EPA Superfund Record of Decision:

LORING AIR FORCE BASE EPA ID: ME9570024522 OU 01 LIMESTONE, ME 09/20/1995



LORING AFB MAINE

ADMINISTRATIVE RECORD COVER SHEET

AR File Number <u>2410</u>

2410 1 File: | 7A-49 P.F. 24/0

FINAL

OPERABLE UNIT 1 (OU 1) RECORD OF DECISION

August 1995



Installation Restoration Program
Loring Air Force Base, Maine

FINAL

Loring Air Force Base

OPERABLE UNIT 1 (OU 1) RECORD OF DECISION

August 1995

Prepared for:

Air Force Base Conversion Agency Limestone, Maine (207) 328-7109

Prepared by:

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OU 1 RECORD OF DECISION LORING AIR FORCE BASE

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DECLARATION FOR THE RECORD OF DECISION

or accommendation of the

SITE NAME AND LOCATION

Loring Air Force Base (LAFB) Operable Unit 1 (OU 1), the Low Level Radioactive Waste Disposal Sites (LLRWDS), Limestone, Maine.

STATEMENT OF BASIS AND PURPOSE

This decision document presents the selected No-Action decision for the LLRWDS, OU 1, at LAFB in Limestone, Maine. OU 1 consists of Areas A-G as identified in Figure 1-2. This decision document was developed in accordance with the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) of 1980, as amended by the Superfund Amendments and Reauthorization Act of 1986, and to the extent practicable, the National Oil and Hazardous Substances Pollution Contingency Plan (NCP), (USEPA, 1990). It is based on the Administrative Record for the site, which was developed in accordance with Section 113(k) of CERCLA and is available for public review at the Air Force Base Closure Agency Office, 5100 Texas Road, Limestone, Maine. The Administrative Record for the LLRWDS, OU 1, includes the memos, letters, reports, and associated information developed during the CERCLA response at OU 1 that provide the basis for selecting No Action.

The State of Maine Department of Environmental Protection concurs with the No-Action under CERCLA remedy for OU 1.

DESCRIPTION OF THE SELECTED REMEDY

The U.S. Air Force and U.S. Environmental Protection Agency (USEPA), with concurrence of the Maine Department of Environmental Protection, have determined that no action under CERCLA is necessary to address the contamination of OU 1 soils, surface water, sediments, and groundwater. Previous response actions relating to radionuclides at OU 1 (Areas A through F) have eliminated the need to conduct a remedial action. OU 1 inorganic groundwater contamination will be addressed in a separate Record of Decision, and the petroleum contamination at Area G will be addressed separately under the Maine Underground Storage Tank Regulations.

DECLARATION

Because this No Action Record of Decision does not result in hazardous substances, pollutants, or contaminants being left at the site above levels that allow for unrestricted exposure, pursuant to CERCLA § 122(c), no five-year review will be undertaken.

DECLARATION

The U.S. Air Force and USEPA, with concurrence of the Maine Department of Environmental Protection, have determined that no remedial action under CERCLA is necessary at OU 1.

By:

Department of the Air Force

Alan K. Olsen

Director

Air Force Base Conversion Agency

By:

nited States Environmental

Protection Agency Linda M. Murphy

Director

Waste Management Division

Region I

Date: Sept , 50 1995

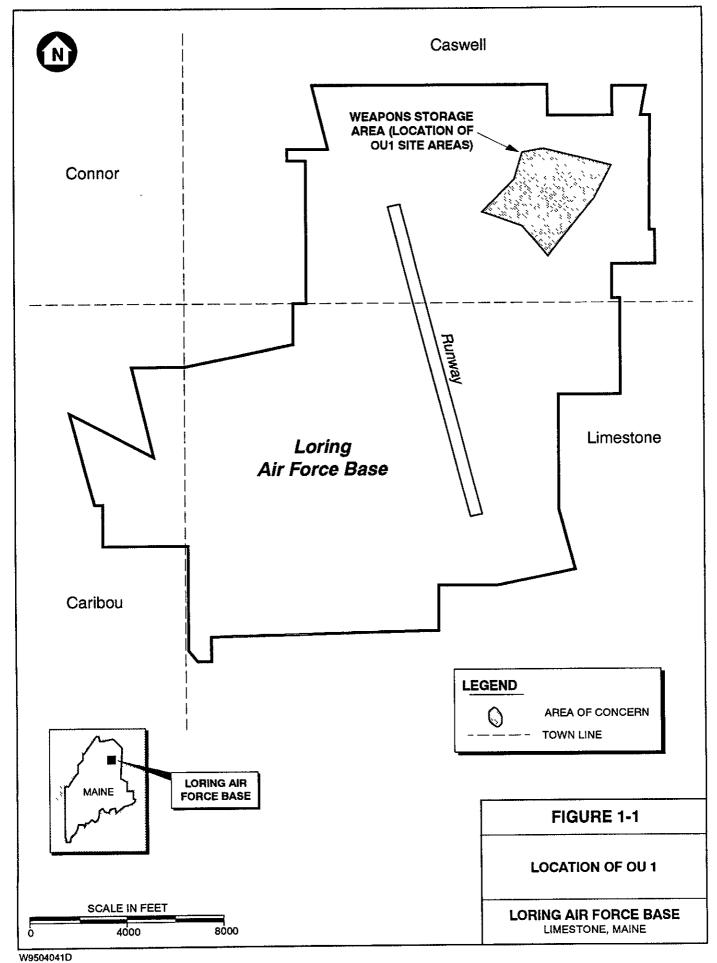
1.0 SITE NAME, LOCATION, AND DESCRIPTION

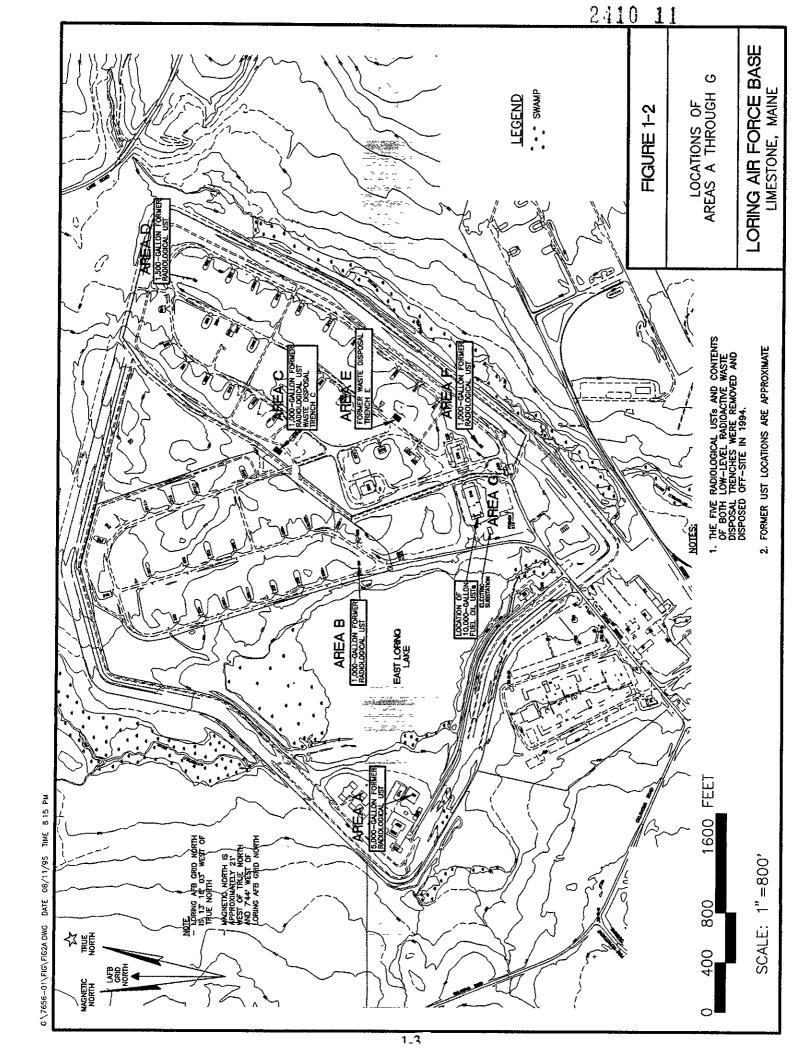
Loring Air Force Base (LAFB), in northeastern Maine, is bordered on the south and east by the Town of Limestone, on the north by the towns of Caswell and Connor, and on the east by the City of Caribou (Figure 1-1). The base is approximately three miles west of the United States/Canadian border and covers approximately 9,000 acres. The base was closed September 1994.

LAFB is a National Priorities List (NPL) site. There are currently a number of areas of concern within LAFB that are under investigation. For purposes of investigation and remedial response, the areas of concern at LAFB have been organized into several operable units (OUs). This Record of Decision (ROD) addresses the former source areas, surface water, sediment, and groundwater at OU 1, the Low Level Radioactive Waste Disposal Sites (LLRWDS). The LLRWDS Areas A through G, identified in Figure 1-2 are discussed further in Subsection 5.1.

Because of its primary mission, LAFB personnel were engaged in various operations, a number of which required the use, handling, storage, and disposal of hazardous materials and substances. In the past, these materials entered the environment through accidental spills, leaks in piping, landfilling operations, burning of liquid wastes during fire-training exercises, and the cumulative effects of operations conducted at the base's flightline and industrial areas. As part of the Department of Defense's (DOD) Installation Restoration Program (IRP), the Air Force has initiated activities to identify, evaluate, and remediate former disposal or spill sites containing hazardous substances.

Since initiation of the IRP, the Base has been placed on the U.S. Environmental Protection Agency's (USEPA's) NPL of sites and will be remediated according to the Federal Facility Agreement (FFA) entered into by U.S. Air Force (USAF), the USEPA, and the Maine Department of Environmental Protection (MEDEP).





2.0 SITE AND INVESTIGATION HISTORY

This section summarizes the uses, response history, and investigation activities at OU 1.

2.1 LAND USE AND RESPONSE HISTORY

The seven LLRWDS in OU 1 are associated with buildings and operations in the Weapon Storage Area (WSA) (Figure 2-1). The WSA was used for the storage and routine maintenance of strategic and conventional weapons from 1952 through 1993. During the 1950s, weapons inspection and maintenance required disassembly and direct handling of radioactive materials. By the mid-1950s, weapon designs had changed, radioactive material was no longer exposed in the new designs, and the earlier type of weapons were progressively phased out of stockpile by 1962. Strategic weapons were removed from the WSA in May 1989. Conventional weapons were progressively removed in 1993 in anticipation of base closure, with the last conventional weapons removed in December 1993.

Five underground storage tanks (USTs) were installed at the WSA LLRWDS to receive and contain potentially radioactive liquids in the event of a release in one of the facilities. USAF records indicated there was never a release of radioactive materials to any of the five USTs. The USTs were excavated and disposed off-site during a removal action in 1994. The USTs were observed to be intact at the time of their removal (Ogden, 1995).

Low-level dry radioactive wastes (e.g., swipes, butcher paper, tape, gloves, protective clothing, respirator cartridges) from maintenance operations were typically placed in cardboard boxes. From 1954 through 1962, the boxes were reportedly disposed of on-site in two waste disposal trenches. During the 1994 removal action, the two waste trenches were delineated, exhumed, and the contents were disposed off-site.

2.2 INVESTIGATION AND RESPONSE HISTORY

The USAF has followed USEPA guidelines for most of the IRP investigations conducted at LAFB since 1983, and for all investigations completed since 1988, when

the IRP investigation process was revised to more closely follow the National Contingency Plan (NCP) used by the USEPA (USEPA, 1990).

The investigation history of OU 1 is summarized as follows:

- In 1983, a Preliminary Assessment (PA) was performed by detailing historical hazardous material usage and waste disposal practices (CH₂M Hill, 1984).
- A Site Inspection (SI) was conducted between 1985 and 1988 to confirm the presence of contaminants at OU 1 (Roy F. Weston, Inc., 1988).
- Between 1988 and 1994, Remedial Investigation (RI) activities were conducted and a Public Health and Ecological Baseline Risk Assessment (RA) was completed (ABB Environmental Services, Inc. [ABB-ES], 1995a).
- LAFB was added to the NPL in February 1990.
- The USAF entered into an FFA in 1991 with the USEPA and MEDEP regarding the cleanup of environmental contamination at LAFB (FFA, 1991).
- In 1994, a removal action was conducted that included excavation of the five radiological USTs and two waste disposal trenches. The USTs and contents of the trenches were disposed off-site (Ogden, 1995).
- The FFA was modified in December 1993 to address base closure-related issues, such as real property transfer and a revised schedule. The FFA was further modified in January 1995 to allow Remedial Project Managers to make minor modification to the FFA, such as schedule adjustments or removal of petroleum-contaminated sites from the agreement.
- Contamination detected at Area G is attributed to fuel oil that leaked from a former UST and pipeline, and as such, future remedial actions should be conducted in accordance with State of Maine UST regulations.

3.0 COMMUNITY PARTICIPATION

Throughout LAFB's history, the community has been involved in base activities. The USAF, USEPA, and MEDEP have kept the community and other interested parties apprised of LAFB IRP activities through informational meetings, fact sheets, press releases, public meetings, site tours, and open houses.

In addition to these activities, during the course of IRP activities at LAFB, there have been regular meetings of the Restoration Advisory Board (RAB) (formerly the Technical Review Committee). The RAB, chaired by the USAF and a representative of the community, is composed of representatives of USEPA, MEDEP, the community, and local officials. The purpose of the RAB meetings has been to ensure clear communication with the public, timely transfer of information, and opportunity for public comment.

The framework for the USAF's approach to community involvement is the LAFB Community Relations Plan (CRP), which was released in August 1991 and subsequently revised in May 1995. The CRP outlines the USAF's program for addressing community concerns and keeping citizens informed and involved during remedial activities.

Documentation of the reports, memoranda, and correspondence that are the basis for IRP remedial response decisions are kept in an Administrative Record. The Administrative Record is open and available for public review at the Air Force Base Conversion Agency Office, 5100 Texas Road, Limestone, Maine.

The following is a summary of the activities the USAF has undertaken to keep the public informed and involved regarding the remedial response at OU 1.

- On June 2, 1994, a RAB meeting was held to discuss the results of the OU 1 investigations and the approach for conducting the UST and radioactive waste disposal trench removal action.
- An IRP Fact Sheet, explaining activities planned for OU 1, was issued in July 1994.

- The USAF published a notice and brief discussion of the proposed removal action in the Aroostook Republican on July 6, 1994 and the Bangor Daily News on July 7, 1994.
- From July 11 through August 10, 1994, the USAF held a 30-day public comment period to accept public input on the Action Memorandum outlining the proposed removal action, and on any other OU 1 documents in the Administrative Record. On July 28, 1994, USAF personnel and regulatory representatives held a public meeting to discuss the Action Memorandum and to accept oral comments.
- During the removal action, the USAF invited the local press to cover the trench removal activities. Information regarding both the trench and UST tank removals was made available to representatives of local media.
- The USAF published a notice and brief analysis of the Proposed Plan in the Bangor Daily News, Aroostook Republican, Fort Fairfield Review, and Presque Isle Maine Star-Herald on July 12, 1995, recommending No Action under CERCLA as the preferred alternative for OU 1.
- From July 17 through August 16, 1995, the USAF held a 30-day public comment period to accept public input on the information presented in the RI/Baseline Risk Assessment and Proposed Plan, and on any other OU 1 documents in the Administrative Record. On August 2, 1995, USAF personnel and regulatory representatives held a public meeting and hearing to discuss the Proposed Plan and to accept oral comments. A transcript of this hearing is included in Appendix A. Comments received during the comment periods and the USAF's response to these comments are included in the Responsiveness Summary in Appendix B.

4.0 SCOPE AND ROLE OF RESPONSE ACTION

The USAF and USEPA have determined that no further Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) action is required at OU 1 because (1) previous response actions conducted at the operable unit have eliminated the need to conduct further remedial action and (2) the petroleum contamination at Area G will be effectively addressed under State of Maine regulations.

USEPA has the authority to revisit the No Action under CERCLA decision even if LAFB is removed from the NPL. This could occur if future conditions indicate that an unacceptable risk to human health or the environment would result from exposure to contaminants at OU 1.

5.0 SUMMARY OF SITE CHARACTERISTICS

The investigation process began at LAFB in 1983 as part of the DOD IRP. The process was revised during 1988 to follow the NCP. Investigations performed to date include a 1983 PA performed to investigate past activities at LAFB (CH₂M Hill, 1984). An SI was initiated in June 1985 to confirm the presence of contaminants at OU 1 (Roy F. Weston, Inc., 1988). In addition, RI activities were conducted from 1988 through 1994 (ABB-ES, 1995b).

There are seven OU 1 sites, Areas A through G, that were grouped together due to their proximity in the former WSA (Figure 2-1). With one exception, the sites were used for low-level radioactive waste disposal. The one exception is Area G, which was inaccurately identified as a low-level radioactive waste disposal site in the Base Master Plan during the 1970s and 1980s. Research and the results of the RI have shown that Area G was not used for low-level radioactive waste disposal. The Base Master Plan was corrected in the 1990s. A more complete description of the site can be found in Section 4 of the Operable Unit (OU 1) Remedial Investigation Report, Volume I (ABB-ES, 1995a).

The site areas comprising OU 1 that potentially received low-level radioactive waste are:

Area A: 5,000-gallon liquid waste disposal UST attached to Building 365 floor drains. Building 365 was the strategic weapon component inspection laboratory that maintained radioactive components. Potential contamination included radioactive materials (uranium oxide) and solvents generated during weapon maintenance activities.

Area B: 1,000-gallon liquid waste disposal UST attached to a floor drain in Building 329. Building 329 was used to store tritium containers. Tritium generated during routine venting of tritium gas during weapon maintenance activities at Building 329 was the primary focus of the investigation at Area B.

Area C: 1,000-gallon liquid waste disposal UST and a dry radioactive waste disposal trench, Trench C. The UST was attached to former Building 309 used to store tritium containers. The waste disposal trench was used in the 1950s and possible early 1960s to dispose of small quantities of low-level radioactive waste, primarily uranium oxides. Potential contamination to be

investigated at Area C included tritium generated during routine venting of tritium gas during weapon maintenance activities, and radioactive materials and solvents generated at Area A disposed in Trench C.

Area D: 1,500-gallon liquid waste disposal UST. This UST was attached to floor drains in Buildings 255 and 284, both used for storage of tritium containers. Tritium was identified as a potential contamination source at Area D.

Area E: Dry radioactive waste disposal Trench E, similar to the trench at Area C in its history and use. The focus of the Area E investigation was radioactive materials and solvents from wastes generated at Area A and disposed of in Trench E.

Area F: 1,000-gallon liquid waste disposal UST. This UST was attached to a floor drain in a weapon assembly and maintenance structure, Building 232. Potential contaminants at Area F included radioactive materials from weapon assembly and maintenance activities at Building 232.

The five radiological USTs were removed from Areas A, B, C, D, and F in 1994. In addition, the contents of both waste trenches (Areas C and E) were removed and disposed off-site in 1994.

Area G is not a radioactive waste disposal site. Building 216, located at Area G, was a weapon assembly building. There were two 10,000-gallon underground fuel storage tanks located at the west end of Building 216. In 1991, both tanks were replaced. During replacement of the tanks, contaminated subsurface soil, attributed to leakage from the tanks or piping, was observed. A former underground fuel pipeline, that supplied the 10,000-gallon tanks, traverses Area G. The pipeline is now abandoned. Investigations at Area G have detected solvents and fuel oil in soil and groundwater.

Other investigations and remedial actions have occurred at the WSA in addition to the PA and RI programs. The five radiological USTs were removed in 1994 as part of a removal action (Ogden, 1995). All five of the tanks were reported to be intact (i.e., not leaking). Based on analysis of UST content samples and confirmatory soil samples collected following UST removal, the Radioisotope Committee and MEDEP acknowledged, through verbal agreement, clean closure of the radiological USTs. Wipe samples from the building floor drains and the cut end of the piping at each UST were also analyzed and reported to be free of radioactive contamination.

During the 1994 removal actions, the dry radioactive wastes disposed in the trenches at Areas C and E were also removed. The contents of both trenches were delineated, analyzed, exhumed, and disposed off-site. Analysis of confirmatory soil samples collected after trench excavation indicated that the radioactive material was successfully removed from both trenches.

In addition to characterizing the sites, radiological building decommissioning surveys were conducted at 56 weapon storage and maintenance structures located within the WSA. No radioactivity above background levels was reported in the surveyed structures.

The University of Maine, on behalf of the MEDEP, conducted supporting radiological investigations to evaluate the possible presence of unidentified or undocumented radioactive waste disposal sites within the WSA. University of Maine personnel performed radiological surveys and laboratory analysis of groundwater, soil, surface water, and sediment samples from OU 1 and vicinity. Their OU 1 data were compared to off-site background samples and data from across the State of Maine. The comparisons indicated that levels of radioactivity across the entire WSA were at background levels, and the study did not identify any undocumented radioactive waste disposal areas. The University of Maine data were not utilized in establishing specific background values for the agreed-upon radionuclides of concern investigated in the RI. However, the University of Maine was involved in the review process for the establishment of these background concentrations developed during the RI.

The following subsections present contamination assessments for various environmental media at OU 1. A more detailed discussion of the contamination assessment is presented in Section 4 of the RI Report (ABB-ES, 1995a).

5.1 ANALYTICAL CHEMISTRY ISSUES

In order to better evaluate the nature and distribution of detected analytes, there are three issues which require preliminary discussion. These include:

- effects of turbidity on groundwater sample inorganic results
- the occurrence of Radium (Ra)-226
- radioactive isotope analytical results

Each of these topics is discussed in the following paragraphs.

