DRAFT Remedial Investigation/Feasibility Study Report

Dakota Creek Industries Anacortes, Washington Ecology Agreed Order No. DE-07TCPHQ-5080

for

Washington State Department of Ecology on Behalf of Port of Anacortes

April 27, 2020



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LIST OF ACRONYMS AND ABBREVIATIONS

Acronym/

Abbreviation Description

AET Apparent Effects Threshold

Agreed Order No. DE-07TCPHQ-5080

ARARS Applicable Regulatory Requirements

ARI Analytical Resources, Inc.

ASTM ASTM International

ASTs above ground storage tanks

bgs below ground surface

bcy in-place cubic yards

BETX benzene, ethylbenzene, toluene and total xylenes

BMPs best management practices

BSAFs biota-sediment accumulation factor

cm centimeter

CAO Cleanup Action Objective

CAP Cleanup Action Plan

CCI CCI Analytical

City of Anacortes

COC Contaminant of Concern

COPC Contaminant of Potential Concern

cPAH carcinogenic polycyclic aromatic hydrocarbons

CSL Cleanup Screening Level

CSO Combined Sewer Outfall

CSM Conceptual Site Model

DAHP Washington State Department of Archaeology and Historic Preservation

DCA disproportionate cost analysis

DCAP Draft Cleanup Action Plan

DCI Dakota Creek Industries



DEA David Evans and Associates

DMMP Dredged Material Management Program

DMMU Dredged Material Management Unit

DNR Department of Natural Resources

DP direct push

DPS Distinct Population Segment

DUA Decision Unit Area

Ecology Washington State Department of Ecology

EDB dibromoethane, 1-2

EDC dichloroethane, 1-2

EIM Ecology's Environmental Information Management System

ELCO electrocoagulation

EPA United States Environmental Protection Agency

ESA Endangered Species Act

ESU evolutionarily significant unit

FS Feasibility Study

FSID Facility Site Identification

ft/day feet per day

F_{oc} fraction organic carbon

GeoEngineers GeoEngineers, Inc.

GPS global positioning system

H:V horizontal to vertical

H_w Head in the well above the equilibrium level at time zero

H_o Initial change in head in the well due to an injection of a volume (V) at time zero

HPAH high molecular weight polycyclic aromatic hydrocarbons

HA hand auger

HSA hollow stem auger

ISGP Industrial Stormwater General Permits

K_{oc} organic carbon partition coefficient

L/kg liters per kilogram

LPAH low molecular weight polycyclic aromatic hydrocarbons



μg/kg micrograms per kilogram

mg/kg milligrams per kilogram

mg/L milligrams per liter

MHHW mean higher high water

MLLW mean lower low water

MS Manufacturing/Shipping (Anacortes Zone Code)

MTBE methyl tertiary-butyl ether

MTCA Model Toxics Control Act ng/kg nanograms per kilograms

NMFS National Marine Fisheries Service (now NOAA Fisheries)

NOAA National Oceanic and Atmospheric Administration

NPDES National Pollution Discharge Elimination System

OHW Ordinary High Water

OnSite OnSite Environmental, Inc.

PAHs polycyclic aromatic hydrocarbons

PCBs polychlorinated biphenyls

PCUL preliminary cleanup level

Pier 1 Port of Anacortes Pier 1 Marine Terminal

Pier 2 Port of Anacortes Pier 2 Marine Terminal

Port of Anacortes

PQL practical quantitation limit

PSEP Puget Sound Estuary Program

PVC polyvinyl chloride

RCRA Resource Conservation and Recovery Act

RI Remedial Investigation

RI/FS Remedial Investigation/Feasibility Study

SAIC Science Applications International Corporation

SCO Sediment Cleanup Objective

SCUM II Sediment Cleanup Users' Manual II

Site Dakota Creek Industries

SM Standard Method



SMS Sediment Management Standard

SPI sediment profile imaging

SQS sediment quality standards

SVOC semi-volatile organic compound

SWPPP Stormwater Pollution Prevention Plan

TBT tributyltin

TDS total dissolved solids

TEE terrestrial ecological evaluation

TEF toxicity equivalency factor

TEQ toxicity equivalency quotients (refers to concentration basis)

