

Final

**United States Army Corps of Engineers
Ravenna Army Ammunition Plant (RVAAP)
Position Paper for the Application and Use of
Facility-Wide Human Health Cleanup Goals**

**Ravenna Army Ammunition Plant
Ravenna, Ohio**

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Prepared by:



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LOUISVILLE DISTRICT

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1.0 PURPOSE AND NEED

The purpose of this Position Paper is to provide *interim* guidance from the United States Army Corps of Engineers, Louisville District (USACE) to Contractors regarding the use and application of facility-wide cleanup goals as part of the path forward in the risk assessment process for

- Determining presence/absence of contamination,
- Assessing data gaps,
- Evaluating nature and extent of contamination, and
- Identifying cleanup requirements.

Ravenna Army Ammunition Plant (RVAAP) has worked closely with the Ohio Environmental Protection Agency (Ohio EPA) and other stakeholders such as the Ohio Army National Guard (OHANG) to develop an acceptable approach to the completion of human health risk assessments. Because of the initial successes of the human health risk assessment program, there was mutual agreement to streamline the process. Streamlining the Human Health Risk Assessment process resulted in the establishment of Facility-wide Cleanup Goals (CUGs). Currently, the CUGs are in draft form and are being developed by Science Applications International Corporation (SAIC) in the document *Preliminary Draft Facility-wide Human Health Remediation Goals, Ravenna Army Ammunition Plant, Ravenna, Ohio, May 2008* (FWCUG Report). The original intent of developing the CUGs was to eliminate the need for baseline risk assessments. Since the development of the CUGs, they also have been recognized as appropriate tools to be used in screening-level assessments.

The CUGs were developed to reduce the level of effort and to limit the amount of time required to make informed risk management decisions regarding sampling locations, delineations of contamination, data gaps, and remediation of contaminants without needing to complete a baseline risk assessment. The selection of chemicals requiring a CUG is based upon the screening process outlined in the *Ravenna Army Ammunition Plant Facility-Wide Human Health Risk Assessor Manual, Amendment 1* (USACE 2005), herein referred to as the Risk Manual.

Besides the screening process, the Risk Manual requires that prior to commencing any risk assessment activities at the Ravenna Army Ammunition Plant (RVAAP), a White Paper should be developed to ensure regulatory agreement with the processes proposed. The White Paper for the development of the CUGs can be found as an attachment to the FWCUG Report. The White Paper provided clarification of technical issues related to developing the CUGs that were not defined in the Risk Manual. The White Paper also included the exposure pathways and parameters pertinent to two newly identified future land uses: the Engineering School use and the Small Arms Range use.

The intent of USACE is to develop a single document that includes pertinent information from the Risk Manual, FWCUG Report (including the White Paper), CUGs, risk assessment information found in previously published documents, and technical guidance and agreements that USACE has developed in conjunction with the Ohio EPA. It is planned that the FWCUG Report will be modified and updated to include all these items. The FWCUG Report will then replace the Risk Manual. The USACE believes that this approach will clarify requirements for the completion of screening level human health risk assessments, selection of appropriate CUGs, and the determination of remediation levels that are risk-

based. Because this unified document will not be available for several months, USACE identified the need for this Interim Guidance.

2.0 DATA EVALAUTION – DETERMINATION OF THE CHEMICALS OF POTENTIAL CONCERN

The first step in using the CUGs and determining which ones should be used depends upon what phase of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) process is being investigated and what decisions will be made using the CUGs. If the data is being evaluated to determine the presence or absence of contamination, nature and extent of contamination, characterization of contamination, sampling locations, or for other reasons where the data identifies potential contamination, then the initial evaluation step should be completed. The end result of the initial data screening process is a list of chemicals of potential concern (COPCs). The COPC screen shall follow the general guidance of the Risk Manual (Sections 3.4 and 3.5):

1.) The concentrations of inorganics shall be compared to the soil background concentrations in the report titled *Phase II Remedial Investigation Report for the Winklepeck Burning Ground at RVAAP, OH* (USACE April 2001). If exceedances above background occur, then the respective metals are retained as COPCs.

2.) Consistent with the Risk Manual (Section 3.5), chemicals identified as essential nutrients will be screened out. Chemicals that are considered essential nutrients (e.g., calcium, chloride, iodine, iron, magnesium, potassium, phosphorous, and sodium) are an integral part of the human food supply and are often added to foods as supplements. The USEPA recommends that these chemicals not be evaluated as COPCs as long as they are: (1) present at low concentrations (i.e., only slightly elevated above naturally occurring levels), and (2) toxic at very high doses (i.e., much higher than those that could be associated with contact at an Area of Concern (AOC).

