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Rocky Mountain Arsenal

Proposed Final Remedial Investigation Report Volume VII Eastern Study Area, Section 2.0 Version 3.2

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PREPARED BY:

EBASCO SERVICES INCORPORATED APPLIED ENVIRONMENTAL, INC. CH₂M HILL DATACHEM, INC. R. L. STOLLER ASSOCIATES, INC.

PREPARED FOR:

U.S. ARMY PROGRAM MANAGER'S OFFICE FOR ROCKY MOUNTAIN ARSENAL CONTAMINATION CLEANUP

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13. ABSTRACT (Maximum 200 words) THE EASTERN STUDY AREA (ESA) REPORT INTEGRATH GEOLOGY, AND HYDROLOGY WITH THE RESULTS OF SOIL, AIR, BIOTA, AND STRUCTURES INVESTIGATIONS TO DEH CONTAMINATION IN THE EASTERN TIER. THE ESA INCLUDES ALL OF SECTIONS 5, 19, 29, 3 SECTIONS 6, 7, 8, 20, 24, AND 25. THE ESA SERVE ALTHOUGH SOME SITES WERE USED FOR STORAGE AND DI CHEMICAL SAMPLING INCLUDED ANALYSIS OF 1457 SAME AT SEVEN SURFACE WATER LOCATIONS, AND MONITORING THE VOLUME OF POTENTIALLY CONTAMINATED SOIL IS E CUBIC YARDS. THIS REPORT IS ORGANIZED IN THREE SECTIONS: 1. CHARACTERIZATION OF THE STUDY AREA - GEC HISTORY	ES THE STUDY AREA , SURFACE WATER, FINE THE NATURE A BO, 31, AND 32 AN ED PRIMARILY AS A (SPOSAL OF MUNITI PLES FROM 751 BOR G OF FORTY WELLS. ESTIMATED TO BE 1 PLOGY, HYDROLOGY,	A HISTORY, GROUND WATER, AND EXTENT OF ND PORTIONS OF A BUFFER ZONE, CONS. THE RINGS, SAMPLING .12 MILLION CLIMATE,			
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Standard Abbreviations used in Eastern Study Area Report

1	Analyte Group					
*•	VUO	Volatile halogenated organic compounds				
	VIIC	Volatile halogenated organic compounds				
	VIC	Volatile hydrocarbons				
	VAU	Volatile aromatic organic compounds				
	OSCM	Organosulfur compounds, mustard-agent related				
	OSCH	Organosulfur compounds, herbicide related				
	OPHGB	Organophosphorous compounds, GB-agent related				
	OPHP	Organophosphorous compounds, pesticide related				
	DBCP	Dibromochloropropage				
	ONC	Organonitrogen compounds				
	DAL	Balumuslaan anomatia hudaaanhana				
	rAn SUO	Polynuclear aromatic hydrocarbons				
	SHO	Semivolatile halogenated organic compounds				
	OCP	Organochlorine pesticides				
	ICP Metals	Metals analyzed for by inductively coupled argon plasma, includes				
		cadmium (Cd), chromium (Cr), copper (Cu), lead (Pb), and zinc				
		(Zn)				
	As	Arsenic				
	Hg	Mercury				
	0	······································				
2.	National Acts & Organizations					
	AMCCOM	Armament Munitions and Chemical Command				
	CERCIA	Comprehensive Environmental Response Compensation and				
	CLACLA	Lishilin, Ant				
	NCD	Liability Act				
	NCP	National Contingency Plan				
	NOAA	National Oceanic and Atmospheric Administration				
	SARA	Superfund Amendments and Reauthorization Act				
	USACOE	United States Army Corps of Engineers				
	USAEWES	United States Army Engineer Waterways Experiment Station				
	USATHAMA	United States Army Toxic and Hazardous Materials Agency				
	USAEWES	United States Army Engineer Waterways Experiment Station				
	LISDA-SCS	United States Department of Agriculture - Soil Conservation				
	000/1000	Samina				
	LICEDA	Juliand Canada David and Destantion Association				
	USERA	United States Environmental Protection Agency				
	USF WS	United States Fish and Wildlife Service				
•						
3.	Local Termino	logy				
	BCF	Bioconcentration Factor				
	BCRL	Below Certified Reporting Limit				
	CAR	Contamination Assessment Report				
	CDH	Colorado Department of Health				
	CDOW	Colorado Division of Wildlife				
	CRL	Certified Reporting Limit				

- CSA
- Certified Reporting Limit Central Study Area Endangerment Assessment Eastern Study Area Electrostatic Precipitator EA
- ESA
- ESP

FS	Feasibility Study
NCSA	North Central Study Area
NBCS	North Boundary Containment System
STP	Sewage Treatment Plant
PMO or	Program Managers Office for the RMA Contamination
PMRMA	Cleanup
RAA	Remedial Action Alternative
RI	Remedial Investigation
RIC	Resource Information Center
RMA	Rocky Mountain Arsenal
RMACCPMT	Rocky Mountain Arsenal Contamination Cleanup Program
	Managers Team
SAR	Study Area Report
SCS	Soil Conservation Service
TPP	Technical Program Plan
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4. Companies

001110011100	
EBASCO	Ebasco Services Incorporated
ESE	Hunter/Environmental Science & Engineering, Inc.
G&M	Geraghty & Miller, Inc.
HLA	Harding, Lawson, & Associates
MKE	Morrison-Knudsen Engineers, Inc.

- Unified Soil Classification System (USCS) Textural Key CL inorganic clay, low plasticity 5.

 - CH inorganic clay, high plasticity
 - GC clayey gravel
 - GP poorly graded gravel
 - inorganic silt MH
 - inorganic silt, low plasticity ML
 - SC clayey sand
 - SM silty sand
 - poorly graded sand well graded sand SP
 - SW

6. Measurements

Measurements	
ac-ft/yr	acre - feet per year
cfs	cubic feet per second
msl	mean sea level
ppm	parts per million
ppb	parts per billion
µg/g	micrograms per gram, equivalent to parts per million (ppm)
μg/1	micrograms per liter, nearly equivalent to parts per billion (ppb)
AA	atomic absorption
Со	ratio of contaminant concentration in an organism
CVAA	cold vapor atomic absorption
Eh	oxidation potential

 f_{oc} soil organic carbon content

xvi

GC/EC	gas chromatography/electron capture
GC/MS	gas chromatography/mass spectrometry
K_d	soil - water coefficient
K_h	Henry's Law constant
K_{∞}	organic carbon partition coefficient
K_{ow}	octanol - water partition coefficient

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2.0 CONTAMINANT DISTRIBUTION

This section discusses analytical methodologies and presents the results of chemical analyses performed on soils, surface water, groundwater, structures, air, and biota samples collected from the ESA. The results from visual inspections of structures for potential contaminants is also presented in this section. Each of the five media, soil, water, air, structures, and biota, are discussed separately. For previously designated sites and nonsource areas studied during the RI program, much of the soils analytical data was presented on a site-by-site basis in the Phase I CARs. Results of Phase II investigations are presented in Phase II data addenda packets for the respective sites and nonsource areas. The CARs used for the compilation of this report were listed in Section 1.0 on Table ESA 1.1-1.

Overall, 634 Phase I soil borings were drilled in the ESA, and 1,135 samples were analyzed for the Phase I suite of analytes. A total of 195 Phase I soil samples were also analyzed for chemical agent degradation products. A total of 117 Phase II borings were drilled, and 322 samples were analyzed for various analytes to further investigate potentially contaminated areas and to evaluate previously unidentified areas. The locations of all borings, as well as all water quality monitoring wells, are presented in Plate ESA 2.0-1. Phase I geophysical surveys were conducted at all sites suspected of metal disposal or burial, and trenching activities were conducted at seven sites to further investigate areas of anomalous geophysical response. Eighteen Phase I programs were conducted at sites in the ESA, and all or portions of twelve nonsource area investigations were within the ESA boundaries. Phase II programs were conducted at 13 sites and in six nonsource areas. Biota and air samples were collected from the ESA and structures were surveyed. Additional information, as ascertained from previous investigations, is discussed in Section 1.1.1, involving the collection and analysis of geologic, hydrologic, and geochemical data from wells and borings in the ESA.

Surface water sampling was conducted at seven locations along First Creek, and stream elevation measurements were taken from two stations. Groundwater samples were analyzed from as many as forty wells per fiscal year quarter in the ESA, and water levels

were obtained from as many as sixty-three wells per fiscal year quarter. Analytical suites for water samples were based on the Tasks 4, 25, and 44 programs, and are discussed in detail in Section 2.2. The water analytical data used in this report were taken from the Water RI Draft Final Report (Ebasco, 1989/RIC 89067R08); the Final Initial Screening Program Report (ESE, 1987b/RIC 87253R01); the Final Screening Program, Third and Fourth Quarters, Final Report (ESE, 1988k/RIC 88173R06); and the Boundary Control Systems Assessment Draft Final Report (ESE, 1988dd/RIC 89024R02).

The potential for contaminants in structures was classified based on historical data and visual inspections; the results are discussed in Section 2.4. Data for this section were taken from the Structures Survey Final Summary of Results Report (Ebasco, 1988w/RIC 88306R02). Air quality in the ESA was monitored for total suspended particulates in one location and is presented in Section 2.5. Data for this section were taken from the Air RI Final Report (ESE, 1988s/RIC 88263R01). Biota samples from several plant and wildlife species were analyzed, and the results are summarized in Section 2.6. Data for this section were taken from the Biota RI Draft Final Report (ESE, 1989a/RIC 89054R01).

Many of the figures in this section present the distribution of contaminants in the different media. Boring, well, surface water, and biota sampling locations on these figures were plotted by computer onto a grid system defined by State Planar Coordinates. These locations were then projected onto a study area base map generated from Basic Information Maps, which were originally developed by the Army Corps of Engineers.

2.1 SOIL AND SEDIMENT SAMPLES

Soil and sediment samples were collected from each of the six site groups within the ESA. A total of 751 borings, yielding 320 composite and 1,137 uncomposited soil and sediment samples, were completed in the ESA during the course of the Phase I and II RIs under Tasks 7, 14, 15, 20, 21, and 22. Composite soil samples were obtained from nonsource areas, and consisted of soil taken from the 0 to 1 and 4 to 5 ft intervals. Uncomposited soil samples were obtained from sites suspected to contain contaminants.

In several instances, auger refusal did not allow the planned depths to be obtained. The objective of Phase I was twofold, to screen for areas of significant contamination, and to collect samples in areas where no historical or physical evidence of activity existed. The second objective provided background analytical information on large areas. The objective of Phase II was to further investigate any specific areas containing contaminants that were identified during Phase I and to better define the horizontal and vertical extent of contaminants.

2.1.1 Sampling Program and Analytical Methods

The soils investigation at RMA was conducted in two phases. Phase I investigations identified potential contaminants and provided a preliminary assessment of the vertical and lateral extent of contaminants. Phase I results also provided the basis for the design of a more quantitative Phase II program. Phase II sampling was conducted at some sites to provide a more quantitative assessment of the vertical and areal extent of detected contaminants. Data from Phase I of the RI were published in the CAR for each site and nonsource area, and Phase II data were published in a corresponding Phase II Data Addendum. These documents are listed and referenced in Section 1.1 of this report.

In the Phase I program, soil borings were drilled to various depths in the vadose zone. Samples were generally collected from these borings at standard sampling depths of 0 to 1, 4 to 5, 9 to 10, 14 to 15, 19 to 20 ft, and at 10 ft intervals below 20 ft. Samples were collected from nonstandard intervals where drilling was difficult or where staining or other evidence of potential contamination was observed in the core. Phase II samples were collected either at standard intervals or in depth intervals above and below Phase I samples in which contaminants were detected.

Samples from the Phase I borings were analyzed for a standard suite of compounds. The Phase I analyses for target compounds are listed in Appendix A and include:

- gas chromatography/mass spectrometry (GC/MS) analysis for volatile organics (VOs);
- o GC/MS for semivolatile organics (SVOs);

- o gas chromatography/electron capture (GC/EC) analysis for DBCP;
- inductively coupled plasma (ICP) screen for the metals cadmium,
 chromium, copper, lead, and zinc;
- o atomic absorption spectroscopy (AA) for arsenic; and
- o cold vapor atomic absorption spectroscopy (CVAA) for mercury.

More complete information on the soil analytical methods may be found in the report titled "Development and Evaluation of Analytical Methodologies Used in RMA Soil Investigations" (Ebasco, 1988k/RIC 88127R02). Some samples were also analyzed by high-performance liquid chromatography (HPLC) for thiodiglycol and agent degradation products.

In the ESA sites, the GC/MS analysis for VOs was applied only to samples from below the 0 to 1 ft depth interval, since these compounds are expected to volatilize from surface soils. VOs were normally not analyzed in samples collected outside of the ESA sites, as the samples from these other areas were composites of the 0 to 1 and 4 to 5 ft intervals.

The GC/MS method provided positive identification and semiquantitative concentration results for target analytes above the Certified Reporting Limit (CRL). The lower CRL is defined as the lowest concentration of analyte in the sample being analyzed that can be reported within a ninety percent confidence interval, using valid precision and accuracy criteria. The upper CRL is the maximum concentration of an analyte in the sample being analyzed that can be reported within a ninety percent within a ninety percent confidence interval, using valid precision and accuracy criteria (USATHAMA, 1987). For the purposes of this report, "CRL" will refer to the lower CRL and "upper CRL" will refer to the maximum quantification limit.

Where GC/MS methods were used, nontarget compounds were tentatively identified by establishing a "best fit" identification using a computer library of spectra. Some of these compounds were been added to the list of target compounds that are evaluated in the

SARs. Non-target compounds were added to the list of target compounds based on the following criteria:

- Fully identified according to analytical technique;
- Arsenal-activity or fuel-component related, rather than naturally occurring;
- Moderate to high carcinogenicity and/or toxicity, as outlined in the Chemical Index (Ebasco, 1988d/RIC 88357R01);
- Frequency of occurrence and concentration;
- Co-occurrence with target compounds on a site-by-site basis.

These significant nontarget compounds are:

VHOs

1,1,2,2-tetrachloroethane trichloropropene

<u>VHCs</u>

2-butoxyethanol 4-hydroxy-4-methyl-2-pentanone 1-methyl-1,3-cyclopentadiene methylcyclohexane 2,2-oxybisethanol 2-pentanone

<u>OPHGBs</u>

phosphoric acid, tributyl ester phosphoric acid, triphenyl ester

<u>ONCs</u>

caprolactam

<u>PAHs</u>

fluoroanthene methylnaphthalene phenanthrene pyrene

SHOs

trichlorobenzene hexachlorobenzene hexachlorobutadiene tetrachlorobenzene pentachlorobenzene

Because the method used to identify these compounds has not been subjected to U. S. Army Toxic and Hazardous Materials Agency (USATHAMA) certification procedures, these compounds have no CRL. The lower limit of detection has been assumed to correspond to ten percent of the internal standard for the GC/MS methods used, which is $0.3 \ \mu g/g$.

The GC/MS analyses for VOs and SVOs were certified by USATHAMA to detect a variety of analytes and to accomplish the Phase I objective of identifying contaminants present in the study area. Phase II methods were developed and certified for use in further quantifying the concentrations of the target compounds identified in Phase I. The Phase II methods were more sensitive GC methods certified for fewer compounds at lower reporting limits. In addition, approximately ten percent of the samples analyzed by GC/MS for confirmation of the GC results.

The Phase I and II target and significant nontarget compounds, analytical methods used, and laboratory CRLs for soil are presented by analyte groups in Table ESA 2.1-1. The analyte groups detected in ESA soil samples include:

o Volatile halogenated organics (VHO);

o Volatile hydrocarbons (VHC);

o Volatile aromatic organics (VAO);

o Organosulfur compounds, mustard-agent related (OSCM);

o Organosulfur compounds, herbicide related (OSCH);

o Organophosphorus compounds, GB-agent related (OPHGB);

o Fluoroacetic acid;

o Polynuclear aromatic hydrocarbons (PAH);

o Semivolatile halogenated organics (SHO);

o Organochlorine pesticides (OCP);

o Arsenic;

o Mercury; and

o ICP metals (cadmium, chromium, copper, lead, and zinc).

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Four laboratories performed analyses on soil and water samples collected during the RI. The analytical and quality assurance techniques employed during certification of analytical methods in the separate laboratories led to the establishment of lower and upper limits of quantification that are method, analyte, laboratory, and instrument specific. Therefore, the reported CRL for a given sample may vary between samples. Lower CRLs for analytes detected during the RI fall within a range established by the most and least sensitive methods from among the four laboratories. This CRL range is presented for each analyte in soil samples in Table ESA 2.1-2. Data values falling below their respective CRLs are reported as below CRL (BCRL). The most and least sensitive methods among the four laboratories also define a range of upper CRLs; however, in some cases it was possible to report a value greater than the upper CRL while maintaining the USATHAMA quality of the data. To accomplish this, samples with higher contaminant concentrations were diluted so that the instrument reading for the diluted sample fell within the upper CRL range, and only exceeded this range when the dilution factor was applied to arrive at the final result. Data for these diluted samples represent the only values above the upper CRL that can be reported with the degree of accuracy and precision required by USATHAMA.

In cases where analytical values exceeded the upper CRL and the samples were not diluted, an attempt was made to recover the actual instrument readings from the laboratory. This information was used to gain a qualitative understanding of the relative level of contamination in the samples. These results that were above the upper CRL are not of USATHAMA quality and cannot be used with the same confidence as those falling within the quantification limits.

To provide a complete review of information pertinent to the contamination assessment at RMA, data from investigations other than the RI have been included in this and previous reports where appropriate, even though the methods used were not USATHAMA certified. This use has set a precedent to consider non-USATHAMA quality data along with the USATHAMA quality results obtained under the RI.

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The inclusion of non-USATHAMA quality data was also done in consideration of the fact that sophisticated statistical manipulations of the data would not be carried out for the purposes of the SARs. Such manipulations were deemed inappropriate considering the large numbers of values that fell below the various CRLs as well as those values (estimated to be approximately one percent of the data set) that fell above the upper CRLs and were not the result of approved dilution procedures. It was determined that statistical manipulations of more limited data sets, such as those obtained from a single site, would be more appropriate and accurate for the FS. In the FS phase, more limited statistical evaluations could then be used to evaluate, on a case-by-case basis, the effects of the non-USATHAMA quality or out-of-range data on the contamination assessment and on the evaluation and selection of appropriate remediai measures.

2.1.2 Analytical Results

In order to present the analytical data in a summary form for this report, contaminant distribution maps are constructed by analyte group rather than by individual analyte. Analyte groups reflect similar chemical composition, physical/chemical properties, and origins; and are discussed in generally decreasing order of mobility. Data were grouped together in continuous depth intervals which encompass the standard sampling intervals. To further simplify the presentation of the analyte levels, ranges of concentrations are shown by dots of different sizes.

Separate maps were used to represent the results for soil samples falling into the 0 to 2, 2 to 5, 5 to 20, and greater than 20 ft depth intervals. Composite samples from the 0 to 1 and 4 to 5 ft intervals are shown on both the 0 to 2 and 2 to 5 ft interval maps. Because the depth intervals used in analytic distribution maps may encompass more than one sample interval, one dot may represent more than one sample from a given soil boring. A single dot may also represent from one to all of the analytes detected in an analyte group at a given location. For these reasons, the number of dots on a map may be less than the number of samples analyzed.

For the organic compounds, the soils data reported for each boring were separated into the depth intervals used in the maps. Next, the data were separated into analyte groups. To compute the total concentration of each group in the boring, the highest concentration of each analyte in the group was summed. The resulting sum is equal to or greater than the total concentration of that group in any one sample. This worst-case concentration is represented on the map by the dot size corresponding to the concentration range attained.

The concentrations for each analyte group were divided into several ranges, and represented by progressively larger dots. The concentration ranges were based upon the following criteria:

Organics: (a) Lowest CRL to 1.0 μ g/g.

- (b) 1.0 μ g/g to next order of magnitude.
- (c) Subsequent ranges based on orders of magnitude, not to exceed four ranges per map.
- Inorganics: (a) Upper end of indicator range (i.e., the indicator level) to nearest order of magnitude.
 - (b) Subsequent ranges increase by a single order of magnitude above the previous concentration range.

Organic compounds associated with RMA activity are not naturally occurring, so detections were generally considered to be of environmental significance. For this reason, the indicator level for these compounds was considered to be the individual CRL. The metals are naturally occurring, so an indicator range of expected natural concentrations was established. Details on the selection of indicator ranges may be found in the Introduction to the CARs (ESE, 1987a/RIC 88204R02). The indicator ranges for metals reflect the concentrations expected to occur naturally in RMA alluvial soils. The upper limits of these ranges are:

Metal Concentration (µg/g) Arsenic10 Cadmium2.0 Chromium40 Copper 35 Lead 40 Mercury0.10 Zinc 80

Arsenic and mercury are mapped individually. The ICP metals, cadmium, chromium, copper, lead, and zinc, are mapped as a group.

Each ICP metal has a different indicator range and therefore cannot be compared directly on the basis of absolute concentration. For this reason, and because these metals tended to occur together in the ESA, the ICP concentrations are not added together to give a total for plotting. Instead, they were compared to their indicator ranges and assigned to a relative range. The highest range attained by any one ICP metal in a given depth interval is represented on the map. The indicator ranges for each metal are listed under Range 1. The metal specific concentration ranges in $\mu g/g$, are based on the indicator ranges, as follows:

<u>Metal</u>	<u>Range 1</u>	Range 2	Range 3	Range 4
Cadmium	BCRL-2.0	2.0-10	10-100	>100
Chromium	BCRL-40	40-100	100-1,000	> 1,000
Copper	BCRL-35	35-100	100-1,000	> 1,000
Lead	BCRL-40	40-100	100-1,000	> 1,000
Zinc	BCRL-80	80-1,000	1,000-10,000	> 10,000

Because metals concentrations at or below the upper limits of the indicator ranges are considered to be consistent with natural conditions, results that do not exceed these indicator ranges are shown on the analyte distribution maps as open circles, like BCRL results. Values above the indicator ranges are shown as solid dots. The significance of

the metals concentrations greater than their indicator ranges is discussed further in the contamination assessment portion of this report (Section 3.0).

In addition to the presentation of these data on the analyte distribution map, a summary list of the compounds detected above the CRLs in ESA soils and sediments is presented by site group and by analyte group in Table ESA 2.1-2. These results represent the sample concentrations of each analyte, less any concentration detected in the method blanks.

2.1.3 Distribution of Analytes

This section discusses the occurrence of target and significant nontarget compounds by analyte group in ESA soil.

2.1.3.1 Volatile Halogenated Organics (VHOs)

In the ESA, 14 of a total of 192 soil samples analyzed contained concentrations of at least one of the VHO compounds above the CRL. Detections ranged from 0 to 30 ft in depth, and 0.3 to 4 μ g/g in concentration. The six compounds of the VHO group present were chloroform, 1,1,1-trichloroethane, 1,1,2,2-tetrachloroethane, trichloroethylene, tetrachloroethylene, and trichloropropene. Methylene chloride was also detected in the ESA, but is discussed separately due to its potential occurrence as a laboratory artifact. Figures ESA 2.1-1 through 2.1-4 illustrate the location, depth ranges, and concentration ranges of VHO detections in the study area.

There are four site groups that contained detections of VHOs in the ESA. ESA-2, the burial trench sites, had one detection in the 2 to 5 ft interval, five detections in the 5 to 20 ft interval, and two detections in the 20 ft or greater interval. These occurred in the refuse cells of ESA-2b, the sanitary landfill (Site 30-4), and in ESA-2a, the Section 32 burn pits (Sites 32-5 and 32-6). The toxic storage sites had one detection in the 0 to 2 ft interval, two detections in the 2 to 5 ft interval, and one in the 5 to 20 ft interval. The tentative identification of 1,1,2,2-tetrachloroethane detected in the 0 to 2 ft interval of ESA-3c, the Section 31 toxic storage yard (Site 31-4), was a suspected misidentification

because this volatile compound was not expected to remain in surface soil for an extended period of time. In the balance of investigations, the nontarget compound trichloropropene was detected in the 0 to 2 ft interval from Section 25. Volatile compounds were not analyzed in this sample.

In the ESA, seven of a total of 162 soil samples analyzed for methylene chloride indicated concentrations above the CRL. Detections ranged from 4 to 33 ft in depth, and were all $1 \mu g/g$ in concentration. Figures ESA 2.1-5 through 2.1-7 illustrate the location, depth ranges, and concentration ranges of methylene chloride detections in the study area.

All detections of methylene chloride occurred in ESA-2b, the Section 30 sanitary landfill (Site 30-4). Four detections were noted in the 2 to 5 ft interval, one in the 5 to 20 ft interval, and two in the 20 ft or greater interval. The two detections in the 20 ft or greater interval occurred in the same boring, and are therefore represented by one dot. For five of the detections, the next deeper sampling interval was not analyzed for methylene chloride.

It should be noted that the detections of methylene chloride were all at a concentration of 1 μ g/g. Methylene chloride is a common laboratory contaminant.

2.1.3.2 Volatile Hydrocarbons and Related Compounds (VHCs)

In the ESA, 30 of a total of 1,120 soil samples analyzed for the VHC group indicated concentrations above the CRL. Detections ranged from 0 to 10 ft in depth, and 0.3 to 4 $\mu g/g$ in concentration. The two members of the group present were the nontarget compounds 2-butoxyethanol and 2,2-oxybisethanol. Figures ESA 2.1-8 through 2.1-10 illustrate the location, depth ranges, and concentration ranges of VHC detections in the ESA. Fifteen of the detections were from samples composited from the 0 to 1 and 4 to 5 ft intervals. These detections are shown on both the 0 to 2 and 2 to 5 ft interval dot maps. Two borings with detections in the 2 to 5 ft interval of ESA-1c, the Section 29 surface burn (Site 29-1), are within close proximity, and appear as one dot.

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Of the 30 VHC detections in the study area, 28 were 2,2-oxybisethanol.

2,2-oxybisethanol is more commonly known as diethylene glycol, a compound used widely in antifreeze. This compound was detected in the following depths and sites: 0 to 10 ft in ESA-1c and ESA-1d, the Section 29 and 30 burn sites (Sites 29-1 and 30-2); 0 to 5 ft in ESA-4a, the Section 30 impact area (Site 30-1); and 0 to 5 ft in the nonsource areas of Section 29 and 30. These detections are suspected field procedure contaminants, because during cold weather, antifreeze was put in the steam cleaner overnight and bled out of the lines before use. The steam cleaner was used to decontaminate equipment between samples. It is strongly suspected that this procedure caused drilling and sampling equipment to be contaminated with antifreeze, which was then transmitted to the samples. All Task 14 borings in Sections 19, 20, 29 and 30, where detections of 2,2oxybisethanol occurred were drilled during cold weather.

The two remaining VHC detections in the ESA occurred in one boring in ESA-3b, the Section 6 toxic storage yard (Site 6-6). The compound 2-butoxyethanol was detected in the 0 to 2 ft interval at 0.3 μ g/g, and in the 2 to 5 ft interval at 0.3 μ g/g, the lowest concentrations reported for the group. Samples were not obtained from this boring below the 2 to 5 ft interval.

