7.5 and 7.6, respectively).

6.3.3.10 *Exposure to Vegetables*

Appendix V (Tables 4.23 and 4.32) provides the dose equations used to estimate chemical intake for vegetable consumption by child resident farmer and adult resident farmer. Where available, the exposure assumptions used in the equations were those suggested in the FWHHRAM.

Intake values for child resident s and adult resident farmer exposure to LL-9 COPCs in vegetables are provided in Appendix V (Tables 7.5 and 7.6, respectively).

6.3.4 **Exposure Point Concentrations**

The exposure point concentration (EPC) is the concentration of a COPC used to best estimate the intake of a COPC detected in an environmental media. Ideally the EPC should be the true average concentration within the exposure unit for the media. However, because of the uncertainty associated with estimating the true average concentration at a site, the 95% UCL of the arithmetic mean is selected as the EPC. The following protocol was used to determine EPCs in the BHHRA for LL-9:

- If there were fewer than 10 samples or fewer than 50% positive detections of chemicals in the samples, the maximum concentration was chosen as the EPC because the upper confidence level (UCL) does not provide a good estimation of the upper bound of the mean concentration for small data sets or for data sets that contain a large number of non-detected values (EPA, 2002c).
- If there were more than 10 samples and more than 50% detections, each data set was evaluated using the Shapiro-Wilk W test (Gilbert, 1987) to determine whether the data



set more closely reflected a normal or log-normal distribution. If results were inconclusive, the data were assumed to be log-normally distributed. The 95% UCL-L and 95% UCL-N were calculated for each analyte in each medium and data set, using one-half the reporting limit for non-detect results and the average for samples with duplicates. Example calculations are found in Appendix V. The 95% UCL-N was used as the EPC if the Shapiro-Wilk W test indicated a normal distribution, and the 95% UCL-L was used as the EPC if the Shapiro-Wilk W test indicated a log-normal distribution or if the distribution was undefined (i.e., where the Shapiro-Wilk W test did not indicate a normal or a log-normal distribution). If the calculated 95% UCL exceeded the maximum detected concentration, the maximum detected concentration was selected as the EPC.

EPCs were calculated for analytes detected in surface soil (0–1 ft bgs), deep surface soil (0–4 ft bgs), subsurface soil, groundwater, surface water, and sediment samples located in LL-9. These EPCs are presented in Appendix V (the RAGS Part D, Tables 3.1 through 3.6).

Route-specific EPCs were also calculated for the inhalation of particulates and vapors migrating from soil and sediment to air, the inhalation of VOCs volatilizing from groundwater, and for consumption of foodstuffs. The methods used for calculation of the route-specific EPCs are presented in Appendix V (Tables 4.1 through 4.32).

6.4 Toxicity Assessment

Information concerning the potential human health effects of exposure to COPCs is examined by the risk assessor during the toxicity assessment. For each COPC, the goal of the toxicity assessment is to provide a quantitative estimate of the relationship between the magnitude and type of exposure and the severity or probability of human health effects. The toxicity values presented in this section are integrated with the exposure assessment (Section 6.3) to characterize the potential for the occurrence of adverse health effects.

6.4.1 Toxicity Information and USEPA Guidance for Non-carcinogens

For non-carcinogens, it is assumed that a dose exists below which no adverse health effects will be seen. Below this “threshold” dose, exposure to a chemical can be tolerated without adverse effects. The potential for non-carcinogenic health effects resulting from exposure to chemicals is assessed by comparing the intake dose with this “threshold” dose, or reference dose (RfD). The RfD is determined using available dose-response data for individual chemicals. Uncertainty factors are applied to the highest no-observed-adverse-effect level (NOAEL) to adjust for inter- and intra-species variation, deficiencies in the toxicological database, and use of subchronic rather than chronic animal studies. Inter-species differences in toxicity responses occur as a consequence of the basic biological differences among the various species. Intra-species differences in toxicity responses occur as a consequence of the varying sensitivities amongst



the individuals within a species. Additional uncertainty factors may be applied to estimate a NOAEL from a lowest-observed-adverse-effect level (LOAEL) if the critical study failed to determine a NOAEL. When chemical-specific data are not sufficient, an RfD may be derived from data for a chemical with structural and toxicologic similarity.

The RfD is expressed in units of mg/kg/day and represents a daily intake of contaminant per kilogram of body weight that is not sufficient to cause the threshold effect of concern. An RfD is specific to the chemical, the route of exposure, and the duration over which the exposure occurs. Separate RfDs are presented for ingestion and inhalation pathways. In particular, reference concentrations (RfCs) in units of [mg/m³] are typically presented for the inhalation pathway. Because characterization of non-carcinogenic effects requires an estimate of dose in units of mg/kg/day, the inhalation RfC must be converted to an inhalation RfD. The conversion is performed by assuming that humans weigh 70 kg and inhale 20 m³ of air per day (i.e., the inhalation RfC [mg/m³] is multiplied by 20 m³/day and divided by 70 kg to yield an inhalation RfD with units of [mg/kg/day]) (USEPA, 1995a). The RfDs used to evaluate the COPCs at LL-9 are presented in Appendix V, Table 5.1. The toxicity values were selected using the hierarchy from the FWHHRAM:

- Tier 1 – USEPA's IRIS
- Tier 2 – USEPA's Provisional Peer-Reviewed Toxicity Values (PPRTVs) – The Office of Research and Development/National Center for Environmental Assessment/Superfund Health Risk Technical Support Center (STSC) develops PPRTVs on a chemical-specific basis when requested by USEPA's Superfund program.
- Tier 3 – Other Toxicity Values – Tier 3 includes additional USEPA and non-USEPA sources of toxicity information. Priority should be given to those sources of information that are the most current, the basis for which is transparent and publicly available, and which have been peer reviewed.

6.4.2 Toxicity Information and USEPA Guidance for Carcinogens

The toxicity information considered in the assessment of potential carcinogenic risks includes a weight-of-evidence classification and a slope factor. The weight-of-evidence classification qualitatively describes the likelihood that a chemical is a human carcinogen, based on an evaluation of the available data from human and animal studies. Historically, a chemical was placed in one of three groups in USEPA's classification system to denote its potential for carcinogenic effects:

- Group A – known human carcinogen;
- Group B1 or B2 – probable human carcinogen; and
- Group C – possible human carcinogen.



Chemicals that cannot be classified as human carcinogens because of a lack of data are placed in Group D, and those for which there is evidence of noncarcinogenicity in humans are in Group E. More recently, the USEPA is evaluating the use of the following categories to describe the carcinogenic potential of a chemical:

- Carcinogenic to humans;
- Likely to be carcinogenic to humans;
- Suggestive evidence of carcinogenic potential;
- Inadequate information to assess carcinogenic potential; and
- Not likely to be carcinogenic to humans.

For carcinogens, risks are estimated as the probability that an individual will develop cancer over a lifetime as a result of exposure to the carcinogen. The cancer slope factor (CSF) is the toxicity value used to quantitatively express the carcinogenic hazard of cancer-causing chemicals. It is defined as the upper-bound estimate of the probability of cancer incidence per unit dose averaged over a lifetime. Slope factors are specific to a chemical and route of exposure and are expressed in units of $[\text{mg/kg-day}]^{-1}$ for both oral and inhalation routes. Inhalation cancer toxicity values are usually expressed as inhalation unit risks in units of reciprocal $\mu\text{g}/\text{m}^3$ $[1/(\mu\text{g}/\text{m}^3)]$. The inhalation unit risk must be converted to an inhalation slope factor. This is done by assuming that humans weigh 70 kg and inhale 20 m^3 of air per day (i.e., the inhalation unit risk $[1/\mu\text{g}/\text{m}^3]$ is divided by $20 \text{ m}^3/\text{day}$, multiplied by 70 kg, and multiplied by $1,000 \mu\text{g}/\text{mg}$ to yield the mathematical equivalent of an inhalation slope factor $[\text{mg/kg-day}]^{-1}$) (USEPA, 1995a). CSFs used to evaluate the COPCs at LL-9 are included in Appendix V, Table 6.1, and were selected as described for the RfDs.

6.4.3 Estimation of Toxicity Values for Dermal Exposure

Dermal RfDs and dermal CSFs are derived from the corresponding oral values. In the derivation of a dermal CSF, the oral CSF is divided by the gastrointestinal absorption factor (GAF) to determine a CSF based on an absorbed dose rather than an administered dose. The oral CSF is divided by the GAF because CSFs are expressed as reciprocal doses. A dermal RfD is derived by multiplying an oral RfD by the GAF. Dermal CSFs and RfDs are shown in Appendix V, Tables 5.1 and 6.1. Values for GAF (or Gastrointestinal Absorption Fraction $[\text{ABS}_{\text{GI}}]$) were taken from *RAGS Part E*, Supplemental Guidance for Dermal Risk Assessment (USEPA, 2004a). For chemicals with greater than 50% ABS_{GI} absorption, it is recommended that the default value of complete (i.e., 100 %) oral absorption be assumed, thereby eliminating the need for oral toxicity-value adjustment. Dermal adjustments to the oral CSF or RfD were made for antimony, cadmium, chromium, manganese, and vanadium.



6.4.4 Toxicity Criteria Assumptions

A provisional RfD is available for aluminum from the National Center for Environmental Assessment (NCEA); however, it is based on typical allowable intakes rather than adverse effect levels and is not considered strictly risk-based.

6.4.5 Chemicals Lacking USEPA Toxicity Criteria

Toxicity criteria are not available for one LL-9 COPC: nitrocellulose. Therefore, cancer and non-cancer risk estimates are not provided for this COPC.

6.5 Risk Characterization

The risk assessor evaluates information obtained through the exposure and toxicity assessments to estimate cancer risks and hazard indices in the risk characterization. Total non-carcinogenic and carcinogenic risk estimates for each exposure route, as well as the cumulative risks for each receptor, are included in Appendix V, Tables 9.1 through 9.7.

The risk characterization is presented in three sections. Methods are listed in Section 6.5.1, and results are included in Section 6.5.2. Remedial Goal Options (RGOs) are listed for potential chemicals of concern in Section 6.5.3.

6.5.1 Methodology

Two methods were employed to evaluate risks: one to estimate potential carcinogenic and another to estimate non-carcinogenic risks.

6.5.1.1 Methods for Estimation of Carcinogenic Risks

Carcinogenic risks can be estimated by combining information on the strength or potency of a known or suspected carcinogen (CSF) with an estimate of the individual exposure doses (or intakes) of a chemical. Carcinogenic risk may be estimated as follows:

$$\text{Risk} = \text{CSF} \times \text{Dose}$$

Where:

CSF = carcinogenic slope factor (mg/kg-day)⁻¹

Dose = amount of a contaminant absorbed by a receptor in mg/kg-day



The equation presented above, however, is valid only at risk levels less than or equal to $1E-02$. When the risk estimate is expected to be greater than $1E-02$, an alternate equation, such as the following one-exceedance equation, may be used to estimate risk (USEPA, 1989a):

$$\text{Risk} = 1 - \exp(-\text{Dose} \times \text{CSF})$$

Where:

exp = exponential

The resultant cancer risk value (i.e., $1E-06$ or a 1-in-1,000,000 chance of developing cancer) can be applied to a given population to determine the increase in the cancer rate that could be expected to result from exposure (e.g., $1E-06$ is one additional case of cancer in 1,000,000 exposed persons).

The total risk resulting from exposure of an individual receptor to multiple compounds in a particular medium is the sum of the cancer risks for the individual contaminants in that medium. Cancer risks are summarized for each receptor and each media.

To interpret the quantitative risk estimates and to aid risk managers in determining the need for remediation, quantitative risk estimates are compared to USEPA and Ohio EPA risk benchmarks. The USEPA has defined a "target cancer risk" range of $1E-04$ to $1E-06$. Incremental lifetime cancer risks (ILCRs) below $1E-06$ are generally considered acceptable risks. ILCRs above $1E-04$ are considered unacceptable risks. Risk management decisions are necessary for ILCRs between $1E-06$ and $1E-04$. The Ohio EPA cumulative cancer risk benchmark is $1E-05$ and will be considered by the risk managers when making risk decisions for sites at RVAAP.

6.5.1.2 Methods for Estimation of Non-carcinogenic Risks

Potential health risks resulting from exposure to non-carcinogenic compounds are estimated by comparing the reasonable maximum daily intake dose calculated for an exposure to an acceptable intake dose, such as a chronic or subchronic reference dose (RfD). The ratio of the exposure dose (intake) to the RfD is referred to as the hazard quotient:

$$\text{Hazard Quotient} = \text{Dose}/\text{RfD}$$

If the hazard quotient (HQ) exceeds unity, there is a potential health risk associated with exposure to that chemical (USEPA, 1989a). The dose/RfD ratio is not a mathematical prediction of the severity or probability of toxic effects; it is simply a numerical indicator of the potential for adverse effects. The summation of HQs for several compounds is referred to as the hazard index (HI).



Conservatively, a total HI for any exposure route is calculated by summing the dose/RfD ratios (HQs) for the individual chemicals of concern (USEPA, 1989a). To provide a better indication of risks, dose/RfD ratios are summed according to the target organ affected. For example, the dose/RfD ratios for those chemicals affecting the liver should be summed separately from those chemicals affecting the central nervous system. An HI greater than 1 indicates potential adverse non-carcinogenic health effects (USEPA, 1989a).

The chemicals of concern (COCs) for a given medium are defined as those contaminants that contribute total cancer risk for a receptor greater than $1E-05$ and/or HIs greater than 1.0 within a land use scenario, and that are not eliminated by risk managers after considering the information in the uncertainty analysis.

6.5.2 Risk Characterization Results

The risk characterization for LL-9 is summarized in this section. Total non-carcinogenic and carcinogenic risks for each exposure route, as well as the cumulative risks for each receptor, are included in Appendix V (Tables 9.1 through 9.7) and summarized in Table 6-11.

The following receptors were evaluated:

- The security guard/maintenance worker;
- The National Guard trainee;
- The National Guard resident trainer;
- The hunter/trapper; and
- The hypothetical future resident farmer (adult and child).

Example calculations and relevant risk calculation spreadsheets are included in Appendix V. It should be noted that although the risk estimates provided in Appendix V are presented using two significant figures, final receptor risk estimates are presented in the following narrative in terms of one significant figure as recommended by *RAGS Part A* (USEPA, 1989a).



Table 6-11
Summary of Cancer Risks and Hazard Indices

Receptor	Media	Exposure Route	Cancer Risk	Chemicals with Cancer Risks > 10 ⁻⁴	Chemicals with Cancer Risks > 10 ⁻⁵ and ≤ 10 ⁻⁴	Chemicals with Cancer Risks > 10 ⁻⁶ and ≤ 10 ⁻⁵	Hazard Index	Chemicals with HI > 1
Security Guard/ Maintenance Worker	Surface Soil	Incidental Ingestion	5E-07	--	--	--	6E-03	--
		Inhalation	3E-09	--	--	--	6E-04	--
		Dermal Contact	7E-06	--	--	--	5E-02	--
		Total	8E-06	--	--	Arsenic, Benzo(a)pyrene, Dibenzo(a,h)anthracene	6E-02	--
National Guard Trainee	Deep Surface Soil	Incidental Ingestion	1E-06	--	--	Arsenic	1E-02	--
		Inhalation	1E-05	--	--	Arsenic, Chromium	2E+00	Manganese
		Dermal Contact	5E-07	--	--	--	3E-03	--
		Total	1E-05	--	--	Arsenic, Chromium	2E+00	Manganese
	Surface Water	Incidental Ingestion	8E-07	--	--	--	2E-02	--
		Inhalation		--	--	--		--
		Dermal Contact	5E-08	--	--	--	3E-02	--
		Total	9E-07	--	--	--	5E-02	--
	Sediment	Incidental Ingestion	0E+00	--	--	--	7E-03	--
		Inhalation	0E+00	--	--	--	1E-03	--
		Dermal Contact	0E+00	--	--	--	2E-03	--
		Total	0E+00	--	--	--	9E-03	--
	Groundwater	Incidental Ingestion	0E+00	--	--	--	3E-01	--
		Inhalation		--	--	--		--
		Dermal Contact	0E+00	--	--	--	2E-02	--
		Total	0E+00	--	--	--	4E-01	--
	Total All Media		2E-05				3E+00	
National Guard Resident	Surface Soil	Incidental Ingestion	8E-06	--	--	Arsenic	9E-02	--
		Inhalation	2E-06	--	--	Chromium	5E-01	--
		Dermal Contact	3E-06	--	--	Arsenic	2E-02	--
		Total	1E-05	--	--	Arsenic, Benzo(a)pyrene, Chromium	6E-01	--
	Surface Water	Incidental Ingestion	5E-06	--	--	Arsenic	2E-01	--
		Inhalation		--	--	--		--
		Dermal Contact	3E-07	--	--	--	2E-01	--
		Total	6E-06	--	--	Arsenic	3E-01	--
	Sediment	Incidental Ingestion	0E+00	--	--	--	4E-02	--
		Inhalation	0E+00	--	--	--	1E-01	--
		Dermal Contact	0E+00	--	--	--	1E-02	--
		Total	0E+00	--	--	--	2E-01	--
	Subsurface Soil	Incidental Ingestion	8E-06	--	--	Arsenic	5E-02	--
		Inhalation	1E-07	--	--	--	2E-04	--
		Dermal Contact	2E-06	--	--	Arsenic	2E-02	--
		Total	1E-05	--	Arsenic	--	7E-02	--
	Groundwater	Incidental Ingestion	0E+00	--	--	--	2E+00	Manganese
		Inhalation		--	--	--		--
		Dermal Contact	0E+00	--	--	--	1E-01	--
		Total	0E+00	--	--	--	2E+00	Manganese
	Total All Media		3E-05				3E+00	



Table 6-11
Summary of Cancer Risks and Hazard Indices

Receptor	Media	Exposure Route	Cancer Risk	Chemicals with Cancer Risks > 10 ⁻⁴	Chemicals with Cancer Risks > 10 ⁻⁵ and ≤ 10 ⁻⁴	Chemicals with Cancer Risks > 10 ⁻⁶ and ≤ 10 ⁻⁵	Hazard Index	Chemicals with HI > 1
Hunter	Surface Soil	Incidental Ingestion	2.7E-08	--	--	--	3E-04	--
		Inhalation	2.8E-10	--	--	--	5E-05	--
		Dermal Contact	5.3E-08	--	--	--	3E-04	--
		Total	8E-08	--	--	--	6E-04	--
	Sediment	Incidental Ingestion	0E+00	--	--	--	1E-04	--
		Inhalation	0E+00	--	--	--	5E-06	--
		Dermal Contact	0E+00	--	--	--	1E-04	--
		Total	0E+00	--	--	--	3E-04	--
	Surface Water	Incidental Ingestion	2E-08	--	--	--	6E-04	--
		Inhalation	--	--	--	--	--	--
		Dermal Contact	5E-09	--	--	--	3E-03	--
		Total	3E-08	--	--	--	3E-03	--
	Venison	Ingestion	2E-08	--	--	--	1E-03	--
	Total All Media		1E-07				4E-03	
Child Resident Farmer	Surface Soil	Incidental Ingestion	na	--	--	--	1E+00	--
		Inhalation	na	--	--	--	5E-02	--
		Dermal Contact	na	--	--	--	7E-02	--
		Total	na	--	--	--	1E+00	--
	Surface Water	Incidental Ingestion	na	--	--	--	1E-01	--
		Inhalation	na	--	--	--	--	--
		Dermal Contact	na	--	--	--	1E-01	--
		Total	na	--	--	--	2E-01	--
	Sediment	Incidental Ingestion	na	--	--	--	1E-01	--
		Inhalation	na	--	--	--	2E-03	--
		Dermal Contact	na	--	--	--	6E-03	--
		Total	na	--	--	--	1E-01	--
	Subsurface Soil	Incidental Ingestion	na	--	--	--	7E-01	--
		Inhalation	na	--	--	--	2E-05	--
		Dermal Contact	na	--	--	--	5E-02	--
		Total	na	--	--	--	8E-01	--
	Groundwater	Incidental Ingestion	na	--	--	--	1E+01	Manganese
		Inhalation	na	--	--	--	--	--
		Dermal Contact	na	--	--	--	4E-01	--
		Total	na	--	--	--	1E+01	Manganese
	Venison	Ingestion	na	--	--	--	5E-03	--
	Beef	Ingestion	na	--	--	--	6E-01	--
	Milk	Ingestion	na	--	--	--	2E+00	--
	Vegetables	Ingestion	na	--	--	--	3E+01	Aluminum, Arsenic, Chromium, Manganese, Mercury
	Total All Media		na				5E+01	



Table 6-11
Summary of Cancer Risks and Hazard Indices

Receptor	Media	Exposure Route	Cancer Risk	Chemicals with Cancer Risks > 10 ⁻⁴	Chemicals with Cancer Risks > 10 ⁻⁵ and ≤ 10 ⁻⁴	Chemicals with Cancer Risks > 10 ⁻⁶ and ≤ 10 ⁻⁵	Hazard Index	Chemicals with HI > 1
Adult Resident Farmer	Surface Soil	Incidental Ingestion	na	--	--	--	1E-01	--
		Inhalation	na	--	--	--	2E-02	--
		Dermal Contact	na	--	--	--	7E-02	--
		Total	na	--	--	--	2E-01	--
	Surface Water	Incidental Ingestion	na	--	--	--	3E-02	--
		Inhalation	na	--	--	--		--
		Dermal Contact	na	--	--	--	6E-02	--
		Total	na	--	--	--	9E-02	--
	Sediment	Incidental Ingestion	na	--	--	--	1E-02	--
		Inhalation	na	--	--	--	8E-04	--
		Dermal Contact	na	--	--	--	7E-03	--
		Total	na	--	--	--	2E-02	--
	Subsurface Soil	Incidental Ingestion	na	--	--	--	8E-02	--
		Inhalation	na	--	--	--	2E-04	--
		Dermal Contact	na	--	--	--	5E-02	--
		Total	na	--	--	--	1E-01	--
	Groundwater	Incidental Ingestion	na	--	--	--	3E+00	--
		Inhalation	na	--	--	--		--
		Dermal Contact	na	--	--	--	2E-01	--
		Total	na	--	--	--	3E+00	Manganese
	Venison	Ingestion	na	--	--	--	1E-03	--
	Beef	Ingestion	na	--	--	--	6E-01	--
	Milk	Ingestion	na	--	--	--	3E-01	--
	Vegetables	Ingestion	na	--	--	--	3E+01	Aluminum, Arsenic, Chromium, Manganese, Mercury
	Total All Media		na				4E+01	
Lifelong Resident Farmer (Child and Adult)	Surface Soil	Incidental Ingestion	4E-05	--	Arsenic	Benzo(a)pyrene, Dibenzo(a,h)anthracene		--
		Inhalation	2E-07	--	--	--		--
		Dermal Contact	1E-05	--	--	Arsenic, Benzo(a)pyrene, Dibenzo(a,h)anthracene		--
		Total	5E-05	--	Arsenic	Benzo(a)pyrene, Dibenzo(a,h)anthracene		--
	Surface Water	Incidental Ingestion	2E-06	--	--	Arsenic		--
		Inhalation		--	--	--		--
		Dermal Contact	2E-07	--	--	--		--
		Total	2E-06	--	--	Arsenic		--
	Subsurface Soil	Incidental Ingestion	4E-05	--	Arsenic	--		--
		Inhalation	1E-07	--	--	--		--
		Dermal Contact	1E-05	--	--	--		--
		Total	5E-05	--	Arsenic	--		--
	Venison	Ingestion	3E-08	--	--	--		--
	Beef	Ingestion	2E-04	Dibenzo(a,h)anthracene	Arsenic, Benzo(a)pyrene	--		--
	Milk	Ingestion	1E-03	Benzo(a)pyrene, Dibenzo(a,h)anthracene	Arsenic	--		--
	Vegetables	Ingestion	3E-03	Arsenic, Benzo(a)pyrene, Dibenzo(a,h)anthracene	--	--		
	Total All Media		5E-03					



1
2
3

Table 6-12
Non-Carcinogenic Risks for LL-9

Receptor⁽¹⁾	Medium	Hazard Index	Table in Appendix V
Security Guard/Maintenance Worker (Current/Future)	Surface Soil	4E-02	9.1
National Guard Trainee (Future)	Deep surface Soil	2E+00	9.2
	Sediments	8E-03	9.2
	Surface Water	4E-02	9.2
	Ground Water	2E-01	9.2
Total (All Media)		3E+00	9.2
Total (Minus Deep Surface Soils)		3E-01	9.2
National Guard Resident	Surface Soils	1E-01	9.3
	Subsurface Soils	7E-02	9.3
	Sediment	5E-02	9.3
	Surface water	3E-01	9.3
	Groundwater	1E+00	9.3
Total (All Media)		2E+00	9.3
Total (All Media Minus GW)		4E-01	9.3
Hunter/Trapper (Current/Future)	Surface Soils	5E-04	9.4
	Sediments	2E-04	9.4
	Surface water	2E-03	9.4
	Venison	4E-04	9.4
Total (All Media)		3E-03	9.4
Hypothetical Child Resident Farmer (Future)	Surface Soils	9E01	9.5
	Subsurface	7E-01	9.5
	Sediments	5E-01	9.5
	Surface water	2E+00	9.5
	Groundwater	6E+00	9.5
Total (All Media)		9E+00	9.5
Total (Minus GW)		3E+00	9.5
Total (Minus GW and SS)		3E+00	9.5



Table 6-12 (Continued)
Non-Carcinogenic Risks for LL-9

Receptor ⁽¹⁾	Medium	Hazard Index	Table in Appendix V
Total (Minus GW and SB)		3E+00	9.5
Hypothetical Adult Resident Farmer (Future)	Surface Soils	2E-01	9.6
	Subsurface Soils	1E-01	9.6
	Sediments	9E-02	9.6
	Surface water	5E-01	9.6
	Groundwater	2E+00	9.6
Total (All Media)		3E+00	9.6
Total (Minus GW)		8E-01	9.6
Total (Minus GW and SS)		6E-01	9.6
Total (Minus GW and SB)		7E-01	9.6

1. More than one total risk estimate has been presented for several receptors to demonstrate the relative contribution of an environmental medium to the total risk for a receptor and because of the potential overestimation of risk that may occur when a receptor is assumed to be exposed to both surface and subsurface soils and sediments at a similar daily ingestion rate (i.e., the National Guard trainee, National Guard resident trainer, the hypothetical future farm resident). Also, risk estimates have been presented for domestic use of groundwater; however, it is very unlikely that the shallow groundwater resources at LL-9 would be used for domestic purposes.

Totals in Table 6-12 are shown for all media, as well as subtotals subtracting one or more media. These subtotals help illustrate which medium or media are responsible for any exceedances and are presented for informational purposes only.

HIs developed for the aforementioned receptors and presented in the referenced tables were as follows:

HIs calculated for the security guard/maintenance worker and the hunter/trapper are less than 1, indicating that adverse non-carcinogenic health effects are not anticipated under the conditions established in the exposure assessment.

The HIs calculated for the National Guard trainee receptor exposed to COPCs in groundwater, sediments, and surface water do not exceed 1. The HI for the National Guard trainee exposed to COPCs in deep surface soils (0 to 4 ft bgs) is 2. Manganese (inhalation of air particulates as the route of exposure) was the only significant contributor to the non-carcinogenic risk (i.e., risk



driver) for the National Guard trainee. The HI calculated by summing the HIs calculated for all other COPCs does not exceed 1; only the HI calculated for the central nervous system (the target organ for manganese) exceeds 1. The non-cancer risk estimates for manganese are strongly affected by the particulate emission factor (PEF) specified for the National Guard Trainee receptor in the RVAAP FWHHRAM. The EPC for manganese (633 mg/kg) in soils is less than the surface soil background value of 1,450 mg/kg. The PEF, background concentrations, and the inhalation reference dose for aluminum and manganese are further discussed in Section 6.6.

Manganese at a concentration equal to RVAAP background would result in an HI of approximately 4, and at a concentration equal to its Region 9 residential PRG would result in an HI of 5.

HIs calculated for the National Guard resident trainer exceed 1, if it is assumed the receptor is using the shallow groundwater resource for domestic purposes. As discussed in Section 4, the use of the shallow groundwater at LL-9 for domestic purposes is considered unlikely currently, although OHARNG has expressed interest in drilling potable and non-potable wells at RVAAP. Manganese in groundwater is the main risk driver. The National Guard resident trainer receptor was not required by the RVAAP FWHHRAM (USACE, 2004). The receptor was added to this HHRA as the result of discussions with the USACE and Ohio EPA because risk estimates for this receptor may assist the risk management team when making risk management decisions for LL-9.

The HI calculated for the adult resident farmer routinely exposed to COPCs in surface soil, surface water, sediment, and groundwater, or routinely exposed to COPCs in subsurface soil, surface water, sediment, and groundwater via the direct-contact pathways, exceed 1 only when it is assumed that the shallow groundwater is used for domestic purposes. Manganese in groundwater is the only risk driver. The HIs calculated for the adult resident farmer routinely exposed via indirect exposure pathways (i.e., consumption of foodstuffs), also presented in Appendix V, Table 9.6, do exceed 1 for vegetable consumption only. However, there are significant sources of uncertainty associated with the risk estimates developed for the ingestion of foodstuffs exposure pathways. Consequently, these risk assessment results are included and discussed in Section 6.6.

The HI calculated for the child resident farmer routinely exposed to COPCs in surface soil, sediment, surface water, and groundwater or routinely exposed to COPCs in subsurface soil, sediment, surface water, and groundwater via the direct-contact exposure pathways exceed 1. The HIs calculated for the COPCs in groundwater and surface water exceed 1. The HIs calculated for sediments or subsurface soils do not exceed 1. Manganese in groundwater is the primary risk driver; HIs calculated for other COPCs evaluated do not exceed 1. The target-organ-specific HI calculated for the central nervous system (the target system for manganese)



exceed 1, as well as the HIs for the target systems for arsenic, skin, and cardiovascular. The HIs calculated for the child resident farmer routinely exposed via indirect exposure pathways (i.e., consumption of foodstuffs) also included in Appendix V, Table 9.5; do exceed 1 for vegetable and milk consumption. However, there are significant sources of uncertainty associated with the risk estimates developed for the ingestion of foodstuffs exposure pathways. Consequently, these risk assessment results are included and discussed in Section 6.6.

6.5.2.1 Carcinogenic Risks for LL-9

ILCR estimates calculated for the aforementioned receptors and listed in the referenced tables are shown in Table 6-13.

Table 6-13

Incremental Lifetime Cancer Risk Results

Receptor ⁽¹⁾	Media	Cancer Risk Estimate	Table in Appendix V
Security Guard/ Maintenance Worker (Current/Future)	Surface Soil	8E-06	9.1
National Guard Trainee (Future)	Deep Surface Soil	2E-05	9.2
	Sediment	0	9.2
	Surface Water	9E-07	9.2
	Groundwater (GW)	0	9.2
	Total All Media	2E-05	9.2
	Total (Minus Deep Surface Soil)	9E-07	9.2
National Guard Resident (Future)	Surface Soil (SS)	1E-05	9.3
	Subsurface Soil (SB)	9.5E-06	9.3
	Sediment (SD)	0	9.3
	Surface Water	6E-06	9.3
	Groundwater	0	9.3
¹ Total (All Media)		2E-05	9.2
¹ Total (Minus Deep Surface Soil)		9E-07	9.2
Hunter/Trapper (Current/Future)	Surface Soil	8E-08	9.5
	Sediment	0	9.5
	Surface Water	3E-08	9.5
	Venison	2E-08	9.5
Total (All Media)		1E-07	9.5
Hypothetical Resident/Trainer (Future)	Surface Soil (SS)	5E-05	9.8
	Subsurface Soil (SB)	5E-05	9.8
	Sediment	0	9.8
	Surface Water	2E-05	9.8



Table 6-13 (Continued)

Incremental Lifetime Cancer Risk Results

Receptor ⁽¹⁾	Media	Cancer Risk Estimate	Table in Appendix V
	Groundwater (GW)	0	9.8
¹ Total (All Media Minus Subsurface Soil)		7E-05	9.8
¹ Total (All Media Minus Surface Soil)		6E-05	9.8

1. More than one total risk estimate has been presented for several receptors to demonstrate the relative contribution of an environmental medium to the total risk for a receptor and because of the potential overestimation of risk that may occur when a receptor is assumed to be exposed to both surface and subsurface soils and sediments at a similar daily ingestion rate (e.g., the National Guard Trainee, National Guard resident trainer, the hypothetical future farm resident). Also, risk estimates have been presented for domestic use of groundwater; however, it is very unlikely that the shallow groundwater resource at LL-9 would be used for domestic purposes.

Totals in Table 6-13 are shown for all media, as well as subtotals subtracting one or more media. These subtotals help illustrate which medium or media are responsible for any exceedances and are presented for informational purposes only.

ILCR estimates calculated for the hunter/trapper do not exceed 1E-06, the conservative end of the USEPA target risk range. The cancer risk estimates for the security guard/maintenance worker are within the USEPA target risk range of 1E-04 to 1E-06 and do not exceed the Ohio EPA cancer risk benchmark of 1E-05. The cancer risk estimates calculated for arsenic, benzo(a)pyrene, and dibenzo(a,h)anthracene exceed 1E-06.