Turbidity. Inorganics analytes were detected at varying levels above natural LAFB background concentrations in bedrock monitoring wells at OU 1. Inorganics were also detected in the two overburden wells. Background concentrations of inorganics in overburden and bedrock groundwater are currently being reassessed as part of the OU 12 basewide groundwater RI. Concentrations of inorganics in groundwater at OU 1 will be compared to the OU 12 background concentrations upon approval and acceptance of those levels. Problems identified during this re-evaluation of OU 1 groundwater inorganic data will be addressed in the OU 12 ROD. As indicated by current and past OU 12 background bedrock and overburden groundwater dissolved and total inorganic analyses, the amount of turbidity in a sample can affect the inorganic concentrations reported by the laboratory. Inorganic concentrations typically decrease in the filtered (dissolved) samples, as compared to the non-filtered (total) samples. Turbidity is often generated during sample collection in both bedrock and overburden monitoring wells.

Soil samples from OU 1, the former radiological UST liquids, and waste samples from the LLRWDS trenches did not contain inorganic concentrations indicative of source areas. Inorganic concentrations in OU 1 groundwater are attributable to natural occurrence, background variation, and/or impacts of turbidity.

Occurrence of Ra-226. Ra-226, one of the most abundant naturally occurring radioactive isotopes, was detected in 80 out of 108 soil samples throughout OU 1. The site-related Ra-226 data have been compared to two sets of off-site background sample concentrations that were developed in 1993 and 1994, respectively. If the site soil samples, collected in 1993 and before, are compared with the 1993 background concentrations, no exceedances of background are observed. If the 1993 and before site soil samples are compared with the 1994 background concentrations, many exceedances of background area observed. However, the exceedances of 1994 background values are a result of analytical method changes between the two years. The reporting limit, or minimum detectable activity, was lower for the 1994 background sample analyses due to increased analytical sensitivity. The background data reported in 1994 therefore had lower and more reliable values than the 1993 background data, with the result that samples collected in 1993 and before exceeded the lower 1994 background values. Based on this fact, and the widespread occurrence of Ra-226, Ra-226 detected at OU 1 is believed to be naturally occurring. Radioactive Isotope Analytical Results. During the RI, several analytical issues regarding the quantification and identification of radioactive isotopes were identified. Issues associated with the analysis for Uranium (U)-235, Americium (Am)-241, and Neptunium (Np)-237 are discussed in the following paragraphs. For a detailed explanation of the specific technical issues associated with the radioactive isotopic analytical program, refer to the OU 1 RI (ABB-ES, 1995a).

Gamma spectroscopy U-235 results are considered questionable due to analytical interference caused by Ra-226. U-235 analyzed by alpha spectroscopy was not subject to this interference and provided more accurate data.

Am-241 was detected once in a surface soil sample at Area A. Am-241 is primarily an alpha-emitting isotope accompanied by low energy gamma rays, however, the low gamma energy is subject to analytical interferences. The laboratory noted that the peak used to identify and quantify Am-241 in this sample had a bad peak shape which indicated an interference. Therefore, the Am-241 result is considered suspect.

Np-237 is also primarily an alpha-emitting isotope accompanied by low energy gamma rays. During the analysis for Np-237 by gamma spectroscopy, interferences were noted by the laboratory, thereby calling into question the identification and quantitation of this isotope. Therefore, the identification and quantification of Np-237 detected by gamma spectroscopy in sediments associated with Area A, Butterfield Brook, and East Loring Lake are questionable.

5.2 SUMMARY OF CONTAMINANTS DETECTED

Results of the RI sampling and analysis are briefly summarized in the following paragraphs. Results are presented for the radiological USTs and waste disposal trenches first, followed by additional results for each site area.

Radiological USTs. Essentially no contaminants were detected in liquid, sediment, or scrape samples collected from the five USTs at Areas A, B, C, D, and F. Analysis of confirmatory soil samples collected from the bottom of the UST excavations also did not detect contamination indicative of a source.

Waste Disposal Trenches. Radiological contamination (enriched uranium) was detected in samples collected from the waste disposal trenches at Areas C and E. Subsequently, removal actions were performed in both trenches in 1994.

SECTION 5

Confirmatory samples collected from the limits of the trench excavations following the removal action indicate that radioactive waste was successfully removed from both Trench C and E.

Arsenic was detected above background in only one of 18, closely gridded, confirmatory soil samples at Trench E. Arsenic is not a documented contaminant associated with OU 1. Detection of arsenic in Area E may be attributable to rodenticides used to control burrowing animals at the trench location.

Area A. Polyaromatic hydrocarbons (PAHs), pesticides, polychlorinated biphenyls (PCBs), lead, and zinc were detected above background in Area A surface soil. The PAHs, PCBs, and metals in surface soil are attributable to non-point source erosion and runoff from nearby parking areas, roads, and former operational areas. The detection of pesticides is consistent with the compounds and concentrations detected at other OUs at LAFB. The presence of these compounds is a result of routine basewide use of pesticides.

Radiological analytes detected above background levels in Area A soils and sediments were Am-241, Np-237, Ra-226, U-235, Thorium (Th)-231, and Th-234. Am-241, Np-237, and U-235 detections are suspect due to analytical difficulties in identification and quantitation as discussed in Subsection 5.1. The detections of Th-231 and Th-234 are considered to be naturally occurring. Ra-226 is an abundant naturally occurring radionuclide and was detected in nearly all OU 1 soil samples.

Concentrations of aluminum, chromium, manganese, and nickel exceeded USEPA Safe Drinking Water Act Maximum Contaminant Levels (MCLs) and MEDEP Maximum Exposure Guidelines (MEGs). These inorganics are naturally occurring and have not been identified as site-related. The detection of these inorganics above background in OU 1 groundwater is assumed to be a result of sample turbidity. Tritium was detected in one groundwater sample at a level approximately 100 times lower than the drinking water standard.

Area B. In general, detected volatile organic compounds (VOCs), semivolatile organic compounds (SVOCs), pesticides, and inorganics were below background levels or at low estimated concentrations. No PCBs were detected at Area B. U-235, Th-231, and Ra-226 were detected above background levels in soil at Area B. As discussed previously, the identification and quantitation of U-235 is suspect due to analytical interferences. Th-231 is believed to be naturally occurring at

Area B. The Ra-226 detection at Area B is typical of Ra-226 concentrations throughout OU 1.

Inorganics (iron, chromium, lead, manganese, and nickel) were detected in groundwater above MCLs and MEGs at Area B. These detections are likely associated with turbidity.

Area C. Three inorganic analytes (calcium, mercury, and sodium) were detected above background in Area C soil. These analytes were detected at estimated concentrations except for calcium, which is considered to be an essential nutrient. U-235, Th-234, and Ra-226 were detected in soil at concentrations above background concentrations. These radionuclides are naturally occurring and their detection above background levels is attributable to natural differences in background and analytical variability. The detections of U-235 are suspect due to analytical difficulties in identification and quantification.

Trace or estimated levels of VOCs and pesticides were detected in groundwater samples collected in 1993, but were not reported in 1994 samples. Inorganics (aluminum, lead, and manganese) were detected in groundwater; however, as in the case of other groundwater samples, the concentrations likely reflect natural variation and the effect of sample turbidity. Th-232 and U-234 were detected at background in groundwater at Area C. Both of these radionuclides are naturally occurring. Tritium was detected in one groundwater sample at a level approximately 100 times lower than the drinking water standard.

Area D. Other than the detection of three pesticide compounds at or just above background levels, no organic or inorganic contamination was detected in soils at Area D. Ra-226 was the only radionuclide detected in soil at Area D at a concentration above background levels.

Aluminum, lead, and manganese were detected in Area D groundwater at concentrations greater than MCLs and MEGs. However, the groundwater samples were turbid. Th-230, U-234, and U-238 were detected above background levels in 1993, but not in 1994. These are naturally occurring radionuclides.

Area E. PAHs were detected below LAFB background levels in surface soil in the drainage swale at Area E. The occurrence of these compounds is attributable to non-point source runoff from the former operations at Area E. No other organic contaminants were detected in soil at Area E. Lead, silver, and sodium were

detected above background concentrations in surface soil samples. In subsurface soil, cadmium (in five samples), zinc (in three samples), and arsenic, cobalt, and lead were detected above background values. The arsenic and lead detections were each localized in a trench confirmatory sample. Their detection is not indicative of widespread residual contamination.

U-235, Ra-226, Ra-228, Th-228, and Th-231 were detected above background levels in Area E soils. These isotopes are naturally occurring and were present at concentrations that slightly exceeded LAFB background concentrations.

Aluminum, chromium, lead, and manganese were detected in groundwater samples at Area E at concentrations exceeding MCLs and MEGs. As with the other sites throughout the OU, the concentrations of inorganics are assumed to reflect the effect of turbidity in the samples. Th-230, U-234, and U-238 were detected in groundwater in 1993 at estimated concentrations above background levels. In the 1994 groundwater sampling round, Th-228, Th-230, and Th-232 (estimated concentration) were detected above background levels. These analytes are naturally occurring, and their detection above background levels is attributable to analytical variability and differences in natural background concentrations. Tritium was detected in one groundwater sample at a level approximately 100 times lower than the drinking water standard.

Area F. No organic compounds were detected in soils at Area F other than one detection of a compound believed to be a laboratory contaminant. Pesticide compounds were detected at concentrations below, or slightly exceeding, background concentrations. The occurrence of these compounds is attributable to the routine application of pesticides at LAFB. Arsenic, lead, and zinc were detected at estimated concentrations and were slightly above background levels. Protactinium (Pa)-234, Th-234, and U-235 were detected in soils at Area F. Th-234, Pa-234, and U-235 are naturally occurring radionuclides. The U-235 result is suspect due to analytical interferences.

No organic compounds were detected in groundwater at concentrations above MCLs or MEGs at Area F. Two pesticide compounds were detected in the 1993 round of sampling, but were not reported in 1994. Aluminum is the only inorganic analyte detected above MCLs and MEGs in groundwater at Area F. In 1994, aluminum was detected below the MEG in the same well. No radionuclides were detected above background concentrations in groundwater at Area F.

Area G. No VOCs were detected in surface soil. PAHs were detected in several surface soil samples. Total PAH concentrations exceeded background levels at one location at the head of a drainage swale that receives runoff from a paved parking area, from Building 216 floor drains, and is located adjacent to and downslope from the fuel pipeline and two fuel oil USTs. In general, inorganics, primarily barium, lead, sodium, and zinc, were detected sightly above background concentrations in several samples at Area G. Most of these samples showed detections of one or two inorganic analytes. One sample, located at the head of the drainage ditch that receives runoff from much of the site, contained 11 inorganic analytes above background concentrations.

TCE and total xylenes were detected once at estimated concentrations in subsurface soil. The concentration of TCE is not indicative of a potential source area. Total xylenes in subsurface soil had been detected in an area where fuel-related contaminants had been detected by field screening. No PCBs were found in subsurface soils at Area G. One inorganic compound (sodium) was detected above background levels.

Pa-234, Th-231, and U-235 were detected in soils at Area G. Th-234, U-235, and Pa-234 are naturally occurring radionuclides. The U-235 result is questionable due to interferences in quantitation and identification. Ra-226 is an abundant, naturally occurring radionuclide and was detected in nearly all OU 1 samples.

In 1993, trichloroethene (TCE) was detected above its MCL and MEG in one downgradient groundwater sample. However, in 1994, TCE concentrations in groundwater were below regulatory limits. Several PAHs, indicative of fuel contaminants, were detected at estimated concentrations in a downgradient monitoring well location. Pesticides were detected at low, estimated concentrations in the samples in 1993, and only in deep bedrock groundwater in 1994. The occurrence of these compounds is attributable to the widespread application of pesticides at LAFB.

U-234, U-235, Ra-226, Th-230, and Th-232 were detected in groundwater above background concentrations. These isotopes are naturally occurring and were detected sporadically during the groundwater sampling rounds. Their detection above background levels is likely the result of natural background differences and analytical variability. Tritium was detected in one groundwater sample at a level approximately 100 times lower than the drinking water standard.

6.0 SUMMARY OF SITE RISKS

Human health and ecological risk assessments were conducted to estimate the probability and magnitude of potential adverse human health and environmental effects from exposure to contaminants at OU 1. The risk assessments followed a four-step process:

- 1) contaminant identification, which identified those hazardous substances that were of significant concern;
- 2) exposure assessment, which identified actual or potential exposure pathways, characterized potentially exposed populations and receptors, and determined the magnitude of possible exposure;
- toxicity assessment, which considered the types and severity of adverse health effects associated with exposure to hazardous substances; and
- 4) risk characterization, which integrated the three earlier steps to estimate the potential risks posed by hazardous substances at the site, including carcinogenic and non-carcinogenic risks.

The methodologies of the baseline human health and ecological risk assessment for the site areas are discussed below, followed by a summary of the conclusions.

6.1 HUMAN HEALTH RISK ASSESSMENT

For the purpose of the human health baseline risk assessment, the areas within OU 1 were segregated as Area A and Areas B through G. Area A is situated west of East Loring Lake (see Figure 1-1) and is isolated from the remaining OU 1 areas, which are located to the east of the lake. The focus of the risk assessment was on both non-radiological (i.e., chemical) and radiological contaminants in soil, sediment, surface water, and groundwater. During the initial evaluation of data, contaminants of potential concern (CPCs) were identified. The rationale for exclusion of selected compounds is included in Tables 6-1 and 6-2. The CPCs were selected to represent potential hazards based on toxicity, concentration, frequency of detection, mobility, and persistence in the environment. A summary of the health effects associated with each CPC can be found in the RI Report (ABB-ES, 1995a).

Name					LORING A	IR FORCE BASE						
Search Company Compa				of	Detected	Detected	of all		MCI	MEG	CPC2	Notes
Semble Land, Collabor College United States 1.5			·		Concombation	concentration	- Campies	GIOGIA	MOL	MEG	OF O	Notes
Assertations										·····	••••	
Americanies 0.350 - 9.4100 1.3 0.050 0.0500 0.15 0.00 - 9.45 0.051 Minimical Philadelines 0.350 - 9.4100 1.3 0.057 0.050 0.15 0.00 - 9.45 Minimical Philadelines 0.350 - 9.4100 1.3 0.057 0.050 0.050 0.050 Minimical Philadelines 0.350 - 9.4100 1.3 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4100 1.3 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4100 1.3 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0.050 0.050 0.050 0.050 0.050 Minimical Philadelines 0.3500 - 9.4500 2.3 0.050 0			0.4400	112	0.005	0.0050	0.15	4/8/8				01
Series June 1980	Anthracene											
Record Description 0.8800 0.4100 1 / 3	Benzo(a)Anthracene	0 3600 -	0 3600	2/3	0 047	0 1600	0 129					
Cabasable 0.9800 0.4100 1 / 2 0.098 0.006 0.147 NOS NO												
Charlestee	Carbazole				0 056							Toxicity Screening ²
Process 9 900 9 4150 1 3 0 0 0 0 145 NDB - Ves Olean California 2 3 2 5 0 145 NDB - Ves Olean California 2 3 2 5 0 145 NDB - Ves Olean California Califo					0 043						Yes	Class ^I
International	Fluorene											
Protection	Indeno(1,2,3-c,d)Pyrene	0 3600 -	0 4100	1/3	0 049	0 049						
Ad - Old Color C												
Ad -DDC	гутеле	0 3600 -	0.3000		00/5	0.20	0 1783	NUB	- -		Yes	Class
Ad-DDT	PESTICIDES/PCBs											
Ancepton 1986 0,986 0,986 1 3 0,981 0,		0.0036	0.0036									
Endostatine 0.0096 0.009	Aroclar=1260	0 0360 -	0 0380									TOXICITY OCTERTING
Enders	Dieldrin Endesulten Sultete					0 0008	0 0016	0 0002			No	
Finds	Endosulfan Sulfate Endrin											
Entire Nations 0.0068 0.0041 1 / 3 0.0005 0.0005 0.0005 0.0007 - No Toxicity Screening* Methodocyclins 0.0150 0.0150 2 / 3 0.0015 0.0026 0.0009 0.010 NDB No Toxicity Screening* Methodocyclins 0.0150 0.0021 1 / 3 0.0002 0.0009 0.0009 0.010 NDB No Toxicity Screening* Methodocyclins 0.0015 0.0021 1 / 3 0.0002 0.0009 0.0009 0.001 NDB No Toxicity Screening* Methodocyclins 0.0015 0.0021 1 / 3 0.0002 0.0009 0.0010 0.0009 0.0009 0.0009 0.0009 Methodocyclins 0.0009	Endrin Aldehyde	0 0036 -	0 0038	1/3	0 0046	0 0046	0 0028	0 0008				
No. Fig. No.												
NUMBER	gamma - Chlordane											
All Principle								***************************************				TOXION DEFORMING
Artenic				2/2	0800	18100	12022	05400			Na	Dankmen und 3
Barlum	Arsenic											
Calculum	Barlum	0.0000	4 0000									Background ³
Chromanis		0 9300 -	1 0000									
Sepper 3 3 12 5 22 1 16 27 65 6 - No Background*	Chromium			3/3	20 2	33 1	27 9					Background ³
Inches												
Lead	Iron											Background ³
Mingranese	Lead			3/3	106	23 4	162	22 6			No	State
Nickel 3 / 3 22 44.1 35.37 73 - No Background'												
Polassium	Nickel											
Vanedrum 3 / 3 13 8 21 16 57 40 - No Background*	Potessium											Background ³ , Essential Nutrient ⁴
Arces B-G' Surface Soil* (0-2 feet bgs) (mg/ks) SEMINOLATILE ORGANIC COMPOUNDS SEMINOLATIC COM												
SEMINOLATILE ORGANIC COMPOUNDS	Zinc											Dueng/ out to
SEMINOLATILE ORGANIC COMPOUNDS	Areas BG: Surface Solls (0-2 feet t	hast Imalkat										
Benzo(a)Anthracene			•••••								•	
Denzelo Pyrene			0.4700									
Denze Dk/F Loranthene												
Chrysene	Benzo(b,k)Fluoranthene	0 3600 -	0 4700	2/9	0 059	0 145	0 3459	NDB				· · · · · · · · · · · · · · · · · · ·
Di-n-butylphthalate												
Phenanthrene	Di-n-butylphthalate											
Pyrene	Fluoranthene											
Design D												
PESTICIDES/PCBs 4,4'-DDD	bis(2-Chloroisopropyl)ether											
4,4'-DDD	bis(2-Ethylhexyl)phthalate	0 3500 -	0 4700	2/10	0 042	0 044	0 1701	NOB	_		No	
4,4'-DDD	PESTICIDES/PCBs											
4.4'-DDT	4,4'-DDD			4 / 10				0 47			Yes	Class ¹
Aldrin 0 0019 - 0 0025 1 / 10 0 0004 0 0001 NDB No Toxicity Screening* Arcolor-1260 0 0360 - 0 0480 4 / 10 0 0009 0 1 0 0385 0 24 Yes Diefdrin 0 0036 - 0 0048 5 / 10 0 0001 0 0006 0 0013 0 0002 No Toxicity Screening* Endosulfian I 0 0019 - 0 0025 2 / 10 0 0001 0 0006 0 0009 NDB No Toxicity Screening* Endosulfian II 0 0036 - 0 0048 1 / 10 0 0004 0 0004 0 0019 NDB No Toxicity Screening* Endosulfian Sulfate 0 0036 - 0 0048 1 / 10 0 0004 0 0004 0 0019 NDB No Toxicity Screening* Endosulfian Sulfate 0 0036 - 0 0048 3 / 10 0 0005 0 0023 0 0019 NDB No Toxicity Screening* Endrin 0 0035 - 0 0048 2 / 10 0 0004 0 0007 0 0017 0 0003 - No Toxicity Screening* Endrin 0 0035 - 0 0048 2 / 10 0 0004 0 0007 0 0017 0 0003 - No Toxicity Screening* Endrin Ndehyde 0 0035 - 0 0048 2 / 10 0 0005 0 0013 0 0018 0 0008 No Toxicity Screening* Endrin 0 0018 - 0 0025 1 / 10 0 0005 0 0013 0 0018 0 0008 No Toxicity Screening* Heptachlor 0 0018 - 0 0025 1 / 10 0 0002 0 0013 0 0008 0 0002 - No Toxicity Screening* Heptachlor Epoxide 0 0018 - 0 0025 3 / 10 0 00002 0 0013 0 0009 0 0001 No Toxicity Screening* Methoxychlor 0 0018 - 0 0025 2 / 10 0 0004 0 0005 0 0084 NDB No Toxicity Screening* Methoxychlor 0 0019 - 0 0025 2 / 10 0 00004 0 0002 NDB No Toxicity Screening* delta-BHC 0 0019 - 0 0025 2 / 10 0 00002 0 0002 No Toxicity Value*	4,4'-DDE											Class ¹
Aroclor - 1280 0 0380 - 0 0480 4 / 10 0 009 0 1 0 0335 0 24 Yes Deterrin 0 0036 - 0 0048 5 / 10 0 0001 0 0006 0 0013 0 0002 No Toxicity Screening² Endosulfan 1 0 0019 - 0 0025 2 / 10 0 0001 0 0006 0 0099 NDB No Toxicity Screening² Endosulfan II 0 0036 - 0 0048 1 / 10 0 0004 0 0004 0 0019 NDB No Toxicity Screening² Endosulfan II 0 0036 - 0 0048 1 / 10 0 0004 0 0004 0 0019 NDB No Toxicity Screening² Endosulfan Sulfate 0 0036 - 0 0048 3 / 10 0 0005 0 0003 0 0019 NDB No Toxicity Screening² Endrin 0 0035 - 0 0048 2 / 10 0 0004 0 0007 0 0017 0 0062 No Toxicity Screening² Endrin Aldehyde 0 0035 - 0 0048 2 / 10 0 0005 0 0013 0 0018 0 0008 No Toxicity Screening² Endrin Aldehyde 0 0035 - 0 0048 2 / 10 0 0005 0 0013 0 0018 0 0008 No Toxicity Screening² Heptachlor 0 0018 - 0 0025 1 / 10 0 0001 0 0001 0 0009 0 0002 No Toxicity Screening² Heptachlor Epoxide 0 0018 - 0 0025 3 / 10 0 0004 0 0005 0 0013 0 0009 0 0001 No Toxicity Screening² Methoxychlor 0 0180 - 0 0250 2 / 10 0 0004 0 0005 0 0084 NDB No Toxicity Screening² Methoxychlor 0 0019 - 0 0025 2 / 10 0 0006 0 0013 0 0022 NDB No Toxicity Screening² deta-BHC 0 0019 - 0 0025 2 / 10 0 00002 0 0002 No Toxicity Screening² deta-BHC	4,4'-DD1 Aldrin											Toxicity Screening ²
Endosulfan 0 0019 - 0 0025	Aroclor-1260	0 0360	0 0480	4 / 10	0 009	01	0 0335	0 24		·	Yes	
Endosulfan	Dieldrin Endosulfen i											
Endosulfan Suffate 0 0036 - 0 0048 3 / 10 0 0005 0 0023 0 0019 0 0062 No Toxicity Screening* Endrin 0 0035 - 0 0048 2 / 10 0 0004 0 0007 0 0017 0 0003 No Toxicity Screening* Endrin Aldehyde 0 0035 - 0 0048 2 / 10 0 0005 0 0013 0 0018 0 0008 No Toxicity Screening* Heptachlor 0 0018 - 0 0025 1 / 10 0 0001 0 0001 0 0009 0 0002 No Toxicity Screening* Heptachlor Epoxide 0 0018 - 0 0025 3 / 10 0 0002 0 0013 0 0009 0 0001 No Toxicity Screening* Methoxychlor 0 0180 - 0 0250 2 / 10 0 0004 0 0005 0 0084 NDB No Toxicity Screening* abpha-Chlordane 0 0019 - 0 0025 2 / 10 0 0006 0 013 0 0022 NDB Yes delta-BHC 0 0019 - 0 0025 2 / 10 0 0002 0 0002 0 0009 0 0002 No Toxicity Value*	Endosulfan II											
Endin Aldehyde 0 0035 - 0 0048 2 / 10 0 0005 0 0013 0 0018 0 0008 - No Toxicity Screening* Heptachlor 0 0018 - 0 0025 1 / 10 0 0001 0 0000 0 0002 - No Toxicity Screening* Heptachlor Epoxide 0 0018 - 0 0025 3 / 10 0 0002 0 0013 0 0009 0 0001 - No Toxicity Screening* Methoxychlor 0 0180 - 0 0250 2 / 10 0 0004 0 0005 0 0084 NDB - No Toxicity Screening* Methoxychlor 0 0019 - 0 0025 2 / 10 0 0008 0 013 0 0022 NDB - Yes delta-BHC 0 0 019 - 0 0025 2 / 10 0 00002 0 0002 - No Toxicity Screening*	Endosulfan Sulfate	0 0036 -	0 0048	3 / 10	0 0005	0 0023	0 0019	0 0062			No	Toxicity Screening ²
Heptachlor 0 0018 - 0 0025 1 / 10 0 0001 0 0001 0 0009 0 0002 - No Toxicity Screening ² Heptachlor Epoxide 0 0018 - 0 0025 3 / 10 0 0002 0 0013 0 0009 0 0001 - No Toxicity Screening ² Methoxychlor 0 0180 - 0 0250 2 / 10 0 0004 0 0005 0 0084 NDB No No Toxicity Screening ² alpha-Chlordane 0 0019 - 0 0025 2 / 10 0 0008 0 013 0 0022 NDB Yes delta-BHC 0 0019 - 0 0025 2 / 10 0 0002 0 0002 0 0009 0 0002 No Toxicity Value ⁶												
Heptachlor Epoxide 0 9018 - 0 9025 3 / 10 0 9002 0 9013 0 9009 0 9001 - No Toxicity Screening* Methoxychlor 0 6180 - 0 9025 2 / 10 0 9004 0 9005 0 9084 NDB - No - No Toxicity Screening* apha-Chlordane 0 9019 - 0 9025 2 / 10 0 9008 0 913 0 9022 NDB Yes delta-BHC 0 9019 - 0 9025 2 / 10 0 9002 0 9002 0 9009 0 9002 No Toxicity Value*	Heptachlor				0 0001							
apha-Chlordane 0 0019 - 0 0025 2 / 10 0 0006 0 013 0 0022 NDB Yes delta-BHC 0 0019 - 0 0025 2 / 10 0 0002 0 0002 0 0009 0 0002 No Toxicity Value ⁵	Heptachlor Epoxide	0 0018 -	0 0025	3 / 10	0 0002	0 0013	0 0009	0 0001			No	Toxicity Screening ²
delta-BHC 0 0019 - 0 0025 2 / 10 0 0002 0 0002 0 0009 0 0002 No Toxicity Value ⁶												Toxicity Screening ²
	delta-BHC		0 0025		0 0002		0 0009	0 0002				Toxicity Value ⁶
	gamma-Chlordane								_			