2.1.3.3 Volatile Aromatic Organic Compounds (VAOs)

In the ESA, 16 of a total of 196 soil samples analyzed contained concentrations of at least one of the VAO compounds above the CRL. Detections ranged from 0 to 30.2 ft in depth, and 0.1 to 5 μ g/g in concentration. The four members of the group present were benzene, toluene, o-xylene, p-xylene, and m-xylene. Figures ESA 2.1-11 through 2.1-14 illustrate the location, depth ranges, and concentration ranges of VAO detections in the ESA. There were three detections in the 2 to 5 ft interval at the sanitary landfill. For one boring in both the Section 30 storage shed plots and the Section 32 burn pits, two detections occurred in the 5 to 20 ft interval. These detections are represented by a single dot at each site at that depth interval.

The highest and lowest concentrations detected were 5 $\mu g/g$ and 0.14 $\mu g/g$, respectively, in the 2 to 5 ft interval from ESA-2b, the sanitary landfill (Site 30-4). Of the 16 VAO detections, six were not analyzed for the same compound in the next interval. One detection of xylene from the 0 to 2 ft interval of ESA-1a, the Section 19 surface burn (Site 19-1), and one detection of toluene from the 0 to 2 ft interval of the Section 20 nonsource area were target analytes identified in nontarget screening.

There are four site groups that contained soil sample detections of VAOs in the ESA. ESA-1, the surface burn sites, had one detection in the 0 to 2 ft interval. ESA-2, the burial trench sites, had five detections in the 2 to 5 ft interval, four detections in the 5 to 20 ft interval, and two detections in the 20 ft or greater interval. ESA-3, the toxic storage sites, contained one detection in the 2 to 5 ft interval and two detections in the 5 to 20 ft interval. The balance of investigations had one detection in the 0 to 2 ft interval of Section 20.

2.1.3.4 Organosulfur Compounds, Mustard-Agent Related (OSCMs)

In the ESA, 20 of a total of 1,138 soil samples analyzed for the OSCM group indicated concentrations above the CRL. Detections ranged from 0 to 10 ft in depth and 0.6 to 330 μ g/g in concentration. The four compounds of the OSCM group detected in the study area were chloroacetic acid, dithiane, 1,4-oxathiane, and thiodiglycol. Figures ESA 2.1-15 through 2.1-18 illustrate the location, depth ranges, and concentration ranges of OSCM detections in this study area. In ESA-3a, the Section 5 storage yard (Site 5-2), and the northwest corner of ESA-3d, the Section 31 storage shed plots (Site 31-7), multiple detections in the 0 to 2 ft interval are represented by what appears to be single dots. This is due to their close proximity. For this reason and the concentration summing protocol, only nine dots representing detections appear in Figures ESA 2.1-15 to 2.1-18.

All OSCM detections in the ESA were in the toxic storage sites. Twelve detections, including the highest concentration, were noted in the 0 to 2 ft interval; four in the 2 to 5 ft interval; and three in the 5 to 20 ft interval. One detection of chloroacetic acid in

the 5 to 20 ft interval of ESA-3a, the Section 5 toxic storage yard (Site 5-2), at a concentration of 120 μ g/g, was in a saturated sample. Samples were not collected below that depth. For all the detections, the compounds detected in the 2 to 5 and 5 to 20 ft depth intervals were not analyzed for in the next sampling interval. The frequency of detections diminish with depth, and detections are located in areas where potential leaks from agent-filled bombs occurred.

2.1.3.5 Organosulfur Compounds, Herbicide Related (OSCHs)

In the ESA, four of a total of 1,138 soil samples analyzed for the OSCH group contained concentrations above the CRL. Detections ranged from 0 to 20 ft in depth, and 0.5 to 9 μ g/g in concentration. The three members of the group present were benzothiazole, chlorophenylmethyl sulfone, and chlorophenylmethyl sulfoxide. Figures ESA 2.1-19 through 2.1-22 illustrate the location, depth ranges, and concentration ranges of OSCH detections in the study area. The dot in the 5 to 20 ft interval (Figure ESA 2.1-21) represents two detections within that interval.

One detection of benzothiazole was tentatively identified in the nontarget screening. The sample was collected from the 0 to 1 ft interval, at a concentration of $0.5 \mu g/g$, in a boring from ESA-1a, the Section 19 burn site (Site 19-1). Phase II samples collected within 50 ft of this boring did not detect benzothiazole even though more sensitive analysis methods were employed. One detection of chlorophenylmethyl sulfoxide was noted in the 0 to 2 ft interval in ESA-3d, the Section 31 toxic storage plots (Site 31-7). The compound was detected on the edge of storage plot 23, in a sample that also contained a mustard degradation product. Two detections of chlorophenylmethyl sulfone occurred in samples collected from one boring in the 5 to 20 ft interval in ESA-2b, the sanitary landfill (Site 30-4). A geophysical survey identified a sanitary landfill waste cell at this boring location.

2.1.3.6 Organophosphorus Compounds, GB-Agent Related (OPHGBs) In the ESA, one of a total of 716 soil samples analyzed for the OPHGB group indicated a concentration above the CRL. The single compound detected was isopropylmethyl

phosphonic acid at a depth of 9 to 10 ft and a concentration of 47 μ g/g. The sample was located in ESA-3c, the Section 31 toxic storage yard (Site 31-4), next to a concrete pad used for demilitarization of VX-filled spray tanks. The concrete pad is west of and isolated from the primary storage yard. Samples analyzed from the intervals above and below this detection tested below the CRL for isopropylmethyl phosphonic acid. A 1973 report suggested that VX was spilled on the ground at this site, although the exact location was not verified (Blackwell, 1973/RIC 81339R20). Figures ESA 2.1-23 and 2.1-24 represent the location of the boring containing isopropylmethyl phosphonic acid in the study area.

2.1.3.7 Fluoroacetic Acid

In the ESA, five of a total of 50 soil samples analyzed for fluoroacetic acid indicated concentrations above the CRL. Detections ranged from 0 to 5 ft in depth and 2.7 to 19 μ g/g in concentration. Figures ESA 2.1-25 through ESA 2.1-27 illustrate the location, depth ranges, and concentration ranges of fluoroacetic acid detections in the study area.

The highest concentration of fluoroacetic acid was noted in one surface sample taken from a man-made mound in ESA-2c, the Section 30 open trenches (Site 30-6). Two detections were noted in samples from trenches in the 2 to 5 ft interval at ESA-4a, the Section 30 impact area (Site 30-1). ESA-5, the demilitarization activity site (Site 30-5), contained one detection in a sample within the 0 to 2 ft interval, next to a building where fuzes were removed from agent-filled bombs. Although not documented, it is conceivable that these bombs leaked during the fuze removal process. The lowest concentration of fluoroacetic acid detected in the study area was in a sample from the 0 to 2 ft interval in the nonsource area investigation of Section 25. This sample was collected from a drainage ditch leading from the North Plants Study Area. For all samples that detected fluoroacetic acid in the ESA, the next deepest sample analyzed was below the CRL for the compound.

2.1.3.8 Polynuclear Aromatic Hydrocarbons (PAHs) In the ESA, six soil samples detected the PAH group in nontarget GC/MS screening.

Detections were only in the 0 to 1 ft sampling interval, and ranged from 0.5 to 20 μ g/g in concentration. PAH detections occurred in ESA-1a, 1b, and 1c, the Section 19, 20, and 29 burn sites (Sites 19-1, 20-1, and 29-1). The two members of the group present were methylnaphthalene and phenanthrene, which are significant nontarget compounds. Figures ESA 2.1-28 and 2.1-29 illustrate the location, depth ranges, and concentration ranges of PAH detections in the study area. The larger dot in the 0 to 2 ft interval of the Section 29 surface burn represents two detections.

2.1.3.9 Semivolatile Halogenated Organics (SHOs)

In the ESA, one of a total of 1,106 soil samples analyzed for the SHO group contained concentrations above the CRL of 0.3 μ g/g. Hexachlorobenzene, a nontarget compound, was detected in a surface sample from the Section 20 nonsource area investigation at a concentration of 0.5 μ g/g. Hexachlorobenzene is also known as "Bunt-cure," a common seed and soil fungicide, which may be related to crops planted in Section 20. Further information may be found in the Contamination Assessment Report for Section 20. Nonsource Area (ESE, May 1988k/RIC 88173R06).

2.1.3.10 Organochlorine Pesticides (OCPs)

In the ESA, four of a total of 1,106 soil samples analyzed for the OCP group detected concentrations above the CRL. Detections ranged from 4 to 15 ft in depth and 0.003 to 20 μ g/g in concentration. The two members of the group present were aldrin and dieldrin. Figures ESA 2.1-30 through 2.1-33 illustrate the location, depth ranges, and concentration ranges of OCP detections in the study area. In the 5 to 20 ft depth interval, one dot in the sanitary landfill represents three detections.

ESA-2b, the sanitary landfill (Site 30-4), and ESA-2c, the Section 30 open trenches (Site 30-6), had the only OCP soil sample detections in the ESA. There was one detection of dieldrin in the 2 to 5 ft interval of a boring adjacent to the open trenches and two of aldrin and one of dieldrin in the 5 to 20 ft interval of the sanitary landfill. Of the two samples from the landfill, the one containing aldrin was not analyzed for the same compound in the next sampling interval.
2.1.3.11 Arsenic

In the ESA, 18 of a total of 1,142 soil samples analyzed for arsenic contained concentrations above the upper indicator level of 10 μ g/g. Detections ranged from 0 to 20 ft in depth and 11 to 270 μ g/g in concentration. Figures ESA 2.1-34 through 2.1-37 illustrate the location, depth ranges, and concentration ranges of arsenic detections above the indicator range in the study area.

Of the detections above its indicator range, the highest concentrations of arsenic, falling within the 100 to 1,000 μ g/g dot size range, were noted in two surface samples collected from ESA-3b, the Section 6 toxic storage yard (Site 6-6). Three samples in this site had concentrations in the 10 to 100 μ g/g range. Other detections in designated sites within the 10 to 100 μ g/g range were: one each for ESA-1c and 1d, the Section 29 and 30 surface burns (Sites 29-1 and 30-2); two in ESA-4b, the Section 29 demolition area (Site 29-4); and one in ESA-2c, Section 30 open trenches (Site 30-6). These concentration ranges for one sample in Section 20, five samples in Section 25, and two samples in Section 30 were also detected during the nonsource area investigations.

2.1.3.12 Mercury

In the ESA, 15 of a total of 1,111 soil samples analyzed for mercury contained concentrations above the upper indicator range of 0.10 μ g/g. Detections ranged from 0 to 20 ft in depth and 0.11 to 0.81 μ g/g in concentration. Figures ESA 2.1-38 through 2.1-41 illustrate the location, depth ranges, and concentration ranges of mercury detections above the indicator range in the study area.

Of the mercury detections above its indicator range, the highest concentrations were noted in two samples collected from ESA-2b, the sanitary landfill (Site 30-4). These occurred in the 0.10 to 1.0 μ g/g range. Other detections at this concentration range were: two in ESA-1c, the Section 29 surface burn (Site 29-1); four in ESA-4b, the Section 29 demolition area (Site 29-4); and one in the Section 30 open trenches (Site 30- δ). The balance of investigations showed this concentration range for one sample in Section 25; four samples in Section 30 (former Site 30-3); and one sample in Section 31.

2.1.3.13 ICP Metals

In the ESA, 150 of a total of 1,188 soil samples analyzed for ICP metals contained concentrations above the indicator level. All of the ICP metals, cadmium, chromium, copper, lead, and zinc, were detected. Detections ranged from 0 to 40 ft in depth, and the maximum concentration detected was 57,000 μ g/g of zinc. Figures ESA 2.1-42 through 2.1-45 illustrate the location, depth ranges, and concentration ranges of ICP metal detections above the upper indicator level in the ESA.

As illustrated by the figures, ICP metals were detected above the upper indicator level in all the ESA site groups. Many of the detections were only slightly above the upper indicator level. However, ESA-2a, the Section 32 burn pits, had the highest concentrations of ICPs in the study area. Concentrations of lead and zinc were as high as 3,400 μ g/g and 57,000 μ g/g in the 0 to 2 ft interval, and 2,100 μ g/g and 15,000 μ g/g in the 5 to 10 ft interval of this site. The lower depth corresponds to the base of the burning pit. Surficial samples from the surface burns in Sections 20, 29, and 30 and the Section 31 storage shed plots also contained high concentrations of ICP inetals.

There were seven ICP metal samples which were well above their upper indicator levels, but were still within the next concentration range. The majority of these samples were collected from the 0 to 2 ft depth interval, and occurred in the following sites: ESA-2a, the Section 32 burn pits (Site 32-6); ESA-2b, the sanitary landfill (Site 30-4); ESA-3c, the Section 31 storage yard (Site 31-4); ESA-3d, the Section 31 storage shed plots (Site 31-6); and an area in the southwest corner of Section 29 (Site 29-5). The remaining high concentration samples were collected from the 5 to 20 ft depth interval, and occurred in the following two sites: a location in Section 30 east of ESA-5, the bomb demilitarization site (Site 30-5); and ESA-1c, the Section 29 surface burn (Site 29-1).

2.1.3.14 Compounds Not Detected

The following compounds were not detected in the soil sampling program at the ESA:

organophosphorus compounds, pesticide related, and DBCP. Organophosphorus compounds were analyzed in 1,048 samples and DBCP was analyzed in 1,122 samples.

2.2 SURFACE WATER CONTAMINANTS

Samples of surface water were collected from seven locations along First Creek (Figure ESA 2.2-1). Sampling was conducted quarterly, from October 1985 to September 1987, as part of the Task 4 and 44 surface and groundwater quality screening programs. Complete surface water analytical data is presented in the Water RI Report (Ebasco, 1989/RIC 89067R08). Both this section, concerning surface water, and the following section, concerning groundwater, summarize the water quality results from Task 4 (ESE, 1988k/RIC 88173R06; ESE, 1987b/RIC 87253R01), and Task 44 (ESE, 1988n/RIC 88244R02). The groundwater discussion also includes information from Task 25 (ESE, 1988bb/RIC 89024R02). Locations of surface water sampling sites were chosen to monitor specific regions of the ESA. Station 8-001 monitors water quality entering the ESA, and Station 24-002 monitors it leaving the ESA. Station 24-007 monitors for the influence of potential runoff from the sewage treatment plant in Section 24. Station 30-002 was located to monitor the influence of surface water entering First Creek from the North Plants complex. Station 31-002 monitors water quality upstream of the Section 31 toxic storage sites, and Station 31-001 checks it downstream. Station 5-001 provides monitoring of water quality in the southern portion of Section 5.

2.2.1 Analytical Methods and Detection Limits

During the two years of surface water and groundwater quality sampling used for this report, the analytical suite expanded from 24 to 66 compounds. This was in response to the ongoing water quality monitoring programs and input from the Army, USEPA, Shell Oil Company, and the State of Colorado. From October 1985 to September 1986, the Task 4 analytical suite was used. From October 1986 to September 1987, the Task 44 analytical suite was used. These suites were originally designed to screen for historically detected contaminants found in the ESA. The site specific sampling onducted in Section 31 and 32 during January 1989 sampled a unique suite of analytes based on compounds detected in nearby soil samples.

Samples collected from October 1985 to September 1986 as part of Task 4 were originally analyzed for 24 target compounds. The analyte list was modified at several stages and at the end of the program consisted of 50 compounds. All analytical methods used were quantitatively certified according to USATHAMA methods to obtain low CRLs and high accuracy. The target compounds, analytical methods, and laboratory CRLs for the final analyte list of Task 4 are presented in Table ESA 2.2-1.

Samples collected from October 1986 to September 1987 as part of Task 44 were analyzed for the same fifty compounds used in Task 4, plus benzothiazole and chlordane. Semiquantitative GC/MS methods were used for an additional list of 42 compounds, 27 of which were also included in the quantitative screening. DBCP was analyzed by both quantitative and semi-quantitative methods. The Task 44 analytical program was applied to both surface and groundwater and was adapted into the Task 25 program, which monitored wells in Section 24 of the ESA. The target compounds, analytical methods, and laboratory CRLs for the final analyte list of Task 44 are presented in Table ESA 2.2-2.

The analytical suite proposed in Tasks 4, 44 and 25 in technical plans varied some that in surface water and groundwater samples collected in the ESA. The specific list of compounds analyzed for in a given sample may be obtained in the reports for Tasks 44 and 25 (ESE, 1988k/RIC 88173R06; ESE, 1987b/RIC 87253R01; ESE, 1988n/RIC 88244R02; and ESE, 1983dd/RIC 89024R02).

2.2.2 Analytical Results

A summary list of the target and significant nontarget compounds detected above the CRL in surface water samples is presented by analyte group in Table ESA 2.2-3. The associated contaminant distribution maps for surface water are presented in Figures ESA 2.2-2 through 2.2-6. The contaminant distribution maps use the dot format to present relative analyte concentrations for surface water and were constructed using the following procedures. No organic compounds were detected more than once at any

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station and ICP metals were detected more than once at only two stations. For a given station, the highest detection of an individual analyte noted during the sampling program was represented on the maps. Within a given organic analyte group, the detections of individual analytes were summed to reach the concentration range represented by the dots. For the ICP metals, the highest individual range attained by an analyte at a station was represented, and no summing procedure was used. Because of these procedures, the data used for the maps represent a worst-case scenario.

2.2.3 Distribution of Analytes

This section discusses the distribution of analytes detected in ESA surface water samples and describes data presented in Table 2.2-3. During the two years that surface water stations were sampled in the ESA, repeated detections of analytes were not common.

2.2.3.1 Volatile Halogenated Organics (VHOs)

In the ESA, chloroform and chlorobenzene were the only VHOs detected during the surface water sampling program. Chloroform was detected at Station 31-002, during the July to September 1986 sampling, at a concentration of $3.5 \ \mu g/l$. It was not detected during the previous sampling period, from April to June 1986; the station was not sampled during the following two fiscal year quarters, from October 1986 to March 1987. Chlorobenzene was detected at Station 31-001, during the April to June 1986 sampling, at a concentration of $1.8 \ \mu g/l$. It was not detected during the previous sampling period from October 1985 to March 1986; the station was not sampled during the following the station was not sampled during the following the station was not detected during the previous sampling period from October 1985 to March 1986; the station was not sampled during the following three quarters, from July 1986 to March 1987. A total of thirty-four surface water samples were collected from the ESA and analyzed for VHO compounds. Figure ESA 2.2-2 illustrates the location and concentration of VHO detections in the study area.

2.2.3.2 Organophosphorus Compounds, GB-Agent Related (OPHGBs)

In the ESA, diisopropylmethyl phosphonate was the only OPHGB compound detected during the surface water sampling program. The compound was detected at Stations 8-001 and 24-002, at concentrations of 11 and 17 μ g/l, during the initial sampling from October 1985 to March 1986. Station 8-001 was sampled during the following six

quarters, from April 1986 to September 1987; and Station 24-002 was sampled during the following quarter, from April to June 1986. Detections of disopropymethyl phosphonate were not repeated at these stations. A total of thirty-four surface water samples were collected from the ESA and analyzed for OPHGB compounds. Figure ESA 2.2-3 illustrates the location and concentration of OPHGB detections in the study area.

2.2.3.3 Organochlorine Pesticides (OCPs)

In the ESA, three detections of aldrin and two detections of dieldrin were the only OCPs noted during the surface water sampling program, as follows:

<u>Compound</u>	<u>Station</u>	Detections/ Samples	Concentration (µg/l)0	Date f Sampling
Aldrin	8-001	1/10	0.20	12/85
Dieldrin	8-001	1/10	0.060	11/85
Aldrin	24-002	1/6	0.20	12/85
Dieldrin	24-002	1/6	0.080	11/85
Aldrin	31-001	1/4	0.080	12/85

Aldrin and dieldrin were detected at Station 8-001, during the initial sampling from October 1985 to March 1986, at respective concentrations of 0.20 and 0.060 μ g/l. The detections were not repeated during the following six sampling quarters, from April 1986 to September 1987. Aldrin and dieldrin were also detected at Station 24-002, during the initial sampling from October 1985 to March 1986, at respective concentrations of 0.20 and 0.080 μ g/l. The detections were not repeated in the following sampling period, from April to June 1986. Finally, aldrin was detected at Station 31-001, during the initial sampling from October 1985 to March 1986, at a concentration of 0.080 μ g/l. The detection was not repeated in the following sampling quarter, from April to June 1986.

A total of thirty-four surface water samples were collected from the ESA and analyzed for OCP compounds. Figure ESA 2.2-4 illustrates the location and concentration of OCP detections in the study area.

2.2.3.4 Arsenic

In the ESA, five detections of arsenic were noted during the surface water sampling program, as follows:

<u>Compound</u>	<u>Station</u>	Detections/ <u>Samples</u>	Concentration $(\mu g/g)$	Date of <u>Sampling</u>
Arsenic	8-001	1/7	6.6	4/86
Arsenic	5-001	1/2	3.8	5/87
Arsenic	31-002	1/2	7.3	5/87
Arsenic	24-007	1/1	5.0	5/87
Arsenic	24-002	1/2	3.5	6/87

One detection occurred at Station 8-001 during the sampling from April to June 1986 at a concentration of 6.6 μ g/l. Arsenic was not detected at this station during the previous sampling, from October 1985 to March 1986; nor during the following five sampling quarters from July 1986 to September 1987. Detections of arsenic occurred in Stations 5-001 and 31-002 during the sampling from April to June 1987, at respective concentrations of 3.8 and 7.3 μ g/l. These stations were not sampled during the previous two quarters from October 1986 to March 1987; they were sampled without arsenic detections during the following quarter from July to September 1987. A detection occurred at Station 24-007 during the April to June 1987 sampling, at a concentration of 5.0 μ g/l. This station was not sampled at any other time during the program. Finally, an arsenic detection was noted at Station 24-002 during the April to June 1987 sampling at a concentration of 3.5 μ g/l.

A total of twenty surface water samples were collected from the ESA and analyzed for arsenic. None of the sample stations contained multiple arsenic detections. Figure ESA 2.2-5 illustrates the location and concentration of arsenic detections in the study area.

2.2.3.5 ICP Metals

Zinc was detected at sample stations located near the east and north boundaries of the ESA. For multiple detections, the highest concentration and its sampling date are presented in this summary. Cadmium and copper were detected at Station 8-001 during

the January to March 1987 sampling period, at respective concentrations of 14 and 21 $\mu g/l$. These compounds were not detected at this station during the previous five sampling quarters from October 1985 to December 1986; nor during the following two sampling quarters from April to September 1987. Chromium and lead were also detected at Station 8-001 during the final sampling from July to September 1987, at respective concentrations of 13 and 22 μ g/l. These detections were not repeated during the previous seven sampling quarters from October 1985 to June 1987. Zinc was detected twice at Station 8-001, first during the April to June 1986 sampling at a concentration of 24 μ g/l and second during the October to December 1986 sampling at a concentration of 25 μ g/l. Zinc was not detected during the intervening sampling period from July to September 1986 nor during the previous period from October 1985 to March 1986. Zinc was not detected during the following three quarters from January to September 1987. Zinc was detected twice at Station 24-002 in two sampling events conducted during April to June 1986, at concentrations of 41 and 30 μ g/l. It was not detected during the previous sampling from October 1985 to March 1986; and the station was not sampled during the following two quarters from July to December 1986. Finally, copper was detected at Station 31-001 during the final sampling from July to September 1987 at a concentration of 10 μ g/l. It was not detected during the previous sampling quarter from April to June 1987. A total of twenty surface water samples were collected from the ESA and analyzed for ICP metals. Figure ESA 2.2-6 illustrates the location and concentration of ICP metal detections in the study area, summarized as follows:

Compound	E <u>Station</u>	Detections/ Samples	Highest Concentration (µg/l)	Date of Sampling
Cadmium	08-001 (08ADD)	1/7	14	3/87
Copper	08-001 (08ADD)	1/7	21	3/87
Chromium	08-001 (08ADD)	1/7	13	10/87
Lead	08-001 (08ADD)	1/7	22	10/87
Zinc	08-001 (08ADD)	2/7	- 25	12/86
Zinc	24-002 (13DCC)	2/3	41	6/87
Copper	31-001	1/2	10	10/87

2.2.3.6 Compound Groups Not Detected

The following compound groups were analyzed for but not detected in the ESA surface water sampling program: VHCs. VAOs, OSCMs, OSCHs, OPHPs, DBCP, SHOs, and mercury. Surface water samples were not analyzed for fluoroacetic acid.

2.3 GROUNDWATER CONTAMINANTS

Samples of groundwater were collected from two distinct hydrologic systems in the ESA, the alluvial aquifer and Denver Formation groundwater. Table ESA 2.3-1 summarizes the detections of target and significant nontarget compounds, by individual analyte, in both hydrologic systems. Table ESA 2.3-2 presents the analytes detected in all sampling events by individual well, for alluvial groundwater samples. Table ESA 2.3-3 presents the analytes detected in all sampling events by individual well, for alluvial groundwater samples. Table ESA 2.3-3 presents the analytes detected in all sampling events by individual well, for Denver Formation groundwater samples. This section will discuss the alluvial and Denver hydrologic systems separately.

2.3.1 Alluvial Aquifer Sampling Network

The sampling network for the alluvial aquifer consisted of the nineteen wells shown in Figure ESA 2.3-1. The first two digits of the well number represent the section, and the last three represent the individual well. The well network was chosen to provide information on compounds historically detected in groundwater in the ESA, as well as to provide overall hydrologic information on the study area. Historical water quality was derived from information summarized in the Water RI Report (Ebasco, 1989/RIC 89067R08). Wells 08002 and 08003 monitor water quality entering the ESA from offpost and were sampled to verify historical concentrations of OCPs and diisopropylmethyl phosphonic acid. Well 07001 is near the southern boundary of the ESA by the Highline Lateral, an area where OCPs were historically detected. Wells 06002 and 06003 were sampled to verify historical concentrations of OCPs and diisopropylmethyl phosphonic acid. Well 31005 is in the vicinity of First Creek and was sampled to verify historical concentration of diisopropylmethyl phosphonic acid and DBCP. Well 31005 provided water quality data for the west-central portion of the ESA. Well 32001 is generally

downgradient of ESA-2a, the Section 32 burn pits (Sites 32-5 and 32-6), and is also within a minor paleochannel which enters the ESA in Section 32. Wells 25011, 25038, and 30009 were sampled to provide verification of historical OCP, DBCP, and diisopropylmethyl phosphonic acid concentrations and to give an indication of general alluvial aquifer water quality in the northeastern portion of RMA. Wells 19001, 24107, 24166, 24183, and 24188 were sampled to provide information on conditions in the vicinity of the North Boundary Containment System. Finally, Wells 31014, 31015, and 31016 were sampled to specifically monitor conditions downgradient from the Section 31 Toxic Storage Area; and Well 32004 was sampled to specifically monitor conditions downgradient from the Section 32 Burn Pits. These wells were installed late in the RI to address site-specific monitoring needs. They were sampled once during January 1989. At the time of publication of this report, the data from these new wells had not completed final quality assurance/quality control procedures. The preliminary data from these wells are included in Appendix ESA-C to assist the characterization of contaminants in the alluvial aquifer. Further sampling of these wells is being carried out under the Comprehensive Monitoring Program.