Cancer risk estimates for the National Guard trainee are within the USEPA target risk range of 1E-04 to 1E-06 but exceed the Ohio EPA cancer risk benchmark of 1E-05. Risk estimates summed for receptor exposure to surface water do not exceed 1E-06. There is no cancer risk from subsurface soil or groundwater. Inhalation of chromium in deep surface soil accounts for 1.0E-05, or 69% of the total cancer risk estimate for the National Guard trainee. Slightly less significant, the inhalation of arsenic in deep surface soil accounts for 2.9E-06, or 19% of the total cancer risk estimate for the National Guard trainee. Inhalation of both chromium and arsenic accounts for 88% of total cancer risk. The estimates for inhalation of chromium and arsenic in deep surface soil are strongly affected by the PEF specified for the National Guard trainee receptor in the RVAAP FWHHRAM. This PEF is further discussed in Section 6.6. The EPC for arsenic in deep surface soil (13 mg/kg) does not exceed the RVAAP background value of 15.4 mg/kg. The EPC of chromium in deep surface soil (17 mg/kg) does not exceed the RVAAP background value of 17.4 mg/kg; nor does it exceed the chromium Region 9 residential PRG of 30 mg/kg.



Cancer risk estimates for the National Guard resident trainer are within the USEPA target risk range of $1E-04$ to $1E-06$ but exceed the Ohio EPA cancer risk benchmark of $1E-05$. The risk estimates for arsenic (surface soil, subsurface soil) and benzo(a)pyrene (surface soils) exceed $1E-06$. The National Guard resident trainer receptor was not required by the RVAAP FWHHRAM. This receptor was added to this HHRA as the result of discussions with the USACE and Ohio EPA because risk estimates for this receptor may assist the risk management team when making risk management decisions for LL-9.

Cancer risk estimates for the hypothetical future resident farmer (combined adult and child) are within the USEPA target risk range of $1E-04$ to $1E-06$ but exceed the Ohio EPA cancer risk benchmark of $1E-05$. Risk estimates developed for surface soils, subsurface soils, and surface water exceed the Ohio EPA $1E-05$ risk benchmark. A review of the media and chemical-specific risk results indicate the following chemicals are primary contributors to the estimated cancer risk:

- Arsenic is the primary risk driver in soil and surface water. Risk drivers are the most significant risk contributors and are typically those COPCs whose contribution to the risk estimates causes the overall receptor or media risk exceeding federal or state risk benchmarks. The risk estimates for arsenic in soils and surface waters exceed $1E-05$. EPCs of arsenic in surface soil (12 mg/kg), subsurface soil (13.8 mg/kg), and deep surface soil (13 mg/kg) were less than their respective RVAAP background concentrations (15.4 mg/kg, 19.8 mg/kg, and 15.4 mg/kg).
- The risk estimates for benzo(a)pyrene in surface soil exceed $1E-06$, but do not exceed $1E-05$.

The cancer risk estimates presented for the environmental media are subject to several significant sources of uncertainty, which are further discussed in Section 6.6. Section 6.6 also presents an evaluation of the ingestion of foodstuffs exposure pathways.

6.5.2.2 Summary of COCs for Each Media/Receptor

Chemicals of concern (COCs) are the COPCs that significantly contribute to a pathway in a use scenario for a receptor that either exceeds the $1E-05$ cumulative cancer risk benchmark or exceeds a non-carcinogenic HI (target organ specific) of 1.0. A summary of COCs for each of the land use/receptor/medium combinations with COCs is listed in Table 6-14. Remedial goal options (RGOs) are calculated and included in Section 6.7 for the potential COCs identified in the risk evaluation.



6.5.2.3 Lead Exposure Analysis

The maximum lead concentration (70 µg/L) in surface water exceeded the SDWA action level of 15 µg/L for residential land use. Concentrations of lead exceeded the action level in two samples. Exposures to lead are evaluated using arithmetic mean concentrations (USEPA, 1994c). The mean concentration of lead in surface water (25.5 µg/L) is slightly above the action level of 15 µg/L, indicating that using the surface water as a drinking water source may be of some concern at LL-9. However, surface water in the ditches at LL-9 will not be used as a drinking water source. Dermal contact (bathing and showering, and washing dishes or clothes) should be safe, even if the water contains lead over the USEPA action level, because human skin does not absorb lead from the water (MWCOG, 2004).

The maximum lead concentration (1330 mg/kg) in surface soil (and, therefore, deep surface soil) exceeded the OSWER soil screening level of 400 mg/kg for residential land use (USEPA, 1994c). Concentrations of lead exceeded the residential screening level in only one soil sample (LL9SS-011-0001-SO). Exposures to lead are evaluated using arithmetic mean concentrations (USEPA, 1994c). The mean concentrations of lead in surface soil (63 mg/kg) and deep surface soil (44 mg/kg) are well below the residential screening level of 400 mg/kg. Consequently, no adverse health effects are anticipated from exposures to lead in surface soil or deep surface soils at LL-9.

6.5.2.4 Sample Location of Concern

Maximum detected concentrations of surface soil and deep surface soil COPCs copper (1,240 mg/kg), lead (1,330 mg/kg), and mercury (882 mg/kg) were at azide sample location LL9SS-011-0001-SO. The next highest detected concentrations were much lower (copper – 170 mg/kg, lead – 320 mg/kg, and mercury – 17 mg/kg). This sample location may be of particular concern when making risk decisions, particularly for exposure to mercury.

6.6 Uncertainty Analysis

A significant uncertainty is the fact that the nature and extent of the contamination in the environmental media at LL-9 has not been completely determined. Therefore, the human health risk assessment presented in Section 6 (and the associated uncertainties) is preliminary and subject to change based on the data collected to resolve the RI data gaps for LL-9. This section summarizes the uncertainties inherent in the BHHRA. It includes a discussion of how uncertainties may affect the quantitative risk estimates and conclusions of the risk analysis. The BHHRA for LL-9 was performed in accordance with current USEPA and RVAAP guidance and the aforementioned White Paper developed for LL-9 (TtNUS, 2004). However, there are varying degrees of uncertainty associated with the BHHRA. The following sections discuss general uncertainties in all risk assessments and uncertainties specific to the BHHRA for LL-9.

Uncertainty in the selection of COPCs can be related to the quality of the analytical data bases, groupings of samples, and procedures used to include or exclude constituents as COPCs.



Table 6-14
Potential Land Use/Receptor/Medium Pathway COCs

Pathway	Surface Soil	Subsurface Soil	Surface Water	Deep Surface Soil		Groundwater	Soil - Beef			Soil - Milk			Soil - Vegetables					
	As	As	As	Cr	Mn	Mn	As	B(a)P	D(ah)A	As	B(a)P	D(ah)A	Al	B(a)P	D(ah)A	As	Cr	Mn
Security Guard/Maintenance Worker		--	--		--	--	--			--			--			--	--	--
National Guard Trainee	--	--		X	X		--			--			--			--	--	--
National Guard Resident				--	--	X	--			--			--			--	--	--
Hunter		--		--	--	--	--			--			--			--	--	--
Child Resident Farmer				--	--	X							X			X	X	X
Adult Resident Farmer				--	--	X							X			X	X	X
Lifelong Resident Farmer	X	X	X	--	--		X	X	X	X	X	X		X	X	X		

No COCs were identified for direct contact with surface water and ingestion of venison.

As - Arsenic
Cr - Chromium
Mn - Manganese
B(a)P - Benzo(a)pyrene
D(ah)A - Dibenzo(a,h)anthracene
"X" indicates chemical is a COC for receptor/medium
"--" indicates pathway not applicable in this BHHRA.
A blank indicates chemical is not identified as a COC for receptor/medium.



1 Uncertainties associated with the exposure assessment include the values used as input
2 variables for a given intake route/scenario, the assumptions made to determine EPCs, and the
3 predictions regarding future land use and population characteristics. Uncertainty in the toxicity
4 assessment includes the quality of the existing toxicity data needed to support dose-response
5 relationships and the weight of evidence used for determining the carcinogenicity of COPCs.
6 Uncertainty in the risk characterization includes that associated with exposure to multiple
7 chemicals and the cumulative uncertainty from combining conservative assumptions made in
8 earlier steps of the risk assessment process.

9 Whereas there are various sources of uncertainty, the direction of uncertainty can be influenced
10 by the assumptions made throughout the risk assessment, including selection of COPCs and
11 selection of values for dose-response relationships. In general, assumptions are made
12 considering safety factors so that the final calculated risks for a receptor are not underestimated
13 and, in fact, such receptor risk estimates tend to be over-estimated.

14 After the risk assessment is complete, the results must be reviewed and evaluated to identify
15 the type and magnitude of uncertainty involved. Reliance on results from a risk assessment
16 without consideration of the uncertainties, limitations, and assumptions inherent in the process
17 can be misleading. For example, to account for uncertainties in the development of exposure
18 assessment, exposure assumptions (e.g., ingestion rates, exposure frequencies) must be
19 estimated conservatively to protect sensitive subpopulations or the maximum exposure
20 individuals. However, if a number of conservative assumptions are combined in an exposure
21 model, the resulting calculations can propagate the uncertainties associated with those
22 assumptions, thereby producing a much larger uncertainty for the final result. This uncertainty
23 is biased toward over-predicting both carcinogenic and non-carcinogenic risks. Thus, both the
24 results of the risk assessment and the uncertainties associated with those results must be
25 considered when making risk management decisions.

26 The evaluation of uncertainty is especially relevant when risk estimates exceed the point of
27 departure for defining “acceptable” risk. For example, when risks calculated using a high
28 degree of uncertainty are less than an acceptable risk level (e.g., 1×10^{-6}), the interpretation of
29 “no significant risk” is typically straightforward. However, when risk calculated using a high
30 degree of uncertainty exceeds an “acceptable” risk level, a conclusion can be difficult unless
31 uncertainty is considered especially when the exceedance of acceptable risk is relatively
32 “marginal.”

33 The following subsections discuss uncertainties associated with each major component of the
34 risk assessment: COPC selection, exposure assessment, toxicity assessment, and risk
35 characterization.



6.6.1 Uncertainties Associated with COPC Selection

The following issues may contribute to uncertainty in COPC selection for LL-9: the quality of the existing databases, inclusion of chemicals potentially attributable to background, toxicity screening levels used, absence of screening levels for a few chemicals detected in the site media, and frequency of detection reported for chemicals detected in the environmental media. A brief discussion of each of these issues is provided in the remainder of this section.

6.6.1.1 Existing Databases

All data used for this evaluation have been validated according to National Functional Guidance (USEPA, 1994a and 1994b). Therefore, uncertainties associated with the quality of the data are considered to be minimal. For most media (groundwater, surface water, sediment, and surface soil and subsurface soil [organic parameters only]), few samples (less than 10) were collected. The availability of small data sets only for these media may result in uncertainty both in terms of COPC selection and in the calculated risks. However, the LL-9 field sampling program was biased toward areas most likely to demonstrate contamination. Thus, receptor risk based on these biased data may be over-estimated rather than under-estimated.

Selected media samples collected during the LL-9 investigation were sent to a second laboratory, GPL, to substantiate the accuracy of the new GPL screening method for explosives utilized during the field investigation program (modified 8330). Because the GPL method is unproven, data results from this screening are presented in appendices but were not used in any risk evaluations. Surface soil data are presented in Appendix F, subsurface soil data are presented in Appendix I, and sediment data are presented in Appendix R. Several explosives detected using the GPL screening method were also detected in samples that were evaluated in the BHHRA and therefore may be present at LL-9. However, because the maximum reported concentrations of these explosives do not exceed their respective Region 9 residential PRGs, the exclusion of these screening samples from the risk evaluation is unlikely to lead to an underestimation of risk.

6.6.1.2 Chemicals Potentially Attributable to Background

The background values for RVAAP were determined as described in RVAAP WBG Phase II Remedial Investigation (SAIC, 2001). Outliers, results exceeding an upper cutoff limit, were removed from the background data set. The upper cutoff limit was the third quartile (the 75th percentile) plus 1.5 times the inter-quartile range. An outlier might indicate a mistake or anomaly in the data set. However, it could also represent a perfectly valid but rare result. Conservatively, some higher background values were removed from the background data set. This conservative approach to the selection of background values and the data discussions presented in the following paragraphs suggest that some chemicals selected as COPCs may marginally exceed background concentrations (if at all).



Arsenic EPCs calculated for surface soil (12 mg/kg), subsurface soil (13.8 mg/kg), and deep surface soil (13 mg/kg) were less than their respective background concentrations (15.4 mg/kg, 19.8 mg/kg, and 15.4 mg/kg)

Only 1 of 46 samples in surface soil and 3 of 85 samples in deep surface soil had aluminum concentrations greater than the background value, and the EPC for aluminum was less than its background concentration in both data sets. Only 2 of 46 surface soil samples and 2 of 85 deep surface soil samples had manganese concentrations greater than the background value. The EPC for manganese was less than its background concentration in both data sets. Only one groundwater sample had a manganese concentration greater than the background value.

The EPCs for chromium in surface soil, subsurface soil, and deep surface soil were less than its background value.

Finally, benzo(a)pyrene was detected in background soil samples at a maximum concentration of 100 µg/kg. The maximum detected concentration in surface soils was 240 µg/kg. As indicated previously, the PAH concentrations detected in LL-9 soil and sediment are within the range of background anthropogenic concentrations reported in the literature.

6.6.1.3 COPC Screening Levels

The use of risk-based screening values should ensure that the significant contributors to risk at a site are not eliminated but are retained for risk evaluation. COPC screening values were based on conservative land use scenarios (e.g., residential land use for soil) and protective levels of risk corresponding to ILCRs of 1×10^{-6} and HIs of 0.1. An exceedance of these conservative screening values is not conclusive evidence that a receptor is at risk. For example, as discussed in Section 6.2, non-carcinogenic chemicals selected as COPCs (e.g., aluminum and copper in surface soil) were not detected in environmental media at concentrations exceeding the Region 9 residential PRGs. Consequently, the aluminum and copper concentrations in the surface soils are unlikely to be significant from a human health perspective.

In addition, the toxicity values used in the derivation of PRGs are subject to change as additional information (from scientific research) becomes available. These periodic changes in toxicity values may cause the PRG values to change as well.

6.6.1.4 Absence of COPC Screening Levels

Essential human nutrients (e.g., magnesium, potassium, calcium, iron, and sodium) do not have toxicity screening levels for COPC selection. These nutrients were eliminated from consideration as COPCs (see Section 6.2.2). Exclusion of these chemicals as COPCs is not expected to add significant uncertainty to the risk estimates.



6.6.1.5 *Frequency of Detection*

No chemicals were eliminated as SRCs or COPCs on the basis of frequency of detection considerations described in Section 6.2.2.

6.6.1.6 *Data Analysis*

Uncertainty can be added to the risk estimates as a result of the limitations in the analytical methods. Some current analytical methods are limited in their ability to achieve detection limits at or below PRGs. Risks may be overestimated when some analyte concentrations are reported as non-detected at the method detection limit, but the actual concentration is much less than the method detection limit. Conversely, risks may be underestimated when some analytes are present, even at concentrations exceeding the Region 9 residential PRG, but are not detected because of limitations in the analytical methods and therefore the analytes are removed from the SRC list.

6.6.2 Uncertainties Associated with the Exposure Assessment

Uncertainty in the exposure assessment arises because of the methods used to calculate EPCs, selection of receptors, selection of land use scenarios to be evaluated, and selection of exposure parameters.

6.6.2.1 *Land Use*

The current land use patterns of LL-9 are established, thereby reducing the uncertainty associated with current land use assumptions. The anticipated future land use patterns of LL-9 are described in the RVAAP FWHHRAM. This BHHRA is based on these known and anticipated future land use scenarios; therefore, the land use assumptions presented herein are not expected to lead to an underestimation of risk.

6.6.2.2 *Exposure Point Concentration*

The maximum concentrations of some COPCs (e.g., all COPCs in surface water and groundwater and organics in soils) were used as the EPCs to quantify potential risks. Risk estimates are likely to be overestimated when the maximum detected concentration is selected as the EPC because it is unlikely that potential receptors would be exposed to the maximum concentration over the entire site for the assumed exposure period.

6.6.2.3 *Exposure Parameters*

The exposure assessment factors (e.g., exposure frequency and duration) utilized in the BHHRA are based on reasonable maximum exposure (RME) assumptions. Generally, these



exposure factors are based on data available as a result of surveys of the physiological and lifestyle characteristics of the general population of the United States. The attributes and activities studied in these surveys generally have a broad distribution. RME exposure factor values were used in this risk evaluation to avoid an underestimation of risk. Risk is not likely to be underestimated for reasonably maximum exposed individuals and is more likely to be overestimated for the general population exposed to chemicals in environmental media at the site when RME assumptions are evaluated in a risk assessment. The following paragraphs exemplify the conservative nature of the exposure factors selected for the BHHRA.

A particulate emission factor (PEF) of 9.24×10^8 m³/kg was calculated based on USEPA's *Soil Screening Guidance* (USEPA, 1996) using site-specific factors for Cleveland, Ohio, the city nearest Ravenna for which USEPA provided parameters to estimate PEF and volatilization factor (VF) values. This value was used to evaluate most receptors in this BHHRA. However, the RVAAP recommends a PEF of 1.67×10^6 m³/kg for the National Guard trainee whose activities are considered more likely to generate more airborne dust than the activities of other receptors. The FWHHRAM states that this PEF was calculated from a dust loading factor (DLF) of 600 µg/m³. This value greatly exceeds the National Ambient Air Quality Standard (NAAQS) PM 10 (particulate matter less than or equal to 10 microns in diameter) 24-hour and annual standards of 150 µg/m³ and 50 µg/m³, respectively. Although it is possible that National Guard activities could generate significant particulate matter less than or equal to 10 microns in diameter (PM10) for short periods of time and at locations adjacent to vehicles, it is very unlikely that these activities would generate ambient air particulate emissions for extended periods of time and across the entire site, especially at concentrations that exceed the NAAQS. As a point of comparison, use of the methods and default values for estimating dust emissions associated with construction activities in the USEPA *Supplemental Guidance for Developing Soil Screening Levels for Superfund Sites*, December 2002, results in a DLF of 364 µg/m³. The HI for the National Guard trainee calculated based on COPC concentrations in the LL-9 deep surface soil and sediment using this guidance would be approximately half the value presented in Section 6.5.

No exposure time term is included in the intake equation for the dermal contact with soil exposure pathway. The underlying assumption is that the receptor does not bathe or remove soil from the skin surface for an extended period (e.g., the full 24 hours of the day). This may overestimate the risk associated with dermal contact with soil. This assumption is especially important when the dermal pathway is the major contributor to the risks and/or hazards, as it is for some receptors at LL-9. For example, the calculated risk from dermal exposure to benzo(a)pyrene in deep surface soil by the National Guard trainee is greater than the risk calculated from ingestion of deep surface soil.

Based on the climatic conditions in the eastern Ohio region, the assumption of 250 days per year and 350 days per year as exposure frequencies for the National Guard resident trainer and



hypothetical future resident farmer, respectively, for soil and sediment likely results in an overestimation of risk because no exposure is likely to occur when these media are covered with snow or frozen solid. Consequently, risks calculated using these exposure assumptions are likely over-estimated. Likewise, the assumption of 250 days per year as an exposure frequency for the National Guard resident trainer contact with COPCs in surface water is very conservative for eastern Ohio and may lead to an overestimation of risk.

6.6.3 Uncertainties Related to Toxicity Information

Uncertainties associated with the toxicity assessment include the derivation of RfDs and CSFs from which the PRGs are calculated and limitations associated with the use of available criteria. These uncertainties are discussed in the following subsections.

6.6.3.1 Derivation of Toxicity Criteria

Uncertainty is associated with hazard assessment and dose-response evaluations. The hazard assessment deals with characterizing the nature and strength of the evidence of causation, or the likelihood that a chemical that induces adverse effects in animals will also induce adverse effects in humans. Hazard assessment of carcinogenicity is evaluated as a weight-of-evidence determination using USEPA methods. Positive animal cancer test data suggest that humans contain tissue(s) that may manifest a carcinogenic response; however, the animal data cannot necessarily be used to predict the target tissue in humans. In the hazard assessment of non-cancer effects however, the positive animal data often suggest the nature of the effects (i.e., the target tissues and type of effects) anticipated for humans.

Uncertainty in hazard assessment arises from the nature and quality of the toxicological information available as a result of both animal and human studies. Uncertainty is reduced when:

- Similar effects are observed across species, strain, sex, and exposure route;
- The magnitude of the response is clearly dose-related;
- Pharmacokinetic data indicate a similar fate in humans and animals;
- Postulated mechanisms of toxicity are similar for humans and animals; and
- The chemical of concern is structurally similar to other chemicals for which the toxicity is adequately characterized.

Uncertainty in the dose-response evaluation is associated with the determination of CSFs for the carcinogenic assessment and derivation of RfDs or RfCs for the non-carcinogenic assessment. Uncertainty introduced from inter-species (animal to human) extrapolation, which, in the absence of quantitative pharmacokinetic or mechanistic data, is usually based on



consideration of inter-species differences in basal metabolic rate. Uncertainty also results from intra-species (i.e., variation within a species). Most toxicity experiments are performed with animals that are very similar in age and genotype so that intra-group biological variation is minimal, but the human population of concern may reflect a great deal of heterogeneity including unusual sensitivity or tolerance to the COPC. Even toxicity data from human occupational exposure reflect a bias because only those individuals sufficiently healthy to attend work regularly (the healthy worker effect) and those not unusually sensitive to the chemical are likely to be exposed to chemicals in the workplace. Finally, uncertainty arises from the quality of the key study from which the quantitative estimate (i.e., the CSF or the RfD) is derived and the supporting database of toxicity information. For cancer effects, the uncertainty associated with dose-response factors is somewhat mitigated by assuming the 95 percent upper bound for the slope factor. Another source of uncertainty in carcinogenic assessment is the method by which data from high doses in animal studies are extrapolated to the dose range expected for environmentally exposed humans. The linearized multistage model, which is used in most quantitative estimations of human risk from animal data, is based on a non-threshold assumption of carcinogenesis. Evidence suggests however, that epigenetic carcinogens, as well as many genotoxic carcinogens, have a threshold below which they are non-carcinogenic (Williams and Welsburger, 1991). Therefore, the use of the linearized multistage model is conservative for chemicals that exhibit a threshold for carcinogenicity.

For non-cancer effects, additional uncertainty factors are often applied in the derivation of RfDs or RfCs to compensate for poor quality data for key study group or gaps in the database. Uncertainty factors are usually applied to the estimate of a no-effects level when lowest-adverse-effect level data only are available. Additional uncertainty arises in estimation of RfDs or RfCs for chronic exposure from subchronic data. Unless empirical data indicate that effects do not worsen with increasing duration of exposure, an additional uncertainty factor is applied to the no-effect level in the subchronic study when deriving toxicity criteria for chronic exposure. Uncertainty in the derivation of RfDs is also addressed by the use of modifying factors that normally range between 3 and 10. The resulting combination of uncertainty and modifying factors are used proportionally to adjust the RfD downward and thereby intentionally often introduce a conservative bias in the RfD by a factor of 1,000 or more.

The derivation of dermal RfDs and CSFs from oral values may also cause uncertainty. This is particularly the case when no gastrointestinal absorption rates are available in the literature or when only qualitative statements regarding absorption are available.

6.6.3.2 Chromium Toxicity Criteria

Some uncertainty is associated with the evaluation of chromium, which was assumed to be present in its hexavalent state. The estimated risks from chromium for the National Guard trainee exceeded 1×10^{-5} only because chromium was assumed to be present in the hexavalent form. Because hexavalent chromium is considered to be more toxic than trivalent chromium



and the latter is more commonly found in the environment, risks for this chemical are probably over estimated.

6.6.3.3 *Toxicity Criteria for Arsenic*

The toxicity criteria for arsenic are a major source of uncertainty in this BHHRA. Although conventional risk assessment methods suggest that there is no “zero risk concentration” for a carcinogen such as arsenic, the human body does have a limited capacity to methylate arsenic, and this limit is not generally reached until the body's intake of arsenic exceeds 500 µg per day. Most environmental exposures result in intakes lower than 500 µg per day. Additionally, the USEPA suggests that an order-of-magnitude adjustment of risk (downward) may be appropriate for arsenic in some cases (USEPA, 1987a).

6.6.3.4 *Toxicity Criteria for Aluminum*

A provisional RfD is available for aluminum (a COPC in soil and sediment) from the NCEA; however, it is based on typical allowable intakes rather than adverse effect levels and is not considered risk based. This is likely to be a significant source of uncertainty in the BHHRA because aluminum is a significant COPC for LL-9 (although the maximum detected concentration does not exceed the Region 9 residential PRG and the metal was detected infrequently at concentrations exceeding the RVAAP background benchmarks). Aluminum at the background soil concentrations would result in an HI of 17.

6.6.4 **Uncertainties and Assumptions in the Risk Characterization**

Uncertainty in risk characterization often results from the lack of toxicity criteria and from assumptions made regarding additive effects of exposure to multiple COPCs from various exposure routes. For example, high uncertainty exists when summing cancer risks for several substances across different exposure pathways. This assumes that each substance has a similar effect, mode of action, or both. Often compounds affect different organs, have different mechanisms of action, and differ in fate in the body; therefore, additivity may not be an appropriate assumption. However, the assumption of additivity is often made to produce a conservative estimate of risk. The risk characterization also does not consider antagonistic or synergistic effects of COPCs. Antagonistic effects (i.e., the toxic effect of a chemical is mitigated by the presence of another chemical) could mean the risk is overestimated, whereas synergistic effects (i.e., the toxic effect of a chemical is magnified by the presence of another chemical) could mean the risk is underestimated.



6.6.4.1 *Unavailable or Provisional Toxicity Criteria*

Risk-based screening levels and/or toxicity criteria are not available for nitrocellulose detected in LL-9 media. According to the USEPA Office of Drinking Water, nitrocellulose is essentially non-toxic (USEPA, 2000d).

6.6.4.2 *Foodstuffs Pathway*

As indicated in Section 6.5.2, there is a significant amount of uncertainty associated with the risk evaluation of the ingestion-of-foodstuffs exposure pathways discussed in the RVAAP FWHHRAM (i.e., ingestion of homegrown beef, milk, and vegetables). Frequently, risk estimates based on background concentrations of chemicals exceed risk benchmarks when these pathways are evaluated. However, conservatively, risk estimates were developed for the hypothetical future resident farmer assuming the receptor is routinely consuming homegrown beef and vegetables and consuming milk from dairy cattle raised on site. COPC concentrations in these media were modeled based on the COPC concentrations detected in surface soil at LL-9 and the equations presented in the RVAAP FWHHRA. The risk estimates are included in Appendix V (Tables 9.6, 9.7, and 9.8) and discussed in the following paragraphs.

Cancer risk estimates for the ingestion of beef, ingestion of milk, and ingestion of vegetables exposure pathways were 2×10^{-4} , 1×10^{-3} , and 3×10^{-3} , respectively. Chemical-specific risk estimates for carcinogenic COPCs selected for surface soil exceed 1×10^{-5} in all cases presented in Appendix V, Table 9.7.

HIs developed for the adult resident farmer and child resident farmer exposed via the ingestion of beef, milk, and vegetables pathways are presented in Appendix V (Tables 9.5 and 9.6.) HIs calculated for the child resident farmer and adult resident farmer exceed 1 for aluminum, arsenic, chromium, and manganese in vegetables.

6.6.4.3 *Summation of Risk*

There are uncertainties concerning the summation of hazards and carcinogenic risk estimates across chemicals and pathways. These include the possibility of synergistic or antagonistic reactions and the varying levels of accuracy and precision of the RfDs or CSFs. In addition, the target organs for carcinogens may be different and therefore, summation of the risk may not be appropriate in all cases.

6.7 Remedial Goal Options

RGOs are developed for each direct-contact exposure COC (i.e., not for COCs identified for the foodstuff exposure pathways). The RGOs are risk-based concentrations that may be used in future risk-based decision making. RGOs are determined using the methods, equations, and



parameters for determining COPC intake shown in Appendix V (Tables 4.1 through 4.32), as well as the CSFs and RfDs shown in Appendix V (Tables 5.1, 5.2, 6.1, and 6.2). The cancer risk (or hazard) equation was rearranged to determine (i.e., to solve for) the concentration that would result in a specified risk or hazard. As discussed in Section 6.5.1, the cancer risk estimate and non-cancer hazard indices are calculated as:

$$\begin{aligned}\text{Cancer Risk Estimate} &= \text{Intake} \times \text{CSF} \\ \text{Hazard Index} &= \text{Intake} / \text{RfD}\end{aligned}$$

Therefore, for a specified risk or hazard, the allowable intake may be determined as follows:

$$\begin{aligned}\text{Intake} &= \text{Cancer Risk Estimate} / \text{CSF} \\ \text{Intake} &= \text{Hazard Index} \times \text{RfD}\end{aligned}$$

The COPC intake equations shown in Appendix V, Tables 4.1 through 4.32, are then rearranged to determine the allowable concentration, or RGO, for a specified risk level (e.g., the RGO associated with the 1E-06 risk level).

RGOs are determined for each medium and land use/receptor scenario. For example, the RGO for arsenic in surface soil at the cancer risk level of 1E-04 for the National Guard trainee receptor is the concentration of arsenic that produces a risk of 1E-04 when using the exposure parameters specified for the National Guard trainee receptor, as shown in Appendix V, Table 4.1.

Direct-contact soil RGOs were not evaluated with regard to residual soil contaminants leaching to groundwater. This pathway is considered not to be a significant pathway, as discussed in Section 5.3.2.

For example, the ingestion of surface soil, the RGO is determined as:

$$C_s = \frac{TR \times BW \times AT}{IR \times FI \times ET \times EF \times ED \times (CSF \text{ or } 1/RfD)}$$

Where:

C_s = concentration in soil (RGO)
 TR = target risk (or hazard)
 BW = body weight
 AT = averaging time
 IR = ingestion rate of soil
 FI = fraction ingested
 ET = exposure time
 EF = exposure frequency
 ED = exposure duration
 CSF = cancer slope factor
 RfD = reference dose



Calculated RGOs that are not physically possible (i.e., the calculated value exceeds 1E+06 mg/kg) are adjusted accordingly. A concentration of 1E+06 indicates a 100% pure substance (i.e., 1,000,000 parts per million). RGOs are calculated for each exposure route (e.g., ingestion), as well as the total risk or hazard for each chemical identified as a COC in an environmental medium. Carcinogenic RGOs are calculated for risk levels of 1E-04, 1E-05, and 1E-06. Non-carcinogenic RGOs are calculated for hazard levels of 0.1 and 1.0.

The potential COCs identified for groundwater, subsurface soil, sediment, and surface soil are shown in Table 6-14. RGOs are calculated for all receptor/medium combinations that have been evaluated in this BHHRA. For example, although manganese is not a surface soil COC for the resident farmer receptor, surface soil RGOs for manganese are calculated for the resident farmer.

RGOs for the direct-contact exposure surface soil COC (arsenic) were calculated for all receptors and included in Table 6-15. RGOs for the subsurface soil COC (arsenic) were calculated for the National Guard resident trainer, and the resident farmer receptors and included in Table 6-16. RGOs for the direct-contact exposure for the National Guard trainee exposed to deep surface soil (0-4 ft bgs) are included in Table 6-17. RGOs for the groundwater COC were calculated for the National Guard trainee, the National Guard resident trainer, and the resident farmer receptors and included in Table 6-18. RGOs for the surface water COC were calculated for the National Guard trainee, the National Guard resident trainer, the hunter/trapper, and the resident farmer receptors and included in Table 6-19.

It should be noted that the COCs discussed in this section should only be considered potential COCs at this time. The final list of COCs for LL-9 will be determined in consultation with the risk management team for RVAAP.

6.8 Summary and Conclusions

This section summarizes the BHHRA and discusses conclusions drawn from the assessment.

6.8.1 Summary

A LL-9 BHHRA was performed using analytical data for groundwater, soil, sediment, and surface water collected from October through December 2003. The LL-9 BHHRA methods are based on the protocol established in the RVAAP FWHHRAM (USACE, 2004) and the *White Paper – Human Health Risk Assessment Approach for Load Lines 6, 9, and 11 Remedial Investigations, Ravenna Army Ammunition Plant, Ravenna, Ohio* (TtNUS, 2004).