					9					
		Frequency	Minimum	Maximum	Mean					
	Range of SQLs	of Detection	Detected Concentration	Detected Concentration	of all Samples*	Back- Ground**	MCL	MEG	CPC?	Notes
		# ************************************	**************************************			7177117				
NORGANIC ANALYTES										-3
Aluminum		10 / 10	13900	20600	16655	25400	<u> </u>	=	No	Background ³
Arsenic Barium	· · · · · · · · · · · · · · · · · · ·	10 / 10 10 / 10	37	10 1 73	6 645 43 525	16 2 93 3			No	Background ³
Bervilium	024 - 12		23 6 0 3	054	0 471	18	- -		No No	Background ³ Background ³
Salcium	024 - 12	10 / 10	659	23500	5192 25	69700			No	Background ³ , Essential Nutrien
Chromium		10 / 10	24 4	33.9	31 41	56 9			No	Background ³
obalt		10 / 10	52	161	11 55	18.5	-		No	Background ³
Copper		10 / 10	39	383	20 545	65 6			No	Background ³
on		10 / 10	18800	32300	28455	47100	-	-	No	Background ³
.ead	149 17	8 / 10	86	33 4	20 145	22 6	-	-	No	State
sagnesium		10 / 10	3460	8950	7261 5	12700	-	.	No	Background ³ , Essential Nutrier
fanganese		10 / 10	248	999	627 05	1400			No	Background ³
Rercury	011 - 014		0 12	26	0 317	0 17	-	-	Yes	
lickef		10 / 10	122	46 5	37 475	73			No	Background ³
otassium	5.65	10 / 10	495	1110	826 15	2900	-	-	No	Background ³ , Essential Nutrier
ilver odium	0 85 - 1 5 37 6 - 57		12	12	0 6415	0 09			Yes	Faces at all bluestes and
anadium	37 6 - 57	8 / 10 10 / 10	57.2 18.2	124 30 4	80 35 23 14	110 40			No	Essental Nutrient ⁶ Background ³
line		10 / 10	182 348	30 4	81 11	83.9	 _		No Yes	packground.
		10710	 	. (3)	01.11	0.70			103	
reas B-G. Surface Soll Sample	JSS-2880* (0-1 feet bgs)	(mg/kg)								······································
EMIVOLATILE ORGANIC COMPOU	INDS									
-Methylnaphthalene		1/ 1	36	36	36	NDB	-	-	Yes	
nthracene		1/1	25	25	25	NDB	-	-	Yes	Class ¹
luoranthene		1/1	3 1	31	31	NDB			Yes	Class ¹
laphthalene		1/ 1	10	10	10	NDB		_	Yes	Class ¹
henanthrane		1/1	12	12	12	NDB	-		Yes	Class'
Pyrene		1 / 1	82	82	. 82	NDB			Yes	Class ¹
PESTICIDES/PCBs (mg/kg)	· · ·									
J.4'-DDT		1/1	0 021	0 021	0 021	0.94			Yes	
Aldrin		1/ 1	0 0036	0 0036	0 0036	NOB			Yes	• • • • • • • • • • • • • • • • • • • •
ndosulfan l		1/ 1	0 0013	0 0013	0 0013	NOB		-	No	Toxicity Screening ²
ndosulfan II		1/1	0 12	012	0 12	NDB			No	Toxicity Screening ²
ndosulfan Sulfate		1/ 1	0 024	0 024	0 024	0 0062	_	_	No	Toxicity Screening ²
ndrin		1/ 1	0 0027	0 0027	0 0027	0 0003	-	-	No	Toxicity Screening ²
ndrin Ketone		1/ 1	0 0052	0 0052	0 0052	0 003	-		No	Toxicity Screening ²
feptachlor Epoxide		1/ 1	0 011	0 011	0 011	0 0001			Yes	
ipha-Chiordane		1 / 1	0 0024	0 0024	0 0024	NDB	-		Yes	
eta-BHC		1/ 1	0 024	0 024	0 024	0 0002	_	_	Yes	
elta-BHC		1/1	0 011	0 011	0 011	0 0002			Yes	Class ^I , Toxicity Value ⁶
amma – BHC (Lindane)		1/ 1	0 024	0 024	0 024	NDB			Yes	
amma-Chlordane		1/1	0 0044	0 0044	0 0044	NDB			Yes	
NORGANIC ANALYTES										
NORGANIC ANALYTES		1/ 1	22000	22000	22000	25400			No	Background ³
Arsenic		1/ 1	22000 4.8	22000	22000 4.8	25400 16.2			No	Background ³
Barium		1/ 1	157	157	157	933	- -		Yes	Duong, Duile
admium	***************************************	1/ 1	118	11.8	11.8	0 31			Yes	
alcium		1/ 1	10700	10700	10700	69700			No	Background ³ , Essential Nutner
hromium		1/ 1	81 4	81.4	81 4	56.9			Yes	
obalt		1/ 1	193	193	193	185			No	Background ³
opper		1/ 1	790	790	790	65 6	-	-	Yes	
on		1/ 1	34400	34400	34400	47100	-		No	Background ³
ead		1/ 1	493	493	493	22 6	_	-	Yes	State ⁵
lagnesium		1/ 1	13500	13500	13500	12700			No	Essential Nutrient ^a
langanese		1/ 1	984	984	984	1400			No	Background ³
lercury		1/ 1	22	22	22	0 17	-	-	Yes	
lickel		.1/ 1	69 5	69 5	69 5	73	-		No.	Background ³
otessium		1/ 1	2170	2170	2170	2900			No	Background ³ , Essential Nutrier
odium		1/ 1	139	139	139	110		_	No	Essential Nutrient
anadium		1/ 1	68 3	68 3	68.3	40			Yes	
inc		1/ 1	1240	- 1240	1240	83 9	-		Yes	

	Range o	of	Frequency of Detection	Minimum Detected Concentration	Maximum Detected Concentration	Mean of all Samples*	Back- Ground**	MCL	MEG	CPC?	Notes
Areas B-G Subsurface Soil* (0-10 fo	eet bas) (ma/k	a)									
	oor Day (mark										
VOLATILE ORGANIC COMPOUNDS 1,2-Dichloroethene (total)	0 011	0 014	1 / 22	0 002	0 002	0 006	NDB			No	Frequency ⁷
2-Butanone	0 011 -	0 014	1 / 22	0 008	0 008	0 006	NDB			No	Frequency ⁷
Acetone	0 011 -	0 044	1 / 22	0 01	0.01	800 0	NDB			No	Frequency ⁷
Methylene Chloride	0 006	0 068	7 / 22	0 005	0 024	0 011	NDB		-	Νo	Toxicity Screening ²
Toluene	0011 -	0 014	1 / 22	0 002	0 002	0 006	NDB			No.	Frequency ⁷
Trichloroethene	0 011 -	0 014	4 / 22	0 001	0 003	0 005	NDB			No	Toxicity Screening ²
SEMIVOLATILE ORGANIC COMPOUNDS		***									
Benzo(a)Anthracene	036 -	0 47	1 / 21	0 11	0 11	0 19	NDB			No	Toxicity Screening ² , Frequency
Benzo(a)Pyrene	036 -	0 47	1 / 20	0 038	0 038	0 186	NDB		<u> </u>	No	Toxicity Screening ² , Frequency ³
Benzo(b,k)Fluoranthene	036 -	0 47	2 / 20	0 059 0 047	0 145 0 14	0 363 0 178	NDB NDB		<u> </u>	No No	Toxicity Screening ² Toxicity Screening ²
Butylbenzylphthalate	035 -	0 47 0 47	3 / 21_ 2 / 21	0 047	012	0 184	NDB	- -	<u>=</u>	No	Toxicity Screening ²
Chrysene Di-n-butylphthalate	035 -	0 47	1 / 21	0 043	0 043	0 187	NDB			No	Frequency ⁷
Fluoranthene	036 -	0 47	3 / 21	0 037	0 077	0 174	NDB		-	No	Toxicity Screening ²
Phenanthrene	035 -	0 47	1 / 21	0 048	0 048	019	NDB	_		No	Toxicity Screening ² , Frequency ²
Pyrene	0.36	0 47	3 / 21	0 049	0 085	0 175	NDB			No	Toxicity Screening ²
bis(2-Chloroisopropyl)ether	035 -	0 47	1 / 13	0 076	0 076	0 184	NDB			No	Frequency ⁷
bis(2-Ethylhexyl)phthalate	0 35 -	0 47	2 / 21	0 042	0 044	0 18	NDB		<u></u>	No	Toxicity Screening ²
PESTICIDES/PCBs											
4,4'-DDD	0 0036	0 019	4 / 21	0 0002	0 011	0 0025	0 47			No	Toxicity Screening ²
4,4'-DDE	0 0036	0 019	7 / 21	0 0003	0 014	0 0029	016	-		No_	Toxicity Screening ²
4,4'-DDT	0 0036 -	0 019	7 / 21	0 0015	0 042	0 005	0 94 NDB	- -		No No	Toxicity Screening ²
Aldrin	0 0018 -	0 0098	1 / 21	0 0004 0 009	0 0004	0 0012 0 0294	0 24			Yes	Frequency ⁷
Aroclor – 1260 Dieldrin	0 0036 -	0 019	5 / 21	0 0001	0 0006	0 0019	0 0002			No	Toxicity Screening ²
Endosulfan I	0 0018 -	0 0098	2 / 21	0 0001	0 0008	0.0011	NDB			No	Toxicity Screening ²
Endosulfan II	0 0036	0 019	1 / 21	0 0004	0 0004	0 0022	NOB	-		No	Toxicity Screening ² , Frequency ⁷
Endosulfan Sulfate	0 0036	0 019	3 / 21	0 0005	0 0023	0 0022	0 0062	-		No	Toxicity Screening ²
Endrin	0 0035 -	0 019	2 / 21	0 0004	0 0007	0 0021	0 0003		<u>=</u>	No	Toxicity Screening ²
Endrin Aldehyde	0 0035 -	0 019	2 / 21	0 0005	0 0013	0 0022	0 0008			No	Toxicity Screening ²
Heptachlor	0 0018 -	0 0098	1 / 21	0 0001	0 0001	0 0011	0 0002			No No	Toxicity Screening ² , Frequency ² Toxicity Screening ²
Heptachlor Epoxide	0 0018 -	0 0098	3 / 21 4 / 21	0 0002	0 0013	0 0011	NDB			No	Toxicity Screening ²
Methoxychlor alpha-Chlordane	0 018 -	0 0098	2 / 21	0 0004	0 0039	0 0017	NDB			No	Toxicity Screening ²
delta-BHC	0 0018 -	0 0098	2 / 21	0 0002	0 0002	0 0011	0 0002			No	Toxicity Value ⁶
gamma-Chlordane	0 0018 -	0 0008	3 / 21	0 0015	0.01	0 0017	NDB			No	Toxicity Screening ²
INORGANIC ANALYTES											
Aluminum			44 / 44	3900	23000	18408	25400	_		No	Background ³
Antimony	78 -	20	1 / 44	30	30	89	NDB		-	No	Frequency ⁷
Arsenic			44 / 44	07	110	10	162			Yes	
Barlum	50 -	50	42 / 44	36	80	55 2	933			No	Background ³
Beryllium	024 -	2	6 / 44	03	0 85	0 852	18			No.	Background ³
Cadmlum	11 -	2	5 / 44 27 / 44	<u>2</u> 659	23500	3486	0 21 69700	_ _		No No	Toxicity Screening ² Background ³ , Essential Nutrient ⁴
Catcium	2000 -	2000	44 / 44	24 4	23500	345	56 9	— <u>-</u>		No	Background ³
Chromium Cobalt	20 -	20	14 / 44	52		13	185	_		No	Toxicity Screening ²
Copper	- <u> </u>	4.0	44 / 44	39		22	65 6			No	Background ³
Iron			44 / 44	18800		33756	47100			No	Background ³
Lead	136 -	17	41 / 44	6.8	270	20 2	22 6			Yes	State
Magnesium			44 / 44	3460		7955	12700	-		No	Background ³ , Essential Nutrient ⁴
Manganese			44 / 44	248		651	1400		-	No	Background ³
Mercury	011 -	02	2 / 44	0 12		0 146	0 17			No	Frequency ⁷
Nicke!			44 / 44	12 2		42	73			No No	Background ³ Essential Nutrient ⁴
Potassium	2000 -	2000	22 / 44	495		1190	2900		 -	No No	Frequency ⁷
Silver	0.85 -	2000	1 / 44	1 2 57 2		1 2 729	110	-		No	Essential Nutrient
Sodium Uranium (total U-234, U-235, U-238)	376	2000	3 / 3			214	3 897			No	Toxicity Screening ²
Vanadium			44 / 44	182		25	40			No	Background ³
Zinc			44 / 44	34 8		94 3	83.9			No	Background ³
MALL OF THE PARTY											

	LORING AIR FORCE BASE												
	Range o		Frequency of Detection	Minimum_ Detected Concentration	Maximum Detected Concentration	Mean of all Samples*	Back- Ground**	MCL	MEG	CPC7	Notes		
tres A 1994 Groundwater* (mg/L)													
SEMIVOLATILE ORGANIC COMPOUNDS Phenol			1 / 1	0 002	0 002	0 002	NA				Taulah Cara iala al		
TRETO!			1/.1	0 002	0.002	0 002	NA.			No	Toxicity Screening ²		
NORGANIC ANALYTES													
duminum Barium			1/1		1 18 0 0219	1 18 0 0219	NA NA	0 05 #	1 43 1 5	No No	Toxicity Value ⁶ Toxicity Screening ²		
alcium			1/1	110	110	110	NA NA			No	Essental Nutrient		
hromium			1/1		0 311	0311	NA.	01	01	Yes	T. 12 2		
opper			1/1	0 0254 4 31	0 0254 4 31	0 0254 4 31	NA NA	13T		No Yes	Toxicity Screening ²		
ead			1/ 1	0 0022	0 0022	0 0022	NA.	0 015 T	0 02	No	State*		
lagnesium langanese			1/ 1	774	7 74 0 174	774 0174	NA NA	0 05 #	- 02	No.	Essential Nutrient		
ickel			1/ 1	0173	0 173	0173	NA NA	01	015				
otassium			1/ 1	0 983	0 983	0 983	NA	_		No	Essential Nutrient		
odium			1/ 1		7 65 0 139	7 65	NA NA			No	Essential Nutrient		
116			1/_1	0 139	0 139	0 139	NA.	5#		No	Toxicity Screening ²		
rea A 1993 Groundwater* (mg/L)	·							· · · · · · · · · · · · · · · · · · ·					
OLATILE ORGANIC COMPOUNDS													
otal Xylenes			1/ 1	0 001	0 001	0 001	NA	10	06	No	Toxicity Screening ²		
											2		
ESTICIDES/PCBs ndosulfan Sulfate			1/ 1	0 0000009	0 0000009	0 0000009	NA.			No	Taviety Caronin-2		
ndrin Aldehyde			1/ 1		0 0000018	0 0000018	NA.			No	Toxicity Screening ² Toxicity Screening ²		
eptachlor			1/ 1		0 000012	0 000012	NA	0 0004	0 00008		Toxicity Screening ²		
ORGANIC ANALYTES													
Juminum			1/ 1	30	30	30	NA	0 05 #	1 43	No	Toxicity Value ⁶		
rsenic			1/ 1	0 009	0 009	0 009	NA	0 05		Yes			
alcium hromium			1/1		128 0 0733	128 0 0733	NA NA	- 01		No	Essential Nutrient		
opper			1/ 1		0 0351	0 0351	NA NA	137	01	Yes	Toxicity Screening ²		
on			1/ 1	45 3	45 3	453	NA	03#	-	Yes			
ead lagnesium			1/ 1		0 0135	0.0135	NA.	0 015 T	0 02		State ⁵		
langanese			1/ 1		19 5 0 683	19.5 0.683	NA NA	0 05 #	- 02	No Yes	Essential Nutrient		
lickel			1/1		0 0687	0 0687	NA	01	0 15				
otassium odium			1/1		6 17	6 17	NA.			No	Essential Nutrient		
inc	••••		1/1		9 54 0 101	9 54 0 101	NA NA	5#		No No	Essential Nutrient ⁴ Toxicity Screening ²		
reas B-F 1994 Bedrock Groundwater	* (mg/L)												
OLATILE ORGANIC COMPOUNDS										· · · · ·	· · · · · · · · · · · · · · · · · · ·		
-Methyl-2-pentanone	0 002 -	0 002	2 / 12		0 0001	0 0001	NDB		-	No	Toxicity Screening ²		
otal Xylenes richloroethene	0 002 -	0 002	1 / 12		0 0002 0 0002	0 0009	NDB	10	0.6		Toxicity Screening ²		
choroenera	0 002 -	0 002	1 / 12	0 0002	0 0002	0 0009	NDB	0 005	0 005	No	Toxicity Screening ²		
EMIVOLATILE ORGANIC COMPOUNDS													
- Methylnaphthalene I-n-butylphthalate	0 01 -	0 01	1 / 12 2 / 12		0 004	0 0049	NDB NDB		0 22	No No	Toxicity Screening ² Toxicity Screening ²		
henol	0 01 -	0 01	1 / 12		0 0007	0 0048	NDB		- 0 22	No	Toxicity Screening ²		
ESTICIDES/PCBs 4'-DDT	0 00001 -	0 00001	1 / 12	0 000011	0 000011	0 00001	NDB		0 00083	No	Tovinhy Company		
DD1	0 00001 -	0 00001	1 / 12	0 000011	0 000011	0 00001	NUB		0 00003	NO	Toxicity Screening ²		
ORGANIC ANALYTES													
luminum	0 0015 ~	0 0015	12 / 12	, , , , , , , , , , , , , , , , , , , ,	33 4 0 0064	6 7631	0 145	0.05 #	1 43		Toxicity Value ⁶		
rsenic arium	0 0015	0 0015	5 / 12 12 / 12		0 222	0 0018	0 0639	0 05	15	Yes	Toxicity Screening ²		
eryllium	0 0003	0 0003	2 / 12	0 001	0 0012	0 0003	NDB	0 004		Yes	-		
alcium	0 0074	0 0074	12 / 12		262	104 7250	163 897	-			Essential Nutrient ⁴		
hromium opper	0 0074	0 0074	7 / 12 8 / 12		0 0404	0 0145 0 0157	NDB NDB	01 13T	01	Yes No	Toxicity Screening ²		
on			12 / 12	0 418	45 8	10 2228	0 313	03#		Yes			
ead	0 0007	0 0007	11 / 12		0 041	0 0084	NDB	0 015 T	0 02				
lagnesium langanese			12 / 12 12 / 12		30 3 1 19	14 7600 0 2361	22 91 0 0249	0 05 #	- 02	No Yes	Essential Nutrient ⁴		
ercury	0 0001 -	0 0001	1 / 12	0 00011	0 00011	0 0001	NDB	0 002	0 002		Toxicity Screening ²		
ickel	0 0226 -	0 0226	3 / 12		0 0447	0 0170	NDB	01	0 15	No	Toxicity Screening ²		
				0 542	5 65	1 6897	0 314	-	_	N٥	Essemal Nutrient		
otessium			12 / 12										
otassium odium anadium	0 012 - 0 0187 -	0 012	12 / 12	2 2 5	23 2 0 0194	6 4933 0 0071	15 213 NDB	<u> </u>	-	No No	Essential Nutrient ⁴ Toxicity Screening ²		

