

INSTALLATION RESTORATION PROGRAM

STAGE 1

KING SALMON, ALASKA 99613

PREPARED BY

CH2M HILL 2550 DENALI STREET — 8TH FLOOR ANCHORAGE, ALASKA



AUGUST 29. 1989

FINAL TECHNICAL REPORT

PREPARED FOR UNITED STATES AIR FORCE ALASKAN AIR COMMAND 5099TH CEOS/CC ELMENDORF AIR FORCE BASE, ALASKA 99506

OCCUPATIONAL AND ENVIRONMENTAL HEALTH LABORATORY (AFOEHL/TS) BROOKS AIR FORCE BASE, TEXAS 78235-5501

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PREFACE

This Final Technical Report for the Installation Restoration Program (IRP) Stage 1 for King Salmon AFS has been prepared under Contract F33615-85-D-4535 by CH2M HILL. This report presents the result of the Phase II/Phase IV, Stage I investigation. Contractor personnel key to the production of this report include Jacques Gusmano, John Martinsen, Paul McBeth, Scott McKinley, and Dixie Simon-Gilles, among others. CH2M HILL acknowledges the contributions of NORTEC, Tracer Research Corporation, and our drilling subcontractors in obtaining specific field data. An extensive appendix, bound separately, includes field data, forms, related reports, and biographies. James F. Williams served as AFOEHL Technical Program Manager.

ERRATA

"King Salmon Air Force Station," "King Salmon AFS," and "station" are names used inaccurately throughout the document. The correct designation should be "King Salmon Airport" for the U.S. Air Force facilities located on a state airport near King Salmon, Alaska.

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EXECUTIVE SUMMARY

PROGRAM HISTORY

The Air Force Installation Restoration Program (IRP) was developed to 1) assess past hazardous waste disposal sites and spill sites at Air Force installations and 2) develop remedial actions consistent with the National Contingency Plan (NCP) for those sites that pose a hazard to human health and welfare or the environment.

The Air Force IRP consists of four phases:

- Phase I--Records Search and Installation Assessment
- o Phase II--Confirmation and Quantification
- o Phase III--Technology Base Development
- o Phase IV--Operation Remedial Actions

The Air Force has combined Phase II and Phase IV to allow the remedial investigation and feasibility study to be conducted in parallel. This report presents the result of the Phase II/Phase IV, Stage 1 investigation conducted by CH2M HILL at the King Salmon Air Force Station (AFS) in King Salmon, Alaska.

SUMMARY OF FINDINGS

Four investigation areas at King Salmon AFS were tested for over 146 inorganic, volatile organic, semivolatile organic, and pesticide contaminants. Of these 146, 31 potential contaminants of concern have been identified. These contaminants are addressed in this report because they exceed federal or state criteria for the protection of human health and the environment, are consistently present in site media (soil, sediment, surface, and groundwaters), and are potentially toxic to humans and the environment.

LOCATION OF INSTALLATION AND INVESTIGATED SITES

King Salmon AFS is located approximately 340 miles southwest of Anchorage, Alaska, on the northwest corner of the Alaska Peninsula.

Past hazardous waste disposal and spill sites have been prioritized and grouped into four areas for the purposes of the Phase II/Phase IV, Stage 1 investigation. The location and identification of each area is shown in Figure 1. A brief description of each is presented in the following subsections.

Eskimo Creek Seep

The Eskimo Creek seep is a petroleum oils and/or lubricants (POL) seep emanating from the toe of a small bluff east of Building 603 along the west bank of Eskimo Creek. The primary source of the seep is unidentified but may be underground POL delivery systems spill sites, underground storage tanks, or POL tanks 11, 12, 13, or 14. Any of the foregoing potential sources could also be minor or secondary sources.

Naknek River Seep

The Naknek River seep is located along the north bank of the Naknek River near the deactivated POL tank farm. The likely source of this seep is soils contaminated by spills and releases from the tank farm, and potentially other unidentified tanks and delivery systems.

North Barrel Bluff

This site is outside the King Salmon AFS installation boundaries. The area has been used in the past to dispose of drums, metal, and wood debris. The most visible type of debris is 55-gallon steel drums. Estimates of the number of barrels at the site range from hundreds of thousands to one million. This debris is visible along the face of the bluff for approximately 2300 feet. Residual liquids within the drums at the time of disposal are a likely source of continuing releases to soil, surface, and groundwater. The contents of the drums have not been fully characterized. However, many of the drum labels indicate fuel oil or other organic products. Because this area was used as a dump, potentially any substance delivered to the base in drums could be present.

South Barrel Bluff

South Barrel Bluff is located south of North Barrel Bluff along the west bank of King Salmon Creek. South Barrel Bluff also includes part of a former landfill. This area has been used to dispose of drums, metal, and wood debris. The most visible debris at South Barrel Bluff is 55-gallon steel drums. The drums and other types of debris are visible along the face of the bluff for approximately 1,000 feet. Contents of the drums would be expected to be similar to those in the North Barrel Bluff.

TIME FRAME FOR FIELD INVESTIGATION ACTIVITIES

Field tasks completed as part of the Stage I investigation and their duration are shown in Figure 2.





| TASK | DESCRIPTION | 1 987 | 1988 JIFIMIAIMIJIJIAIS IOINID | |
|------|--|------------------|--|---|
| 1. | WORK PLAN AND QAPP PREPARATION | | | , |
| 2. | LITERATURE SEARCH | | | 1 |
| 3. | SOIL GAS SURVEY | | | ł |
| 4. | GEOPHYSICAL SURVEY | | | |
| 5. | MONITOR WELL INSTALLATION AND DEVELOPMENT | | | |
| 6. | SEDIMENT AND SURFACE WATER SAMPLING | | | |
| 7. | MONITOR WELL SLUG TESTS AND PUMP TESTS | | • | • |
| 8. | GROUNDWATER SAMPLING | | | ı |
| 9. | LABORATORY ANALYSIS (SOILS, SEDIMENT, SURFACE WATER AND GROUNDWATER) | | + | |
| 10. | STAGE I DATA REPORT | | | |
| | | JIAIS OINID | JIFIMIAIMIJIJIAIS IO IN ID | |
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| | | | | |
| | | Fi | GURE 2 | |
| | | ST PF TII | TAGE I ROJECT SCHEDULE MELINE FOR FIELD ACTIVITIES | |

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KING SALMON AIR FORCE STATION

KING SALMON, ALASKA

1.

FIELD INVESTIGATION, SUMMARY, AND RESULTS

Field work completed as part of the Stage 1 investigation includes:

- Soil gas survey at the Eskimo Creek and Naknek River seeps
- Geophysical surveys at the Eskimo Creek and Naknek River seeps, and North and South Barrel Bluffs
- Groundwater monitoring well installations, slug tests, pump tests, sampling, and analysis at the Eskimo Creek seep site
- Sediment and surface water sampling at the Naknek River seep and North and South Barrel Bluff sites

Table 1 shows the number and type of analyses performed. Summary details of these investigations for each area are provided in the following subsections.

Eskimo Creek Seep

Groundwater Monitoring Wells. Seventeen shallow monitoring wells (30 to 40 feet deep), three deep monitoring wells (95 to 120 feet deep), and one soil boring (35 feet deep) were completed within the King Salmon AFS headquarters area. The three deep wells paired with shallow wells formed well nests, enabling hydrologists to estimate vertical hydraulic gradients. Stage 1 monitoring well and soil boring locations are shown in Figure 3.

The shallow monitoring well screens are 10 feet in length. These were generally positioned so that the center of the well screen intercepted the water table. This installation allowed "floater" constituent types to enter the monitoring well. The deep monitoring well screens are 10 to 20 feet in length. These were positioned in the aquifer where visible observations of hydraulic conductivity and well yield appeared to be the highest.

In situ slug tests were performed on each of the shallow wells. Pumping tests were performed on each of the deep wells to obtain estimates of the lateral hydraulic conductivity in the immediate vicinity of the subsurface zones penetrated by the monitoring well screens. Slug/bail test data were carefully evaluated because of the positioning of the shallow monitoring well screens above and below the water table.

These tests indicated an average hydraulic conductivity of 59 ft/day in the shallow aquifer zone and 25 ft/day in the intermediate aquifer zone. The vertical hydraulic gradient



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between the shallow and intermediate aquifers is estimated at 0.07 ft/day downward.

Shallow and intermediate aquifer groundwater flow directions are primarily from north to south across the site with localized variance in flow direction observed in the shallow aquifer. The vertical hydraulic gradient and groundwater flow directions have been estimated on the basis of a single round of groundwater measurements (May and June 1988) and may, therefore, not depict year-round conditions.

<u>Groundwater Quality</u>. Groundwater sampling and analyses were performed on each of the new monitoring wells, the station's two production wells (No. 4 and No. 5), and on eight existing US Corps of Engineers (USCOE) shallow wells. Table 1 shows the number and type of analyses performed.

Groundwater in the shallow aquifer contained measurable concentrations of aromatic volatiles, which include primarily benzene, toluene, and xylene. Detected concentrations of xylene were generally higher than concentrations of benzene and toluene. Benzene concentrations ranged from <1 to 77 μ g/l, toluene from <1 to 130 μ g/l, and xylene from <1 to 520 μ g/l.

Total petroleum hydrocarbons were also detected in shallow aquifer groundwater at concentrations ranging from <1 to 5,666 mg/1. Figure 4 shows the estimated lateral extent of hydrocarbons detected in shallow groundwater.

Higher concentrations of the aromatic volatile constituents may be present in the USCOE wells. Several of these wells are located in areas where floating product has been observed. However, analyses for aromatic volatiles could not be performed on groundwater samples from these wells because the well construction method included the use of PVC cement, which is partially composed of aromatic volatile constituents.

Groundwater samples collected from the deep monitoring wells did not contain constituents for which analyses were performed. Analyses of groundwater samples collected from the two station production wells indicate total petroleum hydrocarbons concentrations of 6.5 and 27 mg/1.

The source of these constituents is undetermined but may originate from lubricating fluids from the well pumps or from leakage along the well casing from the overlying contaminated portions of the shallow aquifer.

<u>Soil Vapor Gas</u>. The soil vapor gas study identified the presence of a total hydrocarbons plume beneath the headquarters area. Figure 4 shows the estimated lateral extent of this plume. Other localized hydrocarbon occurrences were

| | | Table l | | |
|--------|----|----------|----|------|
| NUMBER | OF | ANALYSES | BY | SITE |

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| Parameter | Analytical Method | Eskimo Creek | USCOE Well | Naknek River Seep | North Barrel Bluff | South Barrel <u>Bluff</u> | Offsite Well | Development Water |
|--|----------------------|-----------------|---------------|-------------------------|--------------------------|---------------------------------|-----------------|----------------------|
| Water Samples | | | | | | | | |
| Alkalinity - Carbonate, Bicarbonate, & Hydroxide (Field Test) | A403 | 23 | 7 | 1 | 3 | 3 | | |
| Common Anions (Chloride, Fluoride, Nitrate, Sulfate, Orthophosphate) | A429 | 23 | 7 | 1 | 3. | 3 | | |
| Specific Conductance (Field Test) | E120.1 | 23 | 7 | 1 | 3 | 3 | | |
| pH (Field Test) | E150.1 | 23 | 7 | 1 | 3 | 3 | | |
| Total Dissolved Solids | E160.1 | 23 | 7 | 1 | 3 | 3 | | |
| Temperature (Field Test) | E170.1 | 23 | 7 | 1 | 3 | 3 | | |
| Lead | SW3005/ SW7421 | 23 | 7 | | | | | 2 |
| Metal Screen (25 metals) | E200.7 | 2 | 0 | 1 | 3 | 3 | 1 | |
| Petroleum Hydrocarbons | E418.1 | 21 | 7 | | | | | |
| Purgeable Halocarbons | E601 | 2 | | 1 | 3 | 3 | 1 | |
| Organochlorine Pesticides | E608 | | | | 3 | 3 | 1 | |
| Extractable Priority Pollutants | E625 | | | | 3 | 3 | 1 | |
| Purgeable Aromatics | SW5030/ SW8020 | 16 | | 1 | 3 | 3 | 1 | 2 |
| Soil Samples | | | | | | | | |
| Petroleum Hydrocarbons | SW3550/ E418.1 | 18 | | | | | | |
| Organochlorine Pesticides and PCBs | SW3550/ SW8080 | | | | 3 | | | |
| Volatile Organic Compounds | SW5030/ SW8240 | 5 | | | 3 | | | |
| EPTOX Metals | 40 CFR 261.24 | 5 | | | | | | |
| Semivolatile Organic Compounds | SW3550/ SW8270 | 5 | | | 3 | | | |
| Soil Moisture Content | ASTM D2216 | 23 | | | 3 | | | |

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FIGURE 4

PLUME INTERPRETATION OF PETROLEUM HYDROCARBONS IN GROUNDWATER KING SALMON AIR FORCE STATION KING SALMON, ALASKA

CH:MHILL-

also identified and are shown on Figure 4. These appeared to be separated from the main plume based on the existence of other soil gas points with low or nondetectable values.

<u>Geophysical Survey</u>. Geophysical survey data did not specifically identify the location of an underground storage tank possibly associated with a deactivated service station west of Building 624. The survey did identify several other "reflective" targets that could be pipes or other buried metallic objects.

Naknek River Seep

Field investigation activities completed at the Naknek River seep were limited to several soil gas sampling locations and a water sample from the seep. A total hydrocarbons concentration of 2,200 μ g/l was observed at the riverbank in one of the soil gas sample locations. Several other soil gas sample locations were reported at less than method detection limits (0.03 μ g/l).

The geophysical survey data collected at this site indicate that a disposal area may exist at the western end of the site. Partially buried drums observed at the site support this conclusion. Water samples collected from the seep on the north bank of the Naknek River identified the presence of xylene and 1,1,1-trichloroethane at concentrations of 15 μ g/1 and 85 μ g/1, respectively.

North Barrel Bluff

Geophysical survey data collected at the North Barrel Bluff site indicate that buried and partially buried debris are concentrated along the face of the bluff in a section 50 to 100 feet wide and 2,300 feet long.

South Barrel Bluff

Data from geophysical surveys performed along the South Barrel Bluff indicate that the primary disposal areas are located along the face of the bluff beginning at a point east of the two residential homes and extending westward for approximately 1,000 feet. Other localized buried debris areas are located in a gully east of the two houses. Chlordane was detected in a surface water sample at a concentration of $0.2 \ \mu g/1$.

RECOMMENDATIONS

Suggested activities and the rationale for each investigation area are presented in Table 2.

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| Suggested Activity | E Rationale | iskimo Creek Seep | Naknek River Seep | North Barrel Bluff | South Barrel Bluff |
|--|---|----------------------|----------------------|-----------------------|-----------------------|
| Review facility as-built drawings and historical aerial photographs | Determine location of burled and/or deactivated structures | x | × | × | × |
| Conduct soil gas survey | Delineate lateral extent of the total hydrocarbons detection observed in the Stage 1 investigation to assist in location of shallow monitoring wells | | × | | |
| Collect samples from shallow soil bor- ings and surface sediment locations | Assess the nature and extent of present contamination in unsaturated and near-surface soils | × | × | × | × |
| Collect and analyze surface water samples | | x | | | |
| Install shallow and/or intermediate monitoring wells and perform single well tests | Determine groundwater flow direc- tions, estimates of aquifer parameters and provide groundwater samples for water quality characterization | × | × | × | × |
| Perform groundwater measurement and water quality sampling | Confirmation of results and investi- gation of seasonal affects | × | × | × | × |
| Analyze soil and water samples for expanded constituent list (priority pollutants) | Identify and quantify individual constituents not tested for in Stage 1 | × | × | × | × |
| Select appropriate analytical methodologies to address data quality objectives | Assure detection limits are appropri- ate for risk analysis and site char- acterization | × | × | × | × |
| Identify and select background soil and groundwater locations | Correlate background concentrations with site concentrations for risk analysis assessment | × | × | × | × |
| Conduct geophysical survey | Identify small disposal areas not pre- viously located and further delineate lateral extent of buried debris iden- tified in Stage 1 reconnaissance investigation | | × | × | × |
| Dig test pits with backhoe | Waste characterization | | | × | × |
| Construct perimeter fence | Reduce potential for human and/or wildlife exposures through physical contact | | | × | × |

| | Table 2 (Continued) Recommendations for King Sa | LMON AFS | | | |
|--|---|----------------------|----------------------|-----------------------|-----------------------|
| Suggested Activity | Rationale | Eskimo Creek Seep | Naknek River Seep | North Barrel Bluff | South Barrel Bluff |
| Include residential water wells in groundwater sampling program | Identify potential health risks associated with a water supply in close proximity to disposal areas | | | | × |
| Conduct shallow aquifer pumping test | Provide estimate of aquifer parameters for "pump and treat" remediation | × | | | |
| CVR193/061 | | | | | |

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I. INTRODUCTION

1.1 AIR FORCE INSTALLATION RESTORATION PROGRAM

The Installation Restoration Program (IRP) is a Department of Defense (DOD) program for assessing and remediating hazardous waste problems on military installations. The program is DOD's response to the requirements of the Comprehensive Environmental Response Compensation and Liability Act (CERCLA or "Superfund") which requires all federal agencies to fully comply with its procedural and substantive requirements. The program was formally designated as the DOD Superfund program on November 21, 1981.

1.1.1 Program Objectives

The objective of the Air Force IRP is to assess past hazardous waste disposal and spill sites on Air Force installations and develop remedial actions consistent with the National Contingency Plan (NCP) for those sites that are determined to pose a threat to human health and welfare or the environment. This program objective is derived from the overall objective of the National Contingency Plan (NCP) (40 CFR -300.68i), i.e., to select "a cost-effective remedial alternative that effectively mitigates and minimizes threats to and provides adequate protection of public health and welfare and the environment." Future actions at King Salmon AFS must meet the requirements of CERCLA, as amended by the Superfund Amendments and Reauthorization Act (SARA), and the NCP.

The NCP is currently under revision, in response to SARA. These revisions are expected to conform to the guidelines established in SARA as well as the Interim Guidance on Superfund Selection of Remedy (December 24, 1986), which supersedes portions of the NCP until its revisions are complete.

Two major provisions of SARA affect selection of treatment technologies. They pertain to regulatory requirements and permanence of treatment. Provisions regarding preference for technologies that will effect permanent remedies are stated in Section 121(b)(1) of SARA:

"...The President shall conduct an assessment of permanent solutions and alternative treatment technologies or resource recovery technologies that, in whole or in part, will result in permanent and significant decrease in the toxicity, mobility, or volume of the hazardous substance, pollutant, or contaminant..." The importance of applicable or relevant and appropriate requirements (ARARs) are defined in SARA and outlined in the Interim Guidance memorandum:

"...Section 300.68 of the NCP specifically refers to ARARs in regard to the development of alternatives. SARA incorporates this requirement into statutory law while adding the provision that remedial actions also attain state requirements more stringent than federal requirements if they are also applicable or relevant and appropriate..."

1.1.2 Organization of Installation Restoration Program

The Air Force has developed its Installation Restoration Program to comply with federal regulations. The IRP was previously organized into four sequential phases, described below:

- O Phase I--Records Search. In this phase, past disposal sites which may be hazardous are identified. This phase constitutes a preliminary assessment of the hazards at each installation. Sites are identified from a review of base records and interviews with current and former base employees. Hazards and potential hazards are assessed based on material disposed and a review of regional geological and hydrogeological factors. Sites are then ranked and those with the highest hazard potential are recommended for additional investigation in Phase II. This phase does not include sampling and analysis.
- o <u>Phase II--Confirmation</u>. In this phase field and laboratory data are collected and analyzed to identify the nature and extent of the contamination at sites identified in Phase I. Sites requiring remedial action are identified and programmed for further work.
- <u>Phase III--Technology Base Development</u>. Contaminants identified in Phase II are further evaluated to determine if they can be treated with known technologies. If no treatments are identified, treatment technologies are developed in this phase.
- <u>Phase IV--Operations</u>. This is the action phase of the IRP. Remedial actions are recommended, then designed, built, and executed.

The Air Force has modified its IRP to make it similar to the EPA's Remedial Investigation/Feasibility Study (RI/FS) program. It has combined Phase II and a portion of Phase IV in order to conduct the remedial investigation and feasibility study in parallel, where possible, instead of in serial fashion. With this approach, preliminary remedial alternatives can be identified and screened as information on the nature and extent of contamination is obtained. The early identification of remedial alternatives that are potentially applicable to the King Salmon sites will contribute to the definition of further data requirements, including data required to assess the feasibility of the preliminary remedial alternatives (e.g., treatability and incineration characteristics).

1.2 DESCRIPTION AND HISTORY OF INSTALLATION

The primary mission of the 5071st Combat Support Squadron (CSS), based at King Salmon AFS, is twofold. The installation is a forward operating base supporting aircraft deployed at the direction of the Commander of the 21st Tactical Fighter Wing (TFW) based at Elmendorf Air Force Base in Anchorage, Alaska, and is also a surveillance radar installation.

The Phase I report provided the following concise narrative of the history of the installation:

King Salmon Airport was constructed by the Civil Aeronautics Authority [now Federal Aviation Administration (FAA)] as part of an overall airfield construction program in Alaska during the early 1940s. The field was completed and turned over to the U.S. Army in 1941. The airfield became an advance staging base and fuel stop for aircraft deploying to and from the Aleutian Islands. It was returned to the control of the FAA in 1945. The FAA turned it over to the State of Alaska in 1959. The State continues to operate the airport with the Air Force as its major tenant.

The Alaskan Air Command (AAC) began using the airport in 1948 as a Forward Operating Base. The installation has provided operational and maintenance support to the aircraft on alert. King Salmon AFS was also one of the original ten aircraft control and warning (AC&W) sites constructed as part of a permanent air defense system in Alaska during the early 1950s. King Salmon became operational as a ground controlled intercept site in 1951. It was converted to a North American Air Defense (NORAD) Control Center in 1953.

Communications were initially provided by a high frequency radio system. A White Alice Communications System, tropospheric scatter and microwave relay equipment site was activated at King Salmon in 1957. It was

deactivated in 1979 and replaced by an Alascom-ownedand-operated satellite earth terminal.

In 1977, the AAC implemented a base support contract with RCA Services as part of an Air Force-wide effort to reduce remote tours. Twenty-eight military positions were eliminated. The remaining 63 positions were primarily in operations. Installation of Joint Surveillance System (JSS) equipment was completed in 1982, enabling radar and beacon data to be transmitted via satellite to the Region Operations Control Center (ROCC). This further reduced the number of personnel required at the installation.

1.3 IRP PHASE I RECORDS SEARCH

The Phase I records search for King Salmon AFS, completed in September 1985 by Engineering-Science, identified 17 sites containing potentially hazardous materials. In addition to the 17 specific sites, the Phase I records search also indicated a need for follow-up monitoring of drinking water supply wells at the installation and in the adjacent community. The Phase I Records Search was performed by completing the following tasks:

- o Identification of Past Waste Management Practices
- o Evaluation of Site-Specific Background Information
- Prioritization of each site using a hazard assessment rating methodology (HARM)
- Selection of sites from the HARM list, that pose the greatest potential hazard to human health and the environment

Each task is discussed briefly in the following subsections.

1.3.1 Past Waste Management Practices

The Phase I investigation reviewed past and present activities at King Salmon AFS resulting in the generation, accumulation, and disposal of hazardous wastes.

The sources of hazardous waste at King Salmon AFS have been grouped into the following categories:

- o Industrial Operations (Shops)
- o Waste Accumulation Areas
- o Fuels Management
- o Spills and Leaks
- o Pesticide Utilization
- o Fire Protection Training

| | | PAST AND PRESENT WASTE MANA | AGEMENT PRACTICES | | |
|--------------------------------------|------------------------|--|--|--|---|
| Category | Location (Building) | Type of Waste or Stored Material | 1985 Rate of Generation | D1sposa Pre-1983 | l Site Post-1983 |
| I NDUSTRIAL OPERATIONS | | | | | |
| POL Maintenance | 642 | Contaminated JP-4 ^a Contaminated Diesel Fuel | 500 gal/yr 400 gal/yr | FPTA FPTA/Road 0111ng | F PTA INC/DRMO ^b |
| Liguid Fuels Operations | 177 | Contaminated JP-4 | 240 gal/yr | FPTA | INC/DRMO ^b |
| Refuel Maintenance | 149 | Contaminated JP-4 Contaminated Diesel Fuel Lubricating Oil & Hydraulic Fluid PD-680 | 300-600 gal/yr 120-180 gal/yr 440 gal/yr 30 gal/hr | Dry Well Dry Well Dry Well Dry Well | INC/DRMO ^b INC/DRMO ^b INC/DRMO ^b INC/DRMO ^b |
| Radar | Mar Tower | Gearbox 011 | 20 gal/yr | Road Oiling | INC/DRMO ^b |
| Vehicle/ Equipment Maintenance | 636 | Lubricating Oil & Hydraulic Fluid Paints & Thinners Battery Acid PD-680 | 500 gal/yr 20 gal/yr 90 gal/yr 55 gal/yr | Road Olling Road Olling Neutralized/ Sanitary Sewer Road Olling | INC/DRMO ^b INC/DRMO ^b Neutralized/ Sanitary Sewer INC/DRMO ^b |
| Power & Heating Plant | 638, 642 | Lubricating Oil Ethylene Glycol/Water Diesel Fuel (Parts Cleaning) | 860 gal/yr 500 gal/yr 200 gal/yr | Road Oiling Road Oiling Road Oiling | INC/DRMO ^D DRMO ^D INC/DRMO ^D |
| Barrier Maintenance | 638 | Lubricating Oil | 60 gal/yr | Road Oiling | INC/DRMO ^b |
| Aircraft Maintenance Shops | 160 | PD-680 Lubricating Oil Compressor Oil Brake Fluid Hydraulic Fluid Contaminated JP-4 | 25-50 gal/yr 300-400 gal/yr 50 gal/yr 2 gal/yr 50 gal/yr 700 gal/yr | FPTA/Road 0111ng FPTA/Road 0111ng FPTA/Road 0111ng FPTA/Road 0111ng FPTA/Road 0111ng FPTA/Road 0111ng | FPTA FPTA FPTA FPTA FPTA FPTA |
| 011-Water Separators | Various Locations | Sluđge | Not reported | Not report e ð | DRMO |

Table 1-1

| | | Table 1-1 (Continued | | | |
|--|---|--|--|---------------------|----------------------------------|
| | Location | Type of Waste or ctored Material | 1985 Rate of Generation | Disposa Pre-1983 | 1 Site Post-1983 |
| Category MASTE ACCUMULATION AREAS | (biit bi ting) | | | į | Shirt |
| Waste Accumulation Area No. 1 | (See F1gure 1-2) | Waste Oils Shop Wastes Spent Solvents | Approximately 5,500 gal. | DRMO | David |
| Maste Waste Accumulation Area No. 2 (Trace ive) | (See Flgure 1-2) | Not Reported | Not Reported | Not Reported | Not Applicable |
| Waste Maste Accumulation Area No. 3 (Inactive) | (See Figure 1-2) | Not Reported | Not Reported | Not Reported | Not Applicable |
| <u>FUELS MANAGEMENT</u> Above Ground & Below Ground Tanks | Various Locations | D1esel, AVGAS, JP-4, MOGAS, Isopropyl Alcohol | 62 Tanks with a total capacity of approx. 5,500,000 gal | Not Applicable | Not Applicable |
| Pipelines | Various Locations | | | Not Applicable | Not Applicable |
| POL Tanks 11, 12, 13, 14 | (See Figure 1-2) | Diesel Fuel | 1,049,000 gal | Not Applicable | Not Applicable Wot Bunlicable |
| POL Tank Farm (Nos. 2, 3, 4, 5, 6, 7, 22, 23, 24, 25, 30, 28) | Naknek River Seep (See Figure 1-2) | Diesel Fuel Isopropyl Alcohol | 450,000 gal 100,000 gal | Not Applicable | NUL APPLICATION |
| Deactivated POL Tank (No. 27) | Unmumbered, near Naknek River Seep (See Figure 1-2) | Unknown | 50,000 gal | Not Applicable | NOL AFFA |

| Category POL Tank (No. 1) POL Tank (No. 1) SPILL AND LEAKS Eskimo Creek Seep N Naknek River Seep N Naknek River Seep N PESTICIDE PESTICIDE PROFINION PROFINION | Location (Building) (Building) Approx. 1,100 ft. SSW of center of SSW of center of SSM of center of SSM of center of SSM of center of Figure 1-2) (Figure 1-2) (Figure 1-2) (Figure 1-2) (Figure 1-2) (Figure 1-2) (Figure 1-2) | Type of Waste or Stored Material Diesel Fuel Diesel Fuel Uncharacterized Petroleum Froduct (Active tanks in vicinity of seep store diesel fuel, isopropyl alcohol, and MOGAS) Malathion fogging in past years Contaminated and noncontam- | 1985 Rate of Generation 427,000 gal all volume over time sufficient to discolor sur- face soil, stress vegetation, and be recoverable in test pit Observable flow in 1984, but 1985 thase 1 report noted sheen only, and May 1987 site visit (CH2M and OEHL personnel) noted soil dis- coloration only Phase I report judged quantities to be small enough to warrant no further invest- igation 50-ftdiameter | Pre-1983 Not Applicable (First observed in 1977) Not Applicable Not Applicable Incineration | Not Applicable Not Applicable Underlying soll, shallow ground- water, Estiments and surface wate ground surface, and surface, and surface wate in seep Not Applicable Not Applicable |
|--|--|--|---|---|---|
| FPTA No. 2 (1979-Present) | (Figure 1-2) | Forming roam (Arfr) Wood debris, contaminated fuels, AFFF, halon, potassium bicarbonate | montn Areal extent/vol- ume of pit not reported, practice burns once/month | Incineration | Incineration ^b |

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|--|

^dContaminated JP-4--Waste fuel that has been contaminated with other materials.

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^bBeginning in 1986, King Salmon no longer incinerates waste POL products. This waste is now shipped to DRMO at Elmendorf AFB. Source: Phase I Report (Engineering Science)

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Table 1-1 summarizes the past and present waste management activities at King Salmon AFS for those activities involving hazardous materials or generating hazardous wastes.

1.3.2 Site-Specific Background Information

After a review of the past waste generation and disposal practices at King Salmon AFS, the Phase I investigation identified the 17 sites (Figure 1-1) as being potentially contaminated with hazardous materials. The Alaskan Air Command (AAC) prioritized these 17 sites based on hazard assessment ranking methodology (HARM) scores (Table 1-2). In ranking each site, the AAC also considered restoration activities accomplished at the site to date and the location sequencing of work tasks over the duration of the King Salmon IRP.

| Phase I Site No | Sito Namo | HARM Score |
|--------------------|-------------------------------------|------------|
| SILE NO. | Site Name | |
| 1 | Fuel Seepage at Eskimo Creek | 86 |
| 2 | POL Tanks No. 11, 12, 13, 14 | 72 |
| 3 | Fuel Seepage at Naknek River | 71 |
| 4 | Building 1959 Dry Well | 65 |
| 5 | Dump | 63 |
| 6 | Road Oiling | 61 |
| 7 | Waste Accumulation Area No. 2 | 61 |
| 8 | Landfill No. 1 | 60 |
| 9 | Landfill No. 2 | 60 |
| 10 | Fire Protection Training Area No. 1 | 59 |
| 11 | Fire Protection Training Area No. 2 | 59 |
| 12 | Fire Protection Training Area No. 3 | 59 |
| 13 | Fire Protection Training Area No. 4 | 59 |
| 14 | Waste Accumulation Area No. 3 | 59 |
| 15 | Landfill No. 3 | 57 |
| 16 | Landfill No. 5 | 57 |
| 17 | White Alice Site | 55 |

Table 1-2 PHASE I SITE IDENTIFICATION AND HARM RANKING SCORE

Six of the highest priority sites from Table 1-2 were selected for investigation during the Stage 1 investigation (Table 1-3).

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| Phase I Site No. | Phase I Site Name | Stage 1 Site Name |
|---------------------|------------------------------|--------------------|
| 1 | Fuel Seepage at Eskimo Creek | Eskimo Creek Seep |
| 2 | POL Tanks No. 11, 12, 13, 14 | |
| 3 | Fuel Seepage at Naknek River | Naknek River Seep |
| 5 | Dump | North Barrel Bluff |
| 8 | Landfill No. 1 | South Barrel Bluff |
| 17 | White Alice Site | North Barrel Bluff |
| | | |

Table 1-3 PHASE I SITES SELECTED FOR INVESTIGATION IN STAGE 1

Figure 1-2 shows the approximate location of each site. These six sites were combined to form four sites, on the basis of location and potential waste characteristics for purposes of the initial efforts performed under this scope of work. The four operable units are identified in Figure 1-3. As additional information is obtained, the boundaries and grouping of these units may be redefined for purposes of design of the initial remedial measure and development of preliminary alternative remedial actions. Table 1-4 identifies the types of waste believed to exist at each of these sites and potential migration pathways for contamination associated with the sites. A discussion of each site follows.

1.3.2.1 Eskimo Creek Seep

The Eskimo Creek seep is a diesel fuel seep located east of Building 603 at Eskimo Creek. The source of this seep may be any of several active facilities in the vicinity of the seep, past spills, and/or associated with active or abandoned portions of the installation's liquid fuel system underground pipelines.

The installation power plant, motor pool, POL Tanks 11, 12, 13, and 14 and fuel lines are all potential sources associated with the management of large quantities of diesel fuel. Leaks and spills from any of these sources may have contributed to fuel oil contamination of the shallow groundwater allowing the oil to migrate along the water table to the point where it emerges as a seep at Eskimo Creek. This seepage of POL was first observed in the mid-1970s when the





| Site | Waste Type | Potential Migration Pathways |
|--------------------|--|--|
| Eskimo Creek Seep | Diesel Fuel " | Shallow groundwater migration; shallow groundwater communication with deeper groundwater/drinking water supply; surface water (Eskim Creek, Naknek River); aquatic life and food chain; direct contact with contaminated surface soils and sediments; direct contact with surface water |
| Naknek River Seep | Petroleum products (not otherwise specified, but possibly diesel fuel, isopropyl alcohol, MOGAS, or unknown contents of unnumbered deactivated POL tank) | Shallow groundwater migration; shallow groundwater communication with deeper groundwater/drinking water supply; surface water (Naknei River); aquatic life and food chain; direct contact with contami- nated surface soils and sediments; direct contact with surface wate: |
| North Barrel Bluff | Discarded drums (mostly empty, but 10 percent of capacity estimated to contain liquid products and wastes); wood and metal debris | Leachate migration to surface water (bog, King Salmon Creek, Naknek River) by surface runoff or shallow groundwater transport; aquatic life and food chain; leachate migration to shallow groundwater; shallow groundwater communication with deeper aquifer and drinking water supply; direct contact with debris; volatilization of contami- nants to ambient air |
| South Barrel Bluff | Discarded drums (mostly empty, but 10 percent of capacity estimated to contain liquid products and wastes); wood and metal debris; shop wastes (paints, thinners, etc.); refuse | Leachate migration to surface water (bog, King Salmon Creek, Naknek River) by surface runoff or shallow groundwater transport; aquatic life and food chain; leachate migration to shallow groundwater; shallow groundwater communication with deeper aquifer and drinking water supply; direct contact with debris; volatilization of contami nants to ambient air |

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Table 1-4 STAGE 1--WASTE TYPES BY SITE

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rate was reportedly as much as 10 gallons per day. The visible seepage was reportedly the greatest when the water table was high, and nearly stopped during the winter lowwater period. Recently the reported flow rate has been lower, approaching 1 to 2 gallons per month during winter months.

. . .

The Phase 1 report indicates that analytical results of samples from the Eskimo Creek site show the POL contamination is a type of diesel fuel that was not used at the installation after 1973, thus implying that the fuel entered the ground prior to that date.

In April 1987, CH2M HILL and USAFOEHL personnel visited the Eskimo Creek seep site and measured 0.6 foot of petroleum product in a monitoring culvert adjacent to Building 648 and 0.3 foot of petroleum product in U.S. Corps of Engineers Monitoring Well 22. In addition, U.S. Corps of Engineers Monitoring Wells 9 and 18 showed noticeable petroleum odors, but no other wells contained measurable floating product.

1.3.2.2 Naknek River Seep

The Naknek River seep is located immediately south of Landfill No. 5 and the nearby POL tank farm. The facility Master Plan dated January 1975 also depicts a deactivated 50,000gallon tank (No. 27) at the POL tank farm and POL tanks and diesel fueling facilities within 400 feet of the Naknek River seep. The Master Plan indicates that five of the tanks at the tank farm were scheduled for deactivation on September 1, 1978. According to the Phase I investigation, the first report of this seep occurred in 1984 when a "fuel/ water mixture" was observed flowing down the embankment toward the Naknek River in the vicinity of the King Salmon AFS POL tanks nearest the river. A 6-foot-deep excavation along the crest of the embankment exposed oil on the shallow groundwater surface, and the Phase I report described an oily film on the water surface at the time of the site visit for that investigation.

When CH2M HILL and USAFOEHL personnel viewed this site in April 1987, some soil discoloration was observed at the embankment, but no flow of fuel/water was observed.

1.3.2.3 North Barrel Bluff

This site is outside the King Salmon AFS installation boundaries and is comprised of two Phase I sites: the dump and the former White Alice site. The dump was described in the Phase I report as consisting of drums, metal, and wood debris extending along the valley of King Salmon Creek from the White Alice site to Landfill No. 1. However, the CH2M HILL Presurvey Report described two separate exposed barrel dumps along the King Salmon Creek Bluff, the northern one being associated with the White Alice site and the southern one associated with Landfill No. 1.

Estimates of the number of barrels at those dumps range from hundreds of thousands to one million, with the number of barrels containing liquid estimated to be approximately 10 percent. During 1984, approximately 2,000 drums were removed from the site, and some of the remaining barrels were covered with soil. The drums were buried at Landfill No. 4 pending permission regarding disposal, and may have contained residual liquids.

1.3.2.4 South Barrel Bluff

This site is southwest of the previously described North Barrel Bluff site, on the east bank of King Salmon Creek. The site includes Landfill No. 1, a closed and covered landfill that was probably operated from the 1940s to the mid--1970s. Similar to the North Barrel Bluff, this dump also consists of drums, metal, and wood debris. The areal extent of the barrel bluff sites cannot be determined by visual inspection, since both sites are partially covered with soil.

1.4 KING SALMON AIRPORT PROJECT OBJECTIVES

The current study is Stage 1 of the combined Phase II/IV remedial investigation and feasibility study for the operable unit sites at King Salmon Airport. In addition to remedial investigation tasks, this Stage 1 effort includes feasibility study tasks through development of preliminary alternative remedial actions. Also included in this study is design of a Conceptual Remedial Measure (CRM) for the petroleum product seep at Eskimo Creek. Following the RI/FS tasks, a work plan will be prepared to determine if any additional confirmation and feasibility studies are needed, as well as remedial actions that can be implemented in 1988-89 work for King Salmon Airport.

This IRP Stage 1 work for King Salmon Airport was performed as authorized by Contract No. F33615-85-D-4535, Delivery Order No. 0006.

The purpose of the Phase II/IV, Stage 1 work conducted in 1987 and 1988 was to (1) collect and evaluate field data to define to site hydrogeology, (2) confirm the presence or absence of contamination within the specified areas of investigation, and (3) determine the extent of contamination at each of the King Salmon Airport operable unit sites and recommend remedial action for each. The specific Stage 1 tasks identified were:

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- 1. Conduct a literature search to determine the geological, hydrogeological, and environmental settings for this investigation
- 2. Perform a soil gas survey and drill monitoring wells, taking soil samples for analyses, to delineate the petroleum, oil, and lubricants (POL) contaminant plume in the vicinity of Tanks 11, 12, 13, and 14 and the Eskimo Creek seep
- 3. Determine the nature and extent of contamination of groundwater in the vicinity of POL Tanks 11, 12, 13, and 14 and the Eskimo Creek seep
- 4. Determine the nature and extent of surface water contamination of Eskimo Creek resulting from the POL seep
- 5. Design a Conceptual Remedial Measure (CRM) to contain/ intercept the POL seep at Eskimo Creek
- 6. Determine the nature of the petroleum product seep at Naknek River
- 7. Define the horizontal and vertical extent of barrels and other debris at the North Barrel Bluff
- 8. Determine whether the wetland area that lies between King Salmon Creek and the face of the North Barrel Bluff and the surface waters of King Salmon Creek downstream of the barrel bluff have been contaminated by liquids from the barrels and debris disposed at this site
- 9. Determine the horizontal and vertical extent of the barrels and other debris at the South Barrel Bluff and Landfill No. 1
- 10. Determine whether surface water in King Salmon Creek is contaminated downstream of the South Barrel Bluff
- 11. Perform a qualitative risk assessment to determine the potential environmental or health hazards associated with those contaminants found
- 12. Develop preliminary alternative remedial actions for the sites addressed in this scope of work
- 13. Develop data quality objectives (DQOs) necessary to perform the detailed evaluation of remedial alternatives and to conduct the quantitative risk assessment required in any future IRP work on these sites

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1.5 PROJECT TEAM ORGANIZATION

The CH2M HILL project team organization and assignments are shown in Figure 1-4. Detailed biographies are provided in Appendix J.

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II. ENVIRONMENTAL SETTING

2.1 <u>GEOGRAPHIC SETTING</u>

2.1.1 Physiography

King Salmon Airport is located on an 86-acre site in the central portion of Section 23, Township 17 S, Range 45 W, Seward Meridian. Figure 2-1 provides a location map and Figure 2-2 shows a detailed vicinity map. The installation is on the northwestern section of the Alaska Peninsula approximately 340 miles southwest of Anchorage, Alaska. This area is generally characterized by glaciated zones bounded by well defined moraines, with some gully development along morainal ridges. Many kettle basins containing lakes are present throughout the area. Most area drainage is not fully integrated.

The Naknek River lies to the south of King Salmon Airport. Eskimo Creek flows through the central portion of the base, and King Salmon Creek lies to the northwest.

Ground surface elevations on the base range from 30 feet mean sea level (MSL) along the banks of the Eskimo Creek, to 68 feet MSL within the central portion of the base headquarters area.

2.1.2 <u>Vegetation</u>

Vegetation at King Salmon consists primarily of scattered white spruce with ericacous shrubs. Trees are sparse, and most of the vegetation is short. The floodplains of Eskimo and King Salmon Creek are wetlands. They contain a variety of shrubs, herbs, grasses, and sedges rooted in a continuous matrix of lichens and mosses. Depressions typically nurture grasses and sedges while raised hummocks and hills foster crowberry, birch, willow, and blueberry. Summer blooms include lousewart, violet, buttercup, and other wild flowers (Alaska Department of Community and Regional Affairs, 1982).

2.1.3 <u>Cultural Geography</u>

2.1.3.1 Land Use

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Land development at King Salmon began in the 1930s when an air navigation silo was built. During World War II, an Air Force base was constructed. The Federal Aviation Administration (FAA) maintained the base throughout the war. The King Salmon Airport remains today one of the major military installations in western Alaska. Other facilities at King Salmon include a road between the town and Naknek, a post office, an Alaska Department of Fish and Game (ADFG)



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office, a weather bureau office, inns, stores, and private businesses.

Three of the largest landowners in the King Salmon area are the federal government, the State of Alaska, and Paug-vik, Inc. (Naknek's native village corporation). The State of Alaska has claimed 1.56 acres for the King Salmon Airport and 12.55 acres on the Naknek River for use by the ADFG. The U.S. Fish and Wildlife Service (USFWS) owns 8.24 acres on the Naknek River adjacent to the parcel occupied by the ADFG. The National Park Service (NPS) has a King Salmon office situated on 11.43 acres. Much of the land is owned by the federal government for use by the Air Force and FAA.

The Alaska Native Claims Settlement Act (ANCSA) (1971) does not acknowledge King Salmon as a native village. Consequently, a native village corporation has not been established for King Salmon. Interim conveyance and patents to land around King Salmon have been provided to Paug-vik, Inc.

Pursuant to the Native Allotment Act of 1906, the Secretary of the Interior may provide up to 160 acres of land to individual natives with the stipulation that the claimant occupies and uses the allotment continuously. By 1982, natives had filed 24 active claims and 2 patented claims near King Salmon. With the passage of ANCSA, new claims were not accepted between 1971 and 1982. Provision for the approval of pending claims was established by the Alaska National Interest Lands Conservation Act (ANILCA) in 1980. Claims are not approvable under ANILCA in cases where protests have been filed. Certificates of claim approval are issued when claims are officially surveyed (Alaska Department of Community and Regional Affairs, 1982).

2.1.3.2 Demography

In 1980, the population at King Salmon was predominantly white and male. The distribution by race was 81.1 percent white, 8.6 percent black, 5.9 percent native, 0.9 percent Asian, and 3.4 percent other races. Males outnumbered females by a ratio of 3.7 to 1 (Alaska Department of Community and Regional Affairs, 1982).

A 1980 Census Bureau survey documented 43 civilian families and 75 households at King Salmon. Since station personnel reside at the installation on unaccompanied assignments, their families remain at home. In 1987, this Air Force station accommodated 275 military personnel and 19 civilians on assignments related to the military (USAF, 1987).

2.1.3.3 Population Density

In 1980 more military personnel resided at King Salmon than civilians: 340 military personnel resided at King Salmon Airport; 205 civilians lived in the village. The civilian population has grown to more than 300, while the military personnel now numbers fewer than 300.

The summer population frequently increases with the seasonal influx of NPS, ADFG, and local hotel and restaurant employees. Additionally, many people pass through King Salmon enroute to fishing and cannery jobs on the coast or to sport fishing and hunting trips (Alaska Department of Community and Regional Affairs, 1982).

2.1.3.4 Age Distribution

A 1980 U.S. Census Bureau survey recorded most of the inhabitants at King Salmon were between the ages of 22 and 44 years. The median age was 27.1 years. Age distribution is shown in Table 2-1.

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2.1.3.5 Education Level

The Bristol Bay School District provides education to residents of North and South Naknek and King Salmon. Children attending preschool through grade 12 commute daily to Naknek. Some students pursue higher education. No information is presently available on the education of King Salmon residents (Alaska Department of Education, 1987).

The education level of Air Force personnel at King Salmon reflects that of the Air Force in general. Air Force personnel typically have high school educations. Officers are required to have a baccalaureate degree.

2.1.3.6 Socioeconomics

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The economy of King Salmon is based primarily on employment related to government, transportation, and fishing. The U.S. government provides the largest, most stable source of employment. Civilians holding government jobs work for the U.S. Air Force, the FAA, USFWS, NPS, and the U.S. Postal Service. The King Salmon installation is self-contained and does not contribute substantially to the local economy. Occasionally, some military personnel seek part-time employment with local flying services, restaurants, and canneries.

| Age | | |
|------------|-------|--------|
| (Years) | Total | Female |
| Under 1 | - | - |
| 1-2 | 5 | 2 |
| 3-4 | 5 | 2 |
| 5 | 5 | 1 |
| 6 | 5 | 5 |
| 7-9 | 18 | 9 |
| 10-13 | 14 | 9 |
| 14 | 4 | 2 |
| 15 | 8 | 6 |
| 16 | 3 | 2 |
| 17 | 1 | - |
| 18 | 5 | 4 |
| 19 | 3 | 1 |
| 20 | 8 | 1 |
| 21 | 37 | 1 |
| 22-24 | 103 | 16 |
| 25-29 | 119 | 15 |
| 30-34 | 81 | 14 |
| 35-44 | 76 | 16 |
| 45-54 | 31 | 6 |
| 55-59 | 4 | - |
| 60-61 | 4 | 2 |
| 62-64 | 3 | 2 |
| 65-74 | 3 | 1 |
| 75-84 | - | - |
| i and over | 1 | 1 |
| | | |

| | Ta | able | 2-1 | |
|------|--------|------|--------------|--|
| KING | SALMON | AGE | DISTRIBUTION | |

Source: U.S. Census Bureau, 1980

The State of Alaska provides year-round and summer jobs in King Salmon. Most state workers are based in King Salmon and work throughout the Bristol Bay Borough. Major state employers include the Department of Public Safety (DPS), ADFG, and the Department of Transportation and Public Facilities (DTPF).

Commercial flying services employ much of the local work force. Airline employers include Peninsula Airways, King Flying Service, and Mark Air. Hotels and restaurants provide a few additional jobs. A few residents seek employment with the Bristol Bay red salmon fishery, sport fishing lodge operations, a construction company, a car rental business, and miscellaneous small enterprises.

2.1.3.7 Public Health and Welfare

The Camai Medical Center at Naknek is the only health clinic serving King Salmon.

2.2 <u>GEOLOGY</u>

2.2.1 <u>Geomorphlogy</u>

King Salmon AFS is located in an area exhibiting the characteristics of past intense glaciation during Pleistocene time (between 10,000 and 1.6 million years ago). Glacial activity is significant on the Alaska Peninsula in that glacier advance and retreat eroded the uplands into block-like groups of mountains with rounded crests separated by U-shaped valleys and low passes. The mountain ridges that rose above the ice sheet remained angular and sharp in appearance. The retreat and melting of the large glaciers produced great quantities of outwash sediment, which has resulted in the filling of many basin and lowland areas.

Alaska's generally cold climatic regime has produced a subsurface condition termed "permafrost," a combination of geological, hydrologic, and meteorologic characteristics that produce perennially frozen ground. Permafrost occurs in both unconsolidated sediments and bedrock. The occurrence of permafrost in Alaska is mapped as areas of continuous permafrost, discontinous permafrost, and no permafrost. King Salmon AFS is located in the zone of discontinous permafrost (Figure 2-3).





2.2.2 Bedrock Geology

Bedrock is not exposed in the vicinity of King Salmon AFS and underlies overburden materials at depths of several hundred feet.

2.2.3 Surficial Geology

In the lowland areas in the vicinity of King Salmon AFS, well-sorted recent alluvium is present, consisting primarily of sand, silt, and clay in stratified layers. Glaciofluvial sediments occur beneath the slopes and terraces overlooking the valleys of streams and rivers. These sediments are poorly to moderately well-sorted outwash and morainal materials (i.e., unconsolidated materials deposited by the melting glacier or by erosion of older moraines). The upland areas bordering or overlooking the site are underlain by glacial moraine and drift materials. These materials are mixed, unsorted, and generally unstratified clay, silt, sand, gravel, cobbles, and boulders arranged in a conspicuous arcuate pattern, usually about the larger lakes. Figure 2-4 shows geologic units which are surficially exposed on the Alaska Peninsula.

At King Salmon AFS, glacial outwash and sand deposits blanket the surface. The outwash sand ranges in thickness from about 20 to 40 feet and overlies silty glacial till.

2.3 HYDROGEOLOGY

2.3.1 Occurrence and Movement of Groundwater

The most productive groundwater sources are the unconsolidated alluvial aquifers of Alaska's major river valleys and the glacial outwash aquifers underlying coastal basins and some lowland areas. King Salmon is underlain by glacial outwash.

The primary source of recharge to the aquifers beneath King Salmon is precipitation. Additional sources of recharge to these aquifers include stream channels that lose flow (influent streams), lakes, topographic high points, and slopes of low hills.

Groundwater discharges from aquifers to springs, effluent streams, lakes, or bays. Groundwater discharge may also occur at topographic low points in the form of seepage, which may be evident by marshes or shallow ponds.

In areas of discontinuous permafrost, localized zones of permafrost may control or influence groundwater movement. Permafrost may act as a barrier to downward percolation and to lateral movement of groundwater and also may act as a confining layer to subpermafrost water. The lower groundwater temperatures in Alaska also influence groundwater flow velocities. Groundwater existing under low temperature ranges (zero to 4.5°C) moves more slowly than groundwater in temperate regions due to increased viscosity.

Groundwater at King Salmon occurs in a near-surface shallow aquifer under unconfined conditions. The shallow aquifer is underlain by a low hydraulic conductivity unit termed the upper aquitard. Beneath the upper aquitard lies an intermediate aquifer where groundwater is present under confined conditions. In the shallow and intermediate aquifers beneath King Salmon Airport, groundwater flows to the south.

A more detailed discussion of the hydrogeology at King Salmon Airport is provided in Section IV.

2.3.2 Surface Water

Figure 2-5 shows the location and flow directions of the predominant surface water features in King Salmon, Alaska. In addition to those identified in Figure 2-5, there are several other unnamed tributaries with intermittent, seasonal flow. These tributaries ultimately discharge to the Naknek River.

Surface water discharge rates have been measured on Eskimo Creek and King Salmon Creek. Periods of measurement, mean annual flow, annual low and annual high flows for each creek are shown in Table 2-2.

Peak flows in Eskimo Creek appear to be associated with both snow-melt runoff and precipitation events. Minimum flows are probably also associated with drought periods between precipitation events and with winter low flows and may represent base flow contributions from the King Salmon shallow aquifer.

2.4 AMBIENT AIR QUALITY

King Salmon Airport is a remote military installation with a limited industrial base. Potential sources of airborne emissions in the King Salmon vici..ity include:

- Exhaust from gasoline, diesel, propane, and aircraft engines
- o Wood stoves
- o Solid waste burning
- o Emissions from cannery facilities

- o Construction
- o Sand dryer

With approximately 20 inches of rainfall per year and an annual average wind velocity of 9 mph, dilution and dispersion of airborne emissions results in overall acceptable air quality.