3.) Chemicals meeting the <5% detection rule may be screened out per Section 3.4.1 of the Risk Manual; however, this screening step is based upon having a statistically valid data set (sample size of at least 20).

4.) For the determination of the COPCs, all chemicals that have not been eliminated should be screened against their specific CUG at the 1.0×10^{-6} cancer risk level and non-carcinogenic risk Hazard Quotient (HQ) using the 0.1 risk value for the Residential Farmer Adult, Residential Farmer Child, and the National Guard Trainee. If there are no CUGs developed for the particular chemical, then the USEPA Regional Screening Levels (RSLs) should be used. The Risk Manual required that the maximum concentrations be compared to the USEPA Region 9 Preliminary Remediation Goals (PRGs); however, the RSLs have replaced the USEPA's Region 9 PRGs. The RSLs should only be used if a CUG is not available for the chemical. The RSL for the Residential Receptor should be used. This Interim Guidance requires that the comparison be completed using the Draft CUGs. Once the CUGs have been finalized, this Interim Guidance will be superseded by the Final FW CUG Report. The steps listed below should be followed for the comparison process to be acceptable and complete when establishing COPCs or characterizing the contamination in an area:

- Use the CUGs developed for the Residential Farmer Adult and Child Receptors and the National Guard Trainee for each chemical. If no CUG is available, use the USEPA's RSL for the chemical. If neither the CUG nor the RSL is available, then a CUG should be developed or another approach must be developed with concurrence from USACE and Ohio EPA.
- Select the CUGs at the 1.0×10^{-6} carcinogenic value and the non-carcinogenic risk value termed Hazard Quotient (HQ) using the 0.1 risk value.

- Report all carcinogenic and non-carcinogenic risk values for each chemical for the Adult and Child Residential Farmer and the National Guard Trainee.
- Complete a comparison of the selected CUG to the Exposure Point Concentration (EPC). The EPC will be either the 95% Upper Confidence Limit (UCL) of the mean for each chemical concentration or the maximum value detected, depending upon whichever value is the lowest. In comparisons where the 95% UCL can not be determined, the maximum concentration of the chemical should be compared to the appropriate CUGs.
- Consider the chemical as a COPC if the EPC exceeds the most stringent risk value for the Adult Resident Farmer, the Child Resident Farmer, or the National Guard Trainee for either one of the 1.0×10^{-6} carcinogenic value and the non-carcinogenic risk value termed Hazard Quotient (HQ) using the 0.1 risk value.

3.0 DETERMINATION OF THE CHEMICALS OF CONCERN

The original application of the CUGs was that they would be used to determine remediation levels and to assist in the completion of remedial design and processes. Once the COPCs are established and all sampling has been completed so that the nature and extent of the contamination is known, the next step is to determine which of the COPCs are Chemicals of Concern (COCs). The determination of the COCs consists of a screening of the chemical concentration to specific CUGs. However, unlike the COPC comparison, the COCs are determined by comparing the chemical concentration to different risk levels and potentially, for different receptors.

The determination of the COCs uses a less stringent risk value but must address the potential for additive effects. To account for the potential additive effects from exposure to multiple chemicals or exposure to multiple chemicals that can cause the same effect (e.g., cancer) or affect the same target organ, then the "Sum of Ratios" approach should be used. This approach compares the chemical concentration (e.g., mean concentration or concentration in confirmation samples) of the COPC to the individual CUG to determine a ratio. The Sum of Ratios method is based upon the principle that a ratio greater than 1 represents unacceptable risk and a ratio less than or equal to 1.0 represents acceptable risk. If there are multiple chemicals in the samples that are being evaluated and they are carcinogens or if there are non-carcinogens that affect the same target organ, then the Sum of Ratios of both the carcinogens and the non-carcinogens, respectively, must be less than or equal to 1.0. If the Sum of Ratios is > 1 and there are chemicals contributing at least 10% to the sum, then they are also considered COCs for the site.

Several examples of the Sum of Ratios approach are presented in the following. Example 1 of the Sum of Ratios is for three chemicals (i.e., A, B, and C) that may affect the same target organ.

Example 1. Sum of Ratios less than 1 and no chemical identified as a COC.

Chemical	EPC or Maximum Concentration	FW CUG	Ratio of EPC to FWCUG	% Contribution to the total sum	COC Yes or No
Chemical A	1	2.1	0.5	88	No
Chemical B	2	56	0.04	7	No
Chemical C	8	320	0.03	5	No
Sum of Ratios			0.57		

In Example 1, Chemical B and Chemical C are eliminated because they do not contribute more than 10% of the total Sum of Ratios. Chemical A is not retained as a COC although it is contributing more than

10% of the total Sum of Ratios, but the total is less than 1. Without doing the Sum of Ratios approach, no consideration would have been given to the chemicals collectively to determine if there was potential for any of them being a COC since their EPCs were less than their CUGs.