Five human receptors were evaluated:

- Security Guard/Maintenance Worker;
- National Guard Trainee;



Table 6-15
Surface Soil (0-1 ft bgs) RGOs

COC	Ingestion					Dermal					Inhalation					Total Across All Pathways				
	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4
Security Guard/Maintenance Worker																				
Arsenic	4.91E+02	4.91E+03	3.05E+01	3.05E+02	3.05E+03	4.42E+01	4.42E+02	2.75E+00	2.75E+01	2.75E+02			2.12E+04	2.12E+05	1.00E+06	4.06E+01	4.06E+02	2.52E+00	2.52E+01	2.52E+02
Hunter/Trapper																				
Arsenic	1.02E+04	1.02E+05	5.30E+02	5.30E+03	5.30E+04	7.47E+03	7.47E+04	3.87E+02	3.87E+03	3.87E+04			2.07E+05	1.00E+06	1.00E+06	4.32E+03	4.32E+04	2.24E+02	2.24E+03	2.24E+04
National Guard Trainer																				
Arsenic	3.04E+01	3.04E+02	1.89E+00	1.89E+01	1.89E+02	1.03E+02	1.03E+03	6.42E+00	6.42E+01	6.42E+02			5.73E+02	5.73E+03	5.73E+04	2.35E+01	2.35E+02	1.46E+00	1.46E+01	1.46E+02
National Guard Fire Dust																				
Arsenic	2.04E+03	2.04E+04	1.27E+02	1.27E+03	1.27E+04	1.72E+03	1.72E+04	1.07E+02	1.07E+03	1.07E+04			7.09E+01	7.09E+02	7.09E+03	9.34E+02	9.34E+03	3.20E+01	3.20E+02	3.20E+03
Resident Child Farmer																				
Arsenic	2.33E+00	2.33E+01				3.56E+01	3.56E+02									2.18E+00	2.18E+01			
Resident Adult Farmer																				
Arsenic	2.17E+01	2.17E+02				3.20E+01	3.20E+02									1.29E+01	1.29E+02			
Resident Farmer																				
Arsenic			3.93E-01	3.93E+00	3.93E+01			1.41E+00	1.41E+01	1.41E+02			3.54E+02	3.54E+03	3.54E+04			3.07E-01	3.07E+00	3.07E+01

RGO = Remedial Goal Option
COC = Chemical of Concern
HQ = Hazard Quotient
HI = Hazard Index
All units in mg/kg



Table 6-16
Subsurface Soil RGOs

COC	Ingestion					Dermal					Inhalation					Total Across All Pathways				
	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4
National Guard Trainer																				
Arsenic	3.04E+01	3.04E+02	1.89E+00	1.89E+01	1.89E+02	1.03E+02	1.03E+03	6.42E+00	6.42E+01	6.42E+02			5.88E+02	5.88E+03	5.88E+04	2.35E+01	2.35E+02	1.46E+00	1.46E+01	1.46E+02
Resident Child Farmer																				
Arsenic	2.33E+00	2.33E+01				3.56E+01	3.56E+02									2.18E+00	2.18E+01			
Resident Adult Farmer																				
Arsenic	2.17E+01	2.17E+02				3.20E+01	3.20E+02									1.29E+01	1.29E+02			
Resident Farmer																				
Arsenic			3.93E-01	3.93E+00	3.93E+01			1.41E+00	1.41E+01	1.41E+02			3.54E+02	3.54E+03	3.54E+04			3.07E-01	3.07E+00	3.07E+01

RGO = Remedial Goal Option
COC = Chemical of Concern
HQ = Hazard Quotient
HI = Hazard Index
All units in mg/kg



Table 6-17
National Guard Trainee Deep Surface Soil (0-4 ft bgs) RGOs

COC	Ingestion					Dermal					Inhalation					Total Across All Pathways				
	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4
National Guard Trainee																				
Chromium	1.95E+03	1.95E+04				4.96E+03	4.96E+04				6.98E+01	6.98E+02				6.65E+01	6.65E+02			
Manganese	9.10E+04	9.10E+05				3.71E+05	1.00E+06				3.49E+01	3.49E+02				3.49E+01	3.49E+02			

RGO = Remedial Goal Option
COC = Chemical of Concern
HQ = Hazard Quotient
HI = Hazard Index
All units in mg/kg



Table 6-18
Groundwater RGOs

COC	Ingestion					Dermal					Inhalation					Total Across All Pathways				
	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4
National Guard Trainee																				
Manganese	1.53E+03	1.53E+04				2.52E+04	2.52E+05									1.44E+03	1.44E+04			
National Guard Trainer																				
Manganese	2.38E+02	2.38E+03				4.08E+03	4.08E+04									2.25E+02	2.25E+03			
Resident Child Farmer																				
Manganese	4.86E+01	4.86E+02				1.36E+03	1.36E+04									4.69E+01	4.69E+02			
Resident Adult Farmer																				
Manganese	1.70E+02	1.70E+03				2.91E+03	2.91E+04									1.61E+02	1.61E+03			
Resident Farmer																				
Manganese																				

RGO = Remedial Goal Option
COC = Chemical of Concern
HQ = Hazard Quotient
HI = Hazard Index
All units in ug/L



Table 6-19
Surface Water RGOs

COC	Ingestion					Dermal					Inhalation					Total Across All Pathways				
	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4	HQ = 0.1	HQ = 1.0	Risk = 10-6	Risk = 10-5	Risk = 10-4
National Guard Trainee																				
Arsenic	1.97E+02	1.97E+03	1.22E+01	1.22E+02	1.22E+03	3.10E+03	3.10E+04	1.90E+02	1.90E+03	1.90E+04						1.85E+02	1.85E+03	1.15E+01	1.15E+02	1.15E+03
National Guard Trainer																				
Arsenic	3.07E+01	3.07E+02	1.91E+00	1.91E+01	1.91E+02	4.70E+02	4.70E+03	3.00E+01	3.00E+02	3.00E+03						2.88E+01	2.88E+02	1.79E+00	1.79E+01	1.79E+02
Hunter/Trapper/Fisher																				
Arsenic	7.67E+03	7.67E+04	3.97E+02	3.97E+03	3.97E+04	3.80E+04	3.80E+05	2.00E+03	2.00E+04	2.00E+05						6.38E+03	6.38E+04	3.32E+02	3.32E+03	3.32E+04
Resident Child Farmer																				
Arsenic	4.70E+00	4.70E+01				1.07E+02	1.07E+03									4.50E+00	4.50E+01			
Resident Adult Farmer																				
Arsenic	2.20E+01	2.20E+02				1.92E+02	1.92E+03									1.97E+01	1.97E+02			
Resident Farmer																				
Arsenic			5.88E-01	5.88E+00	5.88E+01			7.35E+00	7.35E+01	7.35E+02								5.45E-01	5.45E+00	5.45E+01

RGO = Remedial Goal Option
COC = Chemical of Concern
HQ = Hazard Quotient
HI = Hazard Index
All units in ug/L



- National Guard resident trainer;
- Hunter/Trapper; and
- Hypothetical Future Resident Farmer (adult and child).

The National Guard trainee is an important receptor because it is anticipated that the LL-9 area will be used by OHARNG for training activities described in the FWHHRAM as *"Mounted*

Training by Ohio National Guard Soldiers – No Digging Allowed." The planned training activities would involve potential exposure to soils (no deeper than 4 ft bgs), as well as to the other environmental media, at LL-9. The hunter/trapper receptor is an important receptor because a long-term goal for RVAAP is to be able to hunt, fish, and trap anywhere suitable habitat exists for recreational fishing and hunting. (Please note that fishing does not currently occur at LL-9.) The National Guard resident and the hypothetical future resident farmer are included in the BHHRA for purposes of completeness and because risk estimates for these receptors may be useful to the RVAAP risk managers. For example, the need for deed restrictions at LL-9 may be eliminated if no unacceptable risk was determined by the BHHRA for these receptors. The security guard/maintenance receptor is included because of the routine activities (e.g., security patrols) currently occurring by personnel who visit the site for short periods of time.

The following chemicals were selected as COPCs for quantitative human health risk assessment:

Soils	Groundwater	Surface Waters	Sediment
Aluminum	Antimony	Aluminum	Aluminum
Arsenic	Manganese	Arsenic	Mercury
Benzo(a)pyrene		Chromium	Vanadium
Chromium		Lead	
Copper		Manganese	
Dibenzo(a,h)anthracene		Mercury	
Lead		Vanadium	
Manganese			
Mercury			

These COPCs were selected based on chemical concentrations detected in the environmental media at LL-9. The COPC selection protocol included a conservative toxicity screen (based on the Region 9 residential PRGs) and a background screen. The following should be considered when evaluating this list of COPCs:

- Only 1 manganese detection of 5 samples in groundwater exceeded the background benchmark for RVAAP.



- Only 2 detections of manganese of 85 samples in deep surface soil exceeded their respective background benchmarks for RVAAP. The EPC in deep surface soil (634 mg/kg) did not exceed RVAAP background (1,450 mg/kg)
- The benzo(a)pyrene concentrations detected in soils are similar to anthropogenic background concentrations and concentrations reported for background soil samples collected at RVAAP.
- Maximum arsenic concentrations detected in environmental media at LL-9 exceeded background benchmarks established for RIs at RVAAP (10 out of 46 surface soil samples and 5 out of 49 subsurface soil samples). The EPC of arsenic in surface soil (13.8 mg/kg), and in deep surface soil (13 mg/kg) are less than their respective background concentrations (19.8 mg/kg, and 15.4 mg/kg). The historical, widespread application of arsenic-based herbicides across many areas of the United States may have contributed to the concentrations detected in soils. Also, the arsenic concentrations detected at LL-9 are within background concentrations reported by the USEPA (1987a).

A summary of the risk characterization for LL-9 is presented in Table 6-11 and in the following items:

- Non-cancer risk estimates (HIs) developed for the security guard/maintenance worker and the hunter/trapper are less than 1, indicating that adverse non-carcinogenic effects are not anticipated under the exposure conditions considered in the risk assessment. The cancer risk estimates for the security guard/maintenance worker do not exceed $1\text{E-}05$ and for the hunter/trapper do not exceed $1\text{E-}06$. No potential COCs are identified for these receptors.
- The HIs calculated for the National Guard trainee receptor exposed to COPCs in groundwater, sediments, and surface water do not exceed 1. The HI for the National Guard trainee exposed to COPCs in deep surface soils (0-4 ft bgs) is 2. Manganese (inhalation of air particulates being the route of exposure) was the significant contributor to the non-carcinogenic risk (i.e., risk driver) for the National Guard trainee. The estimates for inhalation of manganese in deep surface soil are strongly affected by the PEF specified for the National Guard trainee receptor in the RVAAP FWHHRAM. This PEF is further discussed in Section 6.6. Manganese at a concentration equal to RVAAP background would result in an HI of 4, and at a concentration equal to its Region 9 residential PRG would result in an HI of 5. The EPC for manganese in deep surface soil (633 mg/kg) did not exceed RVAAP background (1,450 mg/kg).
- The total cancer risk estimate for National Guard trainee exposure to all media ($2\text{E-}05$) is within the USEPA target cancer risk range ($1\text{E-}04$ to $1\text{E-}06$) but exceeds the Ohio EPA benchmark of $1\text{E-}05$. Inhalation of arsenic and chromium in deep surface soil account



for 1E-05, or 82%, of the total cancer risk estimate for the National Guard trainee. The estimates for inhalation in deep surface soil are strongly affected by the PEF specified for the National Guard trainee receptor in the RVAAP FWHHRAM. This PEF is further discussed in Section 6.6. The EPCs of arsenic (13 mg/kg) and chromium (17 mg/kg) in deep surface soil do not exceed their respective RVAAP background values of 15.4 mg/kg and 17.4 mg/kg.

- The total HI for the National Guard resident trainer is 2. The HI calculated for receptor exposure to groundwater is 1. The HI for exposure to all other media (not including groundwater) combined is 0.4. Manganese in groundwater is the only chemical identified as a potential COC.
- The total cancer risk estimate for the National Guard resident trainer, (2E-05) is within the EPA target risk range of 1E-04 to 1E-06 but exceeded the Ohio EPA cancer risk benchmark of 1E-05. No risk estimates developed for individual COPCs exceed 1E-05. The cancer risk estimate for surface water is 6E-06. Arsenic is a potential COC for subsurface soil. The EPC of arsenic in subsurface soil (13.8 mg/kg) is less than its background concentration of 19.8 mg/kg.
- The total HI calculated for the adult resident farmer routinely exposed to COPCs in surface soil, surface water, sediment, and groundwater, or routinely exposed to COPCs in subsurface soil, surface water, sediment, and groundwater via the direct-contact pathways, exceed 1 only when it is assumed that the shallow groundwater is used for domestic purposes. Manganese in groundwater is the only risk driver. The HIs calculated for the adult resident farmer routinely exposed via indirect exposure pathways (i.e., consumption of foodstuffs) and also presented in Appendix V, Table 9.6, exceeded 1 for vegetable consumption only. Aluminum, arsenic, chromium, and manganese are potential COCs for the ingestion of vegetables. However, the food chain modeling used to characterize risk is very conservative. HIs calculated for the resident farmer exposed to these metals at site background concentrations would exceed 1.
- The total HI calculated for the child resident farmer routinely exposed to COPCs in surface soil, sediment, surface water, and groundwater, or routinely exposed to COPCs in subsurface soil, sediment, surface water, and groundwater via the direct-contact exposure pathways, exceed 1. The HIs calculated for the COPCs in groundwater and surface water exceed 1. The HIs calculated for sediments and subsurface soils do not exceed 1. Arsenic and manganese are the primary risk drivers; HIs calculated for other COPCs evaluated do not exceed 1. The target-organ-specific HI calculated for the skin and cardiovascular system (the target organs for arsenic) and the central nervous system (the target system for manganese) exceed 1. The HIs calculated for the child resident farmer routinely exposed via indirect exposure pathways (i.e., consumption of foodstuffs) also presented in Appendix V, Table 9.5, do exceed 1 for vegetable and milk consumption. Aluminum, arsenic, chromium, and manganese are potential COCs for



the ingestion of vegetables. However, the food chain modeling used to characterize risk is very conservative. HIs calculated for the resident farmer exposed to these metals at site background concentrations would exceed 1.

- The ILCR estimates for the hypothetical future resident farmer are $5E-05$, $5E-05$, and $2E-05$ for direct-contact exposure pathways for surface soil, subsurface soil, and surface water, respectively. The total cancer risk for the direct-contact exposure pathways did not exceed $1E-04$ when surface soil and subsurface soil risks were considered separately. Arsenic in soils is identified as a potential COC for the direct-contact exposure pathway. The cancer risk estimates calculated for the indirect exposure pathways (i.e., consumption of beef, milk, and vegetables raised on site) exceed $1E-04$. Benzo(a)pyrene, dibenzo(a,h)anthracene, and arsenic are identified as potential COCs for foodstuff consumption. However, the uncertainties highlighted in the previous items are also relevant for the risk estimates presented for the future hypothetical resident farmer.
- Maximum detected concentrations of surface soil and deep surface soil COPCs copper (1240 mg/kg), lead (1330 mg/kg), and mercury (882 mg/kg) were at sample location LL9SS-011-0001-SO. The next-highest detected concentrations were much lower (copper [170 mg/kg], lead [320 mg/kg], and mercury [17 mg/kg]). This sample location may be of concern when making risk decisions, particularly for exposure to mercury.

Table 6-11 presents a summary of all receptors evaluated in this BHHRA and identifies the contaminants that produce risks greater than $1E-06$, $1E-05$, and $1E-04$ or hazards greater than 1.0.

6.8.2 Conclusions

The total cancer risk estimate of $2E-05$ (summarized for each receptor and each media) for the primary receptor of concern (the National Guard trainee) is within the USEPA target cancer risk range ($1E-04$ to $1E-06$), but exceeds the Ohio EPA benchmark of $1E-05$. The cancer risk estimates developed for inhalation of arsenic and chromium in deep surface soils exceed $1E-06$, but not $1E-05$. The total non-cancer risk (HI) estimate (summarized for each receptor and each media) equals 3. Manganese (inhalation of air particulates being the route of exposure) was the significant contributor to the non-carcinogenic risk. However, as discussed in the preceding narrative, significant uncertainties were identified for the risk estimates for arsenic, manganese, and chromium in soils and for the inhalation pathway; these should be considered by the risk management team for RVAAP when making further remedial decisions for LL-9. Manganese at background would result in an HI of 4, and at its Region 9 residential PRG would result in an HI of 5.



1 HIs calculated for the security guard/maintenance worker and the hunter/trapper are less than 1,
2 and total cancer risk estimates for the same receptors do not exceed the Ohio EPA cancer risk
3 benchmark of 1E-05.

4 The total cancer risk estimate (summarized for each receptor and each media) for the National
5 Guard resident trainer exceeds 1E-05. The total HI (summarized for each receptor and each
6 media) for the National Guard resident trainer exceeds 1.0 for the future domestic use of the
7 shallow groundwater resource. HIs for all other media for the National Guard resident trainer do
8 not exceed 1. As noted above, significant uncertainties were identified for the risk estimates for
9 manganese in groundwater and arsenic in soils and should be considered by the risk
10 management team for RVAAP when making further remedial decisions for LL-9.

11 The total cancer risk estimate (summarized for each receptor and each media) for the
12 hypothetical future resident farmer exposed by the direct contact exposure pathways did not
13 exceed 1E-04 when surface soil and subsurface soil risks were considered separately. Cancer
14 risk estimates for soils evaluated for the direct-contact exposure pathways exceed 1E-05 but do
15 not exceed 1E-04. Total non-cancer risk estimates (HI) calculated on a target-organ-specific
16 basis exceed 1 only when future domestic use of the shallow groundwater resource is evaluated
17 for the adult farmer. For the child farmer, total HI calculated on a target-organ-specific basis
18 exceed 1 when future domestic use of the shallow groundwater resource is evaluated and
19 exceed 1 for arsenic's target organs when direct exposure to media other than groundwater are
20 evaluated and only if the HIs for surface soil and subsurface soil are combined. However,
21 cancer and non-cancer risk estimates developed for the indirect exposure pathways (i.e.,
22 consumption of home grown food stuffs) exceed both cancer and non-cancer risk benchmarks
23 (i.e., 1E-04 and HI = 1, respectively) for consumption of foodstuffs. Significant uncertainties
24 were identified for the risk estimates for manganese in groundwater and for the evaluation of the
25 indirect exposure pathways, and these should be considered by the risk management team for
26 RVAAP when making further remedial decisions for LL-9.



7.0 SCREENING-LEVEL ECOLOGICAL RISK ASSESSMENT

7.1 Introduction

The goal of this SERA is to determine whether potential ecological impacts are present as a result of exposure to chemicals released to the environment through past site operations at RVAAP LL9. This SERA provides information to scientists and managers that will enable them to conclude either that ecological risks at the site are most likely negligible or that further information is necessary to evaluate potential ecological risks at the site. The SERA methodology follows the guidance presented in the *RVAAP Facility – Wide Ecological Risk Work Plan* (USACE, 2003) and *Guidance for Conducting Ecological Risk Assessments* (Ohio EPA, 2003).

The ERA process consists of the following eight steps that are required by the RVAAP, Ohio EPA, and USEPA for any ecological risk assessment:

- Step 1 – Screening-Level Problem Formulation and Ecological Effects Evaluation
- Step 2 – Screening-Level Exposure Estimate and Risk calculation
- Step 3 – Baseline Risk Assessment Problem Formulation
- Step 4 – Study Design and Data Quality Objective Process
- Step 5 – Field Verification of Sampling design
- Step 6 – Site Investigation and Analysis Phase
- Step 7 – Risk Characterization
- Step 8 – Risk Management

A SERA typically consists of Steps 1 and 2 and a Baseline Ecological Risk Assessment (BERA) consists of Steps 3 through 7. Step 8, Risk Management, is a distinctly different process from risk assessment. In risk management, the results of the risk assessment are integrated with other considerations to make and justify risk management decisions (USEPA, 1997). This SERA for RVAAP LL9 consists of Steps 1 and 2, along with the first part of Step 3, which is termed Step 3a. Step 3a considers factors other than comparisons of chemical concentrations to screening levels to further refine the list of COPCs (see Section 7.6 for more details). The remaining steps (the rest of Step 3 through Step 7) are conducted only if additional evaluations or investigations are necessary; these steps were not conducted as part of this report. A decision to proceed to a BERA is usually only made after the results of the SERA are evaluated.



7.2 Preliminary Problem Formulation

Problem formulation is the first step of the SERA and includes identification of potential receptor groups, potential contaminants, and the mechanisms for contaminant fate and transport and toxicity. Determination of the complete exposure pathways that exist at a site is accomplished at this point to facilitate proper receptor selection. Site habitats, exposure pathways, and potential ecological receptors are identified as part of the problem formulation. This information, which is gathered from the field investigation and site visits, is used to focus the SERA on the primary chemicals, receptors, and exposure pathways at the site.

7.2.1 Habitat Types and Ecological Receptors

The LL9 site consists of open field habitat with grasses, weeds, and small shrubs (approximately 70% of the site). There are some isolated pockets of mature trees as well (approximately 30% of the site). A visual approximation of the habitat was obtained from the aerial photograph provided in Appendix C. The area surrounding LL9 consists of early successional tree species, as well as trees 15–20 ft tall and woody shrubs. Shallow man-made ditches drain the site, but contain water only during rain events of approximately 1 in or greater based on site observations. However, the amount of rain required to cause the ditches to contain water depends upon various factors such as the dryness of the soil, soil type and frequency of rain events. As can be seen in the photographs in Appendix C, surface water sample locations LL9SW-001-SW, LL9SW-003-SW, LL9SW-004-SW, and LL9SW-005-SW appear to be intermittent drainage ditches with no aquatic habitat, while aquatic receptors may be present at locations LL9SW-002-SW and LL9SW-012-SW. Note that no aquatic plants were present at any of the locations. This could indicate that the ditches do not contain water for significant periods of time. Although all of the surface water and associated sediment samples will be evaluated as aquatic habitat because of the potential for contaminants to migrate downstream where aquatic receptors may be more prevalent, the uncertainties in this evaluation at locations where water is only present during rain events are discussed in the uncertainty analysis section of this SERA (Sections 7.8.2 and 7.8.4). Section 2.8 presents the general ecology for RVAAP. Based on the ecology, and the environmental setting at LL9, it is likely that LL9 is occupied by a variety of mammal and avian receptors. Field observations from a site visit on February 13, 2001 identified crows (*Corvus* sp.), hawks (*Buteo* sp.), bluebirds (*Sialia sialis*), blue jays (*Cyanocitta cristata*), white-tailed deer (*Odocoileus virginianus*), and owls (Tytonidae and/or Strigidae families) (from pellets).

7.2.2 Major Chemical Sources and Migration Pathways

LL9 was used as a detonator processing facility. Fulminate, azide, and tetryl mixing and processing activities were conducted in various buildings at LL9 to support detonator manufacturing processes.



Based on historical site data and sampling, the following parameters are among the site-related chemical contaminants that were detected at LL9. Note that not all the chemicals mentioned in the following bullets were detected or analyzed for in each sample. Figure 7-1 presents a conceptual site model (CSM) that shows the major migration pathways that could occur at the site.

- Explosives (surface soil);
- Metals (surface soil, sediment, and surface water); and
- Semi-volatile organic compounds (SVOCs) (surface soil and sediment).

The results of the surface soil, surface water, and sediment sampling are included in earlier sections of this report. Physical and chemical characteristics of contaminants may affect their mobility, transport, and bioavailability in the environment. These characteristics include the organic carbon partition coefficient, octanol water partition coefficient, and vapor pressure. Table 5-1 presents the physical and chemical characteristics for the chemicals detected at LL9. The fate and transport of these chemicals is discussed in detail in Section 5.0. Although these physical and chemical characteristics were not used directly in this SERA, many of the characteristics were used to develop some of the values that were used in this SERA. For example, K_{ow} values were used to determine some of the biotransfer factors in Appendix W. Also, some of the sediment screening values are based on equilibrium partitioning which utilizes the K_{oc} of the chemical.

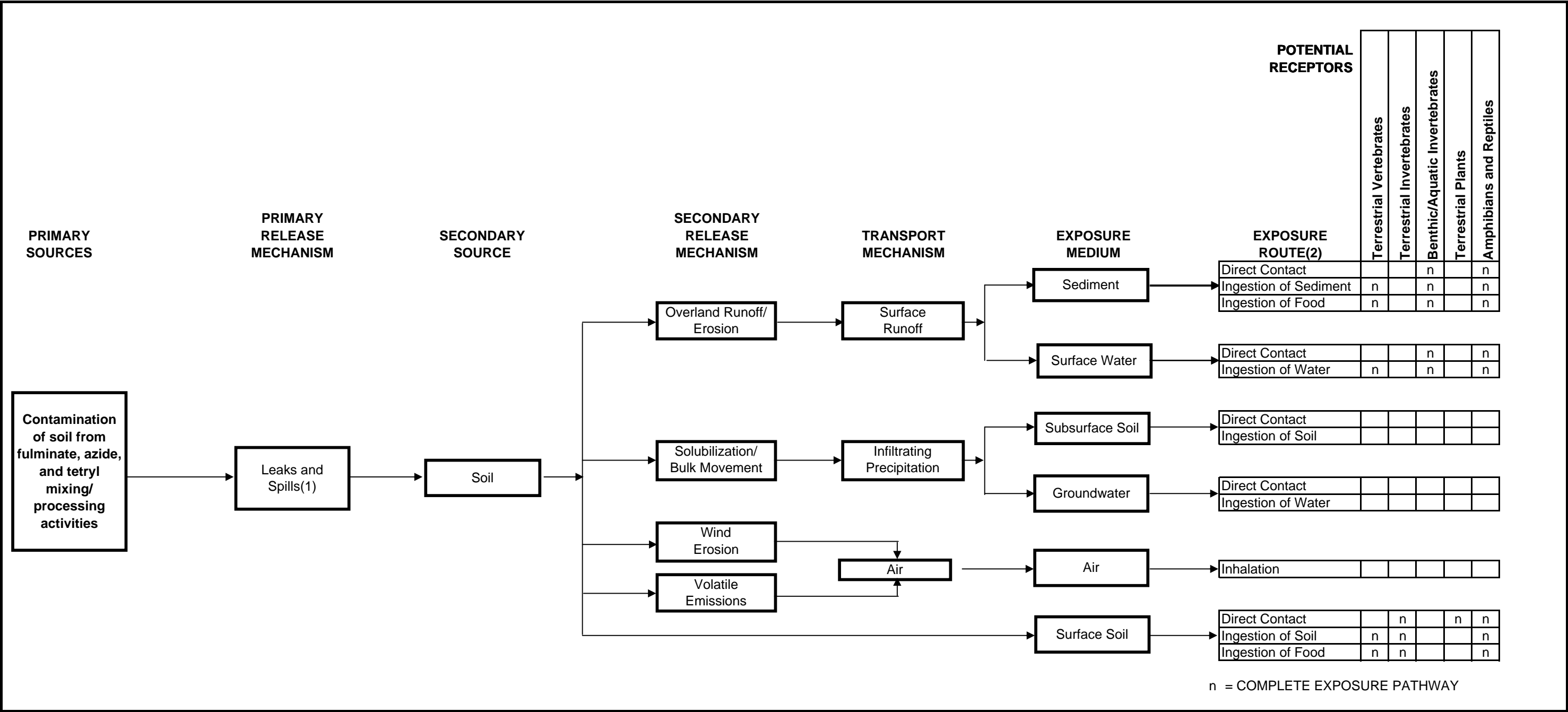
Various bioaccumulation factors (BAFs) and/or bioconcentration factors (BCFs) were used to estimate contaminant loading in plants and soil invertebrates from chemical concentrations in LL 9 surface soil. For soil to plant uptake, two separate BAFs are used to predict the biological transfer of contaminants from soil to vegetative (foliage) or reproductive (berries) and storage (roots) plant parts (Baes et al, 1984; Travis and Arms, 1988). Additional transfer factors include soil to animal (i.e., earthworm) and animal to animal (i.e., earthworm to shrew). Factors were from the RVAAP *Facility-Wide Ecological Risk Guidance* (USACE, 2003). Contaminants that do not have BAFs are assigned a default value of 1. Appendix W presents two tables with the transfer factors for soil to plant, soil to animal, and animal to animal.

7.2.3 Exposure Routes

The potential pathways by which ecological receptors may be exposed to COPCs were identified, along with the species that could be adversely affected by these chemicals. Several potential exposure pathways may exist. For example, terrestrial animals may be exposed to soil contaminants through ingestion of contaminated food items. Animals can also incidentally ingest soil while grooming fur, preening feathers, digging, grazing close to the soil, or feeding on items to which soil has adhered (such as roots and tubers). Terrestrial vegetation may be exposed to contaminants via direct aerial deposition and root translocation. Terrestrial animal receptors may also come into contact with contaminants in surface water by drinking the water,



FIGURE 7-1
Ecological Conceptual Site Model



1. Leaks and spills from explosives processing activities
2. Blank space indicates incomplete exposure pathway or relatively insignificant or not applicable potential exposure.



although this exposure route typically represents a negligible portion of total exposure for most terrestrial receptors because of the relatively low contaminant concentrations in surface water as compared to other media. Aquatic and semi-aquatic organisms may be exposed to contaminants via direct contact with surface water and sediments, incidental ingestion of surface water and sediments, and consumption of contaminated food items. Aquatic and semi-aquatic organisms may also be exposed to constituents from contaminated groundwater that flows into surface water.

7.2.3.1 Surface Soil

Several groups of terrestrial ecological receptors can be exposed to contaminants in the soil (0–4 ft), although most of the exposure is expected to occur in the top 1 ft. Invertebrates, such as earthworms, are exposed to the contaminants as they move through the soil and ingest soil particles while searching for food. Plants are exposed to the contaminants via direct contact as contaminants are absorbed through the roots, which may then translocate to different parts of the plants (i.e., leaves, seeds).

Small mammals may be exposed to contaminants in the soil via several exposure routes. They may be exposed by direct contact as they search for food or burrow into the soil. However, exposure of terrestrial wildlife to contaminants in the soil via dermal contact is unlikely to represent a major exposure pathway because fur, feathers, and chitinous exoskeletons are expected to minimize transfer of contaminants across dermal tissue. The exposure of animals to constituents in soil by dermal contact is likely to be a small fraction of the direct exposure to constituents in soil by incidental ingestion and the indirect exposure by ingestion of contaminated biota. It is true that for amphibians and reptiles, the dermal pathway may be more important. However, there currently are no standard methods for evaluating risks to amphibians and reptiles from dermal exposure to chemicals in soil. Therefore, the dermal pathway is not evaluated in this SERA. Small mammals also may be exposed to contaminants in the soil via incidental ingestion of soil and ingestion of plants and/or invertebrates that have accumulated contaminants from the soil. The soil and food ingestion pathways are evaluated in this SERA.

Larger predatory species, such as the red fox (*Vulpes vulpes*), barn owl (*Tyto alba*) and red-tailed hawk (*Buteo jamaicensis*), can be exposed (indirectly) to site contaminants in the soil by ingesting small mammals that have accumulated contaminants from the soil. The small mammal ingestion pathway is evaluated in this SERA.

7.2.3.2 Groundwater

Ecological receptors are not directly exposed to contaminants in the groundwater, so this exposure pathway is not complete. According to USACE (2003d), because groundwater is more than 2 ft deep, it does not need to be evaluated as surface water.



7.2.3.3 *Surface Water/Sediment*

Contaminants in the soil may enter the drainage ditches via overland flow. The water bodies immediately adjacent to the LL-9 buildings are small drainage ditches that have poor ecological habitat and probably only support a small invertebrate population. Aquatic receptors could be exposed to contaminants in the water or sediment by direct contact and incidental ingestion of water. Because habitat is not present that would support a fish population, piscivorous wildlife were not evaluated in the SERA.

7.2.3.4 *Air*

Although inhalation of particulates may be a complete pathway, it is expected to be insignificant compared to other pathways such as ingestion of food items that have accumulated contaminants from soil. Also, inhalation pathways are typically not evaluated in SERAs because of the uncertainty in exposures and effects concentrations. The exposure of animals to constituents in soil by inhalation is likely to be a small fraction of the direct exposure to constituents in soil by incidental ingestion and the indirect exposure by ingestion of contaminated biota. It is true that for amphibians and reptiles, the inhalation pathway may be more important. However, there currently are no standard methods for evaluating risks to amphibians and reptiles from inhalation exposure to chemicals in soil. Therefore, this pathway was not evaluated.

7.2.4 **Preliminary Assessment and Measurement Endpoints**

One of the major tasks in the screening-level problem formulation is the selection of assessment and measurement endpoints.

7.2.4.1 *Assessment Endpoints*

Assessment endpoints are explicit expressions of the environmental value that is to be protected (USEPA, 1997). The selection of these endpoints is based on the habitats present, the migration pathways of probable contaminants, and the routes that contaminants may take to reach receptors.

The habitat at and adjacent to the site consists of forested areas, open fields with grasses, and drainage ways that contain water only during rain events. For this SERA, the assessment endpoints are the protection of the following groups of receptors from adverse effects of contaminants on their growth, survival, and reproduction (note that although toxicity data may not be available for all of the potential effects for each receptor group, it is assumed that the toxicity data from one endpoint will be sufficiently protective of the other endpoints):

- Soil invertebrates;



- Terrestrial vegetation;
- Herbivorous mammals;
- Invertivorous birds and mammals;
- Carnivorous birds and mammals; and
- Aquatic organisms and benthic invertebrates.

The following paragraphs discuss why the above receptors are selected for this SERA:

Soil Invertebrates – Soil invertebrates are expected to be present in the soil at the site. They aid in the formation of soil and redistribution and decomposition of organic matter in the soil and serve as a food source for higher trophic level organisms. They also can accumulate some contaminants that can then be transferred to the higher trophic level organisms that consume invertebrates.