No records of previous or existing air sampling performed in the King Salmon vicinity are available.

| Table 2-2 SURFACE WATER FLOW DATA | | | | |
|--|--|---|--|--|
| Recording Period | Mean Recorded Discharge (cfs) | Minimum Recorded Discharge & Day of Occurrence (cfs) | Maximum Recorded Discharge (cfs) | |
| Eskimo Creek | | | | |
| Oct 29/87 Oct/83 - Sept/84 Oct/82 - Sept/83 Oct/81 - Sept/82 Oct/80 - Sept/81 Oct/79 - Sept/80 King Salmon Creek | 22.8 ^a 14.1 10.8 11.9 13.0 16.66 | 3.8 (Aug 06) 2.6 (July 20) 3.2 (Feb 18) 2.0 (Dec 13) 1.5 (Jan 13) | 94 (Jun 02) 58 (Oct 01) 103 (Jun 06) 68 (Mar 14) 53 (Nov 08) | |
| Oct 28/87 | 310.5 ^b | | | |
| ^a Discharge measure culvert east of E Appendix D-5.) ^b Discharge measure Bluff sites. (Se | ment performe skimo Creek. ment performe e data sheet | d 100 feet down (See data shee d between South Appendix D-5.) | stream from t and North Barrel | |

cfs = cubic feet per second

2.5 NATURAL RESOURCES

2.5.1 Mineral Resources

Active mining operations are not known to exist in King Salmon. An active sand and gravel operation is located north of King Salmon Airport. This facility produces sand, gravel, concrete,

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and other mixtures for construction and road surface preparation. Estimates of annual production and known reserves at the sand and gravel facility were not available.

2.5.2. <u>Environmental Resources</u>

King Salmon is situated on the Naknek River, a very productive fish and wildlife area, on the Alaska Peninsula. The installation is near the Katmai National Park and Preserve. The Alaska Peninsula is an important state-managed habitat for large and small game animals, fish, and birds (Engineering Sciences, 1985). Wildlife and wildlife habitat are considered an important natural resource in the area.

Commercial and sport fish and their aquatic habitat are important natural resources. The Naknek River and its tributaries (including Eskimo Creek and King Salmon Creek) support an enormous stock of migratory and resident sport and commercial game fish. The Naknek River drainage supports one of the largest sockeye salmon runs in the world. Between 1977 and 1981, the Naknek River system received an average annual return of 1.4 million sockeye salmon. Drainage from the King Salmon installation flows into the Naknek River and on to Bristol Bay, one of the world's most productive fishing grounds, where over 40 million sockeye salmon have been recorded in a single season. Commercial fishermen from Alaska and other states congregate annually in Bristol Bay to harvest salmon and other fish. Additionally, thousands of nonresidents pass through King Salmon on their way to fishing and cannery jobs at Bristol Bay (Alaska Department of Community and Regional Affairs, 1982).

In addition to sockeye salmon, all other species of Pacific salmon return to the Naknek River, King Salmon Creek, and Eskimo Creek. These include pink, chum, coho, and chinook salmon. Eskimo and King Salmon Creeks are utilized by all these salmon species for spawning and rearing, but are used primarily by chinook salmon. The salmon spawning runs in the area begin in late May and extend through late September. Other local fish include trophy-size rainbow trout, arctic char, Dolly Varden, grayling, lake trout, whitefish, burbot, and northern pike. Sport fishermen from around the world come to the Naknek River system to enjoy some of the best fishing in the world. The local economy derives much of its support from sport fishing along the Naknek River and its tributaries and from commercial fishing in Bristol Bay (USAF, 1987).

2.6 <u>CULTURAL RESOURCES</u>

2.6.1 <u>Archaeology</u>

There are many prehistoric archaeological sites along the Naknek River. One site in particular is located at the end of King Salmon's main runway, between the runway and the river bank. The site was excavated by archaeologists in 1973. The remains of a house and other artifacts, estimated to be 1,500 to 2,000 years old, have been discovered. Some local people observed that extensions of the runway had disturbed some native houses, and workers on the runway have reported seeing trade beads and other artifacts (Dumond, 1981).

Eskimo shelters dating back to the early 1900s have been found on the banks of Eskimo Creek. Other local prehistoric sites have been identified on the bank of the Naknek River across from the mouth of King Salmon Creek, and upstream on the Naknek River in the vicinity of the confluence of Big Creek with the Naknek River. Other prehistoric sites are scattered, and abut the banks of the Naknek River between Kvichak Bay and Naknek Lake. These sites are estimated to date back 2,000 years or more (Dumond, 1981).

2.6.2 <u>Human Resources</u>

People residing at the King Salmon installation are primarily military personnel or persons authorized by the military to be there. These personnel are on "unaccompanied assignments," meaning families remain at home away from the base. The human resources at the installation are, therefore, military and largely temporary (USAF, 1987).

Most of the nonmilitary residents of King Salmon hold jobs in government, transportation, or fishing. Government or the public sector is the primary employer of civilians in King Salmon. The FAA has employed in excess of 33 people. While the Air Force station does not contribute substantially to the local economy, some military personnel seek part-time employment with the canneries, restaurants, and flying services. The USFWS, NPS, and U.S. Postal Service are other major governmental employers in King Salmon. The State of Alaska employs seasonal and full-time workers in King Salmon. Most of these employees work for the ADFG, the DTPF, or the DPS.

Part of the civilian work force in King Salmon is engaged in commercial fishing activities. Additionally, various small enterprises provide companies, car rental businesses, fur trapping, and other miscellaneous small businesses.

2.6.3 <u>Existing Cultures</u>

Native groups around the King Salmon and Bristol Bay area include the Athabascan, Kiatagmiut, Aglegmiut, Peninsula Eskimo, and Aleut. The Aglegmiut occupy the lower half of the Naknek River and most of the coastline around the north end of Bristol Bay. The Peninsula Eskimo inhabit the base of the Alaska Peninsula including land along Shelikof Strait, and the upper half of the Naknek River upon which King Salmon is located (Dumond, 1981).

The Peninsula Eskimo natives of King Salmon often claim to be Aleut; however, their dialect is Sugpiaq, the language of the Peninsula Eskimo. Consequently, anthropologists generally consider these natives to be Peninsula Eskimo (Dumond, 1981). Census information gathered in 1980 by the U.S. Census Bureau reflects the natives' desire to be labeled Aleut. The 1980 census recorded two Eskimos and 27 Aleuts at King Salmon.

2.7 BIOLOGY AND ECOLOGY

2.7.1 Flora

The land around the Naknek River resembles the arctic zone of the nearby Katmai National Park and Preserve, but it is classified as transitional between subarctic forest and moist tundra. North of Naknek, trees are more common; but south of Naknek, trees gradually taper off. Lichen and sphagnum heaths are widespread, interspersed with lichens, crowberry, low birches, and marsh blueberry. Lake and river shores have communities of sedges, willow forms, reed grasses, bluejoint, white spruce, balsam poplar, taller birches, and alder (Dumond, 1981). Flowering plants include monkshood, lousewart, violet, buttercup, and other wild flowers (Alaska Department of Community and Regional Affairs, 1982).

2.7.2 <u>Fauna</u>

One of the most prevalent terrestrial mammals is the brown bear, which feeds during the summer months on salmon in the rivers. The brown bear retreats from the Naknek River vicinity in the fall to feed on berries in the high country and, subsequently, to hibernate for the long winter. Moose frequent the area also. In the summer, moose range throughout the region, favoring high, well-drained areas of willow and alder and areas along rivers where forage is good. Deep winter snows typically drive moose out of the high country into lower areas where travel is easier and food is more available.

Other land mammals in the region include caribou, red fox, weasels, mink, beaver, muskrat, lynx, wolves, porcupine, lemmings, shrews, voles, arctic ground squirrels, tundra hares, snowshoe hares, land otter, and caribou.

The Naknek River and its tributaries annually support migrations of anadromous fish. These include sockeye, chinook, silver, chum, and pink salmon species. Resident fish include trophy-size rainbow trout, arctic char, Dolly Varden, grayling, lake trout, whitefish, burbot, and northern pike. King Salmon is a base for world-class sport fishing, and nearby Bristol Bay fosters one of the most productive commercial salmon fishery in the world.

Hundreds of thousands of waterfowl stop at Bristol Bay while migrating to and from distant nesting grounds. Most of the world's black brant population and virtually all of the world's population of emperor and Canada geese fly through the region. Mallards, gadwalls, pintails, green-winged teal, American widgeons, greater scaup, goldeneyes, buffleheads, oldsquaw, harlequins, scoters, and other ducks are known to nest at King Salmon. The Kvichak River is a major flyway for sandhill cranes and whistling swans. Snow geese, whitefronted geese, Canada geese, and loons stop there during their seasonal migrations. Some birds remain at King Salmon to nest and molt. Recent surveys have recorded 141 species of waterfowl, with a breeding population of 32 ducks and 1.2 swans for every square mile of lowlands.

Shorebirds use the same habitats and flyways as other waterfowl. They include rock and western sandpipers, ruddy turnstones, bar-tailed godwits, golden and black-billed plovers, dunlins, and phalaropes. Shorebirds nest on lake shores, coastal beaches, and tundra hummocks. Other species found on coasts and inland waters include glaucous-winged and mew gulls, arctic terns, and black-legged kittiwakes.

The protected bald eagle nests along the Naknek River and its tributaries. Other raptors that frequent the area include American and arctic peregrine falcons, ospreys, and short-eared owls. Many passerines, such as Lapland longspurs, snow buntings, swallows, Savanna sparrows, and dippers, are also present.

Wintering birds at King Salmon include white-tailed, willow, and rock ptarmigan and ravens. Seabirds move offshore in the winter, while most of the other bird species migrate to warmer climates (Alaska Department of Community and Regional Affairs, 1982).

2.7.3 Environmentally Sensitive Areas

The Alaska Peninsula supports a diverse variety of fish, birds, small game, and large game animals. The area is a state-managed habitat for wildlife and is considered to be one of the most significant in the state (Engineering Science, 1985).

King Salmon is located on the bank of the Naknek River. This river and its tributaries (including Eskimo Creek and King Salmon Creek) are rich in sport and commercial fish and ecologically link the project site with Bristol Bay. The Naknek River is one of Alaska's most important sport fishing areas. Drainage from the military installation at King Salmon is conveyed by King Salmon Creek, Eskimo Creek, and the Naknek River to Bristol Bay, site of one of the state's most productive commercial fisheries. The land, aquatic, and marine ecosystems are, therefore, interrelated. Activities at the King Salmon installation can affect the local environment, Naknek River, Eskimo Creek, King Salmon Creek, and possibly downstream environments (Bristol Bay). Because of Alaska's delicate environmental setting, the long-term impacts of pollutant discharge in the area could be severe. The potential impacts on public health are uncertain (Engineering Science, 1985).

2.7.4 <u>Endangered Species</u>

The project area is within the nesting range of the American peregrine falcon and is in the migratory range of the arctic peregrine falcon. Both are considered endangered in Alaska.

2.8 CLIMATOLOGY / METEOROLOGY

Because of its enormous size, Alaska spans four distinct climatic zones, each of which has been established on the basis of temperature and precipitation characteristics. These areas include the arctic, continental, transitional, and maritime zones (Engineering Sciences, 1985). Figure 2-6 shows the location and boundaries of each climatic zone type. Because of its inland location, King Salmon lies on the border between the transitional and continental zones. The Alaska Peninsula has a subpolar maritime climate characterized by high winds, protracted cloud cover, frequent precipitation, and fog.

Rainfall varies significantly in Alaska: 5 inches fall during a typical year in the arctic climatic zone, while 300 inches fall during the average year in the southeastern maritime zone. This considerable spatial variation in rainfall across the state is due primarily to orographic



effects related to topography and exposure. Coastal mountainous areas receive the most rainfall and interior lowlands receive the least (Engineering Sciences, 1985).

Climatic data for King Salmon are summarized in Table 2-3. Climatic variables presented include temperature, precipitation, snowfall, and surface winds.

| Table 2-3 CLIMATIC DATA FOR KING SALMON AIR FORCE STATION | | | |
|---|----------------|--|--------------------------------|
| | Mean Annual | Monthly Recorded Minimum | Monthly Recorded Maximum |
| Temperature | 33°F | -46°F (January) | 88°F (June) |
| Precipitation | 19.8 inches | 0 inches (March, April November) | 7.3 inches (September) |
| Snowfall | 46 inches | 0 inches (March, April November) | 20 inches (March) |
| Surface Winds | 9 mph | 0 | A |

Note: A=High winds associated with severe storms can occur at any time. These high winds are called williwaws. Source: Engineering Science, 1985.

2.81. Precipitation

Precipitation at King Salmon is slightly variable. The average annual precipitation at King Salmon is 19.8 inches. The months of July through October are typically the wettest, averaging above 2 inches of rainfall per month. The maximum monthly precipitation recorded was 7.3 inches in September. The months of March, April, and November have each passed at one time without measurable precipitation being recorded. The maximum 24-hour precipitation was 1.7 inches durint the month of September (11th Weather Squadron, Elmendorf AFB, AK, 1942-1982).

Winter snow depths average 46 inches annually. The highest monthly snowfall depth ever recorded was 20 inches in March. The maximum 24-hour snowfall depth was 11 inches during the month of March (Engineering Sciences, 1985). The position of the modern snowline is determined by the lowest point of firm snow as derived from 1:63,360 scale maps and aerial photographs and from the point where medial moraines on valley glaciers are covered by snow. These data indicate the snowline is lowest, 900 m, along the Pacific side of the peninsula and rises westward across the peninsula by about 100 m every 20 km.

2.8.2 <u>Temperature</u>

The weather at King Salmon is seasonally mild because of its proximity to the coast. The mean annual temperature is 33°F. Monthly average temperatures vary from 55°F in July to 12°F in December. The highest temperature ever recorded in King Salmon was 88°F in June; the coolest temperature was -46°F in January (Engineering Sciences, 1985).

2.8.3 <u>Wind</u>

2.8.3.1 Direction

Wind prevails typically from the north during the winter months and from the south to east in the summer. The annual prevailing wind direction is from the north.

2.8.3.2 <u>Wind Speed/Velocity</u>

Wind speeds at King Salmon are characteristically mild. The annual average wind speed is 9 miles per hour.

2.8.3.3 <u>Seasonal Variability</u>

Average monthly wind speeds are relatively uniform throughout the year. Wind speeds vary on the average between 8 and 10 miles per hour during any month of the year (Engineering Sciences, 1985). However, severe storms are common and can occur at any time, often accompanied by high winds known as "williwaws." These storms are spawned by the Aleutian Low Pressure area, and their northeastward circulation path takes them along the southeast side of the peninsula on their way across the north Pacific to Canada and the northwestern United States.

2.8.4 Evapotranspiration

Information on the seasonal variation of evapotranspiration is not available for the King Salmon area; however, the potential evapotranspiration has been estimated at 17.87 inches. The actual evapotranspiration has been estimated at 15.98 inches (Patrick, et. al., 1968).

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III. FIELD INVESTIGATION PROGRAM

3.1 INTRODUCTION

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Detailed planning efforts concerning the execution and control of the Stage 1 field activities were conducted during September 1987. The results of these efforts were the Work Plan, Quality Assurance Project Plan, and the Site Safety Plan. These documents are referred to frequently in the following text. Exceptions to the documents are discussed in detail.

The field investigation program for this study consisted of the following tasks:

- o Soil gas survey
- o Geophysical survey
- o Monitoring well installation
- o Environmental sampling
- o Aquifer testing

Before proceeding with any task, the King Salmon AFS Point of Contact was notified. Base Civil Engineering Work Clearance Requests, AF Form 103, were prepared, submitted and approved for all soil boring/monitoring well locations.

3.2 SOIL GAS SURVEY

The soil gas survey was conducted on October 21, 1987 through October 31, 1987 by Tracer Research Corporation of Tucson, Arizona under the supervision of a CH2M HILL geologist. Drilling support was provided by M-W Drilling, Inc. of Anchorage, Alaska. Soil gas surveys provide a relatively inexpensive way of identifying areas containing volatile and semivolatile organic contaminants occurring in the soil or which have migrated upward through the soil from underlying contaminated groundwater.

Soil gas samples were collected by driving a hollow steel probe to a depth between 5 and 35 feet below the ground surface. The majority of samples were collected within a few feet of the water table. The probes were driven using the cat-head on a truck-mounted Acker MP4 drilling rig. Probes consisted of either 1.5-inch-diameter steel pipe or E-size drill rods. A volume of gas equal to 5 to 10 times the volume of the probe was evacuated using a vacuum pump. During the soil gas evacuation, samples were collected by inserting a syringe needle through a silicone rubber segment in the evacuation line down into the steel probe. The soil gas samples were analyzed in the field with a Varian 3300 gas chromatograph equipped with a flame ionization detector and a Spectra Physics 4270 computing integrator. The samples were analyzed for benzene, toluene, xylene, and total hydrocarbons.

3.2.1 Eskimo Creek Seep

The soil gas survey at the Eskimo Creek seep site consisted of 49 probes advanced in the general area of the installation structures located west of Eskimo Creek. The locations of the soil gas probes are shown in Figure 3-1, Eskimo Creek Seep Soil Gas Probe Locations. The purpose of the soil gas survey in this area was to locate "hot spots" of volatile organic contamination and assist in determining locations for the monitoring wells.

3.2.2 <u>Naknek River Seep</u>

The soil gas survey at the Naknek River seep site consisted of two probes advanced near the seep. The purpose of the soil gas survey in this area was to assist in determining the lateral extent of contamination associated with the Naknek River seep. Three additional probes were advanced along the highway between the west entrance to the installation and the entrance to the Naknek River tank farm. The purpose of the soil gas survey in this area was to assist in determining whether or not POL tanks 11 through 14 were a possible source for the Naknek River seep.

The soil gas survey for the Naknek River seep was done on a time remaining basis and was not the primary purpose for use of the soil gas survey techniques at King Salmon AFS. The locations of the probes are shown in Figure 3-2, Naknek River Soil Gas Probe Locations.

3.2.3 North Barrel Bluff

No soil gas surveys were conducted at the North Barrel Bluff site.

3.2.4 South Barrel Bluff

No soil gas surveys were conducted at the South Barrel Bluff and Landfill No. 1 site.

3.3 <u>GEOPHYSICAL SURVEY</u>

The geophysical survey was conducted October 21 through 31, 1987 by NORTEC, a division of ERT, of Anchorage, Alaska, under the supervision of a qualified CH2M HILL geologist. The following complementary geophysical techniques were used:

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- Electromagnetic profiling using a Geonics EM-31DL
 Terrain Conductivity Meter with digital recorder
- Magnetic gradiometer surveying using an EDA OMNI IV Magnetometer/Gradiometer
- Ground penetrating radar (GPR) profiling using a GSSI System-GPR with a 120-MHz antenna

A detailed description of these techniques and their application to hazardous waste site investigations is provided by Benson (1982). Geophysical survey lines were established using fiberglass tapes, a Brunton compass, and a right-angle prism. Survey lines were tied to several BLM monuments, building corners, and/or subdivision rebar corner posts. Distances along survey lines are taped distances rather than true horizontal distances. The level of accuracy achieved was considered sufficient for the reconnaissance purposes of the geophysical investigation.

3.3.1 Eskimo Creek Seep

At the Eskimo Creek site, the investigations were directed toward identifying the source or sources of the contaminants observed at the Eskimo Creek seep. The following survey objectives were outlined:

- o Characterize subsurface soils
- o Map contaminant plume, if possible
- Explore for utilities and debris
- Check proposed monitoring well locations for hazards

Geophysical investigations conducted at this site were restricted to the use of the ground penetrating radar system due to cultural interferences (buildings, powerlines, radar dome, etc.) with the electromagnetic and magnetic methods. The radar dome proved to be the greatest interference and virtually eliminated the EM meter from the survey.

Two GPR grids were surveyed at the Eskimo Creek seep site. The geophysical survey grids are shown in Appendix I, Plate 4, Ground Penetrating Radar Trackline Map, Headquarters Area. The first was a detailed grid with a 10-foot line spacing in the vicinity of Buildings 636, 632, and 624. This area was reported by Air Force personnel to have been the location of a previous service station. The purpose of the geophysical survey in this area was to search for a possible buried fuel tank which may have been left in place when the service station was removed. The larger grid was surveyed along several lines that paralleled the soil gas probe transects. The purpose of the geophysical survey in this area was to attempt to characterize subsurface soils, to search for evidence of a contaminant plume, and to explore for buried utilities and debris.

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3.3.2 Naknek River Seep

Initially, a geophysical survey at the Naknek River seep Site was not planned. However, due to the reported presence of rows of buried drums at the site, a brief geophysical survey using both the Geonics EM-31DL Terrain Conductivity Meter and the OMNI IV Magnetometer/Gradiometer was performed. The survey grid is shown in Appendix I, Figure 2, Naknek River Tank Farm, Geophysical Anomaly Map. Instrument readings were taken at 25-foot intervals along the survey lines.

3.3.3 North Barrel Bluff

The geophysical investigation at the North Barrel Bluff site was directed primarily at defining the extent of the waste disposal areas. Most of the investigation was conducted using the conductivity meter and the magnetometer/gradiometer.

Most of the geophysical survey lines were conducted perpendicular to the bluff. Instrument readings were recorded at 25-foot intervals along the survey lines. Survey lines were carried down the face of the bluff when possible and onto the flood plain of King Salmon Creek. Many geophysical lines were terminated at the crest of the bluff because continuation of the line down the face of the bluff would require direct contact with the barrels, violating the site safety plan.

The location of the North Barrel Bluff survey lines are shown in Appendix I, Plate 2, Geophysical Anomaly Map, North Barrel Bluff Site.

Geophysical surveys were conducted at two other reported waste disposal areas near the North Barrel Bluff. The areas are shown in Appendix I, Plate 2, Geophysical Anomaly Map, North Barrel Bluff Site. The first was an area of disturbance south of the road between the north and south barrel bluffs. From aerial photographs, the area appears to have been cleared. The area was reported by a local contractor to have been a waste disposal area. The second area, southeast of the large existing building at the North Barrel Bluff, reportedly was used as a disposal area for debris resulting from demolition of several buildings in this vicinity. An abandoned buried tank was also reported to exist in this area.

Both areas were surveyed with the magnetometer and the conductivity meter.

3.3.4 South Barrel Bluff

The purpose of the geophysical survey at the South Barrel Bluff and Landfill No. 1 site was the same as for the North Barrel Bluff. A more closely spaced grid was established in this area because it is currently being developed as a residential housing area. Two houses have already been constructed within the zone of suspected waste disposal.

Most of the South Barrel Bluff survey lines were oriented in a north-south direction at approximately 45-degree angles to the crest of the bluff. The lines were surveyed with both the conductivity meter and the magnetometer/gradiometer. Instrument readings were recorded at 25-foot intervals.

The locations of the survey lines are shown in Appendix I, Plate 3, Geophysical Anomaly Map, South Barrel Bluff Site.

3.4 SUBSURFACE EXPLORATION AND MONITORING WELL CONSTRUCTION

The subsurface exploration was performed on November 4, 1987 through December 11, 1987 and on May 1, 1988 through June 10, 1988 by M-W Drilling, Inc. of Anchorage, Alaska, under the supervision of a qualified CH2M HILL geologist. Shallow borings were drilled with a truck-mounted Acker MP4 drilling rig equipped with 6-inch inside-diameter hollow-stem augers. Deep borings were drilled with a Schramm TA-60T truck-mounted air rotary drilling rig. Monitoring wells were constructed by M-W Drilling, Inc. under the supervision of a qualified CH2M HILL geologist. Monitoring well construction details are shown in Figure 3-3.

Initially, the subsurface exploration and monitoring well construction was scheduled to be completed in the fall of 1987. After the construction of 12 of the monitoring wells, work was suspended due to extremely cold weather and the nonavailability of an equipment decontamination area. The remainder of the work was completed in the spring of 1988.

Working in extremely cold weather presented unique problems such as:

 Organics in the soil did not volatize and therefore were not detected by the organic vapor analyzer (OVA).



- o Removing frozen soil from the augers and drilling equipment approximately doubled the time required for decontamination.
- Water and bentonite slurry could not be brought to the drill site more than 10 to 15 minutes prior to use. Hoses and pumps had to be returned promptly to the heated decontamination area to avoid freezing and subsequent damage.

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o During decontamination, the area became filled with steam, causing visibility problems. The doors to the area could not be left open because of the extreme cold.

Decontamination of the back of the drill rig, the augers and drill rods, soil sampling equipment, and all tools and equipment was done in accordance with the Statement of Work, Section 5.2.1.5 of the Work Plan and Section 1-7 of the QAPP. After completion of wells in visibly contaminated areas, all drilling equipment was washed with detergent prior to steam cleaning. (Detailed decontamination procedures are described in the QAPP, Section 1.7, and in the Sampling Plan.)

Subsurface soil samples were collected in accordance with the Statement of Work, Section 2.1 of the Work Plan, and Section 5.2.1.5 of the QAPP. Samples were collected at 5-foot intervals above the water table and at points of noticeable change in subsurface materials below the water table in all shallow borings.

Samples were obtained in general accordance with ASTM D1586, Penetration Testing and Split-Barrel Sampling. Exceptions were as follows:

- A 2-1/2-inch inside diameter sampler was used instead of a 1-3/8-inch inside diameter
- o A 330-pound hammer was used instead of a 140-pound hammer

Samples were classified in general accordance with ASTM D2487. Sample classification and the results of the penetration test are shown in the boring logs presented in Appendix C-1.

Split-spoon samples were attempted in the deep borings but no samples were recovered because sample materials were primarily loose, saturated sands that could not be retained by the sampler. The sampler was pushed with the hydraulics on the drill rig instead of being driven with a hammer. Penetration test results were not obtained.

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The split-spoon sampler was decontaminated in accordance with Section 2.1 of the Work Plan and Section 5.2.1.5 of the QAPP.

The monitoring wells were developed by bailing or pumping. Development of the wells continued until the temperature, pH, and specific conductivity measurements stabilized and turbidity was reduced. At least 10 well volumes of water were removed from each of the wells. Development records are provided in Appendix D-3.

Development water from wells suspected to be contaminated was collected in a 1,000-gallon water tank. Sampling and disposal of the development water is discussed under Section 3.9.1

3.4.1 Eskimo Creek Seep

A total of 21 soil borings, 18 shallow and 3 deep, were drilled at the Eskimo Creek seep site. Monitoring wells were constructed in all of the soil borings except MW-35. When drilling the boring for MW-35, an abandoned fuel line was encountered. The boring was therefore abandoned. The locations of the monitoring wells are shown on Figure 3-4.

Shallow wells were constructed at the Eskimo Creek seep site in order to assess the extent of groundwater contamination in the upper aquifer and to further define the hydrogeology of the site. Deep wells were constructed for the purpose of exploring the intermediate aquifer for possible contamination. Deep wells were drilled next to shallow wells in order to determine the vertical gradient and assess the vertical extent of the contamination.

Initially, a shallow and deep well were to be constructed near both of the onsite production wells, KS4 and KS5. This nest of three wells, screened in the upper, intermediate, and deep aquifers, was intended to aid in determining the vertical gradients between the three aquifers. MW-31 and MW-41 were completed near KS5. A second nest was to consist of MW-35 and MW-42. However, during the drilling of MW-35, an abandoned 2-inch-diameter steel fuel line was encountered and ruptured at about 5 feet below the ground surface. The rupture was not discovered until the boring had been drilled to a depth of about 38 feet. The amount of fuel lost was estimated to be about 75 gallons. After base personnel were notified and the rupture repaired, the boring was abandoned by backfilling with tremied neat cement grout.

MW-42 was then moved to nest with MW-33. After completion of drilling at MW-42, the drilling subcontractor was unable to extract the temporary 8-inch-diameter casing. Because of

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damage to the end of the 8-inch casing during drilling of MW-41 without a drive shoe, the drilling subcontractor welded a drive shoe to the 8-inch casing prior to beginning drilling of MW-42. The difficulty in extracting the temporary 8-inch casing may be attributed to the drive shoe. A tool designed to cut the drive shoe from the casing was air freighted from Anchorage, Alaska. Attempts to cut the drive shoe off were unsuccessful. The drilling subcontractor still could not extract the casing even when using jacks capable of delivering 300 tons of lifting force to the casing. A joint decision between the CH2M HILL project hydrogeologist and OEHL was made to construct the well inside of the 8-inch casing. Well construction details for MW-42 are shown in the well construction log in Appendix C-1.

As specified in the Statement of Work, cuttings from MW-30, MW-31, MW-39, MW-40, MW-41, and MW-43 were contained in drums in compliance with <u>Test Methods for Evaluating Solid</u> <u>Waste Physical/Chemical Methods, SW-846, 3rd Edition</u> (US EPA, 1986). Cuttings from these wells showed visible signs of contamination and significant levels of contamination were indicated by the organic vapor analyzer.

3.4.2 Naknek River Seep

No borings or wells were completed at this site.

3.4.3 North Barrel Bluff

No borings or wells were completed at this site.

3.4.4 <u>South Barrel Bluff</u>

No borings or wells were completed at this site.

3.5 MONITOR WELL PROTECTIVE CASINGS, AND GUARD POSTS

Six-inch-diameter, locking, steel protective casings were cemented in place around the top of each 2-inch-diameter monitoring well. The protective casings extend to a depth of 3.5 feet below ground surface in MW-23 through MW-26 and to a depth of 5 feet below ground surface for the remaining monitoring wells. This additional depth was recommended due to the potential frost heave problems.

Monitoring well protective casings for MW-23 through MW-26 originally extended to a depth of 1 foot below ground surface. Frost heave damage was observed in the Spring of 1988. The protective casings had become loose and unstable. These protective casings were removed and reset at a 3.5-foot depth in 1988. The contractor was unable to remove concrete belo a depth of about 3.5 feet. Damage due to frost heave may still occur at these wells. To minimize damage, the PVC

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well casing was wrapped with polytheylene sheeting to prevent the concrete from adhering to the well casing.

During placement of the concrete for the surface seal and protective casing installation, a 2-foot by 2-foot cement surface pad was also constructed, using temporary forms to shape the concrete pads.

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Three guard posts were installed around each well where vehicular damage was considered probable by the King Salmon AFS Point of Contact. The guard posts were constructed from 3-inch-diameter steel pipe. The pipe was placed in a 5-foot hole and backfilled with concrete. Approximately 2.5 feet extended above the ground surface. The pipes were then filled with concrete.

Five guard posts were placed around the three shallowintermediate monitoring well nests.

Twelve-inch-diameter, locking, steel protective casings were placed around each 4-inch-diameter monitoring well. These protective casings also serve as a component of the aquifer seal and extend to depths ranging from 35 to 52 feet. Protective casings were painted bright yellow with the monitoring well identification painted on in black on the side of the protective casing.

3.6 CONSTRUCTION OF AQUIFER SEAL FOR DEEP MONITORING WELLS

Two King Salmon AFS production wells are located in the vicinity of shallow aquifer wells in which floating fuel product has been detected. Intermittent pumping of the production wells may result in a hydraulic head reduction in the deep wells, potentially causing downward leakage of these contaminants into lower aquifers. To determine whether there has been vertical movement of these contaminants from the shallow aquifer into deeper aquifers, three deep monitoring wells were installed at the King Salmon AFS site. These wells were screened in an intermediate aquifer that lies between the shallow aquifer and the deep aquifer in which the production wells are screened. To minimize the potential for migra-tion of contaminants down the well during the installation procedure and after the installation was complete, an aquifer seal was constructed in the upper aguitard that lies between the shallow and intermediate aquifers. Figure 3-5 depicts the construction of the aquifer seal.

The construction of the aquifer seal was perfore using the following methodology. A 12-inch-diameter cased worehole was advanced through the shallow aquifer into the upper aquitard. Drill cuttings from the upper aquitard were monitored for the presence of organic vapors using an OVA and/or HNU photoionization detector. The OVA was calibrated to methane



equivalents and the HNU was calibrated to benzene equivalents. The 12-inch-diameter cased borehole was advanced into the upper aquitard to a final depth at which OVA or HNU measurements on drill cuttings no longer indicated the presence of organic constituents.

The 12-inch borehole was then overreamed below the cased portion of the borehole creating a larger-diameter borehole. The overreaming was completed to a depth approximately 2 feet below the bottom of the 12-inch casing. The borehole was then dewatered and the drill bit and drill pipe removed from the hole. A shrinkage-compensated cement mixture consisting of the following components by approximate percent weight: water 33 percent, Portland Cement 61 percent, gypsum 5 percent, and bentonite 1 percent, was mixed and tremied into the bottom of the borehole. This sealing mixture typically filled the borehole from the bottom to a depth of 6 to 8 feet. After the sealing mixture was tremied into place, the 12-inch casing was advanced into the overreamed portion of the borehole. The seal was then allowed to stabilize for approximately 12 hours.

Construction of the 8-inch borehole then proceeded inside of the 12-inch casing. At total depth, the 4-inch well was built inside the annular space.

Upon completion of the monitoring well, the 8-inch casing was removed and the 12-inch casing left in place.

3.7 SURVEYING

At the conclusion of the Stage 1 field effort, all newly installed monitoring wells, existing USCOE wells, landfill wells, and the two King Salmon AFS production wells were surveyed for horizontal location and elevation to the King Salmon AFS site datum.

The surveying was performed by Coastal Surveyors/Ralph Mancuso and Associates, Naknek, Alaska. Locations for each monitoring well were established by measuring horizontal and zenith angles using an electronic theodolite, and measuring ground distances using electronic distance measuring equipment. Angles and distances were measured between points of known location and each monitor well location. This data was then trigonometrically transformed using standard survey data reduction techniques into a north and east coordinate location. Coordinate locations are given to the nearest 0.01 of a foot.

Elevations were determined for each monitoring well at a position marked at the top of the well casing. Elevation differences between known points and each monitoring well were measured using a differential leveling technique. The

elevation differences were then arithmetically summed to obtain the measuring point elevation for each well. Elevations were calculated to the nearest 0.01 of a foot.

A coordinate and elevation listing for the King Salmon AFS monitoring wells is provided in Appendix E.

3.8 SAMPLING PROGRAM

The surface water and sediment sampling began on October 28, 1987, and was completed on November 2, 1987. Groundwater sampling began on June 5, 1988, and was completed on June 17, 1988. All sampling was performed by trained, experienced CH2M HILL personnel. The location, type of analyses, and number of samples were specified in the SOW. The actual number of samples collected at each site is summarized in Table 3-1, Number of Analyses by Site.

Sampling methodology and protocols are indicated on the analytical reports, in Appendix G, Analytical Data. Sample preservation methods, required containers, and holding times are shown in Table G-2, also in Appendix G.

Sampling was done in accordance with Annex C of the Statement of Work, Sections 1.5, 1.6, and 1.7 and Appendix A, Project Sampling Plan, of the Quality Assurance Project Plan, and Sections 5.2.1.4 and 5.2.1.5 of the Project Work Plan. Exceptions were as follows:

- o A stainless steel bailer with a teflon ball valve was used to sample the USCOE wells. A 1-inchdiameter teflon bailer was not available. The use of the stainless steel bailer should have no affect on the results of the analyses performed on samples from the USCOE wells.
- Sampling of MW-6 and MW-23 was not performed immediately after purging. These wells were purged dry and were allowed to recover overnight prior to sampling. MW-1 did not recover after purging and was not sampled.

Samples were shipped under full chain-of-custody procedures by air freight to the laboratory. Copies of the completed chain-of-custody forms are included in Appendix F, Chain-of-Custody Forms.

Field quality assurance/quality control samples were as specified in Statement of Work, Appendix B, and in Section 1.4 of the QAPP. Exceptions were as follows:

• Trip blanks were not sent with every cooler containing volatile organic compound samples. Had this

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| Parameter | Analytical Method | Eskimo Creek | USCOE Well | Naknek River Seep | North Barrel Bluff | South Barrel Bluff | Offsite Well | Development Water |
|--|----------------------|-----------------|---------------|-------------------------|--------------------------|--------------------------|-----------------|----------------------|
| Water Samples | | | | | | | | |
| Alkalinity - Carbonate, Bicarbonate, & Hydroxide (Field Test) | A403 | 23 | 7 | 1 | 3 | 3 | | |
| Common Anions (Chloride, Fluoride, Nitrate, Sulfate, Orthophosphate) | A429 | 23 | 7 | 1 | 3 | 3 | | |
| Specific Conductance (Field Test) | E120.1 | 23 | 7 | 1 | 3 | 3 | ~= | |
| pH (Field Test) | E150.1 | 23 | 7 | 1 | 3 | 3 | | |
| Total Dissolved Solids | E160.1 | 23 | 7 | 1 | 3 | 3 | | |
| Temperature (Field Test) | E170.1 | 23 | 7 | 1 | 3 | 3 | | |
| Lead | SW3005/ SW7421 | 23 | 7 | | | | | 2 |
| Metal Screen (25 metals) | E200.7 | 2 | 0 | 1 | 3 | 3 | 1 | |
| Petroleum Hydrocarbons | E418.1 | 21 | 7 | | | | | |
| Purgeable Halocarbons | E601 | 2 | | 1 | 3 | 3 | 1 | |
| Organochlorine Pesticides | E608 | | | | 3 | 3 | 1 | |
| Extractable Priority Pollutants | E625 | | | | 3 | 3 | 1 | |
| Purgeable Aromatics | SW5030/ SW8020 | 16 | | 1 | 3 | 3 | 1 | 2 |
| Soil Samples | | | | | | | | |
| Petroleum Hydrocarbons | SW3550/ E418.1 | 18 | | | | | | |
| Organochlorine Pesticides and PCBs | SW3550/ SW8080 | | | | 3 | | | |
| Volatile Organic Compounds | SW5030/ SW8240 | 5 | | | 3 | | | |
| EPTOX Metals | 40 CFR 261.24 | 5 | | | | | | |
| Semivolatile Organic Compounds | SW3550/ SW8270 | 5 | | | 3 | | | |
| Soil Moisture Content | ASTM D2216 | 23 | ~- | | 3 | | | |

| | 1 | Table 3-1 | | |
|--------|----|-----------|----|------|
| NUMBER | OF | ANAL YSES | BY | SITE |

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been done, the number of analyses would have exceeded the maximum number of analyses specified in Tables A-4 and A-5 of the Statement of Work.

Equipment blanks were not collected for every day of groundwater sampling. Had this been done, the number of analyses would have exceeded the maximum number of analyses as specified in Tables A-4 and A-5 of the Statement of Work. Decontamination and sampling procedures were consistent throughout the groundwater sampling program. The results of the analyses on the equipment blanks are likely to be representative.

3.8.1 Sampling of Drummed Soil Cuttings and Development Water

In accordance with the Statement of Work, Section 5.2.1.5 of the Work Plan and Section 2.2 of the QAPP, the drums containing contaminated cuttings from the soil borings were sampled and tested for EP Toxicity Metals 40 CFR 261.24, volatile organic compounds, and semivolatile organic compounds. Exceptions were as follows:

- o The drilling subcontractor supplied ten new DOT approved drums. Additional drums were obtained, with permission, from the King Salmon AFS Point of Contact. These drums had been used but appeared to be clean and in good condition. If the contents must be disposed of as hazardous waste, they may have to be transferred to new drums prior to shipment.
- Composite samples, one from each drum were not collected. In all, 21 drums were filled with contaminated soil from MW-30, MW-31, MW-38, MW-39, MW-41, and MW-43. Each drum could not be sampled without exceeding the maximum number of analyses (5) as specified in Table A-5 of the Statement of Work. If the drums must be disposed of as hazardous waste, the cost of retesting each drum within a group should be compared with the cost of disposing of all the drums. One highly contaminated drum within a group may indicate that the whole group must be disposed of as hazardous waste.

The drums from each well were grouped into one of five groups. Drums from MW-31 and MW-41 were grouped together because these two well are less than 10 feet apart. One composite sample for each group for a total of five samples was prepared. Drums were sampled by compositing discrete samples taken from the top to the bottom of each drum. The composite sample from each drum was composited with other samples

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collected from drums in the same group to form one composite sample per group of drums. The results of the analyses are presented in Section 4.1.4, Soils Investigation. The soils were determined not to be RCRA hazardous waste and were transferred to base engineering for ultimate disposal.

In accordance with the Modification to the Statement of Work, the development water was sampled and tested for purgeable aromatic compounds and lead. As discussed in Section 3.4, the development and purge water was contained in a 1,000-gallon water tank. After the tank was full, the water was sampled by dropping a 5-foot length of tygon tubing through the opening at the top of the tank. After the tube had filled, the top of the tube was plugged and the tube withdrawn. Care was taken to avoid sampling the sediment in the bottom of the tank or the floating product. The results of the analyses are presented in Appendix G, Analytical Data.

3.9 LABORATORY PROGRAM

All analyses were conducted in accordance with the standard methods specified in the Statement of Work. The CH2M HILL environmental laboratory in Corvallis, Oregon, was responsible for overall sample handling, custody, and documentation for the samples collected in the fall of 1987. The CH2M HILL environmental laboratory in Redding, California, was responsible for the samples collected in the spring of 1988. Analyses for total petroleum hydrocarbons were subcontracted to Technical Services, Inc., in Jacksonville, Florida.

Laboratory quality assurance/quality control data are presented in Appendix G, Analytical Data.

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IV. RESULTS AND SIGNIFICANCE OF FINDINGS

4.1 DISCUSSION OF RESULTS

4.1.1 Soil Gas Investigation

This section summarizes the results of the soil gas investigation performed at the Eskimo Creek seep and Naknek River seep sites. The soil gas report is attached as Appendix I.

4.1.1.1 Eskimo Creek Seep

Maps showing soil gas sampling locations and concentrations for total hydrocarbons (Figures 3-1 and 4-1) are attached. Total hydrocarbon values represent the C3-C9 range reported in the attached soil gas report. This range is a hydrocarbon group that is analyzed by Tracer Research and is a function of the calibration of their flame ionization detectors.

Figure 4-1 indicates the distribution of total hydrocarbons in the soil gas. The highest concentration detected was at sampling location SG-6 (820 μ g/l total), northeast and east of POL Tanks 11 and 12.

The extent of contamination from this point appears to include an area southwest of the tanks (west soil gas plume) and a much larger area southeast that includes the Eskimo Creek seep (east soil gas plume). This plume can be traced from the POL tanks through the center of the station to the Eskimo Creek seep. The plume also includes a large area that extends north of the Eskimo Creek seep.

Isolated areas of contamination around sampling locations SG-40, SG-41, and SG-25 appear to be unrelated to the main soil gas plumes. These locations may be isolated sources of groundwater or soil contamination from historically different events.

4.1.1.2 Naknek Seep

Two soil gas samples were analyzed at the Naknek seep (Figure 4 in Appendix I). Sample SGNS-1 located near the Naknek River bank showed a concentration of 2,200 µg/l total hydrocarbons. Sample SGNS-2 was located approximately 400 feet from the riverbank and did not contain detectable hydrocarbons (<0.03 µg/l). Three additional probes located along the abandoned pipeline between the station and Naknek River seep were also at less than detection levels.

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4.1.2 <u>Geophysical Investigation</u>

This section summarizes the results of the 10-day reconnaissance geophysical investigation conducted at King Salmon AFS. Four sites--Eskimo Creek seep, Naknek River seep, North Barrel Bluff, and South Barrel Bluff--were surveyed using combinations of electromagnetic conductivity, ground penetrating radar (GPR), and magnetics. The report is attached as Appendix I. Results for each site are described below.

4.1.2.1 Eskimo Creek Seep

The detailed GPR grid survey over a former service station site (in the vicinity of Buildings 636, 632, and 624) failed to yield any strong radar reflections characteristic of a buried tank. A few hyperbolic reflections were identified, but these are believed to be caused by buried utilities rather than a large buried tank. The locations of these radar targets are indicated in Plate 4 (Appendix I), as well as on the radar records in Plates 15-26. Of these targets, the reflection most suggestive of a buried tank is a target about 10 feet in depth at Station 113N on Line 80E (Plate 23). This feature lies in the center of a 20-foot-wide zone suggesting trenching or disturbance, and Line 70E (Plate 22) just 10 feet to the west shows evidence of a subsurface slab that may be a concrete slab.

A ground penetrating radar (GPR) survey grid was conducted along soil gas probe transects on the Eskimo Creek seep bluff in the hope of providing useful soils data to correlate with the soil gas analyses. Cases are reported in the literature where it has been possible to detect the presence of hydrocarbon contamination in soils by observing a change in the character or "texture" of the GPR records caused by a contrast in subsurface dielectric properties in the zone of contamination. There have also been some sites where petroleum products floating on the water table have resulted in particularly strong radar reflections ("bright spots") at the water table der.

The Headquarters Area GPR records were searched for changes in reflection signal "texture" that might be indicative of zones - contamination, particularly in the areas that indicated high soil gas hydrocarbon concentrations. Such GPR contrasts were not observed. Neither were strong water table reflections observed that might be indicative of the presence of hydrocarbons; over most of the site the water table was apparently deeper than the maximum effective penetration of the GPR signal (about 20 feet). Below that depth, no coherent returns appear on the GPR profiles, only noise and signal reverberations are present.

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4.1.2.2 <u>Naknek River Seep</u>

The results of the reconnaissance geophysical profiling near the Naknek River seep are summarized in Figure 2 of Appendix I. This area was targeted because partially buried drums were identified. Profile plots of the electromagnetic and magnetic data and listings of the data are provided in Appendix I.

The location of the anomalies indicative of major disposal suggest that drums are buried throughout the western end of the site, although additional profiling over a more thorough grid would be required to confirm this interpretation. Lines 100W, 300W, and 500W were surveyed to investigate suspicious cleared areas at the north side of the site, which were thought to possibly contain buried tanks. No metallic features were identified.

4.1.2.3 North Barrel Bluff

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At North Barrel Bluff most of the waste disposal was confined to a narrow strip along the bluff edge, extending south no more than 50 to 100 feet. The grid of geophysical lines surveyed is summarized in Plate 2 of Appendix I. All lines were surveyed using both a terrain conductivity meter and a magnetometer. Selected lines were surveyed with ground penetrating radar.

The rumors of major burial pits extending hundreds of feet back from the bluff are not supported by the geophysical data. Lines 1100W and 634S extended well into the zones of surface disturbance evident on aerial photographs of the site, but no geophysical evidence of major debris burial was found. The baseline was also extended to the west so that a complete transect to the west end of the bluff could be obtained. No evidence of waste burial was observed along this profile.

Two additional areas across which reconnaissance profiles were surveyed did provide some evidence of waste disposal. Line MP-1 (Plate 2 of Appendix I) indicated that the disturbed area southeast of the South Barrel Bluff-North Barrel Bluff road does contain a substantial amount of metallic debris.

The two reconnaissance lines surveyed in the reported disposal area northeast of the school district building (Lines MP-2 and MP-2X, Plate 2) showed little evidence of waste disposal. Rapid profiling of this general area with a terrain conductivity meter indicated scattered occurrences of waste material, but not a large debris area. Additional work would be required in this area in order to determine if a large disposal pit exists. The primary objective of delineating the extent of the major disposal areas at the North Barrel Bluff site was accomplished. The geophysical data provided little direct information toward the secondary objectives of exploring subsurface soil conditions outside the debris boundaries and searching for indications of possible contaminant plumes. Outside the disposal areas, both electromagnetic and magnetic data indicate that uniform soil conditions, as measured by those techniques, exist. The magnetic field data indicated higher total field readings on the terrace above the bluff than down on the flood plain of King Salmon Creek as illustrated by Line 1900W (Figure B4). This suggests that the outwash sands that comprise the terrace are possibly volcanic in origin and contain a significant proportion of magnitite.

4.1.2.4 South Barrel Bluff

The geophysical survey results for the South Barrel Bluff site are summarized in Plate 3 of Appendix I. All lines were surveyed using both the terrain conductivity meter and the magnetometer/gradiometer. In addition, Lines 421E and 500E were also profiled using the ground penetrating radar system.

Beginning at the west end of the South Barrel Bluff site, Lines 548S and 312W were surveyed across a cleared rectangular area, which is identified as "Landfill No. 1" in the presurvey report. The geophysical results indicated very little waste disposal on this portion of the site, except for some disturbance in the vicinity of a reinforced concrete foundation structure.

Plate 3 of Appendix E shows that most of the major waste disposal has apparently been concentrated between Lines 300E and 1280E. The two houses are apparently situated in close proximity to the waste disposal. Local residents report encountering debris when installing septic systems and drilling domestic water wells at these residences (the septic systems and wells are located adjacent to the houses).

A local contractor reported a disposal site between the main road and the bluff just east of the houses. A relatively closely spaced grid of geophysical lines was surveyed in this area to search for this reported deposit [Lines 1280E, 1380E, 1458E, 75N, and 100N (extension)]. The geophysical profiling indicated that the waste disposal in this area is confined to a former gully that cuts through the bluff (Plate 3 of Appendix E, Lines 1458E, 75N, and the Base Line). The geophysical results suggest that the material was deposited in the gully and covered over.

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4.1.2.5 <u>Conclusions of the Geophysical Investigation</u>

The use of the electromagnetic conductivity and ground magnetic geophysical techniques proved to be very effective as rapid reconnaissance methods for delineating the limits of major waste disposal areas at the North Barrel Bluff site, the South Barrel Bluff site, and Naknek River seep. More precise definition of the boundaries of the disposal zones could be achieved at a future date by surveying profiles at a closer line spacing.

Geophysical profiling in the Eskimo Creek seep area using the ground penetrating radar was not successful in identifying the source of contamination at the Eskimo Creek seep. Further general profiling is not recommended in this area, although the GPR would be useful at this site for searching for specific targets such as utility lines, buried tanks, or drilling hazards associated with installation of groundwater monitoring wells.

4.1.3 <u>Hydrogeologic Investigation</u>

The current understanding of the site-specific geology for the Eskimo Creek seep and Barrel Bluff Areas is based on information from four investigations:

- Corps of Engineers oil contamination exploration, 1978
- o Existing well logs (base and local residential)
- o Corps of Engineers foundation report, 1985
- o IRP Stage 1 investigation, CH2M HILL 1987-88

Boring logs from these investigations are presented in Appendix C-5. Well locations are presented in Figures 4-2 and 4-3 and well details are presented in Table 4-1. At the conclusion of the Stage 1 field investigation, existing wells completed during earlier investigations (whose location could be determined) were surveyed for horizontal location and vertical elevation. Each new well installed during this period also was surveyed. A summary of well coordinates and elevations is provided in Appendix E. The new survey indicates that several of the existing Corps well elevations previously reported on well logs were incorrect.

4.1.3.1 <u>Geology</u>

Site stratigraphy is illustrated in two cross sections. Figure 4-4 shows the locations of the cross sections and boring logs are presented in Appendix C-5. Section A-A' (Figure 4-5) illustrates the geology from northwest to south-

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LEGEND

- **EXISTING SOIL BORINGS** 0
- **EXISTING MONITORING WELLS** 0
- EXISTING BASE PRODUCTION WELLS D
- EXISTING LANDFILL MONITORING WELLS Δ
- CORPS OF ENGINEERS CORPS OF ENGINEERS AIR FORCE AH AP
- LF
- мw CH2M HILL

FIGURE 4-3

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EXISTING MONITORING WELL AND SOIL BORING LOCATIONS KING SALMON AIR FORCE STATION KING SALMON, ALASKA - СКМНІШ- Table 4-1 MONITORING WELL CONSTRUCTION DETAILS

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| | | | | | | | | | | Depth | | |
|---------------------------|------------------|-----------|----------|-----------|------------|-----------|-------------|------------|------------|---------|--------------------------------|------------|
| | | | | | | | Hell | Protective | | Below | Screened | |
| | | | Hell | | Total | Weilhead | Casing | Casing | Ground | Ground | Interval Below | Sum |
| Hell. | Date | Installed | Dlameter | Hell | Length | Elevation | Stickup | St 1ckup | Elevation | Surface | Ground Surface | Length |
| Number | Installed | By | (inches) | Materials | (51) | (Et NGVD) | (££) | (EE) | (ft NGVD) | (11) | (ft) | (Et) |
| | L8/L0/11 | NCR.) | ſ | bur | 37 95 | 01.03 | | | 02 23 | 26 35 | 40 CE - 40 CC | • |
| | 10/10/11 | | | | | 50 66 | | | | 03 00 | | |
| | 10/10/11 | | • • | | | 51 DE | | | | 00.01 | 00 FC - 00 FC | ; . |
| | 10/01/11 | | | | 01 92 | 50 Y | | | 60°.72 | | 70.42 - 70.41 FF OF - FF OF | . . |
| | 19/11/11 | | • • | 2110 | 07*0C | 16.00 | 36 1 | | 33.63 | | 10 11 - 20 11 | n u |
| | | | • • | | | 00.00 | | | | | 11.04 11.01 01 01 - 01 01 | • • |
| | 19/01/11 | | | | | 00.00 | | | 16.40 | 67 °C | 67°DC - 67°D7 | n 1 |
| 67-10 | /A//1/11 | | ~ | | 20.02 | 60.9 | 2:1 | دد.، د | 04.40 | 33.82 | 79°97 - 78°87 | " |
| MH-30 | 11/19/87 | CH2N | 7 | PVC | 37.20 | 68.28 | 1.1 | 2.43 | 66.57 | 35.49 | 20.49 - 30.49 | \$ |
| Mi- 31 | 11/20/87 | CH2M | 7 | PAC | 37.63 | 67.08 | 1.77 | 2.42 | 65.31 | 35.86 | 20.86 - 30.86 | 5 |
| Mil-32 | 11/21/87 | CH2M | 7 | PVC | 39.13 | 64.81 | 1.45 | 2.38 | 63.36 | 37.68 | 22.68 - 32.68 | s |
| Mil-33 | 11/23/87 | CH2M | 7 | PVC | 40.71 | 68.57 | 1.82 | 2.21 | 66.75 | 38.89 | 24.89 - 34.89 | • |
| 18-34 1 | 12/09/87 | CH2H | 7 | PVC | 38.00 | 67.55 | 1.75 | 2.33 | 65.8 | 36.25 | 21.25 - 31.25 | ŝ |
| MI-36 | 05/06/88 | CH2M | 2 | PVC | 38.20 | 68.49 | 2.00 | 2.05 | 66.49 | 36.20 | 24.2 - 34.2 | 7 |
| 101-37 101-37 | 05/10/88 | CH2M | ~ | PVC | 40° 90 | 68.93 | 1.91 | 2.36 | 67.02 | 38.99 | 26.99 - 36.99 | 2.2 |
| MW-38 | 05/12/88 | CH2M | 7 | PVC | 39.80 | 69.06 | 1.55 | 2.17 | 67.51 | 38.25 | 25.75 - 35.75 | 2.5 |
| 184-39 | 05/13/88 | CH2M | 7 | PVC | 32.32 | 63.73 | 1.93 | 2.17 | 61.80 | 30.39 | 16.39 - 26.39 | 4 |
| 07-184 | 06/08/88 | CH2M | 7 | PVC | 41.17 | 65.79 | 1.61 | 1.98 | 64.18 | 39.56 | 25.36 - 35.36 | 5.2 |
| 19-19 | 05/13/88 | CH2M | • | PVC | 99.37 | 65.94 | 1.30 | 2.50 | 65.31 | 98.07 | 74.5 - 94.5 | 3.6 |
| HH-42 | 06/11/88 | CH2M | • | PVC | 100.40 | 68.54 | 1.93 | 2.50 | 66.75 | 98.47 | 87.5 - 97.5 | I |
| EF-19 | 06/11/88 | CH2M | - | PVC | 117.57 | 67.25 | 2.15 | 2.50 | 65.20 | 115.42 | 6.91- 9.69 | 5.5 |
| 1-19 | 09/14/83 | USCOE | 1.5 | PVC | 30 | 64.74 | 1.79 | 2.24 | 62.95 | 28.21 | 23.31 - 28.21 | đ |
| 9 ب | 09/11/83 | USCOE | 1.5 | PVC | 35 | 66.34 | 0.99 | 1.55 | 65.35 | 34.01 | 29.01 - 34.01 | 80 |
| 6-MH | 09/15/83 | DSCOE | 1.5 | PVC | 35 | 66.02 | 0.36 | 0.00 | 65.66 | 34.64 | 29.64 - 34.64 | 8 |
| MM-10 | 09/11/83 | USCOE | 1.5 | PVC | 35 | 68.70 | 1.96 | 2.31 | 66.74 | 33.04 | 28.04 - 33.04 | đ |
| Hil-16 | 09/16/83 | USCOE | 1.5 | PVC | 30 | 59.14 | 1.19 | 1.84 | 57.95 | 28.81 | 18.81 - 28.81 | 8 |
| HH-18 | 09/16/83 | USCOE | 1.5 | PVC | 30 | 59.43 | 2.05 | 2.57 | 57.38 | 27.95 | 17.95 - 27.95 | Q |
| MH-21 | C9/16/8 3 | USCOE | 1.5 | PVC | õ | 60.41 | 0.64 | 1.01 | 59.77 | 29.36 | 19.36 - 29.36 | B n |
| MH-22 | 09/18/83 | USCOE | 1.5 | PVC | 35 | 64.30 | 0.93 | 1.27 | 63.37 | 34.07 | 29.07 - 34.07 | 9 0 |
| [- 7] | 8 | USAF | • | PVC | 82.23 | 65.70 | 0.55 | a. | 65.15 | 81.68 | рų | na. |
| LF-2 | An A | USAF | • | PVC | 40.55 | 63.24 | 3.0 | B | 60.34 | 37.55 | e u | Q. |
| LF-3J | B U | USAP | 4 | PVC | 36.6 | 60.75 | 1.25 | na | 59.5 | 35.35 | ŋŋ | Pu |
| KS-4 | 1952 | OSAF | 8 | St eel | 9 0 | 66.62 | na | an | P | 228 | na | P |
| KS-5 | 1952 | USAF | 8 | Steel | | 65.50 | 8 17 | B R | 9 0 | 137 | 227 - 237 | 80 |
| | | | | | | | | | | | | |
| Notes: | | | | | | | | | | | | |
| | | | | | | | | | | | | |

The screened interval given for ACOE wells is estimated from ACOE well logs and may not be accurate because survey data provded in the logs is apparently incorrect.
 na = Not Available.