TOC total organic carbon

TS total solids

TVS total volatile solids

USFWS United States Fish and Wildlife Service

USGS United States Geological Survey

USTs Underground Storage Tanks

UTL upper tolerance limit

VCP Voluntary Cleanup Program

VOCs volatile organic compound

WAC Washington Administrative Code



EXECUTIVE SUMMARY

On behalf of the Port of Anacortes (Port), and in accordance with Agreed Order No. DE-07TCPHQ-5080 (Agreed Order; Ecology 2007), GeoEngineers, Inc. (GeoEngineers) has prepared this Remedial Investigation (RI)/Feasibility Study (FS) Report for the "Anacortes Port of Dakota Creek" site (Site) located along the shoreline of Guemes Channel at the northern terminus of Q Avenue in Anacortes, Washington. The RI/FS was completed using environmental investigation data collected by the Port in general accordance with the Ecology-approved Remedial Investigation/Feasibility Study (RI/FS) Work Plan (GeoEngineers 2008a). Environmental data from previous soil, groundwater and sediment characterization studies completed at the Site were used to identify contaminants of potential concern (COPCs) for evaluation in the RI. The purpose of the RI was to define nature and extent of contamination in affected Site media and to identify and evaluate cleanup actions to address the identified contamination. The purpose of the FS was to develop Cleanup Action Objectives (CAOs), screen potential remedial technologies, develop cleanup action alternatives to address contaminated media of concern, evaluate the cleanup action alternatives relative to the Model Toxics Control Act (MTCA) threshold requirements and identify the cleanup action alternative that achieves the highest level of environmental benefit with a cost that is not disproportionate to the other cleanup action alternatives evaluated. This RI/FS report was prepared under the direction of the Washington State Department of Ecology (Ecology) in accordance with the Agreed Order.

Site Description and Historical Land Use

The Site, located at 115 Q Avenue in Anacortes, Washington, is an active shipyard used for new vessel construction and repair. The Site is comprised of both upland and marine areas and is bounded by the Port of Anacortes Pier 1 to the west and Pier 2 to the east, 3rd Street on the south, and the Guemes Channel to the north. The Site is located in the northwest quadrant of Section 18, Township 35 North, and Range 2 East, has the coordinates of latitude N48.520606° and longitude W122.610640°. Dakota Creek Industries (DCI) currently leases the Site from the Port for vessel construction and maintenance operations. The Site includes a portion of the Port's Pier 1 Marine Terminal (Pier 1), a centrally located outfitting dock (Central Pier), a synchrolift, upland fabrication areas, shops, a sandblast grit storage shed, warehouses and storage areas. The northern portion of Pier 1 (which is a deep-water moorage terminal) is used by DCI to support dry dock operations.

The marine portion of the Site (Marine Area) is located between the Port's Pier 1 and Pier 2 Marine Terminals and has a navigation depth of approximately -35 feet Mean Lower Low Water (MLLW) to support shipyard operations. To the west and south, the Marine Area is separated from the uplands by vertical sheet pile bulkheads. To the east, the Marine Area is bound by the Port's Pier 2 Marine Terminal which is an earth fill structure and a pile supported wharf. The slope of the earth fill is armored with riprap.

The upland portion of the Site (Upland Area) is relatively flat with a ground surface elevation ranging between approximately 13 and 15 feet MLLW. Most of the Upland Area is paved with asphalt or concrete. Limited portions of the Upland Area are unpaved and consist of a crushed gravel working surface for fabrication layout and heavy equipment use. Currently, public access to the shipyard facility and the Port's Pier 1 and Pier 2 facilities is restricted with fencing, signage and security guards.

Since approximately 1879, the Site has been used for shipping, shipbuilding, ship repairs and other maritime-related industrial purposes and has contained various above ground storage tanks (ASTs), a rail spur, and associated buildings including machine shops, welding shops and equipment sheds to support



industrial operations. Prior to 2008, the Marine Area contained multiple piers, docks and two marine railway boat lifts. The west marine railway, located between the East Pier and Pier 1, was removed in the early 1990s. The east marine railway located between the East Pier and Pier 2 was removed in 2008 as part of the Project Pier 1 redevelopment activities. The Project Pier 1 redevelopment activities also included the removal of the L and East Docks and associated marine structures, dredging of approximately 170,000 cubic yards of sediment to achieve the current navigational depth of the Marine Area, installation of 670 linear feet of sheet pile bulkhead to reconfigure the southern shoreline, placement of 250 linear feet of riprap along the basin's east boundary and construction of the Central Pier. Concurrent with the 2008 redevelopment activities, an interim action cleanup was completed in accordance with the Ecology-approved RI/FS Work Plan and Interim Action Work Plan Addendum (GeoEngineers 2008b) to remove approximately 26,000 cubic yards of contaminated sediment from the southern half of the Marine Area and contaminated soil from the Upland Area during excavation activities to install new subsurface utility infrastructure.

Site Characterization

Previous Site Characterization and Cleanup Actions

Multiple environmental studies have been completed at the Site since 1991 to characterize Site conditions. These studies have identified that historical uses including vessel moorage, bulk fuel and oil storage, and shipbuilding activities have resulted in the release of contaminants to soil, groundwater and sediment. Cleanup actions previously completed at the Site include:

- 1991 UST Cleanup Action In 1991, two underground storage tanks (USTs) located near the south end of L dock were removed from the Site for permanent closure. During the removal of these tanks, approximately 20 cubic yards of petroleum impacted soil was removed from this area and transferred from the Site for landfill disposal. Verification samples at the final excavation limits were obtained to confirm the removal of the petroleum impacted soil observed during tank removal activities.
- 2001 Hydraulic Winch Cleanup Action In 2001, a hydraulic winch and its timber frame located near the south end of the east marine railway were removed from the Site. During removal of this structure and associate components, approximately 30 cubic yards of petroleum impacted soil were excavated and transferred from the Site for landfill disposal. Verification samples at the final excavation limits were obtained to confirm the removal of the petroleum impacted soil observed during removal of the hydraulic winch and associated timber frame.
- 2002 Petroleum and Marine Railway Cleanup Actions In 2002, the Port completed cleanup actions to address known soil contamination in the Petroleum Cleanup Action Area extending from the aluminum shop (building formerly identified as the equipment maintenance shed) to the former bulk fuel storage ASTs; and the Marine Railway Cleanup Action Area located near the eastern marine railway structure. Cleanup actions to remove soil contamination (approximately 1,650 cubic yards) in these areas were completed under Ecology's Voluntary Cleanup Program (VCP). Upon completion of the remedial excavation activities, discrete confirmation samples from the excavation sidewalls and base were collected to verify the removal of soil contamination. The previously identified contamination in these areas was successfully removed from the Site as indicated by verification sampling and the excavation areas were backfilled to the original grade with clean imported soil. Although the cleanup actions confirmed the removal of soil contamination in these areas, the work was never finalized under the VCP.



Agreed Order Site Characterization

On December 12, 2007, the Port entered Agreed Order No. DE-07TCPHQ-5080 with Ecology. Under the Agreed Order, the Port is required to complete an interim cleanup action in the marine area, evaluate the nature and extent of contamination in affected media on a Site-wide basis and develop and evaluate cleanup alternatives for addressing the identified contamination. In accordance with the Agreed Order, an RI was completed to further evaluate sediment, groundwater and soil conditions at the Site to define the nature and extent of contamination. RI activities included collection of new environmental data to evaluate the nature and extent of COPCs identified by the previous environmental studies completed at the Site.

Sediment Remedial Investigation

Sediment investigation activities were completed in general accordance with the Ecology-approved RI/FS Work Plan to characterize the vertical extent of sediment contamination in areas previously identified as exceeding the Sediment Management Standard (SMS) criteria, and to evaluate sediment in areas of the basin where no data previously existed.

In October 2008, RI sediment samples were collected from seven locations (G-1 through G-7) within the Marine Area. Sediment samples at locations G1, G2 and G7 were collected using a vibracore deployed from a research vessel. Sediment samples at locations G3 through G6 were collected from the upland area using a limited access direct-push drill rig during low tide. Surface and subsurface sediment samples were collected from sediment cores advanced to depths ranging from approximately 4 to 7 feet below the mudline surface to evaluate sediment quality in areas of the basin where no data previously existed.

Groundwater Remedial Investigation

Groundwater investigation activities were completed in general accordance with the Ecology-approved RI/FS Work Plan to characterize groundwater conditions at the shoreline where groundwater discharges to surface water, evaluate groundwater conditions in the vicinity of the 1991 UST, 2001 Hydraulic Winch and 2002 Petroleum and Marine Railway Cleanup Action Areas, to estimate the hydraulic conductivity of the shallow aquifer, and to evaluate tidal influence on the shallow aquifer.

Initial monitoring activities were completed in June 2008 to evaluate groundwater conditions. Following completion of the 2008 Interim Action and reconfiguration of the DCI shoreline, Ecology required that four additional rounds of quarterly groundwater monitoring be completed to further evaluate groundwater conditions. Quarterly groundwater monitoring events were completed in May 2012, August 2012, November 2012 and February 2013. Due to inconclusive evidence linking contaminant exceedances identified in soil to contaminant exceedances in groundwater, Ecology required that four additional rounds of groundwater monitoring be completed on a semi-annual basis to further evaluate the potential source of soil contamination to groundwater. In addition, Ecology determined that the location of monitoring well MW-1 was not an appropriate location for monitoring the conditional point of compliance and that a new well (MW-8) be installed north of MW-1 to serve that purpose.

Soil Remedial Investigation

Soil investigation activities were completed in general accordance with the Ecology-approved RI/FS Work Plan to characterize soil conditions in the upland portion of the Site to characterize the Site for the purpose of developing and evaluating cleanup action alternatives. Soil investigation included the collection of samples using a combination of hollow stem auger (HSA), direct push (DP), test pit (TP) and hand auger (HA) exploration technologies to evaluate soil conditions. In June 2008, subsurface soil samples were collected from 11 HSA explorations, 10 test pit explorations and three hand auger explorations to meet the



objectives of the RI/FS Work Plan. In 2014, a supplemental soil investigation was completed to further characterize the nature and extent of these contaminants in soil. During this supplemental soil investigation, 43 DP explorations were completed to further evaluate soil conditions. At the request of Ecology, three additional DP explorations were completed in July 2018 to evaluate soil conditions adjacent to and upgradient from monitoring well MW-8 based on the detected concentrations of arsenic and carcinogenic polycyclic aromatic hydrocarbons (cPAHs) in groundwater near this location.