Terrestrial Vegetation – Terrestrial vegetation at the site consists of grasses, shrubs, and trees. These serve as a food source, provide shade and cover for many organisms, and help prevent soil erosion, among other important functions. They also can accumulate some contaminants that can then be transferred to higher trophic level organisms that consume plants.

Herbivorous Mammals – Herbivorous mammals (animals that consume only plant tissue) may be present at the site because of the vegetative habitats. Their role in the community is essential because, without them, higher trophic levels could not exist. They may be exposed to and accumulate contaminants that are present in the plants they consume.

Invertivorous Birds and Mammals – Birds and mammals that eat soil invertebrates (invertivores) are present throughout RVAAP in different terrestrial habitats (i.e., forested, open field). These are considered first-level carnivores, and they serve as a food source for higher trophic level carnivores. The invertivores may be exposed to and accumulate contaminants that are present in the food items they consume.

Carnivorous Birds and Mammals – Carnivorous birds and mammals consist of birds and mammals that consume invertebrates, fish, and other mammals and birds. Carnivorous birds and mammals that feed on other birds and mammals are at the top of the food chain. The top carnivores typically are less densely distributed than the herbivores and first-level carnivores because they require a larger area to hunt for their food. Carnivores may accumulate contaminants that are present in the food items they consume.

Aquatic Organisms and Benthic Macroinvertebrates – Aquatic organisms and benthic macroinvertebrates are similar to the soil invertebrates in that they serve as a food source for higher trophic level organisms (i.e., fish, amphibians, birds, mammals). They may also accumulate some contaminants that can then be transferred to the higher trophic level



organisms that consume them. Based on the poor aquatic habitat at the site, and because the drainage ditches are wet only during and after rain events, it is not likely that significant populations of aquatic organisms or benthic invertebrates exist at the site.

All of the potential receptors are not evaluated in the SERA. As indicated in USEPA guidance (1997), "It is not practical or possible to directly evaluate risks to all of the individual components of the ecosystem at a site. Instead, the receptors included in the risk assessment focus on particular components of the ecosystem that could be adversely affected by contaminants from the site." Therefore, the SERA focuses on the receptors that have the greatest exposure and greatest likelihood of being impacted by site contaminants, which should then account for receptors with lower exposures. For example, to be conservative, large carnivorous mammals (i.e., red fox) and birds (i.e., barn owl and red-tailed hawk) are evaluated in this SERA, although the area of LL9 is small (69 acres) and less than the typical home and feeding ranges of carnivorous animals. The greatest exposure to site contaminants is expected to occur to the small mammals and birds that ingest earthworms or plants. Also, omnivores were not selected as assessment endpoints because exposure to contaminants in plants is greatest for herbivores and exposure to contaminants in animals is greatest for insectivores. Therefore, omnivores were protected by protecting herbivores and insectivores."

The assessment endpoints presented in this section are evaluated using various measurement endpoints as described in the next section. For example, the assessment endpoint "Protection of soil invertebrates from adverse effects of contaminants on their growth, survival, and reproduction" is evaluated using the measurement endpoint in the first bullet in Section 7.2.4.2. Note that at this screening step in the ERA process, the assessment and measurement endpoints are relatively general in nature. However, should the site progress further into a BERA, the assessment and measurement endpoints would become more specific.

7.2.4.2 Measurement Endpoints

Measurement endpoints are estimates of biological impacts (e.g., mortality and reduction in growth or reproduction) that are used to evaluate the assessment endpoints. The following measures of effects are used to evaluate the assessment endpoints in this SERA, where applicable.

- Soil screening values – Mortality, growth, and reproduction of plants and soil invertebrates are evaluated by comparing the measured concentrations of chemicals in the surface soil to screening values designed to be protective of ecological receptors.
- No-observed-adverse effects levels (NOAELs) for surrogate wildlife species – Mortality, reproductive, and/or developmental effects of birds and mammals are evaluated by comparing the estimated ingested dose from contaminants in the surface water, sediment, surface soil, plants, and/or invertebrates to these levels.



- Sediment screening values – Mortality and other adverse effects (e.g., growth, feeding rates, and behavioral changes) of benthic macroinvertebrates are evaluated by comparing the measured concentrations of chemicals in the sediment to screening values designed to be protective of ecological receptors.
- Surface water screening values – Mortality and other adverse effects (e.g., growth, feeding rates, and behavioral changes) of aquatic organisms are evaluated by comparing the measured concentrations of chemicals in the surface water to screening values designed to be protective of ecological receptors.

7.2.5 Selection of Receptor Species for Food Chain Model

To address potential effects to wildlife via food chain modeling (see Section 7.5.5), several species of vertebrates were selected to be evaluated in this SERA. The dietary exposure (dose of contaminants from ingesting and drinking) of these receptors was estimated and then compared to effect levels (threshold values that, if exceeded, may elicit an effect).

For most receptor species, ingestion is the primary route of exposure. Representative species are selected for their preferred habitat, body size, sensitivity, home range, abundance, commercial or sport value, legal status, and functional role (e.g., predators). For conservativeness, representative species may be small and have small home ranges. The availability of exposure parameters, such as body mass, feeding rate, and drinking rate, may also be a factor in selecting representative species. The following representative species are used for the food chain modeling:

- Herbivorous mammal: meadow vole (*Microtus pennsylvanicus*);
- Invertivorous mammal: short-tail shrew (*Blarina brevicauda*);
- Invertivorous bird: American robin (*Turdus migratorius*);
- Carnivorous bird: barn owl and red-tailed hawk; and
- Carnivorous mammals: red fox.

7.3 Preliminary Ecological Effects Evaluation

The preliminary ecological effects evaluation is an investigation of the relationship between the magnitude of exposure to a chemical and the nature and magnitude of adverse effects resulting from exposure. As the first step in the ecological effects evaluation, COPCs screening-level risk calculations are used to compare the conservative exposure estimate to ecological effect values estimated by ESVs. A hierarchy of ESVs has been specified by Ohio EPA and RVAAP (Ohio EPA, 2003; USACE, 2003). The first available screening value is used in the hierarchy, regardless of whether it may be greater or lesser than other screening values.



Surface water chemical concentrations are compared to chemical criteria using a surface water hierarchy of: (1) outside mixing zone maximum (OMZM) criteria for the protection of aquatic life in the Lake Erie Basin from Ohio administrative Codes (OAC) 3745-1 (Ohio EPA, 2004) and (2) USEPA ambient water quality criteria (AWQC) (USEPA, 2002).

Sediment concentrations are screened using the site-specific background concentrations and Ohio-specific sediment reference values (SRVs) (Appendix H in Ohio EPA, 2003), then using a sediment screening hierarchy of: (1) consensus-based threshold effect concentrations (TECs) (MacDonald et al., 2000) and (2) Region 5 ecological data quality levels (EDQLs) (USEPA Region 5, 1999).

Soil concentrations are screened using the site-specific background concentrations, then using a soil screening value hierarchy in the order given in the guidance, as follows: (1) PRGs (Efroymson et al., 1997a); (2) toxicological benchmarks for soil and litter invertebrates (Efroymson et al., 1997b); (3) toxicological benchmarks for terrestrial plants (Efroymson et al., 1997c); and (4) Region 5 EDQLs (USEPA Region 5, 1999).

Although the EDQLs have since been updated by the ecological screening levels (ESLs) (USEPA, 2003a), they were not used as soil or sediment screening levels, based on e-mail correspondence from Laurie Moore of the Ohio EPA on December 17, 2004.

7.4 Preliminary Risk Calculation and Selection of COPC

This initial selection of COPCs uses conservative assumptions to eliminate chemicals detected during the remedial investigation at concentrations below those levels reported in the literature to cause adverse effects on growth, survival, or reproduction. This process involves the combination of a preliminary exposure estimate with a preliminary risk calculation.

The first step in selecting initial COPCs is to select a conservative EPC for comparison with background and ESVs. The maximum concentration is used as the initial EPC for comparison to the ESV. The ratio of the initial EPC to the ESV is called a hazard quotient (HQ) and is defined as follows:

$$HQ_i = EPC_i / ESV_i$$

Where: HQ_i = hazard quotient for analyte "i" (unitless)

EPC_i = exposure point concentration for analyte "i" (mg/L or mg/kg)

ESV_i = ecological screening value for analyte "i" (mg/L or mg/kg)

When the ratio of the EPC to its respective screening value or benchmark exceeds 1.0, adverse impacts are possible, and the chemical is considered for selection as a COPC. The HQ is not probabilistic but a numerical indicator of the extent to which an EPC exceeds an ESV.



7.4.1 Selection of Soil COPCs

The following summarizes the procedures that were used in the SERA for LL9 to select COPCs. Note that calcium, magnesium, potassium, and sodium are not retained as COPCs in any medium because of their relatively low toxicity to ecological receptors and their high natural variability in concentrations. Contaminants without ESVs are retained as COPCs for further evaluation.

- Inorganic contaminants whose maximum concentrations do not exceed the site-specific background concentrations are not selected as COPCs.
- Inorganic and organic contaminants whose maximum concentrations do not exceed soil screening levels are not retained as COPCs, unless the chemicals are bioaccumulative and/or persistent, bioaccumulative, and toxic (PBT). Inorganic and organic contaminants that are bioaccumulative and/or PBT are selected as COPCs even if their maximum concentration does not exceed a screening level. A chemical was considered to be bioaccumulative if it is included in the list of important bioaccumulative chemicals in USEPA (2000a). A chemical was considered a PBT chemical as defined in the Ohio EPA ERA guidance (Ohio EPA, 2003).
- Inorganic and organic contaminants without screening values are selected as COPCs.

Table 7-1 summarizes the screening of soil COPCs. Four analytes (dibenzofuran, nitroguanidine, RDX, and nitrocellulose) were selected as COPCs because no screening values are available. According to the USEPA Office of Drinking Water, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in Section 6.6. Twelve analytes (aluminum, arsenic, chromium, copper, iron, lead, manganese, mercury, nickel, selenium, vanadium, and zinc) were selected as COPCs because their maximum concentrations exceeded their respective ESVs. Eight of those analytes (arsenic, chromium, copper, lead, mercury, nickel, selenium, and zinc) were selected as COPCs for the food chain model because they are bioaccumulative. Although some of the ESVs were developed for the protection of wildlife, 16 bioaccumulative analytes (acenaphthene, acenaphthylene, anthracene, benzo[a]anthracene, benzo[a]pyrene, benzo[b]fluoranthene, benzo[g,h,i]perylene, benzo(k)fluoranthene, chrysene, dibenzo(a,h)anthracene, fluoranthene, fluorene, indeno[1,2,3-cd]pyrene, phenanthrene, pyrene, and cadmium) were selected as COPCs for the food chain model to be conservative, even though their maximum detected concentrations did not exceed soil screening levels.

7.4.2 Selection of Sediment COPCs

The following summarizes the procedures that were used in the SERA for LL9 to select COPCs:

- Inorganic contaminants whose maximum concentrations do not exceed Ohio-Specific SRVs and site-specific background are not selected as COPCs.



- Inorganic and organic contaminants whose maximum concentrations do not exceed screening levels are not selected as COPCs.

- Inorganic and organic contaminants without screening values are selected as COPCs.

Table 7-2 summarizes the initial selection of sediment COPCs. Beryllium, selenium, and nitrocellulose were selected as COPCs because no sediment screening values are available. According to the USEPA Office of Drinking Water, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in Section 6.6. Four analytes [bis(2-ethylhexyl)phthalate, dibenzo(a,h)anthracene, lead, and mercury] were selected as COPCs because their maximum concentrations exceeded their respective sediment ESVs.

7.4.3 Selection of Surface Water COPCs

The following summarizes the procedures that were used in the SERA for LL9 to select COPCs:

- Inorganic contaminants whose maximum concentrations do not exceed background are not selected as COPCs.
- Inorganic and organic contaminants whose maximum concentrations do not exceed water quality criteria in OAC 3745-1 (or the AWQC, as discussed in Section 7.3) are not retained as COPCs.
- Inorganic and organic contaminants without screening values are retained as COPCs.

Table 7-3 summarizes the screening of surface water COPCs. Manganese and nitrocellulose were selected as COPCs because no surface water screening level is available. According to the USEPA Office of Drinking Water, nitrocellulose is essentially nontoxic (USEPA, 2004d). Nitrocellulose is further discussed in Section 6.6. Six analytes (aluminum, copper, iron, lead, mercury, and zinc) were retained as COPCs because they had concentrations that exceeded their respective surface water ESVs.

7.5 Methodology for Refining Conservative Exposure Assumptions

As discussed in Section 7.1, Step 3a is technically the first step of a BERA (although it is included in this SERA) and consists of refining the conservative exposure assumptions/concentrations when evaluating potential risks to ecological receptors (i.e., plants, invertebrates, and wildlife receptors) and re-evaluating the analytical data using benchmarks that may be more appropriate for the assessment endpoints. The objective of the Step 3a evaluation is to reduce the number of chemicals retained as COPCs, if possible, so the SERA can focus on the remaining COPCs that pose the greatest risks to ecological receptors. The Step 3a evaluation is designed to eliminate chemicals from further evaluation for certain groups of receptors.



Table 7-1
Ecological COPC Selection Table of Shallow Soil (0 - 4 ft bgs)

Parameter	Frequency of Detection	Minimum Concentration	Maximum Concentration	Mean Concentration	Average of Positive Detects	Sample of Maximum Detect	95% UCL on Mean ⁽¹⁾	Site Background Criteria ⁽¹⁾
Semivolatile Organics (µg/kg)								
2-METHYLNAPHTHALENE	1/13	15 J	15 J	1.98	15.0	LL9SS-068-0001-SO	2.66	NA
ACENAPHTHENE	4/13	6 J	19 Ja	3.81	10.5	LL9SB-050-0001-SO	10.7	NA
ACENAPHTHYLENE	1/13	50 J	50 J	4.35	50.0	LL9SS-068-0001-SO	5.64	NA
ANTHRACENE	4/13	6.3 J	48 J	7.71	23.9	LL9SS-068-0001-SO	48	NA
BENZO(A)ANTHRACENE	10/13	3.1 J	230	31.3	40.5	LL9SS-068-0001-SO	230	NA
BENZO(A)PYRENE	9/13	6.1 J	240	32.2	46.0	LL9SS-068-0001-SO	240	NA
BENZO(B)FLUORANTHENE	10/13	4.5 J	240 M	34.6	44.6	LL9SS-068-0001-SO	240	NA
BENZO(G,H,I)PERYLENE	10/13	3.3 J	170	23.3	29.9	LL9SS-068-0001-SO	141	NA
BENZO(K)FLUORANTHENE	9/13	4.9 J	200 M	27.5	39.0	LL9SS-068-0001-SO	151	NA
BIS(2-ETHYLHEXYL)PHTHALATE	4/13	14 Ja	150 J	19.3	49.8	LL9SS-014-0001-SO	40.1	NA
CHRYSENE	10/13	3.1 J	250	35.0	45.2	LL9SS-068-0001-SO	250	NA
DIBENZO(A,H)ANTHRACENE	7/13	6.4 J	130	20.6	37.1	LL9SS-068-0001-SO	130	NA
DIBENZOFURAN	2/13	7.9 Ja	13 J	2.98	10.5	LL9SS-068-0001-SO	4.41	NA
FLUORANTHENE	10/13	6.4 J	360	60.8	78.8	LL9SS-068-0001-SO	360	NA
FLUORENE	3/13	8.5 Ja	25 J	4.18	14.9	LL9SS-068-0001-SO	9.81	NA
INDENO(1,2,3-CD)PYRENE	10/13	3.9 J	170	26.6	34.2	LL9SS-068-0001-SO	170	NA
NAPHTHALENE	1/13	19 J	19 J	2.41	19.0	LL9SS-068-0001-SO	3.23	NA
PHENANTHRENE	9/13	4.9 J	280	40.4	58.1	LL9SS-068-0001-SO	280	NA
PYRENE	10/13	4 J	400	56.9	73.6	LL9SS-068-0001-SO	400	NA
Energetics (µg/kg)								
NITROGUANIDINE	1/9	89	89	121	89.0	LL9SS-034-0001-SO	89	NA
RDX	1/21	110	110	37.0	69.5	LL9SB-053-0001-SO	41.5	NA
Inorganics (mg/kg)								
ALUMINUM	85/85	5,700	20,000	10,182	10,182	LL9SS-023-0001-SO	10692	17700
ANTIMONY	21/85	0.19 B	1.4	0.365	0.723	LL9SS-011-0001-SO	0.392	0.96
ARSENIC	85/85	3.1	32	12.3	12.3	LL9SB-019-0001-SO	13.6	15.4
BARIUM	85/85	26	170	63.7	63.7	LL9SS-019-0001-SO	68.5	88.4
BERYLLIUM	85/85	0.16 B	1.8	0.493	0.493	LL9SS-005-0001-SO	0.527	0.88
CADMIUM	20/85	0.095 B	2.9	0.183	0.625	LL9SS-006-0001-SO	0.152	NA
CALCIUM	85/85	11	113,000 H	4,198	4,198	LL9SS-005-0001-SO	3769	15800
CHROMIUM	85/85	8.1	110	15.4	15.4	LL9SS-024-0001-SO	15.9	17.4
COBALT	85/85	1.6	18	8.17	8.17	LL9SS-027-0001-SO	8.92	10.4
COPPER	85/85	5.2	1240	32.7	32.7	LL9SS-011-0001-SO	24.2	17.7
IRON	85/85	6,140 H	76,000	22,357	22,357	LL9SS-031-0001-SO	23706	23100
LEAD	85/85	5.2	1330	46.2	46.2	LL9SS-011-0001-SO	40.2	19.1
MAGNESIUM	85/85	1,100	9,600	2,275	2,275	LL9SS-005-0001-SO	2417	3030
MANGANESE	85/85	50	3,800	559	559	LL9SS-027-0001-SO	634	1450
MERCURY	85/85	0.0057 B	882	10.8	10.8	LL9SS-011-0001-SO	0.433	0.04
NICKEL	85/85	5.7	32	16.1	16.1	LL9SB-015-0001-SO	17.2	21.1
POTASSIUM	85/85	2	1,600	821	821	LL9SB-024-0001-SO, LL9SB-025-0001-SO, LL9SB-052-0001-SO	1350	927
SELENIUM	38/85	0.15 B	1.8 B	0.436	0.648	LL9SS-068-0001-SO	0.484	1.4
SODIUM	24/85	94.3	1,300	246	748	LL9SS-038-0001-SO, LL9SS-046-0001-SO, LL9SB-065-0001-SO	266	123
THALLIUM	1/85	0.33	0.33	0.177	0.33	LL9SB-021-0001-SO	0.19	0.91
VANADIUM	85/85	6.9	29	17.5	17.5	LL9SB-024-0001-SO, LL9SB-025-0001-SO	18.5	31.1
ZINC	85/85	27	780	87.6	87.6	LL9SS-068-0001-SO	89.3	61.8
Miscellaneous Parameters								
NITROCELLULOSE (ug/kg)	9/9	1,100	4,300	2,206	2,206	LL9SB-050-0001-SO	4300	NA



Table 7-1
Ecological COPC Selection Table of Shallow Soil (0 - 4 ft bgs)

Parameter	Detects > Site Background Criteria ⁽²⁾	Max Conc > Bkg Conc	Surface Soil Screening Level		HQ	Bio Chemical	PBT Chemical	Selected as a COPC?	Rationale ⁽³⁾
			Value	Source					
Semivolatile Organics (µg/kg)									
2-METHYLNAPHTHALENE	NA	NA	3,240	d	0.005	NO	NO	NO	BSL
ACENAPHTHENE	NA	NA	20,000	a	0.001	YES	NO	YES	BIO
ACENAPHTHYLENE	NA	NA	682,000	d	0.00007	YES	NO	YES	BIO
ANTHRACENE	NA	NA	1,480,000	d	0.00003	YES	NO	YES	BIO
BENZO(A)ANTHRACENE	NA	NA	5,210	d	0.04	YES	NO	YES	BIO
BENZO(A)PYRENE	NA	NA	1,520	d	0.16	YES	NO	YES	BIO
BENZO(B)FLUORANTHENE	NA	NA	59,800	d	0.004	YES	NO	YES	BIO
BENZO(G,H,I)PERYLENE	NA	NA	119,000	d	0.001	YES	NO	YES	BIO
BENZO(K)FLUORANTHENE	NA	NA	148,000	d	0.001	YES	NO	YES	BIO
BIS(2-ETHYLHEXYL)PHTHALATE	NA	NA	926	d	0.16	NO	NO	NO	BSL
CHRYSENE	NA	NA	4,730	d	0.05	YES	NO	YES	BIO
DIBENZO(A,H)ANTHRACENE	NA	NA	18,400	d	0.007	YES	NO	YES	BIO
DIBENZOFURAN	NA	NA	NA	NA	NA	NO	NO	YES	NTX
FLUORANTHENE	NA	NA	122,000	d	0.003	YES	NO	YES	BIO
FLUORENE	NA	NA	30,000	b	0.0008	YES	NO	YES	BIO
INDENO(1,2,3-CD)PYRENE	NA	NA	109,000	d	0.002	YES	NO	YES	BIO
NAPHTHALENE	NA	NA	99.4	d	0.19	NO	NO	NO	BSL
PHENANTHRENE	NA	NA	45,700	d	0.006	YES	NO	YES	BIO
PYRENE	NA	NA	78,500	d	0.005	YES	NO	YES	BIO
Energetics (µg/kg)									
NITROGUANIDINE	NA	NA	NA	NA	NA	NO	NO	YES	NTX
RDX	NA	NA	NA	NA	NA	NO	NO	YES	NTX
Inorganics (mg/kg)									
ALUMINUM	3/85	YES	50	c	400	NO	NO	YES	ASL
ANTIMONY	4/85	YES	5	a	0.28	NO	NO	NO	BSL
ARSENIC	22/85	YES	9.9	a	3.23	YES	NO	YES	ASL,BIO
BARIUM	11/85	YES	283	a	0.60	NO	NO	NO	BSL
BERYLLIUM	1/85	YES	10	a	0.18	NO	NO	NO	BSL
CADMIUM	NA	NA	4	a	0.73	YES	NO	YES	BIO
CALCIUM	4/85	YES	NA	NA	NA	NO	NO	NO	NUT
CHROMIUM	10/85	YES	0.4	a	275	YES	NO	YES	ASL,BIO
COBALT	15/85	YES	20	a	0.9	NO	NO	NO	BSL
COPPER	44/85	YES	60	a	20.7	YES	NO	YES	ASL,BIO
IRON	33/85	YES	200	b ⁽⁴⁾	380	NO	NO	YES	ASL
LEAD	38/85	YES	40.5	a	32.8	YES	NO	YES	ASL,BIO
MAGNESIUM	13/85	YES	NA	NA	NA	NO	NO	NO	NUT
MANGANESE	2/85	YES	100	b ⁽⁴⁾	38	NO	NO	YES	ASL
MERCURY	37/85	YES	0.00051	a	1,729,000	YES	YES	YES	ASL,BIO,PBT
NICKEL	13/85	YES	30	a	1.07	YES	NO	YES	ASL,BIO
POTASSIUM	25/85	YES	NA	NA	NA	NO	NO	NO	NUT
SELENIUM	1/85	YES	0.21	a	8.57	YES	NO	YES	ASL,BIO
SODIUM	22/85	YES	NA	NA	NA	NO	NO	NO	NUT
THALLIUM	0/85	NO	1	a	0.33	NO	NO	NO	BKG
VANADIUM	0/85	NO	2	a	14.5	NO	NO	YES	ASL
ZINC	49/85	YES	8.5	a	91.8	YES	NO	YES	ASL,BIO
Miscellaneous Parameters									
NITROCELLULOSE (ug/kg)	NA	NA	NA	NA	NA	NO	NO	YES	NTX



Table 7-1
Ecological COPC Selection Table of Shallow Soil (0 - 4 ft bgs)

- Notes:**
- 1 - Shaded cells indicate parameter was selected as a COPC
 - 2 - Background criteria as described in Section 4.
 - 3 - For duplicate pairs, the average chemical concentrations of the original and duplicate samples were compared to the site background criteria.
 - 4 - Rationale Codes
 - Above Screening Levels (ASL)
 - Toxicity information not available (NTX)
 - Maximum detected concentration is below background screening criteria (BKG)
 - Below Screening Levels (BSL)
 - Essential Nutrient (NUT)
 - Bioaccumulative Chemical (BIO)
 - Persistent, Bioaccumulative, and Toxic chemical (PBT)
 - 5 - Value is based on effects to microorganisms.
- Sources of Screening Levels and Heirarchy for Selection:**
- a - Preliminary Remediation Goals from Efroymsen et al. (1997a)
 - b - Toxicological Benchmarks for Screening Contaminants of Potential Concern for Effects on Soil and Litter Invertebrates and Heterotrophic Processes from Efroymsen et al. (1997b)
 - c - Toxicological Benchmarks for Screening Contaminants of Potential Concern for Effects on Terrestrial Plants from Efroymsen et al. (1997c)
 - d - Ecological Data Quality Level (USEPA Region 5, 1999)



Table 7-2
Ecological COPC Selection Table for Sediment

Parameter	Frequency of Detection	Minimum Concentration	Maximum Concentration	Mean Concentration	Average of Positive Detects	Sample of Maximum Detect	Site Background Criteria ⁽¹⁾	Detects > Site Background Criteria ⁽²⁾	Ohio Sediment Reference Value (SRV)	Max Conc > Bkg Conc and Ohio SRV	Sediment Screening Level		HQ	Selected as a COPC?	Rationale ⁽³⁾
											Value	Source			
Semivolatile Organics (µg/kg)															
2-METHYLNAPHTHALENE	1/1	4 Ja	4 Ja	2.70	2.70	LL9SD-015-0001-SD	NA	NA	NA	NA	20.2	b	0.20	NO	BSL
ACENAPHTHYLENE	1/1	1.8 Ja	1.8 Ja	1.33	1.33	LL9SD-015-0001-SD	NA	NA	NA	NA	5.87	b	0.31	NO	BSL
ANTHRACENE	1/1	3.1 Ja	3.1 Ja	1.95	1.95	LL9SD-015-0001-SD	NA	NA	NA	NA	57.2	a	0.05	NO	BSL
BENZO(A)ANTHRACENE	1/1	16 Ja	20 Ja	18.0	18.0	LL9SD-015-0001-SD	NA	NA	NA	NA	108	a	0.19	NO	BSL
BENZO(A)PYRENE	1/1	21 Ja	25 Ja	23.0	23.0	LL9SD-015-0001-SD	NA	NA	NA	NA	150	a	0.17	NO	BSL
BENZO(B)FLUORANTHENE	1/1	20 Ja	24 Ja	22.0	22.0	LL9SD-015-0001-SD	NA	NA	NA	NA	10,400	b	0.002	NO	BSL
BENZO(G,H,I)PERYLENE	1/1	13 Ja	18 Ja	15.5	15.5	LL9SD-015-0001-SD	NA	NA	NA	NA	170	b	0.11	NO	BSL
BENZO(K)FLUORANTHENE	1/1	19 Ja	29 Ja	24.0	24.0	LL9SD-015-0001-SD	NA	NA	NA	NA	240	b	0.12	NO	BSL
BIS(2-ETHYLHEXYL)PHTHALATE	1/1	550	660	605	605	LL9SD-015-0001-SD	NA	NA	NA	NA	182	b	3.63	YES	ASL
BUTYL BENZYL PHTHALATE	1/1	8.8 Ja	8.8 Ja	6.33	6.33	LL9SD-015-0001-SD	NA	NA	NA	NA	4,190	b	0.002	NO	BSL
CHRYSENE	1/1	23 Ja	29 Ja	26.0	26.0	LL9SD-015-0001-SD	NA	NA	NA	NA	166	a	0.17	NO	BSL
DI-N-BUTYL PHTHALATE	1/1	38 Ja	38 Ja	28.3	28.3	LL9SD-015-0001-SD	NA	NA	NA	NA	110.5	b	0.34	NO	BSL
DIBENZO(A,H)ANTHRACENE	1/1	43 Ja	43 Ja	22.4	22.4	LL9SD-015-0001-SD-D	NA	NA	NA	NA	33	a	1.30	YES	ASL
FLUORANTHENE	1/1	37 Ja	48 Ja	42.5	42.5	LL9SD-015-0001-SD	NA	NA	NA	NA	423	a	0.11	NO	BSL
INDENO(1,2,3-CD)PYRENE	1/1	37 Ja	39 Ja	38.0	38.0	LL9SD-015-0001-SD	NA	NA	NA	NA	200	b	0.20	NO	BSL
PHENANTHRENE	1/1	20 Ja	20 Ja	20.0	20.0	LL9SD-015-0001-SD, LL9SD-015-0001-SD-D	NA	NA	NA	NA	204	a	0.10	NO	BSL
PYRENE	1/1	28 Ja	34 Ja	31.0	31.0	LL9SD-015-0001-SD	NA	NA	NA	NA	195	a	0.17	NO	BSL
Inorganics (mg/kg)															
ALUMINUM	13/13	5,500	15,600	11,635	11,635	LL9SD-003-0001-SD	13,900	3/13	29,000	NO	NA	NA	NA	NO	BKG
ANTIMONY	1/13	0.49	0.49	0.347	0.355	LL9SD-001-0001-SD	NA	NA	1.3	NO	NA	NA	NA	NO	BKG
ARSENIC	13/13	4.2	17.3	11.7	11.7	LL9SD-003-0001-SD	19.5	0/13	25	NO	9.79	a	1.8	NO	BKG
BARIUM	13/13	58	140	84.0	84.0	LL9SD-002-0001-SD	123	1/13	190	NO	NA	NA	NA	NO	BKG
BERYLLIUM	13/13	0.31 B	0.99	0.638	0.638	LL9SD-001-0001-SD	0.38	12/13	0.8	YES	NA	NA	NA	YES	NTX
CADMIUM	10/13	0.19 B	0.77	0.340	0.416	LL9SD-004-0001-SD	NA	NA	0.79	NO	0.99	a	0.78	NO	BKG
CALCIUM	13/13	870	11,000	2,266	2,266	LL9SD-016-0001-SD-D	5,510	1/13	21,000	NO	NA	NA	NA	NO	NUT
CHROMIUM	13/13	6.3	22	15.3	15.3	LL9SD-016-0001-SD-D	18.1	3/13	29	NO	43.4	a	0.51	NO	BKG
COBALT	13/13	5.2	9.5	7.67	7.67	LL9SD-016-0001-SD-D, LL9SD-017-0001-SD	9.1	2/13	12	NO	50	b	0.19	NO	BKG
COPPER	13/13	7.6	23	15.0	15.0	LL9SD-003-0001-SD	27.6	0/13	32	NO	31.6	a	0.73	NO	BKG
IRON	13/13	17,000 H	39,000	24,265	24,265	LL9SD-002-0001-SD	28,200	2/13	41,000	NO	NA	NA	NA	NO	BKG
LEAD	13/13	15	100	40.1	40.1	LL9SD-013-0001-SD	27.4	10/13	47	YES	35.8	a	2.79	YES	ASL
MAGNESIUM	13/13	620	3,450	2,149	2,149	LL9SD-003-0001-SD	2,760	2/13	7,100	NO	NA	NA	NA	NO	NUT
MANGANESE	13/13	200	1,400	678	678	LL9SD-017-0001-SD	1,950	0/13	1,500	NO	NA	NA	NA	NO	BKG
MERCURY	13/13	0.026	2.9	0.412	0.412	LL9SD-012-0001-SD	0.06	11/13	0.12	YES	0.18	a	16.1	YES	ASL
NICKEL	13/13	11	22.4	17.5	17.5	LL9SD-001-0001-SD	17.7	7/13	33	NO	22.7	a	0.99	NO	BKG
POTASSIUM	13/13	350	1,820	1,002	1,002	LL9SD-003-0001-SD	1,950	0/13	6,800	NO	NA	NA	NA	NO	NUT
SELENIUM	11/13	0.7 B	2.1	1.19	1.31	LL9SD-004-0001-SD	1.7	2/13	1.7	YES	NA	NA	NA	YES	NTX
SODIUM	3/13	120 B	213	98.1	182	LL9SD-003-0001-SD	112	3/13	NA	YES	NA	NA	NA	NO	NUT
THALLIUM	1/13	0.31	0.31	0.171	0.310	LL9SD-012-0001-SD	0.89	0/13	4.7	NO	NA	NA	NA	NO	BKG
VANADIUM	13/13	7.6	27.2	21.3	21.3	LL9SD-005-0001-SD	26.1	2/13	40	NO	NA	NA	NA	NO	BKG
ZINC	13/13	9	420	134	134	LL9SD-004-0001-SD	532	0/13	160	NO	121	a	3.47	NO	BKG
Miscellaneous Parameters															
NITROCELLULOSE (µg/kg)	1/1	2,800	4,200	3,500	3,500	LL9SD-015-0001-SD-D	NA	NA	NA	NA	NA	NA	NA	YES	NTX
TOTAL ORGANIC CARBON (mg/kg)	13/13	3,500	28,000	15,323	15,323	LL9SD-013-0001-SD	NA	NA	NA	NA	NA	NA	NA	NO	NA