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FIGURE 4-4

LOCATION OF

HYDROGEOLOGIC CROSS-SECTIONS KING SALMON AIR FORCE STATION KING SALMON, ALASKA

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east across the site and includes the Eskimo Creek seep. Section B-B' (Figure 4-6) runs from the north to southwest, in the general direction of regional groundwater flow.

Geology of the site consists of alternating sequences of permeable glacial outwash materials and relatively impermeable tills. The outwash deposits contain a surficial unconfined aquifer and numerous locally-confined aquifers at depth. Glacial tills and finer-grained sediments act as semiconfining or confining layers.

4.1.3.1.1 Shallow Aquifer

The surficial geologic unit is composed of 20 to 40 feet of moderately well sorted outwash sands and silty sands. Some coarser gravel zones are found locally. This unit is extensive throughout the King Salmon area forming terraces that have been partially reworked by wind. Within the site boundary, the unit is laterally consistent as it has been encountered in each boring.

Grain size analyses for shallow aquifer soil samples collected by the U.S. Corps of Engineers are presented in Appendix D-4. Grain size curves indicate the shallow aquifer ranges from clean well-sorted sands (SP) to silty sands (SM).

The shallow unconfined aquifer within this unit is found at depths from 21 to 31 feet below ground surface at the site. The aquifer is separated from lower aquifers by an underlying glacial till and has a saturated thickness ranging from 0 to greater than 9 feet beneath the site.

4.1.3.1.2 Upper Aquitard

Underlying the surficial sands is a till deposit consisting of gravelly sandy silt and sandy silt. The contact between the two deposits is distinct (Figures 4-5 and 4-6). The till deposit acts as an aquitard between the upper unconfined aquifer and the intermediate confined aquifer.

Three borings completed during the Stage 1 investigation penetrated through this aquitard into the underlying intermediate aquifer. The thickness of the till ranged from 7 to 22 feet in these borings. Local water well drilling logs indicate the aquitard is laterally extensive throughout the area (station and barrel bluffs). The thickness of the unit may vary greatly. Several base wells and numerous residential water wells have been identified (Figure 4-7). Associated well logs, where available, are in Appendix C-5.

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KING SALMON AIR FORCE STATION KING SALMON, ALASKA

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Several of the shallow borings [CH2M HILL and U.S. Army Corps of Engineers (USCOE)] penetrated into the upper aquitard. The stratigraphic contact between shallow aquifer and underlying till was interpreted based on descriptions of grain size, density (split-spoon blow counts), and moisture content. These data points were used to construct the structure contour map of the till surface (Figure 4-8). The accuracy of any one point may be t2 feet. However, the data set clearly indicates a high area or thickening of the till beneath the site. The upper contact of the till is at a higher elevation than the shallow groundwater table measured in June 1988. Soil boring SB-35, shown in Figure 4-8, encountered till in the unsaturated zone.

4.1.3.1.3 Intermediate Aquifer

The intermediate confined aquifer is characterized primarily by interbedded sequences of silty sands, sandy gravels, and silty sandy gravels. Gravels are predominantly subrounded to subangular in shape with maximum observed size of 2 inches. The three borings that penetrated this unit during the Stage 1 investigation (MW-41, 42, and 43) indicate a depth to the top of the aquifer ranging from 43 to 48 feet below ground surface and a thickness ranging from 41 to 62 feet. The three borings yielded significant water during drilling throughout this zone.

The intermediate aquifer is generally used for residential water supplies. Nearby wells shown in Figure 4-7 and presented in Appendix C-6 are screened in sandy gravel materials found beneath the upper aquitard. These wells generally produce adequate residential supplies (10 to 20 gpm). This zone could include several small confined aquifers, each contained in a limited extent outwash deposit.

The lateral extent of the intermediate aquifer is unknown. Glacial deposits are very heterogeneous, and it is difficult, therefore, to correlate the intermediate zone at the Eskimo Creek seep site to nearby areas (such as the Barrel Bluffs).

4.1.3.1.4 Lower Aquitard(s)

A second fine-grained deposit was encountered beneath the intermediate aquifer in the three new deep borings at depths ranging from 88 to 110 feet. The geologic logs for two base wells that penetrated this unit, KS-4 and KS-5, indicate a thickness of about 90 to 115 feet. The uppermost portion of the unit is generally characterized by silty sand, sandy/ gravelly clay, and clay. A lower moisture content was observed in drill cuttings. Significant quantities of water were observed in boring MW-42 after penetrating several feet into the lower aquitard. The drilling subcontractor indicated





that a similar water bearing, fractured clay formation was present at several other water well locations in the area. Whether or not these fractures allow communication between the intermediate aquifer and deeper aquifers is unknown.

4.1.3.1.5 Deep Aquifer

Geologic logs from King Salmon AFS production wells KS-4 and KS-5 show the presence of a deep aquifer at a depth ranging from 205 to 225 feet. The aquifer materials consist primarily of sand and sandy gravels.

4.1.3.2 Groundwater Flow

4.1.3.2.1 Shallow Unconfined Aquifer

Groundwater levels in the shallow unconfined aquifer have been recorded in existing wells on six occasions. Water elevation data is presented in Table 4-2. Data from October 1987 and June 1988 were used to interpret and construct the groundwater elevation maps presented as Figures 4-9 and 4-10. These dates were used because they reflect opposite seasons and are the most complete data sets.

Groundwater flow for October 1987 (Figure 4-9) is generally to the south towards the Naknek River and southeast along Eskimo Creek. This flow pattern is somewhat different than previous presentations (CH2M HILL Presurvey Report, 1986) because previous contour maps were based on well elevations from USCOE well logs that were incorrect. These elevations may have been estimated from contour maps.

This aquifer system is extremely anisotropic and heterogeneous because of glacial till silts located within the relatively clean sands. Consequently, the direction of groundwater flow is not interpreted to be parallel to the hydraulic gradient in this area. Flowlines on the groundwater flow maps reflect the effects of the till barrier.

The groundwater flow map presented for the June 1988 data (Figure 4-10) includes the newly installed well data and represents the most detailed interpretation to date. The water level contours indicate the regional groundwater flow direction is from north to south across the site with a major component towards Eskimo Creek to the southeast. Eskimo Creek and the underlying glacial till ridge affect the regional flow to the south. The Eskimo Creek flood plain is 30 to 35 feet lower in elevation than the station. This area is a local discharge zone for the unconfined aquifer. As the groundwater approaches the bluff, flow lines swing to the southeast to reflect this discharge.

Table 4-2 Water level Jata for King Salmon Afs

| } | | | E861 | Octob | er 1985 | April | 1987 | Octob | r 1987 | May | 1988 | | 06) Q88 |
|----------------|-----------------|--------|-----------|--------|-----------|--------|-----------|--------|-----------|--------|------------------|----------------|----------------|
| | He11 | Hater | Mater | Mater | Nater | Mater | Water | Nater | Mater | Hater | Hater | Hater | Mater |
| Well | Elevation | Depth | Elevation | Depth | Elevation | Depth | Elevation | Depth | Elevation | Depth | Elevation | Depth | Elevation |
| ; | 22 22 | | 22 12 | | : | | | | | | | | |
| No. | (feet) | (BTOC) | (feet) | (BTOC) | (feet) | (BTOC) | (feet) | (BTOC) | (feet) | (BTOC) | (feet) | (BTOC) | (feet) |
| 1- 1 | 64.74 | 27.2 | 37.54 | 27.02 | 37.72 | 27.03 | 17.76 | 30.55 | 34.19 | 28.34 | 36.4 | 27.99 | 36.75 |
| Ŷ | 66.34 | 26.9 | 39.44 | 27.16 | 39.18 | 27.9 | 38.54 | 27.19 | 39.15 | 27.9 | 38.44 | 29.833 | 36.507 |
| 6- N | 66.02 | 28.1 | 37.92 | 19.41 | 36.61 | 28.75 | 37.27 | 29.06 | 36.96 | 29.26 | 36.76 | 29.39 | 36.63 |
| 01-MH | 68.7 | 29.0 | 39.7 | 31.78 | 36.92 | 30.91 | 37.79 | 31.38 | 37.32 | 31.65 | 37.05 | 91.74 | 36.96 |
| M-16 | 59.14 | 21.4 | 37.74 | 23.07 | 36.07 | 22.5 | 36.64 | 23.05 | 36.09 | 22.83 | 36.31 | 22.94 | 36.2 |
| 81-18 | 59.43 | | | | | 22.49 | 36.94 | 17.22 | 36.72 | 22.9 | 36.53 | 22.95 | 34. AF |
| 161-21 | 60.41 | 23.3 | 37.11 | 24.47 | 35.94 | 23.8 | 36.61 | 24.18 | 36.23 | 24.12 | 36.29 | 74.37 | 90 YE |
| HH- 22 | 64.3 | 24.4 | 39.9 | 25.3 | 39.00 | 24.48 | 39.82 | 1 24.8 | 39.5 | 25.42 | 38.88001 | 24.875 | 39.425 |
| 161- 23 | 69.1 | | | | | | | | | 11.55 | 35.99 | 33.42 | 35.68 |
| HH-24 | 60.66 | | | | | | | | | 21.77 | 38.89 | 21.48 | 39.18 |
| 164- 25 | 62.05 | | | | | | | | | 22.82 | 39, 23 | 22.54 | 19.51 |
| HH- 26 | 66.92 | | | | | | | | | 27 48 | 30 44 | | 12105 |
| MI- 27 | 64.9 | | | | | | | | | 25 58 | | 67.12 CC 3C | 59.65 90 69 |
| MH-28 | 66.68 | | | | | | | | | 27.96 | 38.72 | 97 7C | 00-00 10 80 |
| 161- 29 | 62.9 | | | | | | | | | ~ ~ ~ | 0.00 | | |
|) | | | | | | | | | | 6.01 | 0.45 | 1/ . 97 | 61.65 |
| 0E-MH | 68.28 | | | | | | | | | 29.36 | 38.92 | 29.19 | 60°6 E |
| 16-191 | 67.08 | | | | | | | | | 27.84 | 39.24 | 27.82 | 39.26 |
| M-32 | 64.81 | | | | | | | | | 25.64 | 39.35 | 25.26 | 39.55 |
| 181-33 | 68.57 | | | | | | | | | 32.59 | 35.98 | 32.78 | 35.79 |
| 101- 34 | 67.55 | | | | | | | | | 30.11 | 37.44 | 30.07 | 37.48 |
| M-36 | 68.49 | | | | | | | | | | | 17.03 | 97 YL |
| 16-37 | 68.93 | | | | | | | | | | | 96.15 | 36.97 |
| 8E~MW | 59.06 | | | | | | | | | | | 30.84 | 38.22 |
| 64- 39 | 63.73 | | | | | | | | | | | 24.13 | 39.6 |
| 07-18 | 65.79 | | | | | | | | | | | 29.03 | 36.76 |
| 17-159 | 65.94 | | | | | | | | | | | 29.73 | 36.21 |
| MI-42 | 68.54 | | | | | | | | | | | 38.56 | 39.98 |
| 5773 | 67.25 | | | | | | | | | | | 34.08 | 71.66 |
| I-J1 | 65.7 | | | | | | | 27.12 | 38.58 | 27.38 | 38.32 | 27.1 | 38.6 |
| LF-2 | 63.24 | | | | | | | 24.42 | 38.82 | 24.80 | 38.44 | | |
| LF-3 | 60.75 | | | | | | | 22.35 | 38.40 | | | | |
| | | | | | | | | | | | | | |
| do 1 Tob | OF VELL CASIF | - 6ı | | | | | | | | | | | |
| BTOCBe | niow top of cas | tng. | | | | | | | | | | | |

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Vertical Datum-King Salmon Air Force Station Site Datum.





OCTOBER, 1987 KING SALMON AIR FORCE STATION KING SALMON, ALASKA

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The shaded area near the center of the site indicates where the upper aquitard surface is higher in elevation than the groundwater level. This is represented by a steepening of the potentiometric contours where finer grained materials impede groundwater flow. Due to the heterogeneous, anisotropic conditions, groundwater flow directions in this area would be expected to swing to the east and/or west around the till feature (not perpendicular to the equipotential lines). As water levels drop seasonally, the till would be expected to have a greater effect on the direction of groundwater flow as a larger area of the aquifer becomes unsaturated.

The aquifer gradient through areas not affected by the till is about 3.5×10^{-3} . Through the till area, the gradient steepens to 1×10^{-2} .

The hydraulic conductivity of the unconfined aquifer was estimated from single well "slug" tests performed on each newly installed shallow well. Parameter values are summarized in Table 4-3. Raw data and plots are presented in Appendix D-1.

Slug and bail tests were attempted in each new well using a solid rod to displace a slug of water. A pressure transducer and data logger were used to record water level and time data. Field data were analyzed with the method presented by Bouwer and Rice (1976), which is applied to unconfined aquifers to determine the hydraulic conductivity principally in the horizontal direction.

Several difficulties were encountered in conducting the tests that resulted from the monitoring well construction and high conductivity of the aquifer. The wells were screened above the saturated interface to allow accumulation of free product. This only left a few feet of water in the wells into which to place a "slug" or withdraw a "bail." Consequently, when the slug or bail was applied, the system often recovered in a very short period of time. This problem was partially compensated for by use of a data logger that could record readings in increments of seconds. However, several wells still responded too quickly (e.g., tenths of seconds) to produce usable results.

Another difficulty arose because the wells were screened above the saturated interface. When a slug was applied, water immediately flowed through the screen into the unsaturated sandpack. On the slug cycle, this took about 1.5 seconds. The slug then decayed through the volume of the borehole. This boundary problem was addressed by using a corrected radius (Bouwer and Rice, 1976) that included the sandpack. On the bail tests, water from the sandpack appeared to drain back into the casing over a much longer period

| | | | SHALLOW A | QUIFER PARAME KING SALM | TERS AND | e 4-3 HYDRAULIC CONDU ING SLAMON, ALA | ICTIVITY ESTIMAT | ш | |
|----------------------------------|---|---|---------------------------------|-----------------------------------|----------------------------------|---|------------------------------------|----------------------------|--|
| Well ID | Casing Radius rc' (ft) | Borehole Radius rw (ft) | Length of Sandpack L (ft) | Saturated Thickness H (ft) | Ratio L/rw | C Dimensionless ParameterD | | Time Lag ^a | Estimated Hydraulic Conductivity Bouwer and Rice |
| MW-23 MW-24 | 0.268 0.268 | 0.416 | 4.53 | 4.53 | I0.9 | 1.3 | 1.72 | To (second) | (ft/day) |
| MW-25 MW-28 | 0.268 | 0.416 | 9, /4 8, 68 | 9.74 8.68 | 23.4 20.9 | 1.7 | 2.37 | 0°27 | 10 9 4 |
| MW-29 | 0.268 | 0.416 | 9.27 8.81 | 9.27 8.81 | 22.3 | 1.5 | 2.37 | 13.0 | 41 61 |
| MW-32 MW-34 | 0.268 0.768 | 0.416 | 10.0 | 10.0 | 24.0 | 1.5 | 2.32 7 38 | 11.5 | 11 |
| MW-36 | 0.268 | 0.416 | 6.17 6.17 | 7.93 | 19.1 | 1.4 | 2.24 | c/ 01 | 69 |
| MW-3 8 | 0.268 | 0.416 | 8.96 | 8.96 | 21.5 | 1.5 | 2.02 2.34 | 36.0 | 28 |
| arime lag | from plots | in Appendix | 0. | | | | | | |
| ^D Used in Reguired | Bouwer and R Equations: | dice model to | calculate ln | (Re/Rw). | | | | | |
| I | | | | | | | | | |
| Bouwer | and Rice (19 | 76): K = <u>rc</u> | ln (Re/rw) 2 L T | ln | k (Re/rw) | = (<u>1, 1, 1</u>) | + C-1 L/rw | | |
| Casing ra | lius correct | ed for sandpi | ick: rc' = [| rc + n(rw ² - | rc1)] ¹ | | | | |
| Where: r r r r | <pre>Effective C = Radius of C = Corrected C = Porosity</pre> | e casing radi f casing d casing radi of sandpack | urs urs | rw = Bore L = Leng To = Lag | hole radju th of sand time | is Ipack H | e = Effective zo = Saturated th | ne of influence ickness | (Bouwer and Rice) |

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Table 4-3

of time (tens of seconds). This interference essentially masked the effects in the aquifer and the bail tests could not be analyzed.

Hydraulic conductivity estimates for nine slug tests ranged from 10 to 94 feet per day. These values are representative of clean fine sands and slightly silty sands. The wells that responded too quickly to produce adequate data may have higher conductivities.

Estimates of the groundwater flow velocity were calculated using a modified form of Darcy's law shown below:

$$V = \frac{Ki}{n_{\rho}}$$

where:

V = average linear velocity K = hydraulic conductivity i = hydraulic gradient n_e = effective porosity

The hydraulic conductivity of the shallow unconfined aquifer was determined from slug test data and ranged from 10 to 94 feet per day, with an average of 51 feet per day.

The effective porosity was assumed to be 0.30, which correlates to a clean, fine sand.

The gradient based on the June 1988 water table elevation contour map for an area unobstructed by the till mound is 3.5×10^{-3} . This gradient can vary seasonally; additional water level measurements and flow system interpretations are recommended.

Estimates of flow velocities are provided in Table 4-4. The travel time downgradient for a distance of 100 feet is also provided. The total flow in gallons per minute through the aquifer, assuming a saturated thickness of 10 feet by 100 feet, is included.

These estimates include a number of simplifying assumptions and, therefore, should only be considered as general indicators of the groundwater flow conditions. For engineering purposes (e.g., design of groundwater extraction system), additional investigation to define aquifer parameters is required. Further investigation would include multiple well aquifer testing (horizontal and vertical) and seasonal groundwater level measurements for flow system interpretations.

Table 4-4 presents estimates of the underflow and groundwater flow velocities for the unconfined aquifer.

| | SE | Tab HALLOW AQUIFER | le 4-4 PARAMETER ESTIMAT | TES | |
|-----------------------------------|---------------------------------------|-----------------------------------|---|--------------------------------------|------------------------------|
| Estimated Thickness, B (ft) | Hydraulic Conductivity (ft/day) | Average Gradient, I (ft/ft) | Underflow, Ω ^a (gpm/100 ft) | Average b Velocity (ft/day) | Travel Time (days/100 ft) |
| 10 | 10 to 94 51 ^C | 3.5×10^{-3} | 0.2 to 1.7 0.9 | 0.12 to 1.1 0.6 | 91 to 833 167 |

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^aUnderflow Q = KIB (100 feet).

ΚI ^bAverage velocity \overline{V} = porosity approximately 0.30

^CAverage conductivity from nine slug tests.

4.1.3.2.2 Intermediate Aquifer

Water levels measured between the shallow aquifer and intermediate aquifer indicate a downward vertical gradient condition of about 7x10⁻⁴ ft/ft. Thus, the shallow unconfined aquifer is a recharge source to the intermediate aquifer(s) (Table 4-5).

| | ESTIMATED | Table 4-5 VERTICAL GRADIENT JUNE 1988 WELL NEST | 'S | |
|----------------|-------------|--|-------------|----------|
| Shallow | Aquifer | Intermediate | Aquifer | Vertical |
| Well | Groundwater | Well | Groundwater | Gradient |
| Identification | Elevation | Identification | Elevation | (ft/ft) |
| MW-31 | 39.26 | MW-41 | 36.21 | 0.06 |
| MW-33 | 35.79 | MW-42 | 29.98 | 0.1 |
| MW-34 | 37.48 | MW-43 | 33.17 | 0.06 |
| | | | | |

The horizontal direction of groundwater flow in the intermediate aquifer was interpreted from water level data from the three newly installed deep wells (Figure 4-11). The flow direction is highly interpretive because only these data points were used and the screened intervals of the wells





| | RESULTS |
|---------------|--------------|
| | TEST |
| e 4- 6 | RECOVERY |
| Tabl | AQUIFER |
| | INTERMEDIATE |

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| Well Identification | Aquifer Materials | Discharge Rate, Q, During Pump Test (ft ³ /day) | Residual Drawdown Per Log Cycle <u>A</u> S (ft) | Aquifer Thickness, B (ft) | Estimated Transmissivity, T (ft ² /day) | Estimated Hydraulic Conductivity (ft/day) |
|-------------------------|---|---|---|---------------------------------|--|--|
| MW-41 | Sandy gravels | 3,850 | 0.355 | 41.0 | 1,985 | 48.4 |
| MW-42 | Sand | 3,850 | 0.855 | 47.5 | 824 | 17.3 |
| MW-43 | Sand and gravel | 2,406 | 0.875 | 62.5 | 503 | 8.1 |
| Theis' Recovery | Y Method: | T = KB = 2.3 Ω 4π ΔS | | | | |
| Where: K ∆S ∆S | = Transmis + Hydrauli = Aquifer (= Discharg | sivity, ft²/day c conductivity, ft/day thickness, ft e rate, ft³/day drawdown per log cycle | | | | |
| | | • | | | | |

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vary. Also, the lateral continuity of these glacial deposits beneath the site is unknown. The direction of flow appears to be to the south, which roughly correlates with the upper aquifer. The apparent horizontal gradient is about 1×10^{-2} ft/ft.

A single well pumping/recovery test was performed on each intermediate well to provide an estimation of the hydraulic conductivity. The recovery data were analyzed using the This Recovery Method (Kruseman and DeRidder, 1970) and results are summarized in Table 4-6. Raw data and plotted data are presented in Appendix D-2. In each of the analyses, late time recovery sections of the plots were used to minimize the effects of partial penetration.

Table 4-7 presents estimates of aquifer parameters for the intermediate zone. These estimations include the assumptions presented in the shallow aquifer section.

| | · · · · · · · · · · · · · · · · · · · | INTERMEDIA | Table 4 TE AQUIFER PAR | -7 RAMETER ESTIMATE | 5 | |
|-------------|---------------------------------------|---|-----------------------------------|---|---------------------------------|------------------------------|
| Well No. | Estimated Thickness, B (ft) | Hydraulic Conductivity,C K (ft/day) | Average Gradient, I (ft/ft) | Underflow, Q ^a (gpm/100 ft) | Average Velocity (ft/day) | Travel Time (days/100 ft) |
| 41 | 41.0 | 48 | 2×10^{-2} | 20.4 | 3.8 | 26 |
| 42 | 47.5 | 17 | 2×10^{-2} | 8.4 | 1.36 | 74 |
| 43 | 62.5 | 8 | 2×10^{-2} | 5.2 | 0.64 | 156 |

Underflow Q = KIB (100 feet).

^bAverage velocity = KI/n (assumed porosity n = 0.25).

Hydraulic conductivity from short term recovery tests.

4.1.3.2.3 Deep Aquifer

The water supply wells of the station are shown in Figure 4-7, and well logs are presented in Appendix C-2. A summary of well data is provided in Table 4-8. Well yields range from 7 to 140 gpm.

4.1.4 Subsurface Soil Contamination Investigation

4.1.4.1 Eskimo Creek Seep

Soils contamination data from previous investigations include oil and grease analyses from soil samples in six borings (AH-1 through AH-6) shown in Figure 4-12. The oil and grease

Table 4-8 KING SALMON AFS WELL DATA

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| | Well No. | Well No. | Well No. | Well No. | Well No. | Well No. | Well No. | Well No. | Well No. | Well No. |
|---|----------------------|-------------------------------|----------------------|-------------------------|-------------------------|----------------------------|-------------------------------|-----------------------|-----------------------|------------------------------------|
| Location | 1 Building 130 | 2 Building 139 | s Building 205 | Building 638 | 5 Building 650 | 6 Building 654 | 7 Building 148 | 8 Building 327 | 9 Building 561 | 10 Building 1001 |
| Installation Date | 5/49 | 3/49 | na | 10/52 | 10/52 | na | Pu | 12/72 | 1/73 | na |
| Depth (feet) | ł | 104 | ł | 228 | 233 | 137 | 168 | 116 | 224 | 135 |
| Casing Dlameter (inches) | ł | ł | œ | ; | 8 | 80 | Q | Q | Q | Q |
| GPM (Open Flow) | ł | 140 | ł | 140 | 55 | ł | 75 | 7 | 10 | ١ |
| Drawdown (feet) | 1 | 120 | ł | 22.7 | 0.6 | 4.0 | 50.3 | 6 | 16 | ١ |
| Static Water Level ^a (feet below grade) | na | 8 | na | 48 | 40 | ł | 30 | ł | 34 | na |
| Pump | L L | Goulds | 1 | Goulds | ł | 1 | Goulds | Jacuzz1 | Jacuzzi | ł |
| Model | ł | Submersible | ł | Submersible | ł | 1 | ubmersible | Submersible | Submersible | 1 |
| Hosepower | 1 | 15 | ł | 15 | 7.5 | ł | 2 | ł | ł | ł |
| Cycle | 1 | ł | ł | ł | ł | 1 | ł | ł | ł | ١ |
| Phase | 1 | ε | ł | æ | en | ł | Ē | ł | 1 | ١ |
| Amperage | ! | 40 | ł | 40 | 23.3 | ł | 15 | ł | ł | 1 |
| RPM | 1 | 3,450 | ł | 3,450 | 3,450 | : | 3,450 | ł | ł | 1 |
| Installation | ł | 1949 | ł | 1952 | 1952 | 1 | 1957 | 1972 | 1972 | 1 |
| Remarks | Abandoned | Potable | Abandoned | Potable | Potable | Nonpotable | Potable | Potable | Potable | Potable |
| Condition & Use | ł | Standby Fire Protection | ł | Main Water Supply | Main Water Supply | Supplies Bldg. 654 J | Standby Fire Protection | Supplies Bldg. 327 | Supplies Blåg. 561 | Supplies White Alice Site |
| ^a Water levels as meas | ured on 10/8 | 38. | | | | | | | | |

na = Not available.
Source: Installation Documents^{*}
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SCALE IN FEET value for the interval 25 feet below ground surface in the unsaturated zone is plotted by each well. The vertical distribution of contamination within each borehole is illustrated in Table 4-9.

One soil sample from each new borehole (MW-23 through MW-40) was selected for total petroleum hydrocarbon analysis. These values are plotted in Figure 4-12 at each well. The soil interval tested in each well was based on the soil sample having the highest volatile organic concentration (soil gas head space) reading with an OVA or HNU. Table 4-10 presents these soil analysis data.

The variation observed in the replicate sample at Boring No. 39 cannot be explained. HNU measurements performed on the discreet and replicate sample at the time of collection correlate reasonably well. Volatile screening values for other intervals are reported on the boring logs in Appendix C-1.

On the basis of available data, hydrocarbon contamination appears to be widespread in the unsaturated zone soils. The oil and grease data suggests contamination has spread vertically in the southern area of the site. The total petroleum hydrocarbons (TPH) data from the newly installed borings indicates contamination is spread laterally across the site at levels well above detection limits shown in Table 4-10.

Soil contamination appears to be highest in the central area of the site at levels above 1,000 mg/kg (dry weight). This correlates with the groundwater and soil gas data identifying a plume that spreads from the POL tanks across the site to the southeast and south towards Eskimo Creek.

As discussed in Section III, cuttings from soil borings exhibiting significant levels of contamination based on visual inspection and/or HNU readings were contained in DOT-approved drums, composited, and then tested for EP Toxicity (metals) and volatile and semivolatile organic compounds. The results of the analyses are presented in Table 4-11.

The drummed cuttings do not meet RCRA criteria for designation as a hazardous waste. The drums are currently in the possession of King Salmon Air Force facility engineers for final disposition.

4.1.5 <u>Water Quality Investigation</u>

Section 3.6 outlines the water quality sampling tasks for this project and procedures used to collect representative groundwater and surface water samples. This section presents the results from this task for each site. Laboratory analytical reports, and QA/QC data are presented in Appendix G.

| Hole No. | Sample No. | Depth (ft BGS) | Grease and Oil mg/kg (dry weight basis) | Water Depth (ft BGS) | Moisture Content % |
|------------------|---------------|-------------------|---|----------------------------|--------------------|
| AH-1 | 1 | 5 | 186.64 | 30 | 9.61 |
| | 2 | 10 | 224.60 | 30 | 4,91 |
| | 3 | 15 | 211.73 | 30 | 3-68 |
| | 4 | 20 | 301.25 | 30 | 3.52 |
| | 5 | 25 | 343.24 | 30 | 3,83 |
| | 6 | 30 | 157.56 | 30 | 13.39 |
| AH-2 | 1 | 5 | 157.50 | 30 | 6.55 |
| | 2 | 10 | 104.88 | 30 | 4.40 |
| | 3 | 15 | 62.14 | 30 | 5.11 |
| | 4 | 20 | 50.80 | 30 | 6.96 |
| | 5 | 25 | 158,24 | 30 | 5.06 |
| | 6 | 30 | 198.46 | 30 | 28.69 |
| ан-з | 1 | 5 | 167.63 | 30 | 11.03 |
| | 2 | 10 | 141.84 | 30 | 5.29 |
| | 3 | 15 | 75.17 | 30 | 10.42 |
| | 4 | 20 | 175.68 | 30 | 9.14 |
| | 5 | 25 | 100.65 | 30 | 4.78 |
| | 6 | 30 | 380.02 | 30 | 23.52 |
| AH-4 | 1 | 5 | 192,15 | 30 | 7.85 |
| | 2 | 10 |)5.06 | 30 | 6,26 |
| | 3 | 15 | 194.89 | 30 | 5.08 |
| | 4 | 20 | 104.49 | 30 | 2.88 |
| | 5 | 25 | 113,20 | 30 | 4.50 |
| | 6 | 30 | 200,50 | 30 | 20.80 |
| AH-5 | 1 | 5 | 271.74 | NE | 9.28 |
| | 2 | 10 | 112.60 | NE | 7.09 |
| | 3 | 15 | 81.36 | NE | 4.93 |
| | 4 | 20 | 83.12 | NE | 4.31 |
| | 5 | 25 | 287.23 | NE | 6.79 |
| | 6 | 30 | 675.17 | NE | 10.21 |
| AH -6 | 1 | 5 | 222.78 | 30 | 14.45 |
| | 2 | 10 | 136.36 | 30 | 13.09 |
| | 3 | 15 | 106.67 | 30 | 7.01 |
| | 4 | 20 | 63.49 | 30 | 6.66 |
| | 5 | 25 | 96.15 | 30 | 10.46 |
| | 6 | 30 | 61.95 | 30 | 10.99 |

Table 4-9 OIL AND GREASE ANALYSES 1978 USCOE SOILS INVESTIGATION

NOTES: BGS = below ground surface. NE = not encountered. The oil and grease tests were conducted in accordance with <u>Standard Methods</u> Section 502D, "Extraction Method for Sludge Samples" which uses the Soxhlet extraction apparatus and 1, 1, 2 trichloro - 1, 2, 2 trifloroethane, freon as the extraction solvent.

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| | | Depth | OVA Head Space | TPH mg/kg | Water Level | Moisture | Detection Limit |
|------------|------------|--------|----------------------|--------------|-------------|----------|--------------------|
| Boring No. | Sample No. | (feet) | (ppm) | (dry weight) | (ft BGS) | (%) | (mg/kg) |
| | | | | | | | |
| 23 | 07 | 28 | NR | 137 | 32 | 4 | 66 |
| 24 | 03 | 13 | NR | 142 | 20 | 8 | 68 |
| 25 | 01 | 18 | NR | 136 | 21 | 4 | 66 |
| 26 | 04 | 23 | NR | 114 | 26 | 9 | 70 |
| 27 | 05 | 23 | NR | 115 | 24 | 9 | 79 |
| 28 | 05 | 23 | NR | <27 | 26 | 6 | 27 |
| 29 | 05 | 23 | NR | <32 | 25 | 23 | 32 |
| 30 | 05 | 23 | 3 | 1,153 | 27 | 12 | 28 |
| 31 | 05 | 23 | 2.6 | 5,177 | 26 | 13 | 29 |
| 32 | 05 | 23 | 0 | <30 | 24 | 18 | 30 |
| 33 | 06 | 28 | 0 | <30 | 31 | 16 | 30 |
| 34 | 06 | 28 | NR | 3,662 | 28 | 16 | 82 |
| 34 (Rep) | 06 | 28 | NR | 4,641 | 28 | 23 | 75 |
| 35 | 06 | 28 | 0 | <55 | NE | 12 | 55 |
| 36 | 06 | 28 | 0 | <57 | 30 | 15 | 57 |
| 37 | 06 | 28 | 0 | 2,231 | 30 | 19 | 229 |
| 38 | 06 | 28 | 130 | 2,694 | 29 | 21 | 166 |
| 39 | 05 | 23 | 100 | <62 | 22 | 16 | 62 |
| 39 (Rep) | 05 | 23 | 80 | 1,330 | 22 | 19 | 268 |
| 40 | 06 | 28 | 0 | <56 | 27 | 14 | 56 |

Table 4-10 TOTAL PETROLEUM HYDROCARBONS ESKIMO CREEK SOIL BORINGS

Test Methods:

TPH--E418.1 Moisture--ASTM D2216

NOTES:

Detection limits for individual samples are dependent on their moisture content.

Sample depths for TPH analyses were selected on basis of OVA/HNu measurements performed in the field on samples at the time of collection. Samples with the highest observed readings were submitted for TPH analysis.

Sample replicates were taken by splitting the sample vertically into four quarters and collecting the first quarter as the discrete sample and the third as the replicate.

BG = Background. NE = Water not encountered to a depth of 35 feet. NR = Not Recorded.

CVR193/018

Table 4-11A EP TOXICITY (METALS) ESKIMO CREEK DRUMMED CUTTINGS

| Parameter | Method | Detection | lintte | | 1 | | DC-MM-41 | | Dr. Miteo |
|---|--|---|--|--|--|--|---|--|--------------------|
| | | | 21112 | LA-FIM-3U | DC-MW-38 | DC-MW-39 | & MW-31 | DCMW43 | DUP MW-43 |
| Arsenic Barium Cadmium Chromium Lead Mercury Selenium Silver | 1310 1310 1310 1310 1310 1310 1310 | 0.5 0.5 0.5 0.5 0.02 0.1 | 1/6m 1/6m 1/6m 1/6m 1/6m 1/6m 1/6m 1/6m | <0.5 <0.5 <0.1 <0.5 <0.5 <0.002 <0.1 <0.5 | <pre><0.5 <0.5 <0.1 <0.5 <0.5 <0.5 <0.5 <0.1 <0.1 <0.1 <0.5 <0.1 <0.5 <0.1 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5</pre> | <pre><0.5 <0.5 <0.1 <0.5 <0.5 <0.5 <0.5 <0.1 <0.1 <0.1 <0.5 <0.1 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5</pre> | <pre><0.5 <0.5 <0.1 <0.1 <0.5 <0.5 <0.5 <0.02 <0.1 <0.1 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5 <0.5</pre> | <pre><0.5 <0.5 <0.1 <0.1 <0.5 <0.5 <0.5 <0.02 <0.1 <0.1 </pre> | <pre><0.5</pre> |
| Soll Moisture | D2216 | I | æ | 8.86 | 18.7 | 16.5 | 13.8 | 18.7 | 16.8 |
| | | | | | | | | | |

IV-43

C¹ 73/033-1

Table 4-11B VOLATILE ORGANIC COMPOUNDS (SW8240)

ESKIMO CREEK DRUMMED CUTTINGS

| | | | | | DC-MW41 | | DC-MW89 |
|----------------------------------|-------|---------------|---------|---------|---------|---------|-----------|
| Parameter | Units | DC-MW30 | DC-MW38 | DC-MN39 | 6. MW31 | DC-MW43 | DUP MM-43 |
| | | | | | | | |
| Chloromethane | ng/kg | 5500 | 5200U | 3600J | 5800U | 6200U | 6000U |
| Bronoethane | ng/kg | 550U | 6200U | 6000U | 5800U | 6200U | 6000U |
| Vinyl chloride | ng/kg | 5500 | 5200U | 60000 | 5800U | 6200U | 60000 |
| Chloroethane | ng/kg | 550U | 6200U | 6000U | 5800U | 6200U | 6000U |
| Methylene chloride | ng/kg | 320 | 16000 | 12000 | 13000 | 15000 | 15000 |
| Acetone | ng/kg | 5500 | 8600 | 6000U | 6400 | 7300 | 6000U |
| Carbon disulfide | ng/kg | 270U | 3100U | 30000 | 2900U | 31000 | 30000 |
| 1,1-Dichloroethane | ng/kg | 270U | 3100U | 30000 | 29000 | 3100U | 30001 |
| 1,1-Dichloroethane | ng/kg | 270U | 3100U | 30001 | 2900U | 3100U | 30000 |
| Trans-1,2-dichloroethene | ng/kg | 270U | 3100U | 3000U | 2900U | 31000 | 3000U |
| Chloroform | ng/kg | 2700 | 3100U | 30000 | 2900Ŭ | 3100U | 30000 |
| 1,2-Dichloroethane | ng/kg | 2700 | 31000 | 3000U | 2900U | 31000 | 3000U |
| 2-Butanone | ng/kg | 5500 | 6200U | 60000 | 5800U | 31000 | 6000U |
| 1,1,1-Trichloroethane | ng/kg | 270U | 3100U | 3000U | 2900U | 3100U | 30000 |
| Carbon tetrachloride | ng/kg | 2700 | 3100U | 3000U | 2900U | 31000 | 30000 |
| Vinyl acetate | ug/kg | 2700 | 3100U | 30000 | 2900U | 3100U | 30000 |
| Bromodichloromethane | ng/kg | 2700 | 3100U | 3000U | 2900U | 3100U | 3000U |
| 1,2-Dichloropropane | ng/kg | 27 0 U | 3100U | 30000 | 2900U | 31000 | 3000U |
| Cis-1,3-Dichloropropene | ng/kg | 2700 | 3100U | 3000U | 2900U | 31060 | 30000 |
| Trichloroethene | ng/kg | 2700 | 3100U | 30000 | 2900U | 31000 | 3000U |
| Dibromochloromethane | ng/kg | 2700 | 3100U | 3000U | 2900U | 31000 | 30000 |
| <pre>l,l,2-Trichloroethane</pre> | ng/kg | 270U | 3100U | 3000U | 2900U | 31000 | 3000U |
| Trans-1,3-dichloropropene | ng/kg | 270U | 3100U | 30000 | 2900U | 3100U | 30000 |
| Benzene | ug/kg | 2700 | 3100U | 30000 | 2900U | 31000 | 30000 |
| 2-Chloroethyl vinyl ether | ug/kg | 5500 | 6200U | 6000U | 5800U | 3100U | 6000U |
| Bronoform | ug/kg | 27 OU | 3100U | 3000U | 2900U | 3100U | 30000 |
| | | | | | | | |

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U = Indicates the compound was analyzed for, but not detected. J = Indicates an estimated trace value. DC = Drum cuttings.

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| Parameter Units DC-MM30 DC-MM39 DC-MM41 4-Methyl-2-pentanone ug/kg 550U 6200U 6000U 5800U 3100U 2-Hexanone ug/kg 550U 6200U 6000U 5800U 3100U 1,1,2,2-Tetrachloroethane ug/kg 270U 3100U 3000U 2900U 3100U Tetrachlorethene ug/kg 270U 3100U 3000U 2900U 3100U Toluene ug/kg 270U 3100U 200U 2900U 3100U Toluene ug/kg 270U 3100U 2900U 2900U 3100U Toluene ug/kg 270U 3100U 2900U 3100U 3100U Toluene ug/kg 270U 3100U 2900U 2900U 3100U EthvIbenzene ug/kg 270U 3100U 2900U 3100U 290UU 3100U | | | | | | | | |
|--|------------------------------|--------|----------|---------|---------|-------------------|---------|----------------------|
| 4-Methyl-2-pentanone ug/kg 550U 6200U 5800U 3100U 3100U <th>Parameter</th> <th>Units</th> <th>DC-MM30</th> <th>DC-MW38</th> <th>DC-MN39</th> <th>DC-MW41 & MW31</th> <th>DC-MW43</th> <th>DC-MW89 DUP MM-43</th> | Parameter | Units | DC-MM30 | DC-MW38 | DC-MN39 | DC-MW41 & MW31 | DC-MW43 | DC-MW89 DUP MM-43 |
| 2-Hexanone ug/kg 550U 6200U 5800U 3100U 1/1/2/2-Tetrachloroethane ug/kg 550U 6200U 5800U 3100U 1/1/2/2-Tetrachloroethane ug/kg 270U 3100U 3000U 2900U 3100U Tetrachlorethene ug/kg 270U 3100U 3000U 2900U 3100U Toluene ug/kg 270U 3100U 3000U 2900U 3100U Chlorobenzene ug/kg 270U 3100U 2900U 3100U Ethylbenzene ug/kg 270U 3100U 2900U 3100U | thvl-2-bentanone | 04/01 | 5 6 O II | | HOOOL | | | |
| Z-Hexanone ug/kg 550U 6200U 5800U 3100U 310U | | 64 /6m | 0000 | 00020 | 00000 | 00085 | 31000 | 6000U |
| 1,1,2,2-Tetrachloroethane ug/kg 270U 3100U 3000U 2900U 3100U Tetrachlorethene ug/kg 270U 3100U 3000U 2900U 3100U Toluene ug/kg 270U 3100U 3000U 2900U 3100U Toluene ug/kg 270U 3100U 200U 2900U 3100U Toluene ug/kg 270U 3100U 200U 2900U 3100U Ethvlbenzene ug/kg 270U 3100U 3000U 2900U 3100U | xanone | ug/kg | 5500 | 6200U | 6000U | 58000 | 31000 | 60000 |
| Tetrachlorethene ug/kg 270U 3100U 3000U 2900U 3100U Toluene ug/kg 330 3100U 1500J 2900U 3100U Toluene ug/kg 370U 3100U 1500J 2900U 3100U Chlorobenzene ug/kg 270U 3100U 3000U 2900U 3100U Ethvlbenzene ug/kg 270U 3100U 3000U 2900U 3100U | 2,2-Tetrachloroethane | ug/kg | 270U | 3100U | 30000 | 2900U | 3100U | 30000 |
| Toluene ug/kg 330 3100U 1500J 2900U 3100U Chlorobenzene ug/kg 270U 3100U 3000U 2900U 3100U Ethylbenzene ug/kg 270U 3100U 3000U 2900U 3100U | achlorethene | ug/kg | 270U | 31000 | 30000 | 2900U | 3100U | 30000 |
| Chlorobenzene ug/kg 2700 31000 30000 29000 31000 Ethylbenzene ug/kg 400 31000 20000 31000 | ene | ng/kg | 330 | 3100U | 1500J | 2900U | 31000 | 30001 |
| Ethylhenzene un/ka 400 suom soom soom | robenzene | ng/kg | 2700 | 31000 | 30000 | 29001 | 31000 | 30001 |
| | lbenzene | ug/kg | 400 | 31000 | 30000 | 29000 | 31000 | 30005 |
| Styrene ug/kg 2700 31000 30000 29000 31000 | ene | ng/kg | 270U | 31000 | 30000 | 29000 | 31000 | 30001 |
| Xylenes (o+p) ug/kg 1800 3100U 3000U 2900U 3100U | nes (o+p) | ng/kg | 1800 | 3100U | 30000 | 2900U | 3100U | 30000 |
| Xylene (p) ug/kg 1500 3100U 2900U 2900U 3100U | ne (p) | ng/kg | 1500 | 3100U | 30000 | 2900U | 31000 | 30000 |

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J = Indicates an estima DC = Drum cuttings.

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Table 4-11C SEMIVOLATILE ORGANICS

(SW8270) ESKIMO CREEK DRUMMED CUTTINGS

| | | | | | DC-MW41 | | |
|----------------------------|-------|----------|--------------|----------|---------|--------------|--------------|
| Parameter | Units | DC-MW-30 | DC-MW38 | DC-MM-39 | £ MW31 | DC-MW43 | DC-MM-89 |
| | | | | | | | |
| N-nitroso-dimethylamine | ng/kg | 360U | 410U | 4000 | 380U | 410U | 4 00U |
| Phenol | ng/kg | 360U | 410U | 4000 | 380U | 41 OU | 400U |
| Antline | ng/kg | 360U | 410U | 400U | 380U | 410U | 400U |
| Bis(2-chloroethyl)ether | ng/kg | 360U | 410U | 4000 | 3800 | 410U | 400U |
| 2-Chlorophenol | ng/kg | 360U | 410U | 4000 | 3800 | 4 10U | 400U |
| 1,3-D1chlorobenzene | ug/kg | 360U | 410U | 400U | 3800 | 410U | 400U |
| 1,4-D1chlorobenzene | ng/kg | 360U | 410U | 400U | 3800 | 410U | 400U |
| Benzyl alcohol | ng/kg | 360U | 410U | 400U | 380U | 41 OU | 400U |
| 1,2-Dichlorobenzene | ng/kg | 360U | 410U | 400U | 380U | 4100 | 400U |
| 2-Methylphenol | ng/kg | 360U | 41 OU | 4000 | 380U | 41 OU | 4000 |
| Bis(2-chloroisopropyl) | ug/kg | 360U | 41 OU | 4000 | 380U | 4100 | 400U |
| 4-Methylphenol | ng/kg | 36 OU | 41 OU | 4000 | 380U | 41 OU | 4 00U |
| N-nitroso-di-n-propylamine | ng/kg | 360U | 4100 | 400U | 380U | 4 10U | 4 00U |
| Hexachloroethane | ng/kg | 360U | 410U | 400U | 380U | 41 OU | 400U |
| Nitrobenzene | ng/kg | 360U | 4 10U | 400U | 380U | 410U | 4 00U |
| Isophrone | ng/kg | 360U | 4100 | 4000 | 380U | 4100 | 400U |
| 2-N1trophenol | ng/kg | 360U | 410U | 400U | 380U | 410U | 400U |
| 2,4-Diethylphenol | ng/kg | 360U | 410U | 400U | 3800 | 4100 | 400U |
| Benzoic acid | ng/kg | 18000 | 20000 | 19000 | 19001 | 2000U | 1900 |
| Bis(2-chloroethoxy)methane | ng/kg | 360U | 410U | 4000 | 3800 | 410U | 400U |
| 2,4-D1chlorophenol | ng/kg | 360U | 410U | 400U | 3800 | 4 10U | 400U |
| 1,2,4-trichlorobenzene | ng/kg | 360U | 410U | 400U | 3800 | 410U | 400U |
| Naphthalene | ng/kg | 360U | 1300 | 490 | 380U | 4 10U | 400U |
| 4-Chloroaniline | ng/kg | 360U | 410U | 400U | 3800 | 410U | 400U |
| Hexachlorobutadiene | ng/kg | 360U | 4100 | 400U | 380U | 410U | 400U |
| 4-Chloro-3-methylphenol | ng/kg | 3600 | 410U | 400U | 380U | 410U | 400U |
| 2-Methylnaphthalene | ng/kg | 8500 | 5100 | 2200 | 3700 | 410U | 400U |
| Hexachlorocyclopentadiene | ng/kg | 360U | 41 OU | 400U | 3800 | 410U | 400U |
| | | | | | | | |

U = Indicates the compound was analyzed for, but not detected. J = Indicates an estimated trace value. DC = Drum cuttings.

| | | | Table 4-11C (Continued) | | | | |
|-------------------------------------|-------|---------------|----------------------------|--------------|---------------|---------|--------------|
| | | | | | DC-MW41 | | |
| Parameter | Units | DC-MM-30 | DC-MW38 | DC-MN-39 | £ MW31 | DC-MM43 | DC-MM-89 |
| 2,4,6-Trichlorophenol | ng/kg | 360U | 4100 | 4000 | 380U | 4100 | 4000 |
| 2,4,5-Trichlorophenol | ng/kg | 1800U | 2000U | 1900U | 1900 | 2000 | 10001 |
| 2-Chloronaphthalene | ng/kg | 360U | 4100 | 4000 | 380U | 410U | 400U |
| 2-Nitroailine | ng/kg | 1800U | 2000U | 1900 | 1900 | 20000 | 1900U |
| Dimethyl phthalate | ng/kg | 360U | 410U | 400U | 380U | 4100 | 400U |
| Acenaphthene | ng/kg | 36 0 U | 410U | 400U | 3800 | 410U | 400U |
| 3-nitroaniline | ng/kg | 18000 | 20000 | 19001 | 19000 | 2000U | 19001 |
| Acenaphthene | ng/kg | 3600 | 4100 | 400U | 380U | 41 OU | 4000 |
| 2,4-Dinitrophenol | ng/kg | 1800U | 2000 | 1900U | 1900 | 20001 | 1900U |
| 4-Nitrophenol | ng/kg | 18000 | 20000 | 1900U | 1900U | 20000 | 10061 |
| Dibenzofuran | ng/kg | 360U | 4100 | 4000 | 380U | 410U | 400U |
| 2,4-Dinitrotoluene | ng/kg | 360U | 410U | 400U | 3800 | 410U | 400U |
| 2,6-Dinitrotoluene | ng/kg | 360U | 4100 | 400U | 380U | 410U | 400U |
| Diethyl phthalate | ng/kg | 360U | 410U | 400U | 380U | 4100 | 4000 |
| 4-Chlorophenyl phenyl | | | | | | | |
| ether | ng/kg | 360U | 410U | 400U | 380U | 410U | 400U |
| Fluorene | ng/kg | 630 | 290J | 4000 | 330J | 410U | 400U |
| 4-Nitroaniline | ng/kg | 18000 | 2000 | 19001 | 19001 | 2000U | 19001 |
| 4,6-Dinitro-2-methylphenol | ng/kg | 18000 | 20000 | 1900 | 19001 | 20001 | 19001 |
| N-nithosodiphenyl a mine | ng/kg | 360U | 410U | 400U | 3800 | 410U | 400U |
| 4-Bromophenyl phenyl | | | | | | | |
| ether | ng/kg | 360U | 4100 | 400U | 38 0 U | 410U | 400U |
| Hexachlorobenzene | ng/kg | 3600 | 410U | 400U | 380U | 410U | 400U |
| Pentachlorophenol | ng/kg | 18000 | 2000U | 1900U | 19001 | 20000 | 1900U |
| Phenanthrene | ng/kg | 210J | 410U | 400U | 3800 | 4100U | 400U |
| Anthracene | ng/kg | 360U | 410U | 400U | 380U | 410U | 400U |
| D1-n-butyl phathalate | ng/kg | 360U | 41 OU | 400U | 380U | 4100 | 400U |
| Fluoranthene | ng/kg | 3600 | 410U | 400U | 3800 | 410U | 400U |
| Benzidine | ng/kg | 18000 | 2000U | 1900U | 19001 | 20000 | 19001 |
| Pyrene | ng/kg | 360U | 410U | 400Ŭ | 380U | 410U | 400U |
| Butyl benzyl phthalate | ng/kg | 360U | 410U | 400U | 1800 | 870 | 1200 |
| 3,3-dichlorobenzidine | ng/kg | 720U | 810U | 800U | 7700 | 8100 | 800U |

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U = Indicates the compound was analyzed for, but not detected. J = Indicates an estimated trace value. DC = Drum cuttings.