		Frequency	Minimum	Maximum	Mean	Bent				
	of	of Detection			ot all Samples*	Ground**	MCL	MEG	CPC?	Notes
r* (mg/L)		*	 							
					0.0002	MOD	0.5		No	Toxicity Screening ²
0 001 -										Toxicity Screening ²
										Toxicity Screening ²
										Toxicity Screening ²
										Toxicity Screening ²
0 001 -	0 001	1 / 12	0 0005	0 0003	0.0000	1000				
						Nen		0.000	No.	Toxicity Value ⁶
0 025	0 025							0 000		Toxicity Screening ²
0 01 -	0.01	1 / 12	0 001	0 001	0 0047	NDB		<u>-</u>		TOAKIN GOLCETHING
0 00002 -	0 00002	2 / 12	0 0000007	0 0000035	0 0000087	NOB		0.00083		Toxicity Screening ² Toxicity Screening ²
	0 00002	1 / 12						0.00083		Toxicity Screening ²
0 00001 -	0 00001	2 / 12	0 000002					0.0000		Toxicity Screening ²
0 00002 -	0 00002	2 / 12	0 0000007							Toxicity Screening ²
0 00002 -	0 00002	3 / 12	0 0000006							Toxicity Screening ²
	0 00001	3 / 12	0 0000004							
		4 / 12	0 0000011	0 0000055						Toxicity Screening ²
										Toxicity Screening ²
			0 0000017	0 0000017						Toxicity Screening ²
			0 0000003	0 0000025			0 002			Toxicity Screening ²
										Toxicity Value ⁶
										Toxicity Screening ²
0 00001	0 00001	3 / 12			0 0000039	NDB	0 002	0 0002	7 No	Toxicity Screening ²
	 	12 / 12								
0.0052 -	0.0052	5 / 12	0 0054							
		2 / 12	0 186	0 226						
			20 3	257						
0.0092	0.0092		0 0127	0 163						
			0.0123	0 127						
00111			0.4	40 6	11 7280					
0.002 =	0.002		0 0023	0 0272			0.015			
0.005			4 95	247	13 1442					
0.0042	0.0043				0 235	8 0 0249	0 05 #			
					0 000	1 NDB				
						3 NDB				
						0 0314		=_		
1/6 ~	1/0									
0 01 -	0 043	2 / 12						*	No	Toxicity Screening ²
water (mg/L	<u> </u>					_,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				
							0.05	# 1.	ia N	Toxicity Value
		1/1								
		1/ 1								
		1/ 1								
		1/ 1								
		1/ 1	10							
		1/ 1	0 20							
				3 0 0673						
					54					
					0.04					
						2 NA				
									15 Ñ	
										Essential Nutrient
		1/ 1							N	
		1/ 1	3 6 0 022						N	
	\$QLe \$QLe \$\frac{1}{2} \text{(mg/L)}\$ 0 001 - 0 001 - 0 001 - 0 001 - 0 001 - 0 002 - 0 001 - 0 001 - 0 0001 - 0 00002 - 0 00002 - 0 00001 -		Range of SQLs Detection **(mg/L) *	Range of SQLs	Range of SqLs	Range of SQLs	Range of Office Detected Detected Samples* Back- Ground** Concentration Samples* Ground** Ground** Concentration Samples* Ground** Grou	Range of soll Defection Defected Defected Samples* Ground** McL.	Range of Detection Samples* Ground*** MCL MEG	Range of Defection Defec

TABLE 6~1 NON-RADIOLOGICAL ANALYTES OF POTENTIAL CONCERN FOR HUMAN HEALTH RISK ASSESSMENT

LORING AIR FORCE BASE												
			Frequency	Minimum	Maximum	Mean						
	Range SQLs		of Detection	Detected Concentration	Detected	of all Samples*	Back→ Ground**	MCL	MEG	CPC	7 Notes	
	OQL8		Detection	Concentration	Concentration	Samples	Ground	MUL	MEG	CFC	Notes	
Areas B-F, 1993 Overburden Ground	dwater* (mg/L)							~~~~				
VOLATILE ORGANIC COMPOUNDS Tetrachioroethene			1/ 1	0 003	0 003	0 003	NA NA	0 005	0 003	No	Toxicity Screening ²	
Toluene			1/1	0 003	0 003	0 003	NA NA	1	14	No	Toxicity Screening ²	
Total Xylenes			1/1	0 003	0 003	0 003	NA.	10	0.6	No	Toxicity Screening ²	
SEMIVOLATILE ORGANIC COMPOUND	<u>s</u>								- 2 200	*1-	T-11-4-Makes	
4-Nrtrophenol			1/ 1	0 002	0 002	0 002	NA.	 	0 083	NO	Toxicity Value ⁶	
PESTICIDES/PCBs												
gamma-BHC (Lindane)			1/ 1	0 0000015	0 0000015	0 0000015	NA	0 0002	0 0002	No	Toxicity Screening ²	
INORGANIC ANALYTES											*	
Atuminum Arsenic			1/ 1	0 011	124 0 011	124 0 011	NA NA	0 05 # 0 05	1 43	No Yes	Toxicity Value ⁶	
Barium			1/ 1	0 468	0 468	0 468	NA.	2	15	No	Toxicity Screening ²	
Calcium			1/ 1	192	192	192	NA.			No	Essential Nutrient	
Chromium			1/ 1	0 19	0 19	0 19	NA	01	01	Yes		
Cobalt			1/1	0 0922	0 0922	0 0922	NA.			No	Toxicity Screening ²	
Copper			1/1	0 119	0 119	0119	NA.	13T	-	No	Toxicity Screening ²	
Iron Lead			1/1	178	176	176	NA NA	03# 0015T	0 02	Yes Yes	State ⁵	
Magnesium			1/1	0 0681 58 2	0 0681 58 2	0 0681 58 2	NA NA	00151	- 002	No	Essential Nutrient	
Manganese			1/ 1	3 43	3 43	3 43	NA NA	0 05 #	02	Yes	ess dilect i terrisold	
Mercury			1/ 1	0 00026	0 00026	0 00026	NA.	0 002	0 002	No	Toxicity Screening ²	
Nickel			1/ 1	0 244	0 244	0 244	NA.	01	0 15	Yes	Exceeds MCL and MEG ⁸	
Potassium			1/ 1	124	12.4	124	NA			No	Essential Nutrient	
Sodium			1/ 1	518	5 18	5 18	NA	-		No	Essential Nutrient	
Vanadium			1/ 1	0 108	0 108	0 108	NA NA	-		Yes		
Zinc			1/ 1	0 346	0 346	0 346	NA	5#		No.	Toxicity Screening ²	
Area G 1994 Groundwater* (mg/L) VOLATILE ORGANIC COMPOUNDS			· · · · · · · · · · · · · · · · · · ·									
1,2-Dichloroethene (total)	0 002	0 002	2/4	0 0007	0 001	0 0009	NDB			No	Toxicity Screening ²	
2-Hexanone Acetone	0 002 - 0 002 -	0 002	1/4	0 011	0 011	0 00225 0 00275	NDB NDB			No No	Toxicity Value ⁵ Toxicity Screening ²	
Benzene	0 002 -	0 002	1 / 4	0 007	0 007	0 0008	NOB	0 005	0 005	No	Toxicity Screening Toxicity Screening	
Bromoform	0 002 -	0 002	1/ 4	0 0002	0 0002	0 0008	NDB	01		No	Toxicity Screening ²	
Chloromethane	0 002 -	0 002	1/4	0 0002	0 0002	0 0008	NDB	_	0 003	No	Toxicity Screening ²	
Ethylbenzene	0 002 -	0 002	2 / 4	~0 0002	0 0004	0 00065	NDB	07	07	No	Toxicity Screening ²	
Total Xylenes	0 002 -	0 002	1 / 4	0 0009	0 0009	0 000925	NDB	10	06	No	Toxicity Screening ²	
Trichforoethene	0 002 -	0 002	2 / 4	0 002	0 004	0 002	NDB	0 005	0 005	No	Toxicity Screening ²	
SEMIVOLATILE ORGANIC COMPOUND	s			·····								
2-Methylnaphthalene	0 01 -	0 01	1/4	0 002	0 002	0 00425	NDB		_	No	Toxicity Screening ²	
Acenaphthene	0 01 -	0.01	1/4	0 004	0 004	0 004875	NDB	-	-	No	Toxicity Screening ²	
Anthracene	0 01 -	0 01	1 / 4	0 0009	0 001	0 0039875	NDB	_	_	No	Toxicity Screening ²	
Dibenzofuran	0 01 -	0.01	1/4	0 002	0 002	0 004625	NOB	-		No	Toxicity Screening ²	
Fluorene Naphthalene	0 01 -	0 01	1/4	0 005	0 005	0 005 0 004625	NDB	=	-	No	Toxicity Screening ² Toxicity Screening ²	
Phenanthrene	001 -	0 01	1 / 4	0 003	0 004	0 004625	NDB NDB	- -		No No	Toxicity Screening ²	
Phenol	001 -	0 01	1/4	0 000	0 007	0 00375	NDB	- -		No	Toxicity Screening ²	
			· · · · · · · · · · · · · · · · · · ·									
PESTICIDES/PCBs												
Aldrin		0 000005	1 / 4	0 000006	0 000006	0 000002938	NDB			Yes		
Endosulfan Sulfate	0 00001 -	0 00001	1/4	0 000015	0 000015	0 00000625	NDB	-		No	Toxicity Screening ²	
Endrin Aldehyde	0 000005	0.000001	1/4	0 000018	0 000018	0.000006625	NDB NDB	0.0004	0 00008	No No	Toxicity Screening ²	
Heptachlor alpha-BHC		0 000005	1 / 4	0 000011	0 000015 0 000007	0 000005125	NDB	0 0004	0 00008	No No	Toxicity Screening ² Toxicity Screening ²	
mprim WIV	3 555555 5	- 000000		V 000001	2 000001	2 000000000	1400			110		
INORGANICS												
Aluminum			4/4	0 323	3 18	2 09075	0 145	0 05 #	1 43	No		
Arsenic	0 0015 -	0 0015	2 / 4	0 0052	0 0058	0 003025	NDB	0 05		Yes		
Barrum			4 / 4	0 0104	0 112	0 0592375	0 0639	2	15			
Calcium	0 0074	0 0074	1 / 4	53 0 0156	149 0 0156	100 175 0 006675	163 897 NDB	- 01	01	No Yes	Essential Nutrient	
Copper	0 0086 -	0 0086	1 / 4	0 0134	0 0134	0 0054375	NDB	13T		No	Toxicity Screening ²	
Iron	V VVVV -	V 0000	4/ 4	0 228	987	3 34575	0313	03#		Yes		
Lead	0 0007	0 0007	3 / 4	0 00098	0 0038	0 00202	NDB	0 015 T	0 02	No		
Magnesium			4 / 4	4 76	126	7 1975	22 91			Nο		
Manganese			4 / 4	0 0099	0 29	0 0948	0 0249	0 05 #	02			
Potessium			4 / 4	0 418	23	6 379	0 314	-		No		
Sodium	·		4 / 4	3 35	92.5	26 16625	15 213			No	Essential Nutrient ⁴	

	Range	of	Frequency of	<i>y</i>	Minimum Detected	Maximum Detected	Mean of all	Back-				
	SQL		Detection	١	Concentration	Concentration	Samples*	Ground**	MCL	MEG	CPC?	Notes
ea G 1993 Groundwater* (mg/L)												
OLATILE ORGANIC COMPOUNDS												
-Hexanone			1/	1	0 032	0 032	0 032	NDB			No	Toxicity Value ⁶
hloroform	0 001 -	0 001	1/	4		0 001	0 000625	NDB	01	-	No	Toxicity Screening ²
thylbenzene	0 001 -	0 001	2 /	4		0 001	0 0006875	NDB	0.7	07	No	Toxicity Screening ²
oluens	0 001	0 001	1/	4		0 0008	0 000575	NDB	1	14	No	Toxicity Screening ²
otal Xylenes	0 001 -	0 001	2/	4	0 002	0 003	0 001625	NDB	0 005	0.6		Toxicity Screening ² Exceeds MCL and MEG ⁸
richloroethene	0 001 -	0 001	3 /	4	0 001	0 006	0 0025	NDB NDB	0 005	0 005		
is-1,2-Dichloroethene	0 001 -	0 001	2 /	- 4	0001	0 002	0 001	NDB	007	001	105	Ciass
EMIVOLATILE ORGANIC COMPOUNDS								NDB				2
- Methylnaphthalene	001 -	0 01	2/	4		0 008	0 005				No No	Toxicity Screening ²
cenaphthene	001	0.01	1/	4		0 007	0 005125	NDB NDB	<u>-</u>		No	Toxicity Screening ² Toxicity Screening ²
luorene	0 01 -	0 01	17	4		0 007	0 005				No	Toxicity Screening ²
laphthalene Phenanthrene	001 -	0 01 0 01	1/	4	0 005	0 003	0 008875	NDB NDB			No	Toxicity Screening ²
ls(2-Ethylhexyl)phthalate	0 024 -	0 046	11	4		015	0 005	NDB	0 006	0 025		ronary octooning
· · · · · · · · · · · · · · · · · · ·	0024 -	0 040			015			1100	0 000	0 020		
EST/CIDES/PCBs	0 00001 -	0 00001	2 /	4	0 0000019	0 0000023	0 000003937	NDB	<u>-</u>		No	Toxicity Screening ²
lidrin Dektrin	0 00001 -	0 00002	2/	4		0.0000023	0 000006825	NDB	_=	0 00002		Toxicity Screening ²
ndosullan II	0 00002 -	0 00002	1/	4		0 0000018	0 000008912	NDB		- 00002	No	Toxicity Screening ²
ndosulian li ndrin Aldehyde	0 00002 -	0 00002		- 4		0 0000025	0.000009062	NDB			No	Toxicity Screening ²
leptachlor	0 00002 -	0 00001		- 4		0 0000024	0 00000435	NDB	0 0004	80000 0		Toxicity Screening ²
bha-BHC	0 00001 -	0 00001	1/	4		0 0000015	0 000004125	NDB		-	No	Toxicity Screening ²
Noha-Chlordane	0 00001 -	0 00001	17	4		0 0000025	0 000004687	NDB	0 002	0 00027	No	Toxicity Screening ²
letta-BHC	0 00001 -	0 00001	2/	4		0 0000081	0 000004662	NDB	_	_	No	Toxicity Value ⁶
jamma-BHC (Lindane)	0 00001 -	0 00001	2/	4		0 0000029	0 000003937	NDB	0 0002	0 0002	No	Toxicity Screening ²
amma-Chlordane	0 00001 -	0 00001	17	4	0 0000061	0 0000061	0 000005137	NDB	0 002	0 00027	No	Toxicity Screening ²
NORGANIC ANALYTES				_				************				
luminum			4 /	4	1 15	32 7	13 65125	0 1450	0 05 #	1 43	No	Toxicity Value ⁶
Arsenic	0 0052 -	0 0052	1/	4	0 0064	0 0098	0 003975	NDB	0.05	-	Yes	
Barlum	0 145 -	0 145	3 /	4	0 0289	016	0 0778125	0 0639	2	15		
Calcium			4./	4		170	91 7	163 8970	-		No	Essential Nutrient ⁴
Chromium			4/	4		0 0212	0 015125	NDB	01	0.1		
Cobalt	0 0136 -	0 0136	1/	_ 4		0 0165	0 0087	NDB	-		No	Toxicity Screening ²
Copper	0 0112 -	0 0112	2/	4		0 0173	0 0094375	NDB	137		No	Toxicity Screening ²
fon			4 /	_ 4		183	10 52475	0 3130	03#		Yes	6 1.1.3
ead	0 002 -	0 002	3/	4		0 0202	0 008175	NDB	0 015 T	0 02		
Magnesium			41	_4		145	9 735	22 9100	0.05.#	- 02	No	Essential Nutrient ⁴
/anganese	0.0440	~~~~	4/	4		0 455 0 0185	0 2761 0 0098	0 0249 NDB	0 05 #	015		Toxicity Screening ²
Nickel	0 0142 -	0 0142	1/	4		15.5	5 72625	0 3140		- 015	No	Essential Nutrient
otassium			4 /	4		15 5 58 2	22 28	15 2130			No	Essential Nutrient
Sodium			*/		4 03	30.2	26 20	104,130				
MISCELLANEOUS PARAMETERS										0.000:-	77.	- Clared
Low Detection Limit Vinyl Chloride	0 0001 -	0 0001	1/_	4	0 0001	0 0001	0 0000625	NDB	0 002	0 00015	Yes	Class ¹
Area G 1992 Groundwater* (mg/L)			·····									·
OLATILE ORGANIC COMPOUNDS												
1,2-Dichloroethene (total)			1 /	1	0 005	0 005	0 005	NDB			Yes	
Acetone			1/		0 018	0 018	0.018	NDB			Yes	
Ethylbenzene			1/			0 001	0 001	NDB	0.7	0.7	Yes	
Total Xylenes	•		- i/		0 003	0 003	0 003	NDB	10	0.6		
Trichloroethene			1/		0 002	0 002	0 002	NDB	0 005	0 005		
INORGANIC ANALYTES												
Jranium (total U-234, U-235, U-238)			1 /	1	1 167	1 167	1 167	NDB	20	-	Yes	
VIG. 10 Kil O 204, O 200, O 200)												