Notes:

- 1 - Shaded cells indicate parameter was selected as a COPC
2 - Background criteria as described in Section 4.
3 - For duplicate pairs, the average chemical concentrations of the original and duplicate samples were compared to the site background criteria.
4 - Rationale Codes
Above Screening Levels (ASL)
Toxicity information not available (NTX)
Maximum detected concentration is less than the site background criteria level or Ohio Sediment Reference Value (BKG)
Below Screening Level (BSL)
Essential Nutrient (NUT)

Sources of Screening Levels and Heirarchy for Selection:

- a - Threshold Effects Concentration from McDonald et al., (2000)
b - Ecological Data Quality Level (USEPA Region 5, 1999)



Table 7-3
Ecological COPC Selection Table for Surface Water

Parameter	Frequency of Detection	Minimum Concentration	Maximum Concentration	Mean Concentration	Average of Positive Detects	Sample of Maximum Detect	Site Background Criteria ⁽¹⁾	Detects > Site Background Criteria ⁽²⁾	Max Conc > Bkg Conc	Surface Water Screening Level		HQ	PBT Chemical	Selected as a COPC?	Rationale ⁽³⁾
										Value	Source				
Inorganics (µg/L)															
ALUMINUM	6/6	339	17,800	5,929	5,929	LL9SW-005-0001-SW	3,370	2/6	YES	87	b	205	NO	YES	ASL
ARSENIC	4/6	1.9 B	9.9	4.24	6.03	LL9SW-003-0001-SW	3.2	3/6	YES	340 ⁽⁵⁾	a	0.03	NO	NO	BSL
BARIUM	6/6	21.4	145	71.2	71.2	LL9SW-005-0001-SW	47.5	3/6	YES	2000 ⁽⁴⁾	a	0.07	NO	NO	BSL
BERYLLIUM	3/6	0.26 B	0.7 B	0.572	0.420	LL9SW-005-0001-SW	NA	NA	NA	15.9 ^(4,6)	a	0.04	NO	NO	BSL
CADMIUM	1/6	1.1 B	1.1 B	0.497	1.10	LL9SW-005-0001-SW	NA	NA	NA	1.23 ^(4,5,6)	a	0.89	NO	NO	BSL
CALCIUM	6/6	9,830 H	61,000	20,772	20,772	LL9SW-003-0001-SW	41,400	1/6	YES	NA	NA	NA	NO	NO	NUT
CHROMIUM	4/6	3.3 B	20.2	7.84	10.3	LL9SW-005-0001-SW	NA	NA	NA	231 ^(4,5,6)	a	0.09	NO	NO	BSL
COBALT	4/6	1.2 B	8	3.80	4.95	LL9SW-003-0001-SW	NA	NA	NA	220 ⁽⁴⁾	a	0.04	NO	NO	BSL
COPPER	4/6	4.4 B	20	9.10	12.2	LL9SW-003-0001-SW	7.9	2/6	YES	4.76 ^(4,5,6)	a	4.20	NO	YES	ASL
IRON	6/6	249	20,000	9,973	9,973	LL9SW-002-0001-SW	2,560	4/6	YES	1000	b	20	NO		ASL
LEAD	5/6	1.2 B	70	25.5	30.4	LL9SW-003-0001-SW	NA	NA	NA	23.8 ^(4,5,6)	a	2.94	NO		ASL
MAGNESIUM	6/6	2,110	9,600	4,578	4,578	LL9SW-003-0001-SW	10,800	0/6	NO	NA	NA	NA	NO	NO	NUT
MANGANESE	6/6	10.1	1,400	638	638	LL9SW-003-0001-SW	391	4/6	YES	NA	NA	NA	NO	YES	NTX
MERCURY	5/6	0.065 B	1.7	0.341	0.389	LL9SW-012-0001-SW-D	NA	NA	NA	1.4 ⁽⁵⁾	a	1.21	YES	YES	ASL,PBT
NICKEL	5/6	4 B	20.1	10.0	11.0	LL9SW-005-0001-SW	NA	NA	NA	184 ^(4,5,6)	a	0.11	NO	NO	BSL
POTASSIUM	6/6	798	5,200	2,366	2,366	LL9SW-003-0001-SW	3,170	2/6	YES	NA	NA	NA	NO	NO	NUT
SODIUM	6/6	966 B	4,300	2,161	2,161	LL9SW-003-0001-SW	21,300	0/6	NO	NA	NA	NA	NO	NO	NUT
VANADIUM	3/6	4.9 B	31.9	10.1	18.6	LL9SW-005-0001-SW	NA	NA	NA	150 ⁽⁴⁾	a	0.21	NO	NO	BSL
ZINC	6/6	19	228	91.7	91.7	LL9SW-005-0001-SW	42	2/6	YES	46 ^(4,5,6)	a	4.96	NO	YES	ASL
Miscellaneous Parameters (mg/L)															
NITROCELLULOSE	1/1	0.1	0.12	0.110	0.110	LL9SW-012-0001-SW	NA	NA	NA	NA	NA	NA	NO	YES	NTX

Notes:

- 1 - Background criteria as described in Section 4.
- 2 - For duplicate pairs, the average chemical concentrations of the original and duplicate samples were compared to the site background criteria.
- 3 - Rationale Codes
Above Screening Levels (ASL)
Toxicity information not available (NTX)
Essential Nutrient (NUT)
Below Screening Levels (BSL)
Persistent, Bioaccumulative, and Toxic chemical (PBT)
- 4 - Outside Mixing Zone Maximum value.
- 5 - Dissolved criteria.
- 6 - Based on a calculated water hardness of 33.2 mg/L (in order to be most conservative, the minimum hardness at location LL9SW-001 was used).

Sources of Screening Levels and Heirarchy for Selection:

- a - Ohio Water Quality Criteria
b - USEPA Recommended Water Quality Criteria



For example, if a chemical concentration is less than a toxicity benchmark for invertebrates, but greater than a benchmark for plants, the chemical may be eliminated as a COPC in soil for risks to soil invertebrates, but it may be retained as a COPC in soil for risks to plants. Therefore, chemicals are evaluated during Step 3a in order of plants/invertebrates, aquatic receptors, and wildlife. The following sections present the methodology for evaluating risks to these receptors.

7.5.1 Surface Soil

Chemicals that were initially selected as COPCs in surface soil were evaluated for risks to: (1) plants, (2) invertebrates, and (3) wildlife (i.e., mammals and birds). This further evaluation was conducted to determine whether there are potential risks to all three receptor groups (i.e., plants, invertebrates, and wildlife), or to only one or two of the receptor groups. This is important because if the site proceeds further in a BERA, the studies in the BERA should focus only on the receptors that are potentially at risk. The first step in the Step 3a evaluation was to compare the maximum and average chemical concentrations in the soil to toxicity benchmarks that are based on effects to earthworms and plants. Tables 7-4 and 7-5 present this comparison for earthworms and plants, respectively.

The following hierarchy was used to select the earthworm benchmark in Table 7-4:

- USEPA ecological soil screening levels (Eco-SSLs) for earthworms (USEPA, 2003a,b and c). These values were selected first because they are USEPA values and are the most recent values.
- The greater of the values from the following two sources. Because both of the following sources are typically used as screening levels in ERAs, the values from either source should be protective of earthworms. For that reason, the greater value was selected to refine the list of chemicals retained as COPCs.
 - Canadian Soil Quality Guidelines (individual documents for each chemical), and
 - Oak Ridge National Laboratory (ORNL), *Toxicological Benchmarks for Contaminants of Potential Concern for Effects on Soil and Litter Invertebrates and Heterotrophic Process*, 1997 Revision (Efroymson, et al., 1997b).

The following hierarchy was used to select the plant benchmark in Table 7-5:

- USEPA Eco-SSLs for plants (USEPA, 2003 a, b, and c) These values were selected first because they are USEPA values and are the most recent values.
- The greater of the values from the following two sources. Because both of the following sources are typically used as screening levels in ERAs, the values from either source should be protective of plants. For that reason, the greater value was selected to refine the list of chemicals retained as COPCs.



Table 7-4
Direct Toxicity to Earthworms

Parameter	Frequency of Detection	Maximum Concentration	Average Concentration	Selected Upper Confidence Level	Earthworm Toxicity Benchmarks	Source	Maximum > TRV?	Average > TRV?
Semivolatile Organics (µg/kg)								
DIBENZOFURAN	2/13	13	2.98	4.41	No TRV	NA	No TRV	No TRV
Energetics (ug/kg)								
NITROGUANIDINE	1/9	89	121	89	No TRV	NA	No TRV	No TRV
RDX	1/21	110	37.0	41.5	No TRV	NA	No TRV	No TRV
Inorganics (mg/kg)								
ALUMINUM	85/85	20,000	10,182	10692	pH ⁽¹⁾	a	NA	NA
ARSENIC	85/85	32	12.3	13.6	60	c	NO	NO
CHROMIUM	85/85	110	15.4	15.9	64	d	YES	NO
COPPER	85/85	1240	32.7	24.2	63	d	YES	NO
IRON	85/85	76,000	22,357	23706	No TRV	NA	No TRV	No TRV
LEAD	85/85	1330	46.2	40.2	1700	b	NO	NO
MANGANESE	85/85	3,800	559	634	No TRV	NA	No TRV	No TRV
MERCURY	85/85	882	10.8	0.433	12	d	YES	NO
NICKEL	85/85	32	16.1	17.2	200	c	NO	NO
SELENIUM	38/85	1.8	0.436	0.484	70	c	NO	NO
VANADIUM	85/85	29	17.5	18.5	130	d	NO	NO
ZINC	85/85	780	87.6	89.3	200	d	YES	NO
Miscellaneous Parameters (µg/kg)								
NITROCELLULOSE	9/9	4,300	2,206	4300	No TRV	NA	No TRV	No TRV

Notes:

- Shaded cells are chemicals that are retained for further evaluation in Step 3a.
- Average concentration includes positive detections and non-detected results. Detection limits are divided by two for non-detected values.
- Frequency of detection refers to number of times compound was detected among all samples versus total number of samples analyzed for that parameter.
- Number of samples may vary based on the number of usable results.
- 1 - Aluminum is only considered a COPC when the pH is <5.5

Sources of Earthworm Toxicity Benchmarks:

- a - Ecological Soil Screening Level (Eco-SSL) from U.S. EPA (2003b).
- b - Ecological Soil Screening Level (Eco-SSL) from U.S. EPA (2005b).
- c - Earthworm toxicity benchmark from Efroymson et al., 1997b.
- d - Canadian Soil Quality Guideline (SQG) from Environment Canada (EC, 1999a-c, e-f).



Table 7-5
Direct Toxicity to Plants

Parameter	Frequency of Detection	Maximum Concentration	Average Concentration	Selected Upper Confidence Level	Plant Toxicity Benchmarks	Source	Maximum > TRV?	Average > TRV?
Semivolatile Organics (µg/kg)								
DIBENZOFURAN	2/13	13	2.98	4.41	No TRV	NA	No TRV	No TRV
Energetics (ug/kg)								
NITROGUANIDINE	1/9	89	121	89	No TRV	NA	No TRV	No TRV
RDX	1/21	110	37.0	41.5	No TRV	NA	No TRV	No TRV
Inorganics (mg/kg)								
ALUMINUM	85/85	20,000	10,182	10692	pH ⁽¹⁾	a	NA	NA
ARSENIC	85/85	32	12.3	13.6	18	b	YES	NO
CHROMIUM	85/85	110	15.4	15.9	64	d	YES	NO
COPPER	85/85	1240	32.7	24.2	100	c	YES	NO
IRON	85/85	76,000	22,357	23706	pH ⁽¹⁾	a	NA	NA
LEAD	85/85	1330	46.2	40.2	120	b	YES	NO
MANGANESE	85/85	3,800	559	634	500	c	YES	YES
MERCURY	85/85	882	10.8	0.433	12	d	YES	NO
NICKEL	85/85	32	16.1	17.2	50	d	NO	NO
SELENIUM	38/85	1.8	0.436	0.484	1	c	YES	NO
VANADIUM	85/85	29	17.5	18.5	130	d	NO	NO
ZINC	85/85	780	87.6	89.3	200	d	YES	NO
Miscellaneous Parameters (µg/kg)								
NITROCELLULOSE	9/9	4,300	2,206	4300	No TRV	NA	No TRV	No TRV

Notes:

- Shaded cells are chemicals that are retained for further evaluation in Step 3a.
- Average concentration includes positive detections and non-detected results. Detection limits are divided by two for non-detected values.
- Frequency of detection refers to number of times compound was detected among all samples versus total number of samples analyzed for that parameter.
- Number of samples may vary based on the number of usable results.
- 1 - Benchmarks for aluminum and iron are based on pH of the soil.

Sources of Plant Toxicity Benchmarks:

- a - Ecological Soil Screening Level (Eco-SSL) from U.S. EPA (2003b,c).
- b - Ecological Soil Screening Level (Eco-SSL) from U.S. EPA, (2005a,b).
- c - Plant toxicity benchmark from Efroymson et al., 1997c.
- d - Canadian Soil Quality Guideline (SQG) from Environment Canada (EC, 1999 a, c-f).



Canadian Soil Quality Guidelines (individual documents for each chemical), and

- ORNL, *Toxicological Benchmarks for Screening Contaminants of Potential Concern for Effects on Terrestrial Plants*, 1997 Revision (Efroymson, et al., 1997c).

If the chemical concentration is less than the benchmarks for earthworms and/or plants, it was concluded that the chemical is not likely causing a risk to that receptor group and the chemical was not evaluated further in Step 3a for that receptor group.

If the chemical concentration was greater than the benchmarks for earthworms and/or plants (or the chemical does not have a benchmark), the chemical was further evaluated in Step 3a to determine whether the risks are great enough to warrant additional evaluations (i.e., proceed to a BERA, develop cleanup levels).

In addition, bioaccumulative chemicals that were retained as COPCs were further evaluated to determine whether there are likely risks to wildlife. If the chemical is not bioaccumulative, it was not carried through the food chain model (see Section 7.5.5), and it was concluded that the chemical is not likely causing a risk to wildlife. If the chemical is bioaccumulative, it was carried through the food chain model. The following bullets outline decisions that were made based on the results of the food chain model:

- If the HQ is less than 1.0, using average contaminant concentrations and the NOAEL as the toxicity organic value (TRV), it was concluded that the chemical is not likely causing a risk to wildlife and the chemical was not evaluated further in Step 3a.
- If the HQ is greater than 1.0, using average contaminant concentrations and the NOAEL as the TRV, the chemical was further evaluated in Step 3a to determine whether the risks are great enough to warrant additional evaluations (i.e., proceed to a BERA, develop cleanup levels).

For chemicals evaluated further in Step 3a, the other Step 3a factors described below in Section 7.5.4 were used to determine whether the risks are great enough to warrant additional evaluations (i.e., proceed to a BERA, develop cleanup levels).

7.5.2 Sediment

Chemicals selected as COPCs in sediment were evaluated to assess potential risks from the chemical to benthic invertebrates. The first step in the Step 3a evaluation was to compare the average chemical concentrations in the sediment to the screening level, and then to compare the maximum and average chemical concentrations in the sediment to higher effects-levels (as described below). Table 7-6 presents this comparison.



Table 7-6
Step 3a Table for Evaluating Risks to Sediment Invertebrates

Chemical of Potential Concern (COPC) ⁽¹⁾	Frequency of Detection	Maximum Concentration	Average Concentration	Screening Level Comparison		Higher Effects-Level Comparison			
				Screening Level ⁽²⁾	Average> Screening Level	Higher Effects Level	Source	Maximum> Higher Effects Level?	Average> Higher Effects Level?
Semivolatile Organics (µg/kg)									
BIS(2-ETHYLHEXYL)PHthalATE	1/1	660	605	182	YES	1300	c	NO	NO
DIBENZO(A,H)ANTHRACENE	1/1	43	22.4	33	NO	1989	b	NO	NO
Inorganics (mg/kg)									
BERYLLIUM	13/13	0.99	0.638	No TRV	No TRV	No TRV	NA	No TRV	No TRV
LEAD	13/13	100	40.1	35.8	YES	128	a	NO	NO
MERCURY	13/13	2.9	0.412	0.18	YES	1.06	a	YES	NO
SELENIUM	11/13	2.1	1.19	No TRV	No TRV	1	c	YES	YES
Miscellaneous Parameters (µg/kg)									
NITROCELLULOSE	1/1	4,200	3,500	No TRV	No TRV	No TRV	NA	No TRV	No TRV

Notes:

- Shaded cells are chemicals that are retained for further evaluation in Step 3a.

1 - This table only presents that chemicals that were retained as COPCs because the maximum detected concentration exceeded the screening level or the chemical did not have a screening level.

2 - Table 7-2 presents the sources of the screening levels.

Sources of Higher Effects Levels:

a - Probable Effects Concentration from MacDonald et al., 2000.

b - Severe Effect Level from OMOE, 1993. Value for dibenzo(a,h)anthracene was calculated using 1.53% TOC.

c - Apparent Effect Threshold from Buchman 1999.



As discussed in Section 7.3, the TEC from MacDonald et al. (2000) was used as the primary source of the sediment screening, followed by the USEPA Region 5 EDQL (USEPA Region 5, 1999). TECs are considered "lower effects levels" because they are intended to identify contaminant concentrations below which harmful effects on benthic organisms are not expected. Probable effects concentrations (PECs) are considered "higher effects levels" because they are intended to identify concentrations above which harmful effects are expected to occur. Therefore, the PECs were selected first as the higher effects level in Table 7-6. Because there are no corresponding higher effects levels for the EDQLs, the severe effects level (SEL) from the Ontario sediment quality guidelines (OMOE, 1993) and the apparent effects threshold (AET) in Buchman (1999), if available, were also used to evaluate risks to benthic organisms.

The chemicals that were selected as COPCs were evaluated further in Step 3a, using the benchmarks described above along with the other Step 3a factors described below, to determine whether the risks are great enough to warrant additional evaluations (i.e., proceed to a BERA, develop cleanup levels).

7.5.3 Surface Water

Chemicals selected as COPCs in surface water were further evaluated using the other Step 3a factors described below to determine whether the risks to aquatic organisms are great enough to warrant additional evaluations (i.e., proceed to a BERA, develop cleanup levels).

7.5.4 Other Step 3a Factors

For chemicals that are evaluated further in Step 3a, the following factors were evaluated, as appropriate, to determine whether the risks are great enough to warrant additional evaluations (i.e., proceed to a BERA, develop cleanup levels).

- Magnitude of criterion exceedance: Although the magnitude of the risks may not relate directly to the magnitude of a criterion exceedance, the magnitude of the criterion exceedance may be one item used in a lines-of-evidence approach to determine the need for further site evaluation. The greater the criterion exceedance, the greater the probability and concern that a possible risk exists.
- Frequency of chemical detection and spatial distribution: A chemical detected at a low frequency typically is of less concern than a chemical detected at higher frequency if toxicity and concentrations and spatial areas represented by the data are similar. Chemicals detected frequently were given greater consideration than those detected infrequently.
- Contaminant bioavailability: Many contaminants (especially metals) are present in the environment in forms that are typically not bioavailable, and the limited bioavailability



was considered when evaluating the exposures of receptors to site contaminants. Contaminants with generally less bioavailability are considered to be less toxic than the more bioavailable contaminants, all other factors being equal.

- Habitat: Although exceedances of criteria may occur, potential risks to ecological receptors may be minimal if there is little habitat for those receptors. Therefore, the extent of habitat was used qualitatively when considering additional evaluation. Areas with little habitat were of less concern than areas with suitable habitat to support the receptors of interest.
- Alternate benchmarks: Use of alternate benchmarks ensures that more realistic exposure assumptions are evaluated. However, some of the alternate benchmarks are overly protective for some receptors and may not have been available in some cases. For example, the EDQLs and PRGs for soil may be based on risks to small mammals. Therefore, an exceedance of that EDQL does not necessarily indicate that potential risks to plants or invertebrates exist, so other more appropriate benchmarks were used to evaluate potential risks to those receptors (as was conducted in Tables 7-4 and 7-5). Use of these alternate benchmarks was case-specific. Additionally, screening values may not have been available for some chemicals in some media. For example, chemicals retained as COPCs in Tables 7-1, 7-2, and 7-3 are evaluated in Step 3a using alternate benchmarks when available.

7.5.5 Terrestrial Food Chain Modeling

Terrestrial intake modeling was used to estimate the exposure of the COPCs to terrestrial wildlife receptors and to evaluate potential risks to terrestrial wildlife ingestion of the soil, surface water, plants, invertebrates, and small mammals. Note that the food chain model incorporates chemical concentrations that are estimated into various food items (i.e., plants, invertebrates), but it does not model the amount of chemical transferred from one trophic level to the next. Chemicals evaluated in the terrestrial food chain model were limited to those identified by the USEPA as bioaccumulative (USEPA, 2000). Although wildlife may be exposed to non-bioaccumulative chemicals via direct ingestion of the media (i.e., soil), the exposure to the chemical will be low compared to the exposure to bioaccumulative chemicals through the food. It is common practice to only include bioaccumulative chemicals in food chain models and this approach results in more chemicals being included in the food chain model than what is required by Ohio EPA (2003) in their risk assessment guidance. In Attachment A, Section 2, of Ohio EPA (2003), it states that "Food-web and direct contact evaluations are required for a Level III ERA and are dependent upon the type of contamination and the affected media. ... Persistent, bioaccumulative and toxic (PBT, see Level II ERA guidance) compounds are also to be evaluated using direct contact and food-web models..." The Ohio PBT list is only a subset of the list of bioaccumulative chemicals. Therefore, only bioaccumulative chemicals were included in the food chain model.



Risk to terrestrial receptors from COPCs in the soil, surface water, and prey is determined by estimating the chronic daily intake (CDI) and comparing the CDI to toxicity reference value (TRVs) representing acceptable daily doses in mg/kg/day. The TRVs are developed from NOAELs and LOAELs obtained from wildlife studies, if available. The majority of the TRVs are obtained from the ORNL *Toxicological Benchmarks for Wildlife*, 1996 revision (Sample et al., 1996). Other sources for NOAELs and LOAELs were used as necessary. Appendix W presents the TRVs that are used in this report and the derivation of the TRVs using the body-weight scaling equation presented below.

For avian species, the NOAEL (or LOAEL) for the test species is used as the NOAEL (or LOAEL) for the surrogate species in accordance with Sample et al. (1996). For mammalian species, the NOAEL (or LOAEL) from one species is adjusted to a NOAEL (or LOAEL) for the surrogate species using the following body-weight scaling equation from Sample et al. (1996):

$$\text{NOAELw} = \text{NOAELt} * (\text{bwt/bww})^{0.25}$$

Where: NOAELw = no-observed-adverse-effect-level for the surrogate wildlife species
NOAELt = no-observed-adverse-effect-level for the test species
bwt = body weight of the test species
bww = body weight of the surrogate test species

The body-weight scaling is done because studies have shown that, for mammals, numerous physiological functions such as metabolic rate, as well as responses to toxic chemicals, are a function of body size (Sample et al., 1996). However, Sample et al. (1996) indicated that physiological scaling factors may not be appropriate for birds. Therefore, no body-weight scaling was conducted for the bird TRVs. No avian TRVs were available for PAHs, so a default of 2 mg/kg/d was used for the NOAEL. Appendix W presents the body weights that are used for the surrogate and potential test species.

The Ravenna ERA guidance provided one set of TRVs that were either based on NOAELs, or based on LOAELs that were divided by 10. Therefore, LOAEL TRVs were estimated by multiplying the TRVs from the Ravenna ERA guidance by 10.

7.5.5.1 Characterization of Exposure

This section describes the potential or actual contact or co-occurrence of the contaminants with the receptors to determine their exposure dose.

Terrestrial soil invertebrates and plants are exposed to contaminants in the surface soil through direct contact, ingestion, or both. Aquatic organisms are exposed to contaminants in the surface water and sediment through direct contact, ingestion, or both. The maximum or selected upper confidence level (UCL) (whichever is lower) and average soil and/or surface



water concentrations are used in the food chain model to obtain a range of exposure from ingestion of soil, surface water, or both.

Exposure of the terrestrial receptors to the COPCs in the surface soil and surface water is determined by estimating the daily doses in mg/kg/day using exposure equations. The contaminant concentrations in the surface soil and surface water are used to calculate the CDI doses. The following equation presents the food chain model that is used for the surrogate species that are selected for modeling:

$$\text{CDI Dose (mg/kg/day)} = \frac{(\text{FI} * \text{FC}) + (\text{WI} * \text{C}_{\text{Sw}}) + (\text{SI} * \text{C}_{\text{Ss}})}{\text{BW}}$$

Where: CDI = chronic daily intake (mg/kg/day)
FI = food ingestion rate (kg/day)
FC = food concentration (mg/kg)
WI = surface water ingestion rate (L/day)
 C_{Sw} = contaminant concentration in surface water (mg/L)
BW = body weight (kg)
SI = incidental soil ingestion rate (kg/day)
 C_{Ss} = contaminant concentration in surface soil (mg/kg)

For inorganic and organic constituents in surface soil, the contaminant concentration of the prey items (i.e., earthworms) is calculated using the following equation:

$$\text{FC} = \text{C}_{\text{Ss}} * \text{BAF}$$

Where: FC = contaminant concentration in food (mg/kg)
 C_{Ss} = contaminant concentration in soil (mg/kg)
BAF = bioaccumulation factor (chemical-specific)

For inorganic and organic constituents in small mammal tissue, the contaminant concentration of the prey items (i.e., small mammal) is calculated using the following equation:

$$\text{FC} = \text{C}_{\text{Ss}} * \text{BAF}_{\text{si}} * \text{BAF}_{\text{dm}}$$

Where: FC = contaminant concentration in food (mg/kg)
 C_{Ss} = contaminant concentration in surface soil (mg/kg)
 BAF_{si} = bioaccumulation factor for soil to invertebrate (chemical-specific)
 BAF_{dm} = bioaccumulation factor for diet to mammal (chemical-specific)

The exposure assumptions (i.e., ingestion rate, body weight) are obtained from the RVAAP ecological risk guidance (USACE, 2003). Appendix W presents the exposure parameters that are used in the SERA. Note that the receptors' home ranges are not used quantitatively in the



food chain model. However, the home ranges are discussed qualitatively in the Step 3a evaluation and uncertainty analysis section, when applicable. The receptors home ranges were not quantitatively used in the food chain model because the home ranges for most of the receptors were less than the size of LL9 (69 acres). Although the home ranges for the owl, fox, and hawk are greater than 69 acres (see Table W-5), the home ranges were not used in the initial food chain to be conservative. If the HQs from the conservative food chain model for the owl, fox, and hawk are greater than 1.0, the home ranges would be quantitatively used for the refinement.

7.5.5.2 Ecological Risk Characterization

The risk characterization is the final phase of a risk assessment that compares the exposure to the ecological effects. It is at this phase that the likelihood of adverse effects occurring as a result of exposure to a stressor will be evaluated. The HQ approach (as described in Section 7.4) is used to characterize the risk to terrestrial receptors. This approach characterizes the potential effects by comparing exposure concentrations with the effects data. An HQ greater than 1.0 is considered to indicate a potential risk. The HQ is not an expression of probability, and the significance of values greater than 1.0 must be interpreted carefully, considering the number of uncertainties associated with the SERA process.

The HQ for the terrestrial wildlife model is calculated as follows:

$$HQ = DOSE / TRV$$

Where: HQ = ecological Effects Quotient (unitless)
Dose = daily intake dose (mg/kg/day)
TRV = toxicity reference value (NOAEL or LOAEL) (mg/kg/day)

7.6 Step 3a Refinement

Subsequent to the initial screening, other factors are considered to further refine COPCs, as discussed in Section 7.5. The following sections present the results of the Step 3a refinement.

7.6.1 Risks to Terrestrial Plants and Invertebrates – Surface Soil

Potential risks to terrestrial plants and terrestrial invertebrates from exposure to COPCs were evaluated using the methodologies described in Section 7.5.1. The following subsections discuss whether the chemicals initially selected as COPCs should be retained for further evaluation of risks to soil invertebrates and terrestrial vegetation. Note that bioaccumulative chemicals that were detected at concentrations less than their respective screening levels are not discussed in the following sections because the chemicals are not COPCs for risks to plants and invertebrates (see Table 7-1).



Tables 7-4 and 7-5 summarize the results of potential direct toxicity of chemicals in soil to earthworms and plants, respectively. The tables list only the chemicals that were retained as COPCs because they were detected at concentrations that exceeded screening levels or they did not have screening levels. Both maximum and average concentrations are compared to the toxicity benchmarks for earthworms and plants. Chromium, copper, mercury, and zinc were the only chemicals with maximum detected concentrations that were greater than the earthworm toxicity benchmarks; however, the average chemical concentrations for these chemicals were less than the benchmarks. Aluminum toxicity is dependent on the pH of the soil. Arsenic, chromium, copper, lead, manganese, mercury, selenium, and zinc were the only chemicals with maximum concentrations that were greater than the plant toxicity benchmarks; the average chemical concentration for manganese was also greater than the benchmarks. Aluminum and iron toxicity are dependent on the pH of the soil. Earthworm and/or plant benchmarks were not available for several of the chemicals.

The following text presents the Step 3a evaluation for the chemicals listed above that were detected at concentrations greater than the earthworm or plant toxicity benchmarks and chemicals that do not have earthworm or plant toxicity benchmarks.

Dibenzofuran

Dibenzofuran was initially selected as a COPC because a screening level was not available. Additional toxicity information for dibenzofuran is not available; therefore, this chemical was further evaluated using the other Step 3a factors listed in Section 7.5.4. This analyte was detected in 2 of 13 sample locations. The maximum concentration is at LL9SS-068-0001-SO, which is a wooded area where surface debris (glass, metal, and other rubbish) was found. The other detection is from a subsurface sample collected under the soil subfloor of a building (LL9SB-050-0001-SO). Because of the lack of toxicity data for dibenzofuran, its relatively low frequency of detection, and its relatively low maximum detected concentration (13 µg/kg), it is not likely that dibenzofuran is significantly affecting plants or invertebrates at LL9. Therefore, risks from dibenzofuran are not likely and it was not retained as a COPC.

Nitroguanidine

Nitroguanidine was initially selected as a COPC because a screening level was not available. Additional toxicity information for nitroguanidine is also not available; therefore, this chemical was further evaluated using the Step 3a factors listed in Section 7.5.4. This analyte was detected in 1 of 9 samples, with the detection found in the surface soil at LL9SS-034-0001-SO. A duplicate sample collected from this location was non-detect for nitroguanidine, so the initial detection could be an anomaly that is not representative of site contamination. Therefore, risks from nitroguanidine are not likely and nitroguanidine was not retained as a COPC.



RDX

RDX was initially selected as a COPC because a screening level was not available. The following alternate toxicity information for RDX was located:

- Plants – 100 mg/kg (Talmage et al., 1999); and
- Lowest-Observed-Effect Concentration for earthworm reproduction – 95 mg/kg (Robidoux et al., 2000).

The plant benchmark of 100 mg/kg from Talmage et al. (1999) is from a study in which an RDX concentration of 100 mg/kg significantly reduced the biomass of cucumber plants. The LOEC for earthworm reproduction was based on the productivity of juveniles (total number of juveniles, biomass, and number of juveniles per hatched cocoon) being significantly reduced by RDX at 95 mg/kg in soil; a no-observed-effect concentration (NOEC) was not generated from the study (Robidoux et al., 2000).

RDX was detected in 1 of 21 samples, with the detection at LL9SB-053-0001-SO. Because a duplicate sample collected from this location was non-detected for RDX, the average soil concentration for LL9 is 37 mg/kg. The average LL9 RDX concentration is less than both the plant and invertebrate benchmarks. Because the average RDX concentration at LL9 is less than the benchmarks and given the low frequency of detection, risks to plants and invertebrates are not likely and RDX was not retained as a COPC.

Aluminum

Aluminum was initially selected as a COPC because the maximum detected concentration (20,000 mg/kg) exceeded the ORNL plant benchmark of 50 mg/kg (Efroymson et al., 1997c). However, the screening value is the toxicological benchmark to protect plants and is based on laboratory toxicity testing using soluble aluminum added to soils. The standard analytical measurement of aluminum in soils is total recoverable aluminum (soluble + fixed). Comparison of total aluminum concentration in soils to soluble aluminum-based benchmarks and screening values is probably not appropriate. According to the Eco-SSL benchmark, aluminum is considered a COPC only when the pH is less than 5.5 (USEPA, 2003b). Although pH data are not available at LL9, the average pH values for LL 2, 3, and 4 are greater than 5.5 S.U. (SAIC, 2003a, 2003b, and 2003c); therefore, it is likely that soil pH at LL9 is also greater than 5.5 S.U. Additionally, the maximum concentration is slightly greater than the background concentration (17,700 mg/kg). With the detected considerations, potential impacts from aluminum are unlikely. Consequently, risks to plants and invertebrates from aluminum are not likely and aluminum is not retained as a COPC. Therefore, aluminum is not considered toxic to plants and invertebrates and was not retained as a COPC.