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| DC-MM-89 4000 4000 4000 4000 4000 4000 4000 40 | 200 |
|---|-----|
| DC-MH43 410U 1100 410U 410U 410U 410U 410U 410U | |
| DC-MM41 E MM31 380U 380U 380U 380U 380U 380U 380U 380U 380U 380U 380U | |
| <u>DC-MM-39</u> 400U 1500 400U 400U 400U 400U 400U 400U 400U | |
| DC-HR438 410U 410U 410U 410U 410U 410U 410U 410U | |
| <u>рс-ни30</u> 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U 36.0U | |
| <u>Units</u> ug/kg ug/kg ug/kg ug/kg ug/kg ug/kg ug/kg | |
| Parameter Benzo(a) anthracene Bis(2-ethylhexyl)phthalate Chrysene Di-n-octyl phthalate Benzo(b)fluoranthene Benzo(b)fluoranthene Benzo(a)pyrene Indeno(1,2,3-cd)pyrene Dibenz(a,h)anthracene Benzo(h,h,1)perylene | |

U=Indicates the compound was analyzed for, but not detected. J=Indicates an estimated trace value, DC=drum cuttings.

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4.1.5.1 Groundwater Results

4.1.5.1.1 Eskimo Creek Seep

Twenty-seven monitoring wells and two base water wells were sampled for water quality in June 1988. In the shallow unconfined aquifer, 20 newly installed wells and 7 existing USCOE wells were analyzed for total petroleum hydrocarbons (TPH) or total organic carbon (TOC), alkalinity, common anions, lead, and total dissolved solids. (Only sample bottles broken in transit were analyzed for TOCs to try and verify the presence of hydrocarbons.) Eight of these shallow wells were also analyzed for aromatic volatile compounds. Five of the eight wells were selected for aromatic volatile analysis by visual contamination inspection and nearness to deep wells. Three of the wells were selected outside the estimated contaminated zone to try and better define the plume of contamination. Table 4-12 presents results for these analyses.

The extent of TPH contamination is presented in Figure 4-13. The highest value (5,666 ppm) is reported at MW-22 where substantial floating product (>1 foot) has previously been measured (April 1987). Other detected values closely correlate with the soil gas plume (Figure 4-1). The extent of hydrocarbon contamination can be traced from well MW-39 through MW-22 and to the southeast toward the Eskimo Creek seep.

A comparison of the TPH concentrations in the soils (Figure 4-12) to the groundwater (Figure 4-13) indicates detected concentrations in the borehole soils at wells where groundwater contained less than the detection limit of 1 ppm. This suggests significant quantities of petroleum contamination may be sorbed to unsaturated soils. Also, the low aqueous solubilities of heavy petroleum hydrocarbons may limit their detection with the TPH analysis.

Aromatic compound data are plotted with benzene, toluene, and xylene concentrations in Figure 4-14. The extent and relative concentrations of aromatics across the site correlate with the TPH data. The highest values are found in MW-30 with 77 ppb benzene, 130 ppb toluene, and 520 ppb xylene. USCOE wells including MW-22 were not analyzed for aromatics because of uncertainty in their construction (i.e., glued joints).

Each of the shallow and intermediate monitoring wells were measured with an oil/water interface probe to check for light and/or dense nonaqueous phase liquids (NAPLs) on June 12, 1988. NAPLs were not detected in any of the wells. However, these measurements cannot be considered entirely representative due to the relatively short period of time between well development (or redevelopment/purging of USCOE wells) and

FIELD PARAMETERS, ALKALINITY, COMMON ANIONS, LEAD, TOTAL DISSOLVED SOLIDS, TOTAL PETROLEUM HYDROCARDONS, AND PURGEABLE AROMATICS ESKIMO CREEK SHALLOM ADUIFER Table 4-12

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| 1 | | | Detection | KS-MM-01 | KS-MM-06 | 60-MM-SX | KS-MM-10 | KS-MW-16 | KS-ERMW |
|-------------------------|-----------|-------|-----------|----------|----------|----------|----------|----------|---------|
| Parameter | Method | Units | Limits | 6/5/88 | 6/7/88 | 6/6/88 | 6/1/88 | 6/6/88 | 6/6/88 |
| Temperature | FIELD | U | N/A | 4.0 | 4.0 | 5.0 | 5.0 | 5.0 | na |
| pH | FIELD | ı | N/A | 5.8 | 6.4 | 6.8 | 6.4 | 6-9 | na |
| Specific Conductivity | FIELD | SOHMU | N/A | 150 | 170 | 420 | 295 | 500 | na |
| Appearance | FIELD | ı | N/A | TURBID | TURBID | TURBID | TURBID | TURBID | na |
| Odor (fuel) | FIELD | ŀ | N/A | NONE | YES | YES | YES | YES | na |
| Lead | 3005/7421 | ug/1 | 2 | na | <5 | 16 | 110 | 17 | <5 |
| Ortho Phosphate | E365.2 | I/gm | 0.5 | na | 0.62 | 0.84 | 2.66 | 0.32 | 10.0 |
| Alkalinity | A403 | l∕2m | 20 | na | 128 | 354 | 100 | 415 | 2.0 |
| Chloride | 9252 | ng/1 | 1 | na | 6.9 | 6,5 | 22.7 | 15.2 | 4 |
| Sulfate | 9038 | 1/bm | ľ | na | 8.5 | 19.4 | 73.5 | 50.3 | 1.0 |
| Fluoride | E340.2 | mg/1 | 0.05 | na | <0.05 | <0.05 | <0.05 | 0.05 | <0.05 |
| Total Diss. Solids | E160.1 | ng/l | 1 | na | 193 | 444 | 316 | 536 | 5 |
| Nitrate-Nitrite | 9200 | ng/1 | 0.05 | na | 2.26 | 0.36 | 10.5 | 0.22 | 0.08 |
| TPH | E418.1 | ng/1 | 1 | na | 23.6 | m | 4.2 | 6 | ę |
| 700 | | 1/2m | J | na | na | na | na | na | na |
| Benzene | SW8020 | 1/Ju | 1 | na | na | Da | na | na | |
| Toluene | SW8020 | 1/bn | 1 | na | na | БЛ | na | па | |
| Ethyl Benzene | SW8020 | 1/gu | 1 | ра | ра | na | na | Пâ | |
| Xy lene | SN802U | 1/6n | l | na | na | Пà | na | ាង | |
| Chlorobenzene | SW8020 | 1/bn | 1 | na | na | na | na | na | |
| 1,4-D1chlorobenzene | SW8020 | ug/1 | 1 | na | na | na | ពង | រង | |
| 1,3-Dichlorobenzene | SW8020 | 1/6n | 1 | na | na | na | ពង | na | |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | 1 | na | na | na | na | na | |
| na = not analvzed/samp | led | | | | | | | | |
| MW = Monitoring Well | | | | | | | | | |
| EB = Equipment Blank | | | | | | | | | |
| TB = Trip Blank | | | | | | | | | |
| AC = Amblent Condition: | s Blank | | | | | | | | |
| N/A = Not Applicable | | | | | | | | | |

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Table 4-12 (Continued)

| | | | Detection | KS-MW-18 | KS-MW-21 | 66-M-SX | KS-ERMW | KS-MW-22 | KS-MW-23 |
|-------------------------|-----------|-------|-----------|----------|----------|--------------|---------|----------|----------|
| Parameter | Method | Units | Limits | 6/6/88 | 6/5/88 | DUP MM-21 | 6/5/88 | 6/8/88 | 6/16/88 |
| Temperature | FIELD | U | N/A | 4.0 | 5.0 | Da | na | 4.0 | 14 |
| Hd | FIELD | ı | N/A | 5.9 | 6.3 | na | na | 6.0 | 7.4 |
| Specific Conductivity | FIELD | SOHMU | N/A | 550 | 460 | Da | na | 500 | 380 |
| Appearance | FIELD | ı | N/A | TURBID | TURBID | na | na | TURBID | YOOUM |
| Odor (fuel) | FIELD | • | N/A | YES | YES | na | na | YES | NONE |
| Lead | 3005/7421 | ug/1 | ŝ | 64 | 85 | 48 | \$\$ | 370 | na |
| Ortho Phosphate | E365.2 | mg/l | 0.5 | 0.10 | 2.67 | 2.22 | 0.01 | 1.71 | na |
| Alkalinity | A403 | mg/l | 20 | 386 | 386 | 391 | 1.8 | 394 | na |
| Chloride | 9252 | ng/1 | 1 | 11.6 | 10.8 | 11.6 | 4 | 6*6 | na |
| Sulfate | 9038 | mg/l | 1 | 39.7 | 27.9 | 28.2 | ₽ | 3.4 | na |
| Fluoride | E340.2 | mg/l | 0.05 | <0.05 | 0.11 | <0.05 | <0.05 | <0.05 | na |
| Total Diss. Solids | E160.1 | mg/1 | 1 | 542 | 558 | 526 | ₽ | 489 | na |
| Nitrate-Nitrite | 9200 | mg/1 | 0.05 | 0.21 | 0.13 | 0.11 | 0.06 | 0.78 | na |
| TPH | E418.1 | mg/l | 1 | 53 | £ | < . 8 | ß | 5666 | na |
| 70C | | mg/l | 1 | na | na | na | na | na | Пâ |
| Benzene | SWB020 | ug/1 | 1 | na | na | na | | na | 4 |
| Toluene | SW8020 | 1/Ju | 1 | na | រាង | лa | | na | 4 |
| Ethyl Benzene | SW8020 | 1/bn | 1 | na | na | na | | na | ₽ |
| Xy lene | SW8020 | ug/1 | 1 | na | na | na | | na | ₽ |
| Chlorobenzene | SW8020 | ug/1 | 1 | na | na | na | | na | ₽ |
| 1,4-Dichlorobenzene | SW8020 | ug/1 | l | na | na | na | | na | ₽ |
| 1,3-Dichlorobenzene | SW8020 | ug/1 | 1 | na | na | na | | na | ₽ |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | 1 | na | na | na | | па | 4 |
| na = not analvzed/samn) | leđ | | | | | | | | |
| MW = Monitoring Well | 5 | | | | | | | | |
| EB = Equipment Blank | | | | | | | | | |
| TB = Trip Blank | | | | | | | | | |
| AC = Amblent Condition: | s Blank | | | | | | | | |
| N/A = Not Applicable | | | | | | | | | |

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Table 4-12 (Continued)

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| | | | (Contin | ued) | | | | |
|--|--------------------|----------|---------------|-------------|----------------|-------------|----------|-----------|
| | | | Detection | KS-MW-24 | KS-MW-25 | KS-MW-26 | KS-MM-27 | Vo-Mi-SX |
| Parameter | Method | Units | Limits | 6/12/88 | 6/10/88 | 6/13/88 | 6/13/88 | DUP MM-27 |
| Temperature | FIELD | IJ | N/A | и С | c | | | |
| Hq | FIELD | • | N/N | | n . | C•7 | 7 | na |
| Specific Conductuates | erer 0 | DOTION - | | P. 0. | 11.0 | 6.25 | 6.3 | na |
| | LIEW | SURAU | N/A | 108 | 2 | 500 | 140 | na |
| | | 1 | N/A | TURBID | TURBID | TURBID | TURBID | na |
| Odor (fuel) | FIELD | I | N/A | NONE | NONE | NONE | NONE | na |
| Lead | 3005/7421 | 1/bn | Ś | 28 | 21 | <u>ر</u> ۶ | 37 | L |
| Ortho Phosphate | E365.2 | 1/Jm | 0.5 | 0.59 | 0.38 | ; ; ; | | 0 X |
| Alkalinity | A403 | mg/l | 20 | 83.8 | 0.94 | 17.0 | 66.U | 0.26 |
| Chloride | 9252 | 1/DM | ¦ - | | | 141 | 111 | 116 |
| Sulfate | 9038 | | 4 | | D•7 | e.02 | 5.7 | 7.5 |
| Fluoride | F340. 2 | | 1 0 | 2.0 V 05 | 2 ° 0 | 10./ | 4.6 | 4.5 |
| Total Diss. Solids | E160.1 | | CD * D | 50.02 | د ں ، ں | 0.06 | 0.05 | 0.05 |
| Nitrate-Nitrite | | 1/51 | 1 0 | 5/T | 145 | 253 | 166 | 173 |
| The second s | 0076 | 7/5m | 0.05 | 1.41 | 1.97 | 1.64 | 0.63 | 0.63 |
| 15.U | E418.1 | mg/1 | - | 4 | 4 | 4 | ₽ | 4 |
| In | | mg/1 | 1 | na | na | na | na | na |
| Benz ene | SW8020 | 1/bn | 1 | ពិធ | ç | đ | ŝ | ł |
| Toluene | SW8020 | 1/J | - | | ' c | | | 2 |
| Ethyl Benzene | SW8020 | na/1 | | | 7 5 | | BI | na |
| Xylene | SW8020 | [/bii | • - | 2 | 7、 | | na | Вũ |
| Chlorobenzene 、 | SW8020 | 1/011 | 4 - | <u>a</u> ; | 7 : | na | ца | na |
| 1.4-Dichlorobenzene | CURD O | 1,20 | 4 - | | 7 : | na | na | Пâ |
| 1.3-Dichlorobenzene | OCOBINS OF COMPANY | 1/5n | -ı, | na | 4 | na | лà | ла |
| | 010000 | | - | ПA | 4 | Пa | na | Da |
| 1/2-DICUTOLODEUZEUE | OZOBWS | ng/1 | 1 | na | 4 | na | na | ກa |
| na = not analyzed/sampled | | | | | | | | |
| MW = Monitoring Well | | | | | | | | |
| EB = Equipment Blank | | | | | | | | |
| TB = Trip Blank | | | | | | | | |
| AC = Amblent Conditions B1 | lank | | | | | | | |
| N/A = Not Applicable | | | | | | | | |

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Table 4-12 (Continued)

| | | | Detection | KS-MW-28 | KS-MW-29 | KS-TBMW | KS-MM-30 | KS-EQMM |
|--------------------------------|-----------|--------|------------------|----------|----------|---------|----------|------------|
| Parameter | Method | Units | Limits | 6/10/89 | 6/12/88 | 6/9/88 | 6/9/88 | 6/9/88 |
| Temperature | FIELD | ပ ပ | N/A | e | 7.5 | na | ŝ | na |
| рн | FIELD | 1 | N/A | 5.59 | 6.31 | na | 6.51 | na |
| Specific Conductivity | FIELD | SOHMU | N/A | 230 | 265 | Bu | 425 | na |
| Appearance | FIELD | ı | N/A | TURBID | TURBID | na | TURBID | na |
| Odor (fuel) | FIELD | • | N/A | NONE | NONE | na | YES | na |
| Lead | 3005/7421 | ug/1 | ĸ | 8 | 21 | na | 22 | <5 |
| Ortho Phosphate | E365.2 | mg/1 | 0.5 | 0.17 | <0.01 | na | 0.51 | 0.01 |
| Alkalinity | 403 | mg/1 | 20 | 174 | 131 | na | 354 | 3.6 |
| Chloride | 9252 | mg/1 | 1 | 3.6 | 10.7 | na | 6.0 | 1> |
| Sulfate | 9038 | 1/20 | 1 | 21.8 | 18.7 | na | 8,9 | 1 > |
| Fluoride | E340.2 | mg/1 | 0.05 | 0.05 | 0.05 | na | 0.07 | <0.05 |
| Total Diss. Solids | E160.1 | mg/1 | 1 | 285 | 331 | ងព | 461 | 2 |
| Nitrate-Nitrite | 9200 | mg/1 | 0.05 | 2.56 | 11.4 | na | 1.0 | 0.15 |
| ТРН | E418.1 | mg/l | 7 | ₽ | 4 | na | 34.2 | 2.6 |
| 700 | | mg/1 | 1 | na | па | na | na | na |
| Benzene | SW8020 | 1/gu | 1 | 4 | na | 4 | 77 | 4 |
| Toluene | SW8020 | ug/1 | 1 | 4 | na | 4 | 130 | ₽ |
| Ethyl Benzene | SW8020 | ug/1 | 1 | 4 | na | ₽ | 130 | 4 |
| Xy lene | SW8020 | 1/gu | l | 4 | na | 41 | 520 | 1> |
| Chlorobenzene | SW80.20 | 1/gu | 1 | 4 | D.B. | 4 | 4 | 4 |
| <pre>l,4-Dichlorobenzene</pre> | SW8020 | ug/1 | - | 4 | na | 4 | ₽ | ₽ |
| 1,3-Dichlorobenzene | SW8020 | 1/6n | 1 | 4 | Па | 4 | 4 | ₽ |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | 1 | 4 | na | 4 | 4 | 4 |
| na = not analvzed/sample | eđ | | | | | | | |
| MW = Monitoring Well | | | | | | | | |
| EB = Equiptment Blank | | | | | | | | |
| TB = Trip Blank | | | | | | | | |
| AC = Ambient Conditions | Blank | | | | | | | |
| N/A = Not Applicable | | | | | | | | |

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Table 4-12 (Continued)

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| | | | (CONCIN | uea) | | | |
|---------------------------|-----------|-------|-----------|-------------|---------|----------|---------|
| | | | Detection | KS-MW-31 | KS-ACMW | KS-MW-32 | KS-EQMW |
| Parameter | Method | Units | Limits | 6/9/88 | 6/9/88 | 6/13/88 | 6/13/88 |
| Temperature | FIELD | U | N/A | 4 | na | 4 | na |
| hq | FIELD | ı | N/A | 6.67 | na | 6.05 | na |
| Specific Conductivity | FIELD | SOHMU | N/A | 185 | na | 162 | pa |
| Appearance | FIELD | ı | N/A | TURBID | na | TURBID | na |
| Odor (fuel) | FIELD | 1 | N/A | YES | na | NONE | រង |
| Lead | 3005/7421 | 1/gu | ß | \$ 5 | រាង | 10 | <5 |
| Ortho Phosphate | E365.2 | mg/1 | 0.5 | 0.10 | na | 0.61 | <0.01 |
| Alkalinity | 403 | mg/1 | 20 | 130 | na | 105 | 2.5 |
| Chloride | 9252 | mg/l | 1 | 14.3 | na | 5.1 | 4 |
| Sulfate | 9038 | mg/1 | 1 | 12.8 | na | 7.2 | 12 |
| Fluoride | E340.2 | mg/1 | 0.05 | 0.06 | na | 0.05 | <0.05 |
| Total Diss. Solids | E160.1 | mg/1 | 1 | 222 | ពន | 206 | 23 |
| N1trate-N1trite | 9200 | mg/1 | 0.05 | 0.46 | na | 6.29 | 0.13 |
| HdL | E418.1 | ng/1 | г | 1.0 | na | 4 | 4 |
| TOC | | mg/1 | I | na | na | Пð | na |
| Benzene | SW8020 | ug/1 | 1 | m | 4 | IJа | ង |
| Toluene | SW8020 | ng/1 | 1 | 14 | 4 | na | na |
| Ethyl Benzene | SW8020 | ug/1 | г | 31 | 4 | na | na |
| Xy lene | SW8020 | ug/l | 1 | 170 | 4 | na | na |
| Chlorobenzene | SW8020 | ug/1 | 7 | ₽ | 4 | na | na |
| 1,4-Dichlorobenzene | SW8020 | ug/l | 1 | 4 | 4 | na | na |
| 1,3-Dictlorobenzene | SW8020 | ug/1 | 1 | 4 | 4 | na | na |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | I | 4 | 4 | na | na |
| na = not analyzed/sample(| ģ | | | | | | |
| MW = Monitoring Well | | | | | | | |
| EB = Equiptment Blank | | | | | | | |
| TB = Trip Blank | | | | | | | |
| AC = Amblent Conditions | Blank | | | | | | |
| N/A = Not Applicable | | | | | | | |

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|-------------------------|-----------|-------|-----------|----------|----------|---------|----------|----------|
| | | | Detection | KS-MN-33 | KS-MW-34 | KD-EQMW | KS-ACMW | KS-MM-36 |
| Parameter | Method | Units | Limits | 6/9/88 | 6/10/88 | 6/10/88 | 6/10/88 | 6/14/88 |
| Temperature | FIELD | υ | N/A | 6.6 | ŝ | ពង | na | 3.5 |
| рн | FIELD | ı | N/A | 6.27 | 7.08 | na | na | 6.5 |
| Specific Conductivity | FIELD | SOHMU | N/A | 210 | 475 | na | na Na | 180 |
| Appearance | FIELD | ı | N/A | TURBID | TURBID | na | na | TURBID |
| Odor (fuel) | FIELD | ı | N/A | NONE | YES | ពង | na | NONE |
| Lead | 3005/7421 | t/bn | ŝ | S | 80 | \$5 | na | Ş |
| Ortho Phosphate | E365.2 | mg/1 | 0.5 | 0.30 | 0.48 | 10.0 | na | 0.32 |
| Alkalinity | 403 | mg/1 | 20 | 131 | 347 | 3.0 | na | 109 |
| Chloride | 9252 | mg/1 | 1 | 5.4 | 14.1 | 4 | na | 13.2 |
| Sulfate | 9038 | 1/Jm | 1 | 16.3 | 16.4 | 4 | na | 8.4 |
| Fluoride | E340.2 | mg/1 | 0.05 | 0.05 | 0.06 | 0.05 | na | 0.07 |
| Total Diss. Solids | E160.1 | mg/1 | I | 270 | 532 | 12 | na | 264 |
| Nitrate-Nitrite | 9200 | mg/1 | 0.05 | 3.34 | 0.22 | . 14 | na | 8.24 |
| ТРН | E418.1 | mg/1 | l | 4 | 31.4 | 4 | па | <1 |
| TOC | | mg/1 | 1 | na | na | na | na | na |
| Benzene | SWB020 | ug/1 | 1 | 4 | 5 | 4 | 4 | na |
| Toluene | SW8020 | ng/1 | 1 | ₽ | 10 | 4 | 4 | па |
| Ethyl Benzene | SW8020 | 1/6n | l | 4 | 4 | 4 | 4 | na |
| Xy lene | SW8020 | 1/5n | 1 | ₽ | 78 | ₽ | 4 | na |
| Chlorobenzene | SW80.20 | 1/bn | 1 | 1> | 41 | 4 | ₽ | na |
| 1,4-Dichlorobenzene | SW8020 | 1/6n | 1 | 1> | 4 | 4 | 4 | na |
| 1,3-Dichlorobenzene | SW8020 | 1/Jn | 1 | 1> | 4 | 4 | 4 | na |
| l,2-Dichlorobenzene | SW8020 | ug/1 | I | ₽ | 4 | 41 | 4 | na |
| | | | | | | | | |
| na = not analyzed/sampl | eđ | | | | | | | |
| MW = Monitoring Well | | | | | | | | |
| EB = Equipment Blank | | | | | | | | |
| TB = Trip Blank | | | | | | | | |
| N/A = Not Applicable | | | | | | | | |

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| | | | (Contin | ued) | | | | |
|--------------------------|-----------|--------------|-----------|------------|----------|----------|---------|----------|
| | | | Detection | KS-WM-37 | KS-MM-38 | KS-MW-39 | KS-TBMW | KS-MM-40 |
| Parameter | Method | Units | Limits | 6/14/88 | 6/11/88 | 6/11/88 | 6/11/88 | 6/12/88 |
| Temperature | FIELD | υ | N/A | 4 | æ | e | na | m |
| pH | FIELD | ı | N/A | 6.6 | 5.88 | 6.52 | na | 6.92 |
| Specific Conductivity | FIELD | SOHMU | N/A | 245 | 410 | 310 | na | 80 |
| Appearance | FIELD | ı | N/A | TURBID | TURBID | TURBID | na | TURBID |
| Odor (fuel) | FIELD | , | N/A | NONE | YES | YES | na | NONE |
| Lead | 3005/7421 | 1/Jn | S | ć 5 | 14 | 10 | na | 6 |
| Ortho Phosphate | E365.2 | mg/1 | 0.5 | 0.16 | 0.95 | 0.85 | na | 0.60 |
| Alkalinity | A403 | 1/gm | 20 | 98.0 | 309 | 264 | па | 56.0 |
| Chloride | 9252 | 1/2m | Ч | 22.1 | 6.6 | 6.0 | na | 4.6 |
| Sulfate | 9038 | mg/1 | ŗ | 55.9 | 16.1 | 4 | na | 3.1 |
| Fluoride | E340.2 | 1/5 u | 0.05 | 0.07 | 0.06 | 0.06 | na | 0.06 |
| Total Diss. Solids | E160.1 | 1/9m | 1 | 311 | 465 | 409 | na | 162 |
| Nitrate-Nitrite | 9200 | 1/2m | 0.05 | 6.20 | 0.22 | 0.17 | 112 | 0.20 |
| Hal | E418.1 | mg/1 | 1 | 4 | 127.7 | 14.9 | D.a | 41 |
| 100 | | mg/1 | I | na | D.C. | na | na | Ла |
| Benzene | SW8020 | 1/gu | 1 | na | Q | 23 | 4 | na |
| Toluene | SW8020 | ug/1 | 7 | Pu | 20 | 7 | 4 | na |
| Ethyl Benzene | SW8020 | ng/1 | 1 | ពង | 21 | 29 | 4 | na |
| Xy lene | SW8020 | 1/5n | 7 | na | 76 | 85 | ₽ | na |
| Chlorobenzene | SW8020 | 1/bn | I | na | 4 | 12 | ₽ | na |
| 1,4-Dichlorobenzene | SW8020 | ug/1 | 1 | na | 4 | 12 | 4 | Bu |
| 1,3-D1chlorobenzene | SW8020 | 1/gu | 1 | na | ₽ | 12 | ₽ | na |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | I | na | 4 | 1> | 4 | na |
| na = not analvzeđ/samole | pe | | | | | | | |
| WW = Monitoring Well | | | | | | | | |
| EB = Equipment Blank | | | | | | | | |
| TB = Trip Blank | | | | | | | | |
| AC = Ambient Conditions | Blank | | | | | | | |
| N/A = Not Applicable | | | | | | | | |

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LEGEND

MW • NEW SHALLOW AQUIFER MONITORING WELL LOCATION AND IDENTIFICATION

30 TOTAL PETROLEUM HYDROCARBON (ppm)

FIGURE 4-13

SHALLOW GROUNDWATER CONTAMINATION, TOTAL PETROLEUM HYDROCARBON

JUNE 1988 KING SALMON AIR FORCE STATION KING SALMON, ALASKA

CHM HILL-



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FIGURE 4-14

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SHALLOW GROUNDWATER CONTAMINATION-AROMATICS JUNE 1988

KING SALMON AIR FORCE STATION KING SALMON, ALASKA

CHM HILL-

conducting the measurements. These measurements should be repeated several weeks after the wells have been disturbed to allow the wells to recover to static conditions. As an example, MW-22 has contained significant NAPLs. However, the well was constructed with the top of the screen slightly below the water table (USCOE, 1978) and accumulation of product will require a drop in the regional water levels to expose the screen.

Inorganic analyses indicate high levels of lead (maximum of 370 ppb in MW-22) correlate with the high organic contamination. Monitoring wells with nondetects for the TPH and aromatics generally contain less than or near detection limit levels of 5 ppb lead.

A cross section showing hydrogeologic conditions at the Eskimo Creek seep is presented in Figure 4-5. Groundwater flow from beneath the station discharges into the Eskimo Creek flood plain. Contaminated groundwater may discharge far into the flood plain through the sand deposits. Floating product would be expected to discharge near the contact between the cliff and flood plain.

Intermediate Aquifer

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Groundwater from the three intermediate wells (MW-41, 42, and 43) did not have any detectable aromatics (Table 4-13). MW-41 is reported at less than the detection limit of 1 ppm for TPH. MW-42 and MW-43 contained 1.7 and 7.2 ppm of TOC, respectively, which are relatively low values. Background sample analyses should be obtained for comparisons.

Inorganic analyses for the intermediate wells did not indicate the aquifer was contaminated. However, the pH for well MW-43 was elevated (10.1); a condition that often results from grout contamination. Future groundwater analyses will indicate if there is a problem with grout in this well.

Deep Aquifer

Base wells KS-4 and KS-5 were sampled and analyzed for aromatic organics and TPHs. Aromatic values were all below the detection limit of 1 ppm. Both wells, however, contained detectable levels of TPH (KS-4 at 27 ppm and KS-5 at 6.5 ppm) (Table 4-14).

As discussed in Section III, the development water was tested for purgeable aromatics (SW8020) and lead (SW7421). The results of the analyses are presented in Table 4-15. The analyses indicated that the contamination was below the MCLs, and subsequently the water was disposed of through the base oil/water separator and treatment facilities.

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Table 4-13 FIELD PARAMETERS, ALKALINITY, COMMON ANIONS, LEAD, TOTAL DISSOLVED SOLIDS, TOTAL PETROLEUM HYDROCARDONS, AND PURGEABLE AROMATICS ESKIMO CREEK INTERMEDIATE AQUIFER

| | | | Detection | KS-MW-41 | KS-EBMW | KS-MM-97 | KS-Mi-42 | KS-MW-43 | K S-T BMM |
|------------------------|-----------|-------------|-----------|------------|------------|------------|----------|-------------|-----------|
| Parameter | Method | Units | Limits | 6/17/88 | 6/17/88 | DUP MM-41 | 6/16/88 | 6/16/88 | 6/16/88 |
| Temperature | FIELD | U | N/A | 4 | na | ŋa | 4.5 | S | na |
| рн | FIELD | ı | N/A | 7.25 | na | Da | 8.05 | 10.1 | na |
| Specific Conductivity | FIELD | SOHMU | N/A | 345 | na | na | 215 | 210 | na |
| Appearance | FIELD | ı | N/A | TURBID | na | na | TURBID | TURBID | na |
| Odor (fuel) | FIELD | ı | N/A | NONE | na | Ъā | NONE | NONE | na |
| Lead | 3005/7421 | ug/1 | 5 | 7 | <5 | 80 | Q | \$ 5 | na |
| Ortho Phosphate | E365.2 | mg/1 | 0.5 | 0.07 | <0.01 | na | 0.51 | 0.04 | na |
| Alkalinity | A403 | mg/l | 20 | 223 | 1.8 | 240 | 157 | 241 | na |
| Chloride | 9252 | mg/l | Ч | 8.3 | 1 > | 8.3 | 8.7 | 11.1 | na |
| Sulfate | 9038 | 1/5m | 1 | 22.3 | 1.2 | 22.3 | 13.5 | 79.7 | na |
| Fluoride | E340.2 | mg/l | 0.05 | 0.11 | 0.05 | 0.10 | 0.09 | 60.0 | na |
| Total Diss. Solids | E160.1 | ng/l | г | 299 | 41 | 282 | 229 | 378 | na |
| Nitrate-Nitrite | 9200 | 1/5m | 0.05 | 0.25 | 0.16 | na | 0.16 | 0.47 | na |
| HAL | E418.1 | ng/l | 1 | 1 > | 4 | пa | пa | na | na |
| 700 | | ng/1 | 1 | na | na | na | 1.7 | 7.2 | na |
| Benzene | SW8020 | ug/1 | г | 4 | 41 | 4 | 12 | 41 | ₽ |
| Toluene | SW8020 | 1/gu | 1 | ₽ | 41 | ¢1 | 41 | ¢ | ¢1 |
| Ethyl Benzene | SW8020 | 1/6n | 1 | <1 | ¢1 | 1 > | 41 | 1 2 | 4 |
| Xy lene | SW8020 | 1/bn | 7 | 41 | 41 | 41 | 41 | ¢1 | 2 |
| Chlorobenzene | SW8020 | 1/gu | 1 | ¢1 | ₽ | 1 2 | 41 | <1> | 4 |
| 1,4-Dichlorobenzene | SW8020 | 1/bn | 1 | 4 | ₽ | 4 | 4 | ¢1 | ≎ |
| 1,3-Dichlorobenzene | SW8020 | t/bn | 1 | ć1 | ¢1 | \$ | ¢1 | 4 | 4 |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | I | 4 | 4 | 1 2 | ₽ | 1 > | 4 |
| | | | | | | | | | |

na = not analyzed/sampled MW = Monitoring Well EB = Equipment Blank TB = Trip Blank AC = Amblent Conditions Blank N/A = Not Applicable

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FIELD PARAMETERS, ALKALINITY, COMMON ANIONS, LEAD, TOTAL DISSOLVED SOLIDS, TOTAL PETROLEUM HYDROCARDONS, AND PURGEABLE AROMATICS ESKIMO CREEK DEEP AOUIFER Table 4-14

| | | | Detection | KS-MW-KS4 | KS-MW-KS5 | KS-MM-KS6 | KS-ACMW |
|----------------------------|-----------|-------------|-----------|-----------|-------------|------------|----------|
| Parameter | Method | Units | Limits | 6/8/88 | 6/8/88 | DUP MW-KS5 | 6/8/88 |
| Temperature | FIELD | U | N/A | | | | Da |
| pH | FIELD | • | N/A | | | | na |
| Specific Conductivity | FIELD | SOHMU | N/A | | | | na |
| Appearance | FIELD | ı | N/A | | | | đ |
| Odor (fuel) | FIELD | I | N/A | | | | na |
| Lead | 3005/7421 | ug/1 | ŝ | 45 | \$ 5 | ŝ | na |
| Ortho Phosphate | E365.2 | mg/1 | 0.5 | 0.67 | 0.71 | 0.70 | na |
| Alkalinity | A403 | mg/l | 20 | 130 | 147 | 147 | Da |
| Chloride | 9252 | <u>mg/1</u> | 1 | 13.0 | 10.1 | 10.6 | na |
| Sulfate | 9038 | mg/1 | 1 | 3.9 | 2.0 | 1.8 | na |
| Fluoride | E340.2 | ng/1 | 0.05 | 0.24 | 0.23 | 0.23 | na |
| Total Diss. Solids | E160.1 | mg/1 | 1 | 177 | 210 | 194 | na |
| Nitrate-Nitrite | 9200 | mg/1 | 0.05 | 0.23 | 0.24 | 0.18 | na |
| TPH | E418.1 | mg/1 | 1 | 27.1 | 6.4 | 6.0 | na |
| TOC | | mg/1 | 1 | па | na | na | na |
| Benzene | SW8020 | 1/gu | 1 | 12 | 4 | 41 | 12 |
| Toluene | SW8020 | 1/bn | 1 | 4 | ₽ | 4 | ₽ |
| Ethyl Benzene | SW8020 | 1/gu | 1 | 4 | 41 | ₽ | ₽ |
| Xy lene | SW8020 | ng/1 | 1 | 4 | 4 | 4 | ₽ |
| Chlorobenzene | SW8020 | 1/bn | 1 | 4 | 4 | 4 | ث |
| 1,4-Dichlorobenzene | SW8020 | ug/1 | 1 | 4 | 4 | 4 | 4 |
| 1,3-D1chlorobenzene | SW8020 | ng/1 | 1 | 4 | 4 | 4 | ₽ |
| 1,2-Dichlorobenzene | SW8020 | ug/1 | 1 | 4 | 4 | 4 | 4 |
| na = not analyzed/sampled | | | | | | | |
| MW = Monitoring Well | | | | | | | |
| EB = Equipment Blank | | | | | | | |
| TB = Trip Blank | | | | | | | |
| AC = Ambient Conditions B1 | lank | | | | | | |
| N/A = Not Applicable | | | | | | | |

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| | PURGEABLE A SW802 ESKIMO CREEK | ROMATICS AND LE 0 AND SW7421 1 DEVELOPMENT WA | AD | |
|---------------------|--------------------------------------|---|-----------------|---------|
| Parameter | Units | Detection Limit | KS-DW 6/8/88 | 6/17/88 |
| Benzene | ug/l | 1 | <1 | <1 |
| Toluene | ug/l | 1 | <1 | 2 |
| Ethyl Benzene | ug/l | 1 | <1 | 5 |
| Xylene | ug/l | 1 | <1 | 20 |
| Chlorobenzene | ug/l | 1 | <1 | <1 |
| 1,4-Dichlorobenzene | ug/l | 1 | <1 | <1 |
| 1,3-Dichlorobenzene | ug/l | 1 | <1 | <1 |
| 1,2-Dichlorobenzene | ug/l | 1 | <1 | <1 |
| Lead | ug/l | 5 | <5 | <5 |

Table 4-15

4.1.5.1.2 Naknek River Seep

A water sample was collected at the Naknek River seep from a small pool that was excavated in the river bank to intercept the seepage face. Analytical results are presented in the following section on surface water.

4.1.5.1.3 North Barrel Bluff

Groundwaters were not collected at the North Barrel Bluff. Results for surface water samples are provided in the following section.

4.1.5.1.4 South Barrel Bluff

A residential well located along the South Barrel Bluff (see Well B Figure 4-7) was sampled for the following water quality parameters:

- o Halogenated volatile organics
- o Aromatic volatile organics
- o Organochlorine pesticides and PCBs
- o Semivolatile organics
- o Metals

Results are reported in Tables 4-15A to 4-15D. There were no detected constituents in these analyses. This well was sampled after information gained from the geophysical survey indicated the well may have been drilled through a landfill area (i.e., drums and other metallic debris) into the intermediate aquifer zone.

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| Table 4-15A |
|-------------------------------------|
| PURGEABLE HALOCARBONS AND AROMATICS |
| E601 AND SW8020 |
| SOUTH BARREL BLUFF GROUND WATER |

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| | | | | KS-MW-SAD | |
|---------------------------|-------|-----------|----------|-----------|---------|
| | | Detection | KS-MW-SA | Duplicate | ACMW-SA |
| Parameters | Units | Limit | 6/8/88 | 6/8/88 | 6/8/88 |
| Chloromethane | ug/l | 1 | <1 | <1 | <1 |
| Bromomethane Vinyl | ug/l | 1 | <1 | <1 | <1 |
| Chloride Methylene | ug/l | 1 | <1 | <1 | <1 |
| Chloride | ug/l | 1 | <1 | <1 | <1 |
| 1,1-Dichloroethene | ug/l | 5 | <5 | <5 | <5 |
| 1,1-Dichloroethane | ug/l | 1 | <1 | <1 | <1 |
| trans-1,2-Dichloroethene | ug/l | 1 | <1 | <1 | <1 |
| Chloroform | ug/l | 1 | <1 | <1 | <1 |
| 1,2-Dichloroethane | ug/l | 1 | <1 | <1 | <5 |
| 1,1,1-Trichloroethane | ug/l | 1 | <1 | <1 | <1 |
| Carbon Tetrachloride | ug/l | 1 | <1 | <1 | <1 |
| Bromodichloromethane | ug/l | 1 | <1 | <1 | <1 |
| 1,2-Dichloropropane | ug/l | 1 | <1 | <1 | <1 |
| cis-1,3-Dichlorpropene | ug/l | 1 | <1 | <1 | <1 |
| Trichloroethene | ug/l | 1 | <1 | <1 | <1 |
| Dibromochloromethane | ug/l | 1 | <1 | <1 | <1 |
| 1,1,2-Trichloroethane | ug/l | 1 | <1 | <1 | <1 |
| trans-1,3-Dichloropropene | ug/l | 1 | <1 | <1 | <1 |
| Bromoform | ug/l | 1 | <1 | <1 | <1 |
| 1,1,2,2-Tetrachloroethane | ug/l | 1 | <1 | <1 | <1 |
| Tetrachloroethene | ug/l | 1 | <1 | <1 | <1 |
| Chlorobenzene | ug/l | 1 | <1 | <1 | <1 |
| 1,3-Dichlorobenzene | ug/l | 1 | <1 | <1 | <1 |
| 1,2-Dichlorobenzene | ug/l | 1 | <1 | <1 | <1 |
| 1,4-Dichlorobenzene | ug/l | 1 | <1 | <1 | <1 |
| Benzene | ug/l | 1 | <1 | <1 | na |
| Toluene | ug/l | 1 | <1 | <1 | na |
| Ethyl Benzene | ug/l | 1 | <1 | <1 | na |
| Chlorobenzene | ug/l | 1 | <1 | <1 | na |
| 1,4-Dichlorobenzene | ug/l | 1 | <1 | <1 | na |
| 1,3-Dichlorobenzene | ug/l | 1 | <1 | <1 | na |
| 1,2-Dichlorobenzene | ug/l | 1 | <1 | <1 | na |

AC-Ambient Conditions Blank

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| | | | | KS-MW-SAD |
|--------------------|-------|-----------|----------|-----------|
| | | Detection | KS-MW-SA | Duplicate |
| Parameters | Units | Limit | 6/8/88 | 6/8/88 |
| a-BHC | ug/1 | 0.05 | <0.05 | <0.05 |
| b-BHC | ug/1 | 0.05 | <0.05 | <0.05 |
| g-BHC | ug/1 | 0.05 | <0.05 | <0.05 |
| d-BHC | ug/1 | 0.05 | <0.05 | <0.05 |
| Heptachlor | ug/1 | 0.02 | <0.02 | <0.02 |
| Aldrin | ug/1 | 0.05 | <0.05 | <0.05 |
| Heptachlor Epoxide | ug/l | 0.05 | <0.05 | <0.05 |
| Endosulfan I | ug/l | 0.05 | <0.05 | <0.05 |
| Dieldrin | ug/l | 0.02 | <0.02 | <0.02 |
| 4,4-DDE | ug/l | 0.1 | <0.1 | <0.1 |
| Endrin | ug/l | 0.06 | <0.06 | <0.06 |
| Endosulfan II | ug/l | 0.1 | <0.1 | <0.1 |
| 4,4-DDD | ug/l | 0.1 | <0.1 | <0.1 |
| Endrin Aldehyde | ug/l | 0.1 | <0.1 | <0.1 |
| Endosulfan Sulfate | ug/l | 0.1 | <0.1 | <0.1 |
| 4,4-DDT | ug/l | 0.1 | <0.1 | <0.1 |
| methxychlor | ug/l | 0.1 | <0.1 | <0.1 |
| Chlordane | ug/l | 0.2 | <0.2 | <0.2 |
| Toxaphene | ug/l | 1 | <1 | <1 |
| PCB-1221 | ug/l | 2 | <2 | <2 |
| PCB-1232 | ug/l | 2 | <2 | <2 |
| PCB-1242 | ug/l | 0.8 | <0.8 | <0.8 |
| PCB-1016 | ug/l | 0.8 | <0.8 | <0.8 |
| PCB-1248 | ug/l | 0.5 | <0.5 | <0.5 |
| PCB-1254 | ug/l | 1 | <1 | <1 |
| PCB-1260 | ug/l | 1 | <1 | <1 |

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TABLE 4-15B ORGANOCHLORINE PESTICIDES AND PCBs E608 SOUTH BARREL BLUFF GROUND WATER

Table 4-15C METALS SCREEN E200.7 SOUTH BARREL BLUFF GROUND WATER

| | | | KS-MW-SAD |
|------------|-----------|----------|-----------|
| | | KS-MW-SA | Duplicate |
| Parameters | Detection | 6/8/88 | 6/8/88 |
| Aluminum | 200 | <200 | <200 |
| Antimony | 300 | <300 | <300 |
| Arsenic | 300 | <300 | <300 |
| Barium | 100 | <100 | <100 |
| Beryllium | 5 | <5 | <5 |
| Cadmium | 10 | <10 | <10 |
| Calcium | 1000 | 19100 | 18500 |
| Chromium | 30 | <30 | <30 |
| Cobalt | 40 | <40 | <40 |
| Copper | 30 | <30 | <30 |
| Iron | 100 | 150 | 120 |
| Lead | 200 | <200 | <200 |
| Magnesium | 1000 | 7300 | 7100 |
| Manganese | 15 | 173 | 167 |
| Molybdenum | 40 | <40 | <40 |
| Nickel | 40 | <40 | <40 |
| Potassium | 1000 | 3800 | 3300 |
| Selenium | 400 | <400 | <400 |
| Silver | 30 | <30 | <30 |
| Sodium | 1000 | 8100 | 7900 |
| Thallium | 500 | <500 | <500 |
| Vanadium | 40 | <40 | <40 |
| Zinc | 20 | <20 | <20 |
| Boron | 30 | <30 | <30 |
| Silica | 43 | 35000 | 35000 |

Note: All values in ug/1

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Table 4-15D EXTRACTABLE PRIORITY POLLUTANTS EPA625 SOUTH BARREL BLUFF GROUND WATER STANDARDS, CRITERIA AND ACTION LEVELS

| | | | KS-MW-SAD |
|------------------------------|-------|----------|-----------|
| | | KS-MW-SA | Duplicate |
| Parameters | Units | 6/8/88 | 6/8/88 |
| | | | |
| n-nitroso-dimethylamine | ug/l | 100 | 100 |
| phenol | ug/l | 100 | 100 |
| aniline | ug/l | 100 | 100 |
| bis(2-chloroethyl) ether | ug/l | 100 | 100 |
| 2-chlorophenol | ug/1 | 100 | 100 |
| 1,3-dichlorobenzene | ug/l | 100 | 100 |
| 1,4-dichlorobenzene | ug/1 | 100 | 100 |
| benzyl alcohol | ug/l | 100 | 100 |
| 1,2-dichlorobenzene | ug/1 | 100 | 100 |
| 2-methylphenol | ug/l | 100 | 100 |
| bis(2-chloroisopropyl) ether | ug/l | 100 | 10U |
| 4-methylphenol | ug/l | 100 | 100 |
| n-nitroso-di-n-propylamine | ug/1 | 100 | 100 |
| hexachloroethane | ug/l | 100 | 100 |
| nitrobenzene | ug/l | 100 | 100 |
| isophrone | ug/l | 100 | 100 |
| 2-nitrophenol | ug/l | 100 | 100 |
| 2,4-diethylphenol | ug/l | 100 | 100 |
| benzoic acid | ug/l | 50U | 50U |
| bis(2-chloroethoxy) methane | ug/l | 100 | 100 |
| 2.4-dichlorophenol | ug/l | 100 | 100 |
| 1,2,4-trichlorobenzene | ug/l | 100 | 100 |
| naphtalene | ug/l | 100 | 100 |
| 4-chloroaniline | ug/l | 100 | 100 |
| hexachlorobutadiere | ug/l | 100 | 100 |
| 4-chloro-3-methylphenol | ug/l | 100 | 100 |
| 2-methylnaphthalene | ug/l | 100 | 100 |
| hexachlorocyclopentadiene | ug/l | 100 | 100 |
| 2.4.6-trichlorophenol | ug/l | 100 | 100 |
| 2.4.5-trichlorophenol | ug/l | 50U | 500 |
| 2-chloronaphthalene | ug/l | 100 | 100 |
| 2-nitroaniline | ug/l | 50U | 500 |
| dimethy1 phthalate | ug/l | 100 | 100 |
| acenaphthylene | ug/1 | 100 | 100 |
| 3-nitroaniline | ug/l | 50U | 500 |
| acenaphthene | ug/1 | 100 | 100 |
| 2.4-dinitrophenol | ug/l | 50U | 500 |
| 4-nitrophenol | ug/1 | 50U | 50U |
| dibenzofuran | ug/l | 100 | 100 |
| 2.4-dinitrotoluene | ug/l | 100 | 100 |
| 2.6-dinitrotoluene | ug/l | 100 | 100 |
| diethyl phthalate | ug/l | 100 | 100 |

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U=Indicates constituent tested for, but not detected.

J=Indicates an estimated trace value.

| | | | KS-MW-SAD |
|-----------------------------|-------|----------|-----------|
| | | KS-MW-SA | Duplicate |
| Parameters | Units | 6/8/88 | 6/8/88 |
| 4-chlorophenyl phenyl ether | ug/1 | 1017 | 101 |
| fluorene | ug/1 | 100 | 100 |
| A-nitroaniline | ug/1 | 501 | 500 |
| 4 6-dinitro-2-methylphenol | ug/1 | 500 | 500 |
| anithogodinhenylamine | ug/1 | 100 | 100 |
| A-bromonbanyl phenyl ather | ug/1 | 100 | 100 |
| 4-Dromognenyi phenyi ether | ug/1 | 100 | 100 |
| nexaciior openzene | ug/1 | 500 | 500 |
| pheneachiolophenol | ug/1 | 300 | 100 |
| phenanchiene | ug/1 | 100 | 100 |
| antiracene | ug/1 | 100 | 100 |
| di-n-bucyi phachalace | ug/1 | 100 | 100 |
| riuoranchene | ug/1 | 100 | 100 |
| Denziaine | ug/1 | 500 | 500 |
| pyrene | ug/1 | 100 | 100 |
| butyl benzyl phinalate | ug/1 | 100 | 100 |
| 3,3-dichlorobenzidine | ug/1 | 200 | 200 |
| benzo(a)anthracene | ug/1 | 100 | 100 |
| bis(2-ethylhexyl)phthalate | ug/l | 100 | 5J |
| chrysene | ug/l | 100 | 100 |
| di-n-octyl phthalate | ug/l | 100 | 100 |
| benzo(b)fluoranthene | ug/l | 100 | 100 |
| benzo(k)fluoranthene | ug/l | 100 | 100 |
| benzo(a)pyrene | ug/l | 100 | 100 |
| indeno(1,2,3-cd)pyrene | ug/l | 100 | 100 |
| dibenz(a,h)anthracene | ug/l | 100 | 100 |
| benzo(g,h,i)perylene | ug/l | 100 | 100 |

Table 4-15D (Continued)

U=Indicates constituent tested for, but not detected. J=Indicates an estimated trace value.

4.1.5.2 Surface Water Results

Surface water samples were collected in the vicinity of the Eskimo Creek seep and the North and South Barrel Bluffs. A sample was also collected at the Naknek River seep. Sample locations were selected at points located upstream and downstream of each disturbance to provide an indication on whether these areas are contributing hazardous constituents to the surface waters. Surface waters and sediment sampling locations are shown in Figure 4-15. Surface water samples were analyzed for aromatic volatiles, purgeable halocarbons, extractable priority pollutants, organochlorine pesticides and PCBs, TDS, alkalinity, common anions, and field parameters.

The results of these analyses are presented in Tables 4-16A through 4-16E.

A detail summary for each investigation area is provided in the following subsections.

4.1.5.2.1 Eskimo Creek Seep

Laboratory analysis of water samples from Eskimo Creek collected at locations upstream and downstream of the Eskimo Creek seep reported no concentrations above method detection limit for aromatic, purgeable, and halocarbon compounds. Concentrations of inorganic constituents did not vary significantly between the downstream and upstream locations.

4.1.5.2.2 Naknek River Seep

Xylene and trans 1,2-dichloroethene were detected at concentrations of 14 ug/l and 85 mg/l in the Naknek River seep sample. Inorganic constituent concentrations were also significantly higher than were observed at other water sample locations.

While collecting the surface water sample, an oily sheen and discoloration were observed.

4.1.5.2.3 North Barrel Bluff

Organic constituent concentrations for surface water samples collected from the King Salmon Creek flood plain in the vicinity of the North Barrel Bluff were reported at less than method detection limits.