Evaluation of Site Outfalls and Catch Basins

During development of the RI/FS Work Plan, Ecology identified sediment in DCI catch basins as a potential source of contamination to the Marine Area sediments due to the configuration of the Site stormwater system at the time. However, the Site stormwater system has been significantly modified since preparation of the RI/FS Work Plan. Most of the Site surfaces have been paved over-time and a new system was installed as part of the Port's Project Pier 1 redevelopment to capture the stormwater and wastewater from the Site for treatment prior to discharge. Stormwater and wastewater captured at the Site is treated prior to discharge to the City's sanitary sewer or to Guemes Channel under National Pollution Discharge Elimination System (NPDES) General Permit WARO45711. DCI performs regular monitoring of the water collection systems to ensure compliance with the discharge requirements of the NPDES permit.

As a result of the ongoing treatment and monitoring activities for stormwater and wastewater from the Site, discharges from the DCI stormwater/wastewater collection systems are no-longer a potential source of contamination to the Marine Area portion of the Site. Potential historical contaminant discharges to the Marine Area prior to the stormwater system upgrade activities have been addressed as part of the 2008 Interim Action, where the known contaminated sediments and underlying clean native sediments were removed from the Site as discussed below.

Interim Action

Interim action dredging and excavation activities as discussed above were completed between July and November 2008 in general accordance with the Ecology-approved RI/FS Work Plan and Interim Action Work Plan Addendum to remove identified contamination in the Marine Area for upland landfill disposal and to remove portions of the known soil contamination in the Upland Area. During the 2008 Interim Action, approximately 26,000 cubic yards (approximately 38,000 tons) of contaminated sediment was removed from the Marine Area and an additional 580 cubic yards (approximate) of contaminated soil was removed from the Upland Area. Contaminated sediment and soil were transported by truck from the Site for upland landfill disposal.

Following verification of the contaminated sediment removal, an additional 230,000 cubic yards of clean sediment (approximate) determined to be suitable for open-water disposal by the Dredged Material Management Program (DMMP) was then dredged from the Marine Area and transported by barge to the Rosario Strait dispersive site to meet the Project Pier 1 redevelopment design grade of -35 feet MLLW.

Sediment samples collected from the base of the interim action dredge surface confirmed removal of contaminated sediments from the Marine Area and were used by the regulatory agencies to confirm that further dredging (beyond the limits of contamination) was in clean materials.



Nature and Extent of Contamination

Based on a review of the Upland and Marine Area RI results, the following COCs were identified for sediment, groundwater and soil at the Site:

- Sediment Metals including arsenic copper, lead, mercury and zinc, tributyltin (TBT), low molecular weight PAHs (LPAHs), high molecular weight PAHs (HPAHs), cPAHs, polychlorinated biphenyls (PCBs), dioxin and furans were identified as COCs for Marine Area sediment. However, interim action dredging in the Marine Area in 2008 resulted in the complete removal of sediment contamination at the Site. As a result, sediment is no longer considered a media of concern.
- **Groundwater** Metals including arsenic and nickel, and cPAHs were identified as groundwater COCs for the Upland Area.
- Soil Metals including arsenic and nickel, and cPAHs were identified as soil COCs for the Upland Area. In addition, results of previous environmental studies identified concentrations of gasoline-, diesel-and/or heavy oil-range petroleum hydrocarbons exceeding PCULs. As indicated above, the Port completed independent cleanup actions (i.e., 1991 UST, 2001 Hydraulic Winch and 2002 Petroleum and Marine Railway Cleanup Actions) to remove the previously identified petroleum contamination from the Site. Confirmation sample results obtained from the limits of these excavations indicated that the petroleum contamination was successfully removed from the Site. However, confirmation soil sample results for these areas could not be independently validated. Therefore, petroleum hydrocarbons within the footprints of the previously completed cleanup action areas are unverified until subsequent sampling result confirm their removal.

Marine Area

As discussed above, interim action dredging and excavation activities completed between July and November 2008 in general accordance with the Ecology-approved RI/FS Work Plan and Interim Action Work Plan Addendum removed the identified sediment COCs exceeding preliminary cleanup level (PCULs) from the Marine Area. Sediment samples collected from the base of the interim action dredge prism and sediment sample results from previous environmental studies within the Marine Area provide the basis for sediment not being considered a media of concern for the Site.

Upland Area

In the Upland area, COCs including arsenic, nickel and cPAHs were detected at concentrations exceeding PCULs in soil throughout the Site. In the eastern portion of the Site, arsenic and nickel exceeded PCULs in fill deposits from the ground surface to a depth of approximately 8 feet below ground surface (bgs). In the north central portion of the Site, arsenic and nickel exceeded PCULs in fill deposits from the ground surface to a depth of approximately 10 feet bgs. In the southwestern portion of the Site, arsenic, nickel and cPAHs exceeded the PCUL in fill deposits from the ground surface to a depth of approximately 13 feet bgs. Results of soil/sediment samples collected at the Site from the underlying native surface show that the Upland Area PCUL exceedances are limited to the overlying fill soil and do not extend to the underlying native surface.