1 Arsenic

2 Arsenic was initially selected as a COPC because the maximum detected concentration (32
3 mg/kg) exceeded the ORNL PRG of 9.9 mg/kg (Efroymson, et al., 1997a). The PRG is based
4 on risks to wildlife (short-tailed shrew) and plants. Alternate benchmarks were used to further
5 evaluate risks to soil receptors. The maximum concentration is less than the ORNL earthworm
6 benchmark of 60 mg/kg (Efroymson et al., 1997b), but is greater than the Eco-SSL plant
7 benchmark or 18 mg/kg (USEPA, 2005a). Although potential risks to plants are possible, the
8 risks are not great enough to warrant retaining arsenic as a COPC for further evaluation in the
9 BERA for several reasons. The maximum detected arsenic concentration of 32 mg/kg is only
10 two times greater than the background concentration (15.4 mg/kg). Therefore, the average
11 arsenic concentration of 12.3 mg/kg was compared to the Eco-SSL plant benchmark to
12 determine whether risks are widespread. Because the average concentration is less than the
13 Eco-SSL plant benchmark and the background concentration, the overall site contamination is
14 low and further evaluation of risks to plants in the BERA is not necessary. Therefore, risks to
15 plants and invertebrates from arsenic are not likely and arsenic is not retained as a COPC.

16 Chromium

17 Chromium was retained as a COPC in soil because the maximum concentration of 110 mg/kg
18 exceeded the ORNL PRG of 0.4 mg/kg (Efroymson et al., 1997a). The Canadian soil quality
19 guideline (SQG) for chromium is 64 mg/kg (EC, 1999a), which is based on direct soil contact.
20 Only the maximum concentration at LL9SS-024-0001-SO exceeded this alternate benchmark.
21 This sample is located adjacent to an unused water tower (Figure 3-1). Because only 1 out of
22 the 85 samples exceeded the Canadian SQG, risks to plants and invertebrates are not likely
23 and chromium was not retained as a COPC.

24 Copper

25 Copper was retained as a COPC in soil because the maximum concentration of 1,240 mg/kg at
26 LL9SS-011-0001-SO exceeded the ORNL PRG of 60 mg/kg (Efroymson et al., 1997a).
27 Alternate benchmarks were used to further evaluate risks to soil receptors. The Canadian SQG
28 for copper is 63 mg/kg (EC, 1999b) and the ORNL plant benchmark for copper is 100 mg/kg
29 (Efroymson et al., 1997c). Only 2 of the 85 sample locations (LL9SS-011-0001-SO and
30 LL9SS-068-0001-SO) exceed both of these benchmarks. Sample LL9SS-011-0001-SO is a
31 subfloor sample collected where a detonator destroying house with a wood frame and a sand
32 floor used to be. The explosives that were destroyed in this building included lead azide and
33 mercury fulminate; but other metals, such as copper, may have been present in these
34 explosives as well. Location LL9SS-068-0001-SO, with a copper concentration of 170 mg/kg,
35 is a wooded area where surface debris (glass, metal, and other rubbish) was found, so it is likely
36 that the copper is associated with this debris. Although only 2 of the 85 samples had copper



concentrations that exceeded the benchmarks, because of the elevated copper concentrations at these locations, copper was retained as a COPC for further evaluation.

Iron

Iron was initially selected as a COPC because both the maximum (76,000 mg/kg) and average (22,357 mg/kg) soil concentrations greatly exceeded the screening level based on toxicity to microorganisms (200 mg/kg) (Efroymson et al., 1997b). According to the Eco-SSL for iron (USEPA, 2003c), iron is essential for plant growth, and is generally considered to be a micronutrient. Because plants regulate its uptake, iron is not expected to be toxic to plants in well-aerated soils with pH levels between 5 and 8 S.U. (USEPA, 2003c). Although soil pH data have not been collected at LL-9, the average soil pH levels at LL 2, 3, and 4 were within the pH range of 5 to 8 S.U. (SAIC, 2003a, 2003b, and 2003c). Therefore, iron is not expected to be toxic to plants at this site. No toxicity data were available to evaluate risks to invertebrates from iron. However, because iron is generally considered a non-toxic metal, it is highly unlikely that soil invertebrates are being affected by iron at the site. For these reasons, risks to plants and invertebrates from iron are not likely and iron is not retained as a COPC.

Lead

Lead was initially selected as a COPC in soil because the maximum concentration of 1,330 mg/kg at LL9SS-011-0001-SO exceeded the ORNL PRG of 40.5 mg/kg (Efroymson et al., 1997a). Alternate benchmarks were used to further evaluate risks to soil receptors. The maximum concentration is less than the Eco-SSL invertebrate benchmark of 1,700 mg/kg (USEPA, 2005b). However, the maximum concentration is greater than the Eco-SSL plant benchmark of 120 mg/kg (USEPA, 2005b). The location of the maximum concentration (LL9SS-011-0001-SO) is a soil subfloor from a building where detonators were destroyed. One of the explosives destroyed in this building was lead azide, which would explain the high lead concentration. Four additional locations (LL9SS-003-0001-SO, LL9SS-006-0001-SO, LL9SS-024-0001-SO, and LL9SS-068-0001-SO) throughout LL9 have lead concentrations that are greater than the plant benchmark with concentrations ranging from 140 mg/kg to 320 mg/kg. Risks to invertebrates from lead are not anticipated, but risks to plants are possible. Therefore, lead was retained as a COPC for further evaluation.

Manganese

Manganese was selected as a COPC in soil because the maximum concentration of 3,800 mg/kg at LL9SS-027-0001-SO exceeded the ORNL screening level of 100 mg/kg based on toxicity to microorganisms (Efroymson et al., 1997b). Alternate benchmarks were used to further evaluate risks to soil receptors. No benchmark was available for earthworms, but the maximum and average concentrations exceed the ORNL plant benchmark of 500 mg/kg (Efroymson et al., 1997c). Of the 85 samples, only four samples had concentrations (2,000 mg/kg and 3,800 mg/kg) greater than the background concentration (1,450 mg/kg). Two



samples were located adjacent to sumps (LL9SS-040-0001-SO and LL9SS-041-0001-SO), and the other two were on the LL-9 boundary (LL9SS-027-0001-SO and LL9SS-031-0001-SO) (Figure 3-1). Because the location with the greatest manganese concentration is several hundred feet from the load line, it is likely that the elevated manganese concentrations greater than background are not related to site activities. Although risks to earthworms cannot be determined because of lack of toxicity data, because manganese does not appear to be site-related, it was not retained as a COPC.

Mercury

Mercury was initially selected as a COPC because the maximum soil concentration of 882 mg/kg at LL9SS-011-0001-SO exceeded the ORNL PRG of 0.00051 mg/kg, which is based on risks to the woodcock. Because this benchmark is based on risks to wildlife, alternate benchmarks were used to evaluate risks to soil receptors. The Canadian SQG of 12 mg/kg (EC, 1999c) was used to evaluate risks to earthworms and plants. The location of the maximum is a soil subfloor from a building where detonators were destroyed. One of the explosives destroyed in this building was mercury fulminate, which would explain the high mercury concentration. One sample (plus a duplicate) was taken approximately 50 ft from this location (LL9SS-034-0001-SO) (Figure 3-1). At this location, the average concentration of mercury is 14.5 mg/kg, which is greater than the Canadian SQG. All other sample concentrations were less than the SQG. Because of the extremely elevated mercury concentration at LL9SS-011-0001-SO and the possibility of contaminant migration, and because risks to plants and invertebrates are possible, mercury was retained as a COPC for further evaluation.

Selenium

Selenium was initially selected as a COPC because the maximum soil concentration of 1.8 mg/kg at LL9SS-068-0001-SO exceeded the ORNL PRG of 0.21 mg/kg (Efroymson et al., 1997a), which is based on risks to mice. Because this benchmark is based on risks to wildlife, alternate benchmarks were used to evaluate risks to soil receptors. The ORNL earthworm benchmark is 70 mg/kg (Efroymson et al., 1997b). The ORNL plant benchmark is 1 mg/kg (Efroymson et al., 1997c). All results were less than the earthworm benchmark. Three locations had concentrations that slightly exceeded the plant benchmark, but only the maximum concentration exceeded the background concentration (1.4 mg/kg). Therefore, most of the selenium concentrations are not likely to be site-related. The location of the maximum detection (LL9SS-068-0001-SO) is a wooded area where surface debris (glass, metal, and other rubbish) was found. Because the maximum concentration only slightly exceeded the background concentration, risks to plants and invertebrates are not likely and selenium was not retained as a COPC.



1 Zinc

2 Zinc was initially selected as a COPC because the maximum soil concentration of 780 mg/kg at
3 LL9SS-068-0001-SO exceeded the ORNL PRG of 8.5 mg/kg, which is based on dietary
4 toxicity to the woodcock. Because this benchmark is based on risks to wildlife, alternate
5 benchmarks were used for risks to soil receptors. The selected alternate benchmark for risks to
6 invertebrates and plants was the Canadian SQG of 200 mg/kg (EC, 1999f). The average zinc
7 concentration (87.6 mg/kg) is less than this benchmark, but a few locations across LL9 have
8 zinc concentrations that exceed the SQG. The two highest concentrations are at LL9SS-011-
9 0001-SO (711 mg/kg) and LL9SS-068-0001-SO (780 mg/kg). Sample LL9SS-011-0001-SO
10 was collected from the subfloor of a building where detonators were destroyed. The explosives
11 that were destroyed in this building included lead azide and mercury fulminate; but other metals,
12 such as zinc, may have been present in these explosives as well. Location LL9SS-068-0001-
13 SO (with a zinc concentration of 780 mg/kg) is a wooded area where surface debris (glass,
14 metal, and other rubbish) was found, so it is likely that the zinc is associated with this debris.
15 The three additional locations had zinc concentrations that slightly exceeded the SQG with
16 concentrations ranging from 210 mg/kg to 290 mg/kg. Although only 2 of the 85 samples had
17 zinc concentrations that significantly exceeded the SQG, because of the elevated zinc
18 concentrations at these locations, zinc was retained as a COPC for further evaluation.

19 Nitrocellulose

20 Nitrocellulose was initially selected as a COPC because a screening level was not available.
21 Although it is probable that nitrocellulose is related to the site, it is typically considered to be a
22 relatively inert compound, so it is unlikely to affect plants or invertebrates. Therefore, it was not
23 retained as a COPC for risks to plants or invertebrates.

24 Summary of Risks to Terrestrial Plants and Invertebrates

25 As presented above, a few sample locations had concentrations of metals that were greater
26 than invertebrate and/or plant benchmarks, indicating that risks are possible to these receptors
27 at those locations. The location with the greatest concentrations of several metals – copper,
28 lead, mercury, and zinc – was LL9SS-011-0001-SO. This was a subfloor sample collected in
29 a detonator destroying house where explosives, including lead azide and mercury fulminate,
30 were destroyed. An elevated, but much lower, mercury concentration was also found at sample
31 location LL9SS-034-0001-SO, which was about 50 ft from LL9SS-011-0001-SO; but the
32 copper, lead, and zinc concentrations were low at that location. It is not known whether the
33 detonator destroying house is the source of mercury at LL9SS-034-0001-SO. The extent of
34 the elevated levels of metals at LL9SS-011-0001-SO is not known because there are no
35 samples to the west and southwest to bound the contamination. Based on a review of the aerial
36 photo in Appendix C, however, the area surrounding LL9SS-034-0001-SO is heavily vegetated
37 and does not appear to be significantly affected by metals at this location.



LL9SS-068-0001-SO had concentrations of several metals that exceeded benchmarks for invertebrates and/or plants. This location was described as being in a wooded area where surface debris such as glass, metal, and other rubbish was present. Therefore, it is likely that the debris is the source of metals in the soil. However, because the location is in a wooded area, where vegetation is present, it is not likely that significant impacts on plants and invertebrates are occurring.

A few other locations across LL-9 had metals with concentrations that slightly exceeded benchmarks for invertebrates and/or plants, but these sporadic exceedances are not likely to significantly affect plants and invertebrates across the site. As can be seen from the aerial photograph and the site photographs in Appendix C, the site is well vegetated except in the areas where there is considerable vehicle traffic.

7.6.2 Risks to Benthic Invertebrates – Sediment

Potential risks to benthic invertebrates from exposure to COPCs were evaluated using the methodologies described in Section 7.5.2. Sediment samples were collected from sewer manholes at LL9SD-009-0001-SD and LL9SD-010-0001-SD. These sewers do not discharge in LL-9, so there is not a direct pathway to ecological receptors. Therefore, these sediment samples were not included in the risk assessment. The following subsections discuss whether the chemicals initially selected as COPCs should be retained for further evaluation of risks to benthic invertebrates.

Table 7-6 summarizes the results of potential direct toxicity of chemicals in sediment to benthic invertebrates. The table lists only the chemicals that were retained as COPCs because they were detected at concentrations that exceeded screening levels or they did not have screening levels. Table 7-6 presents the average chemical concentrations compared to the same screening levels that were used to select chemicals as COPCs because the COPC screening table (Table 7-2) only presented the comparison of the maximum concentrations to the screening levels. Table 7-6 also presents the maximum and average concentrations compared to the higher effects levels (i.e., the PEC, as discussed in Section 7.5.2) for benthic invertebrates to indicate the range of probability of effects. The following text presents the Step 3a evaluation for the chemicals listed in Table 7-6.

Bis(2-ethylhexyl)phthalate

Bis(2-ethylhexyl)phthalate was retained as a COPC because the maximum concentration of 660 µg/kg at LL9SD-015-0001-SD exceeded the screening level. An alternative benchmark was used to further evaluate risks to benthic invertebrates. The detected concentration is less than the AET of 1,300 µg/kg (Buchman, 1999). Bis(2-ethylhexyl)phthalate is also a common laboratory contaminant. Therefore, risks to benthic invertebrates are not likely and bis(2-ethylhexyl)phthalate was not retained as a COPC.



1 Dibenzo(a,h)anthracene

2 Dibenzo(a,h)anthracene was retained as a COPC because the maximum concentration of 43
3 $\mu\text{g/kg}$ at LL9SD-015-0001-SD exceeded the screening level. An alternate benchmark was
4 used to further evaluate risks to benthic invertebrates. The maximum (43 $\mu\text{g/kg}$) and average
5 (22.4 $\mu\text{g/kg}$) concentrations are less than the SEL of 1,989 $\mu\text{g/kg}$ (OMOE, 1993) which was
6 calculated using 1.53% total organic carbon (TOC) (average TOC at the site). Also, the
7 drainage ditches where sediment samples were collected are not wet year-round and are not
8 likely to support benthic invertebrates for most of the year. Therefore, risks to benthic
9 invertebrates are not likely and dibenzo(a,h)anthracene was not retained as a COPC.

10 Beryllium

11 Beryllium was initially selected as a COPC because a sediment screening level was not
12 available and the maximum concentration exceeded the RVAAP background concentration
13 (0.38 mg/kg) and the Ohio SRV (0.8 mg/kg). No toxicity benchmarks for benthic invertebrates
14 were available, but only two sample locations (LL9SD-001-0001-SD and LL9SD-012-0001-
15 SD) had beryllium concentrations that were slightly greater than the Ohio SRV, with
16 concentrations of 0.99 and 0.86 mg/kg, respectively. Because these concentrations only slightly
17 exceed the Ohio SRV, beryllium concentrations are unlikely related to site activities. In addition,
18 all sediment samples were collected from drainage ditches that are wet only during rain events
19 and are unlikely to support benthic organisms for most of the year. Therefore, risks from
20 beryllium to benthic invertebrates at LL-9 are not likely and beryllium was not retained as a
21 COPC.

22 Lead

23 Lead was initially selected as a COPC because the maximum concentration of 100 mg/kg at
24 LL9SD-013-0001-SD exceeded the Threshold Effects Concentration (TEC) of 35.8 mg/kg
25 (MacDonald et al., 2000) and the Ohio SRV of 47 mg/kg. An alternate screening level was used
26 to further evaluate risks to benthic invertebrates. The Probable Effects Concentration (PEC) for
27 lead is 128 mg/kg (MacDonald et al., 2000). Because the maximum concentration is less than
28 this benchmark, and because the drainage ditches where sediment samples were collected are
29 not likely to support benthic organisms for most of the year, risks from lead are not likely and
30 lead was not retained as a COPC.

31 Mercury

32 Mercury was initially selected as a COPC because the maximum concentration of 2.9 mg/kg at
33 LL9SD-012-0001-SD exceeded the TEC of 0.18 mg/kg (MacDonald et al., 2000) and the Ohio
34 SRV of 0.12 mg/kg. An alternate screening level was used to further evaluate risks to benthic
35 invertebrates. The PEC for mercury is 1.06 mg/kg (MacDonald et al., 2000). Because the
36 maximum concentration exceeded the higher effects benchmark, and because the



concentration is much greater than the background concentration and the Ohio SRV, risks to benthic invertebrates are possible at LL9SD-012-0001-SD, so mercury was retained as a COPC for further evaluation. Mercury was also detected at a concentration of 9.7 mg/kg in a nearby surface soil sample location (LL9SS-065-0001-SO), which may be the source of the mercury in the sediment. As discussed above, the drainage ditches are not likely to support benthic invertebrates for most of the year owing to lack of suitable habitat, so the significance of the possible risks to invertebrates is likely to be low.

Selenium

Selenium was initially selected as a COPC because a sediment screening level was not available, and the maximum detected concentration of 2.1 mg/kg at LL9SD-004-0001-SD exceeded the RVAAP background concentration (1.7 mg/kg) and the Ohio SRV (1.7 mg/kg). The AET for selenium is 1.0 mg/kg (Buchman, 1999). The maximum and average selenium concentrations exceed this benchmark, but the concentrations only slightly exceeded the background concentration and Ohio SRV. Therefore, selenium concentrations are unlikely to be site-related. Also, all of the sediment samples were collected from drainage ditches. These ditches are wet only during rain events, so it is unlikely that they can support benthic invertebrates for most of the year. Therefore, risks from selenium to benthic invertebrates at LL-9 are not likely and selenium was not retained as a COPC.

Nitrocellulose

Nitrocellulose was initially selected as a COPC because a screening level was not available. Although it is probable that nitrocellulose is related to the site, it is typically considered to be a relatively inert compound, so it is unlikely to affect benthic invertebrates. Also, because of the general lack of habitat for benthic invertebrates in the drainage ditches (as discussed above), risks from nitrocellulose to benthic invertebrates at LL-9 are not likely and nitrocellulose was not retained as a COPC.

Summary of Risks to Benthic Invertebrates

A few chemicals were initially selected as COPCs in the sediment because they were either detected at concentrations that exceeded screening levels or they did not have screening levels. Based on the Step 3a refinement, mercury, at one location, was the only chemical that may pose a risk to benthic invertebrates. However, as discussed above, the drainage ditches are not wet throughout the year, so benthic invertebrates would be present only at certain times. Therefore, the significance of the possible risks is likely to be low.

7.6.3 Risks to Aquatic Organisms – Surface Water

Potential risks to aquatic receptors from COPCs were further evaluated according to the methodologies described in Section 7.5.3. Surface water samples were collected from sumps



at LL9SW-007-0001-SW and LL9SW-008-0001-SW. These sumps have since been backfilled and are no longer present at the site, so they were not included in the risk assessment. Also, surface water samples were collected from sewer manholes at LL9SW-009-0001-SW and LL9SW-010-0001-SW. These sewers do not discharge in LL-9, so there is not a direct pathway to ecological receptors. Therefore, these surface water samples were not included in the risk assessment. The following subsections discuss whether the chemicals initially selected should be retained for further evaluation of risks to aquatic receptors.

Aluminum

Aluminum was initially selected as a COPC because the maximum surface water concentration of 17,800 µg/L at location LL9SW-005-0001-SW exceeded the AWQC screening level of 87 µg/L (USEPA, 2002) and the RVAAP background concentration (3,370 µg/L). The samples with high aluminum concentrations also had high or medium-high turbidity with water levels approximately 0.3 ft deep when the samples were collected. The actual impact on aquatic receptors at these locations is not known because of the high level of suspended solids (i.e., relatively low bioavailability) and the poor aquatic habitat in the ditches. Risks at most of the sample locations are not expected to be site-related because the concentrations are less than background. However, concentrations of aluminum at two of the locations above the AWQC and the RVAAP background criteria indicate possible risks to aquatic organisms present in the ditches. Therefore, aluminum was retained as a COPC for risks to aquatic organisms in surface water for further evaluation.

Copper

Copper was initially selected as a COPC because the maximum concentration of 20 µg/L at LL9SW-003-0001-SW exceeded the Ohio water quality criteria (WQC) of 4.76 µg/L (Ohio EPA, 2004) and the RVAAP background concentration (7.9 µg/L). The Ohio WQC is based on the minimum water hardness of 33.2 mg/L at LL9SW-001-0001-SW. The maximum hardness occurs at LL9SW-003-0001-SW. The Ohio WQC for LL9SW-003-0001-SW (using a hardness of 191.8 mg/L from this location) is 24.8 µg/L. The maximum copper concentration is less than this benchmark. However, the Ohio WQC for LL9SW-002-0001-SW (using a hardness of 37.1 mg/L from this location) is 5.28 µg/L. The concentration of copper at that location (5.8 µg/L) slightly exceeds this benchmark. Also, the Ohio WQC for LL9SW-005-0001-SW (using a hardness of 56.8 mg/L from this location) is 7.88 µg/L. The concentration of copper at that location (18.6 µg/L) exceeds this benchmark. Although the drainage ditches are unlikely to support aquatic organisms, because the copper concentration at LL9SW-005-0001-SW is greater than two times the Ohio WQC for the location, risks to aquatic receptors from copper are possible and copper was retained as a COPC for further evaluation.



Iron

Iron was initially selected as a COPC because the maximum concentration of 20,000 µg/L at location LL9SW-002-0001-SW exceeded the surface water screening level of 1,000 µg/L (USEPA, 2002) and the RVAAP background concentration (2,560 µg/L). The location of the maximum detection was described as being orange in color, which indicates high iron and mineral content. The other locations with concentrations greater than the screening level had high or medium-high turbidity, so the level of suspended solids is high (relatively low bioavailability). All of the surface water samples were collected from drainage ditches with little habitat for aquatic receptors. However, because four of the six samples contained iron concentrations above the screening level and site background, risks to aquatic organisms from iron are possible and iron was retained as a COPC for further evaluation.

Lead

Lead was initially selected as a COPC because the maximum concentration of 70 µg/L at LL9SW-003-0001-SW exceeded the Ohio WQC of 23.8 µg/L (Ohio EPA, 2004), which is based on the minimum water hardness (33.2 mg/L) at LL9SW-001-0001-SW. The maximum hardness is located at LL9SW-003-0001-SW. The Ohio WQC for lead at LL9SW-003-0001-SW (using a hardness of 191.8 mg/L from this location) is 222 µg/L. The maximum concentration is less than this benchmark. However, the Ohio WQC for LL9SW-005-0001-SW (using a hardness of 56.8 mg/L from this location) is 47.1 µg/L. The concentration of lead at this location (62.9 µg/L) is greater than the benchmark. Although the drainage ditches provide little habitat for aquatic receptors, because the concentration at LL9SW-005-0001-SW is greater than the benchmark, risks to aquatic receptors from lead are possible and lead was retained as a COPC for further evaluation.

Manganese

Manganese was initially selected as a COPC because a surface water screening level was not available and because the maximum surface water concentration of 1,400 µg/L at LL9SW-003-0001-SW exceeded the RVAAP background concentration of 391 µg/L.

The ORNL chronic benchmark (120 µg/L) was developed using the Tier II method described in the USEPA's 1993 Proposed Water Quality Guidance for the Great Lakes System (Suter and Tsao, 1996). Tier II values were developed so that aquatic benchmarks could be established with fewer data than are required for the USEPA water quality criteria. Tier II values are concentrations expected to be higher than WQC in no more than 20% of cases (Suter and Tsao, 1996). In the USEPA's 1986 Quality Criteria for Water (the Gold Book), it states that ions of manganese are found rarely at concentrations greater than 1 mg/L; and because the tolerance values reported range from 1.5 mg/L to over 1,000 mg/L, manganese is not considered to typically be a problem in fresh waters (USEPA, 1986). The ORNL benchmark is



likely lower than the tolerance values cited in the Gold Book because of the numerous uncertainty factors used to calculate the benchmark (Suter and Tsao, 1996).

All of the detections were less than the minimum tolerance value (1,500 µg/L) cited in USEPA (1986), so impacts to aquatic organisms are not expected. Also, as discussed previously in this SERA, there is very little aquatic habitat in the ditches, so there is unlikely to be a significant population of aquatic receptors. Therefore, risks to aquatic receptors from manganese at LL-9 are not likely and manganese was not retained as a COPC.

Mercury

Mercury was initially selected as a COPC because the maximum concentration of 1.7 µg/L at LL9SW-012-0001-SW exceeded the screening level of 1.4 µg/L (Ohio EPA, 2004). A duplicate sample was taken from the location of the maximum, and the average concentration at this location (1.0 µg/L) is less than the screening value. Also, the overall average mercury concentration for LL-9 (0.341 µg/L) is less than the screening level. As discussed above, there is very little aquatic habitat in the ditches where the surface water samples were collected, so there is unlikely to be a significant population of aquatic receptors. Therefore, risks to aquatic receptors from mercury at LL-9 are not likely and mercury was not retained as a COPC.

Zinc

Zinc was initially selected as a COPC because the maximum concentration of 228 µg/L at LL9SW-005-0001-SW exceeded the Ohio WQC of 46 µg/L (Ohio EPA, 2004) and the RVAAP site background concentration (42 µg/L). The Ohio WQC is based on the minimum hardness (33.2 mg/L) at LL9SW-001-0001-SW. The Ohio WQC were recalculated based on the water hardness at the two locations (LL9SW-005-0001-SW and LL9SW-003-0001-SW) with screening level and background concentration exceedances. The Ohio WQC at LL9SW-005-0001-SW (using a water hardness of 56.8 mg/L from this location) is 73 µg/L. The zinc concentration at this location (228 µg/L) exceeds the benchmark. The Ohio WQC at LL9SW-003-0001-SW (using a water hardness of 191.8 mg/L from this location) is 204 µg/L. The zinc concentration at this location (190 µg/L) is less than the benchmark. Although the drainage ditches are not likely to support aquatic organisms, risks from zinc are possible at LL9SW-003-0001-SW, so zinc was retained as a COPC for further evaluation.

Nitrocellulose

Nitrocellulose was initially selected as a COPC because no water quality criteria were available. Nitrocellulose is generally considered to be an inert compound. Also, as discussed above, there is very little aquatic habitat in the ditches where the surface water samples were collected, so there is unlikely to be a significant population of aquatic receptors. Therefore, risks to aquatic receptors from nitrocellulose at LL-9 are not likely and nitrocellulose was not retained as a COPC.



Summary of Risks for Aquatic Organisms

Several chemicals were initially selected as COPCs in the surface water because they were either detected at concentrations that exceeded screening levels or they did not have screening levels. Based on the Step 3a refinement, it was determined that aluminum, copper, iron, lead, and zinc, at some locations may cause a risk to aquatic organisms. There are several uncertainties in this determination, however. Most of the samples with the elevated metals concentrations were turbid. Filtered samples were not collected, so the concentrations of dissolved metals (which is the bioavailable fraction) are not known. Also, because the drainage ditches are not wet throughout the year, significant populations of aquatic organisms are not likely to be present. Therefore, the significance of the possible risks to aquatic organisms is likely to be low.

7.7 Risks to Terrestrial Wildlife

A food chain model was used to evaluate potential risks posed by COPCs to upper-level terrestrial wildlife receptors. Section 7.5.5 describes the food chain model methodology. Chemicals evaluated in the terrestrial food chain model were limited to those identified by the USEPA as bioaccumulative (USEPA, 2000a). Separate discussions are provided below for evaluations of potential risk to insectivorous/herbivorous and carnivorous receptors. The maximum (or 95% UCL) concentration detected in the surface soil and surface water samples is used as the EPC for the conservative food chain model. The average concentration detected in the surface soil and surface water samples is used as the EPC for the average food chain model. Appendix W presents the spreadsheets used to calculate the doses and HQs.

7.7.1 Risks to Insectivorous/Herbivorous Species

Table 7-7 presents the terrestrial wildlife model HQs based on conservative input parameters for terrestrial surrogate species (meadow vole, short-tailed shrew, and American robin).

For PAHs, the NOAEL HQs in the conservative models are less than 1.0. However, the HQs in the conservative scenario exceed 1.0 for the following analytes and receptors:

- Arsenic NOAEL for the vole and shrew;
- Chromium NOAEL for the robin;
- Lead NOAEL for the robin and shrew;
- Lead LOAEL for the robin; and
- Zinc NOAEL for the robin.



Table 7-7
Terrestrial Food Chain Model -
Conservative Scenario (Insectivorous, Herbivorous, and Carnivorous Receptors)

Chemical	Herbivorous Receptor EEQs		Insectivorous Receptors EEQs				Carnovorous Receptors EEQs					
	Meadow Vole		American Robin		Short-tailed Shrew		Barn Owl		Red Fox		Red-Tailed Hawk	
	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL
Semivolatile Organics												
ACENAPHTHENE	8.3E-04	8.3E-05	1.2E-03	1.2E-04	1.0E-03	1.0E-04	NA	NA	NA	NA	NA	NA
ACENAPHTHYLENE	3.5E-04	3.5E-05	5.4E-04	5.4E-05	5.3E-04	5.3E-05	NA	NA	NA	NA	NA	NA
ANTHRACENE	1.8E-03	1.8E-04	3.5E-03	3.5E-04	4.3E-03	4.3E-04	NA	NA	NA	NA	NA	NA
BENZO(A)ANTHRACENE	3.1E-03	3.1E-04	1.2E-02	1.2E-03	2.0E-02	2.0E-03	NA	NA	NA	NA	NA	NA
BENZO(A)PYRENE	2.5E-03	2.5E-04	1.2E-02	1.2E-03	2.0E-02	2.0E-03	NA	NA	NA	NA	NA	NA
BENZO(B)FLUORANTHENE	2.6E-03	2.6E-04	1.2E-02	1.2E-03	2.0E-02	2.0E-03	NA	NA	NA	NA	NA	NA
BENZO(G,H,I)PERYLENE	1.2E-03	1.2E-04	6.6E-03	6.6E-04	1.2E-02	1.2E-03	NA	NA	NA	NA	NA	NA
BENZO(K)FLUORANTHENE	1.6E-03	1.6E-04	7.3E-03	7.3E-04	1.3E-02	1.3E-03	NA	NA	NA	NA	NA	NA
CHRYSENE	3.4E-03	3.4E-04	1.3E-02	1.3E-03	2.1E-02	2.1E-03	NA	NA	NA	NA	NA	NA
DIBENZO(A,H)ANTHRACENE	1.2E-03	1.2E-04	6.1E-03	6.1E-04	1.1E-02	1.1E-03	NA	NA	NA	NA	NA	NA
FLUORANTHENE	7.6E-03	7.6E-04	2.1E-02	2.1E-03	3.1E-02	3.1E-03	NA	NA	NA	NA	NA	NA
FLUORENE	5.4E-04	5.4E-05	8.6E-04	8.6E-05	9.2E-04	9.2E-05	NA	NA	NA	NA	NA	NA
INDENO(1,2,3-CD)PYRENE	1.4E-03	1.4E-04	7.9E-03	7.9E-04	1.4E-02	1.4E-03	NA	NA	NA	NA	NA	NA
PHENANTHRENE	1.0E-02	1.0E-03	2.0E-02	2.0E-03	2.5E-02	2.5E-03	NA	NA	NA	NA	NA	NA
PYRENE	8.4E-03	8.4E-04	2.3E-02	2.3E-03	3.5E-02	3.5E-03	NA	NA	NA	NA	NA	NA
Inorganics												
ARSENIC	1.0E+00	1.0E-01	1.7E-01	1.7E-02	7.2E+00	7.2E-01	NA	NA	NA	NA	NA	NA
CADMIUM	3.9E-03	3.9E-04	7.0E-01	7.0E-02	4.0E-01	4.0E-02	NA	NA	NA	NA	NA	NA
CHROMIUM	2.4E-05	2.4E-06	2.5E+00	2.5E-01	4.1E-04	4.1E-05	NA	NA	NA	NA	NA	NA
COPPER	2.9E-02	2.9E-03	9.6E-02	9.6E-03	1.2E-01	1.2E-02	NA	NA	NA	NA	NA	NA
LEAD	2.8E-02	2.8E-03	4.5E+01	4.5E+00	2.5E+00	2.5E-01	NA	NA	NA	NA	NA	NA
MERCURY	1.2E-02	1.2E-03	2.8E-01	2.8E-02	3.9E-02	3.9E-03	5.3E-01	5.3E-02	1.8E-01	1.8E-02	4.7E-01	4.7E-02
NICKEL	2.6E-03	2.6E-04	4.6E-02	4.6E-03	3.8E-02	3.8E-03	NA	NA	NA	NA	NA	NA
SELENIUM	1.1E-02	1.1E-03	5.0E-01	5.0E-02	5.0E-01	5.0E-02	NA	NA	NA	NA	NA	NA
ZINC	3.3E-02	3.3E-03	7.7E+00	7.7E-01	2.5E-01	2.5E-02	NA	NA	NA	NA	NA	NA

Shaded cells indicate a NOAEL value > 1.0



Although the home ranges of the vole, robin, and shrew are less than 1 acre, average COPC concentrations are more realistic EPCs for most wildlife receptors than maximum concentrations because they are exposed to COPC concentrations throughout the load line, rather than at a single location. Eighty-five surface and shallow subsurface soil samples were collected at soil borings scattered throughout the 69-acre load line, so there is adequate spatial coverage within the exposure area making it appropriate to calculate an average concentration as the EPC.