Inorganic constituent concentrations do not indicate specific trends. Potassium was detected at the North Barrel, but was not observed at the South Barrel Bluff. Overall, inorganic constituent concentrations were generally in the same range when comparing concentrations between the North and South Barrel Bluff.





| | AROMAT | |
|------------|-----------|--------------|
| | PURGEABLE | SEEP |
| | AND | RIVER |
| | SOLIDS | NAKONEK 1 |
| | LVED | AND |
| A | DISSO | BLUFF |
| Table 4-16 | , TOTAL | BARREL |
| | AN IONS | SOUTH 1 |
| | NOM | AND |
| | CO ' | VORTH |
| | KALINITY | CREEK, 1 |
| | S, AL | KIMO |
| | PARAMETER | ន |

FIELD

ICS

KS-SW02-03 11/01/87 CLEAR 2.0 <0.5 15.3 3.81 1.85 <0.5 42 NO 0.49 78 ₫ ₫ ₽ 7777 South Barrel Bluff KS-SW02-02 SL COLOR 10/28/87 1.2 6.42 <0.5 3.98 1.63 <0.5 34.2 0.12 82 NO 17 ₽ ₽ 777777 80 SL COLOR NO KS-SW02-01 10/28/87 1.2 6.64 <0.5 49.7 3.82 1.43 <0.5 111 <0.1 4444 ₽₽ 4 KS-SW01-02 10/28/87 CLEAR <0.5 34.2 0.5 6.63 5.35 1.63 <0.5 171 **60.1** 80 Q ₽₽ ₽ 4444 Eskimo Creek KS-SW01-01 10/28/87 CLEAR 0.5 6.96 <0.5 28.6 5.38 1.67 <0.5 0.17 80 4 4 777 8 55 ₽ ₽ Detection Limit 0.5 20 1 1 1 0.05 0.05 N/A N/A N/A N/A N/A **WHOS** Units լ/նա **m**g/1 mg/l mg/l mg/1 mg/1 ng/1 ug/1 1/bn 1/bn 1/bn ug/1 1/6n 1/bn • • ug/1 ပ ı Method FIELD TELD FIELD SW8020 SW8020 SW8020 SW8020 SH8020 SW8020 SW8020 FIELD FIELD E365.2 E340.2 SW8020 E160.1 A403 9252 9038 9200 N/A = Not Applicable Specific Conductivity = Not Available 1,3-D1chlorobenzene 1,4-Dichlorobenzene 1,2-Dichlorobenzene Total Diss. Solids Parameter Ortho Phosphate Nitrate-Nitrite Ethyl Benzene Chlorobenzene Temperature Appearance Alkalinity Fluoride Chloride Toluene Sulfate Benzene Xy lene Odor na Hq

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KSPB = Ambient Conditions Blank

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Table 4-16A (Continued)

10/28/87 KSPB na na na Пa na na na na na **** River Seep KS-SW04-01 10/31/87 Naknek 3.5 <0.5 3.5 6.14 200 130 **4.**07 2**4**3 <0.1 VILY VES 2 2 2 2 7 0 0 7 DUP SW03-02 KS-SM03-04 10/31/87 <0.5 47.2 5.08 4.71 <0.5 526 1.24 d d d d d d d d na na na na KS-SW03-03 SL COLOR 10/31/87 2.5 6.37 <0**.**5 34.4 3.82 2.45 <0.5 53 0.1 88 BU 4 4 4 4 4 4 ₫ ₫ North Barrel Bluff KS-SW03-02 10/31/87 46.1 4.91 4.69 <0.5 170 2.0 6.34 <0.5 1.46 85 na na **♀ ♀ ♀ ♀** 44 ₫ ₫ 52 SL COLOR NO KS-SW03-01 10/31/87 3.5 6.30 <0.5 4.27 4.18 <0.5 112 0.1 34 Detection Limit N/A N/A N/A N/A N/A 0.05 1 0.05 0.5 20 1 H ы SOHMU mg/l Units mg/l 1/ba 1/60 mg/1 mg/l mg/1 1/bn 1/6n 1/bn ug/1 ug/1 1/bn 1/bn ug/1 1 1 υī E340.2 E365.2 E160.1 9038 A403 9252 9200 SW8020 SW8020 SW8020 SW6020 SW8020 SW8020 FIELD FIELD FIELD FIELD SW8020 SW8020 Method FIELD KSFB = Ambient Conditions Blank = Not Applicable Specific Conductivity = Not Available 1,2-Dichlorobenzene 1,4-Dichlorobenzene 1,3-Dichlorobenzene Total Diss. Solids Nitrate-Nitrite Ortho Phosphate Parameter Ethyl Benzene Chlorobenzene Temperature Alkalinity Appearance Fluoride **Chloride** Sulfate Benzene Toluene Xylene Odor N/A na Ηd

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Table 4-16B PURGEABLE HALOCARBONS E601

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ESKIMO CREEK, NORTH AND SOUTH BARREL BLUFF, AND NAKNEK RIVER SEEP

| | | | Eskimo | o Creek | Sou | th Barrel Bluff | |
|-------------------------------|---------------|--------------------|----------------------------|----------------------------|----------------------------|----------------------------|----------------------------|
| Parameters | Units | Detection Limit | KS-SM01 -01 10/28/87 | KS-SMOI -02 10/28/87 | KS-SM02 -01 10/28/87 | KS-SW02 -02 10/28/87 | KS-SW02 -03 11/01/87 |
| Chloro s ethane | 1/pu | Ч | 4 | 4 | 4 | 4 | 12 |
| Bromomethane Vinyl | uq/1 | | ₽ | 4 | 1 | <u>2</u> | ₽ |
| Dichloridifluormethane | ug/1 | - | \$ | \$ | \$ | \$ | \$ |
| Vinyl Chloride | ug/1 | | | | | | |
| Chloroethane | ug/1 | ŝ | 4 | 4 | 4 | 4 | 4 |
| Methylene Chloride | ug/l | Ч | 2 | 4 | ₽ | 4 | ₽ |
| Trichlorofluoromethane | ug/1 | - | 5 | 4 | 4 | 4 | ₽ |
| 1,1-Dichloroethene | ug/1 | -1 | 4 | 4 | ₽ | 2 | 4 |
| 1,1-Dichloroethane | ug/1 | Ч | 4 | 4 | 4 | ₽ | ≎ |
| trans-1,2-Dichloroechene | 1/gu | н | 4 | 4 | ₽ | ₽ | 4 |
| Chloroform | ug/1 | I | 4 | 4 | 4 | \$ | 4 |
| 1,2-Dichloroethane | ng/1 | | 4 | 4 | \$ | ≎ | 4 |
| 1,1,1-Trichloroethane | 1/bn | - | 4 | 4 | 4 | \$ | ₽ |
| Carbon Tetrachloride | ng/1 | - | ₽ | 4 | 4 | 4 | ₽: |
| Bromodichloromethane | ng/1 | T | 4 | 4 | ₽: | ⊄: | ₽: |
| 1,2-Dichloropropane | ng/1 | -1 | 4 | 4 | ₽. | ₽, | ₽: |
| cis-1,3-Dichlorpropene | 1/bn | -1 | 4 | 4 | ₽: | ₽. | ₽: |
| Trichloroethene | ng/1 | 1 | 1 | 4 | 4 | 4 | ₽ : |
| Dibromochloromethane | 1/gu | -1 | ۵ | ŝ | ۵ | \$ | ŝ |
| 1,1,2-Trichloroethane | ng/1 | 1 | | | | | |
| trans-1,3-Dichloropropene | ng/1 | -1 | | | | ! | ! |
| Bromoform | 1/5n | -1 | 4 | ≎ | 1 | 1 | ۵. |
| 1,1,2,2-Tetrachloroethan | ug/1 | Ч | \$ | \$ | \$ | 4 2 | \$ |
| Tetrachloroethene | u <u>g</u> /1 | 1 | | | | | |
| Chlorobenzene | 1/bn | | 4 | 4 | 4 | ≎ | ₽ |
| 1,3-Dichlorobenzene | ng/1 | -1 | ¢1 | ₽. | ∵ : | ⊄: | ₽: |
| 1,2-Dichlorobenzene | ng/1 | | ₽: | ₽: | ₽; | 4: | 2: |
| l,4-Dichlorobenzene | ng/l | 1 | ₽ | 5 | 5 | 7 | 7 |
| | | | | | | | |

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| | | | Nort | ch Barrel Bluf | Ţ | | Naknek River Seep | |
|----------------------------------|--------------|--------------------|----------------------------|----------------------------|----------------------------|--|----------------------------|------------------|
| Parameters | <u>Un1ts</u> | Detection Limit | KS-SH03 -01 10/31/87 | KS-SW03 -02 10/31/87 | KS-SH03 -03 10/31/87 | KS-SW03-04 DUP SW03-02 <u>10/31/87</u> | KS-SM04 -01 10/31/87 | KSFB 10/28/87 |
| Chloromethane | 1/01 | -1 | 41 | 1 > | 41 | 12 | 12 | 1 |
| Bromomethane Vinvl | 1/on | - | '₽ | '≎' | 12 | . ≏ | ;≏ | . . |
| Dichloridifluormethane | 1/0n | . –4 | ŝ | Ş | ŝ | ŝ | ŝ | ; ¢ |
| Vinyl Chloride | 1/bn | | | | | | | |
| Chloroethane | 1/pu | 2 | ₽ | <u>1</u> | <u>1</u> | : | 4 | 4 |
| Methylene Chloride | 1/bn | -4 | ₽ | 4 | <u>1</u> | ≎ | \$ | 12 |
| Trichlorofluoromethane | 1/bn | - | 12 | 4 | î | \$ | 4 | 4 |
| 1,1-Dichloroethene | ug/1 | - | \$ | 4 | ¢1 | \$ | ≎ | ₽ |
| 1,1-Dichloroethane | 1/bn | 7 | ţ, | <u>1</u> | î | ¢1 | <u>1</u> | 12 |
| trans-1,2-Dichloroethene | ug/1 | - | 4 | 4 | 1 2 | ¢ | 85 | \$1 |
| Chloroform | t/6n | -1 | 2 | ¢ | <u>1</u> | ≎ | 1 | 4 |
| 1,2-Dichloroethane | 1/6n | - | 4 | 4 | 4 | ≎ | ≎ | 4 |
| 1,1,1-Trichloroethane | 1/bn | - | 4 | <u>^</u> 1 | ₽ | ≎ | 1 | 4 |
| Carbon Tetrachloride | 1/6n | | 4 | ₽ | 4 | ₽ | 4 | <u>1</u> |
| Bromodichloromethane | ng/l | Ч | ≎ | 4 | 4 | ≎ | 4 | 4 |
| 1,2-Dichloropropane | ug/l | 4 | 4 | ≎ | 4 | ₽ | ≎ | 1 |
| cis-1,3-Dichlorpropene | 1/5n | -1 | ₽ | 1 | û | ≎ | 1 | 4 |
| Trichloroethene | ng/1 | 1 | ₽ | ≎ | ¢ | î | QN | : |
| Dibromochloromethane | ug/l | - | ŝ | ŝ | ~ 3 | ŝ | \$ | Ω |
| <pre>l,l,2-Trichloroethane</pre> | ug/1 | - | | | | | | |
| trans-1,3-Dichloropropene | t/bn | -1 | | | | | | |
| Bromotorm | ng/1 | -1 | 4 | 4 | û | ≎ | ≎ | î |
| 1,1,2,2-Tetrachloroethan | ng/1 | -1 | \$ | \$ | \$ | \$ | \$ | \$ |
| Tetrachloroethene | 1/gu | - | | | | | | |
| Chlorobenzene | ug/l | ٦ | 7 | <u>1</u> | 1 | ≎ | 2 | 4 |
| 1,3-Dichlorobenzene | ng/1 | - | 4 | 1 | ₽ | ₽ | 1 | î |
| 1,2-Dichlorobenzene | ug/l | - | ≎ | ₽ | <u>1</u> | ≎ | 1 | 4 |
| 1,4-Dichlorobenzene | ug/1 | - | 4 | 4 | 4 | ₽ | ¢1 | ¢ |
| VCPR-Imblent Conditions Blank | | | | | | | | |
| ND-None Detected | | | | | | | | |

Table 4-16B (Continued)

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Table 4-16C Extractable Priority Pollutants E625

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E625 NORTH AND SOUTH BARREL BLUFF SURFACE WATER

| | | | S | outh Barrel Bluf | f | | North Barr | el Bluff | |
|----|-----------------------------------|-------|------------|------------------|------------|------------|------------|------------|-------------|
| | | | KS-SW02-01 | KS-SW02-02 | KS-SW02-03 | KS-SW03-01 | KS-SW03-02 | KS-SW03-03 | DUP SM03-02 |
| | Parameters | Units | 10/28/87 | 10/28/87 | 10/28/87 | 10/31/87 | 10/31/87 | 10/31/87 | 10/31/87 |
| | n-nitroso-dimethylamine | ug/1 | TOU | 101 | TOU | 10U | not | IOU | 10U |
| | phenol | 1/gu | TOU | TOU | TOU | TOU | lou | TOU | TOU |
| | aniline | 1/bn | IOU | 100 | 100 | TOU | IOU | 100 | IOU |
| | bis(2-chloroethy1) ether | t/bn | 100 | 100 | 10L | 100 | lou | IOU | TOU |
| | 2-chlorophenol | 1/Jn | IOU | 100 | TOU | IOU | 100 | 100 | IOU |
| | 1,3-dichlorobenzene | 1/bn | IOU | IOU | IOU | 100 | IOU | IOU | 100 |
| | 1,4-dichlorobenzene | 1/gu | 100 | IOU | TOU | 100 | TOU | TOU | TOU |
| | benzyl alcohol | 1/6n | IOU | IOU | TOU | TOU | IOU | TOU | 100 |
| | 1,2-dichlorobenzene | 1/bn | IOU | 10U | 101 | IOU | IOU | 100 | 101 |
| | 2-methylphenol | ng/1 | 100 | IOU | 100 | 100 | IOU | 100 | 100 |
| I١ | bis(2-chloroisopropyl) ether | 1/5n | lou | TOU | 100 | 100 | 100 | 10U | NOT |
| 7 | 4-methylphenol | 1/gu | IOU | IOU | TOU | IOU | IOU | 100 | 101 |
| 83 | n-nitroso-di-n-propylamine | 1/bn | JOU | IOU | 101 | IOU | IOU | 101 | 10U |
| | hexachloroethane | 1/6n | 100 | 100 | UOT | IOU | 100 | 100 | 101 |
| | nitrobenzene | 1/gu | IOU | TOU | 101 | 101 | TOU | TOU | IOU |
| | isophrone | 1/gu | IOU | IOU | 100 | 100 | 101 | 100 | 100 |
| | 2-nttrophenol | 1/bn | JOU | 100 | 100 | 101 | 10U | 101 | not |
| | 2,4-diethylphenol | 1/bn | lou | lou | 101 | IOU | IOU | TOU | 10U |
| | benzoic acid | 1/gu | 50U | 500 | 500 | 500 | 500 | 50U | 50U |
| | bis(2-chloroethoxy) methane | ug/1 | JOU | 100 | 100 | 10U | 100 | 100 | 100 |
| | 2,4-dichlorophenol | 1/5n | 100 | 100 | 100 | IOU | 100 | 100 | 101 |
| | <pre>l,2,4-trichlorobenzene</pre> | ng/1 | 100 | 100 | 100 | 100 | 10U | 101 | 101 |
| | naphtalene | 1/bn | 100 | 100 | TOU | IOL | 101 | 100 | 100 |
| | 4-chloroaniline | 1/bn | 100 | 100 | IOU | IOU | IOU | IOI | 100 |
| | hexachlorobutadiene | ug/1 | not | 100 | 100 | 101 | 100 | TOU | 100 |
| | 4-chloro-3-methylphenol | ng/1 | lou | 100 | 100 | 100 | 100 | 100 | 100 |
| | 2-methylnaphthalene | ug/1 | not | 10U | 100 | 100 | 100 | 100 | TOU |
| | hexachlorocyclopentadiene | ug/1 | 100 | 100 | 100 | 10U | 100 | IOU | 100 |
| | 2,4,6-trichlorophenol | 1/2n | IOU | 100 | 100 | 10U | 100 | 100 | 101 |
| | | | | | | | | | |

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| | | | Table ((Contir | 4-16C nued) | | | | |
|-----------------------------|---------------|------------------------|------------------------|----------------|------------|------------------------|-------------|-------------|
| | | S | outh Barrel Bluff | | | North B | arrel Bluff | |
| Parameters | Units | KS-SW02-01 10/28/87 | KS-SW02-02 10/28/87 | KS-SW02-03 | KS-SW03-01 | KS-SW03-02 10/31/87 | KS-SM03-03 | DUP SM03-02 |
| | | | | | | | | |
| 2,4,5-trichlorophenol | ug/1 | 500 | 500 | 500 | 50U | 50U | 500 | 500 |
| 2-chloronaphthalene | ug/1 | 100 | 10U | 100 | IOU | 100 | IOL | 100 |
| 2-nitroaniline | 1/gu | 50U | 50U | 50U | 500 | 50U | 500 | 500 |
| dimethyl phthalate | 1/6n | 100 | 100 | 100 | IOU | 100 | 100 | 100 |
| acenaphthylene | 1/bn | IOU | 100 | 100 | IOU | 100 | TOU | 100 |
| 3-nitroaniline | ug/1 | 500 | 50U | 500 | 500 | 500 | 500 | 500 |
| acenaphthene | t/bn | IOU | 100 | 100 | TOU | TOU | IOU | 100 |
| 2,4-dinitrophenol | 1/gu | 500 | 50U | 500 | 50U | 50U | 500 | 500 |
| 4-nitrophenol | 1/gu | 50U | 50U | 500 | 500 | 500 | 500 | 500 |
| dibenzofuran | τ/ 5 π | 100 | 100 | 100 | IOU | TOU | 100 | 100 |
| 2,4-dinitrotoluene | ug/1 | 100 | 100 | 100 | IOU | IOU | 101 | 100 |
| 2,6-dinitrotoluene | 1/6n | 100 | TOU | IOU | TOU | IOU | 100 | 100 |
| diethyl phthalate | 1/6n | IOU | 100 | 100 | 100 | TOU | 100 | 100 |
| 4-chlorophenyl phenyl ether | 1/6n | 100 | TOU | 100 | IOU | TOU | TOU | 100 |
| fluorene | ug/1 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| 4-nitroaniline | ug/1 | 50U | 50U | 500 | 500 | 500 | 500 | 500 |
| 4,6-dinitro-2-methylphenol | 1/6n | 500 | 50U | 50U | 50U | 50U | 500 | 500 |
| n-nithosodiphenylamine | 1/6n | IOU | 100 | 10U | 100 | TOU | TOU | 100 |
| 4-bromopheny1 pheny1 ether | 1/6n | 100 | 100 | 100 | 100 | IOU | 100 | lou |
| hexachlorobenzene | 1/6n | IOU | 100 | 100 | 100 | 101 | 100 | 100 |
| pentachlorophenol | 1/bn | 50U | 500 | 50U | 500 | 50U | 500 | 500 |
| phenanthrene | ug/1 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| anthracene | 1/2u | 100 | 10U | 100 | 100 | IOU | IOU | 100 |
| di-n-butyl phathalate | 1/6n | IOU | 100 | 100 | 100 | 101 | 100 | 100 |
| fluoranthene | 1/gu | 100 | 100 | 100 | IOU | 100 | 100 | 100 |
| benzidine | 1/gu | 500 | 50U | 50U | 500 | 500 | 500 | 500 |
| pyrene | 1/6n | JOU | 100 | 100 | IOU | 100 | 100 | 100 |
| butyl benzyl phthalate | ng/1 | IOU | 100 | IOU | 100 | 100 | IOU | IOU |
| 3,3-dichlorobenzidine | ug/1 | 200 | 20U | 20U | 200 | 200 | 200 | 200 |
| benzo (a) anthracene | ug/1 | 10U | 100 | 100 | 100 | 100 | 10U | 100 |
| bis(2-ethylhexyl)phthalate | ug/1 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| chrysene | ug/1 | 101 | IOU | 100 | 100 | 100 | 100 | 100 |

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| | | | (Cont1 | nued) | | | | |
|--|--------|------------|------------------------|------------------------|------------------------|------------------------|------------------------|---------------------------------|
| | | | South Barrel Blu | ff | | North Bai | rrel Bluff | |
| | lintte | KS-SW02-01 | KS-SW02-02 10/28/87 | KS-SW02-03 10/28/87 | KS-SW03-01 10/31/87 | KS-SW03-02 10/31/87 | KS-SW03-03 10/31/87 | DUP SW03-02 10/ <u>31/87</u> |
| rareters | STITIO | 10 104 104 | | | | | | |
| dismontal whthalate | ua/1 | 100 | TOU | IOU | 101 | TOU | TOU | 100 |
| ut-ii-occy i piiriatace | 1/01 | 100 | TOU | IOU | IOU | TOU | 100 | 100 |
| bell20(b) 1100 all cheric bonno(b) f100; all cheric | - /6m | 100 | IOU | 100 | 101 | TOU | TOU | TOU |
| benzo(x) t tuoj antuene bonzo(z) nistalio | 1/011 | 100 | IOU | 100 | 101 | TOU | IOU | TOU |
| ueutova) pyteme inden∧(1 2 3…rd) nvrene | - 1/DN | not | TOU | IOU | TOU | 100 | 100 | TOU |
| dihenz(a.h) anthracene | 1/bn | IOU | TOU | 101 | IOL | TOU | 100 | 100 |
| benzo (g,h,1) perylene | ug/1 | 100 | 100 | 100 | 100 | 101 | 100 | IOI |
| | | | | | | | | |

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Table 4-16C

(E608) North and South Barrel Bluff Surface Water

ORGANOCHLORINE PESTICIDES AND PCBs

Table 4-16D

KS-SW03-04 DUP-SW03-02 10/31/87 **\$0.1** <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 **ć0.1 ć0.1** <0.1 <0.1 <0.1 <0.5 <0.2 <0.2 <0.1 <0.1 \$0.1 **ć0.1** ₽ Ç Ç 4 7 KS-SW03-02 KS-SW03-03 10/31/87 **6.1** <0.1 \$0**.**1 <0.1 <0.1 ć0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0**.**1 <0.1 <0.1 **ć0.1** <0.5 <0.2 <0.1 <0.1 <0.1 4 \$ \$2 2 ₽ <0.2 North Barrel Bluff 10/31/87 **ć0.1** <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.5 <0.2 <0.2 <0.1 \$0.1 <0.1 <0.1 **ć0.1** <0.1 <0.1 4 Ç Ç 4 4 10-EOMS-SX 10/31/87 <0.1 <0.1 <0.1 <0.1 <0.1 <0.5 <0.2 <0.2 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 ¢0.1 <0.1 <0.1 <0.1 4 ĉ \$ ů 4 KS-SW02-03 10/28/87 ô.1 \$0.1 \$0**.**1 <0.1 <0.1 **ć0.1** <0.1 \$0.1 <0.1 <0.1 **60.1** ć0.1 \$**0.1** <0.1 <0.1 <0.1 \$0.1 Ç <0.5 <0.2 <0.2 0.2 Ļ Ç 4 2 South Barrel Bluff KS-SW02-01 10/28/87 **ĉ.1** <0.1 <0.1 ¢0.1 ć0.1 ć0.1 \$0**.**1 \$0**.**1 <0.1 <0.1 <0.1 <0.2 \$0.1 <0.1 ¢0.1 <0.1 <0.1 \$0.1 <0.1 <0.5 <0.2 7 \Im $^{\circ}$ 4 4 KS-SW0 2-01 10/28/87 <0.1 **ć0.1** <0.1 <0.1 \$0.1 <0.1 <0.1 <0.1 <0.5 <0.1 <0.1 \$0.1 <0.1 \$0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.2 <0.2 4 Ç \$ 4 4 Detection Limit <0.1 \$0.1 <0.1 **ć0.1** <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 <0.1 \$0.1 <0.1 <0.1 <0.1 <0.1 <0.1 ć0.5 <0.2 <0.2 4 ç ₽ \$ ų Units 1/bn ug/1 ug/1 1/6n ng/1 t/bn ug/1 1/bn ug/1 1/bn ī/ɓn 1/bn 1/bn 1/bn 1/bn 1/bn t/bn 1/bn 1/bn 1/bn 1/bn t/bn 1/bn ug/1 ug/1 ug/1 deptachlor Epoxide Endosulfan Sulfate Endrin Aldehyde Parameters Cndosulfan II Endosulfan I nethxychlor Heptachlor Chlordane Toxaphene Dieldrin PCB-1221 PCB-1232 PCB-1242 PCB-1016 PCB-1248 PCB-1254 PCB-1260 4-DDT 4-DDD 4-DDE Aldrin Endrin d-BHC g-BHC a-BHC b-BHC

^aSDWA MCL February 1988.

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ESKIMO CREEK, NORTH AND SOUTH BARREL BLUFF, AND NAKNEK RIVER SEEP METALS SCREEN

Table 4-16E

Naknek <u>River Seep</u> <u>KS-SM04</u> 10/31/87 ទុ KS-SW03-04 DUP SW03-02 10/31/87 North Barrel Bluff KS-SM03 KS-SM03 -02 -03 10/31/87 10/31/87 120 00 12000 12000 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (100 < (1 10/31/87 10/31/87 KS-SM03 ទុ South Barrel Bluff KS-SW02 KS-SW02 KS-SW02 -02 -03 10/28/87 11/01/87 10/28/87 ទុ 10/28/87 Eskimo Creek KS-SW01 KS-SW01 -07 10/28/87 ş **Detection** Limit Units 1/6nn E200.7 E206.2 E206.7 E200.7 E2 E200.7 E200.7 E270.2 E272.2 E200.7 E279.2 E200.7 E200.7 E204.2 E204.2 E200.7 E200.7 E200.7 Method Antimony Silicon Molybdenum Boron ^aSDWA MCLs. Parameters Arsentc Barium Beryllium Cadmium Calcium Chromium Cobalt Copper Iron Lead Manganese Potassium Silver Sodium Thallium Vanadium Aluminum Selenium fercury Nickel 21nc

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^bRCRA MCL.

^CCWA Amblent Water Quality Criteria for Protection of Aquatic Life Freshwater Acute/Chronic.

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While collecting the surface water samples, an oily discoloration in the sediments was observed at sample location SWO3-1.

4.1.5.2.4 South Barrel Bluff

One organochlorine pesticide/PCB compound, chlordane, was reported at 0.2 ug/l. This detection was observed in the King Salmon Creek surface water sample collected at the upstream location.

Concentrations of calcium, iron, magnesium, and boron show an increased trend between the upstream and downstream locations. This may indicate that a source for these constituents exists between the two sample locations.

While collecting the surface water samples at the South Barrel Bluff, water (leachate) seepage from the toe of the bluff appeared to have an oily sheen and a discernible odor (land-fill gas).

4.1.6 Sediment Investigation

This section presents the results of the sediment investigation. Sampling procedures are discussed in Section 3.8, Environmental Sampling. Laboratory analytical reports and QA/QC data are presented in Appendix G, Analytical Data.

4.1.6.1 Eskimo Creek Seep

No sediment samples were collected at the Eskimo Creek seep site.

4.1.6.2 Naknek River Seep

No sediment samples were collected at the Naknek River seep site.

4.1.6.3 North Barrel Bluff

Sediment samples were collected at the same locations as the surface water samples along the toe of the North Barrel Bluff in the flood plain of King Salmon Creek. Samples were initially collected in the fall of 1987. These samples were analyzed for 1) organochlorine pesticides and PCBs, 2) semivolatile organic compounds, and 3) volatile organic compounds. The results of the analyses are shown in Tables 4-17A through 4-17C.

Methylene chloride and acetone were detected in SE05-01 at low concentrations. These constituents are widely used in analytical laboratories and may be the source for the concentrations measured in the sample analysis. Moisture content

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of the sediment samples was not determined, and the concentrations could not be converted to a dry weight basis.

Sampling and analysis for volatile organic compounds were repeated in the spring of 1988. The results of the repeat analyses are shown in Table 4-17D. No constituents were detected.

4.2 SIGNIFICANCE OF FINDINGS

Contaminant concentrations and migrations are discussed below. The significance of the findings may be viewed in terms of the potential exposure pathways (Figure 4-16) and the risk assessment (Section 4.3).

4.2.1 Eskimo Creek Seep

The approximate extent of groundwater contamination at the Eskimo Creek seep has been characterized by the combination of soil gas sampling, groundwater flow interpretations, and water quality sampling. A summary of these data is presented in Figure 4-17. The gross regional groundwater flow direction in the shallow unconfined aquifer is from north to south The flow pattern is locally controlled by across the site. 1) discharge to Eskimo Creek along the east and southeast and 2) the till surface topography south of the POL tanks. Groundwater flow is impeded by the elevated surface till diverting much of the flow, to the southwest toward MW-40 and to the south toward MW-37. A lowering of the water table due to seasonal fluctuations would increase the size and consequently the effect of this till barrier. Groundwater may be diverted further to the east around the till mound to discharge into Eskimo Creek under low water table conditions.

The elevated surface till may be more extensive than current data indicate, and thus have a greater effect on groundwater flow. Local groundwater flow directions will be altered significantly in areas where this till surface or possibly other till mounds intersect the water table.

Hydrocarbon groundwater contamination in MW-22 at POL tanks 11 and 12 (TPH of 5,666 ppm measured in groundwater) moves downgradient and is apparently split into two plumes by the till barrier. The southwest plume is identified by several soil gas points and well MW-39 (TPH of 14.9 ppm). The extent of this plume is defined on the south by nondetect soi? 'as points and well MW-33 (TPH less than 1 ppm). The dc_agradient and west edges of the plume are not fully defined, with distant wells MW-24 and MW-40 showing less than detection limit (1 ppm) of TPH.

The major groundwater plume (east) spreads from the POL tanks across the site to the east and southeast towards Eskimo

Table 4-17A Volatile Organics SW8240

SW8240 North Barrel Bluff Sediment - 1987

| Parameter | Units | KS-SE05-01 11/02/87 | KS-SE05-02 DUP SE05-04 11/02/87 | KS-SE05-03 11/02/87 | KS-SE05-04 11/02/87 |
|---|-------------------|------------------------|---------------------------------------|------------------------|------------------------|
| Chloro n ethane | ug/kg | 100 | 100 | IOU | 100 |
| Bromoethane | ng/kg | 100 | 100 | 100 | 100 |
| Vinyl chloride | ug/kg | IOU | 100 | 100 | 101 |
| Chloroethane | ng/kg | IOU | TOU | IOU | 100 |
| Methylene chloride | ng/kg | 80 0 | 12 | 90 g | 12 |
| Acetone | ng/kg | 20 | 89 | 12 | 69 1 |
| Carbon disulfide | ng/kg | 50 | 02 | 25 | 2 2 1 |
| 1,1-Dichloroethene | ug/kg | 50 | 05 | DC | 25 |
| I,I-Dichloroethane | ng/kg | 00 | | | 0C |
| Itans-1,2-alcolotoetnene | 54/50 | 00 | | | |
| CHIQTOIOLE 1.2-Dichloroethane | נומ/גם נומ/גם | 511 | 2015 | 50 | 50 |
| 2-Butanone | ua/ka | 100 | 100 | 100 | TOU |
| 1,1,1-Trichloroethane | ng/kg | 50 | 50 | 5U | 50 |
| Carbon tetrachloride | ng/kg | 50 | 50 | 50 | 50 |
| Vinyl acetate | ug/kg | 100 | IOU | TOU | 100 |
| Bromodichloromethane | ug/kg | 50 | 50 | 50 | 50 |
| 1,2-Dichloropropane | ng/kg | 50 | 50 | 20 | 5 |
| Cis-1,3-dichloropropene | ng/kg | 50 | 50 | 20 | 05 I |
| Trichloroethene | ng/kg | 50 | 50 | <u>1</u> | 3 8 |
| Dibromochloromethane | ng/kg | 3.1 | 50 | 20 | 3 1 |
| 1,1,2-Trichloroethane | ng/kg | <u>6</u> | 50 | 50 | B 2 I |
| Trans-1,3-dichloropropene | ng/kg | 20 | 20 | 20 | 31 |
| Benzene | ng/kg | 50 | 50 | 50 | 50 |
| 2-Chloroethyl vinyl ether | ng/kg | 100 | 100 | 100 | |
| Bromoform | ng/kg | 00 | nc Not | | 101 |
| 4-Metny1-4-pentanone | ng/ kg | 101 | | 101 | |
| Z"HEXANONE 1 1 2 J_Tetrachlorocthane | 54/5n | | 101 | 511 | 207 |
| t/t/t/t/torrachitotocondic Tatrachloroothone | 54/55 | 511 | 211 | 2115 | 511 |
| Toluene | 5%/50 110/ku | 211 | 3.1 | 511 | 50 |
| [h] orobenzene | 53/201 | 511 | 511 | 211 | 50 |
| Ft by 1 bensene | 647/65 | 511 | 511 | 50 | 50 |
| sturing the second | 54/55 | 215 | 212 | 21 | 50 |
| Vulenes (Atn) | 54/55 | 212 | 212 | 115 | 511 |
| XV)ene (v) XV)ene (v) | 10/kg | 50 | 50 | 50 | 20 |
| | | 1 | 1 | | |
| TB-Trip Blank | | | | | |
| Values in this table based on dry | weight. | | | | |
| U=Indicates constituent tested fo | r, but not detect | .ed. | | | |
| J≂Indicates an estimated trace va | lue. | | | | |

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| | COMPOUNDS | | |
|-------------|-----------------------|--------|--|
| Table 4-17B | SEMI-VOLATILE ORGANIC | SW8270 | |

| I | SEDIMENT |
|------|----------|
| 1270 | BLUFF |
| SWE | BARREL |
| | NORTH |

| | | | KS-SE05-02 | | |
|-----------------------------------|-------------------|---|---------------|---------------|---------------|
| | | KS-SE05-01 | DUP SE05-04 | KS-SE05-03 | KS-SE05-04 |
| Parameter | Units | 11/02/87 | 11/02/87 | 11/02/87 | 11/02/87 |
| h-nitroso-dimethvlamine | ua/ka | 3300 | ijus | 330[] | 330r |
| nherol | 110/kg | IIUEE | 33011 | 33011 | 3301 |
| antitue | ng/kg | 330U | 330U | 330U | 3301 |
| bis(2-chloroethyl) ether | ng/kg | 3300 | 3300 | 330U | 3300 |
| 2-ciılorophenol | ng/kg | 3300 | 3300 | 330U | 3300 |
| 1,3-dichlorobenzene | ng/kg | 3300 | 330U | 3300 | 3300 |
| 1,4-dichlorobenzene | ng/kg | 3300 | 330U | 330U | 3300 |
| benzyl alcohol | ng/kg | 3300 | 330U | 330U | 3300 |
| 1,2-dichlorobenzene | ng/kg | 3300 | 330U | 330U | 330U |
| 2-methylphenol | ng/kg | 3300 | 330U | 33 0 U | 3300 |
| bis(2-chloroisopropy1) | ng/kg | 33 0 U | 330U | 33 0 U | 330U |
| 4-methylphenol | ng/kg | 3300 | 3300 | 330U | 3300 |
| n-nitroso-di-n-propylamine | ng/kg | 3300 | 330U | 33 0 0 | 33 0 U |
| hexachloroethane | ng/kg | 33 0 U | 33 0 U | 3300 | 33 0 U |
| nitrcbenzene | ng/kg | 3300 | 330U | 3300 | 3300 |
| isophrone | ng/kg | 330U | 330U | 330U | 3300 |
| 2-nitrophenol | ng/kg | 3300 | 330U | 3300 | 3300 |
| 2,4-diethylphenol | ng/kg | 3300 | 330U | 3300 | 330U |
| benzoic acid | ng/kg | 1600U | 1600U | 16000 | 1600U |
| bis(2-chloroethoxy) methane | ng/kg | 3300 | 330U | 3300 | 3300 |
| 2,4-dichlorophenol | ng/kg | 3300 | 330U | 3300 | 330U |
| 1,2,4-trichlorobenzene | ng/kg | 33 0 U | 330U | 3300 | 3300 |
| naphtalene | ng/kg | 3300 | 3300 | 330U | 3300 |
| 4-chloroaniline | ng/kg | 3300 | 330U | 330U | 330U |
| Values in this table are based of | on wet we takt | | | | |
| limitates constituent tested f | for but not deter | | | | |
| J=Indicates an estimated trace v | value. | • | | | |
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| | | | KS-SE05-02 | | |
|---------------------------------------|-----------------------|------------|---------------|------------|---------------|
| | | KS-SE05-01 | DUP SE05-04 | KS-SE05-03 | KS-SE05-04 |
| Parameter | Units | 11/02/87 | 11/02/87 | 11/02/87 | 11/02/87 |
| | | | | | |
| hexachlorobutadiene | ng/kg | 330U | 330U | 3300 | 3300 |
| <pre>4-chloro-3-methylphenol</pre> | ng/kg | 3300 | 330U | 3300 | 3300 |
| 2-methylnarhthalene | ng/kg | 330U | 3300 | 330U | 3300 |
| hexachlorc clopentadiene | ng/kg | 330U | 3300 | 3300 | 33 0 U |
| 2,4,6-trichlorophenol | ng/kg | 330U | 3300 | 3300 | 3300 |
| 2,4,5-trichlorophenol | ng/kg | 16000 | 1600U | 1600U | 1600U |
| 2-chloronaphthalene | ng/kg | 330U | 330U | 3300 | 3300 |
| 2-nitroaniline | ng/kg | 16000 | 1600U | 1600Ŭ | 16000 |
| dimethyl phthalate | ng/kg | 330U | 33 0 U | 3300 | 3300 |
| acenaphthene | ng/kg | 330U | 330U | 3300 | 33 0 U |
| 3-nttroaniline | ng/kg | 16000 | 16000 | 16000 | 16000 |
| acenaphthene | ng/kg | 330U | 3300 | 3300 | 33 0 U |
| 2,4-dinitrophenol | ng/kg | 1600U | 16000 | 16000 | 16000 |
| 4-n1trophenol | ng/kg | 1600U | 1600U | 16000 | 1600U |
| dibenzofuran | ng/kg | 330U | 3300 | 3300 | 3300 |
| 2,4-dinitrotoluene | ng/kg | 330U | 3300 | 3300 | 330U |
| 2,6-dinitrotoluene | ng/kg | 330U | 3300 | 3300 | 3300 |
| diethyl phthalate | ng/kg | 330U | 330U | 330U | 33 0 U |
| 4-chlorophenyl phenyl ether | ng/kg | 330U | 3300 | 3300 | 3300 |
| fluorene | ug/kg | 330U | 330U | 3300 | 330U |
| 4-nitrcaniline | ng/kg | 1600U | 16000 | 16000 | 16000 |
| <pre>4,6-dinitro-2-methylphenol</pre> | ng/kg | 1600U | 1600U | 1600U | 1600U |
| n-nithosodiphenylamine | ng/kg | 330U | 3300 | 330U | 3300 |
| 4-bromophenyl phenyl ether | ng/kg | 330U | 3300 | 330U | 3300 |
| hexachlorobenzene | ng/kg | 330U | 3300 | 3300 | 3300 |
| pentacnlorophenol | ng/ky | 16000 | 1600U | 16000 | 16000 |
| phenanthrene | ng/kg | 330U | 3300 | 3300 | 330U |
| anthracene | ng/kg | 330U | 3300 | 330U | 33 0 U |
| d1-n-butyl phathalate | ug/kg | 330U | 330U | 330U | 3300 |
| Values in this table are based o | on wet weight | | | | |
| | con the set determine | | | | |
| U=Indicates constituent tested f | ror, but not detec | rea. | | | |
| J=Indicates an estimated trace v | value. | | | | |

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Table 4-17B (Continued)

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| 4-17B | nued) |
|-------|--------|
| Table | (Cont1 |

| | | | KS-SE05-02 | | |
|--|---------------------------------------|---------------|-------------|------------|---------------|
| | | KS-SE05-01 | DUP SE05-04 | KS-SE05-03 | KS-SE05-04 |
| Parameter | Units | 11/02/87 | 11/02/87 | 11/02/87 | 11/02/87 |
| fluoranthene | ng/kg | 330U | 330U | 330U | 33 0 U |
| benz idine | ug/kg | 16000 | 1600U | 1£ 00U | 1600U |
| pyrene | ng/kg | 3300 | 330U | 3300 | 330U |
| butyl benzyl phthalate | ng/kg | 330U | 330U | 330U | 33 0 U |
| 3,3-dichlorobenzidine | ng/kg | 6600 | 660U | 660U | 660U |
| benzo (a) anthracene | ng/kg | 3300 | 3300 | 3500 | 3300 |
| bis(2-ethylhexyl)phthalate | ng/kg | 3300 | 3300 | 3300 | 3300 |
| chrysene | ng/kg | 33 0 U | 330U | 33CU | 33 0 U |
| di-n-octyl phthalate | ng/kg | 3300 | 330U | 330U | 330U |
| benzo(b)fluoranthene | ng/kg | 330U | 330U | 330U | 330U |
| benzo(k)fluoranthene | ng/kg | 3300 | 330U | 330U | 330U |
| benzo (a) pyrene | ng/kg | 330U | 330U | 3500 | 3300 |
| indeno(1,2,3-cd)pyrene | ng/kg | 330U | 330U | 3300 | 33 0 U |
| dibenz (a , h) anthracene | ng/kg | 330U | 330U | 3301 | 3300 |
| benzo(g,h,i)perylene | ug/kg | 330U | 330U | 3300 | 330U |
| | | | | | |
| Vilues in unis caple are based of U=Indicates constituent tested f | on wet weight. for, but not detect | ted. | | | |
| J=Indicates an estimated trace v | alue. | | | | |

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Table 4-17C ORGANOCHLORINE PESTICIDES AND PCBS E608 NORTH BARREL BLUFF SEDIMENT KS-SE05-04 11/02/87 <0.004 <0.004 <0.04 <0.02 <0.004 <0.004 <0.004 <0.004 <0.004 <0°0 <0.04 <0.04 <0.04 <0.04 0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.01 <0.01 <0.004 <0.004 <0.004 <0.004 KS-SE05-03 11/02/87 <0.004 <0.004 <0.04 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.04 <0.04 <0.04 <0.04 <0.04 <0.02 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.01 <0.01 KS-SE05-02 11/02/87 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.04 <0.04 <0.04 <0.04 <0.04 <0.04 <0.02 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.01 <0.01 KS-SE05-01 11/02/87 <0.004 <0.004 <0.004 <0.004 <0.04 <0.04 <0.04 <0.04 <0.02 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.04 <0.04 <0.01 <0.004 <0.004 <0.01 Detection Limit <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.04 <0.04 <0.04 <0.04 <0.04 <0.04 <0.02 <0.01 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.004 <0.01 mg/kg ng/kg mg/kg mg/kg mg/kg Units mg/kg ∎g/kg mg/kg mg/kg mg/kg mg/kg ng/kg ng/kg Endosulfan Sulfate Heptachlor Epoxide Parameter Endrin Aldehyde Endosulfan II Endosulfan I methxychlor Heptachlor Chlordane Toxaphene PCB-1248 Dieldrin PCB-1221 PCB-1232 PCB-1242 PCB-1260 4,4-DDD PCB-1016 PCB-1254 PCB-1221 4 1-DDE 4,4-DDT Endrin Aldrin b-BHC g-BHC d-BHC a-BHC

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| Table 4-17D | VOLATILE ORGANICS | SW8240 | BARREL BLUFF SEDIMENT - 1988 |
|-----------------|-------------------|--------|------------------------------|
| | - | | BAR |

| Parameter | Units | KS-SE05-01 06/15/88 | KS-SE05-02 06/15/88 | KS-SE05-03 06/15/88 | KS-SE05-04 DUP SE05-02 06/15/88 | KS-TBSE 06/15/88 |
|--------------------------------------|--------------|------------------------|------------------------|------------------------|---------------------------------------|---------------------|
| Chloromethane | ng/kg | 160 | 200 | 140 | 2 4 U | IOU |
| Bromoethane | ng/kg | 16U | 200 | 140 | 240 | IOU |
| Vinyl chloride | ug/kg | 160 | 20U | 140 | 24U | TOU |
| Chloroethane | ng/kg | 160 | 200 | 140 | 24U | TOU |
| Methylene chloride | ug/kg | 20 | 27 | 19 | 44 | 3.7 |
| Acetone | ng/kg | 40 | 32 | 96 | 26 | 101 |
| Carbon disulfide | ug/kg | 80 | 101 | 70 | 120 | 50 |
| 1,1-Dichloroethene | ng/kg | 80 | 101 | 70 | 120 | 50 |
| <pre>1,1-Dichloroethane</pre> | ug/kg | 80 | TOU | 70 | 120 | 50 |
| Trans-1,2-dichloroethene | ng/kg | 80 | TOU | 70 | 120 | 50 |
| Chloroform | ng/kg | 80 | 100 | 0.2 | 120 | 50 |
| 1,2-Dichloroethane | ng/kg | 80 | 100 | 70 | 120 | 50 |
| 2-Butanone | ng/kg | 16U | 20U | 20 | 2 4 U | 100 |
| 1,1,1-Trichloroethane | ng/kg | 80 | 100 | 70 | 12U | 50 |
| Carbon tetrachloride | ng/kg | 80 | IOU | 70 | 120 | 5U |
| Vinyl acetate | ng/kg | 160 | 200 | 140 | 240 | TOU |
| Bromodichloromethane | ng/kg | 80 | IOU | 70 | 120 | 50 |
| 1,2-Dichloropropane | ng/kg | 80 | 100 | 70 | 120 | 50 |
| Cis-l,3-dichloropropene | ng/kg | 80 | 101 | 70 | 120 | 50 |
| Trichloroethene | ng/kg | 80 | 100 | 70 | 120 | 5U |
| Dibromochloromethane | ng/kg | 8U | 100 | 70 | 120 | 50 |
| 1,1,2-Trichloroethane | ng/kg | 80 | 100 | 70 | 120 | 50 |
| Trans-1,3-dichloroproper | ng/kg | 80 | 100 | 70 | 120 | 50 |
| Benzene | ng/kg | BU | 100 | JU | 120 | 50 |
| 2-Chloroethyl vinyl eth | ng/kg | 16U | 200 | 14U | 24U | 100 |
| Bromoform | ng/kg | 80 | 100 | 70 | 120 | 50 |
| 4-Methyl-2-pentanone | ng/kg | 16U | 200 | 140 | 240 | 100 |
| 2-Hexanone | ng/kg | 16U | 200 | 14U | 2 4 U | 100 |
| <pre>1,1,2,2-Tetrachloroethane</pre> | ng/kg | 80 | 100 | 80 | 120 | 50 |
| Tetrachloroethene | ng/kg | 80 | 100 | 70 | 120 | 50 |
| Toluene | ng/kg | 80 | IOU | 4 J | 120 | 50 |
| Chlorobenzene | ng/kg | 80 | 100 | 70 | 120 | 50 |
| Ethylbenzene | ug/kg | 8U | TOU | 70 | 120 | 50 |
| Styrene | ug/kg | 80 | TOU | 70 | 120 | 50 |
| Xylenes (o+m) | ug/kg | 80 | 100 | 70 | 120 | 50 |
| Xylene (p) | ug/kg | 80 | 101 | 7U | 120 | 50 |
| marts Blank | | | | | | |
| Values in this table based on (| dry watcht | | | | | |
| U=Indicates constituent tested | for, but not | detected. | | | | |
| J=Indicates an estimated trace | value. | | | | | |

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GROUNDWATER PLUME INTERPRETATION KING SALMON AIR FORCE STATION

KING SALMON, ALASKA

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Creek. The interpretations of groundwater flow suggest that contaminated groundwater moves to the east around the till, and then southeast towards MW-6 and MW-10. This scenario, however, does not provide a definite source for the Eskimo Creek seep. A second, currently unrecognized, source may exist hydraulically upgradient of the seep that could also account for the large soil gas plume in the northeast portion of the site. Further analysis of groundwater flow patterns (i.e., seasonal fluctuations) additional monitoring wells and water quality sampling would be necessary to further delineate the source of the Eskimo Creek seep contamination.

Hydrocarbon contamination (TPH and aromatics) in the unsaturated soils is highest in areas correlating with the groundwater plume. However, soils contamination (TPH) at lower levels (less than 200 ppm) does appear to be much more extensive than the plume. Past site practices of road oiling for dust control probably spread hydrocarbon contamination across the site. Other potential sources include underground utility corridors and unreported underground tank locations, drum storage areas, and general cleaning/degreasing of equipment in open areas. These localized areas outside of the primary contaminant plume should be considered as potential future sources for continuing releases to shallow groundwater.

Visual evidence of petroleum contamination was observed discharging with shallow groundwater into the wetland area west of Eskimo Creek. The wetland serves as a migration pathway into the creek. However, for the analyses that were performed on the samples of the creek water no constituents were detected.

The nature of soil and groundwater contamination does not appear to consist of the more volatile hydrocarbons. However, the individual constituents could not be identified on the basis of the analyses completed.

Three intermediate wells were constructed next to (nested) shallow wells to investigate the hydraulic relationships between aquifers and to determine if shallow groundwater contamination has migrated vertically beneath the site. A strong downward vertical gradient was found between the unconfined and intermediate aquifers. However, analytical results did not indicate contamination in any of the three intermediate wells. Because the upper aquifer is a recharge source for the intermediate aquifer, this conclusion should be verified with more extensive sampling.

The direction of groundwater flow in the intermediate zone is apparently to the south. Verification of this interpretation will require additional water level measurements. The station's two deep wells produce water from a confined system that is screened at depths greater than 200 feet below ground surface. This zone is about 120 feet below the intermediate aquifer, where it is screened by the new wells. The relationship between these two systems is not well defined. Vertical gradients between the two systems cannot be evaluated because the base wells are actively pumped.

Water quality analyses for the base wells indicate hydrocarbon contamination (TPH of 27.1 and 6.4 ppm) in the wells. The individual constituents and their associated risk to human health are unknown.

Because the nested intermediate wells do not appear to be contaminated, a highly likely migration pathway to the deep aquifer is communication along the well bore annulus of the deep wells. This interpretation should be confirmed with more extensive sampling. If the station wells are acting as a source for deep aquifer contamination, the nature and extent of that contamination should be evaluated, and the wells should be abandoned to prevent further contamination.

4.2.2 Naknek Seep

The soil gas analysis and water sample collected at the Naknek seep verify the presence of hydrocarbon contamination at this location. The data are insufficient to allow further interpretation of the nature, extent, or exact source of contamination. However, the proximity of the contamination to the reported location of an abandoned fuel storage tank and the presence of an existing tank suggest the contamination may be localized in this area.

4.2.3 North Barrel Bluff

Geophysical survey data collected at the North Barrel Bluff indicate that the drums extend approximately 50 to 100 feet from the bluff along the lines surveyed. The vertical extent of the drums could not be determined with geophysical techniques, but visual observations indicate that the vertical extent may span a depth equal to the entire bluff face. Additional areas where metallic debris were buried in trenches were identified. Several more of these small "dumps" could exist within the large disturbed area bordering the bluff.

Sediment samples collected in 1987 at the toe of the bluff contained trace concentrations of organic and inorganic constituents. A resampling of these points did not verify these results. Analysis of surface water samples from the bog at the face of the bluff and from King Salmon Creek did not indicate the presence of constituents above method detection limits.

Many of the drums appeared to be intact and are expected to contain residual fluids. These fluids represent future sources for releases to soil, surface and groundwater. The unsaturated soils beneath the drum and debris area could contain liquids that have already leaked from the drums.

The surface water and sediment samples were not adequate to provide an indication of whether releases are occurring along the bluff. Soil borings into the sediments beneath the drums are required to obtain samples that would be adequate to characterize potentially contaminated soils. Because of uncertainty about the types of waste deposited in this "landfill," the samples should be analyzed for a wide range of constituents.

The greatest potential for adverse affects to human health and the environment results from physical exposure to decomposing materials contained within the bluff area. The uncontrolled areas of buried drums and other debris types pose a potential threat through both physical contact and airborne chemical exposure.

4.2.4 South Barrel Bluff

Geophysical survey data collected along the South Barrel Bluff identified the presence of buried drums and other metallic debris along the bluff face, similar to that observed at the North Barrel Bluff. The large area west of the bluff was determined not to be a landfill. However, an area of smaller extent along the bluff that includes two residential homes does appear to be the site of a landfill. Other small areas within the disturbed areas along the bluff were also identified as "disposal areas."

Visual observations along the toe of the bluff indicated the presence of discolored liquids (i.e., leachate) discharging into the creek. However, for the analysis performed on the surface water samples from King Salmon Creek, no constituents were detected.

The surface water sampling program performed during the Stage I effort was not adequate to provide an indication of whether releases are occurring along the bluff. This sampling represents a limited effort of three sampling locations spaced along the length of the debris area. Additionally, this effort represents a one-time sampling event that does not address the issue of continuing releases and seasonal effects on contaminant transport. Soil borings into the sediments beneath the drums are required to obtain samples adequate to allow characterization of potential contamination. Because of the uncertainty about the types of waste deposited in this "landfill," samples should be analyzed for a wide range of constituents (priority pollutants).

Analysis of drinking water from a residential well located within the landfill area did not indicate hazardous constituents for the parameters tested. Other residential wells in the area were not sampled. The potential must be considered high for contamination in these wells through borehole leakage (contaminant migration along annulus of well).

4.3 <u>RISK ASSESSMENT FOR HEALTH, WELFARE, AND THE ENVIRONMENT</u>

A qualitative risk assessment was conducted for this report. The limited information available characterizing King Salmon AFS wastes made any quantitative estimation of potential risk unfeasible. More complete data (e.g., priority pollutant scan information) will be needed for contaminated site media before a quantitative risk analysis can be conducted. The following sections address the contaminants that have been identified to date. Background data for local water, soil, and sediment are needed to properly assess the potential risks King Salmon AFS wastes contribute to ambient conditions. Also, more complete knowledge of King Salmon AFS wastes (TPH constituents, TOC constituents, etc.) will be needed to adequately address the human health and environmental risks that may be associated with each operable uni'. References to potential human or environmental risks presented in this assessment could change significantly as more contaminant data become available.

4.3.1 Waste Characterization

Discussions in the following sections are based on current knowledge of King Salmon AFS contamination. It should be noted that this knowledge is limited. Background data for local water, soil, and sediment are needed to properly assess the potential risks "ing Salmon AFS wastes contribute to ambient conditions. Also, more complete analyses of TPHs, TOCs, and suspected priority pollutants will be needed to more fully characterize King Salmon AFS contamination.

4.3.1.1 Identifiable Contaminants

The four areas at the King Salmon AFS were tested for over 146 inorganic, volatile organic, semivolatile organic, and pesticide contaminants. Of these, 39 potential contaminants of concern (or contaminant types, i.e. THP, TOC) have been identified (Table 4-18). These contaminants are considered in this report because their concentrations (or detection

| | Soil and Sediment | Groundwater | Surface Water |
|----------------------------|----------------------|-------------|------------------|
| Inorganics | Seatment | GLOUNGWALCL | Hacci |
| <u>a</u> | | | |
| Cadmium | | | |
| Copper | | | |
| Fluoride | | + | |
| Lead | | + | |
| Mercury | | | |
| Nitrate-Nitrite | | + | + |
| Manganese | | + | + |
| Barlum | | | + |
| Zinc Nichal | | | + |
| Nickel | | | + |
| Silver | | | + |
| Boron | | | · |
| Organics | | | |
| Benzene | | + | |
| bis(2-ethylhexyl)phthalate | + | | |
| 2-Butanone | + | | |
| Butyl benzyl phthalate | + | | |
| Ethylbenzene | + | + | |
| Fluorene | + | | |
| 2 Methylnapthalene | + | | |
| Napthalene | + | | |
| Toluene | + | + | + |
| trans-1,2-Dichloroethene | | | + |
| Trichloroethene | | | + |
| Xylenes | + | + | + |
| THP | + | + | |
| roc | | + | |
| Chloromethane | + | | |
| Methylene chloride | + | | |
| Acetone | + | | |
| Phenanthrene | + | | |
| Di-n-octyl phthalate | + | | |
| Chloroform | | | + |
| Pesticides | | | |

Table 4-18 POTENTIAL CHEMICALS OF CONCERN DETECTED AT KING SALMON AFS

Chlordane Dieldrin^a Endosulfan^a Endrin^a Heptachlor^a Methoxychlor^a Toxaphene +

^aChemical not detected put the analytical detection limit was greater than ambient water quality criteria for aquatic toxicity or drinking water.

limits) exceed federal or state criteria for the protection of human health and the environment, they are consistently present in various site media (groundwater, surface water, soils, sediments), or they are potentially toxic to humans and the environment (carcinogenic, noncarcinogenic, bioaccumulation potential).