The number of ESA alluvial aquifer wells sampled in the RI generally decreased during the life of the program. Ten wells were sampled from October 1985 to March 1986, seven wells were sampled from April to June 1986; nine wells were sampled from July to September 1986; three wells were sampled from October to December 1986 and from January to March 1987; thirteen wells were sampled from April to June 1987; three wells were sampled from July to September 1987; and five wells were sampled in January 1989. The reason for the general decrease in sampling over time was the lack of analyte detections. Table ESA 2.3-1 summarizes the detections that occurred in the alluvial aquifer water quality monitoring program.

The dot map format is used in the contaminant distribution maps to represent relative analyte concentrations in alluvial groundwater samples. These maps were constructed using the following procedures. Repeat detections of analytes were not common during the groundwater sampling program in the ESA. For a given well, the highest

concentration of an individual analyte detected in all sampling periods was represented on the maps. Within a given organic analyte group, the highest detections of individual analytes were summed to reach the concentration range represented by the dots. For the ICP metals, the highest individual range attained by an analyte at a well was represented, and no summing procedure was used. Because of these procedures, the data used for the maps represent a worst-case scenario.

2.3.2 Analytical Results of Alluvial Aquifer Sampling

A list of all the target compounds detected above the CRL in alluvial aquifer groundwater samples is presented by analyte group in Table ESA 2.3-2. A total of fiftytwo alluvial aquifer groundwater samples were collected and analyzed for target contaminants during the RI. The associated contaminant distribution maps for the alluvial aquifer are presented in Figures ESA 2.3-2 to 2.3-8. Due to the low frequency of repeat detections, the highest concentration from any sampling period is represented on the figures. Increasing dot sizes indicate increases in the magnitude of contaminant concentrations.

2.3.3 Distribution of Analytes in the Alluvial Aquifer

This section presents the analytes detected in the alluvial aquifer, and their frequency of occurrence. It summarizes data presented in Tables ESA 2.3-1 and 2.3-2.

2.3.3.1 Volatile Halogenated Organics (VHOs)

In the ESA, chlorobenzene, chloroform, and 1,2-dichloroethane were noted in the alluvial aquifer sampling program, as follows:

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Compound	<u>Well</u>	Detections/ Samples	Concentration(µg/1)	Date of Sampling
Chlorobenzene	06002	1/4	1.4	Sp 87
Chlorobenzene	07001	1/4	1.9	Sp 87
Chlorobenzene	08003	1/3	0.74	Sp 87
1,2- Dichloroethane	08003	1/3	0.64	Sp 87
Chloroform	24188	1/5	22	Wi 87
Chloroform	30009	1/4	60	Su 86
Chloroform	31014	1/1	0.58	Wi 89
Chloroform	31016	1/1	3.7	Wi 89

Chlorobenzene was detected in Wells 06002, 07001, and 08003 during the April to June 1987 sampling. Wells 06002 and 07001 were not sampled during the previous two periods from October 1986 to March 1987; Well 08003 was not sampled during the previous three quarter periods from July 1986 to March 1987. None of the three wells was sampled during the one quarter period following the detection, July to September 1987. 1,2-Dichloroethane was detected in Well 08003 during the April to June 1987 sampling. This well was not sampled during the three previous sampling periods from July 1986 to March 1987, nor during the following sampling period from July to September 1987. Chloroform was detected in Well 24188 during the January to March 1987 sampling. There were no detections of chloroform at this well during the previous two quarter periods from July to December 1986 or the following two quarter periods from April to September 1987. Chloroform was detected in Well 30009 during the July to September 1986 sampling. The well was sampled without any detections during the previous three quarter period from October 1985 to June 1986. The well was not sampled during the following two quarter period from October 1986 to March 1987. Chloroform was detected in Wells 31014 and 31016 during the site-specific sampling in January 1989. As noted in Section 2.3.1, data from these wells has not completed the

final quality assurance/quality control procedures, and is therefore considered preliminary. Figure ESA 2.3-2 illustrates the location and concentration of VHO detections in the study area.

2.3.3.2 Volatile Aromatic Organics (VAOs)

In the ESA, benzene and toluene were the only VAOs detected in the alluvial aquifer sampling program, as follows:

,		Detections/ Concentration Date of			
Compound	Well	Samples	<u>(µg/l)</u>	<u>Sampling</u>	
Benzene	06002	1/4	2.4	Sp 86	
Toluene	24188	1/5	8.6	Fa 86	

Benzene was detected in Well 06002 during the April to June, 1986 sampling. There were no detections of benzene during the previous two quarter periods from October 1985 to April 1986 nor the following one quarter period from July to September 1986. Toluene was detected in Well 24188 during the October to December 1986 sampling. The well was sampled with no detections of toluene during the previous one quarter period from July to September 1986 nor any during the following three quarter period from July to September 1986 nor any during the following three quarter period from January to September 1987. Figure ESA 2.3-3 illustrates the location and concentration of VAO detections in the study area.

2.3.3.3 Organosulfur Compounds Herbicide Related (OSCHs)

In the ESA, chlorophenylmethyl sulfone and chlorophenylmethyl sulfoxide were the only OSCHs detected during the alluvial aquifer sampling program, as follows:

Compound	Well	Detections/ Samples	Highest Concentration g/1)	Date of Sampling
Chlorophenyl- methyl sulfone	24183	4/5	3.7	Sp 87
Chlorophenyl- methyl sulfoxide	24183	1/5	6.1	Fa 86
Chlorophenyl- methyl sulfone	24188	3/5	4.3	Fa 86
ESAR\Sec2.Txt		2-3	30	

Chlorophenylmethyl sulfone was detected in Well 24183 four times in the four quarters of sampling from October 1986 to September 1987. The highest concentration of the four detections is reported above although all concentration values are reported in Table ESA 2.3-2. Chlorophenylmethyl sulfoxide was also detected in Well 24183 once during the sampling from October to December 1986. The well was sampled without any detections during the previous quarter from July to September 1986. Chlorophenylmethyl sulfone was detected in Well 24188 three times during the three quarters of sampling from October 1986 to June 1987. The well was sampled with no detections during the previous quarter from July to September 1986. The well was sampled with no OSCH detections during the following quarter from July to September 1987. Figure ESA 2.3-4 illustrates the location and concentration of OSCH detections in the study area.

2.3.3.4 DBCP

In the ESA, DBCP was detected once during the alluvial aquifer sampling program, as follows:

Compound	<u>Well</u>	Detections/ <u>Samples</u>	Concentration (µg/1)	Date of Sampling
DBCP	24183	1/5	0.16	Wi 87

DBCP was detected in Well 24183 during the January to March 1987 sampling. The well was sampled with no detection of DBCP detections during the two previous quarters from July to December 1986, and during the two following quarters from April to September 1987. Figure ESA 2.3-5 illustrates the location and concentration of the DBCP detection in the study area.

2.3.3.5 Organochlorine Pesticides (OCPs)

In the ESA, aldrin, dieldrin, and endrin were the only OCPs detected during the alluvial aquifer sampling program, as follows:

		Highest			
0	117-11	Detections/ C	oncentration	Date of	
Compound	wen	Samples	<u>(µg/1)</u>	Sampling	
Aldrin	32001	1/2	0.11	Wi 86	
Aldrin	31016	1/1	0.11	Jan 89	
Dieldrin	24166	5/5	0.16	Fa 86	
Dieldrin	24183	1/5	0.087	Su 87	
Dieldrin	31016	1/1	0.27	Jan 89	
Endrin	08003	1/3	0.074	Jan 86	
Endrin	31016	1/1	0.18	Jan 89	

Aldrin was detected in Well 32001 during the sampling from October 1985 to March 1986. This was the initial sampling period. There were no detections at this well during the following sampling quarter from April to June 1986. Dieldrin was detected in Well 24166 during the five sampling quarters in July 1986 to September 1987. The well was not sampled during the three previous sampling quarters from October 1985 to June 1986. Dieldrin was also detected in Well 24183 during the sampling period from July to September 1987. No dieldrin detections were reported for this well during the previous four sampling quarters from July 1986 to June 1987. Endrin was detected in Well 08003 during the initial sampling period from October 1985 to March 1986. No detections were noted during the following quarter from April to June 1986. Figure ESA 2.3-6 illustrates the location and concentration of OCP detections in the study area. Aldrin was detected in Well 31016 during the site-specific sampling in January 1989. Dieldrin was detected in Well 31016 during the site-specific sampling in January 1989. Endrin was detected in Well 31016 during the site-specific sampling in January 1989. As noted in Section 2.3.1, data from these wells has not completed the final quality assurance/quality control procedures, and is therefore considered preliminary.

2.3.3.6 Arsenic

In the ESA, arsenic was detected three times during the alluvial aquifer sampling program, as follows:

Compound	Well	Detections/ <u>Samples</u>	Concentration	Date of Sampling
Arsenic	24107	1/2	5.3	Su 86
Arsenic	24183	1/5	4.7	Su 87
Arsenic	24188	1/5	5.1	Su 87

Arsenic was detected in Well 24107 during the sampling from July to September 1986. The well was not sampled during the previous three quarters from October 1985 to June, 1986 nor during the following two sampling periods from October 1986 to March 1987. Arsenic was not detected during sampling of the well during April to June 1987. Arsenic was detected in Wells 24183 and 24188 during the sampling from July to September 1987. The analyte was not detected in either of the wells during the previous four sampling periods from July 1986 to June 1987. Figure ESA 2.3-7 illustrates the location and concentration of arsenic detections in the study area.

2.3.3.7 ICP Metals

In the ESA, all the ICP metals were detected during the alluvial aquifer sampling program, as follows:

Compound	<u>Well</u>	Detections/ Samples	Concentration	Date of Sampling
Zinc	06002	1/1	. 41	Sp 87
Chromium	06003	1/1	7.3	Sp 87
Zinc	06003	1/1	110	Sp 87
Cadmium	07001	1/1	12	Sp 87
Chromium	07001	1/1	37	Sp 87
Copper	07001	1/1	30	S p 87
Lead	07001	1/1	24	S p 87
Zinc	07001	1/1	97	S p 87
Chromium	08003	1/1	11	Sp 87
Chromium	19001	1/1	8	S p 87
Chromium	24107	1/1	26	Sp 87
Zinc	24107	1/1	93	Sp 87
Chromium	25011	1/1	12	Sp 87
Zinc	25038	1/1	53	Sp 87
Chromium	30009	1/1	16	Sp 87
Zinc	30009	1/1	140	Sp 87
Chromium	31005	1/1	23	Sp 87
Zinc	31005	1/1	43	Sp 87

ICP metals were detected only during the sampling from April to June 1987 in the following wells: 06002, 06003, 07001, 08003, 19001, 24107, 25011, 25038, 30009, and 31005. In each case this was the only time during the RI that the samples were analyzed for ICP metals. Figure ESA 2.3-8 illustrates the location and concentration of ICP metals in the study area.

2.3.3.8 Compound Groups Not Detected

The following compound groups were not detected in the alluvial aquifer sampling program in the ESA: VHCs, OSCMs, OPHGBs, OPHPs, DBCP, ONCs, PAHs, SHOs, and mercury. Groundwater samples were not analyzed for fluoroacetic acid.

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2.3.4 Denver Formation Groundwater Sampling Network

The sampling network for the Denver Formation groundwater consisted of the 37 wells shown in Figure ESA 2.3-9. Well locations were chosen to provide information on compounds detected historically in the ESA, as well as to provide information on ESA sites. Wells 07004, 07005, 08004, and 08005 monitored the quality of Denver Formation groundwater as it entered RMA from off-post. Wells 06004, 06005, and 05001 were sampled to verify historical concentrations of diisopropylmethyl phosphonic acid, and Wells 05002 and 05003 were sampled to provide information on water quality entering the ESA from the southeast. Wells 25012, 25013, 25014, 29002, 29003, 30004, 30005. 30010, 30011, 31006, 31007, 31008, 32002, 32003, 06004, 06005, 08004, 08005, 05001, 05002, and 05003 are cluster wells corresponding to alluvial aquifer monitoring wells. These wells provided information on aquifer interactions and any potentia will or groundwater contamination of the alluvial aquifer and Denver Formation groundwater. Wells 19003, 19015, 19016, 19017, 19018, and 19019 were sampled to monitor water quality at the northern border of the ESA. Well 31002 provided information on the western portion of the ESA and potential influence of groundwater from the Central Study Area. Wells 24108, 24109, 24120, 24174, and 24175 provided information on contaminant concentrations in the vicinity of the North Boundary Containment System.

The number of Denver Formation groundwater wells sampled in the RI in the ESA generally decreased during the life of the program. Thirty wells were sampled from October 1985 to March 1986. Sixteen wells were sampled from April to June 1986. Eighteen wells were sampled from July to September 1986. Five wells were sampled from October to December 1986 and from January to March 1987. Eighteen wells were sampled from April to June 1987. Finally, five wells were sampled from July to September 1987. The reason for the general decrease in sampling over time was the lack of analyte detections. Table ESA 2.3-2 summarizes the detections that occurred in the Denver Formation groundwater quality monitoring program.

2.3.5 <u>Analytical Results of Denver Formation Groundwater Sampling</u> A summary list of all the target and significant nontarget compounds detected above the

CRL is presented by analyte group in Table ESA 2.3-3. A total of forty-seven Denver Formation groundwater samples were obtained and analyzed for target contaminants during the RI. The associated contaminant distribution maps for the Denver Formation groundwater are presented in Figures ESA 2.3-10 to 2.3-16. The presentation strategy is similar to that for the alluvial aquifer data, discussed in Section 2.3.2. Detections of the same compound group in different members of a cluster well site resulted in a single dot on the contaminant distribution map.

2.3.6 Distribution of Analytes in Denver Formation Groundwater

This section presents the analytes detected in Denver Formation groundwater and their frequency of occurrence. It summarizes data presented in Table ESA 2.3-3.

2.3.6.1 Volatile Halogenated Organics (VHOs)

In the ESA, chlorobenzene, chloroform, methylene chloride, and trichloroethylene were detected in the Denver Formation groundwater sampling program, as follows:

Compound	<u>Well</u>	Detections/ (Highest Concentration (µg/1)	Date of <u>Sampling</u>
Chlorobenzene	24174	3/5	45	Su 86
Trichloroethylene	24174	1/5	2.4	Su 86
Chlorobenzene	24175	3/5	25	Su 86
Methylene Chloride	24175	1/5	4.3	Fa 86
Chloroform	30011	1/4	20	Su 86

Chlorobenzene was detected in Wells 24174 and 24175 during the one sampling period from July to September 1986 and the two last sampling periods from April to September 1987. The highest concentrations of multiple chlorobenzene detections are reported above, although all concentration values are reported in Table ESA 2.3-3. The wells were not sampled prior to the first detection and were sampled without chlorobenzene

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detections in the two sampling periods from October 1986 to March 1987. Trichloroethylene was also detected in Well 24174 during the sampling period from July to September 1986. No sampling was conducted prior to this, and trichloroethylene was not detected in the following four sampling periods, from October 1986 to September 1987. Methylene chloride was also detected in Well 24175 during the sampling period from July to September 1986. The well was sampled without methylene chloride detections in the prior quarter, April to June 1986 and during the three periods from January to September 1987. Chloroform was detected in Well 30011 during one sampling quarter from July to September 1986. No chloroform was detected in the prior three sampling quarters from October 1985 to June 1986. Well 30011 was not sampled in the following two quarters, from October 1986 to March 1987. Figure ESA 2.3-10 illustrates the location and concentration of the VHO detections in the study area, and Figure ESA 2.3-11 illustrates the methylene chloride detection.

2.3.6.2 Volatile Aromatic Organics (VAOs)

In the ESA, benzene was the only VAO detected in the Denver Formation groundwater sampling program. It was found in the following wells:

		Highest				
Compound	Well	Samples	$(\mu g/1)$	Sampling		
Benzene	06005	1/4	10	Wi 86		
Benzene	07004	1/4	8.9	Su 86		
Benzene	07005	2/3	4.2	Su 86		
Benzene	19015	1/4	7.3	Wi 86		
Benzene	19016	1/3	1.4	Wi 86		
Benzene	24174	2/5	20	Su 86		
Benzene	24175	2/5	4.5	Su 86		
Benzene	25040	1/1	2.1	Wi 86		
Benzene	32002	1/2	1.8	Wi 86		
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Benzene was detected in Wells 06005, 19015, and 19016 during the initial sampling period from October 1985 to March 1986. For these wells, benzene was not detected in the following two sampling periods from April to September 1986. Benzene was also detected in Wells 25040 and 32002 during the initial sampling period from October 1985 to March 1986. These wells were not sampled in the following four quarters from April 1986 to March 1987. Benzene was detected twice in Wells 24174 and 24175, first during the July to September 1986 sampling and second during the final sampling from July to September 1987. The highest concentration of multiple detections is reported above although all concentration values are reported in Table ESA 2.3-3. No benzene detections were noted during the intervening three sampling periods from October 1986 to June 1987. These wells were not sampled during the prior three periods, from October 1985 to June 1986. Benzene was detected in Well 07004 during the July to September 1986 sampling period. No detections were noted during the previous three sampling periods from October 1985 to June 1986. The well was not sampled in the following two quarters from October 1986 to March 1987. Finally, benzene was detected twice in Well 07005 during the two quarters from April to September 1986. No detections were noted in the previous two sampling periods from October 1985 to March 1986. The well was not sampled in the following four quarters from October 1986 to September 1987. Figure ESA 2.3-12 illustrates the location and concentration of benzene detections in the study area.

2.3.6.3 DBCP

In the ESA, DBCP was detected twice in the Denver Formation groundwater sampling program, as follows:

Compound	Well	Detections/ Samples	Concentration (µg/l)	Date of Sampling
DBCP	06005	1/4	0.75	Wi 86
DBCP	24174	1/5	0.15	Su 86

DBCP was detected in Well 06005 during the initial sampling from October 1985 to March 1986. There were no detections in the following two quarters from April to September 1986. It was also detected in Well 24174 during the sampling from July to September 1986. The well was not sampled during the previous three quarters from October 1985 to June 1986. DBCP was not detected in the following four quarters from October 1986 to September 1987. Figure ESA 2.3-13 illustrates the location and concentrations of DBCP detections in the study area.

2.3.6.4 Organochlorine Pesticides (OCPs)

In the ESA, the OCPs aldrin, endrin, and dieldrin were detected in the Denver Formation groundwater samples, as follows: Highest

<u>Compound</u>	Well	Detections/ Samples	Concentration (µg/l)	Date of <u>Sampling</u>
Aldrin	25014	1/4	0.46	Wi 86
Dieldrin	25014	1/4	0.42	Wi 86
Endrin	30005	1/2	0.12	Wi 86
Aldrin	31006	1/3	0.15	Wi 85
Dieldrin	19003	1/1	8.9	Sp 87
Endrin	19003	1/1	0.20	Sp 87
Dieldrin	19019	1/2	0.064	Wi 86
Endrin	19019	1/2	0.092	Wi 86
Dieldrin	24120	5/5	0.19	Fa 86
Aldrin	25013	1/4	0.17	Wi 86

Aldrin was detected in Wells 25013 and 31006, and aldrin and dieldrin were detected in Well 25014 during the initial sampling period from October 1985 to March 1986. There was no detection of these compounds in the following two sampling quarters from April

to September 1986. Dieldrin and endrin were detected in Well 19019, and endrin was detected in Well 30005 during the initial sampling from October, 1985 to March 1986. There were no detections during the following sampling quarter from April to June 1986. Dieldrin and endrin were detected in Well 19003 during the sampling from April to June 1987. This well was not sampled during any other quarter in this program. Dieldrin was detected five times in Well 24120 during the five quarters from July 1986 to September 1987. The highest concentration of the five detections is reported above, although all concentration values are reported in Table ESA 2.3-2. The well was not sampled during the previous three quarters from October 1985 to June 1986. Figure ESA 2.3-14 illustrates the location and concentration of OCP detections in the study area.

2.3.6.5 Arsenic

In the ESA, arsenic was detected ten times in the Denver Formation groundwater sampling program, as follows:

Compound	Well	Detections/ (Samples	Highest Concentration (µg/1)	Date of Sampling
Arsenic	06004	1/4	2.6	Sp 87
Arsenic	08004	1/1	5.9	Wi 86
Arsenic	08005	1/2	2.6	Sp 87
Arsenic	19015	1/4	6.8	Wi 86
Arsenic	19016	1/3	26	Wi 86
Arsenic	24109	1/5	8.6	Su 86
Arsenic	24174	2/5	3.9	Su 87
Arsenic	24175	1/5	4.1	Sp 87
Arsenic	32002	1/2	. 20	Wi 86

Arsenic was detected in Well 06004 during the April to June 1987 sampling. The well was not sampled during the previous two quarters from October 1986 to March 1987, nor during the following quarter from July to September 1987. Arsenic was detected in Well 08004 during the October 1985 to March 1986 sampling period. The well was not sampled in any other quarter. Arsenic was detected in Well 08005 during the April to June 1987 sampling. The well was not sampled during the previous four quarters from April 1986 to March 1987 nor during the following quarter from July to September 1987. Arsenic was detected in Wells 19015 and 19016 during the initial sampling from October 1985 to March 1986. It was not detected in the following two quarters from April to September 1986. Arsenic was detected in Well 24109 during the July to September 1986 quarter. This well was not sampled during the previous three quarters from October 1985 to June 1986. There were no detections during the following four sampling quarters from October 1986 to September 1987. Arsenic was detected twice in Well 24174, during the sampling from January to March 1987, and during the final sampling from July to September 1987. It was not detected in the intervening quarter from April to June 1987, nor was it detected in the previous two quarters from July to December 1986. Arsenic was detected in Well 24175 during the sampling from April to June 1987. It was not detected in the previous three sampling quarters from July 1986 to March 1987, nor in the following sampling quarter from July to September 1987. Finally, arsenic was detected in Well 32002 during the initial sampling period from October 1985 to March 1986. There were no detections of arsenic in the only other sample collected from this well, from April to June 1987. Figure ESA 2.3-15 illustrates the location and concentration of arsenic detections in the study area.

2.3.6.6 ICP Metals

In the ESA, the ICP metals chromium, copper, lead, and zinc were detected in the Denver Formation groundwater sampling program. Detections were as follows:

		Highest			
		Detections/	Concentration	Date of	
Compound	<u>Well</u>	<u>Samples</u>	<u>(µg/1)</u>	<u>Sampling</u>	
Chromium	05001	1/1	16	Sp 87	
Copper	05001	1/1	9.7	Sp 87	
Zinc	05001	1/1	74	Sp 87	
Zinc	08004	1/1	36	Wi 86	
Zinc	08005	1/2	42	Sp 87	
Chromium	19003	1/1	43	Sp 87	
Zinc	19003	1/1	60	Sp 87	
Chromium	19015	1/2	13	Sp 87	
Copper	19015	1/2	17	Sp 87	
Zinc	19015	1/2	43	Wi 86	
Zinc	19016	1/1	33	Wi 86	
Chromium	19017	1/1	13	Sp 87	
Zinc	19017	1/1	300	Sp 87	
Zinc	24120	1/1	35	Sp 87	
Zinc	25013	3/3	82	Sp 86	
Zinc	25039	2/2	23	Sp 87	
Zinc	25040	1/1	34	Wi 86	
Copper	31002	1/1	33	S p 86	
Lead	32002	1/1	65	Sp 87	
Zinc	32002	1/2	34	Sp 87	

Chromium, copper, and zinc were detected in Well 05001; zinc was detected in Well 08004; chromium and zinc were detected in Well 19017; and lead and zinc were detected in Well 32002 during the sampling from April to June 1987. For all these wells, ICP metals were only analyzed for during this sampling. Zinc was also detected in Well 08005 during the sampling period from April to June 1987. This well had also been sampled in the initial sampling period from October 1985 to March 1986, and zinc was not detected. Chromium and zinc were detected in Well 19003 during the April to June

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1987 period. The well was not sampled at any other time in this program. Zinc was detected in Wells 19015 and 19016 during the initial sampling from October 1985 to March 1986. ICP metals were not analyzed for during the following two quarters from April to September 1986. Chromium and copper were also detected in Well 19015 during the sampling from April to June 1987. This well was not sampled during the previous two quarters from October 1986 to March 1987 nor was it sampled during the following quarter from July to September 1987. Zinc was detected in Well 24120 during the sampling from April to June 1987. ICP metals were not analyzed for during the previous three sampling quarters from July 1986 to March 1987 nor during the following sampling quarter from July to September 1987. Zinc was detected three times in Well 25013, twice during the sampling from April to September 1986 and once during the sampling from April to June 1987. Samples from this well were not analyzed for ICP metals during the intervening two quarters from October 1986 to March 1987, nor during the following quarter from July to September 1987. ICP metals were not analyzed for in Well 25013 during the initial sampling from October 1985 to March 1986. Zinc was detected twice in Well 25039, once during the initial sampling from October 1985 to March 1986, and once during the sampling from April to June 1987. The well was not sampled at any other time during the program. Finally, zinc was detected in Well 25040 during the initial sampling from October 1985 to March 1986 and copper was detected in Well 31002 during the April to June 1986 sampling. These wells were not sampled at any other time during the program. Figure ESA 2.3-16 illustrates the location and concentration of ICP metal detections in the study area. Table ESA 2.3-3 lists the Denver wells, dates when groundwater samples were collected, and compounds detected in the samples. ICP metal detections are included in this list, and those samples which were analyzed for ICP metals are indicated by a plus sign on the table.

2.3.6.7 Compound Groups Not Detected

The following compound groups were not detected in the Denver Formation groundwater sampling program in the ESA: VHCs, OSCMs, OSCHs, OPHGBs, OPHPs, ONCs, PAHs, SHOs, and mercury. Groundwater samples were not analyzed for fluoroacetic acid because no certified analysis method for this compound was developed.

2.4 STRUCTURES CONTAMINANTS

On the basis of use history and a visual survey, all ESA structures were classified as to their suspected contamination classification. The locations and identification numbers of structures existing in the ESA are presented in Plate ESA 1.3-1. Four structures in the ESA are suspected to be contaminated, thirty-six are suspected to be contaminated but cleanable, and twenty-two are suspected to be uncontaminated. All four of the structures suspected to be contaminated are located in the Section 31 toxic storage sites. Based on visual observation, 14 structures are suspected to contain asbestos. No structures in the ESA contained process equipment at the time of the Structures Survey for the RI, conducted in the spring of 1987. A summary of the contaminant classification of structures in the ESA is presented in Table ESA 2.4-1. Further information on structures contamination may be found in the Structures Survey Final Summary of Results Report (Ebasco, 1988w/RIC 88306R02).

2.5 AIRBORNE CONTAMINANTS

Twelve air quality monitoring stations were located at RMA during the Air Remedial Investigation (ESE, 1988s/RIC 88263R01). One of these stations, AQ4, was located in the ESA. Station AQ4 was located along the northern edge of Section 5, as shown in Figure ESA 2.5-1. Total suspended particulates were monitored at this station and the other eleven stations at RMA. Particulate matter less than 10 microns (Pm-10), asbestos, and metals were also monitored at selected stations throughout RMA. Volatile and semivolatile organic compounds were evaluated during event monitoring near Basin A and Basin F. Table ESA 2.5-1 summarizes the analytical results from total suspended particulate monitoring at Station AQ4.

At Station AQ4, the analytical geometric average concentration of three individual total suspended particulate samples was 43 micrograms per cubic meter ($\mu g/m^3$). This is below the Federal and State secondary ambient air quality standard of 60 $\mu g/m^3$. Only three individual total suspended particulate samples were collected because of equipment failure. The range of individual 24 hour concentrations for these samples was from 39 to

47 μ g/m³. The maximum 24 hour concentration of 47 μ g/m³ was below the Federal and state standard of 150 μ g/m³ (ESE, 1988s/RIC 88263R01).