TABLE 5-1

NON-RADIOLOGICAL ANALYTES OF POTENTIAL CONCERN FOR HUMAN HEALTH RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

	Range o	of	Frequency of Detection	Minimum Detected Concentration	Maximum Detected Concentration	Mean of all Samples*	Back- Ground**	MCL	MEG	CPC3	Notes
Area A: Surface Water (mg/L)			<u>~</u>	>	GALAN LIKIN		~~~		,		
									-		
PESTICIDES/PCBs			1/1	0 0000011	0 0000011	0 0000011	0 0003	-		Yes	
replacino											
NORGANIC ANALYTES											Background ³ , Essential Nutrient ⁴
Calcium		-	1/1	52 6 0 0123	52 6 0 0123	52 6 0 0123	67200 2.7	- -		No. Yes	Background, Essential Nutrient
Copper			1/1	0 488	0 486	0 486	961			No	Background ³
Magnesium			1/1	2 85	285	2 85	8280			No	Background ³ , Essential Nutrient
langanese			1/1	0 0453	0 0453	0 0453	62 6			No	Background ³
Sodium			1/1	43	43	43	6520			No	Background ³ , Essential Nutrient
Area Ar Sediment (mg/kg)			······								,
EMIVOLATILE ORGANIC COMPOUNDS	•										
- Methylphenol	04-	0 46	2/3	0 097	0 13	0 1 4 7		-	-	No	Toxicity Screening ²
cenaphthene	04	0.51	1/3	016	0 16	0 210		_		Yes	Class
Inthracene Senzo(a)Anthracene	04	0 51 0 46	1/3	0 21 0 072	0 21 0 47	0 227 0 252	····			Yes Yes	Class
senzo(a)Antriracene Senzo(a)Pyrene	04 -	0.51	1/3	038	036	0 252		_ <u>=</u> _		Yes	
Benzo(b,k)Fluoranthene			3/3	0 047	0 67	0 331		-		Yes	
Benzo(g,h,i)perylene	04 -	0 51	1/3	0 13	0 13	0 200		-		Yes	Class ¹ Toxicity Screening ²
Carbazole Chrysene	04 -	051	1/3 3/3	02	02 046	0 223 0 225		 -		Yes	Class ¹
Dibenzofuran	04 -	0 51	1/3	0 072	0 072	0 181				No	Toxicity Screening ²
luoranthene	04 -	04	3/3	0.091	1.3	0 549		-		Yes	Class ¹
luorene	04-	0 51	1/3	0 11	011	0 193				Yes	Class ¹
ndeno(1,2,3-c,d)Pyrene	04 -	051	1/3	0 21 0 063	0 21 0 94	0 227 0 401		-		Yes Yes	Class ¹
Phenanthrene Pyrene	04-	- 04	3/3	0 069	0 72	0315				Yes	Class
PESTICIDES/PCBs	0 0052 -	0 0052	2/3	0 0007	0 012	0 0033	0 077			No	Toxicity Screening ²
1,4'-DDE 1,4'-DDT	0 0052 -	0,0002	3/3	0 0007	0 0018	0 0013	02		-	No	Toxicity Screening ²
Aldrin	0 0021 -	0 0033	1/3	0 0051	0 0051	0 002	0 0003			No	Toxicity Screening ²
Aroclor – 1254	0 045 -	0 064	1/3	0 22	0 22	0 0598	0 33		-	Yes	
Aroclor – 1260	0 052 -	0 052	2/3	0 14 0 0003	0 74 0 0059	0 2387 0 0033	0 14	=		Yes No	Toxicity Screening ²
Dieldrin Endosulfan Sulfate	0 0045 -	0 0052	2 / 3 3 / 3	0 0003	0 0039	0 0033	0 0007			No	Toxicity Screening ²
Endrin	0 004 -	0 0064	2/3	0 0004	0 0025	0 0019	0 0008			No	Toxicity Screening ²
Endrin Aldehyde	0 0052 -	0 0052	2/3	0 0038	0 014	0 0065	0 0058		-	No	Toxicity Screening ²
Heptachlor Epoxide	0 002 -	0 0027	1/3	0 0004	0 0004	0 0009	0 0006	-		No	Toxicity Screening ²
Methoxychlor	0 021 -	0 027	1/3	0 002	0 002	0 0088	0 0013 0 0006			No No	Toxicity Screening ² Toxicity Screening ²
alpha – Chlordane delta – BHC	0 0027 -	0 0033	1/3	0 0016	0 015 0 0004	0 0038 0 0012	NDB	_ <u>-</u> _		No	Toxicity Value
gamma-Chlordane	0 0027 -	0 0033	1/3	0 0004	0 004	0 0019	0 0018	-		No	Toxicity Screening ²
NODCANIO ANALYSTIC							······································				
NORGANIC ANALYTES			3/3	14700	18800	16950	23000		_	No	Background ³
Arsenic			3/3	76		9 1667	167		_	No	Background ³
Barlum			3/3	46 1	150	962	114			No	Toxicity Screening ²
Beryllium	12 -	16	1/3	0 46	0 48	0 6233	0 63	-		No	Background ³
Calcium			3/3	1830		4678 3333 38 5833	17100 50 2			No No	Background ³ , Essential Nutrient Background ³
Chromium Cobalt		-	3/3	30 6 11 6		16 2167	27 8		 -	No	Background ³
Sopper Sopper			3/3	43.2	1200	371 9	43 8		_	Yes	
ron			3/3	25800		38883 3333	42600	_		Yes	
ead			3/3	24 5	256	84 4667	24			Yes	State*
Magnesium			3/3	7280		8580 2555	16800 2990			No Yes	Background ³ , Essential Nutrient
Manganese Mercury	0 12 -	016	3/3	225 0 27	5070 0.67	0 2383	013			No	Toxicity Screening ²
vercury Nickel	012 -	0.10	3/3	40 1	63.6	49 55	16			No	Toxicity Screening ²
Potessium	892 -	892	2/3	958	1140	858 3333	1140		-	No	Background ³ , Essential Nutrient
Sodium			3/3	- 86 7	138	102 7333	84 8	<u>-</u>		No	Essential Nutrient
Uranium (total U-234, U-235, U-236)			1/3	0 0078		NC.	NDB			No Yes	Toxicity Screening ²
Vanadium Zina			3/3	19 7 76 9	54 6 655	33 4 286 3167	39 4 120			Yes No	Toxicity Screening ²
Zinc			3/3	189	692	200 J 10/	120			140	. C. COLL O COLL OF THE PARTY

NOTES

Class' — Although the toxicity screening ratio was less than 0.01, this compound belongs to a class of compounds where at least one compound within this class has a risk ratio greater than 0.01 Toxicity Screening" — Chemicals with low ratios (i.e., less than 0.01) are not considered chemicals of potential concern (CPCs)
Background? — Sample concentrations detected are below background concentrations.
Essential Nutrient" — Analyte is an essential human nutrient (magnesium, calcium, potensium, sodium) and is not considered a CPC
State? — The Maine Department of Environmental Protection (MEDEP, 1990) guidance states lead concentrations less than 15 µg/L in groundwater and 125 mg/kg in soil are not evaluated quantitatively Toxicity Value? — Compound cannot be evaluated quantitively because toxicity values are not available Frequency — Frequency of detection is less than 5 percent
Exceeds MCL/MEG? — Maximum concentration is greater than MCL and/or MEG

- T Action Level

 * If the mean exceeds the maximum concentration, only the maximum concentration will be used in a quantitative evaluation

 ** Background for pesticides/PCBs provided for information only Concentrations of pesticides/PCBs were not screened against background concentrations

 ** Secondary Standard

 SCL Sample Quantitation Limit

 MCL Maximum Contaminant Level, Drinking Water Regulations and Health Advisorles, U.S. Environmental Protection Agency Office of Water, May 1995

 MEG Maximum Exposure Guideline, Maine Department of Human Services, September 1992

 mg miligram

 kg kilogram

 L liter

 ug microgram

 pg microgram

 pg microgram

 pg microgram

 pg microgram

 pg bold ground surface

 NA Background groundwater concentrations are not available for overburden wells

 NDB Background not determined

 NC mean not calculated

 - = No MCL or MEG available

TABLE 6-2 SUMMARY OF RADIOLOGICAL ISOTOPES FOR HUMAN HEALTH RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

Radiological	Range of	Frequency of	Minimum Detected Concen-	Maximum Detected Concen	Back-				•
<u>Analyte</u>	SQLs	Detection	tration	tration	Ground	MCL	MEG	CPC?	Notes
SURFACE SOIL (0-2	feet): AREA A ^a								
AMMA SPECTROSCO			0.177	A F-7-7			•	V	
Americium—241 Radium—226	0 138 -0 155 700 -700	1/3	0 577 1 44	0 577 1 89	0 081 0 94	-	-	Yes Yes	
SURFACE SOIL (0-2								100	
GAMMA SPECTROSCO			4.00	0.67	0.04				
Radium - 226	07 -141	3 / 9	1 86	2 67	0 94	_		Yes	
SUBSURFACE SOIL (0-10 feet): AREA	A *							
GAMMA SPECTROSCO									
Americium – 241	0 138 -0 155	1/3	0 577	0 577	0 081		<u>-</u>	Yes	
Radium – 226	700 - 700	2 / 3	1 44	1 89	0 94			Yes	
SUBSURFACE SOIL	SAMPLES (0-10 fe	et): AREAS B—G	C						
GAMMA SPECTROSCO	PY - 1 - Hour Counts	(pCi/g)							
Radium – 226	0 066 -1 41	49 / 61	0 246	2.67	0 94			Yes	
Radium – 228	0 172 -0.192	42 / 46	0,666	1 11	0 95	-		Yes	
LPHA SPECTROSCOP	Y (pCı/g)								
Plutonium - 239/240	0 013 -0 07	4 / 46	0 02	0 034	0 29				Background ²
rotactinium – 234		46 / 46	0.52	0 83	1.3			No	Background ²
horium – 227	0 015 -0 21	25 / 46 46 / 46	0 018 0 838	0 09	1 6 1,2			No Yes	Background ²
Thorium – 228 Thorium – 230	0 676 -0 941	31 / 46	0 61	1 703	1.4			Yes	
Thorium – 231	0 02 -0 1	30 / 46	0 01	0 08	0 05			Yes	
Thorium – 232	,	46 / 46	0 804	1 227	11	_	_	Yes	
Thorium – 234		46 / 46	0 52	0 83	1,3			No	Background ²
Jranium – 234		46 / 46	0 47	1 38	14				Background ²
Jranium – 235	0 02 -0.1	30 / 46	0.01	0 08	0 05			Yes	
Jranium – 238		46 / 46	0 52	0 83	1.3			No	Background ²
COMPOSITE SAMPLE	S (0-14 feet): AR	EAS B-G ^d							
AMMA SPECTROSCO									
Radium 226	0 901 -1 08	8 / 14	0 938	1 94	0 94			Yes	
GROUNDWATER: AR	EA A, 1994 °								
GROSS BETA (pCi/L)		1/1	18	18	459	50	NA	No	Below MCL ³
TRITIUM (pCi/L)		1/1	538	538	NDB	20000	NA	Yes	Below MCL ³
GROUNDWATER: AR	EA A, 1993 °								
GROSS ALPHA (pCi/L)		1/1	24	24	1655	15	NA	Yes	Exceeds MCL ⁴
GROSS BETA (pCi/L)		1/1	34	34	459	50	NA.		Below MCL ³
ALPHA SPECTROSCOF	V (nCill)								
ALPHA SPECTROSCOP Thorium – 230	T (POI/L)	1 / 1	21	21	0 625	NA	NA	Yes	
		1/1	2	2	1 096		NA		
Jranium – 234									

TABLE 6-2 SUMMARY OF RADIOLOGICAL ISOTOPES FOR HUMAN HEALTH RISK ASSESSMENT

OPERABLE UNIT TRECORD OF DECISION LORING AIR FORCE BASE

Radiological	Range of	F	requency of	Minimum Detected Concen-	Maximum Detected Concen-	Back-				
Analyte	SQLs	D	etection	tration	tration	Ground	MCL	MEG	CPC?	[†] Notes
GROUNDWATER: ARE	AS B-G, 19	94 1								
GROSS ALPHA (pCi/L)	1 -	3 8	7 / 16		61	20,91	15	NA	Yes	Exceeds MCL ⁴
GROSS BETA (pCI/L)	3 -	3	12 / 16	37	55	6	50	NA	Yes	Exceeds MCL ⁴
TRITIUM (pCi/L)	400 -	400	3 / 16	400	497	NDB	20000	NA	Yes	Below MCL ³
EPA METHOD 9320 (pCi/	// \									
Radium - 226	05-	0.5	3 / 4	0.69	1 37	1 767	5	NA	No	Background ² , Below MCL
ALPHA SPECTROSCOPY	(pCı/L)	·		্য						<u> </u>
Protactinium - 234			4 / 4	0.07	0 33	0 376	NA	NA	No	Background ²
Thorrum - 228	0 05 -	0.4	1 / 4	1 28	1 69	0 241	NA	NA	Yes	
Thorium-230	0.14	0.14	3 / 4	0.42	1.79	1.159	, NA	NA	Yes	
Thorium - 232	0 05 ~	0 05	3 / 4	- 0.05	1 37	0.05	NA.	NA	Yes	
Thorium - 234			4/4	0 07	0 33	0 376	NA	NA	No	Background ²
Uranium - 234 Uranium - 238	·····		4 / 4	0 12	0.6	0.541		NA.	Yes	
Oranium-236			4 / 4	007	0 38	0 376		NA	Yes	
GROUNDWATER: ARE	AS B-G, 19	93 [†]								
GROSS ALPHA (pCi/L)			16 / 16	12	50	20.91	15	NA	Yes	Exceeds MCL ⁴
GROSS BETA (pCi/L)	3 -	12	9 / 16	_ ~	52	6	50	NA	Yes	Exceeds MCL ⁴
EPA METHOD 9320 (pCi/	4 Y					·····				
Radium - 226	04 -	11	1/7	1.6	1,6	1 767	5	NA	No	Background ² , Below MCL
				1.0	1,0		-			Davingiouna , Dolow Mor
ALPHA SPECTROSCOPY	(pCi/L)							•		
Thorium - 230			7/7	0.9	6.3	1.159	NA	NA	Yes	
Uranium – 234	06-	06	7/7	07	6 65	0 54		NA	Yes	
Uranium - 238	0 65 -	0 65	6 / 7	0 62	6 73	0 376		NA	Yes	
GROUNDWATER: ARE	AS B-G, 19	92 g								
GROSS BETA	2	2	1/5	14 19	14 19	6	50	NA	No	Below MCL ³
	_						-		-	_
ALPHA-SCAN						-				
Radium - 226 Uranium - 234	0.5 -	0.5	2/5	1 32	2.01	1.767	5_	NA_	Yes	
Uranium - 235	1 -	<u>1</u> 1	4 / 5	⁻ 3,8 1 15	10 78 4.56	0 54 0.05		NA NA	Yes Yes	
Uranium - 238	1 -	-	1/5	3.04	3 04	0.03		NA NA	Yes	
SURFACE WATER: AR										
SOIL ACE WATER, AR	LA A alla OC									· · · · · · · · · · · · · · · · · · ·
GROSS ALPHA (pCi/L)	1 -	26	1 / 5	2.8	28	NC	15	NA	No	Below MCL ³
GROSS BETA (pCi/L)	3 -	3	3 / 5	61	18	5 1	50	NA	No	Below MCL ³
SEDIMENT: AREA A										
GAMMA SPECTROSCOP	Y -1 - Hour C	ounts (pCi/	(g)							
Neptunium – 237	0.45 -	0.5	1/3	0.509	0 509	NC	_		Yes	
Radium-226	07 -	1 28	1/3	2 43	2 43	3.16	_	_	Yes	
Thorium -234	0 78 -	1.48	1 / 3	2 09	2 09	NC	_		Yes	
Uranium 235	0 289 -	0 316	1 / 3	0.0168	0 11	NC	_		Yes	

TABLE 6-2 SUMMARY OF RADIOLOGICAL ISOTOPES FOR HUMAN HEALTH RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

Radiological Analyte	Range of SQLs		Frequence of Detection		Minimum Detected Concen— tration	Maximum Detected Concen— tration	Back Ground	MCL	MEG	CPC? 1	Notes
Analyte	GULB		patector	•	<u> </u>	11444441					
SEDIMENT: OU 13	ı										
GAMMA SPECTROSC	OPY -24-Hour	Counts (pCı/g)								
Radium – 226			4 /	4	0 972	1 51	145		_	Yes	
Thorium – 234	0 37 -	0 486	1/	4	0 92	0 92	NC	-	_	Yes	
Uranium – 235	0 0791 -	0 0966	2 /	4	0 112	0 207	NC			Yes	
ALPHA SPECTROSCO	OPY (pCi/g)										
Neptunium - 237	0.007 -	0 015	1/	4	0.072	0 072	ND	-		Yes	
Uranium – 234	0 304 -	0 531	3 /	4	0 568	0 733	NC	_	_	Yes	
Uranium – 238	0 335 -	0 567	2 /	4	0 704	0 753	NC			Yes	
SEDIMENT: WASTE	WATER TREAT	MENT I	LANT k								
ALPHA SPECTROSCO	OPY (pCi/a)										
Neptunium - 237	v v v v v v v		1/		0 033	0 033	ND			Yes	

NOTES

- 1 For radiological analytes selected as CPCs each detection above background will be quantitatively evaluated,
- with the exception of gross alpha and gross beta results in groundwater for which toxicity values are not available
- 2 Detected concentration does not exceed associated background concentration
- 3 Concentration of isotope or gross radiation does not exceed the associated MCL
- 4 Concentration of isotope or gross radiation exceeds the associated MCL
- 5 Highest 24-hour gamma spectroscopy result for Radium-226 in sediment, data inadequate for a statistical calculation

Sample Locations

- a Based on data from sample locations JSS-2081, -2082, JTB-2060
- b Based on data from sample locations JDT-2480, -2481, JSD-2560, JSS-2680, -2681, -2682, JTB-2260, -2660, JTP-2401
- c Based on data from sample locations JDT-2480 -2481, JSD-2560, JSS-2680, -2681, -2682, JTB-2260 -2660, JTP-2401, TRC01C through TRC23C TRE01C through TRE23C
- d _ Based on data from sample locations MTB-2180 -2181, -2280, -2281, -2282, -2380, -2381, -2480, -2481, -2482, -2580, -2680, -2681, -2682
- e Based on data from sample location JMW-2080
- f Based on data from sample locations JMW-2180 -2181, -2280, -2281, -2282, -2380, -2381 -2480, -2481, -2482 -2580, -2680, -2681, -2682
- 9 Based on data from sample locations JMW-2180, -2280, -2380, -2480, -2682
- h Based on data from sample locations JSW-0041, -0042 -0043, -0073, -2080
- 1 Based on data from sample locations JDT-2080, 2081, JSD-2060
- I Based on data from sample locations JSD-0041, -0042, -0043, -0073
- k Based on data from sample location JSD-0068

Acronyms

- SQL Sample Quantitation Limit
- MCL Maximum Contaminant Level
- MEG Maximum Exposure Guideline
- CPC Chemical of Potential Concern
- mg milligram
- kg kilogram
- L Itter
- μg microgram
- bgs below ground surface
- ND not detected
- NA no MCL/MEG available
- - MCL/MEG not relevent for this medium
- NDB not detected in background

Potential human health risks associated with exposure to the CPCs were estimated quantitatively or qualitatively through the development of hypothetical exposure pathways. These pathways were developed to reflect the potential for exposure to hazardous substances based on present and potential future land uses. Current use exposure scenarios included older child trespasser and groundskeeper. Future use exposure scenarios included resident, construction worker, older child trespasser, groundskeeper, commercial/industrial worker, and forestry worker.

For each pathway evaluated, an average and a reasonable maximum exposure (RME) estimate was generated, corresponding to exposure to the average and the maximum contaminant concentrations detected in that particular medium.

Excess lifetime cancer risks were determined for each exposure pathway by multiplying the exposure level with the chemical-specific cancer factor. Cancer potency factors have been developed by USEPA from epidemiological and animal studies to reflect a conservative upper bound of the risk posed by potentially carcinogenic compounds. That is, the true risk is unlikely to be greater than the estimated risk. The resulting risk estimates are expressed in scientific notation as a probability (e.g., 1x10⁶ or one in a million) and indicate (using this example) that an average individual is not likely to have greater that a one in a million chance of developing cancer over a lifetime of site-related exposure to the compound at the stated concentration. Current USEPA practice considers carcinogenic risks to be additive when assessing exposure to a mixture of hazardous substances.

The hazard quotient (HQ) was also calculated for each pathway as a measure of the potential for noncarcinogenic health effects. An HQ is calculated by dividing the exposure level by the reference dose (RfD) or other suitable benchmark for non-carcinogenic health effects for an individual compound. RFDs have been developed by USEPA to protect sensitive individuals over the course of a lifetime and they reflect a daily exposure level that is likely to be without an appreciable risk of an adverse health effect. RfDs are derived from epidemiological or animal studies and incorporate uncertainty factors to help ensure that adverse health effects will not occur. The HQ is often expressed as a single value (e.g., 0.3) indicating the ratio of the stated exposure to the reference dose value (in this example, the exposure as characterized is approximately one third of an acceptable exposure level for the given compound). The HQ is only considered additive for compounds that have the same or similar toxic effect (e.g., the HQ for a compound known to produce liver damage should not be added to a second compound whose toxic effect is kidney damage). The sum is referred to as the hazard index (HI).

SECTION 6

The results of the human health risk assessment are summarized in Subsection 6.4.

6.2 ECOLOGICAL RISK ASSESSMENT

Following a methodology similar to the human health risk assessment, the ecological risk assessment evaluates potential ecological effects resulting from plant and wildlife exposures to contaminants at OU 1. Ecological CPCs were selected for both non-radiological and radiological analytes detected in surface soil, sediment, and surface water. The rationale for exclusion of selected compounds are included in Tables 6-3 through 6-7.

Representative ecological receptor species were selected for the habitat associated with OU 1. For Area A, five representative wildlife species were selected to quantitatively evaluate the magnitude of potential ecological exposures that may occur. The receptors include:

- short-tailed shrew (Blarina brevicauda); small mammal, omnivore
- American woodcock (Scolopax minor); small bird, omnivore
- maritime garter snake (Thamnophis sirtalis pallidula); reptile, omnivore
- red fox (Vulpes vulpes); predatory mammal, carnivore
- barred owl (Strix varia); predatory bird, carnivore

In addition, potential impacts to terrestrial plants and earthworms, representative of potential exposure to other soil invertebrates, were also selected for risk evaluation.