4.3.1.2 Toxicity of Site Contaminants

Toxicity profiles outlining the health hazards associated with select King Salmon AFS contaminants are presented in Table 4-19. High concentrations of total petroleum hydrocarbons (TPH) and total organic carbon (TOC) have also been identified in the shallow wells and soils at the site. Although no toxicity data have been identified for these classes of compounds, they could be composed of chemicals that adversely affect human health and/or the environment. Critical toxicity values for protection of public health (as recommended by EPA) are summarized in Table 4-19A. Eleven of these chemicals are classified by EPA as carcinogens.

4.3.1.3 Waste Mobility

The physical and chemical properties of the contaminants detected at King Salmon AFS that may influence their environmental fate, transport, or toxicity are summarized in Table 4-20. Many of the contaminants detected have moderateto-high mobility in soils and groundwater. Many of the contaminants are also highly volatile. The TPHs detected at the site showed low mobility in soils and groundwater (based on the extent of the plume identified) and exhibited low volatility when measured in the field.

4.3.2 Human Health/Environmental Criteria Comparison

4.3.2.1 Eskimo Creek Seep

4.3.2.1.1 Shallow Groundwater

The sampling results for the shallow groundwater wells at Eskimo Creek are summarized in Table 4-21. Table 4-21 also presents a comparison of shallow well water quality data to criteria for drinking water and the protection of aquatic organisms. The aquatic organism criteria comparison is valid because contaminants from this shallow aquifer could enter Eskimo Creek. This stream supports fish and wildlife populations. The following paragraphs concern the Eskimo Creek shallow well contaminants (lead, total dissolved solids, benzene, and xylene), which have maximum concentrations that exceed federal or Alaska water quality criteria.

| (and call | Acute Toxicity Summary | Chronic Toxicity Summary | Cancer Potential | Other |
|--------------------------------|---|--|--|---|
| tretone | Generally regarded as having low towicity and, therefore, has not been studied extensively. May produce narcosis, CNS depression, nausea, vomiting, gastric hemorrhage, atoxia, coughing, and bronchial irritation. | Prolonged inhalation of high concentra- tions may produce irritation of the re- spiratory tract, coughing, headache, drywsiness, loss of coordination, in- flammatory bronchitis, conjunctivitis, paryngitis, gastroduodenitis, renal tubular necrosis, and hyperglycemia. | Not applicable. | Possible mutagenicity, can potentiate carbon tetrachloride hepatotoxicity. |
| Bartum | Ingestion of barium salts can chuse prolonged muscular stimulation; gastroenteritis; hypokalemia; and cerdiovascular effects such as ventricular fibrillation and extra systoles. | Prolonged occupational inhalation has resulted in baritosisa benign, reversible pneumonitis. | Not applicable. | Toxicity of compounds depends on solubility. |
| Benzene | Acute exposures (inhalation) to high levels of benzene may lead to depression of the central nervous system, uncon- sciousness, and death or may cause fatal cardiac arrhy'hmias. | Major toxic effect is hematopoietic toxicity (affects formation of blood); chronic exposure of workers to low levels has been associated with blood disorders, such as leukemia and aplastic anemia (depression of all three cell types of the blood in absence of functioning marrow). | Sufficient evidence that is human and animal carcinogen; strong correlation between erposure to benzeme by inhalation and leukemia. | Chromosomal aberra- tions in bone marrow and blood have been reported in experi- mental animals and some workers. |
| Bis (2-ethy lhexy l) phthalate | Low acute fowicity in experimental animals, accidental acute exposure in man resulted in mild gastric disturbance and catharsis. | Chronic exposure at relatively high concentration have retarded growth and resulted in increased liver and kichney weight in experimental animals. | No evidence for carcinogenicity in man. However, is a potential animal carcinogen. Oral administration to rats and mice resulted in increased heptacellular carcinomas or neoplastic nodules. | Some evidence in animals of teratogenic and fetotoxic effects. Poorly absorbed throu skin; repidly metabolized. |
| Boron 2-Butanone | No information reported. The compounds is of relatively low toxicity, but at high doses affects the nervous system and causes irritation of the eys, man skin. The oral L_D value for the rat was 27_{50} mg/kg. Exposures to high concentrations may retard fetal development and cause some teratogenic effects. | | Has not been adequately tested for carcinogenicity. | Although 2-Butanone not strongly neurotoxic on its ow it apparently strong potentiates the neurotoxicity of n-hexanone. |

health effect or target organ may be based on animal studies and does not imply that the results of exposure to humans will be the same.

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| Butyl benzyl phthalate | No Information reported. | No information reported. | No information reported. | |
|------------------------|---|--|--|-------------------------------|
| Cadatus | For acute exposures by ingestion, symptoms of cadmium toxicity include nausee, vomiting, diarrhea, muscular cramps, salivation, spasms, drop in blood pressure, vertigo, loss of consciousness, and collapse. Acute renal failure, liver damage, and death may occur. Exposure by inhalation can cause irritation, coughing, ibbored respiration, vomiting, acute chemical pneumonitis, and pulmonary edema. | Respiratory and remal toxicity are major effects in workers. Chronic oral exposures can produce kidney damage; cadaium accumulates in kidney, and nephropathy results after critical concentration in kidney is reached, probably about 200 ug/g. Inhalation can cause chronic obstructive pulmonary disease, including bronchitis, progressive fibrosis, and emphysema. Chronic exposure affects calcium metabolism and can cause loss of calcium from bone, bone pain, osteomalacia, and associated with hypertension. Cadmium can product testicular atrophy, sterility, and teratogenic effects in experimental animals. | Increased risk of prostate cancer and perhaps respira- tory tract cancer in workers exposed by inhalation. No evidence of carcinogenicity from chronic oral exposure. | A nonessential element. |
| Chlordabe | In clinical cases of humans exposed to chlordame, central nervous system (CNS) effects (irritability, salivation, convulsions, depression, and death) and blood dyscrustas (e.g., aplastic anemia and leukemia) vere observed; routes of exposure included oral, inhalational, and dermal. Neuroblastoma has been associated with exposure to chlordane and heptachlor, as well as an increase in cerebrovascular disease. In animals, CNS disturbances and kidney and liver effects were seen. | Mejor target organ is liver. Animal studies indicate toxicity to liver, kidney, heart, lung, and increased mortality. Decreased fertility in mice and toxicity to mursing rat pups have been reported. | An increased incidence of hepatocellular carcinoma (liver tumors) has been observed in mice fed diets containing analytical grade chlordane. | |
| Chloroform | Anesthetic depresses CHS. Fatalities in hurans may be repid, resulting from cardiac arrest (apparently sensitization to epinephrine) or delayed with kidney and liver damage; respiratory depression, coma, liver and kidney damage are among the symptoms of exposure to chloroform. In laboratory animals, acute toxicity depends on species, strain, sex, and ags: i.ver damage may be cuased of death in rats and mice after acute exposure. | Kidney damage (renal tubular necrosis) can occur in mice; and kidney and liver damage in rats, rabbits, dogs, and guinea pigs exposed by inhalation. | Carcinogenic in mice (hepatomas, hepatocellular caricinomas), male rats (malignant kidney tumors), and demale rats (thyroid tumors). | Tetotoxic in rats rabbits. |
| Chloromethane | No Information reported. | | | |

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Table 4-19 (continued) .

| | | (continued) | | |
|----------------------|--|---|--|--|
| Chenical | Acute Toxicity Summary | Chronic Toxicity Summary | Cancer Potential | Other |
| Copper | Exposure (inhalation) to copper dusts result in symptoms similar to metal fume fever. Exposure to metal fumes results in upper respiratory tract irritation, metallic or sveet taste, neure, metal fume fever, and skin and hair discolora- tion. Exposure to dusts and mists of copper sait result in congestion of nasai mucous membranes, sometimes of pharnyx, and occasional ulcertion with perforation of nasal septum. Acute copper sulfate poisoning in humans (oral) sometimes fatal; includes vomiting, diarrhea, hypotension, coma, and jaundice. | Hemolytic anemia after chronic exposure in some dialysis patients. Sensitive individuals with disorders of metaboli:mHilson's disease and Menke's disease. | Not Applicable | Essential nutrient. Organoleptic threshold in water between 1 to 5 mg/l. |
| Dieldrin | High order of acute toxicity. The oral LD50 for the rat is approximately 50 mg/kg, and the rat dermal LD50 is about 100 mg/kg. | Chronic toxic effects include liver toxicity and CNS abnormalities. Dieldrin is teratogenic and toxic to the reproductive system. Reproductive effects include decreased fertility and increased fertal deaths. Teratogenic effects include cleft palate, vebbed foot, and skeletal anomalies. | Considered a B2 level carcinogen by CMG. A B2 carcinogen is one considered to have sufficient evidence of carcinogenicity in animals, but inadequate data in humans. | Suggestive evidence of Butagenicity avallable. |
| D1-n-octyl-phthalate | No information reported. | | | |
| Endosulfan | Endosulfan acts on the central nervous system, lungs, and skin. Acute exposure has caused death in humans. | No information reported. | No carcinogenic, mutagenic, or teratogenic potential has been reported. | There Is a general absence of tcxic reports concerning Endosuifan. |
| Endrin | Endrin acts on the central nervous system and liver. Symptoms of acute exposure include convulsions, vomiting, abdominal pain, nausea, diziness, and heddche. Acute exposure has caused death in humans. | No information reported. | Potential indicated for oncogenicity and teratogenicity in animals. | EPA filed a rebuttable presumption motice for endrin in 1987 on the basis of omccyenicity, teratogenicity, and reductions it. endan- gered and muntarget species. |
| Ethylbenzene | Ethylbenzene is irritating to eyes, mucous membranes, and skin. It can cause headaches and narcosis. | | | |

^a Health effect or target organ may be based on animal studies and does not imply that the results or exposure to humans will be the same.

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|---|------------|---|---|--|--|
| Hototototototototototototototototototot | Fluoride | Fluoride intoxication from inhelation can produce brochospasm, laryngospasm, polmonary edema, irrepairable damage to lungs, and death. Fluoride intoxication through ingestion can cause nauses, vomiting, cramps, diarthee, convuisions, and coma. | Chronic fluoride exposure can cause nose bleeds, sinus trouble, and irritation of the skin, eyes, mucus membranes, and lungs. Mottling of dental enamel and skeletal fluorosis are other chronic exposure affects. | No carcinogenic potential reported. | The magnitude of toxi effects from fluoride exposure depends on the irritating properties of the compound exposed to. |
| Lot Lot in rougenic lead interaction in montain is interactively upon in the cartion is the montain interactively and interactively the montain interactively and interactively and respiratory artest, and respiratory artest, artest artes | Heptachlor | Acutely toxic doses can cause tremors, convuisions, abnormal behavior, respira- tory collapse, and hyperirritability. Acute exposure by various routes can cause development of hepatic vein thrombl, can affect the CNS, and cause death. The oral LD50 for the rat is 40 mg/kg. | Chronic exposures induce liver changes, affect hepatic microsomal entyme activity, and cause increased mortality in offspring. | Considered a B2 level carcinogen by CMG. A B2 carcinogen is one considered to have sufficient evidence of carcinogenicity in animals, but inadequate data in humans. | |
| | Peg | Acute inorganic lead intoxication in humans is characterized by encephaiopathy, abdominal pain, hemolysis, liver damage, renal tubular necrosis, seizures, comma, and respiratory arrest. | Chronic law levels of exposure to lead can affec. the hematopoletic system, the nervous system, and the cardiovascular system. Lead inhults several ky erzymes involved in heme biosynthesis. One characteristic effect of chronic lead intorication is anomia, by reduced hemoglobin production and shortened erythrocyte surrival. In humons, lead erythrocyte surrival. In humons affect the immune system and produce gingival lead lines. Epidemiological studies have indicated that chronic lead strosure may be associated with increased blood pressure in humons. Exposure to lead is associated with sterility, abortion, neonatal mortality, and morbidity. Organolead compounds are neurotoxic. | Lead salts have some evidence of carcinogenicity in animals. | Children are especially sensitive to low level effects. |

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| | Andre Treite Simmary | (continued) Chronic Toxicity Summary ^a | Cancer Potential | Other |
|--------------|--|--|------------------|---------------------------|
| Chemical | | | | |
| | the state of a second and the very bigh | Chronic manganese poisoning results from | | Manganese is an essen- |
| h anganese | | tohalation to high concentrations of man- | | tial nutrient. Mangan- |
| | concentrations can cause manyanese | reaction of the chronic mandanese polsoning | | ese concentrations in |
| | paeumon1t1s. | Partets output for the second at the second se | | water a 50 ug/l may |
| | | 12 CDGLGCLCELIZED DY PACING LITER | | exhibit undesirable tasts |
| | | such as irritability, difficulty in wain- | | and discloration. |
| | | ing, speech distrubances, and computsive | | |
| | | behavior and by encephalopathy and pro- | | |
| | | gressive deterioration of the central | | |
| | | nervous system. Chronic effects of man- | | |
| | | structure of an and a parkingun's | | |
| | | danese potsuming at a statistic to the statistic of a second statistic of the statistic of the second statistic of the statistic of the second statist | | |
| | | disease. Liver changes are also itrequently | | |
| | | seen. Individuals with an iron deficiency | | |
| | | may be more susceptible to chronic poison- | | |
| | | | | |
| | | * F ₇₁ | | |
| | | | | Mercury crosses |
| Mercury | Inhalation of mercury vapor can cause | according to athendra tentor polo | | placenta. Toxicity |
| | bronchitis and nervous system effects. | metcury can produce ettects on nervous | | denends on chemical |
| | Area accorded to the abdominal | system, including tremors, erethism, | | |
| | TOTO TOTO TOTO TOTO | mismilar weakness personality changes, | | torm. Metallic, |
| | cramps, gastrointestinal errects, | all and horizon and the second s | | organic, and inorganic |
| | ulceration, shock, circulatory collapse, | GINGIVILLS, AND COLORED BYE LETTER. | | compounds can be |
| | and renal failure. | In children, pink disease has been | | be a stranged armod |
| | | reported after ingestion of mercurous | | |
| | | commoninds Exposure to organic mercury | | |
| | | compared and visual dis- | | |
| | | | | |
| | | turbances, lingling, parestuestae, | | |
| | | numbhess, tunnel vision leading to | | |
| | | blindness, weakness in extremities and | | |
| | | Drogressive ataxia, tremor, cerebral | | |
| | | Arroby, and degeneration of nerves; | | |
| | | and management of the state | | |
| | | VISUAL, PETIDNETAL NEUROPACITY, WIL | | |
| | | | | |
| | | No information reported | ot applicable. | Methoxychlor is an |
| Methoxychlor | No human effects reported. In animals, | NO THEOTHEOTHEOTHEOTHEOTHEOTHEOTHEOTHEOTHEO | | insecticide of low |
| | intextcation symptoms include trembling, | | | mammal toxicity used |
| | convulsions, kidney, and liver damage. | | | for home and garden, |
| | | | | domestic animal, and |
| | | | | |
| | | | | COMMETCIAL AGRICUL- |
| | | | | tural pest control. |
| | | | | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |

a Health effect of target organ may be based on animal studies and does not imply that the results of exposure to humans will be the same.

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| | Acute Toxicity Summary | Chronic Toxicity Summary ^a | 1011111 1011111 | Other |
|---------------------|---|---|---|--------------------------------------|
| Methylene Chloride | Methylene chloride acts as a mild marcottc irritating to eyes and upper respiratory passages. At low levels, it may cause elevated carboxyhemoglobin, preventing transfer of oxygen to tissues. | Causes liver and kidney damage, and may substantially increase carboxyhemoglobin levels; preventing the transfer of oxygen to tissues. | The National Toxicological Pro- gram (NTP) classifies methylene chloride us huving clear evi- dence of carcinogenicity in female rais and in male and Agency for Research on Cancer (IARC) crimsiders it a level 3 Agency for Research on Cancer (IARC) crimsiders it a level 3 carcinogen; hovever, this IARC study was released prior to the time the NTP study was made available. A level 3 carcinogen is one that has inadequate evi- dence available to prove its carcinogenicity. | |
| 2-Hethylnaphthalene | No information reported. | No information reported. | No information reported. | |
| Maphthalene | Inhalation or ingestion of dust or vapor can cause intravascular hemolysis, nausea, vomiting, jaundice, hematuria, hemoglobinuria, and acute renal shutdovn. | Chronic effects include skin and eye irritation. | Not applicable. | |
| N1.cke 1 | Sigms of acute nickel toxicity may include headaches, nausea, vomiting, chest pain, sough, hyperpnea, cyanosis, gastrointestinal and central nervous system effects, weakness, fever, pneumonia, respiratory tallure, cerebral edma, and death. Acute exposures to nickel containing dust may result in chemical pneumonitis. | Rhinitis, masal simusitis, and masal mucosal injury are among the effects reported among workers chronically exposed to various nickel compounds. Allergic contact dermatitis and other dermaticidical effects are the most frequent effects of dermal exposure to nickel and nickel-containing compounds. | There is extensive epidemolo- gical evidence indicating excess cancer of the lung and nasal cavity for workers ex- posed to certain nickel com- pounds. Nickel compounds implicated as having carcino- genic potential include in- genic potential include in- soluble dusts of nickel sub- sulfide and nickel carbonyl and soluble aerosols of nickel sulfate, nickel carbonyl. | May or may not t essential elemen |
| Xitrate | Acute toxicity can result when nitrates are reduced to nitrites. Nitrites oxidize hemoglobin, inhibiting ability to carry oxygen in the blood. Anoxia and death may ensue. | No information reported. | Nitrate exposure can lead to the formation of nitrosumines that are known human carcinogens. | |

| Pheuthrene No informatio Silver Argvria (loco | te loxicity summary | | | |
|--|--|--|---|--|
| Silver Argyria (loci | on reported. | | | |
| Impregnation discoloration affect eyes (Gastrointest) with oral ex | <pre>al of generalized of tissue causes n of skin and evel may and respiratory tract. inal irritation may occur posure.</pre> | Kidney and lung damage and possibly arterioscierosis have been reported for industrial or medical exposures. | | |
| Toluene expose mentally or of intentional of tion, then CT heurotoxic et ievels and co ness at high thon have t pulmonary iri exposure. | ed by inhalation experi- occupationally or by abuse may exhibit excita- NS depression and necrosis; ffects include nausea, incoordute nat low onfusion, ataxia, and wek- er levels; in rats, irrita- us membranes and incoordi- been observed, as well as ritation with subchronic | CNS effects have been reported in work- ers, such as disturbances in memory and thinking, psychomotor skills, visual accuracy, sensorimotor speed, and per- formance tests; indications of carebral and cerebellar dysfunction include trem- ors, ataxia, and equilbrium disorders, bizarre behavior and emotional lobility may occur. In cases of abuse, changes in liver and kidney function have been observed. In rats, a decrease in hematocrit has been reported. | Empryotoxicity and possible teratogenicity in mice have been reported in an abstract; in rats, skeletal retardation of offspring has been described. | |
| Toxaphene Acute toxic (contuston, tr unconsciousn | effects include nausea, remors, convulsions, and ess. | Chronic effects include skin and eye irritation. | Toxephene is an animal- positive carcinogen. | Toxaphene is cutr the most heavily insecticide in th USA. |
| Trans-1,2-Dichloroethene Has aneschet. high vapors o weakness, tr | ic properties. Exposure at causes nauses, vomiting, emor, and cramps in humans. | Animals exposes to prolonged inhalation results in loss of appetite, decreased body weight, and pathological changes in liver, kidney, and lungs. May interact with drugs and the metabolizing mono- oxygenase system. | Not Applicable | Very little info available on expo to trans-1,2-dich thene. |

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| | | Table 4-19 (continued) | | |
|--|--|---|--|--|
| Chem Ica I | Acute Toxicity Summary ^a | Chronic Toxicity Summary ^d | Cancer Potential | Other |
| Trichloroethy lene | Exposure to TCE can cause depression of the central nervous system, including distines, headaches, incoordination similar to that induced by alcohol, nausea, vomiting, and unconsciousness. | Long-term inhalation exposure can aifect liver and kidneys in animals. In humans, changes in liver entymes have been associated with TCE exposure. | Exposure of mice (orally and by inhalation) and rats have produced increases in liver or lung or kidney tumors. | "begreasers flush" has been described in "CE-exposed workers Who consume alcohol. |
| Xy lene | Acute exposures to inhaled xylene can depress the central nervous system and irritate mucous membranes. | Changes in behavioral tests, manual coordination, balance, and electro- encephalographic patterns have been reported in humans exposed to xylenes; development of tolerance against some of these effects has been described. Effects on liver of rats have been reported. | | |
| 2 lbc | Acute adverse effects of zinc include acute metal fume fever by the inhalation of fumes. Fever, nausea, vomiting, stomach cramps, and diarthea result from acute ingestions. Zinc from 675 to 2,280 ppm acts as an emetric. Inorganic zinc compounds relatively nontoxic (aside from irritant action). Acute oral toxicity in laboratory animal range from 250 mg/kg (LD in guines pig, for 210 01,190 mg/kg as rat oral LD for zinc nitrate hexahydrate. | Low oral toxicity. Prolonged ingention of zinc at 40 mg/l resulted in irrit- ability, muscular stiffness and pain, loss of appetite, and nausea. High levels of zinc in diet may retard growth and produce defective mineralization of bone. | | Essential nutrient. Taste threshold 15 ppm; 40 ppm soluble zinc salts impart a metallic taste. |
| Sources: Casarett and Doull's <u>Toxicol</u> c IARC Monographs, Vol. 3, 20, Drinking Water and Health, Vo Handbook of Toxic and Hazardo EPA Health Advisories for ino EXperimental and Clinical Neu Pesticides Studied in Man, M. 29 CFR 1910:50412-50499, Dece 40 CFR 1911:25720-25734, July | SY, 3rd edition, ed. C. D. Klassen, M. O. Amd. 23, International Agency for Research on Cance ol. J, National Academy of Sciences, 1977. USE Chemicals, M. Sittig, Noyes Publications, Internations, and pesticides, March, 198, Internations, organics, and pesticides, March, 198, irrotoxicology, ed. P. S. Spencer and H. H. Sch, J. Hayes, Hillians and Hilkins, Baltimore, I ember 10, 1985, OSHA, Occupational Exposure to 8, 1987, EPA, Erinking Hater. | ur, and J. Douli, Macmuilan Publishing Co., New er, Lyon, France, 1973, 1979, 1980. Park Ridge, New Jersey, 1981. 7. Baltimore, 1980. 982. Formaidehyde. | York 1986. | |
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| a Health oriect or target organ | I may be based on animal studies and does not i | umply that the results of exposure to humans wil | l be the same. | |

CVR193/029-6

Table 4: 19A CRITICAL TOXICITY VALUES^a (September 26, 1988)

| $\label{eq:constraints} \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | | | Noncar | cinogen Fact | ors | | Cancer Pote | ncy Factors | |
|---|----|----------------------------|---------------------------|---------------------------|--------------------|--------------------|--------------------------|-----------------------------------|------------------------------------|---|
| $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$ | | | Critical Intake | Critical Intake | 1-Day Child | 10-Day Child | Long-Term Child/Adult | HEA/CAG ^b Ingestion | HEA/CAG ^b Inhalation | |
| $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$ | | | Values for | Values for | Health | Health | Health | Slope | Slope | EPA |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Chemicals of Concern | Ingestion (mg/kg/day) | Inhalation (mg/kg/day) | Advisory (mg/l) | Advisory (mg/l) | Advisory (mg/l) | Factors -1 (mg/kg/day) -1 | Factors (mg/kg/day)-1 | Carcinogenic Classification ^C |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Acetone | 0.1 ^d | : | 1 | 1 | 1 | 1 | E B | |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Bartum | 0.05 | 0.0001 ^e | l i | ł | 1 | ſ | ł | ł |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Benzene | , ¦ | ; | 0.235 | 0.235 | : | 0.029 | 0.029 | A |
| $\begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Bis(2-ethylhexyl)phthalate | 0.02 | ł | 1 | ł | : | 0.014 | 1 | B2 |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | 2-Butanone | 0.05 ^d | ; | ! | ł | 1 | ł | 1 | { |
| $ \begin{array}{ccccccc} {\rm Chlordane} & 0.00005 & & 0.06 & 0.05 & 0.13 & 0.0001 \\ {\rm Dialorotorum} & 0.00005 & & 0.06 & 0.06 & 0.06 & 0.08 & 0.08 \\ {\rm Dialorotorum} & 0.00005 & & & & & 0.05 & 0.0061 & 0.081 & 0.25 \\ {\rm Endatin } & 0.00005 & & & & & & 0.06 & 0.081 & 0.25 \\ {\rm Endatin } & 0.0005 & & & & & & 0.01 & 0.0051 & & & & & & & & $ | | Cadmium | 0.0005 ^u | ł | ł | 1 | 1 | ł | 6.1 | BI |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Chlordane | 0.00005 | 1 | 0.06 | 0.06 | 0.05 | 1.3 | 1.3 | B2 |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Chloroform | 0.01 ^d | ł | : | 1 | ; | 0,0061 | 0.081 | B2 |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Dieldrin | 0.00005 | ł | ; | ł | ţ | 16 | 16 | B2 |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Endosulfan | 0,00005 ^d | : | ł | ł | 1 | ł | ł | ł |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Endrin | 0.0003 | 1 | 0.025 | 0.005 | 0.16 | ł | : | ł |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Ethylbenzene | 0.1 | ! | 32 | 3.2 | 1.0/3.4 | ! | 1 | ۵ |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | | Heptachlor | 0,0005 ^d | ; | 0.01 | 0.01 | 0.005/0.017 | 4.5 | 4.5 | B2 |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ |] | Lead | 0.0014 ^e | 0.00043 ^e | ł | ſ | ; | ł | : | { |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | ΕV | Mercury | 0.002 ^d | ł | ł | 1 | ! | ł | ł | ۵ |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | 7. | Methoxychlor | | ; | 6.4 | 2.0 | 0.5 | ! | { | ł |
| | 1 | Methylene chloride | 0.06 | ł | 13.3 | 1.5 | 0.5/1.75 | 0.0075 | 0.014 | B2 |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | 1: | Naphthalene | 0.4 | } | ł | ! | 1 | 1 | ł | ł |
| $ \begin{array}{cccccccccccccccccccccccccccccccccccc$ | 3 | Selentum | 0.003 | 0.001 ^e | ł | ł | 1 | : | 1 | ł |
| Toluene 0.3^{e} 1.0^{e} $$ | | Silver | 0.003 ^d | ; | f | ł | ł | ; | | 1 |
| Toxaphene 1.1 1.1 $B2$ Trichloroethene 2.0^d 0.4^e 0.011 0.000017 $B2$ Xylenes 2.0^d 0.4^e 0.011 0.000017 $B2$ Zinc 0.21^e 0.011 0.000017 $B2$ Zinc 0.21^e D | | Toluene | 0.3 ^e | 1.0 ^e | 1 | ł | ; | ł | : | ۵ |
| Trichloroethene $\sum_{2,0^d}^{-d}$ $\sum_{2,0^d}^{-e}$ <th< td=""><td></td><td>Toxaphene</td><td>;</td><td>;</td><td>1</td><td>!</td><td>ł</td><td>1.1</td><td>1.1</td><td>B2</td></th<> | | Toxaphene | ; | ; | 1 | ! | ł | 1.1 | 1.1 | B2 |
| Xylenes 2.0 ^d 0.4 ^e D Zinc 0.21 ^e | | Trichloroethene | ; | ; | ! | ł | i | 0.011 | 0.0000017 | B2 |
| Zinc 0.21 ^e | | Xylenes | 2.0 ^d | 0.4 ^e | 1 | 3 | ł | ! | ; | ۵ |
| | | Zinc | 0.21 ^e | ţ | ł | ł | { | ł | ł | ł |

Superfund Public Health Evaluation Manual: December 1986 (list revised July 1988) as amended by IRIS data base as of September 26, ^aSource: 1988.

^bThe cancer potency factors presented are either from Health Effects Assessments (HEA) or evaluations by the Carcinogen Assessment Group (CAG). All values presented were prepared by the USEPA.

^CU.S. EPA Carcinogen Assessment Group (CAG) Classification.

Group A Human carcinogen--Sufficient evidence from epidemiological studies. Group B1 Probable human carcinogen--At least limited evidence of carcinogenicity to humans. Group B2 Probable human carcinogen--Combination of sufficient evidence in animals and inadequate data in humans. Group C Possible human carcinogen--Limited evidence of carcinogenicity in animals in the absence of human data. Group D Not classified--Inadequate animal evidence of carcinogenicity. Group E No evidence of carcinogenicity.

RfD = verified Agency-wide value. dSource:

HEA = Health Effects Assessment. ^eSource:

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Table 4-20 PHYSICAL AND CHEMICAL PROPERTIES FOR ORGANIC CONTAMINANTS DETECTED AT KING SALMON AFS