Table 7-8 presents the HQs based on average exposure input parameters for the insectivorous and herbivorous species. Note that Table 7-8 lists only the chemicals that had NOAEL HQs greater than 1.0 using the maximum (or 95% UCL) concentrations. HQs in the average scenario are greater than 1 for the following analytes and receptors:

- Arsenic NOAEL for the shrew;
- Chromium NOAEL for the robin;
- Lead NOAEL for the robin and shrew;
- Lead LOAEL for the robin; and
- Zinc NOAEL for the robin.

Arsenic was detected in all 85 samples with a 95% UCL of 13.6 mg/kg and an average concentration of 12.3 mg/kg. Although arsenic concentrations in approximately 25 % of the samples (22/85) were greater than the RVAAP background concentration (15.4 mg/kg), most of those detected concentrations only slightly exceeded the background concentration, and the greatest concentration was only twice the background level. Therefore, arsenic may or may not be related to site activities. Also, because the average arsenic concentration that was used in the food chain model is less than background, the risks from arsenic are similar to background risks. For these reasons, it is not likely that potential site-related risks to small mammals are great enough to warrant further evaluation of arsenic in the BERA. Therefore, arsenic is eliminated as a COPC for risks to small mammals.

Chromium was detected in all 85 samples with a 95% UCL of 15.9 mg/kg and an average concentration of 15.4 mg/kg. Although chromium concentrations in approximately 12% of the samples (10/85) were greater than the RVAAP background concentration (17.4 mg/kg), most of the detected concentrations only slightly exceeded the background level. Therefore, chromium may or may not be related to site activities. The NOAEL estrogenic equivalent (EEQ) for the robin (2.4) is only slightly greater than 1.0. Also, because the average chromium concentration that was used in the food chain model is less than the background concentration, risks from chromium are similar to background risks.



Table 7-8
Terrestrial Food Chain Model -
Average Scenario (Insectivorous, Herbivorous, and Carnivorous Receptors)

Chemical	Herbivorous Receptor EEQs		Insectivorous Receptors EEQs				Carnovorous Receptors EEQs					
	Meadow Vole		American Robin		Short-tailed Shrew		Barn Owl		Red Fox		Red-Tailed Hawk	
	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL	NOAEL	LOAEL
Inorganics												
ARSENIC	9.3E-01	9.3E-02	1.6E-01	1.6E-02	6.5E+00	6.5E-01	NA	NA	NA	NA	NA	NA
CHROMIUM	2.2E-05	2.2E-06	2.4E+00	2.4E-01	4.0E-04	4.0E-05	NA	NA	NA	NA	NA	NA
LEAD	2.7E-02	2.7E-03	4.5E+01	4.5E+00	2.5E+00	2.5E-01	NA	NA	NA	NA	NA	NA
ZINC	3.2E-02	3.2E-03	7.6E+00	7.6E-01	2.5E-01	2.5E-02	NA	NA	NA	NA	NA	NA

Shaded cells indicate a NOAEL value > 1.0

NA = No data available



For these reasons, it is unlikely that potential site-related risks to birds are great enough to warrant further evaluation of chromium in the BERA. Therefore, chromium is eliminated as a COPC for risks to birds.

Lead was detected in all 85 samples with a 95% UCL of 40.2 mg/kg and an average concentration of 46.2 mg/kg. Because the average concentration is greater than the 95% UCL, the 95% UCL was used for the average scenario food chain model. The lead NOAEL and LOAEL-based EEQs exceeded 1.0 in the average exposure scenario food chain model for the robin and the NOAEL-based EEQ exceeded 1.0 in the food chain model for the shrew. Lead concentrations in approximately 45% of the samples (38/85) were greater than the RVAAP background concentration (19.1 mg/kg). Many of the detected concentrations greatly exceeded the background concentration. As can be seen in the food chain model calculation spreadsheets in Appendix W, the dose from soil invertebrates accounts for a large portion of the total calculated dose. The estimated earthworm concentration (80.4 mg/kg) was caused by the BAF of 2.0 that was used in the food chain model. USEPA (2005c) presents a soil to earthworm uptake equation for estimating lead concentrations in earthworms from soil. Using the equation from USEPA (2005c), the estimated earthworm concentration in the robin and shrew food chain models is 2.53 mg/kg. Using the earthworm concentration as estimated from the equation in USEPA (2005c), the NOAEL-based EEQ under the average exposure scenario in the robin food chain model is 3.5 and the LOAEL-based EEQ is less than 1.0. Similarly, if the earthworm concentration is calculated using the equation from USEPA (2005c) in the shrew food chain model, both NOAEL and LOAEL-based EEQs are less than 1.0. Because only the NOAEL-based EEQ exceeds 1.0 in the robin food chain model using the estimated earthworm concentration as calculated in USEPA (2005c), the risks to the robin overall are expected to be low. Therefore, lead was eliminated as a COPC for risks to small mammals and birds.

Zinc was detected in all 85 samples with a 95% UCL of 89.3 mg/kg and an average concentration of 87.6 mg/kg. Zinc concentrations in approximately 58% of the samples (49/85) were greater than the RVAAP background concentration (61.8 mg/kg). Similar to lead presented above, the estimated earthworm concentration (158 mg/kg) was caused by the BAF of 1.8 that was used in the food chain model. USEPA (2005c) also presents a soil to earthworm uptake equation for estimating zinc concentrations in earthworms from soil. Using the equation from USEPA (2005c), the estimated earthworm concentration in the robin food chain models is 59.3 mg/kg. Using the earthworm concentration as estimated from the equation in USEPA (2005c), the NOAEL-based EEQ under the average exposure scenario in the robin food chain model is 3.5 and the LOAEL-based EEQ is less than 1.0. Most zinc concentrations were similar to the background concentration with some exceptions. Most notably, two locations (LL9SS-011-0001-SO and LL9SS-068-0001-SO) had concentrations (711 mg/kg and 780 mg/kg, respectively) greatly exceeding background; but the area represented by these few samples is small. Because only the NOAEL-based EEQ exceeds 1.0 in the robin food chain model using the estimated earthworm concentration as calculated in USEPA (2005c), the risks to the robin overall are expected to be low. Therefore, zinc is eliminated as a COPC for risks to birds.



In summary, using average exposure concentrations and more recent BAFs (for lead and zinc), the HQs based on the NOAEL exceeded 1.0 for several chemicals and receptors, but no HQs based on the LOAEL exceeded 1.0. When the additional following factors are considered, it is unlikely that wildlife receptors will be affected and no chemicals were retained as COPCs for further evaluation of risks to small mammals or birds:

- The chemicals in soil will likely be less bioavailable than the form of the chemicals used to conduct the toxicity test to establish the NOAELs, so the risks are overestimated.
- The average concentrations for most metals used in the food chain model were similar to background concentrations, so potential risks would be similar to background risks.

7.7.2 Risks to Carnivorous Species

Table 7-7 presents the carnivorous wildlife model HQs based on conservative input parameters for surrogate species (barn owl, red fox, and red-tailed hawk). Note that only PBT COPCs were carried through the carnivorous wildlife food chain model in accordance with the Ohio EPA guidance (Ohio EPA, 2003). No chemicals had conservative scenario HQs greater than 1.0. Therefore, no chemicals were retained as COPCs for further evaluation regarding risks to carnivorous wildlife.

7.8 Ecological Risk Uncertainty Analysis

This section discusses the uncertainties associated with this SERA at LL-9. A significant uncertainty is the fact that the nature and extent of contamination in the environmental media has not been completely determined. Therefore the results of the ecological risk assessment presented in Section 7 (and the associated uncertainties) are preliminary and subject to change based on the data collected to resolve the RI data gaps for LL-9.

7.8.1 Measurement and Assessment Endpoints

Measurement endpoints are used to evaluate the assessment endpoints that are selected for the SERA. Measures of effects are not the same as the assessment endpoints. The measures are used to predict effects to the assessment endpoints by selecting surrogate species that will be evaluated. For example, the shrew was chosen as a surrogate species to predict effects to the small mammal population. A decrease in reproduction of a shrew is used to assess a decrease in reproduction of the small mammal population. However, because of differences in ingestion rates, toxicity, food preferences, etc. among different species, predicting a decrease in reproduction of a shrew may either under- or over-protect the small mammal population as a whole. Although reptiles and amphibians are likely to be present on the site, risks are not quantitatively evaluated because exposure factors are not established for most species and toxicity data are very limited. Using aquatic organisms as a surrogate species, risks to amphibians exposed to the surface water and sediment are expected to be low based on the Step 3a evaluations.



7.8.2 Exposure Characterization

The contaminant dose to terrestrial wildlife is calculated using an equation that incorporates ingestion rates, body weights, bioaccumulation factors, and other exposure factors. These exposure factors are obtained from literature or are predicted using various equations. Ingestion rates and body weights vary among species, especially among species inhabiting different areas.

Bioaccumulation of contaminants into various biological media (i.e., plants, invertebrates, small mammals) depends on characteristics of the media (soil, sediment, and surface water) such as pH, organic carbon, etc. Therefore, actual bioaccumulation factors at the site may differ from those used in the SERA that were obtained from the guidance. For example, in the case of lead and zinc in the shrew and robin food chain models, the estimated earthworm concentrations were significantly greater when calculated using BAFs from the RVAAP guidance than when the earthworm concentrations were calculated using BAFs from the more recent USEPA guidance (2005c).

Also, the bioavailability of the chemicals is not taken into account in this SERA. All the chemicals are assumed to be 100% bioavailable at the detected concentrations, which is unlikely to occur for contaminants in the environment. For example, surface water samples with elevated concentrations (with respect to the AWQC and background criteria) were noted on the field forms as having “medium-high turbidity” and “high turbidity.” The elevated concentrations reported in these samples are potentially attributable to the suspended solids of these samples and not actual water concentrations. However, the suspended solids do not represent the bioavailable portion of the contaminant in the water column. Typically, uncertainties regarding the bioavailability of contaminants in surface water can be reduced by analyzing dissolved concentrations.

There is uncertainty in the chemical data that are collected at the site. Measured levels of chemicals are only estimates of the true site chemical concentrations. For samples that are deliberately biased toward known or suspected high concentrations, predicted doses probably will be higher than actual doses.

Finally, under the food chain model exposure scenario, terrestrial wildlife are assumed to live and feed only at the site. These assumptions will tend to over predict risk because it is unlikely that most receptors will obtain their food from within the site boundaries and from the most contaminated areas.

7.8.3 Ecological Effects Data

There is uncertainty in comparing the ecological toxicity value to the EPC to determine risk to a receptor. The water quality criteria developed by USEPA in theory protects 95% of the exposed species. Therefore, some sensitive species may be present that are not protected by the use of



these criteria. There also may be situations where the surface water screening levels (SWSLs) are over-predictive of risk if the sensitive species used to develop the criteria do not inhabit the site. Finally, with the exception of hardness for a few metals, SWSLs do not account for site-specific factors, such as TOC or pH that may affect toxicity.

Potential adverse impacts on aquatic receptors from constituents in the sediment are evaluated by comparing the COPC concentration to sediment screening level (SdSLs). SdSLs have more uncertainty associated with them than do SWSLs. The procedures for developing SdSLs are not as well established as the procedures for developing SWSLs. As a result, the sediment screening levels have been developed using different methodologies. Additionally, there are fewer sediment toxicity data than surface water toxicity data. Sediment characteristics (i.e., pH, acid volatile sulfides, and total organic carbon) also will have a large impact on the bioavailability and toxicity of constituents.

Potentially adverse impacts on terrestrial plants and invertebrates from constituents in the surface soil are evaluated by comparing the COPC concentration to shallow soil screening level (SSSLs). SSSLs are similar to the sediment screening levels in that they are less established than SWSLs. Fewer studies and fewer data are available for establishing SSSLs than SdSLs, and many SSSLs are based on the results of only a few studies. In addition, SSSLs are based on different endpoints, depending on the preference of the agency that developed them. Therefore, they have more uncertainty than surface water and sediment screening values.

Ecological screening levels are not available for nitrocellulose detected in LL-9 media. According to the USEPA Office of Drinking Water, nitrocellulose is essentially nontoxic (USEPA, 2004d).

Several soil samples were collected and sent to GPL for explosives analyses. The results from these analyses are not useable for risk assessment purposes and are not included in the quantitative risk assessment because a screening method was used. The results of this screening evaluation are shown in Table 7-9. Six explosives were detected in the soil at LL-9. Three of them (2,4,6-trinitrotoluene, 4-amino-2,6-dinitrotoluene, and RDX) were detected infrequently (1 of 33 samples). 2,4-Dinitrotoluene only exceeded the screening value of 1,280 µg/kg in 1 of 33 samples. A screening value was not available for 3-nitrotoluene, but all results were below the screening value for 2,4-dinitrotoluene. For 2,6-dinitrotoluene, 4 of 33 samples exceeded the screening value of 32.83. Because these data were only used for screening purposes, risks were not assessed.

The NOAELs and LOAELs that were selected for the wildlife endpoint species were based on other than the endpoint species (i.e., rats, mice, ducks). There is uncertainty in the application of toxicity data across species because the contaminant may be more or less toxic to the endpoint species than it was to the test study species.



Table 7-9
GPL Explosives Data Shallow Soil Samples (0-4 ft bgs)

Parameter	Frequency of Detection	Minimum Concentration	Maximum Concentration	Mean Concentration	Average of Positive Detects	Sample of Maximum Detect	Screening Level	
							Value	Source
Energetics (ug/kg)								
2,4,6-TRINITROTOLUENE	1/33	200	200	54.5	200	LL9SS-030-0001-SO-GPL	NA	NA
2,4-DINITROTOLUENE	3/33	120	3700	170	1370	LL9SS-002-0001-SO-GPL	1280	a
2,6-DINITROTOLUENE	4/33	340	3100	188	1190	LL9SB-002-0001-SO-GPL	32.83	a
3-NITROTOLUENE	4/33	90 J	410	115	220	LL9SS-002-0001-SO-GPL	NA	NA
4-AMINO-2,6-DINITROTOLUENE	1/33	120	120	52.1	120	LL9SS-004-0001-SO-GPL	NA	NA
RDX	1/33	220	220	104	220	LL9SS-027-0001-SO-GPL	NA	NA

Source of Screening Level:

a - Ecological Data Quality Level (USEPA Region 5, 1999)



The toxicity of chemical mixtures is not well understood. The toxicity information used in the SERA for evaluating risk to the ecological receptors is for individual chemicals. Chemical mixtures can affect the organisms very differently than the individual chemicals because of synergistic or antagonistic effects.

Toxicological data for a few of the COPCs are limited or do not exist. Therefore, there is uncertainty in any conclusions involving the potential impacts on ecological receptors from these constituents. However, attempts were made to reduce the uncertainties with these COPCs using the other Step 3a factors discussed in Section 7.5.4.

Several alternative benchmark values were used to gain a better understanding of the relationship between the maximum concentrations of the selected COPCs to the overall ecological risks of the site. There is some uncertainty involved when using these alternative benchmarks. The Canadian SQGs, which are used as alternative benchmarks for both plants and invertebrates, are based on effects on either plants or invertebrates and, thus, differentiation of risk to plants versus risk to invertebrates cannot be made using the Canadian guidelines. The ORNL values are separated into guidelines for plants and guidelines for invertebrates. However, the values are limited to only a few chemicals.

7.8.4 Risk Characterization

Risks are possible if an HQ is greater than or equal to unity, regardless of the magnitude of the HQ. However, the magnitude of effects on ecological receptors cannot be inferred based on the magnitude of the HQ. Rather, an HQ greater than 1.0 simply indicates that the dose used to derive the toxicity reference value was exceeded. Finally, there is uncertainty in how the predicted risks to a species at the site translate into risk to the population in the area as a whole.

Several chemicals were selected as COPCs in surface water. There are uncertainties in evaluating these chemicals because most of the drainage ditches at LL9 are not perennial and are wet only seasonally, depending on precipitation and temperature. However, to be conservative, these areas were evaluated as if aquatic receptors were present or could be present for part of the year. Before proceeding further into a BERA based on potential risks to aquatic receptors, a more detailed evaluation of the aquatic habitats will need to be conducted to determine whether sufficient aquatic habitat is present to support aquatic receptor populations.

7.9 Conclusions

Based on the SERA and the first step of the BERA (Step 3a), the following conclusions were drawn.



7.9.1 Terrestrial Plants and Soil Invertebrates

After the initial screening (Table 7-1), 12 metals (aluminum, arsenic, chromium, copper, iron, lead, manganese, mercury, nickel, selenium, vanadium, and zinc), 3 energetics (nitroguanidine, RDX, and nitrocellulose), and 1 SVOC (dibenzofuran) were detected at concentrations that exceeded RVAAP background, and exceeded soil screening levels or did not have soil screening levels. Other chemicals (i.e., PAHs and cadmium) were retained as COPCs only because they were bioaccumulative. These chemicals were not included in the evaluation of risks to plants and invertebrates because they were not detected at concentrations that exceeded soil screening levels.

Tables 7-4 and 7-5 summarize the results of potential direct toxicity of chemicals in soil to earthworms and plants, respectively. Chromium, copper, mercury, and zinc had maximum concentrations that were greater than the earthworm toxicity benchmarks; the average chemical concentrations for these chemicals were less than the benchmarks. Iron, manganese, dibenzofuran, nitroguanidine, RDX, and nitrocellulose did not have earthworm toxicity benchmarks. Aluminum toxicity to earthworms is based on the pH of the soil. Arsenic, chromium, copper, lead, manganese, mercury, selenium, and zinc had maximum concentrations that were greater than the plant toxicity benchmarks; the average chemical concentration for manganese was also greater than the plant benchmark. Dibenzofuran, nitroguanidine, RDX, and nitrocellulose did not have plant toxicity benchmarks. Aluminum and iron toxicity to plants are based on the pH of the soil. Based on the Step 3a factors discussed in Section 7.6.1, all analytes except copper, lead, mercury, and zinc were eliminated as COPCs. The other analytes were eliminated for several reasons, including concentrations below plant and invertebrate benchmarks and background concentrations, low frequencies of detection, and relatively low toxicity of the analytes. Copper, lead, mercury, and zinc are retained as COPCs at LL9SS-011-0001-SO, which had elevated concentrations of these analytes. This was a subfloor sample collected in a detonator destroying house where explosives including lead azide and mercury fulminate were destroyed. It is difficult to determine whether contamination is confined to this area owing to a lack of samples collected around this location. A few other locations across LL9 had metals with concentrations that exceeded benchmarks for invertebrates and/or plants. However, they are not likely to affect invertebrates and plants across the site because the site is well vegetated except where there is heavy vehicle traffic. Therefore, the only significant potential risks that are expected from metals at LL-9 are at LL9SS-011-0001-SO.

7.9.2 Benthic Invertebrates

After the initial screening (Table 7-2), bis(2-ethylhexyl)phthalate, dibenzo(a,h)anthracene, nitrocellulose, and four metals (beryllium, lead, mercury, and selenium) were detected at concentrations that exceeded Ohio SRVs and sediment screening levels or screening levels were not available.



Table 7-6 summarizes the results of potential direct toxicity of chemicals in sediment to benthic invertebrates and includes the average chemical concentrations compared to the same screening levels and the maximum and average concentrations compared to the higher effects levels. Based on the Step 3a factors discussed in Section 7.6.2, mercury, at one location, was the only chemical that may pose a risk to benthic invertebrates. However, because the drainage ditches are not wet year-round, they would not likely support benthic invertebrates for much of the year. Therefore, risks to benthic invertebrates from mercury are likely to be low, and it does not appear that the site needs to proceed further in the BERA because of mercury in sediment.

7.9.3 Aquatic Organisms

After the initial screening (Table 7-3), aluminum, copper, iron, lead, manganese, mercury, zinc, and nitrocellulose were selected as COPCs because maximum concentrations exceeded the screening levels or screening levels were not available. Based on the Step 3a factors discussed in Section 7.6.3, aluminum, copper, iron, lead, and zinc were retained as COPCs because of possible risks to aquatic organisms. However, there is significant uncertainty in the likelihood of potential impacts to aquatic organisms from the above-listed metals. Most of the samples with elevated metals concentrations were turbid, so the metals may be attributable to a high level of suspended solids during the sample collection and may not be dissolved in the surface water. For aquatic organisms, dissolved metals are the bioavailable and potentially toxic fraction of metals in surface water. More significant, however, is the fact that the drainage ditches are intermittent and typically wet only during and after rain events. Therefore, they likely do not support aquatic organisms for much of the year. These factors should be taken into consideration when risk management decisions for LL-9 are made. Because of these factors, it does not appear that the site needs to proceed further in the BERA because of metals in surface water.

7.9.4 Terrestrial Wildlife

The potential risks to mammals and birds associated with the COPCs in the surface soil, surface water, or both were further evaluated to determine whether site-related risks from the chemicals were unlikely or the risks were great enough to warrant retaining the chemicals as COPCs and proceeding further into the BERA.

The conservative assumption of total bioavailability, the use of average exposure concentrations (which are similar to background concentrations), and the use of current BAFs (for lead and zinc) from USEPA were used to conclude that impacts on insectivorous and herbivorous receptors are unlikely. In summary, no chemicals were retained as COPCs for further evaluation of risk to small mammals or birds.



- 1 Based on the conservative scenario and the NOAEL, no carnivorous animals had HQs greater
- 2 than 1.0 for any carnivorous receptor. Consequently, no chemicals were retained as COPCs for
- 3 further evaluation of risks to upper-level carnivorous mammals and birds.



8.0 SUMMARY AND CONCLUSIONS

This section briefly summarizes the existing LL-9 conditions that were found during the RI, the possible fate and transport of contaminants found at the site, and the risk assessments tasks that were completed. The conclusions are based on the outcome of the BHHRA and SERA.

8.1 Summary

The following subsections summarize the existing conditions at LL-9.

8.2 Nature and Extent of Contamination

The concentrations of contaminants are generally low, with a notable exception being a localized spot of high metal concentrations for copper, mercury, lead, and zinc at LL9SS-011-0001-SO in surface soil.

Sumps might have been a primary contamination source from operations water that was routinely diverted to them. However, sumps were removed during the LL-9 demolition and removal of the load line structures. Sewer lines that might have provided a contaminant transport pathway have been plugged. Thus, contaminant transport has been eliminated in these conveyances.

Contamination at LL-9 is sparsely dispersed across the site with some elevated chemical concentrations occurring near operational activities. Most metals appear to be fairly uniformly distributed across the site, with exception of location LL9SS-011-0001-SO noted above.

Metals do not degrade in the environment, but they can be assimilated into minerals. Their oxidation states can change over time as they migrate from one location to another. Metals in general, however, are relatively immobile except for those with predominantly single positive charges, such as sodium and potassium. A few other metals such as calcium and magnesium are also relatively mobile. These four metals, however, generally pose little to no environmental risks to receptors. Receptor exposure and risk are treated more completely in the risk assessment sections (6.0 and 7.0).

Few organic chemicals that are directly related to site operations were selected as COPCs, thus indicating that the concentrations of detected organic chemicals are generally low enough not to pose a significant threat to the health of receptors. However, nature and extent have not been fully determined during this Phase I RI. Additional sampling and analysis may be required to delineate the full lateral and vertical extent of contamination at LL-9.



8.3 Contaminant Fate and Transport

The primary contaminant migration pathways for contaminants at LL-9 are:

- Leaching from soils to groundwater;
- Leaching from sediments to surface water; and
- Transport in surface drainage channels.

Leaching from soils would be expected to occur around sumps if contaminated soil remained in place after the sump removals. This is not expected to be a significant concern. The quantities of contamination present at LL-9 are generally low, and sump removal is completed. Therefore, the mass of contaminants transported by this migration pathway is low. Dilution effects should be large in drainage channels. This is reflected in the low concentrations of contaminants detected in surface waters. Similar effects are observed in groundwater. Nitrocellulose, which has evidently been widely distributed in surface and subsurface soils at low concentrations across LL-9, is notably recalcitrant to dissolution; hence, its persistence is expected to be long.

Any soil or sediment contaminants that leach into groundwater would flow radially away from LL-9. Each contaminant will be retarded in its movement. The degree of retardation depends largely on the adsorption tendency of each contaminant and the amount of organic material in the soils and bedrock.

Surface water drainage channels can transport surface soil contaminants and sediments to downgradient locations. However, the topographical relief is moderate at LL-9 except at the peripheries. Therefore, transport pathways are not expected to move sediments rapidly. An exception to this could be during storm events when flow rates increase significantly. Contaminants detected at LL-9 will tend to adsorb to sediments.

In summary, the low concentrations and small total masses of energetic compounds at LL-9 are consistent with good health and safety practices. In particular, primary explosives such as metal azides (e.g., lead azide) and fulminates (e.g., mercury fulminate), which are very unstable with respect to physical shock, would not be expected to be released indiscriminately to non-operational areas. To do so would result in extreme safety hazards. The small reservoir of contamination in soil provides little total contaminant mass for migration to groundwater or other media. The metals will persist and the organics will degrade over time, albeit some of them (e.g., nitrocellulose) will do so slowly. Given the low concentrations observed at this site, modeling to estimate the degradation rates would not be fruitful.



8.4 Baseline Human Health Risk Assessment

A BHHRA was performed to assess the potential current and future risks associated with human exposure to site-related contaminants found in LL-9. The BHHRA included:

- Data evaluation: selecting COPCs;
- Exposure assessment: identifying potential receptors, evaluating potential exposure pathways, and estimating the chemical intake resulting from exposure;
- Toxicity assessment: evaluating the toxicity of each COPC and summarizing the toxicity criteria that were used;
- Risk characterization: estimating the cancer and noncarcinogenic risk(s); and
- Uncertainty analysis: identifying significant uncertainties that could affect the BHHRA results.

The COPCs selected for quantitative human health risk assessment are summarized in Table 8-1.

Table 8-1

Summary of COPCs Selected for the Quantitative Human Health Risk Assessment

Soils	Groundwater	Surface Water	Sediment
Aluminum	Antimony	Aluminum	Aluminum
Arsenic	Manganese	Arsenic	Mercury
Chromium		Chromium	Vanadium
Manganese		Lead	
Mercury		Manganese	
Benzo(a)pyrene		Mercury	
Dibenzo(a,h)anthracene		Vanadium	

The receptors evaluated were a security guard/maintenance worker, a National Guard trainee, a National Guard resident trainer, a hunter/trapper, and hypothetical future farmer residents (adult and child). The risk assessment calculations are summarized in Table 8-2.



Table 8-2

Human Health Risk Assessment Calculation Summary

Receptor	Non-Carcinogenic	Incremental Lifetime Cancer Risks
Security guard/maintenance worker	HI <1 (no adverse non-carcinogenic health effects)	Within USEPA target risk range of 1E-04 to 1E-06; does not exceed Ohio EPA's target of 1E-05.
Hunter/trapper	HI <1 (no adverse non-carcinogenic health effects)	<1E-06; Does not exceed Ohio EPA's target risk value or USEPA target risk range.
National Guard resident trainer	HI >1 if it is assumed the receptor is using the shallow groundwater for domestic purposes and exposed to surface water. Manganese in groundwater is the risk driver.	Within USEPA target risk range of 1E-04 to 1E-06; exceeds Ohio EPA's target of 1E-05. Arsenic in soil is the major contributor to the risk.
National Guard trainee	Total HI >1 because of inhalation of manganese in deep surface soil. Manganese at background would result in an HI of 4, and at its Region 9 residential PRG would result in an HI of 5.	Within USEPA target risk range of 1E-04 to 1E-06; exceeds Ohio EPA's target of 1E-05 because of inhalation of arsenic and chromium in deep surface soil.
Adult resident farmer	HI >1 only if it is assumed the receptor is using the shallow groundwater for domestic purposes and if the ingestion of foodstuffs pathways is evaluated. Manganese is the primary risk driver in groundwater. See the uncertainty discussion.	Not applicable
Child resident farmer	HI >1. However, target-organ-specific HI > 1 for direct exposure if it is assumed that groundwater is used for domestic purposes and HI > 1 for arsenic's target organs when direct exposure to media other than groundwater is evaluated. Only true if HIs for surface soil and subsurface soil are combined. For indirect exposure via the ingestion of foodstuffs pathways, HI > 1 for ingestion of milk and vegetables. See the uncertainty discussion.	Not applicable



Table 8-2 (Continued)

Human Health Risk Assessment Calculation Summary

Receptor	Non-Carcinogenic	Incremental Lifetime Cancer Risks
Lifelong resident farmer	Not applicable	For direct exposure pathways, within USEPA target risk range of 1E-04 to 1E-06; exceeds Ohio EPA's target of 1E-05. Arsenic in soils is the major contributor to the risk. For indirect exposure via the ingestion of foodstuffs pathways, exceeds target risk range for ingestion of beef, milk, and vegetables.

Several significant uncertainties associated with the risk assessment were identified, as outlined in Section 6.6 and should be considered when making any risk management decisions. In many cases, conservative exposure and toxicity assumptions were applied, which may have resulted in conservative conclusions about potential risks.

Maximum detected concentrations of surface soil and deep surface soil COPCs copper (1,240 mg/kg), lead (1,330 mg/kg), and mercury (882 mg/kg) were at azide sample location LL9SS-011-0001-SO. The next-highest detected concentrations were much lower (copper [170 mg/kg], lead [320 mg/kg], and mercury [17 mg/kg]). This sample location may be of particular concern when making risk decisions, particularly for exposure to mercury.

8.5 Screening Ecological Risk Assessment

A SERA was performed to assess whether adverse ecological impacts are present as a result of site-related contaminants found in LL-9. The SERA included completing Steps 1 through 3a of the eight steps that comprise a Baseline Ecological Risk Assessment. The ecological risk assessment steps are stipulated in U.S. EPA, Ohio EPA, and RVAAP FWERWP (USACE, 2003). The steps that were completed as part of the LL-9 SERA are described below:

- Formulate the preliminary problem by identifying potential receptor groups and complete exposure pathways.
- Compare the contaminant concentrations for the chemicals found in LL 9 surface water, sediment, and surface soil to ecological screening values (ESVs) to initially select COPCs. The ESVs used in this risk assessment were stipulated by Ohio EPA and RVAAP guidance.



- Refine the list of COPCs by comparing exposure point concentrations to toxicity benchmarks established for specific receptor groups, by conducting food chain modeling for mammals and birds, and by evaluating other factors such as presence of suitable habitat, frequency of detection, and spatial distribution.

Ecological impact was evaluated for plants, soil, sediment invertebrates, aquatic organisms, and terrestrial wildlife. Three types of mammals and birds were evaluated: insectivores, herbivores, and carnivores.

The ecological risk calculations are shown in table 8-3.

Table 8-3

Ecological Risk Calculations

Type of Species	Screening Results	Notes
Terrestrial plants and soil invertebrates	Copper, lead, mercury, and zinc were retained as COPCs.	Copper, lead, mercury, and zinc could pose possible risks to terrestrial plants at LL9SS-011-0001-SO. Copper, mercury, and zinc could pose possible risks to soil invertebrates at LL9SS-011-0001-SO. This location was a subfloor sample collected in a detonator destroying house where explosives, including lead azide and mercury fulminate, were destroyed. The contamination needs to be better bounded to determine the extent of the elevated levels of metals. A few other locations across LL-9 had metals with concentrations that exceeded benchmarks for invertebrates and/or plants. However, they are not likely to affect invertebrates and plants across the site because the site is well vegetated except where there is heavy vehicle traffic.
Benthic Invertebrates	Mercury was retained as a COPC.	Mercury, at one location, was the only chemical that may pose a risk to benthic invertebrates. The drainage ditches do not likely support a significant population of benthic invertebrates. Therefore, risks to benthic invertebrates from mercury are likely to be low, and it does not appear that the site needs to proceed further in the BERA because of mercury in sediment.



Table 8-3 (Continued)

Ecological Risk Calculations

Type of Species	Screening Results	Notes
Aquatic Organisms	Aluminum, copper, iron, lead, and zinc were retained as COPCs.	Elevated concentrations are likely the result of high levels of suspended solids in the surface water samples. Also, the drainage ditches at LL-9 provide poor ecological habitat. These factors should be taken into consideration when risk management decisions for LL-9 are made. Based on these factors, it does not appear that the site needs to proceed further in the BERA because of metals in surface water.
Terrestrial Wildlife— Insectivores/Herbivores	No COPCs retained.	Because of the conservative assumption of total bioavailability, no HQs based on the LOAEL were greater than 1.0, and because average concentrations for most metals used in the food chain model were similar to background concentrations, potential risks are not likely.
Terrestrial Wildlife— Carnivores	No COPCs retained.	Because no HQs based on the NOAEL were greater than 1.0 using conservative chemical concentrations, potential risks are not likely.