The highest annual geometric average total suspended particulate concentration at RMA was 47 μ g/m³, along the northwest boundary in Section 27. The highest individual 24 hour total suspended particulate concentration at RMA was 150 μ g/m³, south of the South Plants Study Area (ESE, 1988s/RIC 88263R01). Higher concentrations were generally noted near the boundaries of RMA as compared to levels monitored at interior sites.

2.5.1 Analytical Results

Station AQ4 was located in Section 5, approximately 2,000 ft west of the RMA east boundary as shown in Figure ESA 2.5-1. Total suspended particulate sampling only was conducted at AQ4 due to equipment failure. Particulate matter less than 10 microns (PM-10), asbestos, metals, and organics were not monitored in the ESA. The occurrence and distribution of total suspended particulates is discussed below.

2.5.2 Distribution of Total Suspended Particulates

Three total suspended particulate samples were collected at AQ4 in June and July 1986. After this time, equipment failure did not allow for reliable data collection. The observed total suspended particulate values were 31, 40, and 47 μ g/m³ which are far below the Federal and State 24 hour maximum limit of 150 μ g/m³ and the RMA maximum observed level of 150 μ g/m³. Air quality at AQ4 is expected to be similar to that at AQ1 and AQ2 in the Western and Southern Study Areas, respectively, in that the stations are near the RMA boundary and adjacent to off-post transportation corridors. In general, these stations realize higher total suspended particulate levels in the driest months from dirt roads than do the interior monitoring locations.

2.6 CONTAMINANTS IN BIOTA

A comprehensive biota assessment of all RMA contamination, both on-post and off-post, was initiated in 1985. This program was designed to determine what, if any, RMA

contaminants remained in the environment and constituted hazards to the regional biota. Although the basic approach was to measure contaminant levels in tissues in comparison to levels in off-post controls, valuable information was also obtained from samples collected by chance (e.g., raptors found dead on RMA and salvaged for analysis). The results of these analyses are discussed in conjunction with information on known and potential effects on biota and ecosystems of the ESA in Section 3.0 of this report. A detailed treatment of all aspects of the biota sampling, chemical analysis regime, contaminants considered, and pathways analysis is contained he Biota Assessment Final Technical Plan (ESE, 1988m/RIC 88243R05) and the Biota RI Report (ESE, 1989a/RIC 89054R01).

2.6.1 Contaminants of Concern

Compounds selected as contaminants of concern to biota met the following criteria:

- o Present in the RMA environment above ambient concentrations:
- o Rated at least moderately toxic; and
- o Volume and persistence information indicate that the chemical was present in the environment in sufficient quantity or for a long enough period of time to pose a hazard to biota.

The thirty-nine contaminants finally selected for evaluation based on this approach are presented in Table ESA 2.6-1. Toxicity assessments were prepared for all contaminants of potential concern to biota and are available in the Biota RI Report (ESE, 1989a/RIC 89054R01). These assessments summarize pertinent information on the nature and extent of existing or potential hazards to wildlife. Data from the assessments were incorporated with information on the concentration and distribution of these contaminants in physical media to provide a quantitative evaluation of RMA contaminant hazards to biota for the ESA and are addressed in Section 3.3 of this document.

Seven contaminants identified as major contaminants of concern based on their presence in the biosphere (e.g., in physical media within 20 ft of the ground surface),

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bioconcentration/bioaccumulation potential, and areal extent (more than 5 acres), were selected for detailed pathways analysis. These contaminants were: aldrin/dieldrin, arsenic, DBCP, endrin/isodrin, and mercury. Two major contaminants of concern were not analyzed in biological tissues. DBCP although toxic, does not bioaccumulate significantly, while isodrin, an analog of endrin, is converted to endrin by metabolic processes. Two additional contaminants, DDE and DDT, were analyzed because of their potential implication in adverse biological effects. The pathways approach was used to develop criteria levels in soil, water, and sediment for the protection of regional biota and to evaluate existing levels to determine the nature and extent of contaminant hazards to biota. The results of pathways analysis and bioaccumulation potential and known and potential effects on biota of the ESA are discussed in Section 3.3.

2.6.2 Sampling Program Completed in the ESA

The diversity of wildlife in this study area led to an equally diverse sampling effort, second only to the Southern Study Area. Control samples of sunflowers and morning glory were taken from Sections 19 and 20, respectively. Both earthworms and grasshoppers were collected as on-post control samples for comparison against contaminated sites. Black-tailed prairie dogs, cottontail rabbits, and mule deer were included in the sampling program for mammals, while regular sampling of pheasants and American kestrels occurred as well. Samples of chance found in the ESA were a coyote, a golden eagle, and a great horned owl. A map of biota sample locations in the ESA is presented in Figure ESA 2.6-1.

2.6.3 Contaminant Levels in Species Occurring in the ESA

Biota samples were prepared by homogenation and extraction procedures according to standard certified USATHAMA methods. GC/MS methods were used for the detection of pesticides, while AA methods were used for the determination of arsenic and mercury concentrations. The CRL for each of the seven major contaminants of concern in the biota program is presented in Table ESA 2.6-2.

Contaminant levels in RMA wildlife species found in the ESA are summarized in Tables ESA 2.6-3 and 2.6-4. Table ESA 2.6-3 summarizes the analytical results from plant and animal samples collected in accordance with technical plans. Table ESA 2.6-4 includes biota samples collected by chance and samples collected by the U. S. Fish and Wildlife Services (USFWS). While some of the samples of the wildlife listed in the tables were not taken from the study area, most of these species have been located or spend some time in the area.

2.6.3.1 Plants and Invertebrates

Plant samples and grasshoppers collected from the ESA contained no detectable level of contaminants. Earthworms had concentrations of arsenic in all eight samples, ranging from 0.62 μ g/g to 1.5 μ g/g. Two of eight earthworm samples contained mercury concentrations of 0.25 μ g/g and 0.21 μ g/g and one sample also contained dieldrin and endrin at concentrations of 5 and 0.9 μ g/g, respectively.

Preliminary statistical analysis of earthworm sample data using the Analysis of Variables techniques indicated significant differences among the three sites (on-post controls, off-post controls, on-post contaminated). Probability values were calculated from available data for defining significance. On-post controls were contrasted with off-post controls, and control samples were pooled and compared to samples from contaminated areas. Of all analytes detected in earthworms, only comparisons for arsenic yielded significant differences. On-post control samples collected from Section 5 of the ESA differed from off-post control samples and pooled control areas differed significantly $(0.05 \ge p > 0.01$, where p is probability) from contaminated sites. Due to the low sample sizes, differences between on-post control and contaminated sites may have remained undetected.

Statistical analyses were completed on all grasshopper samples taken from RMA and offpost locations (see Table ESA 2.6-3). Samples collected from Section 26 were contrasted with those collected from Section 36, on-post control in Sections 7 and 8 of the ESA were contrasted with off-post controls, and pooled data from control sites were

contrasted with pooled data from contaminated areas. None of the comparisons for mercury, DDE, and DDT differed significantly.

Arsenic data differed significantly between the two contaminated sites, and approached significance $(0.10 \ge p > 0.050)$ for the comparison of pooled control and contaminated sites. Significance was obtained for comparisons of aldrin levels between contaminated sites but not between pooled contaminated and pooled control sites. Significance was obtained for comparisons of dieldrin levels between contaminated sites, pooled contaminated, and pooled controls, but not between on-post and off-post controls. For endrin, Section 36 values differed significantly from Section 26, but when pooled, these sites did not differ from either on-post or off-post controls.

2.6.3.2 Black-tailed Prairie Dogs

Nineteen samples of prairie dogs were collected from three different areas in the ESA (Figure 2.6-1): four samples were collected from the northwest corner of Section 19, ten from the northeast corner of Section 20, and five from the Section 31 toxic storage yard. Three of the fourteen samples from Sections 19 and 20 (the on-post control samples) contained concentrations of dieldrin at 0.06, 0.1, and 0.3 $\mu g/g$. No other analytes were detected in these samples. Of the five samples taken from the Section 31 toxic storage yard, all samples contained dieldrin from 0.06 $\mu g/g$ to 0.2 $\mu g/g$. One sample from this site also contained 4.2 $\mu g/g$ of arsenic.

Prairie dog samples were statistically contrasted by site and season. Seasonal differences were compared between Section 36, summer versus winter, and between on-post control areas, summer versus winter. Analysis of seasonal differences revealed that only dieldrin had significantly higher contaminant levels in summer than winter. All other seasonal contrasts were not significant. Section 36 (summer and winter combined) was then contrasted to the Section 31 toxic storage yard, while on-post control areas (summer and winter combined) were contrasted to off-post controls. A final comparison was made between pooled contaminated samples (Section 36 and the Section 31 toxic storage yard) and control sites (on and off-post). Dieldrin levels were significantly higher in samples

from Section 36 than in samples from the toxic storage yard and higher in samples from on-post controls than samples from off-post controls. Analysis of pooled samples for dieldrin revealed highly significant differences between contaminated sites and controls. Comparisons for mercury, aldrin, endrin, and arsenic failed to yield any significant differences, although arsenic approached significance $(0.10 \ge p > 0.050)$ for the comparison of pooled control sites with pooled contaminated sites.

2.6.3.3 Cottontail Rabbits and Mule Deer

Seven samples of cottontail rabbit muscle were collected from carcasses obtained in Sections 19 and 20 of the ESA. No analytes were detected. Three mule deer were collected near First Creek in the ESA, but none of the tissue samples from these deer had detectable concentrations of analytes.

2.6.3.4 Kestrels and Pheasants

A total of nine American kestrel eggs were taken from the ESA. Two samples were found to contain 1 and 0.05 μ g/g of dieldrin. No other analytes were detected. No analytes were detected in the three kestrel fledgling carcasses taken from the ESA.

None of the statistical comparisons for aldrin, endrin. DDE, and DDT between contaminated and control sites showed any significant differences for kestrel eggs or carcasses. For mercury, no significant difference was observed between control and contaminated sites, but differences between eggs and juveniles were significant. In contrast, differences between control and contaminated sites for dieldrin were significant for both juveniles and eggs, while no difference was detected between age groups. Both eggs and juvenile kestrel samples contained higher levels of dieldrin on RMA than offpost; in fact, no dieldrin was detected in either egg or juvenile kestrel samples collected off-post.

Three ring-necked pheasant eggs were collected in the ESA, with samples containing 0.2, 0.2, and 0.1 μ g/g of dieldrin; no other analytes were detected. Nine samples of adult pheasant muscle were analyzed (from nine carcasses taken from the ESA), and no

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analytes were detected. Two livers from adult pheasants were analyzed, with one sample containing 0.1 μ g/g of dieldrin.

Ring-necked pheasant samples were statistically contrasted by off-post control versus onpost contaminated areas, and by age groups (egg, juvenile, or adult). No significant differences in levels of mercury, aldrin, endrin, DDE, or DDT were obtained for any contrasts. No significant differences in levels of arsenic were detected between control and contaminated sites within any of the three age groups, but significant differences were revealed among age groups for arsenic. Juvenile pheasant samples collected, both on and off-post, contained significantly higher levels of arsenic. No adult or egg samples contained any detectable arsenic. In contrast, significant differences between control and contaminated sites for eggs and juvenile pheasant samples were obtained for dieldrin. The preliminary Analysis of Variables showed dieldrin was significant. No differences were observed in adult pheasant samples from control and contaminated sites, probably due to the small population sampled.

2.6.3.5 Samples of Chance and Supplemental Samples

Dieldrin was detected at 8 μ g/g in the liver of a coyote found dead in the ESA. No other analytes were detected. Brain and liver tissue was sampled in a great horned owl found dead near First Creek. The owl brain contained 10 μ g/g dieldrin, and 2 μ g/g DDE, while the liver contained 0.057 μ g/g mercury, 9 μ g/g dieldrin, and 5 μ g/g DDE. A golden eagle found on the RMA eastern border was sampled for brain and liver tissue, but no analytes were detected. Supplemental biota samples were provided by the USFWS, and both samples of chance and USFWS samples are summarized on Table ESA 2.6-4.

Complete tables and figures summarizing the contaminant levels found in all samples taken on RMA may be found in the RI Report for Biota (ESE, 1989/RIC 89054R01).

2.7 SUMMARY

Organic compounds detected in the soils of the ESA included a few detections near the

CRL of VHOs and VAOs and OSCM concentrations above the CRL at a few isolated spill sites. VHOs and VAOs were detected in ESA-2b, the sanitary landfill (Site 30-4), and ESA-3d, the Section 31 toxic storage plots (Sites 31-6 and 31-7). OSCMs were detected in ESA-3d, ESA-3b, and ESA-3a, the Section 31 toxic storage plots, the Section 6 toxic storage yard, and the Section 5 toxic storage yard (Sites 31-6 and 31-7, Site 6-6 and Site 5-2, respectively). No apparent concentration trends are associated with VHO or VAO concentrations, and distribution patterns indicative of spills are associated with the OSCM concentrations. Metals were detected throughout the ESA, usually within or slightly above their indicator ranges. Metal concentrations above indicator ranges were found in all the ESA sites except the Section 5 toxic storage yard, although the highest metal concentrations were detected in ESA-4c, the Section 29 trench and mound, ESA-1, the surface burn sites (Sites 19-1, 20-1, 29-1 and 30-2), and ESA-2a, the Section 32 burn pits (Sites 32-5 and 32-6).

Detections of VHOs, OPHGBs, OCPs, arsenic, and ICP metals were found in surface water monitoring of First Creek in the ESA. Concentrations were relatively low level, and no contaminant patterns were noted. Zinc was the only analyte detected twice during the sampling, at both stations 08-001 and 24-002. All other analyte detections were single, nonrepeated hits.

Detections of VHOs, VAOs, OSCHs, DBCP, OCPs, arsenic, and ICP metals were found in the alluvial aquifer groundwater. The majority of organic compound detections occurred in samples collected near the North Boundary Containment System, which is discussed in the North Central Study Area Report. Repeat detections of dieldrin occurred at Well 24166, and repeat detections of chlorophenylmethyl sulfone occurred at Wells 24183 and 24188. All other organic analyte detections were single, nonrepeated hits. In most instances, ICP metals were analyzed for only once in samples collected from each well. Evidence does not indicate contaminant plumes occur in the alluvial aquifer of the ESA.

Detections of VHOs, VAOs, DBCP, OCPs, arsenic and ICP metals were found in the Denver Formation groundwater. Repeat detections of dieldrin occurred at Well 24120, and repeat detections of benzene and chlorobenzene occurred at Wells 24174 and 24175. Benzene was detected in two samples from Well 07005. All other analyte detections were single, nonrepeated hits. In most instances, ICP metals were analyzed for only once in samples collected from each well. Evidence does not indicate contaminant plumes in Denver Formation groundwater in the ESA.

The distribution of contaminants in ESA surface water and groundwater may have been presented differently than the neighboring study areas. Due to the limited numbers of detections in surface water and in the alluvial and Denver Formation groundwater systems in the ESA, the maximum concentrations were presented on contaminant distribution maps. The Southern, South Plants, Central, North Plants, and North Central study areas each presented surface water and groundwater data uniquely, and direct comparison of groundwater contaminant distribution maps among study areas may not be possible.

All structures existing in the ESA were identified and categorized as either contaminated, contaminated but cleanable, or uncontaminated. Of the 62 structures in the ESA, four are suspected to be contaminated, 36 are suspected to be contaminated but cleanable, and 22 are suspected to be uncontaminated. Based on visual observation, 14 structures are suspected to contain asbestos.

Air monitoring showed that the total suspended particulates observed in the ESA were below the Federal and State secondary ambient air quality standards. The maximum twenty-four hour concentrations of total suspended particulates of the ESA monitoring station were also below Federal and State standards.

Generally low levels of contaminants were found in biota sampling in the ESA. Of the few detections of contaminants, dieldrin was the most commonly detected analyte, followed by endrin, aldrin, arsenic, mercury, and DDE.

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Table ESA 2.1-1 Phase I and II Analyte	s and Certified	Reporting L	imits for Eas	tern Study	Area Soil Samples	. Page 1 of 6.			
		Certified R	Phase 1 eporting Li	mits (µg/	6	Certifie	Phase ed Reportin	II ig Limits	(µg/g)
Analytical Groups/Analytes	Method	CAL	ESE	MRI	DataChem	Method	CAL	ESE	DataChem
<u> Volatile Halogenated Organics (VHO</u>	ाज								
1,1-Dichloroethane	GC/MS	6.0	0.3	0.5	2	COHECD	ł	0 040	0.074
1,2-Dichloroethane	GC/MS	0.3	0.3	0.4	0.6	GC/HECD	I	0.050	0.085
1,1-Dichloroethylene	GC/MS	ł	I	I	1	GC/HECD	ŧ	0.047	0.24
1,2-Dichloroethylcne	GC/MS	0.3	0.3	0.8	2	GC/HECD	1	0.051	0.26
1,1,2,2-Tetrachloroethane*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	I	0.3	0.3
1, 1, 1 - Trichloroethane	GC/MS	0.3	0.3	0.5	0.4	GC/HECD	í	0.049	0.088
1,1,2-Trichloroethane	GC/MS	0.3	0.3	0.6	0.4	GC/HECD	ł	0.050	0.26
Carbon tetrachloride	GC/MS	0.3	0.3	0.4	0.3	GC/HECD	ł	0.052	0.12
Chlowbenzene	GC/MS	0.3	0.3	0.3	-	GC/HECD	i	0.051	0.20
Chioroform	GC/MS	0.3	0.3	0.7	0.3	GC/HECD	:	0.052	0.068
Methylene chloride	GC/MS	0.7	0.3	ł	2	GC/HECD	ł	0.50	3.7
Tetrachloroethylene	GC/MS	0.3	0.3	0.5	0.3	GC/HECD	1	0.051	0.27
'frichloroethylene	GC/MS	0.3	0.3	0.6	0.5	GC/HECD	ł	0.050	0.14
Trichloropropene*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	ł	0.3	0.3
<u> Volatile Ilydrocarbons (VHCs)</u>									
2-Butoxyethanol*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	1	0.3	03
4-Hydroxy-4-methyl-2-pentanone*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	ł	0.3	0.3
1-Methyl-1, 3-cyclopentadiene*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	ı	0.3	0.3
2,2-Oxybiscthanol*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	1	0.3	0.3
2-Pentanone*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	ł	0.3	0.3
Bicycloheptadiene	GC/MS	0.3	0.3	0.8	0.4	GC/FID	I	5.1	1.1
Dicyclopentadiene	GC/MS	0.3	0.3	0.3	0.7	GC/FID	I	5.1	0.45
Methylcyclohexane*	GC/MS	0.3	0.3	0.3	0.3	GC/MS	1	0.3	0.3
Methyli:sobutyl ketone	GC/MS	0.3	0.5	0.4	0.7	GC/FID	1	5.2	0.64

 Certification not received
 Significant nontarget compound without a CRL. The lower limit of detection is 10% of the internal standard for the method used. For the purpose of this report, a CRL" of 0.30 µg/g was used
 CRL" of 0.30 µg/g was used
 Ortho- and para- (o- and p-) Xytenes coelute under the GC conditions specified in this method 1.*

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Table ESA 2.3-1/ESA-1 5/10/89 10:00 AM

	•	Certified R	Phase I teporting Li	mits (µg/	g)	Certifi	Phase Phase	e II 	
Analytical Groups/Analytes	Method	CAL	ESE	MRI	DataChem	Method	CAL	ng Limits F.S.F.	(µg/g) DataChin
<u> </u>								101	DataChem
Berzene Ethylbenzene m-Xylene o- and p-Xylene** Toluene	GC/MS GC/MS GC/MS GC/MS GC/MS	0.3 0.3 0.3 0.3	0.3 0.3 0.5 0.3 0.3		0.3 5 8 0.3 0.3	GC/PID GC/PID GC/PID GC/PID GC/PID	1111	0.081 0.043 0.053 0.086	0.085 0.16 0.26 0.39
<u>Oreanosulfur Compounds.</u> Must <u>ard–Agent Related (OSCMs)</u>							I	0.096	0.19
1,4-Oxadhiane Chloroacctic acid Dithiane Thiodiglycol	GC/MS GC/MS	1 - 1 - 0	0.3 - 0.3	0.5 - -	0.3 - 0.4	GC/FPD HPLC GC/FPD	1 1 1	0.90 18 0.60	1.7 36 1.4
Organosultur Compounds. Uerbicide Related (OSCHs)							I	0.7	4.2
Benzoshiazole Chlorophenylmethyl sulfide Chlorophenylmethyl sulfone Chlorophenylmethyl sulfoxide Dimethyldisulfide	GC/MS GC/MS GC/MS GC/MS	4 + 0.6 0.8 0.8	0.3 0.3 0.4 0.3	- 0.3 - 4 - 4	0.9 0.3 20	GC/FPD GC/FPD GC/FPD GC/FPD GC/FPD	11111	0.53 1.1 2.4 2.3 0.70	9.4.0 9.6.0 9.1.8

Table ESA 2.1-1 Phase I and II Analytes and Certified Reporting Limits for Eastern Study Area Soil Samples. Page 2 of 6.

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 Certification not received
 Significant nontarget compound without a CRL. The lower limit of detection is 10% of the internal standard for the method used. For the purpose of this report, a CRL" of 0.30 µg/g was used
 Ortho- and pura- (o- and p-) Xylenes coelute under the GC conditions specified in this method *

Table ESA 2.3-1/ESA-1 5/10/89 10:00 AM

		Certified R	Phase 1 eporting Li	imits (µg/	(g	Certifie	Phase d Reportin	II ig Limits	(β/8I)
Analytical Groups/Analytes	Method	CAL	ESE	MRI	DataChem	Method	CAL	ESE	DataChem
Organophophorous Compounds, GB-Agent Related (OPHGBs)									
Diisopropylmethyl phosphonate Dimethylmethyl phosphonate Isopropylmethyl phosphonic acid Methylphosphonic acid Phosphoric acid, tributyl ester* Phosphoric acid, triphenyl ester*	GC/MS GC/MS GC/MS GC/MS	0.3 - - 0.3 0.3	0.5 2 - 0.3 0.3	0.3 0.3 0.3	1 0.3 0.3	GC/FPD GC/FPD HPLC IONCHROM GC/MS GC/MS	0.050 0.050 4.7 0.3 0.3	0.11 2.6 0.3 0.3 0.3	1111
Organonhosphorous Compounds, Pesticide Related (()PHPs)									
Atrazine Malathion Parathion Supona Vapona	GC/MS GC/MS GC/MS GC/MS	0.3 0.3 0.3 0.3 0.3	0.7 0.6 0.3 0.3	0.5 2 0.9 0.3	0.3 0.6 3	GCANPD GCANPD GCANPD GCANPD GCANPD		0.25 0.25 0.25 0.70	1111
DBCP	GC/MS	0.3	0.3	0.6	0.3	GC/EC	0.014	0.0050	0:0050
Organonitrogen Compounds (ONCs) Caprolactum* Hydrwine Methythydrazine n-Nitrosodimethylamine n-Nitrosodi-n-propylamine Unsymmetrical dimethyl hydrazine		1 1 1 1 1 1				GC/MS IONCHROM IONCHROM GC/NPD GC/NPD GC/NPD			0.3 50 0.10 200 200

Table ESA 2.1-1 Phase I and II Analytes and Certified Reporting Limits for Eastern Study Area Soil Samples.

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Page 3 of 6.

= Certification not received
 = Significant nontarget compound without a CRL. The lower limit of detection is 10% of the internal standard for the method used. For the purpose of this report, a CRL" of 0.30 μg/g was used
 = Ortho- and para- (o- and p-) Xylenes coelute under the GC conditions specified in this method

Table ESA 2.3-1/ESA-1 5/10/89 10:00 AM

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Table ESA 2.1-1 Phase I and II Analytes	s and Certified	Reporting L	imits for East	tern Study	Area Soil Samples	. Page 4 of 6.			
	Ŭ	Certified R	Phase 1 eporting Lin	mits (µg/	8)	Certifie	Phase d Reporti	e II ng Limits	(J)(g/g)
Analytical Groups/Analytes	Method	CAL	ESE	MRI	DataChem	Method	CAL	ESE	DataChem
Eluaroscetic Acid		I	ł	ł	I	IONCHROM	I	2.0	ł
<u>Polynuclear Aromatic Hydrucarbons (</u>	PAILS								
Fluoranthene* Methylnapthalene* Phenanthrene* Pyrene*	GC/MS GC/MS GC/MS GC/MS	0.3 0.3 0.3 0.3	0.3 0.3 0.3 0.3	0.3 0.3 0.3	0.3 0.3 0.3 0.3	GC/MS GC/MS GC/MS	0.3	0.3 0.3 0.3 0.3	0.3 0.3 0.3
Semivolatile Nalogenated Organics (S	(50115	•					2	2	2
Hexachlorobenzene* Hexachlorobutadiene* Hexachlorobenzene* Tetrachlorobenzene* Trichlorobenzene*	GC/MS GC/MS GC/MS GC/MS GC/MS	0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0	0.3 1.3 0.3 0.3 0.3	0.3 0.3 0.3 0.3	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	GC/MS GC/MS GC/MS GC/MS GC/MS		0.3 0.3 0.3 0.3 0.3	0.3 0.018 0.3 0.3 0.3 0.3

= Certification not received
 = Significant nontarget compound without a CRL. The lower limit of detection is 10% of the internal standard for the method used. For the purpose of this report, a CRL" of 0.30 µg/g was used
 = Ortho- and pura- (o- and p-) Xylenes coelute under the GC conditions specified in this method

Table ESA 2.3-1/ESA-1 5/10/89 10:00 AM

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	-	Certified 1	Phase 1 Reporting Li	mits (µg/	g)	Certifi	Phas ed Reporti	e II ng Limits	(Jug/g)
Analytical Groups/Analytes	Method	CAL	ESE	MRI	DataChem	Method	CAL	ESE	DataChem
Organochlorine Pesticides (OCPs)									
Aldrin	GC/MS	0.3	0.9	0.5	0.3	GC/FC	ł	0.0010	
Chlordane	GC/MS	0.6	-	Q	2	CCEC	1		0.013
Dichlorodiphenylethane (DDE)	GC/MS	0.3	0.3	0.5	0.6	GC/EC	ł	0,0010	0 000
Dichlorodiphenyltrichloroethane (DDT)	GC/MS	0.6	0.4	2	0.5	GC/EC	ł	0.050	0,0000
Dicklin	GC/MS	0.3	0.3	0.6	0.3	GC/EC	I	0.0012	0.0013
Endrin	GC/MS	0.3	0.7	4	0.5	GC/EC	I	0100.0	0.0058
lsotrin	GC/MS	0.3	0.3	0.6	0.3	GC/EC	I	0.0011	1100.0
Arsenic	GFAA	5.0	4.7	2.5	2.5	GFAA	5.0	4.7	2.5
Mercury	CVAA	0.060	0.050	0.070	0.050	CVAA	0.060	0.050	0.050
ICP Metals									
Cadmium	ICP	0.66	06.0	0.51	0.74	ICP	0.66	06.0	0 74
Chromium	ICP	5.2	7.2	7.4	6.5	ICP	5.2	7.2	6.5
Copper	ICP	4.9	4.8	4.9	4.7	ICP	4.9	4.8	4.7
L CM	ICP	13	17	16	8.4	СЪ	13	17	8.4
Zinc	ICP	9.5	16	28	8.7	ICP	9.5	16	8.7

Table ESA 2.1-1 Phase I and II Analytes and Certifiert Reporting Limits for Eastern Shireb Area Soil Sa

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Certification not received
 Jignificant nontarget compound without a CRL. The lower limit of detection is 10% of the internal standard for the method used. For the purpose of this report, a CRL" of 0.30 µg/g was used
 Ortho- and para- (o- and p-) Xylenes coelute under the GC conditions specified in this method

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 Table ESA
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Table ESA 2.1-1 Phase 1 and 11 Analytes and Certified Reporting Limits for Eastern Study Area Soit Samples. Page 6 of 6.