| Acctone 67-64-1 56 1.006-06 2.706+02 2.066-05 2.2 0.24 Baraten 740-39-3 13 1.550-03 9.525+01 5.595-03 9.32 2.12 0.024 Baraten 740-37-3 11 2.350,00 9.525+01 5.595-03 9.32 2.12 0.23 Paratene 77-41-7 31 1.24 3.39 2.112 2.560 0.000400 9.525+01 5.595-03 9.32 2.12 0.33 2.113 0.32 2.113 0.32 2.113 0.31 3.169 3.13 2.112 2.560 0.0005+00 9.515-00 3.13 | Actone 67-64-1 58 1.00E+06 2.70E+02 Barium 740-39-3 137 1.755403 9.522+01 Barium 740-39-3 78 1.755403 9.522+01 Barium 71-43-2 78 1.755403 9.522+01 Barium 71-43-2 78 1.755+03 9.522+01 Batybenzyl phthalate 71-43-3 72 353,000 0.0005+00 Batybenzyl phthalate 740-39-3 112 2.292+00 0.0005+00 Batybenzen 65-66-3 112 8.206-01 1.78E-07 Copper 67-66-3 119 8.206+00 1.778E-07 Deldrin 1.552-01 1.78E-01 1.78E-07 Edidorin 740-57-1 381 1.95E-01 1.78E-07 Edidorin 67-66-3 119 8.20E+03 1.00E+00 Edidorin 740-57-1 381 1.95E-01 1.78E-07 Edidorin 100-41-4 106 1.52E+02 7.00E+04 Edidorin <td< th=""><th>Pressure Constant Koc (mm Hd) (atm-m³/mol) (ml/d)</th><th>Fish BC Log Kow (1/kg)</th></td<> | Pressure Constant Koc (mm Hd) (atm-m ³ /mol) (ml/d) | Fish BC Log Kow (1/kg) |
|--|---|---|---------------------------|
| Action Barren Action Settom Action Barren $0.764-1$ 59 $1.000+66$ $2.706-05$ 2.23 0.23 2.23 0.23 0.23 0.23 2 | Acetone 6/-04-1 58 1.00E+06 2.70E+02 Bartum 71-32-3 78 1.75E+03 9.52E+01 Bartum 71-32-3 78 1.75E+03 9.52E+01 Bartum 71-32-3 78 1.75E+03 9.52E+01 Bartum 710-33-3 73 1.75E+03 9.52E+01 Datylbenzyl phthalate 740-43-9 112 2.95E+00 0.00E+00 Butylbenzyl phthalate 740-43-9 112 2.95E+00 0.00E+00 Chlorrdare 740-43-9 113 8.20E+03 1.51E+02 Chlorrdare 66-5-3 119 8.20E+03 1.51E+02 Copper 0 9.55E+01 1.78E-01 1.78E-01 Didrin Edition 60-5+1 381 1.95E-01 1.78E-01 Didrin Edition 60-5+1 1.152E+02 1.00E+00 0.00E+00 Didrin Edition 60-5+1 1.300E+01 1.78E-01 1.78E-01 Didrin Edition 60-5+1 < | | |
| Description $71.75-73$ 73 $1.75-73$ 3.95 0.4 $5.59E-03$ 83 2.112 $2.59E-03$ 83 2.112 $2.59E-03$ 1.3 3.33 2.112 $2.59E-03$ 1.3 3.32 2.112 $2.59E-03$ 1.31 3.32 2.112 $2.59E-03$ 1.37 3.32 $2.59E-03$ 1.37 3.32 $2.59E-03$ 1.370 3.32 2.12 $2.59E-03$ 1.370 3.32 2.12 $2.59E-03$ 1.370 3.32 2.12 $2.59E-03$ 1.700 3.25 $2.59E-03$ 1.700 3.25 $2.59E-03$ 1.700 3.25 2.212 2.212 2.212 2.212 2.212 2.212 2.212 2.212 2.212 2.212 2.212 | Matrix Ti-3-2 bit (1-bit) Ti-3-2 (1-bit) Ti-3-2 (1-b | 2.70E+02 2.06E-05 2.2 | -0.24 |
| Reference $1,7,2,2,2$ $3,0$ $1,7,2,2,0$ $3,2,2,0,0$ $1,3,4,0,2$ $3,1,0,0,0,0$ $3,1,2,0,0,0,0$ $3,1,2,0,0,0,0,0$ $3,1,2,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0,0$ | Definition 0.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1 | | |
| This is a second and the second an | Distribution 2.9 E-00 0.00E+00 Distribution 85-68-7 72 35,000 Distribution 85-68-7 72 2.9 E+00 0.00E+00 Chloroform 67-66-3 112 2.9 E+00 0.00E+00 Chloroform 67-66-3 119 8.20E+03 1.51E+02 Chloroform 67-66-3 119 8.20E+03 0.00E+00 Chloroform 67-57-1 301 1.95E-01 1.78E-07 Diseldin 115-29-7 407 1.55E-01 1.78E-07 Diseldin 115-29-7 301 1.95E-01 1.78E-07 Diseldin 11 115-29-7 407 1.65E-01 1.78E-07 Diseldin 11 115-29-7 301 1.95E-01 1.78E-07 Diseldin 11 115-29-7 301 1.95E-01 1.78E-07 Diseldin 11 11 1.55E+02 7.00E+04 3.00E+00 End 11 1.66 1.55E+02 7.00E+04 3.65E+02 Postencie 7439-97-6 201 1.95E+03 3.62E+02 | 9.02E+UI 0.09E+U3 03 | °C 71°7 |
| Transmort Transmort <thtransmort< th=""> <thtransmort< th=""> <th< td=""><td>Further 91.2010 0.006+00 0.006+00 Survision 67-65-3 112 2.95+00 0.006+00 Cadmium 740-43-9 112 2.95-00 0.006+00 Cadmium 67-67-3 113 5.60E 1.51E+02 Cadmiun 67-67-3 113 5.60E 1.51E+02 Copper 60-57-1 381 1.95E-01 1.78E-07 Diedarin 67-57-1 381 1.95E-01 1.78E-07 Diedarin 66-57-1 381 1.95E-01 1.78E-07 Diedarin 100-41-4 106 1.52E+02 0.006+00 Ethylbenzene 86-73-7 374 1.80E-01 3.00E-04 Mangares 76-44-8 374 1.80E-01 3.00E-04 Flowrene 86-73-7 166 1.55E+02 0.00E+00 Mangares 7439-95-5 55 2.00E+00 3.00E-04 Mangares 7439-97-6 201 1.75E+01 3.65E+02 Mangares 7439-97-6 201 1.75E+01 3.65E+02 Mangares 7440-02-3</td><td>123,027</td><td></td></th<></thtransmort<></thtransmort<> | Further 91.2010 0.006+00 0.006+00 Survision 67-65-3 112 2.95+00 0.006+00 Cadmium 740-43-9 112 2.95-00 0.006+00 Cadmium 67-67-3 113 5.60E 1.51E+02 Cadmiun 67-67-3 113 5.60E 1.51E+02 Copper 60-57-1 381 1.95E-01 1.78E-07 Diedarin 67-57-1 381 1.95E-01 1.78E-07 Diedarin 66-57-1 381 1.95E-01 1.78E-07 Diedarin 100-41-4 106 1.52E+02 0.006+00 Ethylbenzene 86-73-7 374 1.80E-01 3.00E-04 Mangares 76-44-8 374 1.80E-01 3.00E-04 Flowrene 86-73-7 166 1.55E+02 0.00E+00 Mangares 7439-95-5 55 2.00E+00 3.00E-04 Mangares 7439-97-6 201 1.75E+01 3.65E+02 Mangares 7439-97-6 201 1.75E+01 3.65E+02 Mangares 7440-02-3 | 123,027 | |
| Condition $770-43-6$ 112 $5.60E$ $0.000+00$ M $140,000$ 3.50 Condition $67-66-3$ 119 $8.206+00$ M $140,000$ 3.50 Condentie $67-66-3$ 119 $8.206+00$ $1.51E+02$ $2.48E-07$ 1.700 3.50 Copert $66-57-1$ 301 $1.95E-01$ $1.51E+02$ $2.48E-07$ 1.700 3.50 Dedosulfan and II $1.78E-07$ $4.58E-07$ 1.700 3.50 Dedosulfan and II $1.95E-01$ $1.95E-01$ 1.7700 3.50 Dedosulfan and II $1.00-41-4$ 106 $1.51E+02$ $2.075-03$ 1.7100 3.10 Dedosulfan and II $1.00-41-4$ 106 $1.58E-01$ 1.700 3.10 Dedosulfan $8e^{-73-7}$ $1.00-700$ $3.00E-04$ $8.19E-07$ 1.700 3.10 Dedosulfan $8e^{-73-7}$ $1.80E-01$ $3.00E-04$ $8.19E-04$ 1.400 | Cadmium $7740-43-9$ 112 $5.60E$ 0.00E+00 Chloroform $67-65-3$ 119 $5.60E$ 1.51E+02 Chloroform $67-65-3$ 119 $8.20E+03$ 1.51E+02 Coportin $67-65-3$ 119 $8.20E+03$ 1.51E+02 Coper Coper $67-57-1$ 381 1.95E-01 1.78E-07 Dieldrin Endylbenzee $67-57-1$ 381 1.95E-01 1.78E-07 Endosulfan I and II $115-29-7$ 407 $1.52E+02$ $0.00E+00$ Endylbenzee $86-73-7$ 166 $1.55E+02$ $1.78E-07$ $0.00E+00$ Fullylbenzee $86-73-7$ 166 $1.50E-01$ $1.78E-07$ $0.00E+00$ Fullylbenzee $7439-92-6$ $7439-92-6$ 207 $1.80E-01$ $0.00E+00$ Manganese $7439-97-6$ 201 $1.77E+01$ $0.00E+00$ $7.10E-04$ Metroyresie $7439-97-6$ 201 $1.77E+01$ $0.00E+00$ $2.00E+01$ Metroyresis $7439-97-6$ 201 $1.77E+01$ $2.00E+01$ | | 99 |
| $ \begin{array}{c} \mbox{Choreane} & \begin{tabular}{ c c c c c c c c c c c c c c c c c c c$ | Chlordane Chlordane5.60E 5.60E5.60E 5.60EChloroform ChloroformChloroform Chloroform $5.60E$ 6100000116an $1.51E+02$ 0.0000000Copper Copper Copper DieldrinCopper 0.0000000 $67-6-3$ 61000000 119 61000000 $8.20E+03$ 0.0000000 $1.51E+02$ 0.0000000Dieldrin Endosulfan $115-29-7$ 1000000 407 6100000 $1.95E-01$ 10000000 $1.78E-07$ 10000000Ethylbenzene Ethylbenzene Flourene Heptachlor $100-41-4$ 766-44 7439-97-6 7439-97-6 7439-97-6 7439-97-6 7439-97-6 7439-97-6 $1.006-40$ 7.006-00 7.006-00 7.006-00FI Definition Nethnylene Nethnylene Nickel Silver $1.735E+03$ 7.00E-01 7.00E-01 7.00E-02 $1.735E+02$ 7.00E-03FI Demogravior Nethnylene Nickel Silver $1.736+03$ 7.00E-00 $2.00E+00$ 7.00E-00Nickel Silver $91-20-3$ 7.00E-01 81100 $1.740-22-4$ 7.00E+00 $0.00E+00$ 7.00E+00Nickel Silver $91-20-3$ 860-04 81100 $1.740-22-4$ 7.00E+00 $0.00E+00$ 7.00E+00Nickel Silver $91-20-3$ 860-04 81100 $1.740-22-4$ 7.00E+00 $0.00E+00$ 7.24402Nickel Silver $91-20-3$ 860 81100 $1.740-22-4$ 92 7.2000000 $0.00E+00$ 7.24402 <td< td=""><td>0.00E+00 NA</td><td>, œ</td></td<> | 0.00E+00 NA | , œ |
| $ \begin{array}{c} \mbox{Chlorotorm} & \mbox{6-3} & \mbox{119} & \mbox{8.20E+03} & \mbox{1.51E+02} & \mbox{2.87E-03} & \mbox{3.1} & \mbox{1.97} & \mbox{3.50} & \mbox{3.1} & \mbox{3.50} & \mbox{3.51} & \mbox{3.50} & \mbox{3.51} & \mbox{3.51} & \mbox{3.51} & \mbox{3.52} & \mbox{3.51} & \mbox{3.51} & \mbox{3.52} & \mbox{3.51} & \mbox{3.52} & \mbox{3.51} & \mbox{3.52} & 3.$ | Chloroform 67-66-3 119 8.20E+03 1.51E+02 Copper Copper 64 1.95E-01 1.78E-07 Dieldrin Dieldrin 115-29-7 407 0.00E+00 0.00E+00 Endosulfan I 115-29-7 407 1.78E-07 1.78E-07 Endosulfan I 115-29-7 407 1.55E+01 1.78E-07 Endosulfan I 115-29-7 407 407 407 Endosulfan I 115-29-7 407 407 407 Endosulfan I 100-41-4 106 1.58E+00 7.10E-04 Endosulfan 86-73-7 166 1.58E+00 7.10E-04 Heptachlor 76-44-8 374 1.80E-01 3.00E-00 Flourene 86-73-7 166 1.58E+00 7.10E-04 Manganese 7439-97-6 50 1.80E-01 3.00E-01 Mectury (inorganic) 7439-97-6 201 1.75E+03 3.62E+02 2.00E+00 Methylane 7439-97-6 20 1.77E+01 3.00E-01 3.62E+02 | 9.63E-06 140,000 | 3.32 14,00 |
| Copper Decontant740-50-8 Decontant64 $115-29-7$ 0.00E+00 $115-29-7$ M.M.Disclation Decontant100-41-41061.52E+027.00E+006.43E-031.7003.15Endosulfan I Decontant111-29-74011.52E+027.00E+006.43E-031.71003.15Endosulfan I Decontant111-29-74061.52E+027.00E+006.43E-031.71003.15Endosulfan I Decontant111-29-71661.52E+027.00E+006.43E-031.71003.15Endosulfan I Endosuce75-44-83741.80E-010.00E+006.43E-057.3004.40Amagarese Magarese75-44-83741.80E-010.00E+008.19E-0412.0004.40Amagarese Magarese75-99-62011.75E+022.00E+048.19E-0412.0004.40Amagarese Magarese75-09-2852.00E+041.7004.464.40Amagarese Methylene75-09-2852.00E+041.7604.464.46Amagarese Methylene75-09-2852.00E+041.7769623.30Amagarese Methylene75-09-2921.75E+022.00E+041.4004.46Amagarese Methylene75-09-2852.00E+041.9623.302.73Amagarese Amphthalene740-22-31283.17E+010.00E+041.464.46Amagarese Maphthalene7440-22-31283.17E+014 | Copper 64 0.00E+00 Dieldrin Dieldrin 381 1.95E-01 1.78E-07 Dieldrin Indosulfan I and II 60-57-1 381 1.95E-01 1.78E-07 Didosulfan I and II Didosulfan I and II 60-57-1 381 1.95E-01 1.78E-07 Endosulfan I and II Didosulfan I and II 100-41-4 106 1.52E+02 7.00E+00 Endosulfan I and II Diotene 86-73-7 166 1.52E+02 7.00E+00 Endosulfan Noutrene 86-74-8 374 1.80E-01 3.00E-04 Manganese 7439-97-6 55 200 1.60E+00 3.00E-04 Methylene 7439-97-6 201 1.75E+03 3.62E+02 3.62E+02 Methylene 7439-97-6 201 1.75E+03 3.62E+02 3.62E+02 Methylene 7439-97-6 201 1.75E+03 3.62E+02 3.62E+02 Methylene 7.00E+00 7439-97-6 201 1.75E+03 3.62E+02 Methylene | 1.51E+02 2.87E-03 31 | 1.97 3.7 |
| Dieldrin Dieldrin (6^{-57-1}) 381 $1.95E-01$ $1.78E-07$ $4.58E-07$ $1,700$ 3.50 Endosulfan Endosulfan 1 and 11 $100-41-4$ 106 $1.52E+02$ $7.00E+00$ $6.43E-03$ $11,100$ 3.15 Endosulfan Endosulfan $100-41-4$ 106 $1.52E+02$ $7.00E+00$ $6.43E-03$ $11,100$ 3.15 Endosulfan Endosulfan $100-41-4$ 106 $1.52E+02$ $7.00E+00$ $6.43E-03$ $7,300$ 4.20 Flourene Endosuch $86-73-7$ 166 $1.55E+02$ $7.10E-04$ $6.43E-03$ 1.100 4.20 Reptachlor $7.399-92-15$ 50 $0.00E+00$ $6.43E-03$ 1.100 4.20 A Manganese $739-97-6$ 50 $0.00E+00$ $8.19E-04$ 12.000 4.40 A Manganese $7339-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.8 8.3 2.12 Marganese $7-09-2$ 85 $2.00E+04$ $3.62E+02$ $2.00E+03$ 8.8 1.30 Marcury (increate Metholace $7-09-2$ 8.8 $3.17E+01$ $0.00E+00$ 4.40 Mathalane $7-09-2$ 8.8 $3.17E+01$ $0.00E+00$ $4.40-22-4$ 9.2 Mathalane $91-20-3$ $12,000+00$ $4.302-03$ 8.8 1.300 2.73 Mathalane $91-20-3$ $12,000+00$ $6.30E+00$ $1.402-02-03$ 8.6 $1.400-00$ Mathalane $91-20-3$ $12,000-00$ $4.02E-03$ 8.6 9.2 <td>Dieldrin 60-57-1 381 1.95E-01 1.78E-07 Endosulfan I and II Endosulfan I and II Endosulfan I and II 100-41-4 106 1.52E+02 7.00E+00 Endosulfan I and II Endosulfan I and II Endosulfan I and II 100-41-4 106 1.552E+02 7.00E+00 Endosulfan I and II Endosulfan I and II Endosulfan I and II 100-41-4 106 1.552E+02 7.00E+00 Flourene 86-73-7 166 1.552E+02 7.10E-04 Flourene 86-73-7 166 1.552E+02 7.10E-04 Amaganese 76-44-8 374 1.80E-01 3.00E-04 Amaganese 7439-97-6 201 1.75E+03 2.00E+04 Mertury (inorganic) 7439-97-6 201 1.75E+03 3.6.2E+02 Methoration 7439-97-6 201 1.75E+03 3.6.2E+02 3.6.2E+02 Methoration 7439-97-6 201 1.75E+03 3.6.2E+02 3.6.2E+02 Methoration 7440-02-0 128 3.17E+01 0</td> <td>0.00E+00 NA</td> <td>20</td> | Dieldrin 60-57-1 381 1.95E-01 1.78E-07 Endosulfan I and II Endosulfan I and II Endosulfan I and II 100-41-4 106 1.52E+02 7.00E+00 Endosulfan I and II Endosulfan I and II Endosulfan I and II 100-41-4 106 1.552E+02 7.00E+00 Endosulfan I and II Endosulfan I and II Endosulfan I and II 100-41-4 106 1.552E+02 7.00E+00 Flourene 86-73-7 166 1.552E+02 7.10E-04 Flourene 86-73-7 166 1.552E+02 7.10E-04 Amaganese 76-44-8 374 1.80E-01 3.00E-04 Amaganese 7439-97-6 201 1.75E+03 2.00E+04 Mertury (inorganic) 7439-97-6 201 1.75E+03 3.6.2E+02 Methoration 7439-97-6 201 1.75E+03 3.6.2E+02 3.6.2E+02 Methoration 7439-97-6 201 1.75E+03 3.6.2E+02 3.6.2E+02 Methoration 7440-02-0 128 3.17E+01 0 | 0.00E+00 NA | 20 |
| Endosulfan115-29-7407Findosulfan I and IIEndosulfan I and IIEthylbenzene $100-41-4$ Ethylbenzene $100-41-4$ Manganes $7439-97-6$ Manganes $733-97-6$ Mercury (Inorganic) $7439-97-6$ Methylane $7-100-2$ Methylane $91-20-3$ Nambhhalene $91-20-3$ Name $1000-00$ Nambhhalene $91-20-3$ | Endosulfan 115-29-7 407 Endosulfan I and II Endosulfan I and II 115-29-7 407 Endosulfan I and II Endosulfan I and II 100-41-4 106 1.52E+02 7.00E+00 Ethylbenzene 86-73-7 166 1.69E+00 7.10E-04 Flourene 86-73-7 166 1.69E+00 7.10E-04 Reptachlor 76-44-8 374 1.80E-01 3.00E-04 Amaganese 7439-97-5 55 7.00E+00 0.00E+00 Amaganese 7439-97-6 201 1.75E+03 3.62E+02 Mercury (inorganic) 7439-97-6 201 1.75E+03 3.62E+02 Methylene 7430-92-0 85 2.00E+04 3.62E+02 Nethylene 740-02-0 85 2.00E+04 3.62E+02 N | 1.78E-07 4.58E-07 1,700 | 3.50 4,76 |
| Endosulfan I and IIEndosulfan I and IIEndosulfan I and IIEndosulfan I and IIEthylbenzene $100-41-4$ Ethylbenzene $1.52E+02$ Tourene $7-30-44-8$ Flourene $7-30-44-8$ Reparence $7-30-2-44-8$ Reparence $7-30-2-4-8$ Manganese $7-30-2-4-8$ Manganese $7-30-2-4-8$ Manganese $7-30-2-4-8$ Manganese $7-30-2-4-8$ Manganese $7-30-2-6-5-5$ Manganese $7-30-2-6-5-5-5-5-5-5-5-5-5-5-5-5-5-5-5-5-5-5$ | Endosulfan I and II Endosulfan I and II Endrin Endrin Flouree 86-73-7 Flouree 86-73-7 Flouree 86-73-7 Flouree 86-73-7 Flouree 7439-92-1 Amanganese 7439-97-6 Manganese 7439-97-6 Manganese 7439-97-6 Mercury (inorganic) 7439-97-6 Methylene Chloride 7-5-09-2 Bethoxychior 7-50-2 Second 3.17E+01 Mapthalene 91-20-3 Naphthalene 91-20-3 Naphthalene 91-20-3 Naphthalene 7440-22-4 Naphthalene 7440-22-4 Nockel 7440-22-4 Silver 7440-22-4 Silver 108 Toluene 540-59-0 | | |
| Endrin Endrin Endrin FloureneEndrin Endrin Flourene100-41-4 (101-41-4)106 1.52E+02 $1.52E+02$ (100-41-4) $1.00E+04$ (1.66E+00) 1.100 (1.00E-04) 1.100 (1.100) 1.100 (1.100)< | Endrin EndrinEndrin Ethylbenzene100-41-4106 $1.52E+02$ $7.00E+00$ Fluorene Flourene $86-73-7$ 166 $1.52E+02$ $7.00E+00$ Fluorene Flourene $86-73-7$ 166 $1.55E+02$ $7.00E+00$ A Manganese Floar $7439-92-1$ 207 $1.80E-01$ $3.00E-04$ A Manganese Manganese $7439-97-6$ 55 55 $0.00E+00$ A Manganese Manganese $7439-97-6$ 201 $1.78E+03$ $2.00E+04$ A Manganese $7439-97-6$ 201 $1.75E+03$ $2.00E+04$ Metroxy (inorganic) $7439-97-6$ 201 $1.75E+03$ $3.62E+02$ Methylene Chloride $75-09-2$ 85 $2.00E+04$ $3.62E+02$ Naphthalene $740-22-3$ 128 $3.17E+01$ $0.00E+00$ Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+00$ Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+00$ Naphthalene $7440-22-4$ 108 92 $5.35E+02$ $2.81E+01$ Naphthalene $7440-22-4$ 108 92 $5.35E+02$ $2.81E+01$ Toluene $740-22-4$ 108 92 $5.35E+02$ $2.81E+01$ Towene $79-01-6$ 178 $1.00E+03$ $3.28+02$ $2.78E+02$ Trans 1, 2-dichloroethene $59-02-0$ 97 $6.00E+02$ $3.28+02$ Trans 1, 2-dichloroethene $79-01-6$ 131 $1.00E+03$ $3.78+02$ Trans 1, 2-dichloroethene $79-01-6$ 131 < | | 1,00 |
| Ethylbenzene $100-41-4$ 106 $1.52E+02$ $7.00E+00$ $6.43E-03$ 1.100 3.115 ToureneHeptachlor $7-10E-04$ $6.42E-05$ $7,7300$ 4.40 Heptachlor $76-44=8$ 374 $1.80E-01$ $3.00E+00$ $6.43E-05$ $7,7300$ 4.40 Andamese $76-44=8$ 374 $1.80E-01$ $3.00E+00$ $6.43E-05$ $7,7300$ 4.40 Anganese $7439-97-6$ 55 $0.00E+00$ $6.43E-05$ $7,7300$ 4.40 Amganese $7439-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.8 1.300 Amganese $7439-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.8 1.300 Amganese $7439-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.8 1.300 Methylane $75-09-2$ 85 $2.00E+04$ $3.62E+02$ $2.03E-03$ 8.8 1.300 Methylane $91-20-3$ 128 $3.17E+01$ $0.00E+00$ $6.30E-04$ 962 3.30 Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+00$ 4.46 962 3.30 NickelNickel $1.440-02-0$ $5.35E+02$ $2.03E-04$ 962 3.30 NickelNickel $1.440-02-0$ $5.35E+02$ $2.01E+01$ $6.37E-03$ 3.00 2.73 NickelNickel $1.740-02-0$ $5.35E+02$ $2.81E+01$ 962 3.30 NickelNickel $1.240-59-0$ 97 $6.00E+02$ $2.81E+01$ $6.37E-03$ <td< td=""><td>Ethylbenzene 100-41-4 106 1.52E+02 7.00E+00 Flourene 86-73-7 166 1.52E+02 7.00E+00 Flourene 86-73-7 166 1.52E+02 7.00E+00 A Manganese 7439-92-1 207 0.00E+00 7.10E-04 A Manganese 7439-92-1 207 0.00E+00 7.10E-04 Mendpanese 7439-97-6 50 7.39-97-6 50 0.00E+00 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 7.00E-04 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 7.00E-04 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 7.00E-04 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 3.62E+02 Methoxychlor 7440-22-9 85 2.00E+04 3.62E+02 3.62E+02 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 6.80E-04 Nickel 7440-22-4 108 178 1.00E+01 0.00E+00 Nickel 74</td><td></td><td>13,00</td></td<> | Ethylbenzene 100-41-4 106 1.52E+02 7.00E+00 Flourene 86-73-7 166 1.52E+02 7.00E+00 Flourene 86-73-7 166 1.52E+02 7.00E+00 A Manganese 7439-92-1 207 0.00E+00 7.10E-04 A Manganese 7439-92-1 207 0.00E+00 7.10E-04 Mendpanese 7439-97-6 50 7.39-97-6 50 0.00E+00 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 7.00E-04 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 7.00E-04 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 7.00E-04 Methoxychlor 7439-97-6 201 1.75E+03 3.62E+02 3.62E+02 Methoxychlor 7440-22-9 85 2.00E+04 3.62E+02 3.62E+02 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 6.80E-04 Nickel 7440-22-4 108 178 1.00E+01 0.00E+00 Nickel 74 | | 13,00 |
| Flourene $86-73^{-7}$ 166 $1.69E+00$ $7.10E-04$ $6.42E-05$ $7,300$ 4.20 AHeptachlor $739-92-1$ 207 $0.00E+04$ $8.19E-04$ $12,000$ 4.40 AManganese $7439-92-1$ 507 $0.00E+04$ $8.19E-04$ $12,000$ 4.40 AManganese $7439-97-6$ 507 $0.00E+04$ $8.19E-04$ $12,000$ 4.40 Hercury (inorganic) $7439-97-6$ 501 $1.75E+03$ $3.00E-04$ $8.19E-04$ $12,000$ 4.40 Hercury (inorganic) $7439-97-6$ 201 $1.75E+03$ $3.00E-04$ 8.18 1.302 Methoxychlor $7439-97-6$ 201 $1.75E+03$ $3.62E+02$ $2.03E-03$ 8.8 1.302 Methoxychlor $75-09-2$ 85 $2.00E+04$ $3.529E-03$ 8.8 1.302 Methoxychlor $740-02-0$ $12,702$ $3.17E+01$ $0.00E+00$ 4.46 Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+00$ $4.22E-04$ 962 3.30 Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+00$ 4.46 4.6 Nickel 8.8 $1.00E+00$ $0.00E+00$ 4.46 4.6 $5.559E-03$ 8.8 1.40 Naphthalene $91-20-3$ 128 $1.00E+00$ $4.22E-04$ 962 3.30 2.73 Nickel 8.8 $1.00E+00$ $0.00E+00$ $0.00E+00$ 4.46 3.6 2.73 Nickel 8.8 $1.00E+00$ <t< td=""><td>Flourene 86-73-7 166 1.69E+00 7.10E-04 Heptachlor 76-44-8 374 1.80E-01 3.00E-04 Manganese 76-44-8 374 1.80E-01 3.00E-04 Manganese 7439-92-1 207 0.00E+00 3.00E-04 Manganese 7439-97-6 55 55 0.00E+00 Mercury (inorganic) 7439-97-6 201 1.75E+03 3.00E-04 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Nethylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Nickal 7440-02-0 59 1.00E+04 0.00E+00 Nickal 7440-02-0 59 0.00E+00 0.00E+00 Nickal 7440-22-4 108 1.78 0.00E+00 Nickal 7440-22-4 108 0.00E+00 0.00E+00 Silver 7440-22-4 108 5.35E+02 2.81E+01 <t< td=""><td>7.00E+00 6.43E-03 1,100</td><td>3.15 37.</td></t<></td></t<> | Flourene 86-73-7 166 1.69E+00 7.10E-04 Heptachlor 76-44-8 374 1.80E-01 3.00E-04 Manganese 76-44-8 374 1.80E-01 3.00E-04 Manganese 7439-92-1 207 0.00E+00 3.00E-04 Manganese 7439-97-6 55 55 0.00E+00 Mercury (inorganic) 7439-97-6 201 1.75E+03 3.00E-04 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Nethylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Nickal 7440-02-0 59 1.00E+04 0.00E+00 Nickal 7440-02-0 59 0.00E+00 0.00E+00 Nickal 7440-22-4 108 1.78 0.00E+00 Nickal 7440-22-4 108 0.00E+00 0.00E+00 Silver 7440-22-4 108 5.35E+02 2.81E+01 <t< td=""><td>7.00E+00 6.43E-03 1,100</td><td>3.15 37.</td></t<> | 7.00E+00 6.43E-03 1,100 | 3.15 37. |
| Heptachlor $76-44-8$ 374 $1.80E-01$ $3.00E-04$ $8.19E-04$ $12,000$ 4.40 AHeptachlor $739-92-1$ 207 $0.00E+00$ NA NA NA AManganese $739-97-5$ 55 55 NA NA AManganese $739-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.3 2.12 AMethoxychior $7439-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.3 8.3 2.12 AMethoxychior $7439-97-6$ 201 $1.75E+03$ $2.00E-03$ 8.3 1.30 AMethoxychior $7439-97-6$ 201 $1.75E+03$ $5.55E-03$ 8.3 1.30 AMethoxychior $7430-02-0$ 85 $2.00E+04$ $3.62E+02$ 3.30 3.30 Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+04$ $1.60E+04$ 1.46 Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+04$ 1.46 3.30 Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+04$ 1.600 4.46 Nickel $91-20-3$ 128 $1.00E+00$ $1.82E-03$ 8.8 1.40 Nickel $91-20-3$ 128 $1.00E+00$ $1.82E-03$ 8.8 1.40 Naphthalene $91-20-3$ 128 $0.00E+00$ $1.82E-03$ 8.8 1.40 Nickel $100E+00$ $1.00E+00$ $1.82E-03$ 8.8 1.40 $1.82E-03$ 8.8 Nickel $108-$ | Heptachlor $76-44-8$ 374 $1.80E-01$ $3.00E-04$ I. Lead $7439-92-1$ 207 55 55 $0.00E+00$ Manganese $7439-97-6$ 55 55 $0.00E+00$ Mercury (alkyl) $7439-97-6$ 201 $1.75E+03$ $0.00E-03$ Methoxychlor $7439-97-6$ 201 $1.75E+03$ $2.00E+04$ Methoxychlor $7439-97-6$ 201 $1.75E+03$ $2.00E+04$ Methylene Chloride $75-09-2$ 85 $2.00E+04$ $3.62E+02$ Nethylene Chloride $7-00-20-3$ 128 $3.17E+01$ $0.00E+00$ Naphthalene $91-20-3$ 128 $3.17E+01$ $0.00E+00$ Nickel $91-20-3$ 178 $1008+00$ $0.00E+00$ Phenanthrene $91-20-3$ 92 $5.35E+02$ $2.81E+01$ Toluene $7440-22-4$ 108 92 $5.35E+02$ $2.81E+01$ Toluene $740-50-0$ 97 $6.00E+01$ $3.264+02$ Trans 1,2-dichloroethene $540-59-0$ 97 $6.00E+02$ $3.724+02$ Trichloroethene $79-01-6$ 131 $1.09E+02$ $1.00E+01$ Niese $1330-20-7$ 106 131 $1.00E+02$ | 7.10E-04 6.42E-05 7,300 | 4.20 1,30 |
| I Lead 7439-92-1 207 0.00E+00 NA A Manganese 7439-97-6 55 NA NA Hercury (alky1) 7439-97-6 50 NA NA A Manganese 7439-97-6 50 NA NA Mercury (alky1) 7439-97-6 201 1.75E+03 83 2.12 Methoxychlor 7439-97-6 201 1.75E+03 5.59E-03 83 2.12 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 3.30 8.8 1.30 2-methylnapthalene 91-20-3 128 3.17E+01 4.82E-04 962 3.30 Naphhalene 7440-22-4 178 1.00E+00 A 4.82E-04 962 3.30 Naphhalene 7440-22-4 178 1.00E+00 A 4.82E-04 962 3.30 Naphhalene 700E+01 1.00E+00 1.55E-04 14,000 4.46 Nickel 85 1.00E+00 6.86E-04 1.48 | I Lead 7439-92-1 207 0.00E+00 A manganese 7439-97-6 55 55 55 0.00E+00 T Manganese 7439-97-6 55 55 55 0.00E+00 T Manganese 7439-97-6 201 1.75E+03 2.00E-03 T Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Aethylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Ampthalene 75-09-2 85 2.00E+04 3.62E+02 Naphthalene 7440-02-0 59 1.00E+04 3.62E+02 Naphthalene 7440-02-0 59 1.00E+00 6.80E-04 Naphthalene 7440-22-4 108 5.35E+02 2.81E+01 Silver 7440-22-4 108 5.35E+02 2.81E+01 Toluene 7440-22-4 108 5.35E+02 2.81E+01 Toluene 7440-22-4 108 5.35E+02 2.81E+01 Towerphene 7440-22-0 | 3.00E-04 8.19E-04 12,000 | 4.40 15,70 |
| C Manganese 7439-96-5 55 NA manganese 7439-97-6 55 NA manganese 7439-97-6 201 1.75 F+03 2.00 F-03 NA | C Manganese 7439-96-5 55 H Mercury (alkyl) 7439-97-6 201 1.75E+03 2.00E-03 H Mercury (inorganic) 7439-97-6 201 1.75E+03 2.00E-03 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 Naphthalene 740-20-3 128 3.17E+01 0.00E+00 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 Nickel 7440-22-4 108 1.00E+00 6.80E-04 Phenanthrene 7440-22-4 108 0.00E+00 0.00E+00 Silver 70luene 108-88-3 92 5.00E-01 3.2E4+02 Toxaphene 730-20-0 97 6.00E+02 3.2E4+02 3.2E4+02 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3.2E4+02 3.2E4+02 Xylenes 1330-20-7 106 1.98E+02 3.2E4+02 3.2E4+02 | 0.00E+00 NA | 4 |
| $ \begin{array}{llllllllllllllllllllllllllllllllllll$ | H Mercury (alkyl) 7439-97-6 201 2.00E-03 H Methoration 7439-97-6 201 2.00E-03 Methoxychlor 7439-97-6 201 1.75E+03 2.00E-03 Methylape Chloride 75-09-2 85 2.00E+04 3.62E+02 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 Nickel 91-20-3 128 3.17E+01 0.00E+00 Nickel 7440-02-0 59 1.00E+00 6.80E-04 Nickel 7440-22-4 108 1.00E+00 6.80E-04 Silver 7440-22-4 108 0.00E+00 0.00E+00 Phenanthrene 7440-22-4 108 0.255560 2.81E+01 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toluene 79-01-6 131 1.00E+02 3.2E4+02 Trichloroethene 79-01-6 131 1.98E+02 3.7E4+02 | NA | |
| ItMercury (inorganic)7439-97-62012.00E-03NA \mathbf{W} methoxychlor $739-97-6$ 201 $1.75E+03$ $5.59E-03$ 83 2.12 \mathbf{W} methoxychlor $75-09-2$ 85 $2.00E+04$ $3.62E+02$ $2.03E-03$ 8.3 1.30 \mathbf{W} methor $75-09-2$ 85 $2.00E+04$ $3.62E+02$ $2.03E-03$ 8.3 1.30 \mathbf{W} methor $791-20-3$ 128 $3.17E+01$ $4.82E-04$ 962 3.30 \mathbf{N} mapthalene $7440-02-0$ 59 $1.00E+00$ $4.82E-04$ 962 3.30 \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} nuckel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} | Instruct Marcury (inorganic) 7439-97-6 201 2.00E-03 A methoxychlor Methoride 75-09-2 85 2.00E+04 3.62E+02 A methylnapthalene 75-09-2 85 2.00E+04 3.62E+02 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 Nickel 7440-02-0 59 1.00E+00 6.80E-04 Nickel 7440-22-4 108 0.00E+00 6.80E-04 Phenanthrene 7440-22-4 108 0.00E+00 0.00E+00 Silver 108 89-3 92 5.35E+02 2.81E+01 Toluene 108 92 5.35E+02 2.81E+01 Tosubne 108-88-3 92 5.35E+02 2.81E+01 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3.2E4+02 Trichloroethene 79-01-6 131 1.09E+02 1.00E+01 1.00E+01 | NA | 3 , 75 |
| \mathbf{F} Methoxychlor $\mathbf{S}_{2.59E-03}$ $\mathbf{S}_{3.3}$ $\mathbf{S}_{3.12}$ $\mathbf{S}_{3.59E-03}$ $\mathbf{S}_{3.3}$ $\mathbf{S}_{3.13}$ \mathbf{M} methylene Chloride $75-09-2$ \mathbf{S}_{5} $\mathbf{S}_{2.00E+04}$ $3_{6.5E+02}$ $\mathbf{S}_{2.03E-03}$ $\mathbf{S}_{3.3}$ $\mathbf{S}_{3.3}$ \mathbf{M} methylene Chloride $91-20-3$ 128 $3_{117E+01}$ $0_{0.00E+00}$ $4_{4.82E-04}$ 962 $3_{3.3}$ \mathbf{N} mapthalene $7440-02-0$ 59 $1.00E+00$ $6_{6.80E-04}$ $1.59E-04$ $14,000$ $4_{4.46}$ \mathbf{N} henanthrene $7440-22-4$ 108 $1.00E+00$ $6_{6.80E+04}$ $1.59E-04$ $14,000$ $4_{4.46}$ \mathbf{N} rickel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} rest \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} rickel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} rest \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} rickel \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} \mathbf{N} rest \mathbf{N} rest \mathbf{N} rest \mathbf{N} \mathbf | A Methoxychlor 1.75E+03 Methylene Chloride 75-09-2 85 1.75E+03 Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 2-methylnapthalene 91-20-3 128 3.17E+01 0.00E+00 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 Nickel 7440-02-0 59 1.00E+00 6.80E-04 Nickel 7440-22-4 108 0.00E+00 6.80E-04 Silver 7440-22-4 108 0.00E+00 6.80E+00 Silver 100-22-4 108 0.00E+00 0.00E+00 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene 108-88-3 92 5.35E+02 2.81E+01 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3.2E4+02 Trichloroethene 79-01-6 131 1.098E+02 1.00E+01 1.00E+01 | 2.00E-03 NA | 5,50 |
| Methylene Chloride $75-09-2$ 85 $2.00E+04$ $3.62E+02$ $2.03E-03$ 8.8 1.30 $2^{-methylnapthalene}$ $91-20-3$ 128 $3.17E+01$ $4.82E-04$ 962 3.30 $Naphthalene$ $91-20-3$ 128 $3.17E+01$ $4.82E-04$ 962 3.30 $Naphthalene$ $91-20-3$ 128 $3.17E+01$ $4.82E-04$ 962 3.30 $Nickel$ $7440-02-0$ 59 $0.00E+00$ $1.59E-04$ $14,000$ 4.46 $Nickel$ $7440-22-4$ 108 $1.00E+00$ $6.80E-04$ $14,000$ 4.46 $Nickel$ $7440-22-4$ 108 $0.00E+00$ $0.00E+00$ 1.46 3.3 $Nickel$ $700E+01$ $6.80E-04$ $1.59E-04$ $14,000$ 2.73 $Nickel$ $7440-22-4$ 108 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 $Nickel$ $100-802$ $2.81E+01$ $6.37E-03$ 300 2.73 700 $Naxaphene79-01-61311.10E+035.72E+02590.48Trans 1, 2-dichloroethene79-01-61311.10E+035.79E+019.10E-032126NickelNickel1.90E+017.00E+017.00E+017.04E-032.40NickelNickel1.00E+017.00E+017.00E+017.00E+017.00E+01NickelNickel1.00E+023.2E4+029.10E-032.202.03Nickel1.00E+01$ | Methylene Chloride 75-09-2 85 2.00E+04 3.62E+02 2-methylnapthalene 91-20-3 128 3.17E+01 0.00E+00 Naphthalene 91-20-3 128 3.17E+01 0.00E+00 Nickel 7440-02-0 59 1.00E+00 6.80E-04 Nickel 7440-22-4 178 1.00E+00 6.80E-04 Silver 7440-22-4 108 0.00E+00 6.80E+00 Toluene 7440-22-4 108 5.35E+02 2.81E+01 Toluene 7440-22-4 108 5.35E+02 2.81E+01 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene 108-88-3 92 5.00E-01 3.2E4+02 Trans 1,2-dichloroethene 79-01-6 131 1.00E+03 5.79E+01 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 | 5.59E-03 83 | 2.12 1,00 |
| 2-methylnapthalene $2-methylnapthalene4.82E-049623.30Naphthalene91-20-31283.17E+014.82E-049623.30Naphthalene7440-02-0593.17E+010.00E+0014,0004.46Nickel7440-02-0591.00E+001.59E-0414,0004.46Phenanthrene7440-22-41081.00E+006.80E-0414,0004.46Silver708-88-3925.35E+022.81E+016.37E-033002.73Toluene108-88-3925.35E+022.81E+016.37E-033002.73Toluene540-59-0976.00E+023.244+024.2E-02590.48Trans 1,2-dichloroethene79-01-61311.10E+035.79E+019.10E-032.20Xylenes1330-20-71061.98E+021.00E+017.04E-032.403.26$ | $\begin{array}{c ccccccccccccccccccccccccccccccccccc$ | 3.62E+02 2.03E-03 8.8 | 1.30 |
| Naphthalene $91-20-3$ 128 $3.17E+01$ $4.82E-04$ 962 3.30 Nickel $740-02-0$ 59 $3.17E+01$ $0.00E+00$ $4.82E-04$ $14,000$ 4.46 Nickel $7440-02-0$ 59 $1.00E+00$ $1.59E-04$ $14,000$ 4.46 Phenanthrene $7440-22-4$ 108 $1.00E+00$ $6.80E-04$ $14,000$ 4.46 Silver $701ere$ $108-88-3$ 92 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 Toluene $108-88-3$ 92 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 Toluene $540-59-0$ 97 $6.00E+02$ $3.264+02$ $4.2E-02$ 59 0.48 Trans 1,2-dichloroethene $79-01-6$ 131 $1.10E+03$ $5.79E+01$ $9.10E-03$ 126 2.38 Xylenes $1330-20-7$ 106 $1.98E+02$ $1.00E+01$ $7.04E-03$ 240 3.26 | Naphthalene 91-20-3 128 3.17E+01 Nickel 7440-02-0 59 0.00E+00 Phenanthrene 7440-02-0 59 0.00E+00 Silver 85-01-8 178 1.00E+00 6.80E-04 Silver 7440-22-4 108 0.00E+00 0.00E+00 Toluene 7440-22-4 108 0.00E+02 0.00E+00 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene 5.00E-01 97 5.00E-01 3,2E4+02 Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 Xylenes 136 1.98E+02 1.00E+01 1.00E+01 | | 40-30 |
| Nickel $7440^{-0.2}$ -0 59 $0.00E^{+00}$ 0.0 Phenanthrene $7440^{-0.2}$ -1 178 $1.00E^{+00}$ $0.00E^{+00}$ $14,000$ 4.46 Phenanthrene $7440^{-2.2}$ -4 108 $1.00E^{+00}$ $0.00E^{+00}$ $1.59E^{-04}$ $14,000$ 4.46 Silver $7440^{-2.2}$ -4 108 $0.00E^{+00}$ $0.37E^{-03}$ 300 2.73 Toluene 108^{-88-3} 92 $5.35E^{+02}$ $2.81E^{+01}$ $6.37E^{-03}$ 300 2.73 Towaphene 79^{-01-6} 131 $1.10E^{+03}$ $3.2E4^{+02}$ $4.2E^{-02}$ 5.964 3.3 Trans 1, 2-dichloroethene 79^{-01-6} 131 $1.10E^{+03}$ $5.79E^{+01}$ $9.10E^{-03}$ 240 3.26 Xylenes 1.30^{-20^{-7} 1066 $1.98E^{+02}$ $1.000E^{+01}$ $7.04E^{-03}$ 240 3.26 | Nickel $740-0.2-0$ 59 $0.00E+00$ Phenanthrene $740-0.2-0$ 59 $500E-04$ Silver $85-01-8$ 178 $1.00E+00$ $6.80E-04$ Silver $7440-22-4$ 108 $0.00E+00$ $0.00E+00$ Toluene $108-88-3$ 92 $5.35E+02$ $2.81E+01$ Toxaphene $1,2-dichloroethene$ $540-59-0$ 97 $6.00E+02$ $3,2E4+02$ Trichloroethene $79-01-6$ 131 $1.10E+03$ $5.79E+01$ Xylenes $130-20-7$ 106 $1.98E+02$ $1.00E+01$ | 4.82E-04 962 | 3.30 40-30 |
| Phenanthrene $85-01-8$ 1781.00E+00 $6.80E-04$ $1.59E-04$ $14,000$ 4.46 Silver $7440-22-4$ 108 $0.00E+00$ $0.07E+00$ 37 300 2.73 Toluene $108-88-3$ 92 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 Toluene $108-88-3$ 92 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 Trans 1, 2-dichloroethene $540-59-0$ 97 $6.00E+02$ $3.2E4+02$ $4.2E-02$ 59 0.48 Trans 1, 2-dichloroethene $79-01-6$ 131 $1.10E+03$ $5.79E+01$ $9.10E-03$ 126 2.38 Xylenes $1330-20-7$ 106 $1.98E+02$ $1.00E+01$ $7.04E-03$ 240 3.26 | Phenanthrene 85-01-8 178 1.00E+00 6.80E-04 Silver 7440-22-4 108 0.00E+00 0.00E+00 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene 540-59-0 97 6.00E+02 3,2E4+02 Trains 1,2-dichloroethene 79-01-6 131 1.00E+03 5.79E+01 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 | 0.00E+00 NA | 4 |
| Silver $7440-22-4$ 1080.00E+00NAToluene $108-88-3$ 92 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 Toluene $108-88-3$ 92 $5.35E+02$ $2.81E+01$ $6.37E-03$ 300 2.73 Toraphene $7-01-6$ $5.00E-01$ $4.36E-01$ 964 3.3 Trans 1,2-dichloroethene $540-59-0$ 97 $6.00E+02$ $3,2E4+02$ $4.2E-02$ 59 0.48 Trichloroethene $79-01-6$ 131 $1.10E+03$ $5.79E+01$ $9.10E-03$ 126 2.38 Xylenes $1330-20-7$ 1066 $1.98E+02$ $1.00E+01$ $7.04E-03$ 240 3.26 | Silver 7440-22-4 108 0.00E+00 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toluene 108-88-3 92 5.35E+02 2.81E+01 Toraphene 540-59-0 97 6.00E+02 3,2E4+02 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 Xylenes 131 1.00E+03 5.79E+01 Xylenes 1.00E+02 1.00E+01 | 6.80E-04 1.59E-04 14,000 | 4.46 2,63 |
| Toluene 108-88-3 92 5.35E+02 2.81E+01 6.37E-03 300 2.73 Toxaphene 108-88-3 92 5.00E-01 4.36E-01 964 3.3 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 4.2E-02 59 0.48 Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 9.10E-03 126 2.38 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 7.04E-03 240 3.26 | Toluene 108-88-3 92 5.35E+02 2.81E+01 Toxaphene Toxaphene 5.00E-01 5.00E+01 3,2E4+02 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 Trichloroethene 79-01-6 131 1.00E+03 5.79E+01 Xylenes 1330-20-7 106 1.99E+02 1.00E+01 | 0.00E+00 NA | 3,06 |
| Toxaphene 5.00E-01 4.36E-01 964 3.3 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 4.2E-02 59 0.48 Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 9.10E-03 126 2.38 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 7.04E-03 240 3.26 | Toxaphene 5.00E-01 Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 Xylenes 136-20-7 106 1.98E+02 1.00E+01 | 2.81E+01 6.37E-03 300 | 2.73 10. |
| Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 4.2E-02 59 0.48 Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 9.10E-03 126 2.38 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 7.04E-03 240 3.26 | Trans 1,2-dichloroethene 540-59-0 97 6.00E+02 3,2E4+02 Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 | 4.36E-01 964 | 3.3 13,1C |
| Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 9.10E-03 126 2.38 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 7.04E-03 240 3.26 | Trichloroethene 79-01-6 131 1.10E+03 5.79E+01 Xylenes 1330-20-7 106 1.98E+02 1.00E+01 | 3,2E4+02 4.2E-02 59 | 0.48 1. |
| Xylenes 1330-20-7 106 1.98E+02 1.00E+01 7.04E-03 240 3.26 | Xylenes 1330-20-7 106 1.98E+02 1.00E+01 | 5.79E+01 9.10E-03 126 | 2.38 10. |
| | | 1.00E+01 7.04E-03 240 | 3.26 |
| VN 0.042UU-0, C0 0.207 /440-04 /440-04 /440-04 /440-04 /440-04 /440-04 /440-04 /440-04 /440-04 /440-04 /440-04 | Zinc 7440-66-6 65 0.00E+00 | 0.00E+00 NA | 4 |

Note: NA = Information not available.

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Source: Water-Related Environmental Fate of 129 Priority Pollutants. EPA 440/4~79-029a. 1979. Vershueren, 1983. Superfund Public Health Evaluation Manual, EPA 54D/1-86-060 Oct. 1986.

CVR109/022

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| | GROUNDWATER | Tabl CONTAMINANT COMPA (ESKIMO CREEK SEE | e 4-21 RISONS TO WATER (P, SHALLOW AQUIF | JUALITY CRITERIA ER) | | | |
|--|---|--|--|---|---|---|---|
| Surface Water Contaminants ^a | Number of Wells Sampled vs. Contaminant Detections | Maximum Detected Concentration (ug/l) | Mean ^b Detected Concentration (ug/l) | Federal Drinking Water Criteria (ug/l) | Alaska ^C Drinking Water Criteria (ug/l) | Aquatic ^d Acute Toxicity (ug/1) | Aquatic ^d Chronic Toxicity (ug/l) |
| Lead | 23/19 | 370 | 37 | 50 ^e | 50 | 82 ^f | 3.2 ^f |
| Fluoride | 23/20 | 110 | 50.3 | | 2,400 | | |
| Total Dissolved Solids | 23/23 | 558,000 | 336,000 | 250 ,000⁹ | 500,000 | | |
| Nitrate-Nitrite | 23/23 | 11,400 | 2,630 | 10,000 ⁹ | 10,000 | | |
| Total Petroleum Hydrocarbons | 23/12 | 5,670,000 | | | | | |
| Total Organic Carbons | 2/2 | 7,200 | 4,450 | | | | |
| Benzene | 8/4 | 11 | 11.6 | 5 ^h | | 5,300 ¹ | |
| Toluene | 8/4 | 130 | 21.8 | 2 , 000 ^j | | 17,000 ¹ | |
| Ethyl benzene | 8/4 | 130 | 25.1 | 680 ^j | ., | 32 , 000 ¹ | |
| Xylene | 8/4 | 520 | 107 | 440 ^j | | | |
| <pre>aChlorobenzene, 1,4-dichlorobe detection (detection 'imit = bArithmethic mean of detected</pre> | nzene, 1,3-dichloro 1.0 ug/1). concentrations. | benzene, and 1,2- | dichlorobenzene | were also analyse | d. Their conce | ntrations w | ere below |
| ^C State of Alaska Drinking Wate | er Regulations, Depa | rtment of Environ | mental Conservat | lon (1982). | | | |

^fHardness dependent criterion (100 mg/l used). ^gUSEPA Water Quality Criteria for Drinking Water (1986).

^eRCRA MCL for drinking water (May, 1988).

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^dFederal Clean Water Act: Ambient Water Quality Criteria for the Protection of Aquatic Life (EPA, 1986).

^hSDWA MCL for drinking water (May, 1988).

¹Lowest observed effect level (LOEL).

^JProposed SDWA MCL for drinking water (May, 1988).

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The maximum level of lead identified (370 ppb) exceeds drinking water (50 ppb), aquatic organism acute (82 ppb), and aquatic organism chronic toxicity (3.2 ppb) criteria. The mean level of lead (37 ppb) meets drinking water and acute aquatic organism toxicity criteria. However, this average lead level exceeds the chronic toxicity criterion for the protection of aquatic organisms.

The maximum total dissolved solids (TDS) level (558,000 ppb) exceeds the federal (250,000 ppb) and Alaska (500,000 ppb) criteria for drinking water. The mean concentration (336,000 ppb) meets the Alaska drinking water criterion but still exceeds the federal criterion. Because this is a shallow aquifer that is close to the coast, these TDS levels are more likely due to the influence of the ocean, rather than King Salmon AFS operations. There are no total dissolved solids criteria for the protection of aquatic organisms.

The maximum (77 ppb) and mean (11.6 ppb) levels of benzene in the shallow wells exceed the federal drinking water criterion (5 ppb). These levels do not exceed the acute toxicity criterion for the protection of aquatic organisms (5,300 ppb). There is no chronic toxicity criterion for benzene for the protection of aquatic organisms.

The maximum level of xylene (520 ppb) exceeds the federal drinking water criterion (440 ppb). The mean level of xylene (107 ppb) meets the federal arinking water criterion. There are no acute or chronic xylene criteria for the protection of aquatic organisms.

4.3.2.1.2 Intermediate Groundwater

The intermediate depth wells were tested for the same contaminants as the shallow wells (see Table 4-21). Of these contaminants, only the maximum (378,000 ppb) and mean (302,000 ppb) TDS levels exceeded the federal drinking water quality criterion (250,000 ppb). These total dissolved solids levels meet the Alaska drinking water criterion. As with the shallow aquifer, the TDS levels in the intermediate depth wells are more likely due to the influence of the ocean, rather than King Salmon AFS operations.

None of the contaminants measured in the intermediate depth wells exceeded the acute or chronic criteria for the protection of aquatic organisms.

4.3.2.1.3 Deep Groundwater

The King Salmon AFS wells were analyzed for selected inorganics, PCBs, and pesticides. The inorganic concentrations identified meet federal and Alaska drinking water quality criteria. No PCBs or pesticides were detected in the King

Salmon Airport well samples. High levels of TPHs (27,100 ppb and 6,400 ppb) were detected in King Salmon Airport wells drawing water from this deep aquifer. The source of these constituents is uncertain and could be from lubricating fluids contained within the well pumps or from other potential sources, such as leakage along the well bore from the overlying contaminated shallow aquifer.

4.3.2.1.4 Surface Water

The surface water data collected for Eskimo Creek are summarized in Table 4-22. Of the contaminants analyzed, only zinc exceeded the federal water quality criterion for the protection of aquatic organisms from chronic toxicity. Cadmium, copper, and mercury were analyzed at detection limits greater than the criteria for protection of aquatic organisms. Therefore, they could be present at levels that are hazardous to aquatic organisms.

4.3.2.1.5 Soils

Soil data collected for the Eskimo Creek seep site are summarized in Table 4-23. There are no federal or Alaska criteria for contaminated soil or sediment.

4.3.2.2 Naknek River Seep

The inorganic contaminants detected at this site met federal and Alaska criteria for drinking water and acute aquatic organism toxicity. Several of the inorganic constituents analyzed were below detection limits; however, the detection limits exceeded the criteria for chronic aquatic organism toxicity. These inorganics included cadmium (detection limit = 5 ppb; chronic criteria = 1.1 ppb), lead (5 ppb; 3.2 ppb), mercury (0.5 ppb; 0.012), and antimony (5 ppb; 1.6 ppb). Copper was also below detection (30 ppb) but the detection limit exceeds both the acute (18 ppb) and chronic (12 ppb) criteria for the protection of aquatic organisms.

All the organic contaminants detected in the Naknek River met federal and Alaska drinking water criteria. None of the detected organic contaminants exceeded acute or chronic water quality criteria for the protection of aquatic organisms.

4.3.2.3 North Barrel Bluff

4.3.2.3.1 Surface Water

Samples from King Salmon Creek were analyzed for selected inorganic, organic, PCB, and pesticide contaminants. All the organics tested were below detection limits and the

| | SURFACE WAT | T: ER CONTAMINANT C((ES) | able 4-22 Omparisons to Way Kimo Creek) | rer Quality Crite | RIA | | |
|---|---|--|---|---|---|---|---|
| Surface Water Contaminants ^a | Number of Wells Sampled vs. Contaminant Detections | Maximum Detected Concentration (ug/1) | Mean Mean Detected Concentration (ug/l) | Federal Drinking Water Criteria (ug/l) | Alaska ^C Drinking Water Criteria (ug/l) | Aquatic Aquatic Acute Toxicity (ug/1) | Aquatic Aquatic Chronic Toxicity (ug/l) |
| Lead | 2/2 | 70 | 50 | | | 320 ^e | 47 ^e |
| Chloride | 2/2 | 5,380 | 5,360 | | 250,000 | | |
| Sulfate | 2/2 | 1,670 | 1,650 | | 250,000 | | |
| Fluoride | 2/0 | <500 | <500 | | 2,400 | | |
| Total Dissolved Solids | 2/2 | 171,000 | 113,000 | 250 , 000 ^e | 500,000 | | |
| Nitrate-Nitrite | 2/1 | 170 | 110 | 10,000 ^e | 10,000 | | |
| | | | | | | | |

^aBenzene, toluene, xylene, ethylbenzene, chlorobenzene, 1,4-dichlorobenzene, 1,2-dichlorobenzene, 1,3-dichlorobenzene and priority pollutant metals were also analyzed. Organic concentrations were below detection (detection limit = 1.0 ug/1). Metals levels were also below their respective detection levels, which did not exceed federal or state action levels for human health or the environment (see Table 4-16E).

^bArithmetic mean of detected concentrations.

^CState of Alaska Drinking Water Regulations, Department of Environmental Conservation (1982).

^dFederal Clean Water Act: Ambient Water Quality for the Protection of Aquatic Life (EPA, 1986).

^eUSEPA Water Quality Criteria for Drinking Water (1986).

^fHardness dependent criterion (100 mg/l used).

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| Contaminant | Maximum Concentration (ug/kg) |
|----------------------------|-------------------------------------|
| Toluene | 1,500 |
| Ethylbenzene | 400 |
| Xvlene (o and m) | 1,800 |
| Xylene (p) | 1,500 |
| Chloromethane | 3,600 |
| 2-butanone | 3,100 |
| 2-methylnaphthalene | 8,500 |
| Fluorene | 630 |
| Phenanthrene | 210 |
| Naphthalene | 1,300 |
| Bis(2-ethylhexyl)phthalate | 1,500 |
| Butyl benzyl phthalate | 1,800 |
| Di-n-octyl phthalate | 210 |
| Methylene chloride | 16,000 |
| Acetone | 8,600 |

Table 4-23 ESKIMO CREEK SOILS DATA

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detection limits used did not conflict with drinking water or aquatic organism water quality criteria.

All the pesticides and PCBs analyzed were below detection limits. However, the detection limits used for heptachlor, Endosulfan I, dieldrin, endrin, methoxychlor, chlordane, and toxaphene were greater than the chronic criteria for the protection of aquatic organisms. In addition, the maximum toxaphene detect (1 ppb) exceeded the acute (0.73 ppb) aquatic organism toxicity criterion.

The inorganic contaminants analyzed met federal and Alaska drinking water criteria. However, as with the Naknek River seep, the detection limits for cadmium, copper, lead, mercury, and antimony exceeded the chronic toxicity criteria for aquatic organisms. Also, the copper detection limit used exceeded the acute toxicity criterion for aquatic organisms. Therefore, these constituents could be present at hazardous levels.

4.3.2.3.2 Sediments

Selected organic analyses were conducted on eight sediment samples from King Salmon Creek. Two organic contaminants were detected in the samples. Toluene (3 and 4 ppb) and 2-butanone (20 ppb) were detected in only one of the four samples collected. There are no federal or Alaska human health or environmental criteria for exposure to contaminated sediments.

4.3.2.4 South Barrel Bluff

4.3.2.4.1 Surface Water

None of the organic contaminants analyzed were above their detection limits and the detection limits do not conflict with federal or Alaska water quality criteria for drinking water or the protection of aquatic organisms.

The inorganic contaminants analyzed met federal and Alaska drinking water criteria. As with the Naknek River seep, the detection limits for cadmium, copper, lead, mercury, and antimony exceeded the chronic toxicity criteria for aquatic organisms. The copper detection limit also exceeded the acute toxicity criterion for aquatic organisms.

All the pesticides and PCBs analyzed were below detection limits. However, the detection limits used for heptachlor, Endosulfan I, dieldrin, endrin, methoxychlor, chlordane, and toxaphene were greater than their chronic criteria for the protection of aquatic organisms.

4.3.2.4.2 Groundwater

No organics, pesticides or PCBs were detected in the groundwater well tested at this site. There were no conflicts between the detection limits used at this operable unit and federal or Alaska water quality criteria.

All the inorganic contaminants analyzed for the well at this operable unit were below detection limits. However, the detection limits for arsenic (300 ppb), lead (200 ppb), and selenium (400 ppb) greatly exceed the federal criteria for drinking water (50 ppb, 50 ppb, and 10 ppb, respectively). Therefore, these contaminants could be present at hazardous levels.

4.3.3 Human Receptors/Human Health Effects

This section identifies the potential human receptors (base personnel, local residents, etc.) that could be adversely affected by site contaminants. It also addresses the potential for human exposure to site contaminants. The presence of contaminated media and the potential for human exposure to those media suggest that the potential exists for adverse human health effects. A summary of the potential human exposure pathways considered for King Salmon AFS is presented in Figure 4-16.

4.3.3.1 Eskimo Creek

4.3.3.1.1 Shallow Groundwater

The shallow groundwater aquifer at King Salmon AFS is not known to be used as a water source (private, municipal, industrial) for the base or surrounding areas. The principal pathways for exposure to the organic and inorganic contaminants identified in the shallow groundwater would be dermal contact or incidental ingestion of water in locations downstream from the seep where it enters Eskimo Creek. The potential also exists for the consumption of potentially contaminated fish (resident trout or salmon).

The section of Eskimo Creek associated with the shallow groundwater is not known to be used for drinking water. Therefore, a drinking water exposure pathway is not considered an immediate hazard for Eskimo Creek.

Potential receptors for shallow groundwater contamination could include King Salmon AFS personnel, local residents or visitors who fish, hunt, boat, or hike downstream from where the shallow groundwater seep enters Eskimo Creek.

4.3.3.1.2 Intermediate Groundwater

No contamination (except total dissolved solids) has been identified in this groundwater source. It is not believed that the TDS levels in the intermediate depth groundwater is associated with King Salmon Airport operations. Based on the information available, the intermediate groundwater aquifer is not believed to represent a substantial immediate human exposure pathway. This aquifer is not currently used as a potable water source. However, a more thorough chemical analysis of this aquifer is needed to confirm that it does not represent a potential human health hazard.

4.3.3.1.3 Deep Groundwater

The deep groundwater aquifer at King Salmon Airport is used as the primary water source for the base. This aquifer is also believed to be the primary water source for the King Salmon community. Therefore, potential receptors for any contaminants in this aquifer would include King Salmon Airport personnel, local residents, and visitors who use the water for drinking, cooking, or bathing.

No contaminants in excess of federal or Alaska drinking water standards have been identified for this deep aquifer. However, total petroleum hydrocarbons (TPHs) have been detected. TPHs may be due to oil in the well pumps or other sources outside the well bore. It is possible that the TPHs contain toxic components.

4.3.1.1.4 Surface Water

Potential human receptors for surface water contamination in Eskimo Creek could include base personnel, local residents, or area visitors. Exposure pathways for Eskimo Creek could include dermal contact and incidental water ingestion while using the area for fishing, hunting, boating, or hiking. The possibility also exists for the consumption of potentially contaminated fish. The section of Eskimo Creek associated with this operable unit is not known to be used for drinking water. Therefore, a complete drinking water exposure pathway has not been identified for Eskimo Creek.

4.3.3.1.5 Soils

Potential receptors for soil contamination on the Eskimo Creek Operable Unit would primarily include base personnel working in the area. Exposure to the soil contaminants identified could occur through dermal contact, incidental soil ingestion, or contaminated dust inhalation.

4.3.3.2 Naknek River Seep

4.3.3.2.1 Surface Water

Potential receptors for surface water contamination at the Naknek River seep could include base personnel, local residents, and area visitors. However, because of the rugged and remote nature of the area near the seep, it is unlikely that significant human exposure to contaminants would occur. The Naknek River seep is adjacent to a tidal mud flat. The water in this area is not a potential potable water source because it is saline. Further, the area is not popular for hiking, clamming, fishing, crabbing, or hunting. Therefore, significant human exposure through ingestion (water, sediment, potentially contaminated fish, etc.) or dermal contact is not considered likely.

4.3.3.3 North Barrel Bluff

4.3.3.3.1 Surface Water

Potential receptors for surface water contamination at North Barrel Bluff could include base personnel, local residents, or area visitors. The organic, inorganic, pesticide, and PCB data for surface water meet federal and state criteria for drinking water. Based on current information, the principal human exposure pathway of concern for surface water is by consumption of potentially contaminated fish.

4.3.3.3.2 Sediments

Potential receptors for sediment contamination on the North Barrel Bluff could include base personnel, local residents, or area visitors. Exposure to the toluene and 2-butanone in the sediments at this site could occur through dermal contact or incidental ingestion while using this section of King Salmon Creek (or downstream areas) for fishing, hunting, boating, or hiking. Unfiltered King Salmon Creek water used as a drinking water source could also represent a potential human exposure pathway to contaminated sediments. Further, the potential exists for consumption of potentially contaminated fish.

4.3.3.3.3 Barrel Contents

The exact contents of exposed or buried barrels at this area are not known. However, potentially hazardous materials are believed to be contained in many of the barrels. If the barrels leak, these hazardous materials may pose potential human health risks for base personnel, local residents, and area visitors.

4.3.3.4 South Barrel Bluff

4.3.3.4.1 Surface Water

Potential receptors for surface water contamination on the South Barrel Bluff could include base personnel, local residents, and area visitors. Potential exposure to surface water contaminants could occur through water use for drinking or through dermal contact and incidental ingestion while fishing, hunting, boating, or hiking.

The data available for this site indicate that no inorganic, organic, pesticide, or PCB contaminants were detected at levels exceeding state or federal drinking water quality criteria. Chlordane, a carcinogen, was detected $(0.2 \ \mu g/l)$ but there are not drinking water criteria for it. Based on current information, the principal exposure pathway of concern for surface water is the potential consumption of potentially contaminated fish.

4.3.3.4.2 Groundwater

The well tested at this site is used by local residents as their potable water source. Receptors for contamination in this well would include the residents and their guests. Human exposure to potential well contaminants could occur through drinking, bathing, cooking, and other domestic water uses.

The data available for this well indicate compliance with drinking water criteria for organics, pesticides, and PCB. The inorganic results for lead, arsenic, and selenium are suspect because their detection limits exceed the drinking water quality criteria for these contaminants. Therefore, it is possible that these three contaminants pose a current health threat for the people using the well.

4.3.3.4.3 Barrel Contents

The exact content of exposed or buried barrels at this area unit are unknown. However, potentially hazardous materials are believed to be contained in many of the barrels. If the barrels leak, these hazardous materials pose potential human health risks for base personnel, local residents, and area visitors.

4.3.4 Carcinogenic Risks

This section of the report identifies the carcinogenic contaminants detected to date at each operable unit. The presence of media contaminated with carcinogens and the potential for human exposure to the contaminated media suggests a potential carcinogenic risk exists. Expanded media analyses (e.g., priority pollutant scans) and additional chemical characterizations of TPHs and TOCs could reveal the presence of additional carcinogens at the King Salmon AFS operable units. Cancer potency factors for chemicals of concern were listed in Table 4-19A.

4.3.4.1 Eskimo Creek Seep

Carcinogens have been detected in the shallow groundwater (benzene, a known human carcinogen) and soils (methylene chloride and bis (2-ethylhexyl) phthalate, probable human carcinogens) at this operable unit. The potential human exposure pathways for these contaminated media include incidental ingestion (soil, groundwater), dermal contact (soil, groundwater), and dust inhalation (soil). The primary receptors are base personnel (soil, groundwater), local residents (groundwater), and area visitors (groundwater).

4.3.4.2 Naknek River Seep

Trichloroethene (a probable human carcinogen) has been detected in the Naknek River seep. The potential for substantial human exposure to contamination in this rugged remote area is not considered likely.

4.3.2.3 North Barrel Bluff

No carcinogens have been verified at this operable unit to date. However, it is possible that several pesticides that are carcinogenic (Endosulfan, dieldrin, methoxychlor, chlordane, and toxaphene) are present. Sampling at reduced detection limits will allow their presence to be confirmed or discounted.

4.3.4.4 South Barrel Bluff

No carcinogens have been verified for this operable unit. However, it is possible that arsenic (a known human carcinogen) is present at levels exceeding the drinking water criterion in the well sampled at this site. The data show arsenic to be below detection but the detection limit was above drinking water quality criteria. The receptors of this potential contaminant include the residents using the well and their guests. More extensive sampling is needed here to verify the presence or absence of arsenic in the well.

Similarly, several pesticides that are carcinogenic (Endosulfan, dieldrin, methoxychlor, chlordane, and toxaphene) could be present in local surface waters and sediments. Human exposure to these contaminants could occur while hiking, hunting, or fishing in the area. Analysis with reduced detection limits is needed to verify their presence or absence.

4.3.5 Threat to the Environment

This section presents information concerning the potential for adverse contaminant effects on fish and wildlife resources.

4.3.5.1 Eskimo Creek Seep

Shallow groundwater lead contamination poses a potential environmental threat in areas downstream from where the seep enters Eskimo Creek. The maximum lead concentrations detected in the shallow groundwater samples exceed the acute and chronic criteria for the protection of aquatic organisms. The average lead concentration exceeds the chronic aquatic organism toxicity criteria. Lead also has a moderate bioconcentration factor and represents a potential threat to predators (resident trout, eagles, bears, waterfowl) that eat potentially contaminated prey (insects, fish, etc.).

Several organic contaminants have been detected in the soils at this operable unit. Those present in surface soils could pose a threat to fish and wildlife through surface runoff entering Eskimo Creek, incidental soil ingestion during feeding or preening, and dermal contact. Three of these contaminants (xylene, naphthalene, bis (2-ethylhexyl) phthalate have moderate-to-high bioconcentration factors and could present a threat to the top predators (resident trout, eagles, bears, waterfowl) that eat contaminated prey.

4.3.5.2 Naknek River Seep

No immediate environmental threat has been identified to date at this operable unit. However, the detection limits for cadmium, lead, copper, mercury, and antimony were greater than the chronic aquatic organism toxicity criteria. The detection limit for copper also exceeded the acute toxicity criteria for aquatic organisms. Therefore, even though these contaminant concentrations were below detection, they could be present at chronically toxic levels. Further, all these contaminants have moderate-to-high bioconcentration factors and could represent a threat to predators (resident trout, eagles, bears, waterfowl) that eat potentially contaminated prey.

4.3.5.3 North Barrel Bluff

The maximum concentration detected for toxaphene at this operable unit (1 ppb) exceeds the acute toxicity criteria for aquatic organisms. Therefore, localized toxaphene concentrations may currently present an immediate environmental threat to aquatic organisms. The detection limits for cadmium, lead, copper, mercury, and antimony were greater than the chronic aquatic organism toxicity criteria, and the detection limit for copper was higher than the acute toxicity criterion for aquatic organisms at this operable unit. Therefore, even though these inorganic contaminant concentrations were below detection, they could be present at toxic levels. Furthermore, all these contaminants have moderate-to-high bioconcentration factors and could represent a threat to predators (resident trout, eagles, bears, waterfowl) that eat potentially contaminated prey.

Similarly, the detection limits for heptachlor, Endosulfan I, dieldrin, methoxychlor, chlordane, and toxaphene were higher than the chronic water quality criteria for the protection of aquatic organisms. Therefore, even though these pesticide concentrations were below detection, they could be present at toxic levels. These contaminants have high bioconcentration factors and could represent a threat to predators (resident trout, eagles, bears, waterfowl) that eat potentially contaminated prey.

4.3.5.4 South Barrel Bluff

Based on current information, no immediate environmental threat is apparent at this operable unit. However, the detection limits for cadmium, lead, copper, mercury, and antimony were greater than the chronic aquatic organism toxicity criteria. The detection limit for copper also exceeded the acute toxicity criterion for aquatic organisms. Therefore, even though these contaminant concentrations were below detection, they could be present at toxic levels. Furthermore, all these contaminants have moderate-to-high bioconcentration factors and could represent a threat to predators (resident trout, eagles, bears, waterfowl) that eat potentially contaminated prey.

Similarly, the detection limits for heptachlor, Endosulfan I, dieldrin, methoxychlor, chlordane, and toxaphene were greater than the chronic water quality criteria for the protection of aquatic organisms. Therefore, even though these pesticide concentrations were below detection, they could be present at toxic levels. All these contaminants have high bioconcentration factors and could represent a threat to predators (resident trout, eagles, bears, waterfowl) that eat potentially contaminated prey.

4.3.6 Finding of No Significant Impact

Given the current limited data base for site contaminants, it is not possible to conclude that the contamination present at the four operable units identified has no significant impact.

The pesticide detection limits should be lowered so it can be determined if their levels comply with water quality criteria for the protection of aquatic organisms or human health.

Deep wells should be resampled with reduced arsenic, selenium, and lead detection limits to determine if water from this well poses a human health threat.

King Salmon Creek sediments should be sampled and analyzed for inorganics, organics, and pesticides to determine if they may pose human or environmental threats.

Detection limits for all carcinogens, particularly volatile organics, should be at least low enough to be equivalent to a 10- excess cancer risk.

4.4 APPLICABLE OR RELEVANT AND APPROPRIATE REQUIREMENTS

As presented earlier in this document, the Stage 1 field investigation is for identified contamination at all four sites at King Salmon AFS. A preliminary assessment of Applicable or Relevant and Appropriate Requirements (ARARs) was conducted to identify any existing conditions which violate federal or state ARARs. The results of this assessment were also used in the development of remedial alternatives (Section V), and recommendations (Section VI).

The concept of ARARs was developed as part of the Comprehensive Environmental Response Compensation and Liability Act (CERCLA) program for site cleanups. Under CERCLA, as amended by the Superfund Amendments and Reauthorization Act (SARA), remedial actions on CERCLA sites must comply with all federal and state ARARs unless one of six conditions are met (waivers).

The Installation Restoration Program (IRP) is the Department of Defense's (DOD) response to the requirements of CERCLA. All federal agencies must comply with the procedural and substantive requirements of the Superfund program. The IRP objective is derived from the overall objective of the National Contingency Plan (NCP).

4.4.1 Definitions

ARARs are federal and duly promulgated state environmental and public health laws, requirements, and regulations. In evaluating potential ARARs for a site, a determination is made as to whether a requirement is applicable, relevant, or neither.

Applicable requirements are those cleanup standards, standards of control, and other substantive environmental protection requirements promulgated under federal or state law that specifically address the hazardous substance, pollutant, contaminant, remedial action, location, or other circumstance at the site.

For a requirement to be applicable, the remedial action or the circumstance at the site must satisfy all of the jurisdictional prerequisites of that requirement. For example, the minimum technology requirement for hazardous waste landfills under RCRA would apply only if a new hazardous waste landfill (or an expansion of an existing hazardous waste landfill) were to be built on the site.

Relevant and appropriate requirements are those cleanup standards, standards of control, and other substantive environmental protection requirements promulgated under federal or state law that, although not "applicable" to a hazardous substance, pollutant, contaminant, remedial action, location, or other circumstance at the site, address problems or situations sufficiently similar to those encountered, that their use is well suited to the site. In some circumstances, a requirement may be relevant but not appropriate to the specific situation.

The relevance and appropriateness of a requirement can be judged by comparing factors such as the characteristics of the remedial action, the hazardous substances in question, and the physical characteristics of the site with those characteristics addressed by the requirement. For example, RCRA hazardous waste management requirements would not be applicable to wastes that could not be strictly classified as hazardous wastes. However, if those wastes are similar to hazardous wastes, the RCRA requirements could be relevant and appropriate to their management.

ARARS are divided into three categories: chemical-specific, location-specific, and action-specific. Chemical-specific ARARS include those requirements that regulate the release to the environment of materials possessing certain chemical or physical characteristics or materials containing specific chemical compounds. These requirements generally set health or risk-based concentration limits or discharge limitations for specific hazardous substances. If, in a specific situation, a chemical is subject to more than one discharge or exposure limit, the more stringent of the two requirements is generally used.