In addition, results of previous environmental studies identified concentrations of gasoline-, diesel- and/or heavy oil-range petroleum hydrocarbons exceeding preliminary cleanup levels (PCULs) in historical fill deposits from the ground surface to a depth of approximately 8 feet bgs in the central and eastern portions of the Site. Between 1991 and 2002, the Port completed independent cleanup actions in these areas to



remove the previously identified petroleum contamination from the Site. These cleanup actions also likely resulted in the removal of arsenic, nickel and cPAH contamination from the Upland Area further limiting their nature and extent. Confirmation sample results obtained from the limits of these excavations indicated that the petroleum contamination was successfully removed from the Site. However, confirmation soil sample results for these areas could not be independently validated. A supplemental soil investigation is proposed as a pre-remedial design activity for the Site to verify the completeness of the previous remedial actions and would be reported in the Cleanup Action Plan (CAP). Sampling and analysis for this investigation will be completed under an Ecology-approved addendum to the RI/FS Work Plan in support of the selected preferred cleanup action alternative (discussed below) to address Site contamination.

Between 2015 and 2016, DCI replaced a significant portion of their gravel working surface with asphalt pavement which acts to prevent stormwater infiltration through the soil column. RI Groundwater monitoring results show that the paved surfaces are limiting the infiltration, leaching and subsequent migration of contaminants through the soil column to groundwater. In addition, this data show that contaminants that remain in place in saturated zone soils have stabilized and are not migrating downgradient toward the Guemes Channel since paving was completed.

At shoreline monitoring well locations, the following COCs were detected at concentrations greater than the groundwater PCUL since completion of the Upland Area paving activities:

- Dissolved nickel detected at a concentration of 8.3 micrograms per liter (μg/L) marginally exceeded the groundwater PCUL of 8.2 μg/L at shoreline monitoring well location MW-2B during the February 2017 monitoring event. However, total nickel at this location was not detected at a concentration exceeding the groundwater PCUL during this event and dissolved nickel did not exceed the PCUL in subsequent monitoring events at this location.
- Total cPAH toxicity equivalency (TEQ) concentrations and/or dissolved arsenic exceeded the groundwater PCUL at shoreline monitoring well location MW-8 during one or more semi-annual monitoring events between February 2016 and August 2017. However, cPAHs and arsenic concentrations at monitoring well location MW-1 (upgradient monitoring well location) were less than the PCUL for each of these monitoring events. Supplemental soil sampling and analysis to further evaluate soil conditions in the vicinity of MW-8 did not identify potential source materials for cPAHs or arsenic in saturated soil adjacent to or upgradient of this location.

At upgradient monitoring well locations, PCUL exceedances were only observed at monitoring well location MW-7. At location MW-7, total and dissolved arsenic and nickel were detected at concentrations exceeding groundwater PCULs during one or more semi-annual monitoring events. However, at downgradient monitoring well locations MW-3A and MW-6, total and dissolved arsenic and nickel either were not previously detected or were detected at concentration less than the groundwater PCUL for the last four semi-annual groundwater monitoring events.

Preferred Cleanup Action Alternative Selection

Potentially applicable response actions and associated remediation technologies were identified and screened for the development of cleanup action alternatives to address COCs in soil and groundwater discussed above. The screening process determined the most appropriate technologies and process options based on their expected implementability, reliability, effectiveness, and relative cost. Screening also considered modifying criteria associated with current and future land uses, consideration of potential



historical and archaeological remains, and impacts to existing habitat resources. Cleanup action alternatives were then developed by combining technologies retained through the screening process to meet the Site cleanup standards. The design parameters used to develop the alternatives were based on both engineering judgment and the current knowledge of Site conditions and are conceptual-level designs for the implementation of the individual technologies. In accordance with the requirements of WAC 173-340-350 and WAC 173-340-360, cleanup action alternatives were evaluated against the following criterion:

- Compliance with cleanup standards and applicable laws;
- Provision for a reasonable restoration time frame; and
- Use of permanent solutions to the maximum extent practicable by comparison of the following:
 - Protectiveness:
 - Permanence:
 - Cost:
 - Effectiveness over the long term;
 - Short-term risk management;
 - Net environmental benefit;
 - Technical and administrative implementability; and,
 - Consideration of public concerns.

A MTCA disproportionate cost analysis (DCA) was then completed to determine which cleanup action alternative that otherwise meets the threshold requirements achieves the highest level of environmental benefit while not being disproportionate in cost relative to the other alternatives. As a result of this evaluation, Cleanup Action Alternative 2 emerged as the preferred alternative which meets the minimum threshold requirements, achieves a high level of environmental benefit and is not disproportionate in cost relative to the other alternatives evaluated. Implementation of Cleanup Action Alternative 2 will result in contaminant mass reduction in the southeast portion of the Site and will be used in conjunction with containment technologies and institutional controls to prevent direct human contact and reduce the potential for leaching and migration of residual COCs contained within the fill soil within a reasonable restoration time frame.