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CAL ESE MRI DutaChem	Е И И И	Califonia Analytical Laboratory, W. Sacramento, California. Environmental Science and Engineering, Inc., Gainesville, Florida. Midwest Research Institute, Kansas City, Missouri. formerly UBTL, Salt Lake City, Utah.
Methodi		
GC/MS GC/FID GC/FID GC/FID GC/FID GC/FID GC/FD GC/FD GC/FD CVAA CVAA CVAA CVAA CVAA CVAA CVAA CVA	11 N H H H H H H H H H H	Gas chromatography/mass spectrometry. Gas chromatography/flame ionization detector. Gas chromatography/flame ionization detector. Gas chromatography/flame phouxmetric detector. Gas chromatography/flame phouxmetric detector. Gas chromatography/nitrogen phosphorous detector. Cold vapor atomic absorption. Inductivity coupled plasma. Ion chromatography. High performatee fiquid chromatography.
Source: EBA	SCC.	April 1989.

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 Table ESA 2.3-1/ESA-1

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			ESA-1. Surface	Burn Sites		
Total Borings Total Samples	Phase	I Analyses 32 113		<u>е</u>	hase II Analyses 2	
	Frequency of	Range	CRL Range	Frequency of	Rame	ਲੂ
Analytical Groups and Analytes Detected	Detections/1	(6/6rl)	^{2/(6/61)}	Detections ^{/1}	(b/bn)	fin/n/2
<u> Volatile Halovenated Organics (VHOs)</u>						IRACI
1,1-Dichloroethane	8/0	BCRL	0.3-0.5	CN		
1,2-Dichloroethane	8/0	BCRL	0.3-0.4	CN CN	1]
1,1-Dichloroethylene	QZ	I	1	QN		ļ
1,2-Dichloroethylene	8/0	BCRL	0.3-0.6	QN	1	ł
1,1,2,2-Tetrachloroethane*	Q	ł	J	QN	1	ł
1,1,1-Trichlomethane	8/0	BCRL	0.3-0.5	QN	1	ł
1,1,2-Trichtoroethane	8/0	BCRL	0.3-0.6	ÛN		1
Carbon Tetrachloride	8/0	BCKL	0.3-0.4	ON N		1
Chlorobenzene	8/0	BCRL	0.3	Q	1	
Chloroform	8/0	BCRL	0.3-0.7	QN	I	
Tetrachloroethylene	8/()	BCRL	0.3-0.5	ND	- 	
Trichloroethylene	8/0	BCRL	0.3-0.6	Q	I	
Trichloropropene•	QN	1	I	QN	ł	
Methylene Chloride	()/4	BCRL	0.3 /	Q	1	I
<u>Volaule Hydrocarbons (VHCs)</u>						
2-Butoxyethanol*	QN	I	I	QN	1	
4-Hydroxy-4-methyl-2-pentanone*	QN	I	1		[]	1
I-Methyl-1,3-cyclopentadiene*	0N	ł	I	Ē	1	ł.
2,2-Oxybisethanol*	13	0.5-1.0	1	QZ	: 1	
2-Pentanone*	QN	ł	I	CZ		
Bicycloheptadiene	8/0	BCRL	0.5-0.8	QN	1	

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 1 of 30.

Below Certified Reporting Limit. Ħ BCRL

Indicator Range H X

At or Below Upper Indicator Level Micrograms per gram. 11 BUIL

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= Analysis not requested for original target compounds/not detected for nontarget compnunds subsequently considered in analyte groups. 2

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Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.
Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used.

			ESA-1 Surfec				
Total Borings Total Samples	ξ (ase I Analyses 32 113			Phase II Analyses 2		
Analytical Groups and Analytes Detected	Frequency cf Detections ^{/1}	Range (µq/q)	CRL Range Iun/o/2	Frequency of	4 Range	CRL Range	
Dicyclopentadiene	0/113	RCRI	76,64	Detections'	(6/6n)	^{2/(6/61)}	1
Methylcyclohexane* Methylicoburd Process	QN	1	0-C.U	1/0 1/0	BCRL	0.3	
wichly itsobuly! Actone	8/0	BCRL	0.4-0.5	2 Q	i I	1	
Volattle Aromatic Organics (VAOs)				•	I	ı	
Benzene Ethotheorean	8/0	BCRL	0.3-1	Ę.			
m-Xvlene	8/0	BCRL	0.3-0.4	2 Z	I	I	
0- and n-Xylene	8/0	BCRL	0.3	2	j	1	
Toluene	<u>8</u>	0.7 BCDI	0.5	DN		1 1	
Organosulfur Communds		DUNG	0.3	QN	1	1	
Mustard - Avent Related (OSCMs)							
1,4-Oxathiane	0/113	BCRL	0.3-0.5	20	-		
Cinoroaceue acid Dithiane	QN	ţ .		ND 4	BCRL	0.3-0.9	
Thiodiglycol		BCRL	0.3-2	0/4	BCRL	0.3-0.6	
Organosulfur Compounds	l ,	ł	ł	QX	đ	1	
Herbicids: Related (OSCHs)							
Benzothiazole	-	0.5	0.3	CIN			
Curtorphenylmethyl sulfage Chloronhenylmethyl sulfage	0/113	BCRL	0.3	0/4	 BCBI	0	
Chlorophenylmethyl sulfoxide		BCRL	0.3-0.4	0/4	BCRL	0.4-2	
Dimethyldisulfide	8/0	BCRL	0.3-4	0/4	BCRL	0.3-2	
Date					BCKL	0.7	
BCKL = Below Certified Reporting Limit. IR = Indicator Range							- 1
BUIL = At or Below Upper Indicator Level							
$\mu g/g = Micrograms per gram.$							

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Table ESA 2.1-2. Summary of Soil EUTing Analytical Results in the Eastern Study Area. Page 2 of 30.

We analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.
 Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.
 I = Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. .

			ESA-1. Surface	e Burn Sites		
lotal Borings Iotal Samoles	Phas	e I Analyses 32 113			hase II Analyses 2 4	
thatical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g/ ²	Frequency of Detections ^{/1}	Range (µg/g)	САL Rarge (µg/g) ^{/2}
Xranophosphorous Compounds/ GB-Agent Related (OPHGBs)		ia.cu	ç			
Dusopropyincuryi prospriorate Dimethylmethyl phosphonate	0/113	BCRL	2.3	1/0	BCRL	0.3-0.11 2
Isopropylmethyl phosphonic acid Methylphosphonic acid	Q Q			a a	11	11
Phosphoric acid, tributyl ester* Phosphoric acid, triphenyl ester*	QN NN	11	1 !	an Un	11	1 1
Drgunophosphorous Compounds/ Pesticide: Related (OEHPs)						
Atrazine Malathion	0/113 0/113	BCRL BCRL	0.5-0.7 0.6-2	1/0	BCRL	0.5 0.6
Parathion Supona Version	0/113 0/113	BCRL BCRL BCRI	0.7-2 0.5-0.9 0.3	1/0	BCRL BCRL BCRI	0.7 0.5 0.3
Dibromochloropropane	0/113	BCRL	0.3-0.6	1/0	BCRL	0.3
Drganonitrogen Compounds (ONCs) Hydrazine Methylhydrazine n-Nitrosodi-n-propylamine n-Nitrosodimethylamine	<u> 2</u> 2 2 2 2 2		1111	88888	1 []]	
		l	1	2	l	ļ

Table ESA 2.1-2. Summery of Soil Boring Analytical Results in the Eastern Study Area. Page 3 of 30.

Below Certified Reporting Limit. 11 BCKL

Indicator Range H Ř

At or Below Upper Indicator Level Micrograms per gram. H BUIL

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. n Iŧ 8/84 N

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= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

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Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 4 of 30.

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			ESA-1. Surface	Burn Sites		
Total Borings Total Samples	Phas	e I Analyses 32 113			hase II Anaiyses 2 4	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µ9/9)	CRL Range (µ9/9/ ²	Frequency of Detections ^{/1}	Range (µq/q)	CPL Range (uo/o//2
Fluorcacetic Acid	QN	}	ł	Q		
Polymicken Aromatic Hydrocarbons (PAHs) Fluoranthene* Methylnapht-hlene* Phenanthrene* Pyrene*	ND 2 2 ND	0.5-1 1-20	0.3 0.3 1	8888	111	1111
Semivolutic Halogenated Organics (SHOs) Hexachlorobenzene* Hexachlorobutadiene* Hexachlorobenzene* Pentachlorobenzene* Tetrachlorobenzene*		BCRL		Q Q 50 Q Q Q	BCRL	
Organschlorine Pesticides (OCPs) Aldrin Chlordane Dichlorodiphenylethane Dichlorodiphenyltrichloröcthane Dichlorin Endrin Isodrin	0/113 0/113 0/113 0/113 0/113 0/113	BCRL BCRL BCRL BCRL BCRL BCRL	0.5-0.9 1-6 0.3-0.5 0.3-0.6 0.3-0.6 0.7-4 0.3-0.6	1/0 1/0 1/0 1/0	BCRL BCRL BCRL BCRL BCRL BCRL BCRL	0.3 0.3 0.3 0.3 0.3 0.3

Below Certified Reporting Limit. BCRL =

Indicctor Range n ĸ

At or Below Upper Indicator Level 9 BUIL.

Micrograms per gram. n 8/81 9/81

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. H 7

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution 2 ٠

effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit hased on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

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Results in th
ng Analytical
ol Soil Borin
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ble ESA 2.1-2
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			ESA-1. Surface	Burn Sites		
Total Borings Total Samples	Pha	se I Analyses 32 113		8	Phase II Analyses	
Analytical Groups and Analytes Detected	Frequency of Detections/1	(µg/g)	CRL Range (µ9/9/ ²	Frequency of Detections/1	Range (uɑ/ɑ)	CRL Range Iunin/2
Arsenic (IR=CKL-10)	2/113	15-20	4.7-5.2	QN		16,84
Mercury (IR=CRL-0.10)	2/113	0.16-0.32	0.050-0.070	QN	I	i
ICP Metals						I
Cadmium (IR=CRL-2.0)	11/113	2.1-5.1	0.51-0.00	ÛN		
Chromium (IR=25-40)	5/113	44-56	000 1000 V L-C L		I	ł
Copper (IR=20-35)	13/113	36-130	48.40		1	I
Lead $(IR=25-40)$	14/113	41-110	16-17	2 Q	ł	I
Zinc (IK=60-80)	18/113	81-120	16-28	2 Q		1

Below Certified Reporting Limit. H BCRL

Indicator Range 8 R BUIL

At or Below Upper Indicator Level ĸ

H 1ł 18/8 1

⁼ Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. Micrograms per gram. Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

⁼ Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. 2 *

Total Borings Total Samples	LE I	iase I Analyses 48 126	ESA-2. Burlel	Irench Sites	Phase II Analyses 35	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g) ^{/2}	Frequency of Detections/1	105 Range	Crt. Range
Yolaufe Halegenated Organics (YHO2) 1,1-Dichloroethane 1,2-Dichloroethane 1,2-Dichloroethane 1,1,2-Tetrachloroethane 1,1,2-Trichloroethane 1,1,2-Trichloroethane 1,1,2-Trichloroethane 1,1,2-Trichloroethane 1,1,2-Trichloroethane Carbon Tetrachloroethane Carbon Tetrachloroethane Carbon Tetrachloroethane Trichloroethylene Tr	071 071 071 071 071 071 071 071 071 071	BCRL BCRL BCRL BCRL BCRL BCRL BCRL BCRL	$\begin{array}{c} 0.3-2\\ 0.3-0.6\\ 0.3\\ 0.3-2\\ 0.3-0.6\\ 0.3-0.6\\ 0.3-0.6\\ 0.3-0.6\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3\\ 0.3$	0/37 0/37 0/37 0/37 0/37 0/37 0/37 0/37	BCRL BCRL BCRL BCRL BCRL BCRL BCRL BCRL	0.07-2 0.08-0.6 0.2 0.3-2 0.3-0.4 0.1-0.3 0.1-0.3 0.1-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.3 0.2-0.4 0.0-2 0.2-2 0.2-0.4 0.0-2 0.2-0.4 0.0-2 0.0-4 0.0-2 0.0-4 0.0-2 0.0-4 0.0-2 0.0-4 0.0-2 0.0-2 0.0-4 0.0-2 0.0-2 0.0-4 0.0-2 0.0-2 0.0-4 0.0-2 0.0-3 0.1-0.3 0.0-2 0.0-2 0.0-4 0.0-2 0.0-2 0.0-4 0.0-2 0.0-2 0.0-4 0.0-0-4 0.0-2 0.0-2 0.0-2 0.0-4 0.0-2 0.0-2 0.0-2 0.0-2 0.0-4 0.0-0-4 0.0-0-3 0.0-2 0.0-2 0.0-2 0.0-2 0.0-2 0.0-4 0.0-0-3 0.0-2

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 6 of 30.

Below Certified Reporting Limit. 9 RCKL

Indicator Range B ž

At or Below Upper Indicator Level H BUIL

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đ 8/8rd 7 N

Micrograms per gram Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

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= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. ٠

= There is no CRL for tentausely identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of idencitions is given, but the number of samples is not.

			ESA-2. Burlet	Trench Sites			
Total Borings Total Samrias	Phase	I Analyses 48 126			hase II Analyses 35 105		
AnaMtical Groups and AnaMtes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µ9/9) ^{/2}	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g/2	
Dicyclopentadiene	0/125	BCRL	0.3-6	0/29	BCRL	0.3-0.6	r i
Methylcyclohexane* Methylisobutyl Ketone	CIN 11/0	- BCRL	0.3-2	UN 81/0	– BCRL	0.3-0.7	
Volatile Aromatic Organics (VAOs)							
Benzene	4/11	0.3-0.5	0.3	5/40	0.014-0.5	0.085-0.3	
Ethylthenzene	11/0	BCRL	0.3-0.4	0/40	BCRL	0.2-0.4	
m-Xylene	1/11	0.8	0.7-0.8	0/40	BCRL	0.3-0.7	
o- and p-Xylene	1/1	5	0.3-5	0/40	BCRL	0.3-5	
Toluene	1.1/0	BCRL	0.3-0.6	0/40	BCRL	0.2-0.3	
Organosulfur Compounds Mustard - Arent Related (OSCMs)							
1,4-Oxathiane	0/125	BCRL	0.3-6	0/20	BCRL	0.3-6	
Chloroacetic acid	86/0	BCRL	40	0/22	BCRL	20	
Dithiane	0/125	BCRL	0.3-7	0/20	BCRL	0.3-7	
і пихнідіусої	vc/n	DURL	Ŧ	7710	DUND		
Organosulfur Compounds Herbicide Related (OSCHS)							
Benzothiazole	QN	1	ł	6/0	BCRL	-	
Chlorophenylmethyl sulfide	0/125	BCRL	0.34	0/20	BCRL	0.3-4	
Chlorophenylmethyl sulfone	2/125	0.9-2	0.3-0.6	0/20	BCRL	0.4-7	
Chlorophenylmethyl sulfoxide	0/125	BCRL	0.3-0.6	0/20	BCRL	0.3-2	
Dimemylaisuitae	1/10	BURL	N7-C'N	(7)N	DUNE	07-01-0	

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 7 of 30.

Below Certified Reporting Limit. 11 BCKL

Indicator Range H ¥

ł BUIL

At or Below Upper Indicator Level Micrograms per gram. 11 8/81 ND

Analysis not requested for original target compounds/not detected for nontarget comprunds subsequently considered in analyte groups. 0

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. -

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Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

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Total Borings Total Samples	Phase	I Analyses 48	ESA-2. Burlal	Trench Siles	Phase I Analyses	
		136	đ		35 105	
Analytical Groups and Analytes Detected	Frequency of Detections/1	Range (µg/g)	Range (uo/o)/2	Frequency of Detections/1	Range	CRL Range
<u>Orgattophosphorous Compounds/</u> <u>GB-Arent Related (OPHGRs)</u>			18 5 1	CALACIOUS	(6/61)	(н <u>9</u> /д)/2
Diisopropylmethyl phosphonate	30175	4				
Dimethylmethyl phosphonate	620	BCRL	0.3-3	0/12	BCRL	
tsopropytmethyl phosphonic acid Methylphysnhonic acid	0/18	BCRL	2-3 0.6-5	11/0	BCRL	0.1-0 0.1-2
Physphoric acid, tributyl ester-	Q A	ł	1	0/22	BCRL	2
Phosphoric acid, triphenyl ester+	2 Q	1	I	Q		2-10
Organophosphorous Compounds/		ł	ŀ	QN	ł	11
<u> Festicide Related (OPLIPs)</u>						
Malathion	0/126	BCRL	0.3-0.7	C 170		
Parathion	0/126	BCRL	0.3-2	0/12	BCRL	0.3-0.7
Supona Varveo	0/126	BCRL	0.3-2	0/12	BCRL	0.3-0.6 0.4-0.6
	0/126	BCRL	0.3-3	0/12	BCRL	0.3-0.5
Libromox hloropropane	0/125	BCRL	01-0	71/0	BCRL	0.3
Organemitrogen Compounds (ONCs)			7-0-0	67/0	BCRL	0.3-2
Hydrazine Methylhydraeiee	QN	I	1	<u>í</u>		
n-Nitrosxdi-n-pronylamine	2 !	I	1	2 F	I	1
n-Nitrosodimethylamine		ł	1	2 Q	I	1
Unsymmetrical dimethyl hydrazine	Q N	I	ł	Ð		I
•		1	ł	ŊŊ	1	11
BCRL = Below Certified Reporting Limit						
IR = Indicator Range Free Summer Bins						
12012 = At of Below Upper Indicator Level 148/8 = Micrograms per gram.						
N) = Analysis not requested for original to	åfeet communiction dat	acted for				

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 8 of 30.

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. Not a Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. I = Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. ٠

			ESA-2. Burlat	Trench Sites		
[otal Bovieus	Phas	e i Analyses 48			hase II Analyses	
fotal Samples		126			105	
	,		ਦਸ਼			ਲੋ
	Frequency of	Range	Range	Frequency of	Range	Range
Analytical Groups and Analytes Detected	Detections' ¹	(6/6n)	5/(6/6 л)	Detections'	(<u>Б</u> /Б л)	<mark>շ /</mark> б/бл)
Eluoroacetic Acid	QN	ł	ł	1/22	19	2.0
Polynuclear Aromatic Hydrocarbons (PAHs)						
Fluoranthene*	QN	I	I	Q	1	ł
Methylnaphtahlene*	QN	ł	ł	QN	I	ł
Phenanthrene*	QN	1	1	QN	I	1
Pyrene*	ND			QN	**	ł
Semivolatile Halogenated Organics (SHOs)						
Hexachlorotenzene*	QN	1	I	Q	ł	1
Hexachlorobutadiene•	QN	1	ł	QN	I	1
Hexachlorocyclopentadiene	0/125	BCRL	0.3-1	0/22	BCRL	0.002-1
Pentachlorobenzene*	QN	ł	I	QN	-	I
Terrachlorobenzene*	QN	ł	1	QN	I	
Trichlorobenzene*	QN	1	1	QN	1	1
Organochlorine Pesticides (OCPs)						
Aldrin	1/125	2	0.3	1/22	0.5	0.002-0.9
Chlordane	0/125	BCRL	0.6-6	0/22	BCRL	0.02-0.2
Dichlorodiphenylethane	0/125	BCRL	0.3-0.6	0/22	BCRL	0.001-0.3
Dichlorodiphenyltrichloroethane	0/125	BCRL	0.4-2	0/22	BCRL	0.002-0.6
Dieldrin	1/125	20	0.3-0.6	1/22	0.003	0.001-0.3
Entrin	0/125	BCRL	0.3-4	0/22	BCRL	0.001-0.7
Isodrin	0/125	BCRL	0.3-0.6	0/22	BCRL	0.001-0.3
			•			

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 9 of 30.

Below Certified Reporting Limit. H BCRL

At or Below Upper Indicator Level Indicator Range R

Micrograms per gram. n BUIL

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Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 1 µ 8/8 2

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include

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multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution 2 .

effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

1

			ESA-2. Burlal	Trench Sites		
Total Borings Total Samples	Phase	e I Analyses 48 126			hase II Analyses 35	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (uq/q) ^{/2}	Frequency of Detections ^{/1}	Range Mande	CRL Range
Arsenic (IR=CRL-10)	2/125	14-32	2.5-5.2	0/16	BUIL	- 16/641 - 17
Mercury (IR=CRL-0.10)	3/125	0.15-0.81	0.050-0.06	6/0	BUIL	0.050
ICP Metals Cadmium (IR=CRL-2.0)	501/8	31.50	01150			
Chromium (IR=25-40)	0/125	BUIL	0.1-10.0	1/33	59 11 III	0.70 (
Copper (IR=20-35)	23/125	36-340	2.0-4.7		200	0.0
Lead (IR=25-40)	8/125	43-3,400	5-16	EVI	46	V.T +
Zinc (1R=60-80)	28/125	81-57,000	8.7-9.5	5/33	83-130	8.0 8.0

Pana 10 of 20 Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Fastern Study Area ۰.

Below Certified Reporting Limit. n BCRL

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Indicator Range H R

BUIL

8/81

 At or Below Upper Indicator Level
 Micrograms per gram.
 Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 7

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

2

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the inter standard for the method used. The number of detections is given, but the number of samples is not. ٠

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 11 of 30.

			ESA-3. Toxic	Storage Sites		
Total Borings	Phas	 I Analyses 86 			Phase II Analyses	
Total Samples		539			101 101	
			ਲੱ			æ
	Frequency of	Range	Range	Frequency of	Range	Rance
Analytical Groups and Analytes Detected	Detections ^{/1}	(6/6ಗ)	^{2/(б/бл)}	Detections/1	(б/бл)	(µg/g)/2
Yolatile Halocenated Organics (VHOS)						
1,1-Dichloroethane	0/19	BCRL	0.9-2	200		
1,2-Dichloroethane	61/0	BCRL.	0 3.0 6			7-6.0
1,1-Dichloroethylene	QN			NDN	BURL	0.3-0.6
1,2-Dichlorocthylene	QN	ł	ł			
1,1,2,2-7 etrachloroethane*	QN	I	ł		DCAL	2-0.0
1,1,1-Trichloroethane	61/0	BCRL	0.3-0.4	101	 	
I, I, 2-Trichloroethane	61/0	BCRL	0.3-0.4	121	2 BCR1	0.3-0.4
Carbon Tetrachloride	61/0	BCRL	0.3	0/27	BCRL	0.2.0.3
Chlorobenzene	61/0	BCRL	0.3-2	0/27	BCRL	0.1-2
Chloroform	0/19	BCRL	0.3	0/27	BCRL	13
Tetrachloroethylene	1/19	0.3	0.3	0/27	BCRL	0.2.03
Trichloroethylene	61/0	BCRL	0.3-0.5	1/27	0.8	0.3-0.5
Trichloropropene*	(IN	***	ł	QN	I	1
Methylene Chloride	61/0	BCRL	0.7-2	0/27	BCRL	0.7-2.0
<u>Volatile Hydrocarbons (VHCs)</u>						
2-Butoxyethanoi•	2	0.3	0.3	QN	ł	Ŧ
4-Hydroxy-4-methyl-2-pentanone	QN	1	1	QN	I	1
I-Methyl-1,3-cyclopentatiene*	()N	ł	I	QN	•	ł
2,2-Oxybisethanol*	<u>(</u> N	I		QN		1
2-Pentanone•	QN	1	I	QN	ł	
Bicycloheptadiene	61/0	BCRL	0.3-0.4	0/27	BCRL	0.3-0.4

= Below Certified Reporting Limit. BCRL

Indicator Range H ž

At or Below Upper Indicator Level Ħ BUL

Micrograms per gram. ı H8/8

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 2

7

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution .

effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of

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	1		ESA-3. Toxic	Storada Sites		
Total Borinos Total Samples	Pha	se I Analyses 86 29			Phase II Analyses	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/q)	CPL Rarge fun/n/2	Frequency of	127 Range	CRL Range
Dicyclopentadiene Methylcyclohexane* Methylisobutyl Ketone	0/143 ND 0/19	BCRL - ncei	0.4-1	0/65 ND	(1999) BCRL 	с,(р9/9/2
Volatile Aromatic Organics (VAOs)			1.0-0.0	0/27	BCRL	0.3-0.7
Benzene Ethylbenzene m-Xylene	61/0 61/0	BCRL	0.3 0.3-0.4	0/27 1/2/0	BCRL RCRI	0.2-0.3
o- and p-Xylene Toluene	91/0 3/19	BCRL BCRL 0.9-1	0.7-0.8 0.3-5 0.3	0/27	BCRL	0.3-0.4 0.7 0.3-5
<mark>Organosulfur Compounds</mark> Mustard - Agent Related (OSCMs)				1710	BCKL	0.2-0.3
1,4-Oxathiane Chloroacetic acid Dithiane	2/139 0/37	0.6-0.9 BCRL	0.3-6 40	2/99 3/99	2.1-9.8	1.7
Thiodiglycol Organosulfur Communes	851/c 851/b	1-90 6-30	0.3-7 3-4	4799	14-330 BCRL	00 1.5 4
Herbicide Related (OSCHs) Benzothiazole	ũ					
Chlorophenylmethyl sulfide Chlorophenylmethyl sulfone	0/139	BCRL BCRL	0.5-4	0/34 0/99	BCRL	2 0.9-4
Cinoropiconymethyl sulfoxide Dimethyldisulfide	0/139 0/19	BCRL BCRL	0.3-0.7 0.8-20	66/0 66/1	BCRL 9.3 BCRL	0.3-5 0.48 0.8-20
BCKL = Below Certified Reporting Limit.						

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Indicator Range At or Below Upper Indicator Level H

Micrograms per gram.

л 8/8 N 1

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

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Cettified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

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Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 13 of 30.