Example 2 shows the Sum of Ratios follows for three chemicals (i.e., E, F, and G) that may affect the same target organ where the Sum of Ratios is greater than 1.0 and several COCs are identified.

Chemical	EPC or Maximum Concentration	FW CUG	Ratio of EPC to FW CUG	% Contribution to the total sum	COC Yes or No
Chemical E	1	2	0.5	0.32	Yes
Chemical F	57	56	1.02	0.66	Yes
Chemical G	8	320	0.03	0.02	No
Sum of Ratios			1.55		

In this example, Chemical G is eliminated because its ratio does not contribute more than 10% of the total Sum of Ratios. Chemicals E and F are retained as a COC because their ratios are contributing more than 10% of the total Sum of Ratios. Without the Sum of Ratios approach, only chemical F would have been considered a COC since its EPC exceeds its CUG value.

The determination of the COCs should follow this Interim Guidance until the Final FW CUG Report is available. The screening process is as follows:

- Select the CUGs developed for the Resident Farmer Adult and Child Receptors and the receptor for the planned future land use by the Ohio Army National Guard.
- Select the CUGs at the 1.0×10^{-5} carcinogenic value and the non-carcinogenic risk value termed Hazard Quotient (HQ) using the 1.0 risk value.
- Report all carcinogenic and non-carcinogenic risk values for each chemical for all receptors.
- Report critical effect and target organ for each of the non-carcinogenic risk values.
- Complete a comparison of the selected CUG to the EPC. The EPC will be either the 95% UCL of the mean for each chemical concentration or the maximum value detected, depending upon whichever value is the lowest. In comparisons where the 95% UCL can not be determined, the maximum concentration of the chemical should be compared to the appropriate CUGs.
- For non-carcinogens, compare the chemical-specific concentration to the target risk CUG. Sum the ratios of chemicals that affect similar target organs.
- For carcinogens, compare the chemical-specific concentration to the target risk CUG. Sum the ratios of all carcinogens.
- Consider the chemical as a COC if the EPC exceeds the most stringent risk value for either the Adult Resident Farmer or the Child Resident Farmer, and/or the OHARNG planned future use receptor, for either one of the 1.0×10^{-5} carcinogenic value and the non-carcinogenic risk value termed Hazard Quotient (HQ) using the 1.0 risk value. The Sum of Ratios for all carcinogens and all non-carcinogens that may affect the same organ must be less than or equal to 1.0 as well.

If the Sum of Ratios for all carcinogens and all non-carcinogens (that may affect the same organ) are greater than 1 then the chemicals contributing at least 10% to the sum are considered COCs.

4.0 USE OF CUGS DURING REMEDIATION AND CONFIRMATION

In general, the CUGs for each of the COCs identified are the actual remediation levels unless there are additive effects. In some circumstances there may be a risk management analysis such as a “Weight of Evidence” approach that may allow the COC to be re-assessed. As described in the previous section, the Sum of Ratios is used to account for the potential additive effects from exposure to multiple chemicals that can cause the same effect (e.g., cancer) or affect the same target organ. This approach compares the chemical concentration (e.g., mean concentration or concentration in confirmation samples, the EPC) of the COC to the individual CUG to determine a ratio.

This final application of CUGs would generally occur during the Feasibility Study (FS) or during remediation. During the determination of COCs, and accounting for potential additive effects, the numbers that were obtained are essentially the remediation levels and would be used for the confirmation samples. The target risk values are the same for remediation levels as they are for the determination of the COCs. The cancer risk is 1.0×10^{-5} and the non-carcinogenic HQ is 1.0.

Chemicals that are identified in confirmation samples that were not identified as COCs need to be considered in the overall estimation of the success of the remediation. For example, if a chemical is detected and is shown to affect the same target organ as one that has a remediation level established, then the chemical should be considered using a Sum of Ratio approach to determine if there are any risks remaining for the designated end user.

5.0 REFERENCES

Science Applications International Corporation (SAIC). 2001. Facility-Wide Sampling and Analysis Plan for Environmental Investigations at the Ravenna Army Ammunition Plant, Ravenna, Ohio.

United States Army Corps of Engineers (U.S. Army Corps of Engineers). 2004. Focused Feasibility Study for the Winklepeck Burning Grounds at the Ravenna Army Ammunition Plant, Ravenna, Ohio, DACA62-00-D-0001, D.O. CY08, March.

USACE. 2005. Ravenna Army Ammunition Plant Facility-Wide Human Health Risk Assessor Manual, Amendment 1, Prepared by the U.S. Army Corps of Engineers, Louisville District, November.