Location-specific ARARs are those requirements that relate to the geographical or physical position of the site, rather that the nature of contamination or the proposed remedial actions. These requirements may limit the type of remedial
actions that can be implemented, or may impose additional constraints on the cleanup action. Flood plain restrictions and protection of endangered species are among the potential location-specific ARARs.

Action-specific ARARs are requirements that define acceptable treatment and disposal procedures for hazardous substances. These ARARs generally set performance, design, or other similar action-specific controls or restrictions on particular kinds of activities related to management of hazardous substances or pollutants. These requirements are triggered by the particular remedial activities that are selected to accomplish a remedy. Since there are usually several alternative actions for any remedial site, very different requirements may be ARARs. The action-specific requirements do not in themselves determine the remedial alternative; rather, they indicate how or to what level cleanup will be achieved.

State standards and requirements must satisfy five criteria in order to be considered ARARs. The requirements must:

- 1. Be promulgated standards
- 2. Be more stringent than federal requirements
- 3. Be identified by the state in a timely manner
- 4. Not result in a statewide prohibition on land disposal
- 5. Be consistently applied statewide.

Under CERCLA, it is EPA's policy that state ARARs will be achieved to the greatest extent practicable.

4.4.2 Location-Specific ARARs

There are a number of location-specific ARARs that may affect the remedial actions at the King Salmon AFS. Many of these requirements will be verified during the Stage II efforts. Table 4-24 lists the potential location-specific ARARs, along with their prerequisites and comments. The most significant of the potential requirements are the flood plain requirements under RCRA and the Executive Order 11988 Protection of Floodplains, the wetlands requirements under Executive Order 11990 Protection of Wetlands, and the Fish and Wildlife Coordination Act.

Under RCRA, any of the four sites at King Salmon AFS that are located in a 100-year flood plain, and can be classified as hazardous waste disposal sites, will have to be closed in a manner that prevents washout of wastes during a 24-hour, 25-year flood event. Remedial actions at any of the sites

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that are located in a flood plain (100-year or otherwise) will be subject to the Executive Order 11988, Protection of Floodplains. This order requires that actions in flood plains avoid adverse effects, minimize potential harm, and restore and preserve natural and beneficial values.

If any remedial actions affect the wetlands located on King Salmon AFS (the bog), those actions will have to comply with the Executive Order 11990, Protection of Wetlands, which requires action to minimize the destruction, loss, or degradation of wetlands. No dredged or fill materials may be disposed in a wetland without a Clean Water Act 404 Permit from the Army Corps of Engineers.

The Fish and Wildlife Coordination Act requires that the U.S. Department of Fish and Wildlife be consulted regarding any action that modifies a stream or river, or other water of the United States. The intent of this regulation is to protect fish and wildlife that may be adversely affected by changes in the quality or quantity of water in a river, stream, or other water body. This requirement could affect the remedial actions for the Naknek River seep, and the Eskimo Creek seep.

4.4.3 Chemical-Specific ARARs

The potential chemical-specific ARARs for the King Salmon AFS include the Safe Drinking Water Act Maximum Contaminant Levels (MCLs), the RCRA Maximum Concentration Limits (RCRA MCLs), and the Clean Water Act Ambient Water Quality Criteria (AWQC). Evaluation of these potential ARARs cannot be completed with the currently available water quality data. Potential chemical-specific federal and state applicable or relevant and appropriate requirements (ARARs) for particular contaminants identified to date at King Salmon AFS are given in Table 4-24A.

Some of the detection levels for inorganic constituents were above the corresponding MCL or AWQC. This will require confirmation sampling and priority pollutant metals analysis at lower detection levels to verify compliance, or identify noncompliance with ARARS. In addition to the inorganic parameters, there is a need for more specific data on petroleum hydrocarbons. Total petroleum hydrocarbons (TPH), the parameter used in this preliminary investigation, is useful as an indicator of rotential contamination by POLS. ARARS evaluations, however, require quantification of the individual components of the waste. Confirmation sampling and analysis for volatile organics, base/neutral and acids extractable, and polynuclear aromatic hydrocarbons (PAHs) will clarify the nature of the TPH contamination.

| | | | Table 4-24 POTENTIAL LOCATION-SPECIF | IC ARARS | |
|------|--------|---|--|---|--|
| | | Loc.vt ion | Reguirement | Prerequisite(s) | Citation |
| | 1. | Within 61 meters (200 feet) of a fault displaced in Holocene time | New treatment, storage, or dis- posal of hazardous waste prohib- ited | RCRA hazardous waste; treat- ment, storage, or disposal | 40 CFR 264.18(a) |
| | 2. | Within 100-year floodplain | Facility must be designed, con- structed, operated, and main- tained to avoid washout | RCRA hazardous waste; treat- ment, storage, or disposal | 40 CFR 264.18(b) |
| | , m | Within floodplain | Action to avoid adverse effects, minimize potential harm, restore and preserve natural and bene- ficial values | Action that will occur in a floodplain, i.e., lowlands, and relatively flat areas adjoining inland and coastal waters and other flood prone areas | Executive Order 11988, Pro- tection of Floodplains, (40 CFR 6, Appendix A) |
| | 4. | Within salt dome formation, underground mine, or cave | Placement of noncontainerized or bulk liquid hazardous waste pro- hibited | RCRA hazardous waste; placement | 40 CFR 264,18(c) |
| IV-1 | £. | Within area where action may cause irreparable harm, loss, or destruction of significant artifacts | Action to recover and preserve artifacts | Alteration of terrain that threatens significant scien- tific, prehistorical, histori- cal, or archaeological data | National Archaeological and Historical Preservation Act (16 U.S.C. Section 469); 36 CFR Part 65 |
| 32 | 6. | Historic project owned or controlled by federal agency | Action to preserve historic prop- erties; planning of action to minimize harm to National His- toric Landmarks | Property included in or eligi- ble for the National Register of Historic Places | National Historic Preserva- tion Act Section 106 (16 USC 470 et seq.); 36 CFR Part 800 |
| | ٦. | Critical habitat upon which endangered species or threatened species depends | Action to conserve endangered species or threatened species, including consultation with the Department of the Interior | Determination of endangered species or threatened species | Endangered Species Act of 1973 (16 USC 1531 et <u>seq</u> .); 50 CFR Part 200, 50 CFR Part 402 |
| | 8. | Wetland | Action to minimize the destruc- tion, loss, or degradation of wetlands | Wetland as defined by Executive Order 11990 Section 7 | Executive Order 11990, Protection of Wetlands, (40 CFR 6, Appendix A) |
| | | | Action to prohibit discharge of dredged or fill material into wetland without permit | | Clean Water Act Section 404; 40 CFR Parts 230, 231 |
| | •6 | Wilderness area | Area must be administered in such a manner as will leave it unim- paired as wilderness and to pre- serve its wilderness character | Federally owned area designated as wilderness area | Wilderness Act (16 USC 1131 et seg.); 50 CFR 35.1 et seg. |
| | | | | | |

A

| (Continued | |
|------------|--|
| 4-24 | |
| Table | |

| | location | Requirement | Prerequisite(s) | Citation |
|-----|--|--|--|--|
| 10. | Wildlife refuge | Only actions allowed under the provisions of 16 USC Sec- tion 668 dd(c) may be under- taken in areas that are part of the National Wildlife Refuge | Area designated as part of National Wildlife Refuge System | 16 USC 668 dd <u>et seg</u> .; 50 CFR Part 27 |
| 11. | Area affecting stream or river | orseem Action to protect fish or wild- life | Diversion, channeling, or other activity that modifies a stream or river and affects fish or wildlife | Fish and Wildlife Coordina- tion Act [16 U.S.CC. 661 et seq.); 40 CFR 6.302 |
| 12. | Within area affecting national wild, scenic, or recreational river | Avoid taking or assisting in action that will have direct adverse effect on scenic river | Activities that affect or may affect any of the rivers specified in Section 1276(a) | Scenic Rivers Act [16 U.S.C. 1271 et seg. Sec- tion 7(a)); 40 CFR 6.302(e) |
| 13. | Within coastal zone | Conduct activities in manner con- sistent with approved State man- agement programs | Activities affecting the coast- al zone including lands there- under and adjacent shorelands | Coastal Zone Management Act (16 U.S.C. Section 1451 et seq.) |
| 14. | Oceans or waters of the United States | Action to dispose of dredge and fill material into ocean waters is prohibited without a permit | Oceans and waters of the United States | Clean Water Act Section 404 40 CFR 125 Subpart M; Marine Protection Resources and Sanctuary Act Section 103 |
|] | | | | |

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| | Primary SDWA (EPA) Maximum | CWA Ambient Criteria for Freshwater | Water Quality r Protection of Aquatic Life |
|--|----------------------------------|---|--|
| Constituent | Contaminant Levels (MCL) | Acute | Chronic ^a |
| Benzene | 5 ug/l | 5.3 mg/l | |
| Dieldrin | | 2.5 ug/l | 1.9 ng/l |
| Endosulfan | | 220 ng/l | 56 ng/l |
| Ethylbenzene | | 32 mg/l | |
| Heptachlor | | 520 ng/l | 3.8 ng/l |
| Methyl ethyl ketone (2-Butanone) | | | |
| Toluene | | 17.5 mg/1 | |
| Trichloroethylene | 5 ug/l | 45 mg/l | 21 mg/l |
| Xylene(s) | | | |

Table 4-24A POTENTIAL CHEMICAL-SPECIFIC FEDERAL ARARS KING SALMON AFS, ALASKA

^aFederal Ambient Water Quality Criteria, more stringent than an SDWA MCL, may be found relevant and appropriate when there are environmental factors that are being considered at a site, such as protection of aquatic organisms.

Sources: U.S. EPA, <u>Superfund Public Health Evaluation</u> <u>Manual</u>, EPA 540/1-86/060 (OSWER Directove 9285.4-1) October 1986 and U.S. EPA, <u>Quality Criteria for Water 1986</u>, EPA 440/5-86-001, May 1986 (51 Federal Register 43665).

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The contamination in the seeps at the Naknek River and Eskimo Creek may violate AWQC. Resampling of the seeps and analysis for inorganics and organics as discussed above will resolve this issue.

Groundwater contamination, as evidenced by the results of the TPH and benzene-toluene-xylene (BTX) analyses, needs to be evaluated in more detail, as discussed for surface seep samples. The potential ARARs for groundwater include the MCLs, which are applicable to groundwater that could potentially be a source of public drinking water, and the RCRA MCLs, which are standards for groundwater at RCRA hazardous waste treatment, storage or disposal (TSD) sites. The RCRA MCLs are a subset of the drinking water MCLs, and include primarily inorganic constituents.

4.4.4 Action-Specific ARARs

The action-specific ARARs cannot be evaluated effectively until the wastes at the sites are more thoroughly delineated and classified. Action-specific requirements are derived primarily from RCRA, and therefore, are applicable to hazardous wastes. Some nonhazardous wastes may be sufficiently similar to hazardous wastes to cause the RCRA requirements to be relevant and appropriate to a site or situation.

Although the King Salmon AFS is not strictly classified as a licensed RCRA TSD facility, it would be held to the RCRA operating requirements if hazardous wastes were disposed on the station. Disposal is defined by RCRA to include unintentional spills or leaks. Therefore, if any of the material that is being detected at the Naknek River seep, or the Eskimo Creek seep was derived from hazardous waste, RCRA cleanup standards would be applicable to the contaminated groundwater and soil. An investigation into the source(s) of the contamination at the Naknek River seep and the Eskimo Creek seep will be necessary to determine whether the RCRA standards are applicable.

If the results of the confirmation sampling and analysis indicate that the contaminants in either seep are sufficiently similar to characteristic or listed hazardous wastes, the RCRA standards could be considered relevant and appropriate to the remedial actions at the site.

The North Barrel Bluff and the South Barrel Bluff were the sites of disposal of a large number of drums and other debris. It is reported that, while the majority of the drums were empty, some contained liquid wastes. The ARARs for the bluffs are dependent on the contents of the drums, and the presence or absence of contaminants in the groundwater, soil and surface water surrounding and underlying the sites. These sites have not been fully characterized by the preliminary investigation. Additional sampling and analysis is being recommended for these sites, as described in Section VI, Recommendations.

The disposal of liquid or solid hazardous wastes is strictly regulated by RCRA. If any of the drums disposed of at the bluffs contained hazardous wastes, the site would be classified as an unpermitted hazardous waste disposal facility, and would be subject to RCRA closure standards. If the site has resulted in groundwater contamination, it would be subject to RCRA groundwater standards (if the drums contained hazardous waste) or MCLs (if the contamination exceeded the MCLs and the groundwater is a potential source of public drinking water).

The determination of whether the sites contain hazardous wastes could be made by sampling some of the drums to determine what wastes remain at the site. In addition, an investigation of the sources of the drums at the sites, and the operations or processes that generated the wastes would help to determine the appropriate classification. As described for the seep sites, wastes that are very similar to hazardous wastes may cause the hazardous waste disposal and management requirements to be relevant and appropriate to the remedial actions at the bluff sites.

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V. REMEDIAL ALTERNATIVES

The selection of site remedial alternatives usually involves several steps. Initially, general classes or categories of response actions are considered. As more detailed information about the site and the contaminants becomes available, the selection process advances to the consideration of different technologies, and then to identification and evaluation of processes. Each step involves more detailed analyses and is more tailored to the site and the wastes to be managed.

As discussed in Section 4.4, Applicable or Relevant and Appropriate Requirements, more site investigation work will be required to delineate the nature and extent of contamination at King Salmon AFS. The additional characterization will allow determination of which action-specific ARARs may be applicable or relevant to the site, and will influence the selection of remedial alternatives.

5.1 IDENTIFICATION OF GENERAL RESPONSE ACTIONS

General response actions are generic classes of potential remedial actions that can be considered for a given situation. Each of the general response actions can be seen as meeting one or more of the objectives of the overall remedial measure. Within each of the general response actions, there may be several alternative technologies that may be evaluated. Table 5-1 identifies seven general response actions that may be considered for the sites at the King Salmon AFS. A number of alternative remedial technologies are identified for the response actions that will be discussed in more detail in the next section.

These general response actions will be evaluated for each of the four sites at King Salmon AFS during Stage 2 of the IRP. Alternative technologies that are considered unproven, or that incorporate a high degree of risk, were not included in the table. The technologies presented are primarily conventional technologies, which will be evaluated in more detail after collection of the confirmation sampling data.

The first of the general response actions identified in Table 5-1 is No Action. Under the CERCLA process the No-Action alternative is carried through the entire selection process. The No-Action alternative serves as a basis for comparison of the remaining alternatives. It represents the existing situation, and is evaluated for risk and comparison with ARARs, along with the other alternatives.

Institutional Controls serve to prevent public access to the site or the contaminants, but it does not attempt to remove or modify the contamination. This alternative is typically

TABLE 5-1

IDENTIFICATION OF GENERAL

GENERAL RESPONSE ACTION

REMEDIAL TECHNOLOGY

DESCRIPTION







NO ACTION; REMEDIAL RES

EXAMPLES INCLUDE LEGAL FENCES 10 REGULATE SITE

ALTERNATIVE WATER SOURC WATER SYSTEM, ETC.

RESIDENTS ARE RELOCATE RESIDENCE) BASIS.

IMPERMEABLE COVER MAIL INFILTRATION AND PREVEN

WALLS THAT ARE LESS PER MAY BE USED TO ENCIRCU

SURFACE SEALING, GRADIN TO REDUCE SITE RUNON/R

A PHYSICAL BARRIER, FIL[®] REMOVAL.

TARPS, SOIL COVERS, St

DRUMS AND/OR DEBRIS AN EQUIPMENT.

CONTAMINATED SOIL, SEDA

GROUNDWATER AND LEACH

CONTAMINANTS THAT ARE

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NERAL RESPONSE ACTIONS

?TION

REMEDIAL RESPONSES INITIATED AT SITE ARE ABANDONED; NO MONITORING.

NGLUDE LEGAL RESTRICTIONS ON DEEDS TO REGULATE LAND USE; PERMITS TO REGULATE GROUNDWATER USE; REGULATE SITE ACCESS.

'E WATER SOURCES MIGHT INCLUDE CISTERNS, ABOVEGROUND TANKS, DEEPER OR UPGRADIENT WELLS, MUNICIPAL TEM, ETC.

ARE RELOCATED ON EITHER A TEMPORARY (APARTMENT OR MOTEL) OR A PERMANENT (PURCHASE OF BASIS.

LE COVER MATERIALS ARE PLACED OVER CONTAMINATED SOILS AND LANDFILLS TO REDUCE OR ELIMINATE IN AND PREVENT EROSION.

T ARE LESS PERMEABLE THAN THE IN SITU SOIL ARE PLACED UPGRADIENT OF WASTE, OR DOWNGRADIENT OF WASTE, OR ED TO ENCIRCLE WASTES.

EALING, GRADING, SOIL STABILIZATION, REVEGETATION, AND DIVERSION/COLLECTON PROCESSES ARE IMPLEMENTED SITE RUNON/RUNOFF, SURFACE WATER INFILTRATION AND EROSION, AND TO STABILIZE SURFACE SOILS.

BARRIER, FILTER SCREEN, OR CAP IS CONSTRUCTED TO MINIMIZE THE SPREAD OF CONTAMINANTS DURING SEDIMENT

L COVERS, SPRAYS, OILS, ETC. ARE APPLIED TO SUPPRESS DUST AND NON-POINT-SOURCE VAPOR.

J/OR DEBRIS ARE REMOVED BY VARIOUS MECHANICAL MEANS, E.G., CRANES AND OTHER STANDARD CONSTRUCTION

VIED SOIL, SEDIMENT, SLUDGES AND OTHER SOLIDS ARE REMOVED WITH STANDARD CONSTRUCTION EQUIPMENT.

IFR AND LEACHATE ARE COLLECTED IN WELLS OR DRAINS AND PUMPED SO THEY CAN BE TREATED AND DISPOSED.

INTS THAT ARE PHYSICALLY OR CHEMICALLY BOUND TO SOILS ARE FLUSHED OR PUMPED FROM THE SOILS.

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IDENTIFICATION OF GENERAL

GENERAL RESPONSE ACTION REMEDIAL TECHNOLOGY

DESCRIPTION



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ERAL RESPONSE ACTIONS (CONT.)

PTION

MENTS, SLUDGES, AND OTHER SOLIDS ARE PHYSICALLY OR CHEMICALLY TREATED TO REMOVE THE HAZARDOUS 'TS OR CONVERT THE CONSTITUENTS TO NONHAZARDOUS SUBSTANCES.

ON AND FIXATION PROCESSES MAINTAIN HAZARDOUS CONSTITUENTS IN THEIR LEAST TOXIC AND/OR LEAST SOLUBLE IFICATION PRODUCES A MONOLITHIC BLOCK OF TREATED WASTE WITH HIGH STRUCTURAL INTEGRITY.

D SEPARATION CLARIFIES LIQUID STREAMS, RECOVERS SOLIDS AND LIQUIDS, OR REMOVES EXCESS LIQUID FROM OR TO DISPOSAL.

; LIQUIDS ARE TREATED TO CONVERT THE HAZARDOUS CONSTITUENTS TO A MORE EASILY HANDLED FORM THROUGH ON, CONCENTRATION, AND/OR PHASE CHANGE.

; LIQUIDS ARE ALTERED BY CHEMICAL REACTIONS TO DETOXIFY HAZARDOUS COMPOUNDS OR TO CONVERT THE 3 TO A MORE EASILY TREATED FORM.

OF MICROORGANISMS METABOLIZES BIODECRADABLE ORGANIC COMPOUNDS.

THEMICAL, AND BIOLOGICAL PROCESSES ARE EMPLOYED TO TREAT CONTAMINATED SOIL, SOLID WASTES OR TER IN PLACE.

ERIAL IS EXPOSED TO HIGH TEMPERATURES TO TRANSFORM THE HAZARDOUS COMPOUNDS INTO OR LESS HARMFUL SUBSTANCES.

SE IMPURITIES ARE REMOVED FROM GAS STREAMS THROUGH PROCESSES RANGING FROM ONCE-THROUGH ATIONS TO MULTIPLE-STEP RECYCLE SYSTEMS.

E MATERIALS CAN BE CONVERTED TO ENERGY

5 MATERIAL IS TEMPORARILY STORED IN SURFACE IMPOUNDMENTS, WASTE PILES, OR CONTAINERS/TANKS.

SOLID WASTES THAT ARE PRIMARILY ORGANIC ARE INCORPORATED INTO THE UPPER SOIL HORIZON SO THEY CAN BE TRANSFORMED, OR IMMOBILIZED.

TREAMS ARE DISCHARGED TO A POTW, SURFACE WATER, OR SHALLOW OR DEEP WELLS.

POUNDMENTS ARE USED TO CONTAIN TREATED OR UNTREATED WASTEWATER OR GROUNDWATER UNTIL IT S.

ES ARE PERMANENTLY DISPOSED OF IN A LANDFILL. LANDFILLS CANNOT ACCEPT LIQUID WASTES.

/OR LONG-TERM MONITORING IS IMPLEMENTED TO RECORD SITE CONDITIONS AND CONTAMINATION LEVELS.

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selected when contamination is widespread and difficult or impossible to remove.

Containment actions serve to isolate the contamination and prevent it from migrating offsite or into currently uncontaminated areas or media. Containment is often selected when the volume of waste is too large to permit removal and treatment, or when there is no practicable treatment for the waste. Containment options require ongoing monitoring to ensure the effectiveness of the containment system.

Removal actions involve the excavation, extraction, or physical removal of wastes and contaminated media for treatment or disposal. Groundwater extraction and treatment systems are considered removal actions. Removal actions may result in clean closure of a site, with no requirements for ongoing monitoring or maintenance.

Treatment actions may involve some sort of removal process, or may be in-situ operations. The goal of treatment actions is to reduce the toxicity, mobility, or volume of wastes requiring final disposal. Treatment processes may generate a residue or waste stream, which would also require treatment and disposal. The type of treatment selected is dependent on the characteristics and volume of the waste to be treated.

Disposal actions may occur onsite or offsite, and may involve treated or untreated wastes and contaminated materials. Some disposal actions may also fall into other general response action categories. For example, landfilling is both a containment action and a disposal action.

5.2 IDENTIFICATION AND SCREENING OF TECHNOLOGIES

Table 5-2 is a screening table for remedial technologies. For each of the general response actions discussed in Section 5.1, a number of remedial technologies were identified. In this table, each of the technologies are further defined by a group of potential process options. A brief description is included for each of the process options. Other process options currently considered experimental or unproven, or that incorporate a high degree of risk, have not been included in the table.

During Stage 2 of the IRP, data from the confirmation sampling will be used to evaluate the general response actions and remedial technologies. Process options will then be evaluated in order to identify alternative remedial measures for each of the four sites. The remedial technologies and process options will be evaluated on the bases of feasibility, implementability, effectiveness, permanence, and cost.

TABLE 5-2

REMEDIAL TECHNOLOGY SCREENING





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GENERAL RESPONSE

ACTIONS

DESCRIPTION PROCESS OPTION REMEDIAL TECHNOLOGY

Other Chemical Modifications Flow and Strength Equalization Dewatering and Drying Beds Coogulation /Flocculation Screens, Hydrouilc Closeifiers, and Scolpers **Chemical Reduction** Solvent Evaporation Beit Filter Presses Pozzolunic Agente **Gravity Thickening** Vacuum F#tration Solvent Leaching Water Leaching Sedimentation Filter Presses Neutralization Encopeulation Sludge Dryers Centrifuges Oxidation Sorption Solicitication, Fixation, Stobilization Physical Treatment Solide Treatment Solids Dewotering J: \INDUST\K24173\5-2-5 Treatment

Acid added to an akaine waste ar base added to an acidic waste to adjust the pH.

Dxidizing agents added to waste for oxidation of organic contaminants to less toxic axidation

Reduction agents added to wastes to stabilize metals by converting them to a leve aduple, more stable form.

Contaminated solids are treated with appropriate reagents under suitable conditions to convert the contaminants into less toxic forms.

Mater Infiltrates conteminated solid, leaching out water-soluble contemination.

Solvent infitrates contaminated solid, leaching but contaminants that are soluble in the selected

Contaminated sludge or soll heated in equipment which allows mixing. Organics volatilized from solds into the gas stream.

In inst. nondegradable, nonreactive sorbent add to itquid or sludge to sock up moleture and sroduce a soli-like material.

Lms and fine-grained allocous material added to outecus were to produce a contractile addi-diversal is encopsuited by either costing the dividual practices or enclosing artite mass in a polated of meet material. Applicable for extremely dangerous or radioactive wasted.

uspended solids removed from liquid by gravity in tonk or lagoon. Sadimentation aften preceded by hemical precipitation and/or by coogulation and occulation.

later removed from eludge by gravity in thickening anks. Chemicals can aid in sludge dewatering.

Sudge dewotered by centrifugal force. Centrifuge are used when space is limited or where sludge scineration/disposal is required.

Coarse and/or gross solids removed from raw wastewater by a screening eurlace in a classifie

Sudga dewatered by on andless fliter belt that uns own a drive and guide roller of each and. A reas belt runs in the same drection and speed bove the fliter belt.

Sudge passed into press under pressure, woter passes through filter cloth while solids form a cake on the filter cloth surface.

A partially submarged cylindrical drum rotatas in a vot or pan of conditional sudda. A vacuum is builed inside the drum, solida collect on outside and are continuously ecraped off.

Sudge placed on sand layered over gravel. Water in the sludge evaporates and drame.

Contaminated sludge or soll heated in equipment which allows mixing. Aqueous liquids evaporated from solids into the gas stream.

Equalization basin requiates flow through a treatment facility. Mechanical mixing equalizes concentration.

Suspanded porticle aggregation that results from semical additions to the watewater. Once particles have aggregated the large porticles, threation, floration, floration, x fliration.





GENERAL RESPONSE

ACTIONS

PROCESS OPTION REMEDIAL TECHNOLOGY



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J: \INDUST\K24173\5-2-9

Selection of process options is one of the last steps in the development of a remedial measure. The selection must incorporate detailed information on the characterization and quantity of waste involved, the treatment standards or other restrictions identified by the ARARs analysis, and site specific details such as space availability, climate, etc.

5.3 DEVELOPMENT OF REMEDIAL ALTERNATIVES

Remedial alternatives are developed following the evaluation of general response actions, remedial technologies, and process options. Typically, several alternatives are developed for each site that represent a range of cleanup levels.

The data obtained during Stage 1 of the IRP primarily served to identify areas of contamination and to indicate the types of contaminants present. The confirmation sampling and analysis to be done during Stage 2 will attempt to delineate the sources of the contamination at each site, and to identify the constituents of the wastes. This information is essential to the identification of ARARs, and to the evaluation of appropriate treatment technologies. Therefore, the development of remedial alternatives will be done as part of Stage 2.

The following discussion outlines a range of general remedial alternatives for each site. The purpose of developing general alternatives at this time is to aid in the identification of additional data requirements to be satisfied during Stage 2. For example, if a removal action is considered, it is important to be able to estimate the total volume of material to be removed. This would involve determination of the extent of contamination, which could be done during Stage 2.

The general alternatives described in this section do not reflect site-specific data or conditions. The alternatives were derived from experience at similar sites, and incorporate general response actions and remedial technologies from Tables 5-1 and 5-2.

5.3.1 North Barrel Bluff and South Barrel Bluff

The following general remedial alternatives were developed for the North Barrel Bluff and the South Barrel Bluff. The similarities of the two sites makes the dévelopment of separate alternatives impractical at this time. After additional sampling and analysis during Stage 2, sufficient data will be available to allow screening of general response actions, remedial technologies, and process options for each site. The alternatives are ordered by increasing level of protection, which usually corresponds to increasing level of cost and complexity.

5.3.1.1 No Action Alternative

The No-Action alternative is carried through the evaluation and screening process as a basis for comparison of other alternatives. At the North and South Barrel Bluff sites, no action would allow any existing contamination of the groundwater or surface water to continue. The wastes that may be contained in the drums would remain at the sites. In the future, the drums can be expected to fail due to exposure to the weather. Wastes could be expected to leak from the drums and contaminate surrounding and underlying soils. Groundwater or surface water contamination could also result.

Current levels of human and environmental exposure to the wastes at the sites would be expected to continue. Future increases or decreases in habitation or use of the area would change the exposure and risk levels.

5.3.1.2 Institutional Controls

Institutional controls for the North and South Barrel Bluff sites would include restriction of access to the area, and possibly a moratorium on groundwater use (in the event that the groundwater is determined to be contaminated). As discussed for the No-Action alternative, this would not result in the removal of any waste, and would not prevent migration of the wastes from the site. Institutional controls would result in some decrease in the level of human exposure and risk. No change would occur in environmental exposure or risk.

5.3.1.3 Containment

The goal of a containment action is to prevent migration of wastes or waste components from the site. Containment of the wastes and contaminated media at the North and South Barrel Bluff sites would include one or more of the following: capping, surface water controls, and groundwater barriers. The type of cap to be installed, and the treatment or disposal requirements for wastes, drums, and contaminated soils would be dependent on the classification and characterization of the wastes. If the wastes at the sites are hazardous wastes, RCRA capping, waste management, and disposal requirements would be applicable.

Capping of the site would prevent direct contact between humans or wildlife and the site wastes. Capping of a site containing a large number of barrels is exceedingly difficult due to the potential for subsidence and cap failure. It would probably be necessary to remove or crush a number of drums to achieve a surface contour that would sustain capping. Treatment or disposal of the wastes and drums removed from the site would be a separate concern. Surface water control would be a requirement of the capping program, in order to prevent damage to the cap during storm events. Any contaminated surface soils not covered by the cap would have to be isolated. The surface runoff from these areas would require containment and possible treatment. The surface water control system would be designed to minimize the quantity of storm water requiring containment and treatment by directing offsite storm water around the site, and thereby preventing run-on.

Containment of contaminated groundwater, if necessary, would require extensive site work. Containment of the shallow aquifer could be accomplished by construction of a slurry wall or grout curtain. In order to be effective, the wall or curtain would have to be keyed into an impermeable strata below the shallow aquifer. Local site conditions may make this option impossible.

In the event that contaminated soil is excavated during the construction of the wall or curtain, treatment, or disposal of the soil would be necessary. The appropriate level of treatment would be defined by the classification and characterization of the wastes. Containment of groundwater does not remove contaminants from offsite groundwater. Some risk to offsite users of the shallow groundwater would remain.

5.3.1.4 Removal of Wastes

Removal alternatives offer the benefit of reduced risk and exposure level following the remedial action. During the action, however, the risk to site workers is usually higher than the risk during construction of containment alternatives.

Removal of the wastes from the North and South Barrel Bluffs would be complicated by the condition of the drums. Some of the drums are reported to contain liquid wastes. The source and identity of these wastes is not known. A number of the drums have probably corroded or have been ruptured by the disposal method. These drums would be hard to handle and would present an unknown risk to site workers. Drums containing liquid or solid waste would have to be sampled, and would probably require overpacking to secure the contents. Treatment or disposal of these wastes could trigger RCRA hazardous waste management requirements.

The soil surrounding and underlying the sites is probably contaminated to some depth by the former contents of the drums. Total removal of the contamination would require excavation of the contaminated soil, followed by treatment or disposal. During implementation of the removal alternative, provisions would have to be made to collect, contain, and treat any storm water runoff that contacts the area of the contamination. As discussed under the containment alternative, the storm water control system would be designed to prevent run-on, and thus minimize the quantity of storm water requiring treatment and disposal.

Groundwater removal is most often accomplished using extraction wells. If the groundwater surrounding or underlying the sites is contaminated, extraction of the contaminated plume, followed by treatment and disposal, would be included in the removal alternative. The level of treatment and the method of disposal would be determined on the basis of evaluations of the waste constituents and concentrations in the groundwater, and the available disposal methods.

In the event that a removal alternative results in the complete removal of all wastes, contaminated soils, and contaminated groundwater, the site would be considered to have undergone clean closure. No further monitoring or maintenance of the site would be required.

Variations in a removal action include the quantities of waste removed, or the level of cleanup, and the treatment or disposal alternatives incorporated in the alternative. Removal with offsite disposal may be appropriate for some wastes. Disposal can range from landfilling at a municipal landfill, to landfilling at a licensed RCRA landfill, to disposal in a licensed RCRA incinerator. The selection of disposal technique will depend on the type and quantity of the waste. Some wastes may be amenable to treatment prior to disposal. If the wastes can be rendered nonhazardous by the treatment, lower cost disposal options may be considered.

5.3.2 Naknek River Seep and Eskimo Creek Seep

The following general remedial alternatives were developed for the Naknek River seep and the Eskimo Creek seep. The similarities of the two sites makes the development of separate alternatives impractical at this time. After additional sampling and analysis during Stage 2, sufficient data will be available to allow screening of general response actions, remedial technologies, and process options for each site. The alternatives are ordered by increasing level of protection, which usually corresponds to increasing level of cost and complexity.

5.3.2.1 No Action Alternative

The No-Action alternative is carried through the evaluation and screening process as a basis for comparison of other alternatives. At the Naknek River seep and the Eskimo Creek

seep, no action would allow contamination of the groundwater and surface water to continue. There would be no impact on the sources of the seeps, and current levels of human and environmental exposure to the contaminants at the sites would continue. Future increases or decreases in habitation or use of the area would change the exposure and risk levels.

5.3.2.2 Institutional Controls

Institutional controls for the Naknek River seep and the Eskimo Creek seep would include restriction of access to the area, and a possible moratorium on use of the Naknek River and Eskimo Creek for fishing and recreational use in the area immediately downstream of the seep areas. As discussed for the No-Action alternative, this would not result in the removal or containment of the source of the contamination, and would not prevent continued release of the contaminants to the Naknek River and Eskimo Creek. Institutional controls would result in some decrease in the level of human exposure and risk. No change would occur in environmental exposure or risk.

5.3.2.3 Containment

Containment of the wastes and contaminated media at the Naknek River seep and Eskimo Creek seep would include one or more of the following: groundwater barriers, some capping, and surface water controls to protect the capped area.

In order to contain the contaminants being released at the seeps, the source of each of the seeps would have to be determined. It is suspected that the seeps resulted from the leaking or spilling of POL over a period of time. The sources of the seeps could be extensive areas of contaminated soil in contact with the shallow groundwater. Containment of a large area would require slurry wall construction to isolate the contaminated soil and groundwater, and capping to prevent infiltration of precipitation and surface runoff, which would add to the volume of contaminated groundwater.

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Containment of the seep areas themselves would be difficult due to the presence of flowing water. Excavation of some of the contaminated soil could be required, followed by construction of a channel to prevent contact of the river or creek with the contamination.

Surface water controls would be necessary in the area of the cap to protect the cap from storm water damage. The surface water controls would be designed to minimize the quantity of water flowing over the cap by diverting water around the cap and preventing run-on. If the contaminants at the sites are classified as hazardous wastes, the cap and the surface water control system would have to comply with RCRA requirements. Capping and containment of the sites would prevent direct contact of humans or wildlife with the site contaminants. Ongoing monitoring and maintenance of the sites would be required, since the contamination would remain onsite.

5.3.2.4 Removal

Removal alternatives offer the benefit of reduced risk and exposure level following the remedial action. During the action, however, the risk to site workers is usually higher than the risk during construction of containment alternatives.

Removal of the wastes from the Naknek River seep and Eskimo Creek seep would be complicated by the nature of the suspected sources of the seeps. However, compared to the problems associated with containment of the contaminants at the seep site, a removal alternative would probably be preferred.

Assuming that the current sources of the seeps are large areas of contaminated soil in contact with shallow groundwater, two types of removal alternatives are possible. The first alternative would be to remove the entire source of contamination. This would involve excavation and removal of the soil in the area of contamination. Since much of the soil is below the water table, a significant quantity of dewatering would be required during excavation. The contaminated groundwater, as well as the excavated soil, would have to be contained for treatment or disposal.

Obviously, for all but the smallest areas of contamination, this alternative would be difficult and costly to implement.

Another alternative would involve the removal of the contaminated groundwater without excavation of the soils. Extraction wells could be constructed surrounding the area of contamination. By withdrawing groundwater in the contaminated zone, contaminants could be prevented from entering the river or creek.

The contaminants in the soil would be leached into the groundwater over a period of time. The construction of groundwater barriers upgradient of the contaminated area would limit the quantity of groundwater to be treated, but would extend the duration of treatment. Enhanced leaching of the contaminants from the soil could be accomplished by flushing the soil with water, which would be collected in the extraction wells. Soil vapor extraction could also be used to remove volatile organic contaminants from the soil.

All of these removal alternatives would require long-term operation, monitoring, and maintenance. Human and environmental exposure and risk would be reduced but not eliminated at the site. It would be difficult to project the length of treatment time required to remove the source of the contaminants, or the lowest attainable concentrations in the soil or groundwater.

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VI. RECOMMENDATIONS

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6.1 DIRECTION AND APPROACH FOR FUTURE WORK

6.1.1 Eskimo Creek Seep

Additional investigative efforts at the Eskimo Creek seep should be directed towards the following:

- Expanding the analyses performed on groundwater samples to define the constituents that contribute to the TPH results, especially for the base drinking water wells. Risk and ARARs analyses should be refined based on the new data.
- Attempting to identify the source of the Eskimo Creek seep and other groundwater contamination.
- Defining the vertical and lateral extent of the hydrocarbon plume moving to the west away from the POL tanks.

Recommendations for addressing these concerns include the following tasks:

- Review historical site-specific practices including facility operations, tank locations and storage records, drum storage or disposal records, and contents. Task includes additional focused records search, aerial photography analyses, interviews, and field reconnaissance.
- Expand the analyses of base drinking water from KS-4 and KS-5 to identify and quantify individual constituents from TPH results (priority pollutants).
- Consider deep well rehabilitation or abandonment if the contamination is found to be the result of interaquifer communication along the well bore hole.
- Collect additional samples and expand analyses of groundwater from shallow monitoring wells should be performed to characterize the individual constituents. Samples from wells MW-39, MW-32, MW-30, MW-38, MW-34, and MW-9 would provide a distribution between the POL tanks and Eskimo Creek seep that would indicate a "worst case" for potential risk analyses. Analytical results would also provide a basis for determining a "target" list of

constituents for additional groundwater, surface water and sediment samples.

- Sample and analyze Eskimo Creek soils between the seep and Eskimo Creek for a target list of potential contaminants. Sample locations and depths should be adequate (10 locations) to characterize this potential groundwater discharge area.
- o Expand sampling and sample analyses of Eskimo Creek surface water to provide a broader data base and allow a comprehensive assessment of the human and environmental risks this area presents.
- Include sampling of background conditions for each media (soil, surface water, and groundwater) to provide input for risk analyses.
- o Perform seasonal water level measurements and groundwater flow system interpretations on a quarterly basis to provide a refinement of the current hydrologic model. Lowering of water levels could have a major affect on groundwater flow directions in the area of the till mound.
- Install three to five additional groundwater monitoring wells at locations shown in Figure 6-1. Rationale for each well is provided in Table 6-1. Sample all wells for target list of constituents. Wells should be resampled after 3 to 6 months for verification of analyses and investigation of seasonal effects.
- Develop each well adequately to provide representative groundwater samples. A single well slug/ bail or recovery test should be performed to provide an estimate of aquifer parameters.
- Due to the difficult sampling conditions, update the sampling and analysis plan based on actual site sampling experience to provide quality assurance in future field techniques.
- Conduct a shallow aquifer pumping test to provide a more accurate estimate of aquifer parameters if remediation includes a pump-and-treat scenario. This test could be conducted during a predesign stage of remedial activities.

6.1.2 Naknek Seep

Hydrocarbon contamination was detected at the Naknek seep. In addition, two adjacent areas with potential hydrocarbon





FIGURE 6-1

PROPOSED MONITORING WELL LOCATIONS KING SALMON AIR FORCE STATION

KING SALMON, ALASKA

Table 6-1 RATIONALE FOR ADDITIONAL SHALLOW MONITORING WELLS

| Well Number | Rationale |
|----------------|--|
| A | Extent of west plumegroundwater quality Water level data Till surface elevation |
| В | Extent of west plumegroundwater quality Water level data Till surface elevation |
| C | Extent of major plume Water level data Till surface elevation at critical area to determine flow path from POL tanks towards Eskimo Creek seep |
| D | Water quality upgradient of Eskimo Creek seep and within soil gas plume Water level data Till surface elevation |
| Е | Water quality data upgradient of Eskimo Creek seep and along discharge route to Eskimo Creek Water level data Till surface elevation |
| F | Background water quality Upgradient water level data |

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were identified. One area includes buried and partially buried drums 1,500 feet northwest of the Naknek seep. The other area includes a POL seep near the large POL tanks, located 1,500 feet north of the Eskimo Seep.

The detection of aromatic and halogenated volatile organic constituents at the site suggests a potential for adverse public health and environmental effects. Additional investigations recommended to evaluate this potential at the three sites are outlined below:

- Review historical site-specific practices including facility operations, tank locations and storage records, drum storage or disposal records, and contents. Task includes additional focused records search, aerial photography analyses, interviews, and field reconnaissance.
- Perform additional soil gas survey work to assist in delineating the lateral extent of the hydrocarbon plume(s) and to aid in placement of shallow monitoring wells.
- Perform additional reconnaissance geophysical surveys to delineate the extent of drums or buried tanks.
- Drill three to five shallow soil borings at each site (depending on soil gas results) with vertical sampling of soils for chemical analysis.
- Install three to five shallow monitoring wells (in soil borings) to allow hydraulic and chemical analyses of the shallow aquifer.
- Develop each well adequately to provide representative groundwater samples. A single well slug/ bail or recovery test should be performed to provide an estimate of aquifer parameters.
- Complete two groundwater measurement and sampling events at a 3-month interval to characterize the flow system and confirm the presence of hazardous constituents.
- Collect and sample seepage fluids along Naknek
 River for chemical analyses. Background water
 quality should be established by a sample upstream
 of the seep.
- Analyze soil and water samples for an expanded constituent list (Priority Pollutants) to identify and quantify potential constituents. Data quality objectives, including detection limits, need to be adequate to determine if concentrations comply with water quality criteria for the protection of

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aquatic organisms. Background water quality must be established for correlation with risk analyses data.

6.1.3 North Barrel Bluff

The North Barrel Bluff poses a potential human health and environmental hazard resulting from physical and chemical exposures. Installation of a locking gate and perimeter fence is recommended to reduce this potential and to establish a buffer zone. Further investigations are recommended to better evaluate human health and environmental consequences associated with this debris area. These investigations should include:

- Review historical site-specific practices including facility operations, tank locations and storage records, drum storage or disposal records, and contents. Task includes additional focused records search, aerial photography analyses, interviews, and field reconnaissance.
- o Collect soil samples from the area beneath the toe of the bluff. Ten shallow borings should be completed 10 to 15 feet into the soils along the bluff to indicate whether leakage of residual fluids from drums has entered into the soils. Piezometer casing should be installed in the boreholes to provide sampling points for water quality and water level measurements.
- Install six shallow aquifer monitoring wells along 0 the bluff and adjacent to the drums (based on boundaries established in the geophysical survey) to obtain samples for soils and groundwater characterization. Four of the wells should be located within the landfill and two placed as background (upgradient water quality and hydraulic data). The wells should be sampled for the priority pollutants as well as any other target constituents identified in the historical review. Groundwater sampling results should be verified with a second round of sampling the following guarter. Water levels should be monitored seasonally for 1 year (four guarters) to determine shallow aguifer conditions.
- o Investigate the intermediate aquifer for direction of groundwater flow and presence of contaminants if groundwater contamination is found in the shallow aquifer. This would initially include three intermediate wells; two would be located within the landfill and one as a background. Sampling should follow the same schedule as the shallow wells.

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- Develop each well adequately to provide representative groundwater samples. A single well slug/ bail or recovery test should be performed to provide an estimate of aquifer parameters.
- Include sampling of background conditions for each media (surface water, soils, and groundwater) to provide correlation with risk analyses data.
- Conduct four shallow soil borings near the White Alice building. Samples should be analyzed for priority pollutants.
- O Use additional reconnaissance geophysical profiling to identify small "dump" areas that are apparently common within the disturbed ground along the top of the bluffs. Any future remedial actions that include a delineation of the extent of the drums along the bluff will require extensive geophysical surveys to fill in the existing reconnaissance grid.
- o Conduct backhoe explorations at small "dump" areas identified in the previous (and any future) geophysical surveys to provide soil samples for chemical analyses. Analyses should be a target list (subset of priority pollutants) that reflects the type of debris found in the excavation.

6.1.4 South Barrel Bluff

Drum and other debris types present at the South Barrel Bluff present similar human health and environmental hazards to those present at North Barrel Bluff. The South Barrel Bluff presents an additional problem because of its proximity to residential housing. In two instances, housing has been located near to or on top of fill material placed over the debris area. Installation of a locking gate and fence that encompasses the known perimeter of the debris area, which is not presently inhabited, is recommended to assist in reducing the potential for human and wildlife contact with the deteriorating drums and debris.

To adequately evaluate the potential hazards to human health and the environment associated with this area, the following are suggested:

 Review historical site-specific practices, including facility operations, tank locations and storage records, drum storage or disposal records, and contents. Task includes additional focused records search, aerial photography analyses, interviews, and field reconnaissance.
- O Collect soil and leachate samples from beneath the toe of the bluff. Ten shallow auger borings 10 to 15 feet into the soil horizon may provide an indication whether residual liquids that have leaked from deteriorating drums have entered into bluff soils or into King Salmon Creek. Piezometer casing should be installed in the boreholes to provide sampling points for water quality and water level measurements.
- o Install five shallow aquifer monitoring wells along the bluff adjacent to the drums and within the landfill to obtain samples for soils and groundwater characterization. Groundwater analyses for the shallow wells and piezometers should include priority pollutant analyses. Sampling results should be verified with a second sampling round the following quarter. Water levels should be monitored quarterly for 1 year to determine shallow aquifer conditions.
- o If groundwater contamination is found in the shallow aquifer, investigate the intermediate aquifer for direction of groundwater flow and presence of contaminants. This would initially include three intermediate wells; two would be located within the landfill and one as a background. Sampling should follow the same schedule as the shallow wells.
- Include sampling of background conditions for each media (surface water, soils, and groundwater) to provide correlation with risks analyses data.
- Develop each well adequately to provide representative groundwater samples. A single well slug/ bail or recovery test should be performed to provide an estimate of aquifer parameters.
- Use additional reconnaissance geophysical profiling to identify small "dump" areas that are apparently common within the disturbed ground along the bluffs. Any future remedial actions to be conducted at the South Barrel Bluff will require a detailed delineation of the lateral extent of buried debris. This determination can be made by completing additional geophysical surveys along the bluff.
- Conduct backhoe explorations at small "dump" areas identified in previous geophysical surveys to provide soil samples for chemical analysis.

 Include residential wells in the area into a periodic, extended groundwater monitoring schedule and/or remove them as drinking water supplies.
Wells located within landfill boundaries (two identified) should be removed and alternate water supplies secured. Potential health risks for residents along the bluff should be evaluated and future use of the area controlled. Land use controls may include removal of the two houses located on the landfill.

6.2 CATEGORIZATION OF SITES

At the conclusion of the Stage 1 process, the Department of Defense IRP divides sites into three categories. Category 1 includes sites where no further IRP action is required. In order for a site to be included in Category 1, the data gathered during Stage 1 must be sufficient to conclude that no significant threat to human health or the environment exists at the site. This may be appropriate for sites that are fully characterized and are in attainment with environmental standards. Sites at which access to contaminants is very limited and natural biological processes are expected to degrade the contaminants without producing a threat to human health or the environment are also included in Category 1.

Category 2 includes sites requiring additional IRP effort, such as additional sampling and analysis to complete characterization of contaminants. Sites at which risk assessments have not been completed belong in Category 2, as well as sites for which an evaluation of remedial alternatives is incomplete.

Category 3 includes sites for which the feasibility study is complete and for which a remedial alternative has been recommended.

All four of the King Salmon AFS sites included in this report-the North Barrel Bluff, the South Barrel Bluff, the Naknek River seep, and the Eskimo Creek seep--have been determined to be classified in Category 2.

6.3 METHODOLOGY FOR ABANDONMENT OF TEST AND WATER WELLS

6.3.1 Introduction

Wells that are no longer in use or are in a state of physical deterioration need to be carefully sealed to:

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 Prevent contamination of the groundwater source by contaminants entering the well from the ground surface

- o Eliminate physical hazards
- o Conserve aquifer yield
- o Maintain confined head conditions
- Minimize the potential for contaminated groundwater from one aquifer to enter another aquifer

The abandonment procedure should restore as far as possible the hydrogeologic conditions that existed prior to the installation of the well.

Well abandonment procedures are specific to the type of well being abandoned. There are four known well types at King Salmon AFS currently being used for the Eskimo Creek seep groundwater investigation. Prior to the abandonment of any wells, written approval of the abandonment procedure should be obtained from the Director of the Alaska Department of Environmental Conservation. The four known well types at King Salmon AFS are:

- O Cased wells under unconfined conditions (MW-1, 6, 9, 10, 16, 18, 21, and 22)
- O Cased wells under confined conditions (KS-4 and KS-5)
- Gravel/sand filter pack wells under unconfined conditions (MW-23 through MW-40)
- Gravel/sand filter pack wells under confined conditions (MW-41 through MW-43)

Abandonment procedures specific to each well type are discussed in the following subsections.

6.3.2 Cased Wells Under Unconfined Conditions

The geologic log for USCOE wells MW-1, MW-9, MW-10, MW-16, MW-18, MW-21, and MW-22 did not indicate the placement of a sand or gravel pack around the well casing. The well casing consists of 1-1/2-inch diameter PVC with glued, coupled joints. This type of construction is generally not strong enough to allow extraction of the well casing. Additionally, these wells have been constructed in areas where groundwater contamination is known to exist. The abandonment procedure may consist of the following.

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- 1. The well is measured to record the static water level in the well and the total depth. The measured depth is compared with the construction depth to ascertain whether there are obstructions that may interfere with placement of sealing materials. Obstructions are removed. The volume of the well casing should be calculated to assure that sufficient sealing material is emplaced.
- 2. The well casing should be pressure grouted throughout the perforated section. The sealing materials should include cement grout, concrete, puddling clay or bentonite. Sealing materials placed below the static water level in the well shall be placed from the bottom up by methods that avoid segregation or dilution of material. When a tremie pipe is used to place the sealing material, the discharge end of the tremie pipe should be submerged in the grout to avoid breaking the seal while filling the annular space. The remainder of the well casing shall be filled in a similar manner.
- 3. The protective casing and guard post should be extracted and the exposed portion of PVC casing cut off at ground level.
- 4. A cement or concrete patch should be constructed over the surface exposure of the well.

6.3.3 Cased Wells Under Confined Conditions

Geologic logs for base production wells KS-4 and KS-5 indicate that these wells are cased wells that may have been constructed with a telescope well screen. The abandonment procedure for this well type consists of the following steps.

- 1. The well is measured to record the static water level in the well and the total depth. The measured depth is compared with the construction depth to ascertain whether there are obstructions that may interfere with placement of sealing materials. Obstructions are removed.
- 2. If the well construction method included the placement of a telescoped well screen, removal of the well screen is recommended. This is performed using a method called sand locking. A pipe of smaller diameter, generally one-half the diameter of the well screen, is lowered inside of the well screen. 'Angular sand particles are then placed in the annular space between the pulling pipe and the well screen. The sand particles form the sand joint that serves as a structural connection between the pulling pipe and the well screen.

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3. Removal of the well casing is also recommended. This can be accomplished by pulling up the well casing using a cable tool drilling rig, vibration hammers, or hydraulic jacks. If the well construction method includes leaving the casing drive shoe in place, a casing cutter may be used to separate the drive shoe from the casing to facilitate removal of the well casing.

If the well casing cannot be removed, the casing should be perforated from the bottom to within 5 feet of the land surface. Perforations should be at least four equidistant cuts per row, with one row per foot of well casing. Each cut should be a minimum of 1-1/2 inches long. The perforations are necessary to allow contact of the sealing material with the walls of the borehole.

4. As the well casing is being withdrawn, sealing materials such as cement grout, neat cement, or bentonite are placed concurrently into the borehole using a dump bailer or tremie pipe. The sealing material is placed from the bottom up to avoid segregation or dilution of these materials. When a tremie pipe is used, it is submerged at the discharge end to avoid breaking the seal while filling the annular space.

6.3.4 <u>Gravel/Sand Filter Pack Wells Under Unconfined</u> Conditions

These wells have been constructed using 2-inch-diameter flushthreaded PVC well casing. The construction also includes a 2- to 5-foot bottom sump with a fitted end cap. This type of installation is not compatible with a concurrent casing extraction, seal material emplacement procedure.

Abandonment procedures are the same as those presented for casing wells under unconfined conditions, Section 6.3.2.

6.3.5 <u>Gravel/Sand Filter Pack Wells Under Confined Con</u>ditions

These well types have been constructed using 4-inch-diameter, flush-threaded PVC well casing. The construction also includes a 1- to 6-foot bottom sump with fitted end cap. This well installation type cannot be extracted while concurrently placing sealing materials.

The abandonment procedure should consist of the following steps.

 The static water level and total depth should be measured. The measured depth is compared with the construction depth to ascertain whether or not there are obstructions that may interfere with placement of sealing

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material. Obstructions are removed. The volume of the well casing should be calculated to assure that sufficient sealing material is emplaced.

- A cement grout or concrete plug should be placed in the confining stratum overlying the confined zone to minimize subsurface leakage from the confined zone. The remainder of the well should be filled with cement grout, concrete, or bentonite.
- 3. The protective casing and guard posts should be extracted, and the exposed portion of PVC casing cut off at ground level.
- 4. A cement or concrete patch should be constructed over the surface exposure of the well.

Records of the abandonment procedure are kept to provide future reference and to demonstrate to federal or state agencies that the abandonment was properly performed. Information to be recorded should include:

- o The depth of each layer of all sealing materials
- o The quantity of sealing materials used
- o Physical changes made to the well casing (e.g., perforations)

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