			ESA-3. Toxic	itorade Sites		
Total Borings Total Samples	Phase	i I Analyses 86 239			Phase II Analyses 44 127	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g)/2	Frequency of Detections ^{/1}	Range (uɑ/ɑ)	CRL Range (un/n/2
Organophosphorous Compounds/ GB-Agent Related (OPHGBs) Diisopropylmethyl phosphonate Dimethylphosphonic acid Methylphosphonic acid Phosphoric acid, tributyl ester* Phosphoric acid, triphenyl ester*	67/0 07/0 UN UN	BCRL BCRL 	0.05-1 0.05-2 	0/69 0/43 ND ND ND	BCRL BCRL 47	0.05-1 0.05 4.7
Organophosphorous Compounds/ Pesticide Related (OPHPs Atrazine Malathion Parathion Supona Vapona	0/139 0/139 0/139 0/139	BCRL BCRL BCRL BCRL	0.3-3 0.3-0.7 0.4-0.9 0.3-0.6 0.3-3	0/65 0/65 0/65 0/65	BCRL BCRL BCRL BCRL BCRL	0.3 0.9 3
Dibromochloropropanc	0/143	BCRL	0.3-2	0/65	BCRL	0.3-2
Organonitrozen Compounde (ONCs) Hydrazine Methylhydrazine n-Nitrosodi:n-propylamine n-Nitrosodi:nethylamine Unsymmetrical dimethyl hydrazine				88888		1111

14

Below Certified Reporting Limit. Ħ BCRL

At or Below Upper Indicator Level Indicator Range ព BUIL Ľ

Micrograms per gram. в ш

H 18/8 N

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. Ħ

= Certified Reporting Limit (CFL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. 2

There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. 11

Table ESA 2.1-2. Summary of Soil Boring Analytical Rosults in the Eastern Study Area. Page 14 of 30.

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			ESA-3. Toxic :	Stot. 18 Sites		
Total Borings Total Samples	Рлаз	e i Analyses 86 209			Fhase II Analyses 44	
Analytical Groups and Analytics Detected	Frequency of Detections/1	Range	CRL Range	Frequency of	127 Range	CRL Barne
Eluoroacetic Acid	NN	(6/64)	z,(5/61)	Detections/1	(Б/Бт)	2/(6/6rl)
Polynuckar Annatic Hydrocarlywis (PAHe)	2	I	ļ	QN	ł	1
Fluoranthene*	QN]		:		
Methylnaphtahkene•	Q	!	1	£ !	I	ł
Phenanthrene*	QN		1	<u>e</u>	I	ł
Pyrene*	UN	1]		I	I
Semivolatile Halogenated Organics (SHOs)				N	ł	1
Hexachlorobenzene*	UN ND					
Hexachlorobutadiene.	CIN CIN	ł	1	Q	1	I
Hexachlorocyclonentatiene		1 2	1	Q	ł	ļ
Pentachlorobenzene•		HCRL	0.3-0.6	0/65	BCRL	2
Tetrachlorobenzene*	2 CZ	1	ł	CIN	1	; ;
Trichlorobenzene*	(IN	1 1	ł	0N	ł	ł
Organochlorine Pesticides (OCPs)	1	1	I	<u>n</u>	ł	i
Aldrin Chlordrae	661/0	BCRL	0.3-0.6	0/65	1a,Ja	4
Dichloravtinhenvisihana	651/0	BCRL	0.6-2	0/65	BCRL	0.J
Dichlorodipheny lurichloroethane	0/139	BCRL BCRI	0.3-0.9	0/65	BCRL	2 0.6
Dieldrin Eadria	0/139	BCRL	0.3-0.8	(0/0 29/0	BCRL	0.5
Isodrin	0/139 01/0	BCRL BCBL	0.3-0.6	0/65	BCRL	0.3
		DCKL	C. 0- C .0	0/65	BCRL	0.3

Below Certified Reporting Limit. II BCRL

Indicator Range H ×

At or Below Upper Indicator Level Ħ BUIL

Micrograms per gram. N 9/81 N

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 7

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E Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. .

Tatle ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 15 of 30.

			ESA-3. Toxic S	torade Sites		
Total Borings Total Samples	Phase	I Analyses 86 239			hase II Analyses 44 127	
	Frequency of	Banne	CH			ਲ
Analytical Groups and Analytes Detected	Detections/1	(6/6rl)	ебири 2/(Бл)	Frequency of Detections ^{/1}	Hange (µq/q)	Range //n//2
Arsenic_(IR=CRL-10)	2/130	75-270	2.5-5	4/74	12-130	2.0
Mercury (IR=CRL-0.10)	0/130	BUIL	0.050-0.50	0/47	BUIL	0.050
ICP Metals						
Cadmium (IR=CRL-2.0)	1/130	4.3	0.66-0.74	2/15	71.26	01.0
Chromium (IR=25-40)	0/130	BUIL	5.2-6.5	1/75	41	0. V
Copper (IR=20-35)	0/130	BUIL	4.7-4.9	0/75	BL III.	0.0
Lead (IR=25-40)	4/130	42-140	8.4-13	2/15	45-53	0.8
Zinc (IR=60-80)	6/130	81-550	8.7-9.5	8/75	82-180	8.0

Below Certified Reporting Limit. 11 BCRL

Indicator Range Ř

At or Below Upper Indicator Level 11 BUIL

Micrograms per gram. Ħ 8/84 0N 1/

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. R

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= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. 2

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 16 of 30.

];		ESA-4. Munitions	Activity Stree		
Total Borings	Phas	e i Analyses			Phee I Andress	
Total Sampies		90 20 20				
	Ļ		ਲ		29	
Analytical Groups and Analytes Detected	Frequency of Detections/1	Range	Range	Frequency of	Ranne	E S S S S S S S S S S S S S S S S S S S
	Celecinits .	(6/6rl)	2/6/6r/)	Detections/1	(u/uii)	range
YOIJUIG Halogenated Organics (VHOs)					Isaset	- 16/611
1, 1-Dichloroethane	51/0					
1,2-Dichloroethane	110	DCKL	0.3-0.5	9/0	BCBI	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
1,1-Dichloroethylene		BCKL	0.3-0.4	0/0	BCBI	0.5
1.2-Dichloroethylene		1	I	QN		0.3
1,12,2-Tetrachloroethane	(1/n)	BCKL	0.3-0.8	0/6	BCPI	15
1.1.1-Trichlorischane		4	0.3	Z	PONE	0.3
1 1 2. Tricklossething	0/13	BCRL	0.3-0.5		1	I
	0/13	BCRL	20-E U	0/0	BCKL	0.3
	0/13	RCRI		0/0	BCRL	0.3
Chlorobenzene	0/13	BCBI	4.0-C.0	9/()	BCRL	0.3
Chloroform	0/13	BCBI	C.U FOFO	9/0	BCRL	0.3
Tetrachlorocthylene	0/13	DCDI	0.3-0.7	9/0	BCRL	20
Trichloroethylene	21/0	DCAL	0.3-0.5	9/0	BCRU	
Trichloropropene*		BLKL	0.3-0.6	0/6	RCRL	C.D
		I	ł	CIN		C .0
Methylene Chloride	9/0	BCRL	6.0		I	I
Volatile Hydrocarbons (VHCs)			C.V	0/0	BCRL	0.3
2-Butoxycthanol*	EN					
4-Hydroxy-4-methyl-2-pentanone*		ł	1	QN	ļ	ļ
I-Methyl-1,3-cyclopentadiene*	CIN CIN	ł	1	QN	ļ	
2.2-Oxybisethanol*			1	Q	i	ł
2-Pentanone*		0.0-0.0	0.3	QN	: 1	ł
Bicyclohentadiene]	ł	QN		I
	61/D	BCRL	0.3-0.8	0/0	BCRI	1 2
						C-D

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Below Certified Reporting Limit. H BCRL

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Indicator Level. н BUIL 2

At or Below Upper Indicator Level и µ8/8

= Micrograms per gram. 2

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 7

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10⁴⁷, of the internal standard for the method used. The number of detections is given, but the number of samples is not. = Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2 .

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Table ESA

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			ESA-4. Munition	Activity Stree			
Total Borings Total Samoles	Phas.	e l Analyses 66 17			Phase II Analyses		
		N/	· ·· ···		29		
Anabrical Gmune and Anabrice Detected	Frequency of	Range	Range	yes A mamilian, y	flarnju	CRL Range	
MIRINICAL CLORDS AND MIRINES DELECTED	Defections.	(6,61)	(6.64)	[leterterrs]	(E. E.A)	(µg/g)/2	
Dicyclopentadiene	0/161	BCRL	0 1 6	1 4 5 1 1			
Methylcyclohexane*	QN	ļ				0.3	
Methylisobutyl Ketone	0/13	BCRL	0403	23	IN NI		
Volatile Aromatic Organics (VAOs)							
Benzene	0/13	BCRL	0 1 1	1146.	10, 70		
Ethylbenzene	0/13	IR, BI	0 1 0 1		I I I		
m-Xylene	51/0	BCDI			IX XI	0.3	
o- and p-Xylene	0/13	IN YU		9/0		0.3	
Toliene				C D	IN KI	0.5	
	CIAN	IN KL		13/61	IN 'RL.	0.3	
<u>Organosulfur Compounds</u> <u>Mustard - Arent Related (OSCMs)</u>							
1,4-Oxathiane	0/159	BCRL	0.1.0.5	1.2.0	14.78	10	
Chloroacetic acid	QN	1	ì	1/14	IN. NI	(.) (6	
Dithiane	0/159	BCRL	0.3-2	1.70	BIC'RI,	0.3	
I hiodigiycoi	QN	1	1	61/13	IK'RL) m	
Organosulfur Compounds Herbivide Related (OSCHs)	!						
Benzouniazole		1	ł	N N	I	I	
Chlorophenylmethyl sulfide	0/159	BCRL	0.3	0/23	BCRL	0.3	
Chlorophenylmethyl sulfone	0/159	BCKL	0.4.1	0/23	BCRL	₹U	
Chlorophenylmethyl sulfoxide	0/159	BCRL	0.3-0.4	0/23	BCRL	10	
Dimethyldisulfide	0/13	BCKL	0.3-4	9/0	BCRL	0.3	
BCRL = Below Certified Reporting Limit.							

Below Certified Reporting Limit. н

Indicator Level. II Ę

At or Below Upper Indicator Level 11 BUIL

Micrograms per gram. Ħ 12/8 ND 1/

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 15

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

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Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

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			ESA-4. Munitions	Activity Sites		
Total Borings Total Samples	Phase	e I Anelyses 66 170			Phase II Analyses	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µq/q)	CRL Range (uo/o)/2	Frequency of Detections /1	29 Range	CRL Range
Organophosphorous Compounds/ GB-Agent Related (OPHGBs) Diisopropylmethyl phosphonate	05170				(5/61)	- <u>7</u> (6/64)
Dimethylmethyl phosphonate Isopropylmethyl phosphonic acid	0/159 UN	BCRL	0.5-3 2-3	0/23 0/23	BCRL BCRL	0.5 2
Methylphosphonic acid Phosphoric acid, tributyl ester	<u>a</u> g			0/14 0/14 ND	BCRL	7 7
Phosphoric acid, triphenyl ester* Organozhosybotous Compounds/	QN	ł	ł	Q		11
Pesticide Related (OPHPs)						
Arazıne Malathion	0/159 0/159	BCRL BCRL	0.5-0.7	0/23	BCRL	0.7
Parathion	0/159	BCRL	0.7-2	0/23	BCRL	0.6
Vapona	0/159	BCRL BCRL	0.5-0.9 0.3	0/23 0/23	BCRL	0.5
Dibromochloropropane	0/161	BCRL	0.3-0.9	0/23	BCRL	C.0 F U
<u>Organonitrogen Compounds (ONCs)</u>						
Hydrazine Methylbydrazine	Q Q	ł	ł	QN	ł	ł
n-Nitrosodi-n-propylamine		11	1		I	1
n-Nitrosodimethylamine	QN	I			ł	1
Unsymmetrical dimethyl hydrazine	ND	1	ł	Q	1	
						-

Below Certified Reporting Limit. н BCRL

Indicator Level. 11

 At or Below Upper Indicator Level
 Micrograms per gram. BUIL

8/8 f 2

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 1

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. .

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	HA	ase I Analyses	ESA-4. Munitions	Activity Sites	Phase I Analyses	
l otal Borings Total Samples	يو ان ان ال	66 170			ridae ii Anaryueu 11 29	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µq/q) ^{/2}	Frequency of Detections ^{/1}	Range traces	CRL Range
Eluoroacetic. Acid	QN	-		2/14	19491 3 0.8 0	- /6/6ਜ)
Pelvnuckar Aromatic Hydrocartions (PAHs)				Ì		0.7
Fluoranthene•	QN	ł	I	ÛN		
Methylnaphtahlene•	QN	ł	ł	2	1	
Phenanthrenc	Q	I	1	QN		1
Pyrenc•	QN	I	ł	QN	1	1
Semivelatile Halorenated Organics (SHOs)						l
Hexachlorobenzene"	QN	!	ł	Q		
Hexachlorobutadiene•	Q	1	1		1	i
Hexachlorocychypentadiene	0/159	BCRL				1.
Peatachlorobenzene•	QN	1	• 1	ND	PUAL .	-
Tetrachlorobenzene*	QN	I	ł	Q]	1
Trichle-obenzene•	ŊŊ	i	i	Q	1	1
Creancehlorine Pesticides (OCPs)						1
Aldrin	0/159	BCRL	0.5-0.9	540	BCDI	00
Chlordane	0/159	BCRL	1-6	120	BCBI	۲.U
Dichlorodiphenylethane	0/159	BCRL	0.3-0.5	0/23	BCRI	7 7
Dichlorodiphenyltrichlorocthane	0/159	BCRL	0.4-2	0/23	BCRI	
Dieldrin	0/159	BCRL	0.3-0.6	0/23	BCRI	+. C
Eadrin	0/159	BCRL	0.7-4	0/23	BCRL	
Iscolnia	0/159	BCRL	0.3-0.6	0/23	BCRL	0.3
BCRI = Below Confided Develope 1 and						
IL = Indicator Level.						
BUIL = At or Below Upper Indicator Level		-				

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 19 of 30.

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MICTOGRAMS per gram. ii 4 р8/8 N

Anal/sis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. ٠

			ESA-4. Munitions	L Activity Sites		
Total Borings Total Samples	ш	hase I Analyses 66 1/0		-	Phase II Analyses	
Analytical Groups and Analytes Detected	Frequency of Detections/1	Range (uo to)	CRL Range Incru/2	Frequency of	29 Range	CRL Range
Arsene (IR=CRL-10)	2/166	=	4.7.52	Delections	(б/бл)	_Z /Б/Бл)
Mercury (JR=CRL-0.10)	4/166	0.11-0.14	0.050-0.070	500	BUIL	4.7
ICP Metals				67/0	BUIL	0.050
Cadmium (IR=CRL-20)	0/1/9	RIN	0 50 0 00			
Chromium (IR=25-40)	0/1/0	BUIL	UK-0-00	11/0	BUIL	0.92
Copper (18=20-35)	0/1/51	36-500	0.4.8.4	110	BUIL	7.2
	0/1/1	41-200	16-17	2012	5	5.0
2.1MC (1N =00-80)	18/170	82-160	16-28	4/17	4 3	17
				•	•	
BCRL = Below Certified Reporting Limit.						
BUIL = At or Below Upper Indicator Level						
HEVE * MINORGAME PET PLAN. NO * Analysis not removable for minimal						
/1 Effection terreterite the total mumber		mot detected to hontarge	t compounds subsequer	uly considered in analyte		

multiple detectumes of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 12 * Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. • There is no CRL for tentauvely identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

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		ESA.	5. Demilitarizat	ion Activity Str.		
Total Borings	Phase	Analyses		d	hase II Analvana	
Total Sumplies		o 81			7	
Analytical Groups and Analytes Detected	Frequency of Datactions ^{/1}	Range	CRL Range	Frequency of	Range	CR Rance
Yolatic Halosenated Organics (VHO)		16.64	- 7 6/64)	Detections ^{/1}	(6/6rl)	2/(6/6π)
1,1-Dichloroethane	1/0					1
1,2-Dichlorrethane		BURL	0.5	Q	ł	ł
1.1-Dichloroethylene		BURL	0.4	QN	ſ	1
1.2-Dichlorric thylene			ł	QN	ł	1
1.1.2.2.Tetrachlance		BCRL	0.8	QN	ł	!
1.1.1.Trehenebare	23	1	ł	QN	1	ł
112-Trichlowethane	1/1	BCRL	0.5	ŊŊ	I	1
Carbon Terresburge		BCRL	0.6	ŊŊ	1	I
Children reas		BCRL	0.4	QN		1
China contaction		BCRL	0.3	CZ.	I	1
	141	BCRL	0.7		I	ł
I cummioncelly icre	0/1	BCRL	0.5		1	I
I nchiorcethylene	1/0	BCRL	0.6			ł
Trehkropresee	ND				I	1
Methalene Chlorida			I	nn N	ł	1
	ON ON	1	ł	QN]	
<u>Volatile Hydrocarbons (VHCs)</u>				1	ļ	ł
2-Buloxycthamol•	űz					
4-Hydroxy-4-methyl-2-pentanone	QZ		1	ÔŽ	1	3
I-Methyl-1.3-cyclopentatione*	C2	l	1	QN	1	I
2.2. (Dryhierhand)		1	1	ND	ł	Ĩ
7. Pentarcone	2	ł	1	CIN	ł	n an
	N	ł	ł	ND		ł
Bicyclohepuxdiene	0/1	BCRL	8.0	2 2	ł	1
					1	1
BCRL = Active Confired Dominics Linit						
In I						

Indicator Range Ħ

At or Below Upper Indicator Level 11

Mkrograms per gram. Ħ

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include

multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. に

2

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit hased on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is joint. .

	į	ESI	A-5. Demilitarize	tion Activity Site		
Total Borings Total Sampies	steny	i i Analyses 6 18			hase II Analysee 7	
	Frequency of	Range	CPI. Range	Frequency of	Bame	CR
Analytical Groups and Analytes Detected	Detections/1	(Б,Бл)	^{2/(6/6л)}	Detections ^{/1}	(5/5rl)	2/6/6/)
Dicyclopentadiene	0/18	BCRL	ç	0/14	BCRL	11
Methykcyclohexane*	QN	ł	I	QN		5 1
Methylisobutyl Ketone	0/1	BCRL	0.4	Ð	ł	1
Volatile Aromatic Organics (VAOs)						
Benzene	1/0	BCRL	-	QN	1	I
Ethylbenzene	1/0	BCRL	0.4	Q	1	
m-Xylene	1/0	BCRL	0.3	QZ	I	! 1
o- and p-Xylene	1/0	BCRL	0.5	2	1	1
Toluene	1/0	BCRL	0.3	QN	ł	1
Organosulfur Compounds Musturd - Avent Related (OSCMs)					۰.	•
1,4-Oxathune	0/18	BCRL	0.5	F1/0	Iaca	5.0
Chloroacetic acid	QN	1	; 1	CN CN	DCNL I	<u>.</u> 1
Duthiane	0/18	BCRL	2	0/14	BCRL	10
Thiodigiycol	ND	ł	ł	QN	. 1	1
Organosulfur Compounds Herbouck Related (OSCHs)						
Benzothiazole	Ð	ł	I	Ę]	
ChlorophenyImethyl sulfide	0/18	BCRL	0.3	0/14	BCRL	12
Chlorophenyimethyl sulfone	8170	BCRL	-	0/14	BCRL	4.0
Chlorophenylmethyl sulfoxide Dimethyldisulffale	81/0	BCRL	4 .0	0/14	BCRL	0.3
	5		r	2	ł	ł

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Easiern Study Area. Page 22 of 30.

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Below Certified Reporting Limit. Ħ BCR R

Indicator Range 54

At ur Below Upper Indicator Level Micrograms per gram. BUIL

84 92

Analysis not requested for original larget compounds/not detected for nontarget compounds subsequently considered in analyte groups. Ħ

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 1

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution 2

effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit hased on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. .

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 23 of 30.

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		ESA-5.	Demilitarizati	on Activity Site			
	Phase	Analyses		Чd	iase II Analyses		
Total Borings		6 18			7 14		
1 Oldi Sainpies			ਲ			ਲ	
	Frequency of	Range	Range	Frequency of	Ranga	Range	
AnaMical Groups and AnaMes Detected	Detections'	(6/61)	- (6/6d)	Defections' -	(6/64)	15/610	
Organophosphorous Compounds/ CB. A new Belated (OPHCBs)							
Diisonmovimethyl nhosphonate	81/0	BCRL	£	0/14	BCRL	0.5	
Dimethyl methyl phosphonate	0/18	BCRL		0/14	BCRL	6 0	
Isomovimethyl phosphonic acid	QN	ł	I	0/14	BCRL	. 2	
Methylphosphonic acid	QN	1	I	0/14	BCRL	2-20	
Phosphoric acid, tributyl ester	QN	1	ł	Q	1	ł	
Phosphoric acid, triphenyl ester	ND	ł	1	QN	ł	1	
Organophosphorous Compounds/ positivite Referent (OPHPs)							
A trazine	0/18	BCRL	0.5	0/14	BCRL	0.7	
Malathion	0/18	BCRL	2	0/14	BCRL	0.6	
Parathion	0/18	BCRL	2	0/14	BCRL	0.0	
Supona Viscona	0/18 0/18	BCRL	0.3	0/14	BCRL	0.3	
Dibromochloropropane	0/18	BCRL	0.6	0/14	BCRL	0.3	
Organoniuogen Compounds (ONCs)	ÛN	1	1	QN	۱	1	
n yu azuk Methylhydrazine	Q	1	1	QN	1	I	
n-Nitrosodi-n-propylamine	QN	1	1	2	ł	ł	
n-Nitrosodimethylamine	DD	i	I	Q i	1	I	
Unsymmetrical dimethyl hydrazine	QN	t	I	CN NO	•		

= Below Certified Reporting Limit. BCRL

Indicator Range H e,

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. BUIL = At or Below Upper Indicator Level µg/g = Micrograms per gram. ND = Analysis not requested for original

7

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution 2

effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. .

		ESA	V-5. Demliitariza	tion Activity Site		
Total Borings Total Samples	Phase	i 1 Analyses 6 18			hase II Analyses	
AnaMical Groups and AnaMies Detected	Frequency of Detections ^{/1}	Range (µg/q)	CRL Range (III0/0/ ^{/2}	Frequency of Datactions ^{/1}	Range Incol	CRL Range
Elucroacetic Acid	QN	-		1/14	15/64	- /6/61)
Polynuckar Aromatic Hydrocarbons (PAHs)						0.4
Fluoranthene*	QN	I	ł	ŰZ		
Methylnaphtahlene*	QN	!	!		1	ł
Phenanthrene*	QN	ł	I		ł	ł
Pyrene*	QN	I	1	29	1	I
Semivolatile Halogenated Organics (SHOs)				8		1
Hexachlorobenzene*	GN	ļ				
Hexachlorobutadiene*	Q	1	1		ł	ł
Hexachkorocychopentadiene	0/18	BCRL	!	FIN		۱.
Pentachlorobenzene*	QN	1	- 1		BURL	_
Tetrachlorobenzene*	QN	i	1	Ĩ	ł	I
Trichlorobeazene*	QN	I	I	Q	1	1 1
Organochlorine Pesticides (OCPs)						
Aldrin	0/18	BCRI	0.5	V114		
Chlordane	0/18	BCRL	6	0/14	BCBL	۷.0
Dichlorodiphenylethane	0/18	BCRL	0.5	0/14	BCRL	۲ د د
Dichlorodiphenyltrichloroethane	0/18	BCKL	2	0/14	BCRL	40
Deldrin	0/18	BCRL	0.6	0/14	BCRL	10
Endin	0/18	BCRL	4	0/14	BCRL	0.7
Isodrin	0/13	BCRL	0.6	0/14	BCRL	0.3

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 24 of 30.

Below Certified Reporting Limit. n BCRL

Indicator Range 11 R

At or Below Upper Indicator Level N BUIL

8/81 NO

 Micrograms per gram.
 Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 1

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of ٠

detections is given, but the number of samples is not.

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 25 of 30.

		ESI	A-5 Demilitarize	lon Astronom Ob		
Total Borings Fotal Samples	Phase	I Analyses 6 18		P P	hase li Analyses 7	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g)/2	Frequency of Detections/1	14 Range (µg/g)	CRL Range fun/nV2
VISCUIC (IK=CKT-10)	81/0	BUIL	5.2	QN		Index
Aercury (IR=CRL-0.10)	0/18	BUIL	0.050-0.070	ũ	ł	I
<u>CP Meals</u>) -	1	1
Cauttuni (IK=CKL-2.0) Chromium (IR=25.40)	0/18	BUIL	0.51	QZ	ł	
Copper (IR=20-35)	81/0	BUIL	7.2-7.4	QN	5	i I
Lead (IR=25-40)	0/18	BUIL BUIL	4.7	QZ	I	ł
	6/18	82-110	28	Q	11	11

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Below Certified Reporting Limit. 11 BCRL

Indicator Range 11 R

At or Below Upper Indicator Level Micrograms per gram. я 11 BUIL

N N N N

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

Eraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. •

Tetal Deriver	,	Phase Ansiver	Belence of Ir	ive*'igations		
rorai borings Totai Samples	ľ	32 113			Phase II Analyses 21	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (uo/o)	CRL Range	Frequency of	51 Range	CRL Ranoe
Yolatile Halogenated Organics (VHOs)		IR.E.I	- 76/61)	Detections ^{/1}	(б/бл)	2/(6/6n)
1,1-Dichloroethane 1,2-Dichloroethane	10	BCRL	0.3-0.9	5/0		·
1,1-Dichloroethylene	â	BCRL 	0.3-0.4	0/5	BCRL	0 7 0 6
1,4-Dichloroethylene 1,1,2,2-Tetrachlorocthane*	9/0	BCRL	0.3-0.8	Q X		31
1,1,1-Trichloroethane	0N 010	ł	1	â	PCKL	2
1,1,2-Trichloroethane Carten Terrechloride	10	BCRL	0.3-0.5 0 3-0.6	C.S	BCRL	0.4
Chlorobenzene	50	BCRL	0.3-0.4	c/0 2/0	BCRL BCRL	0.4
Chloroform Terret Land	LI0	BCRL	0.3	0/5	BCRL	0.2 2
renation of the second se	<i>L</i> 0	BCRL	0.3-0.5	0/5 2/0	BCRL	0.3
Trichloropropene*	10	BCRL	0.3-0.6	0/5	BCRL	0.2
Methylene Chloride.	0/5	RCP1	0.5	QN	ł	31
<u>Volatile Hydrocarbons (VHCs)</u>	l T	PUNL	0.3-0.7	0/5	BCRL	2
2-Butoxyethanol*	QN	**	,	-		
1-11.3	2	1	1	a a	I	1
2,2-Oxybisethanol*	15/15		1	Q	1 1	1
2-Pentanone*	Q		1	Ð	ł	
Dicycloneptadiene	<i>LI</i> 0	BCRL	0.3-0.8	0/5 0/5	BCRL	
BCRL = Below Certified Reporting Limit.						

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 26 of 30.

= Indicator Range ž

At or Below Upper Indicator Level H BUIL

= Micrograms per gram. µ8/8

2

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. 7

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution = Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

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effect for some sample analyses. = There is no CRL for tentaively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not.