Based on a habitat evaluation for Areas B through G, the following five representative species were selected for the ecological exposure evaluation:

- meadow vole (Microtus pennsylvanicus); small mammal, herbivore
- American robin (Turdus migratorius); small bird, omnivore
- maritime garter snake; reptile, omnivore
- red fox; predatory mammal, carnivore
- American kestrel (Falco sparverius); predatory bird, carnivore

Five representative species were also selected to evaluate the risks associated with potential exposure of wildlife to radiological contaminants in sediment, including:

TABLE 6-3 CHEMICALS OF POTENTIAL CONCERN FOR THE AREA A SURFACE SOIL [a] ECOLOGICAL RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

		FREQUENCY	MOMIXAM	,	
AVERAGE		OF	BACKGROUND		
(mg/kg) [b]	(mg/kg)	DETECTION	CONCENTRATION (mg/kg) [c]	CPC?	NOTES
		173		Yes	
		STORE OF THE PROPERTY OF THE P		Yes	
				Yes	
		3	NA NA	Yes	
0.329 *	0,218 ``````	173	NA NA	Yes	, , ,
	0.056	1/3	NA NA	Yes	
·	0.150	2/3	NA	Yes	
0.237	0,420	^ ^ 27/3	NA	Yes	
0.145 *	0,050		, NA	Yes	
"0.14\$ *	0,049	7/3		Yes	5
0.210	0,3 6 0 11	2/3	· '''		`w` .
0.178	0.280	2 / 3			,
		Same Salaharan	•		
	S. S. 111.	umanan umum Maranan maranan umum	(97) (35) 3.3	*	,
0 0327					•
		202 20302 20 00000000			

		2/3	• •		
		1/3			
		2/3		Yes	
			NA (d)	Yes	
** * * * * * * * * * * * * * * * * * * *		173	NA [d]	Yes	
		3	(NA (d)	Yes	
0.0045 *	0 0028	2 / S	NA	Yes	
		Marine A			
13 033	16 100		05.400	No	tal
			_		[e]
		3 77			[e]
		x - Lewis C	*		[e]
					[e]
•	•				[e,f]
		2 × 10 3 × 12 × 12 × 12 × 12 × 12			(e)
			. = , =		(e)
				No	[e]
•	2 2 2 22 22 22 22 22 22 22 22 22 22 22		The ends	No	[e,f]
,	23;4 🗥 📜	3/3	22,6	Yes	
6,460	7,490	373	12,700	No	[e,f]
430	504	3/3	1400	No	[e]
35 4	44 1	<u> </u>	73	No	[e]
831	986	3/3	2,900	No	[e,f]
57 3	85 4	3/3	110	No	[e,f]
186	21.0	3 / 3		No	[e]
65,0	89.9	200 40 40 40 40 40 40 40 40 40 40 40 40 4	*	Ŷes `	(-)
	AVERAGE (mg/kg) [b] 0 150 * 0.150 * 0	(mg/kg) [b] (mg/kg) 0 150 * 0.065 0.150 * 0.065 0.129 0.160 0.129 0.160 0.329 * 0.218 0.147 * 0.056 0.124 0.150 0.237 0.420 0.145 * 0.050 0.145 * 0.049 0.210 0.360 0.178 0.280 0 0027 0.0610 0.0010 * 0.0003 0.0010 * 0.0003 0.0016 * 0.0008 0.0025 0.0031 0.0013 * 0.0002 0.0028 0.0046 0.0014 * 0.0005 0.0015 * 0.0028 13,933 16,100 5 37 6.20 30 4 36 6 0.0014 * 0.0028 13,933 16,100 5 37 6.20 30 4 36 6 0.40 * 0.23 2,127 2,830 27 9 33.1 9 97 11.6 18 3 22 1 26,167 30,200 16.2 23,44 6,460 7,490 430 504 35 4 44 1 831 986 57 3 85 4 18 6 21 0	AVERAGE (mg/kg) DETECTION 0 150 * 0.065 1/3 0.150 * 0.065 1/3 0.129 0.166 2/3 0.161 * 0.056 1/3 0.329 * 0.218 1/3 0.147 * 0.056 1/3 0.124 0.150 2/3 0.145 * 0.050 1/3 0.145 * 0.050 1/3 0.145 * 0.049 1/3 0.210 0.380 2/3 0.178 0.280 2/3 0.010 * 0.0009 0.0009 0.0009 0.0019 0.0035 0.0016 * 0.0008 1/3 0.0015 0.0016 0.0008 1/3 0.0016 * 0.0008 1/3 0.0016 * 0.0008 1/3 0.0018 * 0.0002 1/3 0.0014 * 0.0005 1/3 0.0014 * 0.0005 1/3 0.0014 * 0.0005 1/3 0.0014 * 0.0005 1/3 0.0015 0.0016 0.0008 0.0014 * 0.0005 1/3 0.0015 0.0016 0.0005 0.0016 * 0.0008 0.0016 0.0017 0.0017 0.0018 0.0005 1/3 0.0018 0.0018 0.0018 0.0019 0.0018 0.0018 0.0019 0.0018 0.0018 0.0019 0.0018 0.0018 0.0019 0.0018 0.0018 0.0010 * 0.0008 0.0018 0.0011	AVERAGE (mg/kg) [b] (mg/kg) DETECTION CONCENTRATION (mg/kg) [c] 0.150 * 0.065	AVERAGE (mg/kg) (b) (mg/kg) DETECTION CONCENTRATION (mg/kg) [c] CPC ? 0 150 * 0.065

[[]a]Based on samples JSS-2081, JSS-2082 and JTB-2060

⁽b) Average concentration is the arithmetic mean of all sample results with 1/2 the SQL used for non-detects. Some averages may exceed maximum concentrations due to elevated SQLs.

[[]c]Base-wide surface soil background concentrations

[[]d]Analyte has been detected in background samples, however, these concentrations are not being used to screen for CPCs

Consideration of background levels of pesticides will be discussed in the risk uncertainty section

[[]e]Maximum concentration of analyte is below maximum surface soil background concentration

[[]f]Analyte is an essential nutrient, and is considered to be hazardous via ingestion in the terrestrial food web only at very high concentrations

^{*}Average concentration exceeds maximum due to elevated SQLs

NA = not available

Shaded analytes are CPCs

${\it TABLE~6-4} \\ {\it CHEMICALS~OF~POTENTIAL~CONCERN~FOR~THE~AREAS~B-F~SURFACE~SOIL~[a]~ECOLOGICAL~RISK~ASSESSMENT} \\$

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

	CONCEN	TRATION	FREQUENCY	MAXIMUM		
	AVERAGE	MAXIMUM	OF	BACKGROUND		
ANALYTE	(mg/kg) [b]	(mg/kg)	DETECTION	CONCENTRATION (mg/kg) [c]	CPC ?	NOTES
EMIVOLATILES						
Benzo(b,k)Fluoranthene	0,341 *	0.082	1 / 4	NA	Yes	
bis(2-Ethylhexyl)phthalate	0.145 *	0.044	2/5	NA	Yes	
Chrysene	0 172 *	0,054	1/5	NA	Yes	
Fluoranthene	0,178 *	0 077	1 / 5	NA	Yes	
Phenanthrene	0.188 *	0,048	1 / 5	NA	Yes	
Pyrene	0.174 *	0 057	1 / 5	NA	Yes	
PESTICIDES/PCBs						
Aroclor-1260	0,0191 *	0.0090	1 / 5	NA [d]	Yes	
delta-BHC	0.0009 *	0 0002	1 / 5	NA [d]	Yes Yes	
4,4'-DDD	0.0012 *	0,0010	3 / 5	na (d) na (d)	yeş Yes	
4,4'DDE	0.0024	0.0045	5 / 5	NA [d]	Yes	
4,4'-DDT	0,0044	0.0095	4 / 5	NA [d]	Yes	
Dieldnn	0.0016 *	0 0006	2 / 5		Yes	
Endosulfan Sulfate	0,0019 *	0 0005	1 / 5	NA [d]		
Endกก	0,0018 *	0.0007	1 / 5	NA [d]	Yes	
Endrin Aldehyde	0,0017 *	0.0005	1 / 5	NA [d]	Yes	
Heptachlor Epoxide	* 6000	0 0002	1 / 5	NA [d]	Yes	
NORGANICS					No. To	,
Aluminum	16,020	17,800	5 / 5	25,400	No [e	
Arsenic	7 21	10 1	5 / 5	162	No [e	-
Barium	44 4	59 9	5 / 5	93 3	No [e	
Beryllium	0 52	0 54	3 / 5	1 80	No [e	-
Calcium	4,394	17,800	5 / 5	69,700	-	if]
Chromium	31 4	33 9	5 / 5	56 9	No [6	-
Cobalt	126	161	5 / 5	18 5	No (e	
Copper	20 3	27 2	5 / 5	65 6	No [e	
Iron	29,430	32,300	5 / 5	47,100	-	:,f]
Lead	21.7	32.1	5 / 5	22.6	Yes	
Magnesium	7,680	8,950	5 / 5	12,700		e,f]
Manganese	735	998	5 / 5	1,400	No [6	*]
Mercury	0.57	2.60	1 / 5	0.17	Yes	
Nickel	40 7	46 5	5 / 5	73 0	No [6	e)
Potassium	823	1,110	5 / 5	2,900	No [6	∍,f]
Silver	0.767	1,20	1 / 5	0,090	Yes	
	100	124	5 / 5	110	No [f	1
Sodium	22 0	248	5 / 5	40 0	No [e	
Vanadium Zinc	85 5	141	5 / 5	83.9	Yes	

[[]a]Based on samples JDT-2480, JDT-2481, JSD-2560, JTB-2260, JTP-2041

[[]b]Average concentration is the arithmetic mean of all sample results with 1/2 the SQL used for non-detects. Some averages may exceed maximum concentrations due to elevated SQLs

[[]c]Base-wide surface soil background concentrations

[[]d]Analyte has been detected in background samples, however, these concentrations are not being used to screen for CPCs

Consideration of background levels of pesticides will be discussed in the nsk uncertainty section

[[]e]Maximum concentration of analyte is below maximum surface soil background concentration

[[]f]Analyte is an essential nutrient, and is considered to be hazardous via ingestion in the terrestrial food web only at very high concentrations

^{*}Average concentration exceeds maximum due to elevated SQLs

NA = not available

Shaded analytes are CPCs

CHEMICALS OF POTENTIAL CONCERN FOR THE AREA G SURFACE SOIL [a] ECOLOGICAL RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

	CONCENTE		FREQUENCY	MAXIMUM		
×	avërăgë 🐃	MAXIMUM	OF	BACKGROUND	• •	``
ANALYTE `	(mg/kg) [b]	(mg/kg)	DETECTION	CONCENTRATION (mg/kg) [c]	CPC ?	NOTES
SEMIVOLATILES			-2014FF-1			
2-Methylnaphthalene	6.16	36,0	incidental () 6	NA NA	Yes	
Anthracene	4.33	25.0 1	1/6	ÑA	Yes	
Benzo (a) Anthracene	0.935 *	0,110 ° · · · · · · · ·	1 / 6	NÀ	Yes	,
Benzo(a)Pyrene	0.923 *	0.038	1/6	NA	Yes	,
Benzo(b,k)Fluoranthene	1,86 *	0.145	1/6	ΝA	Yes	× .
bis(2-Chlorosopropyl)ether	0.924 *	0 076	******* / 6	NA	Yes	
Butylbenzylphthalate	0.912 *	0,140	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	NA NA	Yes	
Chrysene	0.937 *	0.120		NA	Yes	
Di-n-butylphthalate	0.923 * *	0.043	176	" NA	Yes	
Fluoranthene	0,631	3.10	mmanajajajaj	`, NA	Yes	
Naphthalene	1,83	10.0	. 176 . 176	NA NA		
Phenanthrene	2,16	12.0	> 5' '2' '		Yes	*
Pyrene		varanja	1/6	NA	Yes	CR 53 C
ryrene	1,49	8.200	,`, 3 / 6	, NA	Yes	
ECTIONE CIDAD						
PESTICIDES/PCBs	* ***		Sandar W. A.	,		
Aldrin	0,0013	0,0036	7 3 / 6 7 3 / 6	NA	Yes	
Aroclor – 1260	0 0480	0 1000	3 / 5	NA (d)	Yes	
beta-BHC	0.0048	0.0240	1/6	NA	Yes	
delta~BHC	0.0026	0.0110	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	NĂ [d]	Yes	
gamma-BHC (Lindane)	0.0048	0.0240	176	NA	Yes	
alpha-Chlordane	0 0032	0.0130	SEASTERN STATE OF THE	NA NA	Yes	,
gamma-Chlordane	0.0035	0.0100	4/6	NA	Yes	
4,4'DDD 4,4'DDE	0 0038	0.0110	minimum 5	" "ŅĄ [d]	Yes	
4,4'-DDT	0.0042	0.0140	476	ŅĂ [d]	Yes	
	0.0127	0.0420	60000000000000000000000000000000000000	NA (d)	Yes	
Dieldrin	0.0010 *	0.0004	375	NÀ [d]	Yes	
Endosulfan I	0.0009	0 0013	3/6	NA NA	Yes	
Endosulfan II	0.0214	0.1200	276	NA	Yes	**
Endosulfan Sulfate	0.0055	0.0240	3/6	NA [d]	Yes	
Endrin	0.0018	0.0027	2/6	NA [d]	Yes	
Endrin Aldehyde	0 0018 *	o oom	3333	NA [d]	Yes	
Endrin Ketone	0 0025	0,0052	176	NA [d]	Yes	
Heptachlor	0.0008 *	10000.0	75	NA [d]	Yes	
Heptachlor Epoxide	0.0026	0 0,110	3/6	NA [d]	Yes	
Methoxychlor	0.0062 *	0 0005	^```2`/`5	NA	Yes	
IORGANICS						
Aluminum	40.075					
	18,075	22,000	6/6	25,400	No	(e)
Arsenic	5 87	8 60	6/6	16,2	Nο	(e)
Barlum	61,8	157	6/6	93.3	Yes	
Beryllium	0 54 *	0.30 ‴	1/6	1 80	No	[e]
Cadmium	2.46	11,8 1 🛒	1)6	` ` ` 0.21	Yes	
Calcium	6,775	23,500	6 / 6	69,700	No	[e,f]
Chromium	39 7	81.4	ningerialistication /	569	Yes	*
Cobalt	11,9	19,3	6/6	18,5	Yes	
Copper	149	790	8/6	65,6	Yes	
Iron	28,633	34,400	6/6	47,100	No	[a]
Lead	97,7	493	^^^476			[e,f]
Magnesium	7,953	13,500	•	22 60 12 700	Yes	fen
-			, 6/6	12,700	No	[f]
Manganese	597	999	6/6	1,400	No	[e]
Mercury	0 42	2.20	ຼິ ຶ2 / 6	0.17	Yes	
Nickel	40 1	69 5	6/6	73 0	No	[e]
Potassium	1,053	2,170	6/6	2,900	No	[e,f]
Sodium	74 0	139	4/6	110	No	[f]
Vanadium	31,6		6/6	40.0	Yes	r.1
Zinc	271	1,240	6/6	83.9	Yes	

[[]a]Based on samples JSS-2680, JSS-2681, JSS-2682, JTB-2660, JTB-2680, JTB-2681, and JTB-2683

[[]b]Average concentration is the arithmetic mean of all sample results with 1/2 the SQLTused for non-detects. Some averages may exceed maximum concentrations due to elevated SQLs

[[]c]8ase-wide surface soil background concentrations

[[]d]Analyte has been detected in background samples, however, these concentrations are not being used to screen for CPCs

Consideration of background levels of pesticides will be discussed in the risk uncertainty section

[[]e]Maximum concentration of analyte is below maximum surface soil background concentration

[[]f]Analyte is an essential nutrient, and is considered to be hazardous via ingestion in the terrestrial food web only at very high concentrations

^{*}Average concentration exceeds maximum due to elevated SQLs

NA = not available

Shaded analytes are CPCs

TABLE 6-6 CHEMICALS OF CONCERN FOR THE AREA A (DRAINAGE DITCH) SURFACE WATER [a] ECOLOGICAL RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

ANALYTE	DETECTED CONCENTRATION (ug/L)	FREQUENCY OF DETECTION	MAXIMUM BACKGROUND CONCENTRATION (UQIL) [b]	SCREENING BENCHMARK (Ug/L)	CPC ?	NOTES
PESTICIDES/PCBs						
Heptachlor	0 0011	1 / 1	NA(d)	0 0038	No	[d]
NORGANICS						
Calcium	52,600	1/1	67,200	NA	No	[e,f]
Copper	12,3	1/1	27	0 205	Yeş	
Iron	486	1 / 1	961	16	No	(e)
Magnesium	2,850	1 / 1	8,280	NA	No	[e,f]
Manganese	45 3	1 / 1	62 6	112	No	[d,e]
Sodium	4,300	1 / 1	6,520	85,049	No	[d,e]

NOTES

- [a] Based on samples JSW-2080
- [b] Base-wide surface water background concentrations
- [c] Analyte has been detected in background samples, however, these concentrations were not used to screen for CPCs Consideration of background levels of pesticides is discussed in the risk uncertainty section
- [d] Maximum concentration of analyte below screening benchmark
- (e) Maximum concentration of analyte below maximum surface water background concentration
- [f] Analyte is an essential nutrient and is not known to adversely impact aquatic organisms except at very high concentrations NA = Not available

Shaded analytes are CPCs

TABLE 6-7 CHEMICALS OF CONCERN FOR THE AREA'A' (DRAINAGE DITCH) SEDIMENT (a) ECOLOGICAL RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

	CONCENTRATI		FREQUENCY	MAXIMUM	SCHEENING	,	
	AVERAGE	MUMIXAM	OF	BACKGROUND	BENCHMARK		
ANALYTE	(mg/kg) [b]	(mg/kg)	DETECTION	CONCENTRATION (mg/kg) [c]	(mg/kg)	CPC ?	NOTE
SEMIVOLATILES							
2-Methylphenol	0.147 *	0.130	273	NA NA	0,063	Yes	
Acenaphthene	0.210 *	0 160	773	NA	0.016	Yes	
Anthracene	0.227 *	0.510	773	NA.	0 0853	Yes	
Benzo(a)Anthracene	0.252	0.470	~~ <u>~~~~</u>	, NA	0.261	Yes	
Benzo(a)Pyrene	0,277	0 360	173	NA	0,43	No	[d]
Benzo(b,k)Fluoranthene	0 308	0 670	3/3	NA	45	No	[d]
Benzo(g,h,i)perylene	0 200 *	0 130	1/3	NA	0 78	No	[d]
Carbazole	0 223 *	0.200	1 / a	ΝA	NA	Yes	
Chrysene	0 225	0.450	373	NA.	0,384	Yes	
Dibenzoturan	0 181 *	0 072	173	NA	0 58	No	[d]
Fluoranthane	0.549	1,300	3/3	NA ()	0.6	Yes	, ,
Fluorene	0.193 *	0.110	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	NA	0.019	Yes	
Indeno(1,2,3-c,d)Pyrene	0.227 *	0 210	773	NA NA	0.88	No	(d)
Phenanthrene	0,401	0 940	373	NA	0.24	Yes	1-3
Pyrene	0.315	0 720	3/3	NA `	0,665	Yes	
•			42 (,,,	31776	,	
PESTICIDES/PCBs			/ ///				•
Aldrin	0 0020	0 0051		NA M	0 002	Yes	
Aroclor-1254	O 0598	0 2200	Sagarania (S. C. S.	NA (Í)	0.06	Yes	
Aroclor-1260	0.2387	0.7400	273	NA [f]	0.005	Yes`	
delta-BHC	0.0012 *	0 0004	` 1 /-3	NA	0 003	No	[d]
alpha-Chlordane	0 0038	0.0150	1/3	NA M	0.007	Yes	,
gamma-Chlordane	0 0019	0 0040	17~3	NA [f]	0 007	No	[d]
- _4,4′DDE	0.0033	0.0120	· · · · · · · · · · · · · · · · · · ·	NA [f]	0.005	Yes	[~]
4,4'-DDT	0,0013	0 0018	3 / 3	NA [f]	0 007	No	(d)
Dieldrin	0.0033	0 0059	* ((M) AN	0 007	Yes	(u)
Endosulfan Sulfate	0 0033	0 0039	S 10 Hadifida jana m			7	
Endrin	0 0019		a designation	NA [f]	0.002	Yes	
		0 0025	2// 3	NA (f)	0 003	No	[q]
Endrin Aldehyde	0.0065	0.0140	THE STATE OF THE S	NA II)	0,003	Yes	
Heptachtor Epoxide	0 0009 *	0 0004	1/3	NA [f]	0 005	No	[d]
Methoxychlor	0 0088 *	0.0020	f / 3	NA [f]	0 005	No	[d]
NORGANICS							
Aluminum	16,950	18,800	3°/3°	23,000	NA	No	[f]
Arsenic	9 17	10.4	3 / s	16.7	6	No	[1]
Barium	962	150	150,00° 55,00° 50° 3 7 3	114	20	Yes	UJ
Beryllium	0 62 *	0 48	1/3	0 63	05	No	64.63
Calcium	4,678	7,060	ີ 3້/ື3້	17,100			[d,f]
Chromium	38 6	48 4	_3 / 3 _3 / 3		NA 20	No	[d,g]
Cobalt	162	22 3		50 2	26	No 	[f]
	372		\	27.8	50	No	[d,f]
Copper		1,200	2.350	44	16	Yes	
Iron	38,863	56,500	3 7 3	42,600	50,000	Yes	(h]
Lead	84 5	256	`_````````````````````````````````````	24 0	`31	Yes	
Magnesium	8,580	10,000	3/3	16,800	NA	No	[d.g]
Manganese	2,555	5,070	3 7 3	2,990	460	Yes	
Mercury	0 24	0 67	<u>2</u>	0,13	0.5	Yes	
Nickel	49.6	63 6	3/3	16 0	16	Yes	
Potassium	858	1,140	2 / 3	1,140	NA	No	[d,g]
Sodium	103	138	3 / 3	848	NA	No	(g)
Uranium	0.057 *	0 051	1/3	NA NA	NA	Yes	
Vanadium	33 4	54 `6 **	3	39.4	ÑA	Yes	
Zínc	286	655	373	120	120	Yes	
Total Organic Carbon	3,400	3,400	- 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1	NA	NA	NA	

TABLE 6--7 CHEMICALS OF CONCERN FOR THE AREA A (DRAINAGE DITCH) SEDIMENT [a] ECOLOGICAL RISK ASSESSMENT

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

NOTES

- [a] Based on samples JDT-2080, JDT-2081 and JSD-2060
- (b) Average concentration is the arithmetic mean of all sample results with 1/2 the SQL used for non-detects. Some averages may exceed maximum concentrations due to elevated SQLs
- [c] Base-wide sediment background concentrations
- [d] Maximum concentration of analyte below screening benchmark
- [e] Analyte has been detected in background samples, however, these concentrations were not used to screen for CPCs

 Consideration of background levels of pesticides is discussed in the risk uncertainty section
- [f] Maximum concentration of analyte below maximum sediment background concentration
- (g) Analyte is an essential nutrient, and is not known to adversely impact aquatic organisms except at very high concentrations
- (h) Analyte is a CPC for aquatic exposures only

*Average concentration exceeds maximum due to elevated SQLs

NA = Not available

Shaded analytes are CPCs

6-20

- muskrat (Ondatra zibethicus); small mammal, herbivore
- belted kingfisher (Ceryle alcyon); medium-sized bird, piscivore
- maritime garter snake; reptile, omnivore
- great blue heron (Ardea herodias); large bird, omnivore
- mink (Mustela vison); predatory mammal, omnivore

With the CPCs and receptors selected, the evaluation of exposure pathways, toxicity of CPCs, and resulting risks followed an approach similar to that of the human health risk assessment.

Results of the ecological risk assessment are summarized in Subsection 6.4.