1.0 INTRODUCTION

This report presents the results of the Remedial Investigation (RI) and Feasibility Study (FS) completed for the Port of Anacortes (Port) Dakota Creek Industries (DCI) shipyard facility (Site) located along the southern shoreline of Guemes Channel at the northern terminus of Q Avenue in Anacortes, Washington (Figure 1). The RI/FS was completed pursuant to the Washington State Department of Ecology (Ecology) Agreed Order DE-07TCPHQ-5080 (Agreed Order; Ecology 2007) and in accordance with the Washington State Model Toxics Control Act (MTCA) cleanup regulations (Chapter 173-340 Washington Administrative Code [WAC]). The Site is listed in the Washington State Department of Ecology (Ecology) Cleanup Site Database as Facility Site Identification (FSID) No. 2670 and Cleanup Site Identification No. 5174 and is formally referred to as Anacortes Port Dakota Creek. Ecology is managing the Site under the Puget Sound Initiative as part of their regional cleanup efforts on Fidalgo Island.

This RI/FS presents:

- The results of the investigation to define the nature and extent of contamination in media of concern at the Site and provides the data needed to complete an evaluation of cleanup actions to address the identified contamination; and
- The development and evaluation of cleanup action alternatives for addressing contamination identified at the Site and to select a preferred cleanup action alternative utilizing information gathered during the RI and previous environmental studies.

This RI/FS was completed in accordance with the requirements of the MTCA Cleanup Regulation, Chapter 173-340 WAC and the Sediment Management Standards (SMS), Chapter 173-204 WAC.

1.1. General Site Information

1.1.1. Site Description

The Site is located at 115 Q Avenue in Anacortes, Washington (Figure 1) and is an active shipyard used for new vessel construction and repair. The Site is comprised of both upland and marine areas and is bounded by the Port's Pier 1 Marine Terminal to the west and Pier 2 Marine Terminal to the east, 3rd Street on the south, and the Guemes Channel to the north. The Site is located in the northwest quadrant of Section 18, Township 35 North, and Range 2 East and has the coordinates of latitude N48.520606° and longitude W122.610640°.

DCI currently operates a shipyard at the Site and leases the property from the Port. DCI uses the facility for vessel construction and maintenance activities. The Site includes a portion of the Port's Pier 1 Marine Terminal, a centrally located outfitting dock (Central Pier), a synchrolift, upland fabrication areas, shops, a sandblast grit storage shed, warehouses and storage areas. The northern portion of Pier 1 (which is a deep-water moorage terminal) is used by DCI to support dry dock operations. Features of the Site and surrounding area are shown on Figure 2.

The offshore area of the Site (henceforth referred to as the Marine Area) is located between the Port's Pier 1 and Pier 2 Marine Terminals (Pier 1 and Pier 2) and is maintained with a navigation depth of approximately -35 feet Mean Lower Low Water (MLLW) to support shipyard operations. To the west and south, the Marine Area is separated from the uplands by vertical sheet pile bulkheads. To the east, the



Marine Area is bound by Pier 2 which is an earth fill structure and a pile supported wharf along the northern most part of the facility. The slope of the earth fill is armored with large rock (riprap).

The parts of the Site above Ordinary High Water (OHW) or upland area portion of the Site (henceforth referred to as the Upland Area) is relatively flat with a ground surface elevation of approximately 15 feet MLLW. Most of the upland area is paved with asphalt or concrete. The limited unpaved parts of the Upland Area consist of a crushed gravel working surface that is maintained for fabrication layout and heavy equipment. Currently, public access to the shipyard facility and the Port's Pier 1 and Pier 2 facilities is restricted with fencing, signage and security guards.

1.1.2. Legal Description

Tax parcel numbers and legal descriptions containing the Site are summarized in the following table. Tax parcel boundaries are shown on Figure 2.