-			Balance of h	nvestigations		
Total Borings Total Samples	Phae	se i Analyses 32 113			hase II Analyses 21 53	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g)/2	Frequency of Detections ^{/1}	Aange Range (un/o)	CPt Range fundrov2
Dicyclopentadiene Methyliveloherane*	0/417 ND	BCRL	0.3-6	0/15	BCRL	0.3-1
Methylisobutyl Ketone	<i>L</i> 0	BCRL	0.3-0.5	QN 570	BCR1.	16
<u>Volatile Aromauc Organics (VAOs)</u> Benzene	01	BCRL	0.1.1	YU.		
Ethylbenzene m-Xvlene		BCRL	0.3-0.4	0/5 2/0	BCRL	0.2
o- and p-Xylene Totuene	L/0	BCRL 0.5	0.3-0.7	6/0 2/0 2/0	BCRL BCRL	5.0 5
Organosulfur Compounds Mustard - Arent Related (OSCMs)		1			-	7.0
1,4-Oxathiane Chloroacetic acid	0/417 ND	BCRL —	0.3-6	0/15 0/1	BCRL	0.3
Dithiane Thiodiglycol	0/117 0/16	BCRL BCRL	0.3-7 3	0/15	BCRL	0.3-0.4 4
Organosul fur Compounds Herbicide Related (OSCHs)				<u>.</u>		
Benzothiazole Chlorophenylmethyl sulfide	ND 0/417	BCRL		QN S1/0		
Chlorophenylmethyl sulfone Chlorophenylmethyl sulfoxide	0/417 0/417	BCRL	0.3-7	0/15	BCRL	0.3-0.4
Dimethyldisulfide		BCRL	0.3-4	61/0 2/0	BCRL	20.3
BCRL = Below Certified Reporting Limit. IR = Indicator Range BUIL = At or Below Upper Indicator Level µg/g = Micrograms per gram. M) = Analysis port requested for activity						
I = Fraction represents the total number	r of detections of an a	uncertaint for momentation to the	e number of analyses	nuy considered in anaiyic conducted in a distinct s	e groups. amhle This value doe	e not include

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 27 of 30.

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used.

2

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. +

Section 2.

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Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 28 of 30.

			Belance of Ir	Vestigations		
Total Borings Total Samples	Phas	 I Analyses 32 113 			Phase II Analyses	
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (µg/g)	CRL Range (µg/g)/2	Frequency of Detections/1	21 Range Anorit	CRL Range
Organophosphorous Compounds/ GB-Agent Related (OPHGBs) Diisopropylmethyl phosphonate Dimethylmethyl phosphonic acid Methylphosphonic acid Pho 'phoric acid, tributyl ester* Phosphoric acid, tributyl ester*	0/417 0/234 ND ND ND	BCRL BCRL	0.1-3 2-3 	0/15 0/15 0/2	BCRL BCRL BCRL -	- 40/94 0.5-1 0.05-2 -
Organophosphorous Compounds/ Pesticide Related (OPHPs) Atrazine Matathion Parathion Supona Vabona	0/417 0/417 0/417 0/417	BCRL BCRL BCRL	$\begin{array}{c} 0.3-0.7\\ 0.3-2\\ 0.4-2\\ 0.3-0.9\\ 0.3-0.9\end{array}$	ND 0/15 0/15 0/15	BCRL BCRL BCRL BCRL	0.3-0.7 0.6-0.9 0.6-0.9 0.5-0.6
Dibromochloropropane	0/419	BCRL	0.3-3 0.005-0.6	0/15 0/15	BCRL BCRL	0.3
Organoni-rogen Compounds (ONCs) Hydrazine Methylhydrazine n-Nitrosodi-n-propylamine n-Nitrosodimethylamine Unsymmetrical dimethyl hydrazine	<u> </u>	1111	1111	2 2 2 2 2 2	11111	
BCRL = Below Certified Reporting Limit. IR = Indicator B-no.						

At or Below Upper Indicator Level Ħ

Micrograms per gram. Ņ 108 8/81 NB/81

= Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups.

= Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when more than one analytical method has been used. 2

 Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses.
 There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. *

			Balance of Ir				
Total Borings Total Samples	Phase	I Analyses 32 113			hase i Analyses 21		
Analytical Groups and Analytes Detected	Frequency of Detections ^{/1}	Range (uq/q)	CRL Range fun/n/2	Frequency of Datasized/1	21 Range	CRL Range	
Eluoroacetic Acid	ND			1/4	(5/5il)	- <u>Д</u> б/бл)	1
Polynuckar Aromatic Hydrocarbons (PAHs)					t.1	2.0	
Fluoranthene*	QN	:					
Methylnaphtahlene*	QN	!	ł	an	I	1	
Phenanthrene*		ł	ł	QN	I	ł	
Purne*		ł	ł	QN	I	I	
		i	1	QN	i	ł	
Semivolatile Halogenated Organics (SHOs)						ł	
Hexachlorobenzene*	_	50		Ĩ			
Hexachlorobutadiene•	CN		C.U	n	ł	1	
Hexachlorocyclonentadiene			!	N	1	ł	
Pentachlorohenzene*		BLAL	0.3-1	0/15	BCRL	0.6-1	
Tetrachlarohenven	2	ł	i	QN		1	
Trichlorobanaanat		1	ł	QN	1	1	
	0N	I	1	QN	ł	5	
Organochlorine Pesuicides (OCPs)							
Aldrin	0/417	BCRL	0.3-0.9	0/15	IaJa	0010	
Chicrdane	0/417	BCRL	0.6-6	51/0	DCNL	0.3-0.9	
Dichlorodiphenylethane	0/417	BCRL	03.06			7 2 2 2	
Dichlorodiphenyltrichloroethane	0/417	BCRL.	04.2		BCRL	0.5-0.6	
Dieldrin	0/417	BCRL	0.3-0.6	51/0	BCBL	0.4-0.5	
Erkirin	0/417	BCRL	0.3-4	51/0	BCDI	(.) ()) (
Isodrin	0/417	BCRL	0.3-0.6	0/15	BCRL	1.0-0.0	

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area. Page 29 of 30.

ļ

Below Certified Reporting Limit. н BCRL

0.3

BCRL

Indicator Range 11 X

At or Below Upper I vdicator Level IJ BUIL

Micrograms per gram. Ĥ µ8/8

Analysis not requested for original target compounds/not detected for nontarget compounds subsequently considered in analyte groups. H 2 1

Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct sample. This value does not include multiple detections of a specific analyte in the same sample, which occasionally has occurred when mere than one analytical method has been used.

= Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution 2

effect for some sample analyses. = There is no CRL for tentatively identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not. .

fotal Borings fotal Samples	Рна	se I Anklyses 52 113		c.	hase II Analysea 21	
unalytical Groups and Analytes Detected	Frequency of Detections/1	Range (µg/g)	CPL Range (µg/g) ^{/2}	Frequency of Detections ^{/1}	Lange Lange (uo to)	CAL Range
Arsenic (IR=CRL-10)	5/470	11-21	2.5-5.2	221	25	2.0-4.7
<u> </u>	5/464	0.12-0.45	0.050-0.070	1/23	BUIL	0.050
CP Metals Cadmium (IR=2.0)	1/435	6.6	0.51-0.92	500	Ela	
Chromium (IR=40)	3/470	46-62	52.12		DU IL	0.10-0.52
Copper (IR=35)	1/470	40-120	4.74.9		307	4 5.4 0
l ead (IR=40)	3/470	11-60	8.4-17	101	IKO	80-17
Zinc $(IR=80)$	0/4/51	081-18	8.7-28	18/1	88-120	16

Table ESA 2.1-2. Summary of Soil Boring Analytical Results in the Eastern Study Area Pane 30 of 30

Below Certified Reporting Limit. Ħ HCRL

Indicator Range Ħ ž

BUIL = At of Below Upper Indicator Level µg/g = Micrograms per gram.

^{*} Analysis not requested for original target compounds/not detected for nontarget compounds subsequently consudered in analyte groups. 2 ~

⁼ Fraction represents the total number of detections of an analyte in relation to the number of analyses conducted in a distinct semple. This value does not include multiple detections of a specific analyte in the same sample which occasionally has included when more than one analytical method has been used.

Certified Reporting Limit (CRL), or detection limits used among laboratories conducting analyses for specific sites shown on table. Variability also may reflect dilution effect for some sample analyses. 2

⁼ There is no CRL for intrinvely identified compounds. The value shown is a detection unit based on 10% of the internal standard for the method used. The number of detections is given, but the number of samples is not •
Analysis/Analytes	Level of	Analytical Method	Cartified Decenter-
	Certification		Crimed Acporting Limit (μg/l)
<u> Yolatile Halozenated Organics</u>	Quantitative	PACK-GC/Hall	
1,1 - Dichloroethane			•
1,2 - Dichloroethane			1.2
 I.I Dichloroethytene 			10.0
1,2 - Dichloroethylene			1.1
1,1,1 - Trichlonethane			5-7 L 1
I.I.Z - Inchloroethane			
Carbon utrachloride			0.1
Chlorobenzene			850
Chloroform			
Methy kine chloride			5.0
I ctrachlartethy icne			11
Trichloraethylene			1.2
<u>ćolatile ilydrocarbons</u>	Quantitative	CAP-GC/FID	
Detectorentelione			
Methylisobutyl ketone			9.3 13
colatile Aromatics	Quantitative	PACK-GC/PID	
Berucne Eulylberucne Tolucne Xulune (m.)			1.3 1.0 1.2
Xykree (0-, p-)			1.4 2.5
Dreanosultur Compounds Austard-Arent Related	Quanticute	PACK-GC/FPD-S	
Dithiane 1,4 - Oxathiane		Ē	2.0

Table ESA 2.2-1. List of Chemical Analyses Conducted on Task 4 Water Samples. Page 1 of 3.

ISA Table 2.2-1/ESA 1 tev. 5/10/89

Analysis/Analytes	Level of Certification	Analytical Method	Certified Reporting Limit (µg/l)
<u> Leanosulfue Compounds</u> lechicute Related	Quantum V	PACK-GC/FPD-S	
Dunethykulfake Chkwaptenyimethyl sulfake (CPMS) Chkwoptenyimethyl sulfane (CPMS) Chkwoptenyimethyl sulfaxide (CPM	(02) (50)		1.8 4.2 1.3
lteenoodosodotous Compounds iB-Azent Related	סאוזרוואוניוע)	PACK-GC/FPD-P	
Dristynopy Irweidyl phreistwaare Drimethy Imethyl płasętwaare			10 15
TRUE	Quantative	CAP-GC/ECD	
Debneschlercyscyme			0.13
Dreanachlarine Pesticides	Quantitative	CAP-GC/ECD	
Aktna DOE DOT Debtua Endra Hexa Akaricyckyentatiene istaina			0.070 0.050 0.070 0.050 0.050 0.070 0.050
Arsenic	ON-DEPENDENCE OF	AA-hythde fumace	3.9
Mercara	Quantitative	Cold vapry atomic advertation	0.20

Table ESA 2 2-1. List of Chemical Analyses Conducted on Task 4 Water Samples. Page 2 of 3.

ESA Table 22 HESA 1 Rev. 5, HEBV

Certified Reportiug Limit (µg/1) 20 9 7 9 20 20 1 9 20 20 20 Inductively coupled plasma Analytical Method Table ESA 2.2-1. List of Chemical Analyses Conducted on Task 4 Water Samples. Page 3 of 3. Level of Certification Quantitative Quantitute Analysis/Analytes Cartmuun Chromuun **Other Analytes** Contra Line ICP Meials

SPIN/SEPAN Magnesium Cakium Chlonde Fluonde

Polassium Sodium Sulfate

Inductively coupled plasma fon chromatography fon chromatography Inductuvely coupled plasma fon chromatography Inductively coupled plasma Auto analyzer AA Ilame

4,800 500 1,200 1,300 1,300 10,000

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Analysis/Analytes	Certification Level	Method	Certified Reporting Limit (µg/l)
Organochlorine Pesticides	Quantitative	CAP-GC/ECD	
Aldrin Endrin Dieldrin Isodnin Hexachlorocyclopentadiene DDE DDT Chlordane			0.083 0.060 0.055 0.056 0.083 0.046 0.059 0.15
Volatile Organohalogens	Quantitative	PACK-GC/Hall	
Chlorobenzene Chloroform Carbon Tetrachloride 1,2 - Dichloroethylene Trichloroethylene 1,1 - Dichloroethylene 1,1 - Dichloroethylene 1,2 - Dichloroethane 1,2 - Dichloroethane 1,1,2 - Trichloroethane 1,1,2 - Trichloroethane 1,1,2 - Trichloroethane			1.8 1.7 4.9 1.8 1.3 2.8 1.9 1.9 2.1 1.1 1.6 2.5
Organosulfur Compounds	Quantitative	PACK-GC/FPD-S	
Chlorophenylmethyl sulfone (CPMSO ₂) Chlorophenylmethyl sulfoxide (CPMSO) Chlorophenylmethyl sulfide (CPMS) Dithiane 1,4-Oxathiane Dimethyldisulfide Benzothiazole			2.2 2.0 1.1 1.6 1.4 1.2 1.1
olatile_Aromatics	Quantitative	PACK-GC/PID	
Benzene Ethylbenzene Toluene m-Xylene o-, p-Xylene			1,9 0.62 2.1 1.0 1.3

Table ESA 2.2-2. List of Chemical Analyses Conducted on Task 44 Water Samples. Page 1 of 3.

ESA Table 2.2-2 5/10/89 11:12 AM

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Table ESA 2.2-2. List of Chemical Analyses Conducted on Task 44 Water Samples. Page 2 of 3.

Quantitative	CAP-GC/FID	
Quantitative	CAP-GC/FID	
		9.3 13
Quantitative	PACK-GC/FPD-P	
		10 19
Quantitative	CAP-GC/ECD	
		0.13
Quantitative	Inductively Coupled Plasma	
	AA-Hydride EPA 245 Cold Vapor Ion Chromatograph	500 500 760 1,300 5.2 6.0 7.9 19 20 2.5 0.36 1,600 1,000 5,000
	Quantitative Quantitative	Quantitative CAP-GC/ECD Quantitative Inductively Coupled Plasma AA-Hydrid: EPA 245 Cold Vapor Ion Chromatograph

ESA Table 2.2-2 5/10/89 10:32 AM

Certification Level	Method	Certified Reporting Limit (µg/l)
Samiauastitatius	CONS	
Semiquantizave	GC/MS	
		2 3 1 4 1 0.7 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2
Semiquantitative	GC/MS	
·		5 6 5 8 20 10 20 5 5 10 5 6 10 5 6 10 8 10 6 8 10 6 8 10 7 9
	Certification Level Semiquantitative	Certification Level Method Semiquantitative GC/MS

Table ESA 2.2-2. List of Chemical Analyses Conducted on Task 44 Water Samples. Page 3 of 3.

ESA Table 2.2-2 5/10/89 10:32 AM

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					UT AIRE SUIIACE WAIEL Pag	6 1 01 2.	
Compound Group	Samplin Locatio	80 E	Compound Detected	Frequency of Detections	Concentration Range (μg/l)	Date of Detection	
<u>Yolatile Italogenated</u>	Organics						
	05-001 08-001 24-007 24-007	08ADD* 13DCC*		0/4 0/10 0/6	BCRL BCRL BCRL		·
	30-002 31-001 31-002		Chlorobenzene Chloroform	0/5 1/4 1/4	BCRL 1.8 3.5	— 6/30/86 07/01/86	
Organophosphorous.	Compounds.	GB-Acent B	clated				
	05-001 08-001 24-002 30-002 31-001 31-001	0%ADD• 13DCC•	Dimp Dimp	0/4 1/10 1/6 0/1 0/4 0/4	BCRL 11 BCRL BCRL BCRL BCRL	. 12/20/85 12/20/85 	
Organochlorine Pest	cides						
	05-001 08-001 08-001 24-007 30-002 31-001	08ADD* 08ADD*	Akhrin Dickdrin Akhrin	0/4 1/10 1/10 0/5 1/4	BCRL 0.20 0.060 Σ=0.26 BCRL BCRL 0.080	• 11/22/85 	
	31-002 24-002 24-002	13DCC• 13DCC•	Akhin Dicidin	()/4 1/6 1/6	BCRL 0.20 0.080 Σ=0.28	11/22/85	

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BCRL = Below certified reporting limit DIMP = Diisopropylmethyl plusphonate * = Alternate name for sumpling locations. ** = Summution of the group.

Table ESA 2.2-3/ESA-1 5/10/89 10:09 AM

Table ESA 2.2-3 Summary of Target and Significant Nontarget Compounds Detected in Eastern Study Area Surface Water

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				area Derevied III Eastern Stu	dy Area Surface Water Pa	age 2 of 2.	
Compound Group	Sampli Locatio	8 E	Compound Detected	Frequency of Detections	Concentration Dance (1.20)	Date of	
Arsenic					vange (J.g/1)	Detection	•
ICE Metals	05-001 08-001 24-002 30-002 31-002 31-002	(%ADD• 13DCC•	Arsenic Arsenic Arsenic Arsenic Arsenic	2222882	3.8 6.6 3.5 5.0 BCRL 7.3	05/15/87 04/02/86 06/17/87 05/15/87 05/27/87	
	05-001 08-001 08-001 08-001 08-001 08-001 24-007 31-002 31-002 31-002	08ADD+ 08ADD+ 08ADD+ 08ADD+ 08ADD+ 13DCC+	Zinc Cadmium Copper Chromium Lead Zinc Copper	82222233828	BCRL 24, 25 14 13 13 21 13 21 21 21 81, 30 BCRL 10 BCRL	04/02/86, 12/16/86 03/26/87 03/26/87 10/16/87 10/16/87 10/16/87 10/16/87	

able ESA 2.2.3/ESA-1 /10/89 10:09 AM

³CRL = Below certified reporting limit DIMP = Diisopropylmethyl phosphonate • = Alternate name for sampling locations. •• = Summation of the group.

Die ESA 2.3-1 Summary of Target and Significant Nontarget Communication Imalytical Total Wetts Attuvial Total Samples 52 Total Samples 61 Total Samples 52 Total Samples 61 Total Groups/Analytes Frequency of Concentrati Total Latile Halogenated 0145 BCRL Latile Halogenated 0145 BCRL 0.64 Li Dickloroethane 0/45 BCRL 0.64 Li Li 0/45 BCRL 0.64 Li Li 0/45 BCRL 0.74 - 15 Chorobenzene 0/45 0.745 <	oound Detections in Easterr (in Range (µg/l) 1.1 - 1.9 1.1 - 1.9 1.1 - 1.9 1.1 - 1.7 1.1 - 1.7 1.0 - 5.0 1.0 - 5.0 1.3 - 2.8 0.50 - 1.9 2.5 - 5.0 1.3 - 2.8	Study Area Groundwi Frequency of Detections 0/94 0	aler. Page 1 of 5. Denver 37 95 Concentration Range (µg/l) BCRL BCRL BCRL BCRL BCRL BCRL BCRL BCRL BCRL BCRL BCRL 20 4.3 2.4 2.4	CRL Range (µg/l) 0.73 - 1.9 0.61 - 2.1 1.1 - 17 0.76 - 1.8 0.76 - 3.0 0.78 - 5.0 1.0 - 2.4 0.78 - 5.0 1.0 - 2.4 0.75 - 2.8 0.56 - 5.5
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Volatile Hydrocarbons (VHCs)

I QN	ND	I QN) ON	I QN	0/46 BCRL	ND	
+ Construction C	2-buwycumu 4 Uudawy A-methyl-2-methol	4-nyuruxy-1-1 acvelopentadiene			Bicyclohepiatiene Disserbrastatiene	Methylcyclohexane*	Michightsought activity

9.3 - 22 4.9 - 13

BCRL BCRL

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* = Significant nontarget compound
 BCRL = Below Certified Reporting Limit
 ND = None Detected

Table ESA 2.3-1/ESA-1 5/19/89

Total Wells All1 Total Samples 5	Analytes Frequency of Concen Detections Range	czanics (YAOs) - 1/41 - 2 - 0/41 - BC -	0/46 BC ND 0/46 BC 0/46 D	OSCH5) auffide 0/19 BCI auffide 0/46 BCI auffone 7/46 3.0 -
Alluvial 19 52	oz ncentration inge (μg/l)	2.4 BCRL BCRL BCRL 8.6	BCRL 	BCRL BCRL 3.0 - 4.3
	CRL Range (µg/l)	1.3 - 1.9 0.62 - 1.3 1.0 - 1.4 1.3 - 2.5 1.2 - 2.1	1.4 - 2.0 	1.1 1.0 - 1.3 2.2 - 4.7
	Frequency of Detections	12/91 0/91 0/91 0/91	0/92 ND 0/92 ND	0/28 0/92 0/92
Denver 37	95 Concentration Range (µg/1)	1.4 - 20 BCRL BCRL BCRL BCRL	BCRL 	BCRL BCRL NCRI
	CRL Range (110/1)	1.3 - 1.9 0.62 - 1.3 0.62 - 1.3 1.0 - 2.0 1.3 - 3.2 1.2 - 2.8	1.4 - 14 1.1 - 16 	1.1 0.1 - 10

ii C E Table ESA 2.3-1 Summary of Target and Significant Nontarget Compound Detections in Easte

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Significant nontarget compound
 BCRL = Below Certified Reporting Limit
 ND = None Detected

Table ESA 2.3-1/ESA-1 5/19/89

Total Total	Wells Samples	Alluviat 19 52			Denver 37 95	
Analytical Groups/Analytes	Frequency of Detections	Concentration Range (μg/l)	CRL Range (µg/1)	Frequency of Detections	Concentration Range (µg/l)	CRL Range (μg/l)
Organophophorous Compounds. GB-Azent Related (OPUGBs)						
Diisopropylmethyl phosphonate Dimethylmethyl phosphonate Isopropylmethyl phosphonic acid Methylphosphonic acid, Phosphoric acid, Phosphoric acid, tributyl ester* Phosphoric acid, tributyl ester*	0/46 0/46 0 N N N N N N N N N N N N N N N N N N N	BCRL 	10 - 11 15 - 30 	16/0 16/0 0 N N N N N N N N N	BCRL BCRL	10 - 11 15 - 30
Organophosphorous Compounds. Pesticide Related (OPHPs)					•.	
Atrazine Malathion Parathion Supona Vaponu	222 222			<u> </u>] [[]]
DBCP	1/46	0.16	0.13	2/95	0.15 - 0.75	0.13 - 0.20
Organonitrogen Compounds (ONC Caprolactum* Hydrazine Methylhydrazine n-Nitrosodimethylamine n-Nitrosodi-n-propylamine Unsymmetrical dimethyl hydrazine	6 1 1 1 1 1 1 1 1 1 1	1 1 1		<u> </u>		

Significant nontarget compound
 BCRL = Below Certified Reporting Limit
 ND = None Detected

Table ESA 2.3-1/ESA-1 5/19/89

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I dove ESA 2.3-1 Summary of Ta	Irget and Significant	Nontarget Compound	d Detections in Eastern	Study Area Groundw	ater. Page 4 of 5.	
Т.	tal Wells tal Samples	Alluvial 19 52			Denver 37	
Analytical Groups/Analytes	Frequency of Detections	Concentration Range (µg/l)	CRL Range (µg/l)	Frequency of Detections	Concentration Range (µg/l)	CRL Range (119/1)
Eluoroscetic Acid	QN	-	-	Q		
<u>Polynuclear Aromatic Hydrocar</u>	bons (PAHs)]
Fluoranthenc* Methylnapthalene* Phenanthrenc*	Q Q Q			882	11	11
r yitike	Q	1	ļ	QN		
Semi-Volatile Halogenated Org	anics (SHOs)					
Hexachkorobenzene* Hexachkorobutadiene*	QN QN	11	ł	Q		1
Hexachlorocyclopentadiene Pentachlorobenzene*	0/42 ND	BCRL	0.070 - 0.21	11/0	BCRL	0.070 - 0.15
Tetrachlorobenzene* Trichlorobenzene*	2			Q Q		1
	ND	I	*	ŊŊ	1	

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* = Significant nontarget compound
 BCRL = Below Certified Reporting Limit
 ND = None Detected

Table ESA 2.3-1/ESA-1 5/19/89

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	Tol Tol	tal Wells Laf Samples	Alluvial 19 52			Denver 37 95	
Analytical	Groups/Analytes	Frequency of Detections	Concentration Range (µg/l)	CRL Range (µg/l)	Frequency of Detections	Concentration Range (μg/l)	CRL Range (ug/l)
Organochlo	cine Pesticides (OCI	ার্					
Aldrin		2/48	0.11	0.050 - 0.083	3/94	0.16 - 0.46	0.051 - 0.088
Dichlorodi	phenylethane (DDE)	0/48	BCRL	67.0 - 61.0 0.046 - 0.16	0/33 0/94	BCRL	0.15 - 0.23 0.046 - 0.14
Diektrin	ipnenyltrichloroethane (L	0/1) 0/48	BCRL	0.059 - 0.070	500	BCRL	0.050 - 0.13
Endrin		2/48	0.074 - 0.18	0.050 - 0.063	3/94	0.090 - 0.20	0.054 - 0.11 0.050 - 0.32
UUDOSI		81/0	BCRL	0.056 - 0.072	0/94	BCRL	0.056 - 0.14
Arsenic		3/26	4.7 - 5.3	2.5 - 3.9	10/46	2.6 – 26	2.5 - 4.0
Mercury		0/10	BCRL	0.36	0/22	BCRL	0.10 - 0.36
ICP Metals							
Cadmium		01/1	12	5.2	0/25	BCRL	. 5.2 – 8.4
Copper	-	01/1	16 - 6.1 30	0.0	3/25	13 – 43 9.7 – 33	6.0 - 24 7.9
Zinc		01/1	24 41 - 140	19 20	1/25	65 33 300	19 - 74
				07	(7/01	MC - 77	77 - 77

Table ESA 2.3-1 Summary of Target and Significant Nontarget Compound Detections in Eastern Study Area Groundwater. Page 5 of 5.

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Significant nontarget compound
 BCRL = Below Certified Reporting Limit
 ND = None Detected

Table ESA 2.3-1/ESA-1 5/19/89

Well ID	10/85 - 3/86 ISP	4/86 - 6/86 SP86	7/86 - 9/86 SU86	10/86 - 12/86 FA86	1/87 - 3/87 W187	4/87 - 6/87 SPR7	7/87 - 9/87 SUR7	1/89 - 3/8 WIE9
06002	٠	C6H6 2.4	•			+ CIC ₆ H ₅ 1.4 2n 41		
0090	•	•	•			+ Cr 7.3 72,110		
10020	•	•	•			- Cd 12 + Cd 12 ClCeH5 19 Cr 37 Cr 30		
						Pb 24 2n 97		
08002	•							
08003	ENDRN 0.070	•				+ 12DCLE 0.64 CTC 6H5 0.74 CT 11		
10061						+ Cr 18		
24107			As 5.3			+ Cr 26 Zn 93		
24166			DI.DRN 0.060	DLDRN 0.16	DLDRN 0.10	DLDRN 0.12	DLDRN 0.080	
24183			•	CPMSO 6.1 CPMSO ₂ 3.2	CPMSO ₂ 30 DBCP 0.16	CPMSO ₂ 3.7	As 4.7 CPMSO ₂ 3.6 DLDRN 0.087	
24188			•	CPMSO2 4.3 MEC6H5 8.6	CHCh, 22 CPMSO ₂ 4.0	CPMSO ₂ 3.4	As 5.1	
25011	•	•	•			+ (7 12		
25038	•					+ Zn 53		

Table ESA 2.3-2. Analytes Detected in Eastern Study Area Alluvial Well Samples (units in µg/l). Page 1 of 3.