6.3 UNCERTAINTY EVALUATION

Quantitative estimates of risk are based on numerous assumptions, which are intended to be protective of human health and the environment (i.e., conservative). The interpretation of risk estimates is subject to a number of uncertainties as a result of the multiple layers of conservative assumptions inherent in risk assessments. As such, risk estimates are not truly probabilistic estimates of risk, but are conditional estimates, given a series of conservative assumptions about exposure and toxicity. While it is true that there are some uncertainties inherent in the risk assessment methodology that might lead to an underestimation of true risks, most assumptions bias the evaluation in the direction of overestimation of risk. This results in more conservative clean-up criteria, more protective of human health and the environment.

The possibility of underestimation of true risks may be caused by the exclusion of exposure pathways from quantitative evaluation (i.e., ingestion of homegrown produce from backyard garden plots) or through the exclusion of compounds from the risk assessment through the CPC selection procedure. However, the CPC selection procedure evaluated compounds that constituted more than 99 percent of the risk; therefore it is unlikely that the risks will be underestimated by a substantial amount.

Other sources of uncertainty that could cause overestimation of risks include the use of purposive sampling (biased targeting of "hot spots" or visible contamination); the estimation of exposure concentrations by the use of maximum detections (while assuming no degradation or dilution); the use of the 95 percent (or upper-bound 90 percent) exposure parameter values such as contact rate and exposure frequency and

duration; the use of conservatively derived toxicity values such as RfDs (incorporating multiple safety factors); and cancer slope factors, which are based on experimental animal data used in a multi-stage model.

6.4 RISK ASSESSMENT CONCLUSIONS

Summaries of both human health and ecological risk assessments are presented in the following paragraphs. The discussion begins with the radiological USTs and waste disposal trenches and ends with conclusions for Area A and Areas B through G.

Radiological USTs. Based on the UST data, analysis of confirmatory soil samples, and downgradient groundwater quality, the USTs were not sources of radiological or non-radiological contamination.

Waste Disposal Trenches. No human health radiological risks above regulatory target risk levels were associated with the Trench C and E confirmatory soil samples following the removal action.

Arsenic was detected above background concentrations in only one out of 18 confirmatory soil samples at Trench E. Based on this result, subsurface soil non-radiological human health carcinogenic and non-carcinogenic risks were predominantly attributable to arsenic in combined Areas B through G. However, arsenic is not a documented contaminant associated with OU 1 strategic weapons maintenance, nor was there widespread detection of this analyte. The single detection of arsenic may be the result of rodenticide application at the former Trench E location.

Area A Soils, Surface Water, and Sediments. No human health non-radiological risks have been identified at Area A in soils, surface water, or sediments above the regulatory target risk levels. No ecological radiological risks have been identified in Area A soils and sediments.

Total maximum cancer risks associated with exposure to radionuclides detected in soil above established background concentrations range from $5x10^4$ to $9x10^6$. Maximum radiological risks identified for sediment $(1x10^{-5})$ are less than the established background risks for that medium $(2x10^{-5})$. These risks represent a minimal incremental cancer risk above the LAFB background risks of $2x10^4$ to $8x10^6$

and are less than published total natural radiological background risks for the United States of $1x10^{-2}$ to $3x10^{-3}$ (Shleien, 1992).

A portion of the radiological human health risks is attributable to Am-241 associated with a single surface soil sample adjacent to the former Area A radiological UST. As discussed in Section 5.0, this data is suspect due to analytical uncertainties in identifying and quantifying these radionuclides. To be conservative, this radionuclide was included in the risk assessments. It constitutes only a minimal risk as compared to total natural background levels for the United States (1x10⁻² to 3x10⁻³).

Elevated human health risks from Ra-226 (maximum cancer risk of 2x10⁴) were also associated with surface soils and one ditch sediment. Ra-226 is above established 1994 background levels at these locations. Ra-226 is ubiquitous at OU 1 and is considered to be part of natural background. At LAFB background levels, naturally occurring Ra-226 alone contributes a maximum cancer risk of 2x10⁻⁴. Significant reduction of risk attributable to radioactive isotopes is not possible due to the high levels of naturally occurring radioactive isotopes.

Analytical data for the surface water collected from the Area A drainage ditch were evaluated, and only copper was detected at concentrations in excess of the aquatic benchmarks. A review of the toxicological data for copper suggests that organisms that would likely use this ephemeral habitat (such as amphibians) would not be impacted at the concentration reported. The data and rationale for this conclusion are presented in the OU 1 RI Report (ABB-ES, 1995a). No impacts to plants growing in Area A surface soil or to other terrestrial receptors were identified in the ecological risk assessment.

Area A Groundwater. No human health radiological risks above regulatory target risk levels have been identified associated with potential residential groundwater exposures at Area A.

Background concentrations of inorganics in overburden and bedrock groundwater are currently being revised as part of the OU 12 basewide groundwater RI. Concentrations of inorganics in groundwater detected at OU 1 will be compared to the OU 12 background concentrations upon approval and acceptance of those levels. Groundwater inorganic data for OU 1 will be addressed in the OU 12 ROD.

Areas B through G Soils. Total maximum cancer risks associated with exposure to detected radionuclides in soil at levels above established background concentrations

range from $5x10^4$ to $2x10^{-5}$. These risks represent a minimal incremental cancer risk above the LAFB soil background risks of $2x10^4$ to $6x10^{-6}$, and are well below published total natural radiological background risks of $1x10^{-2}$ to $3x10^{-3}$ (Shleien, 1992).

The maximum radiological human health risk of $5x10^4$ is based on Ra-226 detected in surface, subsurface, and composite soil samples. As discussed in Subsection 5.1, Ra-226 is naturally occurring at OU 1. At LAFB off-site background levels, a cancer risk of $2x10^4$ is associated with naturally occurring Ra-226. The radiological human health risks at Areas B through G are considered acceptable because they are a result of naturally occurring Ra-226.

No non-radiological human health carcinogenic or non-carcinogenic risks above regulatory target risk levels were identified in surface soils at Areas B through G, except for a single surface soil sample at Area G (JSS-2680). The surface soil sample analysis indicated a non-carcinogenic risk due to inhalation of barium for both the forestry worker and construction worker scenarios. JSS-2680 was the only surface soil sample location out of 17 collected at OU 1 in which barium was detected above background levels.

No ecological radiological risks were indicated at Areas B through G. Ecological non-radiological risks at Areas B through F were indicated due to an elevated mercury result in one Area C surface soil sample. The mercury concentration suggested risk to the red fox, and exceeded the screening benchmark for terrestrial plants. Mercury was detected only once out of six surface soil samples at Areas B through F. Zinc exceeded screening benchmarks to terrestrial invertebrates and to plants due to one surface soil result at Area G.

Ecological non-radiological risk at Area G was calculated for zinc and mercury in surface soil for lethal effects to the robin and red fox, respectively. Concentrations of 2-methylnaphthalene, chromium, copper, and zinc also exceeded the screening toxicological benchmarks for terrestrial invertebrates. Concentrations of cadmium, chromium, copper, lead, mercury, vanadium, and zinc exceeded the screening benchmarks for terrestrial plants. Maximum concentrations of all risk-contributing ecological CPCs were detected at sample location JSS-2680, which is located at the head of the drainage ditch at Area G. Potential ecological impacts are likely to be spatially limited, and it is unlikely that mobile wildlife would be impacted.

Area B through G Groundwater. A total maximum radiological risk of 1x10⁻⁵ was identified for potential residential exposure to overburden groundwater. The risk does not exceed USEPA's target risk range or MEDEP's cancer risk guidance value. The site-specific risk level represents a minimal incremental cancer risk above the LAFB groundwater background risk level of 9x10⁻⁷ and is below published total natural radiological background risks of 1x10⁻² to 3x10⁻³ (Shleien, 1992).

Total maximum radiological risks of $4x10^{-5}$ to $4x10^{-6}$ were identified for potential residential exposure to bedrock groundwater. Groundwater samples from one well out of the four at Area G indicated radiological risk due to Ra-226. The site-specific Ra-226 concentration is only slightly above the LAFB background concentration and represents a minimal incremental cancer risk as compared to published total natural backgrounds risks.

Non-radiological Area G bedrock groundwater data were separated from Areas B through F during risk assessment because fuel oil USTs at Area G have influenced groundwater quality. Area G non-carcinogenic risks range from HIs of 0.06 to 7. Those above the target HI of 1 were attributable to arsenic, iron, and manganese. Bis(2-ethylhexyl)phthalate (BEHP) and arsenic were identified as the carcinogenic risk drivers from Area G groundwater with a maximum risk of $3x10^4$ and $2x10^4$. BEHP is a common laboratory contaminant, and not likely to be site-related.

Evaluation of Radionuclides and Inorganics Detected at OU 1. Two summary tables have been developed to present conclusions with respect to radionuclides and inorganics, Tables 6-8 and 6-9, respectively. These tables summarize the radionuclides and inorganics detected above background, the site areas where they were detected, and present discussion and conclusions. The purpose of these tables is to put into perspective the detections above background within OU 1.

TABLE 6 - 8 EVALUATION OF RADIONUCLIDES ABOVE BACKGROUND AT 0U1

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE BASE

MEDIUM Subsurface Soil	ANALYTICAL	СРС	AREAS	CONCTUSIONS
	METUDO			
	METHOD		DELECTED > BKG	
	ALPHA-SPEC	Thorum-228	C,D,E	Radium-226 was the primary radioisotope responsible for elevated radiological risks at 0U 1 Radium-226 was
₹	ALPHA-SPEC	Thorum-230	၁	detected at greater that the established packground value in sols at all areas of OO 1. Insulant-220 is one of the most abundant naturally occurring radionuclides on the OU 1 isotope list. Radium-226 was added to the
¥	ALPHA-SPEC	Thorum-231	C,E	analyte list based on the possibility that radium paint associated with aircraft instrumentation dials might have
4	ALPHA-SPEC	Thonum-232	A,C,E	been buried in the LLRWD trenches. No dials were reported during the trench removals. The background value
4	ALPHA-SPEC	Uranium-235	C,E	Tof hadiumy-20 was established in 1994 in 1995 background soll semples were also collected and alialyzed. But not used in establishing background values. The 1993 background samples indicated levels of Radium-226
9	GAMMA-SPEC	Radium-226	B,C,D,E,G	between 3 44 to 4 49 pC//g, which are higher than the maximum detected site-related Radium-226 result
9	GAMMA-SPEC	Radium-228	A,B,C,D,E,F	The widespread detection of Radium-226 at OU 1 is indicative of natural occurrence.
Surface Soil	GAMMA-SPEC	Americium-241	V	Amendanizer was detected in one when A surface son senior. The testin is rightly questionation as to identification due to analytical interference (Note: The laboratory reported bad peak shape).
	GAMMA-SPEC	Radium-226	A,E,G	in summary, subsequent to the LLRWD trench removals, soil results do not indicate a source of base-related radiological contamination. No human health radiological risks above requistory risk levels were identified for
				the LRWD trench and tank confirmatory soil samples. Target isotopes detected in soil are all naturally
				occurring with the exception of Americium-241 which is mentioned above
Surface water		NONE		There were no ecological or human health risks above regulatory thresholds indicated by radiological surface
	ALPHA-SPEC	Neptunium-237	0013	water or sedement results. The isotopes associated with the sedements are all naturally occurring with the accordance of Nicotuming 227. Nicotuming 1947 results obtained by resonance of the results and a farrance of
N. C.	GAMMA-SPEC	Neptunium-237	A,0U13	uncertainty due to analytical interferences. Neptumum-237 was detected in sediment background samples by
9	GAMMA-SPEC	Radium-226	A,0U13	both gamma-spectroscopy and alpha-spectroscopy. A positive detection of Neptunium-237 by alpha
9	GAMMA-SPEC	Thorum-234	A,0U13	spectroscopy was also identified at a background location. Based on this intomation, Neptunium-237 is believed to be a false positive caused by the analytical procedures
A	ALPHA-SPEC	Uranium-234	0U13	
9	GAMMA-SPEC	Uranium-235	A,0U13	
4	ALPHA-SPEC	Uranum-238	0U13	The state of the s
Groundwater	GROSS-ALPHA	Gross Alpha	C,D,E,G	Groundwater radiological results did not indicate risks of concern at OU 1, except for Radium-226 in one
9	GROSS-BETA	Gross Beta	B,C,D,E,F,G	Bariphe at Afea G. 1718 naghtiff-2.20 detection is beneved to be attributable to turbany in the groundwater Sample
4	ALPHA-SCAN/SPEC	Radium-226	g	
A	ALPHA-SPEC	Thorum-228	E	turbidity. This is supported by background data collected for both dissolved and total radioisotopes. Gross appearant on an assemble to the reference and inside the refere
4	ALPHA-SPEC	Thonum-230	A,B,D,E,G	arplic and gloss Lord baranters were first used in the last assessment, however, miner entres of these parameters exceeded primary drinking water standards (MCLs), further isotope-specific analysis was
4	ALPHA-SPEC	Thonum-232	C,E,G	performed. The isotope-specific analyses were performed to determine what impact, if any, base-related
L	TRITIUM	Tritium	A,C,E,G	activities may have had on local groundwater quality. A risk assessment was performed using the target
4	ALPHA-SPEC/SCAN	Uranium-234	A,B,C,D,E,G	isotope-specific data In summary, groundwater radiological data do not indicate a base-related source of contamination All isotopes
4	ALPHA-SCAN/SPEC	Uranium-235	B,C,D,G	detected occur naturally at varying levels
	ALPHA-SPEC/SCAN	Uranium-238	B,D,E,G	
NOTES:				
MICE B WAXANGE CONTENTION LOVO	Jungan Can			
ALPHA-SPEC = Apha Spectroscopy	reposes or one			
GAMMA-SPEC = Gamma Spectroscopy				
CPC = Compound of Potential Concern				
> BKG = Greater than extablished background values	ound values			

TABLE 6 - 9 EVALUATION OF INORGANICS DETECTED ABOVE BACKGROUND AT OU 1

OPERABLE UNIT 1 RECORD OF DECISION LORING AIR FORCE RASE

			LORING AIR FORCE BASE	INCE BASE
MEDIUM	ANALYTICAL	CPC	AREAS	CONCENSIONS
	METHOD		DETECTED > BKG	
Subsurface Soil	CLP TAL-INOR	Arsenic	E,F	The inorganic analytes in soil contributing to the elevated risks at OU 1 are primarily arsenic, barturn, mercury,
100		Lead	ш	and zinc. Arsanic was detected at greater than the background value in only 2 out of 75 soil samples. The
Surface Soil	CLP TAL-INOR	Barium	G	maximum arsenic detection was in 1 out of 18 closely gridded (equally spaced). Trench E confirmatory soil
		Cadmium	9	oanipos - Danuni was ustected at greater tran the background concentration in only 1 out of 75 samples. The sole elevated banum detection was located at the head of the drainage dirch at Area G. Mercury was
		Сһгопнит	9	detected at greater than the background value in 2 out of 75 soll samples. The maximum mercury result is
		Cobatt	9	also located at the head of the dramage ditch at Area G A second Area G dramage ditch sample collected
		Copper	9	approximately 30 feet away (downslope) did not contain banum or mercury greater than background. Zinc
		Lead	A.E.F.G	Was detected at greater than the background concentration in 9 out of 75 soil samples. These sporadic
7.77		Mercury	0,0	Unificially of Histograms are that makening of source areas
		Silver	п	
		Vanadium	9	
		Zino	A,F,G	
Surface Water(Area A only)	CLP TAL-INOR	Copper	4	The inorganic analytes contributing to elevated ecological risks for surface water and sediment at Area A were
Sediment[Area A only]	CLP TAL-INOR	Barium	A	primarily copper and zinc. One surface water and three sediment samples were collected in the drainage ditch
The state of the s		Copper		at Area A Copper in both surface water and sediment produced elevated ecological risks. Zinc in the
		Iron		equinous was also responsible for an increased ecological risk. A review of the textcological data for copper-
		Lead		exceeded the screening benchmark associated with plant receptors. However, the screening benchmarks used
		Manganese	A	in the ecological risk assessment (ERA) were primarily below established LAFB background concentrations, and
		Mercury	A	corresponding conservative assumptions used in the ERA suggest that impacts to wildlife are unlikely
		Nickei	A	l hese inorganic analyte concentrations are attributeble to overfand runoff and accumulation
		Vanadum	A	
		Zinc	٨	
Groundwater	CLP TAL-INOR	Arsenic	A,B,C,D,E,G	The inorganic analytes contributing to elevated risks for groundwater at OU 1 were arsenic, iron, manganese,
		Banum	A,B,C,D,E,G	chromium, and beryllium. Each of these elements was likely detected at greater than background values due
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		Berylkum	B,C,E	to the impacts of turbidity. These inorganic analytes are naturally occurring in the soil and can cause elevated arctional water concentrations when turbididy is present in comples. This is amounted by the household
		Chromium	A.B.C.D,E,F	grounwater concentrations when turning is present in eartiples. This is supported by the background overburden groundwater samples collected under 0U 12. Iron and manganese are responsible for up to 90%.
		Iron	A,B,C,D,E,F,G	manganese have promulgated drinking water
		Lead	A,B,C,D,E,F,G	standards (MCLs)
	,	Manganese	A,B,C,D,E,F,G	Arsenic and betylittim are responsible for up to 100% of the carcinogenic risks from groundwater. Arsenic is unimitiate in arctinguistic strange of the Center of Manager and the comments of the decaded and the comments of
		Nickel	A,B,C,D,E,G	conjuisces in groundwater unoughout the state of mains size is extranonly detected groundwater element. Absence concentrations found in OU 1 groundwater were all wall below the MCI of 50urs?
		Vanadium	B,c	Beryllum was detected in 4 groundwater samples out of 40 collected. All the detections were below or at the
				CRDL of 10 ug/l. All detections of beryllum were below the MCL of 40 ug/l.
				In summary, these detections of inorganics do not indicate any base-related inorganic source areas at OU 1
NOTES:				
CPC = Compound of Potential Concern	Concern			
>BKG = Greater than established background values	ed background values			
= Background values are for bedrock groundwater only	sedrock groundwater only	Overburden backgr	ound concentrations have	Overburden background concentrations have not been established to date for LAFB.
Ct P TAI-INOR = Contract 1ab Program Target Apalyta 1st of Ingressive	entrations listed may be to Program Target Apalyte I	om overburden grou et of homenoore	indwater samples and here	Some of the maximum concentrations listed may be from overburden groundwater samples and hence a proper comparison to background is not possible TAI-INOR = Contract I ab Program Pariat Analyta List of Increases
MCL = Maximum Contaminant Level	Level			The state of the s
CRDL = Contract Required Detection Limit	ction Limit			
ERA = Ecological Risk Assessment	9nt			
LAFB = Loring Air Force Base				
LIGHT - Mineragement non like				

ug/L = Micrograms per liter

7.0 DESCRIPTION OF THE NO ACTION ALTERNATIVE

Sampling conducted after the removal actions were completed at the LLRWDS confirmed that no significant radiological or non-radiological contamination above background concentrations remained at the former UST or disposal trench locations. Analysis of groundwater sampled from monitoring wells installed downgradient of the USTs and disposal trenches did not consistently detect contamination above MCLs or MEGs, other than that attributable to background variation or sample turbidity.

In accordance with USEPA guidance, additional monitoring and five-year reviews are not necessary for sites where no hazardous substances, pollutants, or contaminants remain at levels that would necessitate restricted use or access (USEPA, 1991). Because the USTs and waste disposal trenches were removed during the removal action and results of the RI indicate no substantial contamination remains on-site, additional monitoring and five-year reviews will not be conducted.

Based on these results, and the baseline risk assessment, no further remedial action under CERCLA is considered necessary for OU 1 at LAFB. Areas A through F of OU 1 will be removed from the IRP. Area G will also be removed from the IRP and be redesignated as a non-CERCLA site that will be managed in accordance with the Maine UST regulations.

Remediation of the contaminated soil and groundwater associated with the former fuel oil UST and abandoned pipeline is best addressed as a non-CERCLA action conducted under Maine UST regulations. The authority of CERCLA is limited to the hazardous substances defined in Section 101(14) of the law. Under both Sections 101 and 104 of CERCLA, petroleum products are excluded from regulation under CERCLA. Remediation of the contaminated soil and groundwater associated with the former fuel oil UST and abandoned pipeline will be addressed as a non-CERCLA action conducted under the Maine UST regulations.

Section 12 of the Maine UST regulations (06-096 CMR 691) outlines requirements for leak investigation, response, and corrective action. Many of the requirements for response and investigation have been met during the course of replacing the Building 216 USTs and conducting the RI. Further response at Area G, in accordance with Section 12 requirements, potentially includes soil remediation, groundwater treatment, and monitoring.

SECTION 7

If during the course of the UST remedial response, CERCLA-regulated wastes are identified at concentrations that pose risk to human health or the environment, Area G of OU 1 will be managed under the IRP and CERCLA.

8.0 DOCUMENTATION OF NO SIGNIFICANT CHANGES

The USAF prepared a Proposed Plan for OU 1 (ABB-ES, 1995b). The Proposed Plan describes the USAF's recommendation to pursue no further action under CERCLA at OU 1. There have been no significant changes made to the No Action under CERCLA decision stated in the Proposed Plan.

9.0 STATE ROLE

MEDEP, on behalf of the State of Maine, reviewed the RI Report and Proposed Plan and indicated its support for the selected remedy. MEDEP concurs with the selected remedy for OU 1. A copy of the declaration of concurrence is included in Appendix C.

GLOSSARY OF ACRONYMS AND ABBREVIATIONS

ABB-ES ABB Environmental Services, Inc.

Am Americium

BEHP bis(2-ethylhexyl)phthalate

CERCLA Comprehensive Environmental Restoration, Compensation, and

Liability Act

CPC contaminants of potential concern

CRP Community Relations Plan

DOD Department of Defense

FFA Federal Facilities Agreement

HI hazard index

HQ hazard quotient

IRP Installation Restoration Program

LAFB Loring Air Force Base

LLRWDS Low Level Radioactive Waste Disposal Sites

MCL Maximum Contaminant Levels

MEDEP Maine Department of Environmental Protection

MEG Maximum Exposure Guidelines

NCP National Contingency Plan

Np Neptunium

NPL National Priorities List

OU operable unit

Ogden Environmental and Energy Services, Inc.