Tax Parcel Number	Legal Description
P32866	Anacortes Tide Lands Tracks 2 and 3, Plate 9 Together with a Portion of Adjacent Vacated Q Avenue (Ord No. 1728) (1.3 acre).
P32867	Tracks 4 and 16, Plate 9 Including Vacated Portions of 2 nd and Broadway Street Adjacent and West 15 feet of Vacated R Street Together with East Half Vacated Q Avenue Adjacent to Track 4 (Ord No. 1707, AF No. 862268) Less Following Described Track on East Line of R Avenue 40 Feet North of North Line 3 rd Street Then East 10 Feet Then North 150 Feet Then West 10 Feet to East Line R Avenue Then South 150 Feet to Point of Beginning (2.1 acre).
P32903	Anacortes Tide Lands Tax 24A Beginning at the Intersection of North Line 3 rd Street with East Line R Avenue Then North Along East Line R Avenue 190 Feet Then West 65 Feet Then North to Inner Harbor Line Then East Along Said Line 165 Feet Then South to North Line 3 rd Street Then West 100 Feet to True Point of Beginning Less Portion Tax 24B and Easement to City Less Roll Tract 0-041-01 (0.41 acre).
P32904	Portion Block 296 City of Anacortes Together with Portion Plate 9 Tide and Shore lands Defined as Follows Beginning at a Point 25 Feet West Centerline R Avenue and 190 Feet North of North Line 3 rd Street Then North Parallel to Centerline R Avenue to Intersection West Inner Harbor Line Then East Along Said Line to a Point 100 Feet East of East Line R Avenue Then South to a Point Which Lies 190 Feet North of North Line 3 rd Street Then West to Point of Beginning (1.08 acre).
P32905	Anacortes Tide Lands Tax 24B Beginning on East Line R Avenue 40 Feet North of North Line 3 rd Street Then East 10 Feet Then North Parallel to East Line R Avenue 150 Feet Then West 10 Feet to East Line R Avenue Then South Along R Avenue 105 Feet to Point of Beginning (0.03 acre).
P32906	Anacortes Tide Lands Tax 25 then Portion West Half R Avenue Lying Between a Line 40 Feet North of and Parallel to North Line 3 rd Street and a line 190 Feet North of and Parallel to North Line of 3 rd Street (0.03 acre).
P32907	Anacortes Tide Lands Tax 26 Then Portion East Half Vacated R Avenue Lying Between a Line 40 Feet North of and Parallel to North Line 3 rd Street and a Line 190 Feet North and Parallel to North line 3 rd Street (0.13 acre).
P54924	Anacortes Block 3 Together with Vacated Alley through Block (Ord No. 1775) (0.74 acre).
P55030	Anacortes All Block 26 Together with Vacated Alley through Said Block (Ord No. 1708, AF No. 862269) (1.39 acre).



Tax Parcel Number	Legal Description
P55031	Anacortes Block 27 and Portion Northerly Extended Through Block 27 Along Vacated Alley of Block 27 Except Any Portion Lying within Tract 3, Plate 9 of Anacortes Tide Lands (1.21 acre).
P56539	Anacortes Lot 13 Block 296 11 to 13 (0.1 acre).

1.2. Historical Operations and Use

The Site has been used for shipping, shipbuilding, ship repairs and other maritime-related industrial purposes since approximately 1879. Historically, various above ground storage tanks (ASTs), a rail spur, and associated buildings including machine shops, welding shops and equipment sheds were located at the Site to support maritime operations. The historical features are shown on Figure 3. Aerial photographs presented in Appendix A show historical operations and development of the Site and surrounding area since the early 1900s.

Sanborn maps show that a bulk oil storage and distribution facility with at least six ASTs was in operation in the central upland portion of the Site. Historical records indicate that Pacific Tow Boat leased this portion of the Site to Standard Oil in late 1946 who operated the bulk oil storage and distribution facility until 1969 after which it was sold to the Dillingham Corporation. By the time that the Site was purchased by the Port in 1975, all structures associated with the bulk oil storage and distribution facility had been removed. The location of these tanks is visible on circa 1946 and 1960s aerial photographs (Appendix A).

The southwest portion of the Site was historically used for residential purposes from the early 1900s until the late 1960s based on a review of historical Sanborn maps and aerial photographs. The ground surface in this area was historically lower that the surrounding areas by several feet and that following the purchase of this area by the Port in 1975, the grade was raised to match the surrounding area using dredged sediments from Guemes Channel. In about 1976, DCl began to lease the Site from the Port and has continued to operate the shipyard since this time.

Prior to 2008, the Marine Area contained multiple piers and docks and two marine railway boat lifts (Figure 3). The west marine railway, located between the East Pier and Pier 1, was removed in the early 1990s. The east marine railway located between the East Pier and Pier 2 was removed in 2008 as part of the Project Pier 1 redevelopment activities. The Project Pier 1 redevelopment activities also included the removal of L and East Docks, the east marine railway and associated marine structures, dredging of approximately 170,000 cubic yards of sediment to achieve the current navigational depth of the Marine Area, installation of 670 linear feet of sheet pile bulkhead (open cell bulkhead) to reconfigure the southern shoreline, placement of 250 linear feet of riprap along the Marine Area's east boundary and construction of the Central Pier. The layout of the shipyard facility following redevelopment activities is shown on Figure 4. Concurrent with the redevelopment activities, an interim action cleanup was completed in accordance with the Ecology-approved RI/FS Work Plan (GeoEngineers 2008a) and Interim Action Work Plan Addendum (GeoEngineers 2008b) to remove approximately 26,000 cubic yards of contaminated sediment from the southern portion of the Marine Area and the removal of approximately 580 cubic yards of contaminated soil from the Upland Area to install new subsurface utility infrastructure. The Interim Action activities are further discussed in Section 4.0.



1.3. Current and Future Use

The Site and adjacent area are zoned by the City of Anacortes (City) for industrial use (Manufacturing/Shipping [MS]) and is characterized by marine shipping, warehousing, bulk material storage, transportation, and other industrial uses. The Port currently leases the Site to DCI who operates a shippard for vessel construction and maintenance activities as described above. Public access to the Site (including the Port's adjacent Pier facilities) is restricted with by fencing, signage and guards.