9.0 RECOMMENDATIONS

The conclusions of the risk assessments are based on available data. If data collected in the Phase II RI alters the conceptual site model presented in this Phase I RI, then the risk assessment findings presented in this report will require revision. Although the BHHRA indicated potential risk to a National Guard trainee due to exposure to arsenic, chromium, and manganese, conservative toxicity and exposure values (such as PEF) were used to calculate the potential risk. Thus, the risk estimates presented for those chemicals likely overestimate the potential for non-carcinogenic risk. Additionally, the concentrations detected in the environmental media may represent background conditions. Manganese at background would result in an HI of 4, and the Region 9 residential PRG for manganese would result in an HI of 5. Arsenic at background would result in a risk of $5E-06$. Chromium at background would result in a risk of $1.1E-05$, and at the Region 9 residential PRG for chromium would result in a risk of $1.8E-05$. It is recommended that the risk management team consider the need for any further remedial action based on the risk assessment results presented for this receptor.

Concentrations of target analytes (primarily metals and propellants) were detected in surface soil, subsurface soil, and sediment samples at concentrations exceeding RVAAP installation background (USACE, 2001b) concentrations, Region 9 residential soil PRGs, or both. Copper, lead, mercury, and zinc in surface soil require further evaluation to determine the extent of contamination because these chemicals may pose a risk to plants and invertebrates. Most of the contaminants were detected in very few samples and there are insufficient occurrences to provide statistically valid analysis for contaminant distribution. Therefore, the nature and extent of the contamination in soil, sediment, surface water and groundwater at LL-9 have not been fully determined and additional sampling will be necessary to fully identify the lateral and vertical extent of the contamination at LL-9 for metals and propellants. A Phase II RI should be conducted to fill data gaps identified in this Phase I RI. If data collected in the Phase II RI alters the conceptual site model presented in this Phase I RI, then the conclusions of the risk assessments presented in this report will require revision.



10.0 REFERENCES

- Abromwicz, 1993. Presented at the Conference on Biodegradation: Its Role in Reducing Toxicity and Exposure to Environmental Contaminants, 26–28 April. Research Triangle Park, North Carolina.
- Agency for Toxic Substance and Disease Registry (ATSDR), 1995. Toxicological Profile for Polycyclic Aromatic Hydrocarbons.
- ATSDR, 2004a. Case Studies in Environmental Medicine, Polychlorinated Biphenyl Toxicity, Agency for Toxic Substances and Disease Registry, downloaded from the Intranet at: http://www.atsdr.cdc.gov/HEC/CSEM/pcb/biologic_fate.html, December 21.
- ATSDR, 2004b, Toxicological Profile for Cyanide (Draft for Public Comment), September. Downloaded from the Intranet at: <http://www.atsdr.cdc.gov/toxprofiles/tp8-c1.pdf>, December 21.
- Baes, D.F. III, R.D. Sharp, A.L. Sojoreen, and R.W. Shor, 1984. *A Review and Analysis of Parameters for Assessing Transport of Environmentally Released Radionuclides through Agriculture*. ORNL-5786. September. Oak Ridge National Laboratory, Oak Ridge, TN.
- Battelle and Exponent, 2000. *Guide for Incorporating Bioavailability Adjustments into Human Health and Ecological Risks Assessments at U.S. Navy and Marine Corps Facilities, Part 1: Overview of Metals Bioavailability*. Prepared for Naval Facilities Engineering Service Center, Port Hueneme, California. July.
- Bechtel Jacobs Company (BJC), 1998a. *Biota Sediment Accumulation Factors for Invertebrates: Review and Recommendations for the Oak Ridge Reservation*. Prepared for U.S. DOE, Office of Environmental Management. Prepared by BJC, Oak Ridge National Laboratory, Oak Ridge, Tennessee. BJC/OR-112.
- Buchman, M.F., 1999. *NOAA Screening Quick Reference Tables*. NOAA HAZMAT Report 99-1, Seattle WA, Coastal Protection and Restoration Division, National Oceanic and Atmospheric Administration.
- Canadian Council of Ministries of the Environment (CCME), 1997. *Recommended Canadian Soil Quality Guidelines*. ISBN 1-895-925-92-4.
- Census Bureau (U.S. Bureau of Census) 1992. Results of 1990 Census.



-
- 1 Condor, J.M., T.W. LaPoint, J.A. Stevens, and G.R. Lotupo, 2004. *Recommendations for*
2 *the Assessment of TNT Toxicity in Sediment*. Environmental Toxicology and Chemistry,
3 23(1):141–149.
- 4 Census Bureau (U.S. Bureau of Census), 1992. *Results of 1990 Census*.
- 5 Condor, J.M., T.W. LaPoint, J.A. Stevens, and G.R. Lotupo, 2004. Recommendations for
6 the Assessment of TNT in Sediment. *Environmental Toxicology and Chemistry*, 23
7 (1):141–149.
- 8 Daniels, J.I., Ed. 1988. *Evaluation of Military Field–Water Quality. Volume 4. Health*
9 *Criteria and Recommendations for Standards. Part 1. Chemicals and Properties of Military*
10 *Concern Associated with Natural and Anthropogenic Sources*. AD UCRL–21008.
- 11 Efroymsen, R.A., G.W. Suter II, B.E. Sample, and D.S. Jones, 1997a. *Preliminary*
12 *Remediation Goals for Ecological Endpoints*. Prepared for U.S. DOE, Office of
13 Environmental Management. Prepared by Lockheed Martin Energy Systems, Inc. Oak
14 Ridge National Laboratory, Oak Ridge, Tennessee. ES/ER/TM–162/R2.
- 15 Efroymsen, R.A., M.E. Will, and G.W. Suter II, 1997b. *Toxicological Benchmarks for*
16 *Contaminants of Potential Ecological Concern for Effects on Soil and Litter Invertebrates*
17 *and Heterotrophic Process*: 1997 Revision. Prepared for U.S. DOE, Office of
18 Environmental Management. Prepared by Lockheed Martin Energy Systems, Inc. Oak
19 Ridge National Laboratory, Oak Ridge, Tennessee. ES/ER/TM–126/R2.
- 20 Efroymsen, R.A., M.E. Will, G.W. Suter II, and A.C. Wooten, 1997c. *Toxicological*
21 *Benchmarks for Screening Contaminants of Potential Concern for Effects on Terrestrial*
22 *Plants*: 1997 Revision. Prepared for USDOE, Office of Environmental Management by
23 Lockheed Energy Systems, Inc., Oak Ridge National Laboratory, Oak Ridge, Tennessee.
24 ES/ER/TM–85/R3.
- 25 Emsley, Jr., 2001. *Nature's Building Blocks, An A–Z Guide to the Elements*, Oxford
26 University Press, 2001.
- 27 Environment Canada (EC), 1999a. *Canadian Soil Quality Guidelines for Chromium.*
28 *Scientific Supporting Document*. National Guidelines and Standards Office, Environmental
29 Quality Branch, Ottawa.
- 30 Environment Canada (EC), 1999b. *Canadian Soil Quality Guidelines for Free Cyanide.*
31 *Scientific Supporting Document*. National Guidelines and Standards Office, Environmental
32 Quality Branch, Ottawa.



-
- 1 Environment Canada (EC), 1999c. *Canadian Soil Quality Guidelines for Mercury.*
2 *Scientific Supporting Document.* National Guidelines and Standards Office, Environmental
3 Quality Branch, Ottawa.
- 4 Environment Canada (EC), 1999d. *Canadian Soil Quality Guidelines for Vanadium.*
5 *Scientific Supporting Document.* National Guidelines and Standards Office, Environmental
6 Quality Branch, Ottawa.
- 7 Environment Canada (EC), 1999e. *Canadian Soil Quality Guidelines for Zinc. Scientific*
8 *Supporting Document.* National Guidelines and Standards Office, Environmental Quality
9 Branch, Ottawa.
- 10 Environment Canada (EC), 1999f. *Canadian Soil Quality Guidelines for PCBs. Scientific*
11 *Supporting Document.* National Guidelines and Standards Office, Environmental Quality
12 Branch, Ottawa.
- 13 Environment Canada (EC), 1999g. *Canadian Soil Quality Guidelines for Arsenic.*
14 *Scientific Supporting Document.* National Guidelines and Standards Office, Environmental
15 Quality Branch, Ottawa.
- 16 Environment Canada (EC), 1999i. *Canadian Soil Quality Guidelines for Iron. Scientific*
17 *Supporting Document.* National Guidelines and Standards Office, Environmental Quality
18 Branch, Ottawa.
- 19 Environment Canada (EC), 1999h. *Canadian Soil Quality Guidelines for Thallium.*
20 *Scientific Supporting Document.* National Guidelines and Standards Office, Environmental
21 Quality Branch, Ottawa.
- 22 Erickson, 2001. *PCB Properties, Uses, Occurrence, and Regulatory History, pp. xi-xxx in:*
23 *PCBs—Recent Advances in the Environmental Toxicology and Health Effects of PCBs*
24 *(L.W. Robertson, and L.G. Hansen, Eds.). University Press of Kentucky, Lexington.*
- 25 Gilbert, Richard O., 1987. *Statistical Methods for Environmental Pollution Monitoring.*
26 Von Nostrand Reinhold Company, New York.
- 27 HAZWRAP (Hazardous Waste Remedial Action Program), 1994. Loring Air Force Base
28 Ecological Risk Assessment Methodology.
- 29 Hirsch, M.P. 1998. *Toxicity of Silver Sulfide-Spiked Sediments to the Freshwater*
30 *Amphipod (Hylella azteca). Environmental Toxicology and Chemistry.* Vol. 17, No. 4, pp.
31 601-604.



-
- 1 Kammer, H.W. 1982. AHydrogeologic Study of the Ravenna Arsenal, Eastern Portage and
2 Western Trumbull Counties, Ohio, Masters Thesis, Kent State University.
- 3 Lang, 1985. *Lang's Handbook of Chemistry*, J. A. Dean, editor. Thirteenth edition.
4 McGraw-Hill Book Company, New York.
- 5 Lide, D.R., Ed. 1991. *CRC Handbook of Chemistry and Physics, 1991-1992*. 72nd ed.
6 Boca Raton, FL, pp. 3-96.
- 7 Long, E.R. and L.G. Morgan, 1991. *The Potential for Biological Effects of Sediment-*
8 *Sorbed Contaminants Tested in the National Status and Trends Program*. NOAA
9 Technical Memorandum NOS OMA 52. National Oceanic and Atmospheric
10 Administration.
- 11 Long, E.R., D.D. MacDonald, S.L. Smith and F.D. Calder, 1995. Incidence of Adverse
12 Biological Effects Within Ranges of Chemical Concentrations in Marine and Estuarine
13 Sediments. *Environmental Management* 19(1):81-97.
- 14 MacDonald, D.D., C.G. Ingersoll, and T.A. Berger, 2000. *Development and Evaluation of*
15 *Consensus-Based Sediment Quality Guidelines for Freshwater Ecosystems*. Arch.
16 Environ. Contam. Toxicol. 39:20-31.
- 17 Ministry of Housing, Spatial Planning and Environment (MHSPE), 1994. *Intervention*
18 *Values and Target Values -Soil Quality Standards*. Directorate-General for
19 Environmental Protection, Department of Soil Protection, The Hague, Netherlands.
- 20 MKM Engineers, Inc. 2001a. *Sampling and Analysis Plan Addendum for the Phase II*
21 *Remedial Investigation at the Upper and Lower Cobbs Ponds (AOC 29)*, Ravenna Army
22 Ammunition Plant. Prepared for US Army Operations Support Command (OSC) Rock
23 Island, IL 61299-6000. July.
- 24 MKM Engineers, Inc. 2001b. *Work Plan for the Phase II Remedial Investigation at the*
25 *Upper and Lower Cobbs Ponds (AOC 29)*, Ravenna Army Ammunition Plant. Prepared
26 for U.S. Army Operations Support Command, Joliet, IL 61299-6000. July.
- 27 MKM Engineers, Inc. 2003a. *FWSAP Addendum for the Remedial Investigation at Load*
28 *Line 9 (RVAAP 42)* Ravenna Army Ammunition Plant. Ravenna, Ohio
- 29 MKM Engineers, Inc. 2003b. *Final Work Plan Appendix B*, Ravenna Army Ammunition
30 Plant. Ravenna, Ohio



-
- 1 NIH, 2004. Internet download from: [http://toxnet.nlm.nih.gov/cgi-](http://toxnet.nlm.nih.gov/cgi-bin/sis/search/f?/temp/~Y2cDf9:2:enex)
2 [bin/sis/search/f?/temp/~Y2cDf9:2:enex](http://toxnet.nlm.nih.gov/cgi-bin/sis/search/f?/temp/~Y2cDf9:2:enex), December, 16, 2004.
- 3 NJ, 2003. *New Jersey Comparative Risk Project*, Final Report, Rubenstein, D.I. and
4 Telford, S., co-chairs of the Steering committee, March.
- 5 ODNR (Ohio Department of Natural Resources), 1993. *Species and Plant Communities*
6 *Inventory. Ravenna Army Ammunition Plant.* ODNR The Nature Conservancy,
7 Columbus, Ohio.
- 8 ODNR (Ohio Department of Natural Resources), 2002a.
9 <http://www.dnr.state.oh.us/wildlife/resources/mgtplans/endangered.htm>
- 10 ODNR (Ohio Department of Natural Resources), 2002b.
11 <http://www.dnr.state.oh.us/wildlife/resources/mgtplans/specofconcern.htm>
- 12 ODNR (Ohio Department of Natural Resources), 2002c.
13 <http://www.dnr.state.oh.us/wildlife/resources/mgtplans/special%20interest.htm>
- 14 ODNR (Ohio Department of Natural Resources), 2004.
15 http://www.ohiodnr.com/dnap/heritage/Rare_Species2004.htm
- 16 Ohio Environmental Protection Agency (Ohio EPA), 2003. *Guidance for Conducting*
17 *Ecological Risk Assessments*. DERR-00-RR-031. Division of Emergency and Remedial
18 Response, Columbus, Ohio. February.
- 19 Ohio Environmental Protection Agency (Ohio EPA) 2004. *Recommendations for Interim*
20 *Toxicity Criteria for Trichloroethylene (TCE)*. Office of Federal Facilities Oversight.
21 January.
- 22 Ontario Ministry of Environment and Energy (OMOE), 1993. *Guidelines for the Protection*
23 *and Management of Aquatic Sediment Quality in Ontario*. Prepared by D. Persaud, R.
24 Jaagumagi, and A. Hayton for OMOE, ISBN 0-7778-9248-7. August.
- 25 ORNL, 2001. Risk Assessment Information System, http://risk.lsd.ornl.gov/rap_hp.shtml,
26 Oak Ridge National laboratory, Oak Ridge, TN, April 2001.
- 27 *Portage County Soil and Water Conservation District Resources Inventory*, 1985. Pp. 6-8.
- 28 Ratte, H.T., 1999. Bioaccumulation and Toxicity of Silver Compounds: A Review.
29 *Environmental Toxicology and Chemistry*, Vol.18, No. 1, pp. 89-108.



-
- 1 Rowe, B.L., S.J. Landrigan, and T.J. Lopes, 1997. *Summary of Published Aquatic Toxicity*
2 *Information and Water Quality Criteria for Selected Volatile Organic Compounds*. US
3 Geological Survey Open File Report 97-563.
- 4 Sample, B.E., D. M. Opresko, and G. W. Suter II, 1996. *Toxicological Benchmarks for*
5 *Wildlife*, 1996 Revision. Prepared for U.S. DOE, Office of Environmental Management.
6 Prepared by Risk Assessment Program, Health Sciences Research Division, Oak Ridge
7 National Laboratory, Oak Ridge, Tennessee. ES/ER/TM-86/R3.
- 8 Science Applications International Corporation (SAIC) 2001, *Phase II Remedial*
9 *Investigation Report for Winklepeck Burning Grounds at the Ravenna Army Ammunition*
10 *Plant, Ravenna Ohio*. Oak Ridge, TN, April.
- 11 Science Applications International Corporation (SAIC), 2003a. *Phase II Remedial*
12 *Investigation Report for Load Line 2 at the Ravenna Army Ammunition Plant*. Prepared for
13 the U.S. Army Corps of Engineers. May.
- 14 Science Applications International Corporation (SAIC), 2003b. *Phase II Remedial*
15 *Investigation Report for Load Line 3 at the Ravenna Army Ammunition Plant*. Prepared for
16 the U.S. Army Corps of Engineers. May.
- 17 Science Applications International Corporation (SAIC), 2003c. *Phase II Remedial*
18 *Investigation Report for Load Line 4 at the Ravenna Army Ammunition Plant*. Prepared for
19 the U.S. Army Corps of Engineers. May.
- 20 Suter, G.W., II and C.L. Tsao, 1996. *Toxicological Benchmarks for Screening Potential*
21 *Contaminants of Concern for Effects on Aquatic Biota* 1996 Revision. Lockheed Martin
22 Energy Systems at Oak Ridge National Laboratory, Oak Ridge, Tennessee. ES/ER/TM-
23 96/R2.
- 24 Talmage, S.S., D.M. Opresko, C.J. Maxwell, C.J.E. Welsh, F.M. Cretella, P.H. Reno, and
25 F.B. Daniel, 1999. Nitroaromatic munition compounds: Environmental effects and
26 screening values. *Rev. Environ. Contam. Toxicol* 161:1-156.
- 27 ToxNet, 2004. *Environmental Health Criteria* Vol:131 (1992) 127 p, downloaded from the
28 Internet on December 21, 2004.
- 29 Travis, C.C. and A.D. Arms, 1988. *Bioconcentration of Organics in Beef, Milk and*
30 *Vegetation*. *Environ. Sci. Technol* 22(3):271-274.



-
- 1 Trust, Kimberly A., A. Fairbrother, and M.J. Hooper, 1994. "Effects of 7,12-
2 Dimethylbenz(a,h)anthracene on Immune Function and Mixed-Function Oxygenase
3 Activity in the European Starling." *Env. Toxicol. Chem.* 13: 821-830
- 4 TtNUS. 2004. *White Paper – Human Health Risk Assessment Approach for Load Lines 6,
5 9, and 11 Remedial Investigations.* Ravenna Army Ammunition Plant, Ravenna, Ohio.
6 April.
- 7 UN, 1992. *Environmental Health Criteria, Diethylhexyl Phthalate.* United Nations
8 Environment Programme, the International Labour Organisation, and the World Health
9 Organization International Programme on Chemical Safety, Geneva.
- 10 UN, 1998. *Environmental Health Criteria, Selected Non-Heterocyclic Polycyclic Aromatic
11 Hydrocarbons.* United Nations Environment Programme, the International Labour
12 Organisation, and the World Health Organization International Programme on Chemical
13 Safety, Geneva.
- 14 USACE, 1996a. *Facility-Wide Sampling and Analysis Plan for the Ravenna Army
15 Ammunition Plant, Ravenna, Ohio.* DACA62-94-D-0029, D.O. 0009, April.
- 16 USACE, 1999. *Phase II Remedial Investigation Report for the Winklepeck Burning
17 Grounds at Ravenna Army Ammunition Plant, Ravenna, Ohio.* Prepared for the U.S. Army
18 Corps of Engineers Louisville District by SAIC, August.
- 19 USACE, 2001a. *Revised Facility-Wide Sampling and Analysis Plan for the Ravenna Army
20 Ammunition Plant, Ravenna, Ohio.* DACA 62-00-D-0001, D.O. CY02, March.
- 21 USACE, 2001b. *Phase II Remedial Investigation Report for the Winklepeck Burning
22 Grounds at Ravenna Army Ammunition Plant, Ravenna, Ohio.*
- 23 USACE, 2003a. *Phase II Remedial Investigation Report for Load Line 2 at the Ravenna
24 Army Ammunition Plant, Ravenna, Ohio.* (Preliminary Draft). Prepared for the U.S. Army
25 Corps of Engineers Louisville District by SAIC, May.
- 26 USACE, 2003b. *Phase II Remedial Investigation Report for Load Line 3 at the Ravenna
27 Army Ammunition Plant, Ravenna, Ohio.* (Preliminary Draft). Prepared for the U.S. Army
28 Corps of Engineers Louisville District by SAIC, May.
- 29 USACE, 2003c. *Phase II Remedial Investigation Report for Load Line 4 at the Ravenna
30 Army Ammunition Plant, Ravenna, Ohio.* (Preliminary Draft). Prepared for the U.S. Army
31 Corps of Engineers Louisville District by SAIC, May.



-
- 1 USACE, 2003d. *RVAAP Facility-Wide Ecological Risk Work Plan*. Louisville District,
2 ESSAYONS, U.S. Army Corps of Engineers.
- 3 USACE, 2004. *RVAAPs Facility-Wide Human Health Risk Assessor Manual*. January
- 4 U.S. Army Center for Health Promotion and Preventive Medicine (USACHPM), 1998.
5 *Relative Risk Site Evaluation for Newly Added Sites at the Ravenna Army Ammunition*
6 *Plant, Ravenna, Ohio*. Hazardous and Medical Waste Study No. 37-EF-5360-99,
7 October. 19-23
- 8 U.S. Army Toxic and Hazardous Materials Agency (USATHMA), 1978. *Installation*
9 *Assessment of Ravenna Army Ammunition Plant*. Records Evaluation Report No. 132.
- 10 USDA (U.S. Department of Agriculture), 1978 *Soil Survey of Portage County, Ohio*.
- 11 USDA (U.S. Department of Agriculture), 1992 *Soil Survey of Trumbull County, Ohio*.
- 12 U.S. Department of the Army, 1993 *Environmental Assessment*.
- 13 USEPA (U.S. Environmental Protection Agency), 1986. *Quality Criteria for Water- 1986*.
14 Office of Water Regulations and Standards. Washington, D.C. USEPA/440/5-86-001.
15 May.
- 16 USEPA (U.S. Environmental Protection Agency), 1987a. *Special Report on Ingested*
17 *Arsenic: Skin Cancer; Nutritional Essentiality*. Prepared for the Risk Assessment Forum,
18 Washington, D.C. USEPA/625/3-87/013.
- 19 USEPA (U.S. Environmental Protection Agency), 1987b. *Hazardous Waste Treatment,*
20 *Storage, and Disposal Facilities (TSDF) – Air Emission Models*, Office of Air and
21 Radiation, Research Triangle Park, N.C.
- 22 USEPA (U.S. Environmental Protection Agency), 1989a. *Risk Assessment Guidance for*
23 *Superfund, Vol. 1: Human Health Evaluation Manual (Part A)*, USEPA/540/1-89/002,
24 Washington, D.C.
- 25 USEPA (U.S. Environmental Protection Agency), 1989b. *Statistical Analysis of Ground-*
26 *water Monitoring Data at RCRA Facilities, Interim Final Guidance*. Office of Solid Waste.
27 February 1989.
- 28 USEPA (U.S. Environmental Protection Agency), 1990. *Guidance on Remedial Actions for*
29 *Superfund Sites with PCB Contamination*. OSWER Directive 9355.4-01



1 USEPA (U.S. Environmental Protection Agency), 1991a. *Risk Assessment Guidance for*
2 *Superfund, Vol. 1: Human Health Evaluation Manual (Part B, Development of Risk-based*
3 *Preliminary Remediation Goals)*, OSWER Directive 9285.7-01B, Office of Emergency and
4 Remedial Response, Washington, D.C.

5 USEPA (U.S. Environmental Protection Agency). 1991b. *Risk Assessment Guidance for*
6 *Superfund, Vol. 1: Human Health Evaluation Manual, Supplemental Guidance, Standard*
7 *Default Exposure Factors*, Interim Final, OSWER Directive 9285.6-03, Office of
8 Emergency and Remedial Response, Washington, D.C.

9 USEPA (U.S. Environmental Protection Agency), 1992a. *Dermal Exposure Risk*
10 *Assessment: Principles and Applications*, Interim Report, USEPA/600/8-91/011B,
11 including Supplemental Guidance dated August 18, 1992, Office of Research and
12 Development, Washington, D.C.

13 USEPA (U.S. Environmental Protection Agency), 1992b. *Supplemental Guidance to*
14 *RAGS: Calculating the Concentration Term*, Office of Solid Waste and Emergency
15 Response, Washington, D.C.

16 USEPA (U.S. Environmental Protection Agency), 1993a. *Wildlife Exposure Factors*
17 *Handbook, Volumes I and II*. Office of Research and Development, Washington, DC.
18 December. USEPA/600/R-93/187.

19 USEPA (U.S. Environmental Protection Agency), 1993b. *Water Quality Guidance for the*
20 *Great Lakes System and Correction: Proposed Rules*. *Federal Register* 58(72):20802-
21 21047.

22 USEPA (U.S. Environmental Protection Agency), 1994a. *USEPA Contract Laboratory*
23 *Program National Functional Guidelines for Organic Data Review*, USEPA-540/R-94-012
24 (PB94-963501), Office of Emergency and Remedial Response, Washington, D.C.

25 USEPA (U.S. Environmental Protection Agency), 1994b. *USEPA Contract Laboratory*
26 *Program National Functional Guidelines for Inorganic Data Review*, USEPA-540/R-94-
27 013 (PB94-963502), Office of Emergency and Remedial Response, Washington, D.C.

28 USEPA (U.S. Environmental Protection Agency), 1994c. *Guidance on Residential Lead-*
29 *based Paint, Lead-contaminated Dust, and Lead-contaminated Soil*. OSWER Directive
30 9355.4-12. Office of Prevention, Pesticides, and Toxic Substances, Washington, D.C.

31 USEPA (U.S. Environmental Protection Agency), 1994d. *Risk Reduction Engineering*
32 *Laboratory (RREL) Treatability Database*, Ver. 5.0. Office of Research and Development,
33 Cincinnati, Ohio.



-
- 1 USEPA (U.S. Environmental Protection Agency), 1995a. *Supplemental Guidance to Risk*
2 *Assessment Guidance for Superfund: Region 4 Bulletins, Human Health Risk*
3 *Assessment*, Waste Management Division, Atlanta, GA.
- 4 USEPA (U.S. Environmental Protection Agency), 1995b. *Groundwater and Leachate*
5 *Treatment Systems*, Office of Research and Development, Washington, D.C.
- 6 USEPA (U.S. Environmental Protection Agency), 1995c. *Policy for Risk Characterization*
7 *at the U.S. Environmental Protection Agency*. Carol Browner, Washington, D.C., March.
- 8 USEPA (U.S. Environmental Protection Agency), 1996. *Soil Screening Guidance:*
9 *Technical Background Document*. Office of Solid Waste and Emergency Response,
10 Washington, D.C.
- 11 USEPA (U.S. Environmental Protection Agency), 1997a. *Exposure Factors Handbook*,
12 USEPA/600/p-95/002, Office of Health and Environmental Assessment, Washington, D.C.
- 13 USEPA (U.S. Environmental Protection Agency), 1997b. *Ecological Risk Assessment*
14 *Guidance for Superfund: Process for Designing and Conducting Ecological Risk*
15 *Assessments*. Edison, New Jersey. June 5.
- 16 USEPA (U.S. Environmental Protection Agency), 1997c. *The Incidence and Severity of*
17 *Sediment Contamination in Surface Waters of the United States. Volume 1: National*
18 *Sediment Quality Survey*. Office of Science and Technology, Washington, D.C.
19 September. USEPA 823-R-97-006.
- 20 USEPA (U.S. Environmental Protection Agency), 1998. *Technical Protocol for*
21 *Evaluating Natural Attenuation of Chlorinated Solvents in Ground Water*. Office of
22 Research and Development, Washington D.C., USEPA/600/R-98/128, September.
- 23 USEPA (U.S. Environmental Protection Agency), Region 5, 1999. *Ecological Screening*
24 *Levels for RCRA Appendix IX Hazardous Constituents*. Working Draft. See also
25 <http://www.epa.gov/Region5/rcraca/edql.htm>.
- 26 USEPA (U.S. Environmental Protection Agency), 2000a. *Bioaccumulation Testing and*
27 *Interpretation for the Purpose of Sediment Quality Assessment: Status and Needs*.
28 Bioaccumulation Analysis Workgroup. February.
- 29 USEPA, (U.S. Environmental Protection Agency), 2000b. *Ecological Soil Screening Level*
30 *Guidance. (for Arsenic and Copper)*, Draft, USEPA, OERR, Washington D.C. July 10.



-
- 1 USEPA (U.S. Environmental Protection Agency), 2001a. *Risk Assessment Guidance for*
2 *Superfund, Vol. 1: Human Health Evaluation Manual, Part D.* Office of Solid Waste and
3 Emergency Response. Washington, D.C.
- 4 USEPA, (U.S. Environmental Protection Agency). 2002a. *National Recommended Water*
5 *Quality Criteria: 2002.* Office of Water, OST, Washington D.C. USEPA-822-R-02-047.
6 December.
- 7 USEPA, (U.S. Environmental Protection Agency). 2002b. *Supplemental Guidance for*
8 *Developing Soil Screening Levels for Superfund Sites.* December
- 9 USEPA, (U.S. Environmental Protection Agency). 2002c. *Calculating Upper Confidence*
10 *Limits for Exposure Point Concentrations at Hazardous Waste Site.* OSWER 9285.6-10
- 11 USEPA (U.S. Environmental Protection Agency), 2003a. *Ecological Screening Levels.*
12 Region 5 (<http://www.epa.gov/reg5rcra/ca/edql.htm>). August.
- 13 USEPA (U.S. Environmental Protection Agency), 2003b. *Ecological Soil Screening Level*
14 *for Lead, Interim Final.* Office of Emergency and Remedial Response. OSWER Directive
15 9285.7-70. November.
- 16 USEPA (U.S. Environmental Protection Agency), 2003c. *Ecological Soil Screening Level*
17 *for Iron.* Interim Final. U.S. Environmental Protection Agency, Office of Emergency and
18 Remedial Response, Washington, D.C. OSWER Directive 9285.7-69. November.
- 19 USEPA, (U.S. Environmental Protection Agency), 2003d. *Ecological Soil Screening Level*
20 *for Aluminum.* Interim Final. Washington D.C. OSWER Directive 9285.7-60. November.
- 21 USEPA, (U.S. Environmental Protection Agency), 2003e. *Ecological Soil Screening Level*
22 *for Barium.* Interim Final. Washington D.C. OSWER Directive 9285.7-63. November.
- 23 USEPA, (U.S. Environmental Protection Agency), 2003f. *Ecological Soil Screening Level*
24 *for Cobalt.* Interim Final. Washington D.C. OSWER Directive 9285.7-67. November.
- 25 USEPA (U.S. Environmental Protection Agency), 2004a. *Risk Assessment Guidance for*
26 *Superfund, Vol. 1: Human Health Evaluation Manual, Part E: Supplemental Guidance for*
27 *Dermal Risk Assessment.* Interim Guidance, Office of Solid Waste and Emergency
28 Response. Washington, D.C.
- 29 USEPA (U.S. Environmental Protection Agency), 2004b. *Integrated Risk Information*
30 *System (IRIS).* Environmental Criteria and Assessment Office, Office of Health and
31 Environmental Assessment, Cincinnati, Ohio.



-
- 1 USEPA (U.S. Environmental Protection Agency), 2004c. *Region 9 Preliminary*
2 *Remediation Goals (PRGs)*, 1998. Created by Stanford J. Smucker and found on the
3 world wide web address <http://www.epa.gov/region09/waste/sfund/prg/index.html>.
- 4 USEPA (U.S. Environmental Protection Agency), 2004d. *Drinking Water Standards and*
5 *Health Advisories*. USEPA 822-B-00-001, Washington, D.C.
- 6 USEPA (U.S. Environmental Protection Agency), 2005.
- 7 US Navy, 1998. *Technical Guidelines for Evaluating Monitored Natural Attenuation of*
8 *Petroleum Hydrocarbons and Chlorinated Solvents in Groundwater at Naval and Marine*
9 *Corps Facilities*. Prepared for Southwest Division Naval Facilities Engineering Command,
10 Southern Division Naval Facilities Engineering Command, and Engineering Field Activity,
11 Northwest. Prepared by: Parsons Engineering Science, Inc. Pasadena, California and
12 United States Geological Survey, Columbia, South Carolina, September.
- 13 Van Nostrand, 1976. Van Nostrand's Scientific Encyclopedia, fifth edition. (E. Considine,
14 editor), Van Nostrand Reinhold Company, New York.
- 15 White, G. W. 1982. *Glacial Geology of Northeastern Ohio*. Bulletin 68, Division of
16 Geological Survey, Ohio Department of Natural Resources.
- 17 Williams, G.M. and J. H. Welsburger, 1991. "Chemical Carcinogenesis." Cassarett and
18 Doull's *Toxicology, The Basic Science of Poisons*, 4th Edition. Pergamon Press, New
19 York, New York.
- 20 World Health Organization (WHO). 1984a. *Guidelines for Drinking Water Quality*.
21 *Vol. 2. Health Criteria and Other Supporting Information*.
- 22 World Health Organization (WHO). 1984b. *Guidelines for Drinking Water Quality*.
23 *Vol. 1. Recommendations*.