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3009 • CIICly 60 + Cr 16 Aug Mis 3105 • - Cr 13 - Cr 16 - Cr 13 3201 A.DSKN 011 • + Cr 23 - Cr 12/3 31014 11014 • • Cr 23 - Cr 16/3 131014 • • • Cr 13/3 - Cr 16/3 131015 • • • Cr 16/3 • Cr 16/3 131015 • • • Cr 16/3 • Cr 16/3 131015 • • • Cr 16/3 • Cr 16/3 33044 • • • Cr 16/3 • Cr 16/3 33044 • • • Cr 16/3 • Cr 16/3 33044 • • Cr 10/3 • Cr 10/3 • Cr 10/3 33044 • • Cr 10/3 • Cr 10/3 • Cr 10/3 • Cr 10/3 33044 • • Cr 13/3 • Cr 13/3 • Cr 13/3 • Cr 10/3 • Cr 13/3 • Cr 13/3 • Cr 13/3 • Cr 10/3 • Cr 13/3 • Cr 13/3	Well 1D	10/85 - 3/86 1SP	4/86 - 6/86 SP86	7/86 - 9/86 SU86	10/86 — 12/86 Faxá	1/87 - 3/87	4/87 - 6/87 5 0 0 7	7/87 - 9/87	1/89 - 3/89
3008 •							1010	1800	W189
3105 • • • • • • • • • • • • • • • • • • •	3000	a	•	CHCI ₃ 60			+ Cr 16 Zn 140		
3301 ALDRN 0.11 • • • • • • • • • • • • • • • • • •	31005	٠					+ Cr 23 7n 43		
31014 31015 31016 31016 3304 3304	32001	ALDRN 0.11	•						
31015 31016 31016 31016 ALDRIN PLURIN PLURIN PLURIN PLURIN PLURIN PLURIN	31014								
31016 CHCl ₃ O CHCl ₃ O BLDRN ENDRN ENDRN	31015								
TORNO	31016								
ALDRIN DLDRNO	11/1/1								CIICE 0.31
									ALDRIN 0.11 DLDRN 0.27 ENDRN 0.18

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Table ESA 2.3-2. Analytes Detected in Eastern Study Area Alfuvial Well Samples (units in µg4). Page 3 of 3.

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- Spring \$2.5 x
 - Summer
 - Fall
- Winter
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- Sample analyzed, all analyzes helow. CRL. ICP metals in clucked in analyzed at suite .
 - Inual Sere ling Program. SP
- Berniene.
- Dibramen hlorigitigune. 1.2 - Dichlonichane. Chlonhenrene, 646 CCLE CCLE CCLE CCLE FCCLE FCCLE FCCLE FCCLE FCCLE FCCLE FCCLE FCCLE
 - Methylene uhlande. Tru hkaruthy kine.
- Chkwophenyl sulfoxude. Chkwaphenylmethyl sulfone. Tolucne.
 - Chloroform.

С'нкотит, C.Mmum. Copper LexaL Zunc, Akten, Dickten, Ekten, Arving. ALDRN DUDRN ENDRN ×588£\$

Table ESA 2.3-3. Analytes Detected in Eastern Study Area Denver Well Samples (units in µg/). Page 1 of 4.

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Vell ID	10/85 - 3/86 1SP	4:86 - 6/86 SP86	7/86 - 9/86 Sti86	10/86 - 12/86 Fa86	1/87 - 3/87 W187	4/87 - 6/87 SP87	7/87 - 9/87 SU87
0000	•					+ Cr 16 Cu 9.7 Zn 74	
05002	٠						
65003	٠						
06004	•	•	•			+ As 2.6	
8005	C ₆ H ₆ 10 DBCP 0.75	•	• ,			• +	
1004	•	٠	C ₆ H ₆ 8.9			• +	
20070	٠	C ₆ H ₆ 3.0	C ₆ H ₆ 4.2				
8004	+ As 5.9 Zn 36						
36005	• •					+ As 2.6 Zn 42	
19003						+ Cr 43 DLDRN 8.9 ENDRN 0.20 Za 60	
19015	+ As 6.8 C ₆ H ₆ 7.3 Zn 43	•	•			+ Cr 13 Cu 17	
91061	+ As 26 C ₆ H6 1.4 Zn 33	•	•				
19017	•					+ Cr 13 Zn 300	

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ESA Tab 2.3-3/ESA-1 5/19/x0 12:50 PM

	10/46 3/46	7817 7814	7146 0146	10/01	note nate		
Well ID	100% - COM	5P86	5U86	10/30 - 12/30 FA86	1/8/ - 5/8/ W187	4/87 - 6/87 SP87	7/87 - 9/87 SU87
81061	•						
61061	DLDRN 0.06 ENDRN 0.09	•					
24108			•	•	•	•	•
24109			As 8.6	•	•	•	•
24120			DLDRN 0.14	DLRN 0.19	DLDRN 0.12	+ DLDRN 0.12 Zn 35	DLDRN 0.16
24174			C6H6 20 CIC6H5 45 DBCP 0.15 TRCLE 2.4	•	As 3.3	CIC6H5 10	As 3.9 C6H6 2.5 CIC6H5 14
24175			C6H6 4.5 CIC6H5 25	CH2Cl24.3	٠	As 4.1 CIC ₆ H ₅ 17	CIC ₆ H ₅ 4.8 C ₆ H ₆ 4.0
25012	•						
25013	ALDRN 0.17	+Zn K2	+Zn 69			+ Zn 22	
25014	ALDRN 0.46 DLDRN 0.42	•	•			*	
250.9	+ Zn 23					+ Zn 23	
25040	+ C ₆ H ₆ 2.1 Zn 34						
29002	•						
29003	•						

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ESA Tab 2.3-3/ESA-1 5/19/89 12:50 PM

3004 • 3005 ENDRN 0.12 • 3001 • • • 3001 • • • 3001 • • • 3001 • • • 3001 • • • 3001 • • • 31002 • • • 31003 • • • 31004 • • • 31005 • • • 31006 • • • 31008 • • • 31008 • • • 31008 • • • 31008 • • • 31008 • • • 31008 • • • 31008 • • • 31008 • • • 31008 • • • 31009 • • •	Well ID	10/85 - 3/86 1SP	4/86 - 6/86 SPR6	7/86 - 9/86 SU86	10/86 - 12/86 FA86	1/87 - 3/87 W187	4/87 - 6/87 SP87	7/87 - 9/87 SU87
3003 ENDRN 0.12 • 3001 • • 3011 • • 3012 • • 3102 • • 31005 ALDRN 0.15 • 31007 • • 31007 • • 31008 • • 31009 • • 31001 • • 31002 • • 31003 • • 31004 • • 31008 • • 31009 • • 31001 • • 31003 • • 2.025 • • 2.033 • • 32003 • • 2.034 • •								
3005ENDRN 0.12 \cdot 3010 \cdot \cdot 3011 \cdot \cdot 3012 \cdot \cdot 3102 $+Cu 33$ 3102 $+Cu 33$ 3103 \cdot 3104 \cdot \cdot \cdot 3105 $-Cu 133$ 3107 \cdot	30004	•						
30010 • <td>30005</td> <td>ENDRN 0.12</td> <td>•</td> <td></td> <td></td> <td></td> <td></td> <td></td>	30005	ENDRN 0.12	•					
3001 • • CHCL ₃ 20 • • 31002 • • Cu 33 • • Cu 33 • • • • 31008 ALDRN 0.15 • • • • • • • 31007 • <td>30010</td> <td>•</td> <td>•</td> <td>•</td> <td></td> <td></td> <td></td> <td></td>	30010	•	•	•				
31002 + Cu 33 31006 ALDRN 0.15 + Cu 33 31007 • • • • • • • • • • • • • • • • • •	30011	•	•	CHCL ₃ 20			• +	
31006 ALDRN 0.15 • • • • • • • • • • • • • • • • • • •	31002		+ Cu 33					
31007 • • • • • • • • • • • • • • • • • •	31006	ALDRN 0.15	•	•				
31008 • • • • • • • • • • • • • • • • • •	31007	•	•	•				
32002 C6H6 I.8 + As 20 2.n 25 2.n 34 32003 + •	31008	•	•	٠				
- 175 65 Zn 34 32003 + •	32002	C6H6 1.8 + As 20 Zn 25					2	
32003 + •							+ Pb 65 Zn 34	
	32003	•						

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ESA Tab 2.3-3/ESA-1 5/19/89 12:50 PM Table ESA 2.3-3. Summary of Analytes Detected in Eastern Study Area. Denver Wells (units in µg/l). Page 4 of 4.

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Noles:

Sample analyzed, all analytes below CRL. ICP Metals included in analytical suite Methylene chloride. Chlorophenylmethyl sulfoxide. Chlorophenylmethyl sulfone. Initial Screening Program. Dibromochloropropane. Trichkoroethykene. 1.2 - Dichloroethane. Chlowbenzene. Arsenic. Chromium. Chloroform. Cadmium. Aktrin. Dickdrin. Endrin. Copper. Lead Zinc. Benzene. Tolucne. Summer Spring Winter Fall MEC₆H5 CHCl3 CIC6H5 12DCLE DBCP CPMSO2 ALDRN DLDRN ENDRN TRCLE CPMSO CH₂Cl₂ C6H6 ISP **ಬ**ಶರೆ SU FA SP M As 25 +

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ESA Tab 2.3-2/ESA-1 5/19/89 12:50 PM

CONTAMINATION CLASSIFICATION*	STRUCTURE NUMBER	SECTION	STRUCTURE FUNCTION
1	1736	31	Toxic Storage Yard
1	NN3103	· 31	Storage Building
1	NN3106	31	Shed
1	NN3109	31	Shed
2	395	6	Toxic Storage Yard Sewage Plant
2	785	6	Drum Storage Warehouse
2	792	31	Drum Storage Warehouse
2	853	30	Observation Pit/Mortar Range
2	864	6	General Structure
2	865	6	Warehouse
2	867A	6	Toxic Yard Metal & Wood Shop
2	867B	6	Flammable Materials Storehouse
2	871A	6	Magazine
2	871B	6	Magazine
2	871C	6	Magazine
2	871D	6	Magazine
2	8 72A	6	Magazine
2	872B	6	Magazine
2	872C	6	Magazine
2	872D	6	Magazine
2	873A	6	Magazine
2	873B	6	Magazine
2	873C	6	Magazine
2	874A	6	Magazine
2	874B	6	Magazine
2	874C	6	Magazine
2	874D	6	Magazine
2	883	6	Igloo Storage
2	1735	31	Loading Dock
2	NN0601	6	Loading Dock
2	NN0602	6	Long Metal Shed
2	NN0603	6	Metal Shed
2	NN2001	20	Antenna Installation

Table ESA 2.4-1 Contaminant Classification of Structures in the Eastern Study Area. Page 1 of 2

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CONTAMINATION CLASSIFICATION*	STRUCTURE NUMBER	SECTION	STRUCTURE FUNCTION
2	NN3001	30	Metal Shed
2	NN3002	· 30	Meial Shed
2	NN3102	31	3 Sets Shed Siding
2	NN3104	31	Shack
2	NN3105	31	Shed
2	NN3107	31	Antenna Station
2	NN3108	31	Shed
3	786	6	Drum Storage Warehouse
3	787	6	Drum Storage Warehouse
3	788	6	Drum Storage Warehouse
3	791	31	Drum Siorage Warehouse
3	793	31	Drum Storage Warehouse
3	794	31	Drum Storage Warehouse
3	795	31	Drum Storage Warehouse
3	796	31	Drum Storage Warehouse
3	797	31	Drum Storage Warehouse
3	798	31	Drum Storage Warehouse
3	851	19	Pistol Range House
3	866	6	Toxic Yard Office and Change House
3	881	6	Igloo Storage
3	882	6	Igloo Storage
3	884	6	Igloo Storage
3	885	6	Igloo Swrage
3	886	6	Igloo Storage
3	1730	31	Sentry Station/Gatchouse
3	1734	31	Change House
3	NN0501	5	Abandoned Schoolhouse
3	NN2002	20	Tank Pad
3	NN3101	31	Metal Shed

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Table ESA 2.4-1 Contaminant Classification of Structures in the Eastern Study Area. Page 2 of 2

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1 = Suspected to be contaminated
 2 = Suspected to be contaminated but cleanable
 3 = Suspected to be uncontaminated

ESA 2.4-1/ESA 1 Rev. 5/10/89

Table ESA 2.5-1. Airborne Contaminant Distribution. Page 1 of 1.

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Parameter	Station AO4	
Total Suspended Particulates		
Number of Sampling Events	3	
Annual Geometric Average (µg/m3)	43	
Range of Individual 24-Hour Samples (µg/m3)	39-47	

Source: ESE, 1988 (Air Media Rpt)

ESA 2.5-1/ESA-1 Rev. 5/10/89

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Table ESA 2.6-1. Contaminants of Concern to Biota in Analyte Groups. Page 1 of 1.

GI	ROUP	CONTAMINANT
1.	<u>Volatile Halogenated</u> <u>Organics</u> (VHOs)	Chloroform Chlorobenzene Trichloroethylene Allyl chloride*
2.	Volatile Aromatic Organics (VAOs)	Ethylbenzene Toluene Xylene
3.	Volatile Hydrocarbons (VHCs)	Dicyclopentadiene (DCPD)
4.	Organochlorine Pesticides (OCPs)	Aldrin Chlordane Dichlorodiphenylethane (DDE) Dichlorodiphenyltrichloroethane (DDT) Dieldrin Endrin Heptachlor* Heptachlor epoxide (HE)* Isodrin Oxychlordane* Polychlorinated Biphenyls (PCB)*,**
5.	Organophosphorous Compounds. Pesticide Related (OPHPs)	Atrazine Azodrin* Malathion Methyl parathion* Parathion
6.	Organophosphorous Compounds. GB-Agent Related (OPHGBs)	Diisopropylmethyl phosphonate (DIMP) Dimethylmethyl phosphonate (DMMP) Methylphosphonic acid (MPA)
7.	Organosulfur Compounds, Herbicide Related (OSCHs)	Chlorophenylmethyl sulfide Chlorophenylmethyl sulfone Chlorophenylmethyl sulfoxide
8.	Organosulfur Compounds. Mustard-Agent Related (OSCMs)	Dithiane Mustard* 1,4-Oxathiane
9.	Organonitrogen Compounds (ONCs)	n-Nitosodimethylamine
10.	DBCP	
11.	Arsenic	
12.	Mercury	
13.	ICP Metals (ICPs)	Cadmium Copper

Compounds not included in the contaminant groups of other media.
 Non-pesticide organochlorine

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ESA 2.6-1/ESA-1/Rev. 5/10/89

ISATHAMA			Certified Rep	orting Limits
Method Code	Matrix Type	Analyte	Lower CRL	Upper CRL
B-6	Animals and Plants	Arsenic	0.25	5.0
C-6	Animals and Plants	Mercury	0.050	0.40
D-6	Plants	Aldrin Dieldrin Endrin	0.02 0.04 0.04	0.3 0.3 0.6
E-6A	Animals	Aldrin Dieldrin Endrin	0.02 0.03 0.04	0.3 0.3 0.6
F-6A	Anime's	DDE DDT	0.09 0.3	2 4

Table ESA 2.6-2. Certified Reporting Limits for Biota Analysis Methods. Page 1 of 1.

Source: ESE, 1988m

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ESA 2.6-2/ESA-1 Rev. 5/10/89

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Table ESA 2.6-3. Contaminant Levels in Terrestrial Ecosystems --- Samples of Species Ranging Across RMA. Page 1 of 3.

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opecies	Tissue	Location	Arsenic (n/nt)	Mercury (n/nt)	Aldrin (n/nt)	Dieldrin	ht Basis (Range/mr Endrin	can*) p,p-DDE	D.p-DDT
Invertebrat	ฮ					(111)	(n/nt)	(n/nt)	(n/nt)
Earthworms	Whole	RMA, South Plants	BDL (1)	<0.050->2.4 (1/2)	BDL (1)	2 (1)	BDL (1)	BDL (1)	BDL (1)
	Whole	RMA Control 0 (Section 5 - ESA)	1.618-1.53(8/8) 1.03	<0.050-0.25 (2/8)	BDL (7)	<0.06-5 (1/7)	<0.08-0.9 (1 <i>1</i>)	BDL (8)	BDL (8)
	Whole	Offpost Control	BDL (2)	BDL (2)	BDL (1)	BDL (1)			
Grasshoppers	Whole	RMA Section 26	BDL (4)	BDL (4)	0.05.6 (4/4)	0.5-7 (4/4)	c0.06-2 (3/4)	BDL (1)	BDL (1) BDL (1)
		Section 36 0	.91-6.6 (4/4) 3.2	<0.050-0.11 (2/4) 0.058	ء BDL (4)	3 0.3-0.4 (4/4)	0.5 BDL (4)	BDL (4)	BDL (4)
	(Se	RMA Control ction 7 and 8 - ESA	BDL (3) \)	BDL (3)	BDL (3)	0.4 BDL (3)	BDL (3)	BDL (3)	BDL (3)
		Offpost Control	RDL (2)	BDL (2)	BDL (2)	BDL (2)	BDL (2)	B DL (2)	BDL (2)

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= Mean is calculated when 50 percent or more of samples have detectable contaminant levels. If less than 50 percent of samples have detectable contaminant levels, only the range of values are presented. When calculating the mean, values of 1/2 the detection limit are substituted for samples that are below the detection limit. •• = MKE sample.

BDL = Below Detection Limit. NRQ = Not requested. n = Number of samples analyzed that contain detectable contaminant levels. n = Number of samples. TSY = Toxic Storage Yard bource: ESE, 1088m

Source:

ESA Tab 2.6-3 Species Rev. 5/10/89

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Table ESA 2.6-3. Contaminant Levels in Terrestrial Ecosystems --- Samples of Species Ranging Across RMA. Page 2 of 3.

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				Contamir	nant Level in	n me/ko Wet Weicht	Rasis (Rance)	/mean*)	
Species	Tíssue	Location	Arsenic (n/nt)	Mercury (n/nt)	Aldrin (n/nt)	Dieldrin (n/nt)	Endrin (n/nt)	p,p-DDE (n/nt)	p,p-DDT (n/nt)
Vertebrates									
Ring-necked Pheasant	Juvenile	RMA <	:0.25 - 1.8 (3/11)	BDL (11)	BDL (12)	<0.03 - 1 (5/12)	BDI. (12)	BDL (11)	BDL (11)
	Adult	RMA	BDL (4)	BDL (4)	BDL (4)	<0.03 - 3 (3/4) 0.8	BDL (4)	BDL (3)	BDL (3)
	Egg	RMA	NRQ	0.050 - 0.41 (8/34)	BDL (33)	<0.03 - 4 (1773) >0.5	BDL (33)	<0.09 - 1 (1/29)	BDL (29)
	Egg	Offpost	NRQ	0.050 - 0.057 (1/11)	(11) TOB	BDL (11)	BDL (11)	<0.09 - 1 (2/11)	BDL (11)
Prairie Dog	Carcass Sec	RMA < tion 36 Summ	:0.25 - 0.74 (2/9) ICT	BDL (9)	BDL (9)	0.2 - 10 (9/9) 2	BDL (9)	NRQ	NRQ
	Carcass Sc	RMA ction 36 Winte	BDL (5) ST	BDL (5)	BDL (5)	0.1 - 6 (5/5) 1	BDL (5)	NRQ	NRQ
	Carcass	ESA Toxic Storage Yard	<0.25 - 4.2 (1/5)	BDL (5)	BDL (5)	0.06 - 0.2 (5/5) 0.1	BDL (5)	NRQ	NRQ
	Carcass	RMA Control Summer	BDL (9)	BDL (9)	BDL (9)	<0.03 - 0.3 (2/9)	(6) HUL (9)	NRQ	NRQ

• = Mean is calculated when 50 percent or more of samples have detectable contaminant levels. If less than 50 percent of samples have detectable contaminent levels, only the range of values are presented. When calculating the mean, values of 112 the detection limit are substituted for samples that are below the detection limit.

** = MKE sample. BDL = Below Detection Limit. NRQ = Not requested.

n = Number of samples analyzed that contain detectable contaminant levels. nt = Number of samples. TSY = Toxic Storage Yard ource: ESE, 1988m

Source:

ESA Tab 2.6-3 Species Rev. 5/10/89

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				Contani	nant Level in	me/ke Wet Weieht	Basis (Range/1	mean•)	
Species	Tissue	Location	Arsenic (n/nt)	Mercury (n/nt)	Aldrin (n/nt)	Dieldrin (n/nt)	Endrin (n/nt)	p,p-DDE (n/nt)	p.p-DDT (n/nt)
	Carcass	RMA Control Winter	BDL (5)	BDL (5)	BDL (5)	<0.03 - 0.1 (1/5)	BDL (5)	NRQ	NRQ
	Carcass	Offpost Control Summer	BDL (9)	BDL (9)	BDL (8)	BDL (8)	BDL (8)	NRQ	NRQ
	Kidneys	RMA, Section 36 Winter	BDL (5)	<0.10 0.36 (3/5) 0.18	BDL (5)	<0.2 - 2 (2/5)	BDL (5)	NRQ	NRQ
Cottontail	Muscle	RMA, Section 36	BDL (7)	RDL (7)	RDL (7)	<0.03 · 0.09 (371)	BDL (7)	NRQ	NRQ
	Muscle	RMA, Control	BDL (7)	BDL (7)	BDL (7)	BDL (7)	BDL (7)	NRQ	NRQ
	Muscle	RMA, Offpost Control	BDL (7)	RDL (7)	(1) 108	BDL (7)	BDL (7)	NRQ	NRQ
Mule Deer	Liver	RMA	BDL (14)	BDL (14)	BDL (14)	<0.01 - 0.2 (1/14)	BDL (14)	NRQ	NRQ
	Liver	Offpost	RDL (2)	RDL (2)	RDL (2)	BDL (2)	BDL (2)	NRQ	NRQ
	Muscle	RMA	BDL (14)	BDL (14)	RDL (14)	BDL (14)	BDL (14)	NRQ	NRQ
	Muscle	Offpost	BDL (2)	RDL. (2)	RDL (2)	RDL (2)	BDL (2)	NRQ	NRQ

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Table ESA 2.6-3. Contaminant Levels in Terrestrial Ecosystems --- Samples of Species Ranging Across RMA. Page 3 of 3.

Mean is calculated when 50 percent in more of samples have detectable contaminant levels 15 less than 50 percent of samples have detectable contaminant levels, only the range of values are prevented. When calculating the mean, values of 1/2 the detection limit are substituted for samples that are below the detection limit. N .

MKE sample
 Below Denection Limut
 RDL = Relow Detection Limut
 NRQ = Not requested.
 n = Number of samples analysed that contain detectable contaminant levels
 n1 = Number of samples
 TSY = Tosic Storage Yard
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ESA Tab 2 6-3 Species Rev. 5/10/89

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Species	Tissue	Location	Arsenic (n/nt)	Mercury (n/nt)	Aldrin (n/nt)	1 mg/kg Wet Weight Dieldrin (n/nt)	Basis (Range/m Endrin	p.p-DDE	P.P-DDT
							(1u/u)	(n/nt)	(u/nt)
Mouming Dove	Carcass	RMA	BDL (2)	BDL (2)	<0.6 - 2 1	6 - 60 (2/2) 30	<0.8 - 3 (1/2) 2	BDL (2)	BDL (2)
	Liver	RMA	BDL (1)	BDL(I)	BDL (1)	7 (1)	4 (1)	BIN 232	
Bald Eagle	Egg	Barr Lake	HDL	0.10	BDL (1)	0.8 (1)	BDLOD	10,000	BUC (I)
àolden Eagle	Liva	RMA	NRQ	<0.050 - 0.22 (1/2) 0.12	BDL (2)	<0.031 0.22 (1/2) 0.12	BDL (2)	BDL (2)	BDL (1)
	Brain	RMA	BDL (2)	<0.098 - 0.26 (2)	RDL (2)	BDL (2)	RDL (2)	RDI.(7)	
Terruginous Hawk	Liva	RMA	BDL (5)	<0.050 - 0.29 (1/5)	BDL (5)	0.26 - 4.8 (5/5) 2.7	BDL (5)	BDL (5)	BDL (5)
	Brain	RMA	BDL (5)	<0.050 - 0.15 (1/5)	8DL (5)	<0.24 - 1.0 (4/5)	BDL (3)	BD1 (5)	
Red-tailed Hawk	Liva	RMA	BDL (1)	(UI) SE 0 · 050 0>	BDL (3)	0.5 - 7 (3/3) 4	BDT (3)	<0.3 - 0.8 (2/3)	BDL (3)
	Brain	RMA	BDL (3)	<050.050 - 050.050 (1/3)	RDL (3)	<0.8 - 9 (2/1) 6	BDL (3)	BDL (3)	BDL (3)
irrat hornod Owl	Liva	RMA	BDL (4)	<0.0% - 0.0%6 (2/4) 0.047	BDL (4)	0.1 - 30 (4/4) 1.0	BDL (4)	<0.09 - 20 (3/4) 6	BDL (4)

Mean is calculated when 50 percent or more of samples have detectable contaminant levels. If less than 50 percent of samples have detectable contaminant levels, only the range of wilnes are presented. When calculating the mean, values of 1/2 the detection limit are substituted for samples that are below the detection limit. •

MKE sample.
 MKE sample.
 BOL = Beiow Denetion Limit.
 NRQ = Not requested
 NRQ = Number of samples analyzed that contain detectable containing levels.
 NSY = Twee Storage Yard
 Source - ENE. 1988m

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Species	Tissue	Location	Arsenic (n/nt)	Mercury (n/nt)	Aldrin (n/nt)	Dieldrin (n/nt)	Endrin (n/nt)	p.p-DDE (n/nt)	p.p-DDT (n/nt)
	Brain	RMA	BDL (4)	BIX (4)	BDL (4)	<0.2 · 20 (3/4) 9	BDL (4)	<0.5 - 10 (3/4) 3	BDL (4)
Northern Harrier	Egg	RMA	BDL (2)	BOL (2)	BDL (2)	0.3 - 0.7 (2) 0.5	BDL (2)	BDL (2)	BDL (2)
Coyote	Liver	RMA	BDL (1)	BDL (1)	BDL (1)	8 (1)	BDL (1)	BDC (1)	(I) JOB
Bake	Live	RMA	BDL, `	8DF (1)	BDL (1)	2 (1)	BDL (1)	NRQ	NRQ
	Kidneys	RMA	NRQ	08N	BDL (1)	0.8 (1)	BDL (I)	NRQ	NRQ

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ESA 2 6-4/ESA-1 Rev. 5/10/89

⁼ Mean is calculated when 50 percent or more of samples have detectable contaminant levels. If less than 50 percent of samples have detectable contaminant levels, only the range of values are presented. When calculating the mean, values of 112 the detection limit are substituted for samples that are below the detection limit. •

MKE sample.
 Below Detection Limit.
 Not requested
 Not requested
 Number of samples unalysed that contains detectable containant levels.
 Number of samples
 Source E.S., 1905m

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EASTERN STUDY AREA LOCATION AT ROCKY MOUNTAIN ARSEMAL Methylene Chloride[®] in Soils in the 2-5 ft. Depth Interval 2000 Rocky Mountain Arsenol Cleanup Program Manager's Office for 1" = 4 MILES ÷ 2 * -2 F E T 8 FIGURE ESA 2.1-5 2 * \$ ~ 2 Rocky Mountain Arsenat] 2 2 * • Prepared for: \$ 2 • • $\overline{\mathbf{n}}$ -----1 ï 5 0 9 0 22 0 14 Ò ā, -**0**0 õ 0 0 £. Ŷ Ŧ 0 0

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