Pa Protactinium

PA Preliminary Assessment
PAH polyaromatic hydrocarbons
PCB polychlorinated biphenyls

Ra Radium

GLOSSARY OF ACRONYMS AND ABBREVIATIONS

RAB Restoration Advisory Board

RfD reference dose

RI Remedial Investigation

RME reasonable maximum exposure

ROD Record of Decision

SI Site Inspection

SVOC semivolatile organic compounds

Th Thorium

TCE trichloroethene

U Uranium

USAF U.S. Air Force

USEPA U.S. Environmental Protection Agency

UST underground storage tank

VOC volatile organic compounds

WSA weapons storage area

- ABB Environmental Services, Inc. (ABB-ES), 1995a. "Operable Unit (OU 1) Remedial Investigation Report"; Installation Restoration Program; prepared for HAZWRAP; Portland, Maine; April 1995.
- ABB Environmental Services, Inc. (ABB-ES), 1995b. "Operable Unit (OU 1) Proposed Plan"; Installation Restoration Program; prepared for HAZWRAP; Portland, Maine; July 1995.
- CH₂M Hill, 1984. "Records Search Report"; Installation Restoration Program; prepared for HAZWRAP; Limestone, Maine, January 1984.
- Federal Facility Agreement (FFA), 1991. Under CERCLA Section 120, the Matter of Loring Air Force Base by U.S. Environmental Protection Agency Region I, State of Maine, and the U.S. Department of the Air Force, January 3, 1991.
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- U.S. Environmental Protection Agency (USEPA), 1990. "National Oil and Hazardous Substances Pollution Contingency Plan (National Contingency Plan)"; Code of Federal Regulations, Title 40, Part 300; Federal Register, Volume 55, Number 46, pp. 8666 et seq.; March 8, 1990.
- U.S. Environmental Protection Agency (USEPA), 1991. "Structure and Components of Five-Year Reviews"; OSWER Directive 9355.7-02; Office of Solid Waste and Emergency Response, Washington, DC; May 23, 1991.

W0049530.080 7656-16

TRANSCRIPT OF THE PUBLIC MEETING (AUGUST 2, 1995)

W0049530.080 7656-16

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STATE OF MAINE

AROOSTOOK, ss.

LORING AIR FORCE BASE OPERABLE UNIT 1

CARY MEDICAL CENTER
VAN BUREN ROAD
CARIBOU, MAINE
8:03 P.M.

Philip R. Bennett, Jr., Court Reporter 13 Vaughn Street Caribou, Maine 04736 (207)498-2729

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PETER FORBES

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LORING AIR FORCE BASE, OPERABLE UNIT # 1

3.

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August 2, 1995

PETER FORBES: Good

evening. Welcome to the public hearing to receive comments on the proposed plan for Operable Unit 1 at Loring Air Force Base, the Low Level Radioactive Waste Disposal Sites.

Today's date is August 2nd, 1995. My name is Peter Forbes, the Remedial Project Manager for the Installation Restoration Program at Loring. And seated with me are Michael Nalipinski of the U.S. Environmental Protection Agency and Naji Akladiss of the Maine Department of Environmental Protection. They will assist me in receiving your comments tonight.

This hearing is being held in accordance with the provisions of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA), as amended in 1986, also known as Superfund. The act requires federal facilities on the National Priorities List to present clean up proposals to the local community for comment and consideration before the final clean up decisions are made. The purpose of this hearing is to receive comments on the Proposed Plan for Operable Unite 1.

Mr. Phil Bennett from Aroostook Legal Reporters will

LORING AIR FORCE BASE, OPERABLE UNIT # 1

3.

serve as the court reporter tonight, preparing a verbatim record of the proceedings. The verbatim record will become a part of the final clean up plan. The court reporter will be able to make a complete record only if he is able to hear and understand what you say. With that in mind, please follow a few ground rules. Speak only after I recognize you and please address your remarks to me. State your name and the organization you represent and present your statement. Please do not state your address or any other personal information which you do not want to become a matter of the public record. Do not begin speaking until you have reached the podium. Speak slowly and clearly into the microphone. If you have prepared a statement beforehand, you may read it aloud or you may describe it and place it on this table.

Now are there any individuals who would like to make a comment or question or statement at this time?

Okay. Well, ladies and gentlemen, it's 8:05 p.m.,

August 2nd, 1995. I declare the public hearing to receive

comments on the Proposed Plan for Operable Unit 1 at Loring

Air Force Base closed. Thank you for coming.

- END OF HEARING

CERTIFICATION

I HEREBY CERTIFY THAT the foregoing is a true

and correct transcript of the record of proceedings held

on the aforementioned date.

3.

 STATE OF MAINE

Philip R. Bennett, J

Court Reporter

AROOSTOOK, ss.

RESPONSIVENESS SUMMARY

W0049530.080 -- - - 7656-16

FINAL

Loring Air Force Base

OU 1 RESPONSIVENESS SUMMARY

AUGUST 1995

Prepared for:

Air Force Base Conversion Agency Limestone, Maine (207) 328-7109

Prepared by:

Service Center: Hazardous Waste Remedial Actions Program
Oak Ridge, Tennessee

Contractor: ABB Environmental Services, Inc. Portland, Maine

Project No. 7656-16

OU 1 RESPONSIVENESS SUMMARY LORING AIR FORCE BASE

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The U.S. Air Force (USAF) held a 30-day comment period from July 17 to August 16, 1995, to provide an opportunity for the public to comment on the Proposed Plan and other documents developed for Operable Unit No. 1 (OU 1) at Loring Air Force Base, Maine. The Proposed Plan is the document that identifies remedial action objectives, evaluates remedial alternatives, and recommends the alternative that best meets the evaluation criteria for OU 1. The USAF made preliminary recommendations of its preferred alternative for remedial action at OU 1 in Section 6.0 of the Proposed Plan, which was issued on July 17, 1995. All documents on which the preferred alternative was based were placed in the administrative record for review. The administrative record is a collection of the documents considered by the USAF while choosing the remedial action for OU 1. It is available to the public at the following location:

Air Force Base Conversion Agency 5100 Texas Road Limestone, ME 04751 (207) 328-7109

The purpose of this Responsiveness Summary is to document USAF responses to the questions and comments raised during the public comment period regarding the proposed OU 1 preferred alternative. The USAF considered all comments in this document before finalizing the preferred remedy for OU 1.

This Responsiveness Summary is organized into the following sections:

- 1.0 Overview of the Preferred Alternative. This section briefly outlines the preferred alternative presented in the Proposed Plan for OU 1.
- 2.0 Background on Community Involvement and Concerns. This section provides a brief history of community interest in OU 1 and concerns regarding these areas.
- 3.0 Summary of Comments Received During the Public Comment Period and USAF Responses. This section summarizes and provides the USAF's responses to all written and oral comments received from the public during the public comment period.

1.0 OVERVIEW OF THE PREFERRED ALTERNATIVE

The following paragraphs outline the preferred alternative presented in the Proposed Plan OU 1.

Based on the results of the RI, no further remedial action under CERCLA is considered necessary for OU 1 at LAFB.

Areas A through F: In 1994, removal actions were conducted for the five radiological USTs and the contents of the former waste disposal trenches. Completion of these removal actions has eliminated the need for any further remedial action at Areas A through F.

Area G: The contamination detected at Area G is primarily attributed to a former leaking UST and possibly the fuel oil pipeline. The tanks were replaced and the pipe was abandoned. Because the release involved only petroleum product, the USAF will address the petroleum contamination as a non-CERCLA action under the Maine UST regulations.

Section 12 of the Maine UST regulations (06-096 CMR 691) outlines requirements for leak investigation, response, and corrective action. Many of the requirements for response and investigation have been met during the course of replacing the Building 216 USTs and conducting the RI. Further response at Area G, in accordance with Section 12 requirements, potentially includes soil remediation, groundwater treatment, and monitoring.

If during the course of the UST remedial response, CERCLA-regulated wastes are identified at concentrations that pose risk to human health or the environment, Area G of OU 1 will be managed under the IRP and CERCLA.

2.0 BACKGROUND ON COMMUNITY INVOLVEMENT AND CONCERNS

Throughout LAFB's history, the community has been involved in base activities. The USAF, USEPA, and MEDEP have kept the community and other interested parties apprised of LAFB IRP activities through informational meetings, fact sheets, press releases, public meetings, site tours, and open houses.

In addition to these activities, during the course of IRP activities at LAFB, there have been regular meetings of the Restoration Advisory Board (RAB) (formerly the Technical Review Committee). The RAB, chaired by the USAF and a representative of the community, is composed of representatives of the USEPA, MEDEP, the community, and local officials. The purpose of the RAB meetings has been to ensure clear communication with the public, timely transfer of information, and opportunity for public comment.

A Federal Facilities Agreement (FFA) between USEPA Region I, MEDEP, and USAF, signed January 30, 1991, governs environmental activities being conducted at LAFB. The FFA provides the framework for addressing environmental effects associated with past and present activities so that appropriate investigations and remedial actions are implemented to protect human health, welfare, and the environment. Since the signing of this agreement, LAFB was placed on Congress' Base Closure List and closed in September 1994. The FFA was amended in December 1993 to address base closure-related issues such as transfer of real property. The FFA was further modified in January 1995 to allow Remedial Project Managers to make minor modifications to the FFA, such as schedule adjustments or removal of petroleum-contaminated sites from the agreement.

The framework for the USAF's approach to community involvement is the LAFB Community Relations Plan (CRP), which was released in August 1991 and subsequently revised in May 1995. The CRP outlines the USAF's program for addressing community concerns and keeping citizen informed and involved during remedial activities. To ensure the public was informed about the IRP program, the USAF held three public information meetings in the towns of Limestone, Caribou, and Fort Fairfield in February and March, 1993. The purpose of the meetings was to introduce the IRP program and respond to any questions from the public.

Documentation of the reports, memoranda, and correspondence that are the basis for IRP remedial response decisions are kept in an Administrative Record. The

Administrative Record is open and available for public review at the Air Force Base Conversion Agency Office, 5100 Texas Road, Limestone, Maine.

The following is a summary of the activities the USAF has undertaken to keep the public informed and involved regarding the remedial response at OU 1.

- On June 2, 1994, a RAB meeting was held to discuss the results of the OU 1 investigations and the approach for conducting the UST and radioactive waste disposal trench removal action.
- An IRP Fact Sheet, explaining activities planned for OU 1, was issued in July 1994.
- The USAF published a notice and brief discussion of the proposed removal action in the Aroostook Republican on July 6, 1994 and the Bangor Daily News on July 7, 1994.
- From July 11 through August 10, 1994, the USAF held a 30-day public comment period to accept public input on the Action Memorandum outlining the proposed removal action, and on any other OU 1 documents in the Administrative Record. On July 28, 1994, USAF personnel and regulatory representatives held a public meeting to discuss the Action Memorandum and to accept oral comments.
- During the removal action, the USAF invited the local press to cover the trench removal activities. Information regarding both the trench and UST tank removals was made available to representatives of local media.
- The USAF published a notice and brief analysis of the Proposed Plan in the Bangor Daily News, Aroostook Republican, Fort Fairfield Review, and Presque Isle Maine Star-Herald on July 12, 1995, recommending No Action under CERCLA as the preferred alternative for OU 1.
- On July 17, 1995, the Proposed Plan for OU 1 was made available for public review at the Air Force Base Conversion Agency Office, 5100 Texas Road, Limestone, Maine.
- From July 17 through August 16, 1995, the USAF held a 30-day public comment period to accept public input on the recommendations in the

RI/Baseline Risk Assessment and the No Action preferred alternative presented in the Proposed Plan, and on any other documents included in the Administrative Record. On August 2, 1995, USAF personnel and regulatory representatives held a public meeting and hearing to discuss the OU 1 RI and Proposed Plan. During the public meeting, the USAF answered questions informally from the public. Immediately following the public meeting, a public hearing was held to accept oral comments. Based on the public comments, the public is in agreement regarding the preferred alternative for OU 1 as presented in the Proposed Plan.

3.0 SUMMARY OF COMMENTS RECEIVED DURING THE PUBLIC COMMENT PERIOD AND USAF RESPONSES

This Responsiveness Summary addresses comments received by the USAF and USEPA during the public comment period from July 17 to August 16, 1995 relative to the Proposed Plan for OU 1. The only comments received were those received in writing from a RAB member. The comments and corresponding responses are included herein.

1. <u>Comment</u>: The commenter asked what was the purpose of the five (5) radiological USTs attached to weapon maintenance facilities.

Response: The purpose of the five radiological USTs was to receive and contain potentially radioactive liquids in the event of a release in one of the buildings. Further information can be obtained from the OU 1 Remedial Investigation Report which is part of the Administrative Record.

2. <u>Comment:</u> The commenter asked what radioactive isotopes were to be transported to these radiological USTs.

Response: The radiological USTs at Areas A and F supported Buildings 365 and 232, respectively. Strategic weapons components were reportedly installed and inspected within these buildings, with the UST backups in the event of a release of radioactive materials. A radioactive release from these buildings could have potentially been composed of enriched uranium, plutonium, americium, or tritium. There were no documented releases to these tanks, which is supported by the analysis of the tank liquids, sediments, and scrape samples. Further information can be obtained from the OU 1 Remedial Investigation Report which is part of the Administrative Record.

The remaining three radiological USTs at Areas B, C, and D supported the "short igloos" where the tritium containers were stored. The "short igloos" contained floor drains which were connected to the USTs to receive washdown liquids in the event of a tritium release. There were no documented releases to these radiological USTs, which is supported by the analysis of the tank liquids.

3. <u>Comment</u>: The commenter asked if there are no documents showing release of any radioactive isotopes into these radiological USTs, why were these tanks tested.

Response: The tanks were sampled because they did contain liquid and documentation on the origin of the liquid could not be located. To confirm that the tanks did not contain chemical or radioactive contaminants, liquid, sediments, interior scrape samples, and soil samples from beneath the tanks were collected and analyzed for the target radioisotopes for all five USTs prior to their removal in 1994. Further information can be obtained from the OU 1 Remedial Investigation Report which is part of the Administrative Record.

4. <u>Comment:</u> The commenter asked if any radioactive isotopes had been found in the UST, would it have been necessary to have disposed of these at the Repository in Utah.

<u>Response</u>: Depending on the levels and radioisotopes found, it might have been necessary to have disposed of these USTs in Utah. However, based on the lack of contamination in the tanks, they were simply disposed of as scrap metal.

5. <u>Comment</u>: The commenter asked why tritium is found all over the Loring WSA if tritium is a very light gas and, when released either by accident or purposeful venting, should have risen into the Stratosphere and Ionosphere.

Response: Tritium is found in background due to atmospheric weapons testing in the 1960s, more recently from nuclear power plant releases, and naturally occurring interactions with cosmic rays and gases in the upper atmosphere. The tritium detections in the University of Maine and ABB-ES analyses indicated levels of tritium at the Weapons Storage Area (WSA) which are consistent with background levels. Further information can be obtained from the OU 1 Remedial Investigation Report which is part of the Administrative Record.

6. <u>Comment</u>: The commenter asked why are the areas of tritium concentration at the WSA not related to the weapon maintenance facilities.

Response: As discussed, the tritium detected at the WSA is at background levels with normal local variation. There are no significant areas of "tritium concentration" at the WSA.

7. <u>Comment</u>: The commenter asked why tritium radiation background was not established at Loring, since a great deal of effort was made to establish background radiation of certain isotopes around the Loring WSA.

Response: Tritium background was not established due to the low levels detected and because of tritium's relatively low health risks. Tritium detections from within the WSA were what would be expected in background. Detections of tritium in groundwater and surface water were all less than USEPA's drinking water standard for tritium.

8. <u>Comment</u>: The commenter asked whether the southern area was mentioned in the plan, with reference to tritium, around the Nuclear Power Plant, at Wiscasset.

Response: No reference to the "southern area" was made in the Proposed Plan. However, in the University of Maine report, there is a discussion of samples collected from Southern Maine. In 1972, tritium analyses were performed around the "then being constructed" nuclear power plant at Wiscasset (which is in Southern Maine). The data were collected prior to the power plant receiving any nuclear fuel to establish a baseline against which future monitoring data could be compared.

9. <u>Comment</u>: The commenter asked why tritium would be defined as a contaminant at Area D, and, when found at other areas, not be acknowledged as a contaminant.

Response: Tritium is acknowledged as a potential contaminant at Areas B, C and Area D, based on known site history.

10. <u>Comment:</u> The commenter asked why there is such a reluctance to acknowledge tritium as a radioactive substance throughout this whole plan.

Response: It was certainly not the intent of the Air Force to appear reluctant to address tritium. Tritium has been carefully addressed throughout the RI process by the USAF, the University of Maine, the MEDEP, and USEPA.

Tritium was identified as one of the WSA's target radioactive isotopes and therefore was included in analyses of OU 1 environmental samples. There is no detailed discussion of tritium, in particular, because (1) the purpose of the Proposed Plan is to present the Air Force's preferred alternative and a general overview of the IRP activities conducted to date, and (2) the results of the radiological investigation did not identify tritium at other than naturally occurring levels.

- 11. <u>Comment</u>: The commenter asked whether the following is a correct paraphrase of the last paragraph on Pages 4-5 and 4-6:
 - (1) Background radiation at Loring and its Weapon Storage Area (WSA) may pose a natural health risk.
 - (2) Background radiation at Loring and its WSA is lower than other areas throughout the United States.
 - (3) That even though the WSA at Loring is contaminated with weapons-grade radioactive isotopes, tritium, the human health risk here due to radiation is still lower than risk typically associated with naturally occurring radiation throughout the United States.

Response: There are several inaccuracies in this interpretation of the referenced paragraph. To clarify, risk calculations were performed using (1) concentrations of naturally occurring radiation throughout the United States, (2) background concentrations of radioactive isotopes established for Loring, and (3) concentrations of radioactive isotopes detected at the WSA. The risks associated with background radiation at Loring and at the WSA were lower than risks associated with published naturally occurring levels of radiation throughout the U.S. Further information can be obtained from the OU 1 Remedial Investigation Report which is part of the Administrative Record.

These comparisons were made to illustrate that while the human health risks calculated for the radioactive isotopes at the WSA are higher than the USEPA target risk range (1x10⁻⁴ to 1x10⁻⁶), naturally occurring radiation also has a risk higher than the USEPA target risk level. Following the trench removal action, the risks associated with radioactivity at the WSA are consistent with naturally occurring radiation.

The statement that "Loring is contaminated with weapons-grade radioactive isotope, tritium", is somewhat misleading. Tritium is tritium, whether it is included in a weapon or a result of natural reactions in the atmosphere, and the levels of tritium detected are consistent with background levels in Maine.

LETTERS OF CONCURRENCE (TO BE INCLUDED IN ROD FOR SIGNATURE)

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STATE OF MAINE

DEPARTMENT OF ENVIRONMENTAL PROTECTION

ANGUS S. KING, JR. GOVERNOR

EDWARD O. SULLIVAN COMMISSIONER

August 16, 1995

Mr. Peter Forbes Air Force Base Conversion Agency Operation Location "M" RR # 1 Box 1719 Limestone, Maine 04750

RE: Loring Air Force Base Superfund Site, Maine

Dear Mr. Forbes:

The Maine Department of Environmental Protection (MEDEP) has reviewed the May 1995 Draft Record of Decision (ROD) regarding Operable Unit 1 (OU1) for the Loring Air Force Base Superfund Site located in Limestone, Maine.

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Based on that draft, the MEDEP concurs with the Air Force's determination that no action under CERCLA is necessary to address the contamination at OU1. The MEDEP also concurs with the following recommendations:

- That Areas A through F of OU1 be removed from the U.S. Air Force's CERCLA 1. response under Installation Restoration Program.
- That Area G be redesignated a Non-CERCLA site to be managed in accordance with the 2. State of Maine regulations for underground storage tanks.

Clean Up Levels

The remedial alternative selected for the site must achieve goals for reducing contamination at OU1. Clean-up goals for Area G have been set for contaminated soil, sediment, and groundwater based either on background concentration, analytical detection limits, or on risk calculation.

Compounds and elements for which remedial goals have been set are listed in Table 10-1 through 10-6 of this ROD.

Description of No Action Alternative

The following paragraph describes the no action remedial alternative developed for Operable Unit 1 at Loring:

Serving Maine People & Protecting Their Environment

Sampling conducted after the response actions were completed at the LLRWDS, Areas A through F of OU1, confirmed that essentially no radiological or non-radiological contamination, above background concentrations, remained at the former UST or disposal trench locations. Analysis of groundwater sampled from monitoring wells installed downgradient of the USTs and disposal trenches did not consistently detect radiological or non-radiological contamination above MCLs or MEGs, other than that attributable to background variation or sample turbidity.

Based on these results, no further remedial action under CERCLA is considered necessary for OU1 at LAFB and no further remedial action under State law is considered necessary for Areas A through F of OU1. Sampling has shown fuel-related contamination of soils and groundwater at Area G. It is, therefore, recommended that Areas A through F of OU1 be removed from the IRP for closure of federal facilities. It is further recommended that Area G also be removed from the IRP and be redesignated as a non-CERCLA site that will be remediated in accordance with the Maine UST Regulations. Because no significant residual contamination, attributable to the LLRWDS, remains on site, additional monitoring and five-year reviews are not recommended.

The State's concurrence in the selected remedy, as described above, should not be construed as the State's concurrence with any conclusions of law or findings of fact which may be set forth in the Record of Decision (for OU1). The State reserves any and all rights to challenge any such finding of fact or conclusion of law in any other context.

This concurrence is based upon the State's understanding that the MEDEP will continue to participate in the Federal Facilities Agreement and in the review and approval of operational, design and monitoring plans.

The MEDEP looks forward to working with the Department of the Air Force and the USEPA to resolve the environmental problems posed by this site. If you need additional information, do not hesitate to contact myself or members of my staff.

Sincerely,

Edward O. Sullivan, Commissioner

Department of Environmental Protection

pc: Mark Hyland, MEDEP

Mike Nalipinski, EPA Hank Lowman, BCA

COMSUPER/dlb

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