Although the specific future uses of the Site will depend on the operations of the Port's lessees, the anticipated future use of the Site and surrounding area will continue to be for industrial purposes including shipbuilding, ship repairs and other maritime-related industrial business. The property is currently leased to DCI for an additional 37 years.

1.4. Environmental Setting

Key elements of the environmental setting of the Site, including physical conditions, geologic setting, natural resources and cultural resources are summarized in the following sections.

1.4.1. Climate

Anacortes temperatures are relatively mild. Summer daytime mean temperatures are in the 70s with night-time temperatures in the 50s. Maximum temperatures reach 80 to 85 degrees, with a few 90- to 100-degree days recorded. The highest temperatures and lowest relative humidity are recorded during periods of easterly winds. December and January are the coldest months, with average minimum temperatures in the upper 30s.

The prevailing wind direction is from the southeast in winter and southwest in summer. During late spring and summer, a prevailing westerly and northwesterly flow of air into Puget Sound brings a dry season beginning in May which reaches a peak in July. In late fall and winter, a prevailing southwesterly and westerly air flow from the Pacific Ocean results in a wet season beginning in October which lasts until the beginning of the dry season in May. During winter, the combined influence of low-pressure systems off the Pacific Ocean and cold air from the Fraser River Canyon produce strong northeasterly winds. Although it is not uncommon to have 30- to 40-knot winds under these conditions, the short fetch in the Anacortes area usually limits wind generated wave heights to no more than six feet. Wind gusts up to 73 miles per hour and sustained westerly velocities up to 54 miles per hour have been recorded.

Mean annual precipitation for Anacortes is 26.2 inches, most of which falls as rain. Average monthly precipitation varies from a low of 0.93 inch in July to a high of 3.79 inches in December.

1.4.2. Sea Level Rise

Since the time of the last glacial maximum about 20,000 years ago, sea level has been on the rise at varying rates. Global sea level has been rising over the past century, and the rate has increased in recent decades. In 2014, global sea level was 2.6 inches above the 1993 average and continues to rise at a rate of about $\frac{1}{8}$ of an inch per year (NOS 2019).

Global warming is thought to cause the two main mechanisms contributing to sea level rise which include: 1) thermal expansion (ocean water expands as it warms); and 2) melting stores of ice sheets and glaciers. Local application of global projections of sea level rise are complicated by multiple factors such as



atmospheric circulation patterns and tectonic movement. Considering these variables, the National Research Council has made projections of anticipated sea level rise for California, Oregon, and Washington. For the coast of Washington, the projected rise is up to 9 inches by 2030, up to 19 inches by 2050, an up to 56 inches by 2100 (NAP 2012).

To evaluate extreme high tide levels that are currently anticipated, graphs provided by the National Oceanic and Atmospheric Administration (NOAA) compare 10 percent and 1 percent exceedance probability levels, which correspond to tide levels that would be exceeded ten times and one time per century (i.e., the probability of an extreme tide level occurring on a 10-year interval and the probability of an extreme tide level occurring on a 100-year interval). Extreme levels are a combination of the astronomical tide, storm surge, and limited wave setup caused by breaking waves. NOAA has developed tide projections for Anacortes based on the Port Townsend tide gauge corrected for Anacortes. Currently, mean higher high water (MHHW) for Anacortes is 8.2 feet. NOAA tide predictions for a 10-year tidal level exceedance is 2.8 feet and 3.2 feet for a 100-year tidal level exceedance.

1.4.3. Topography and Bathymetry

The Site is located along the southern shoreline of Guemes Channel (Figure 1) and includes the Marine and adjacent Upland Area to the south (Figure 2). The Upland area is generally flat with elevations ranging from approximately +13 to +15 feet MLLW. Prior to 2008, the working surface of the Upland Area primarily consisted of crushed gravel. Following redevelopment of the shoreline in 2008, DCI began to pave the Upland Area with up to 6 inches of asphalt. Currently, most of the Upland Area is paved with asphalt while limited unpaved parts of the Upland Area consisting of a crushed gravel working surface are being maintained for fabrication layout and equipment storage. The extent of the gravel working surface prior to 2008 is shown on Figure 3. The current approximate extent of asphalt pavement in the Upland Area is shown on Figure 4.

The eastern bank of the Marine Area is armored with riprap which extends at an approximate 2H:1V slope to approximate elevation of -35 feet MLLW. To the south and west, sheet pile bulkheads separate the Upland and Marine Areas which extend vertically from approximately +15 feet to -35 feet MLLW. In the Marine Area, the navigation area is approximately -35 feet MLLW. Near the outer harbor line, the mudline surface rapidly drops off toward Guemes Channel.

Recent topographic and bathymetric contours at the Site are shown on Figure 4 and are referenced from a June 2014 bathymetric survey completed by David Evans and Associates (DEA) and Lidar imagery for Anacortes completed in May 2009.

1.4.4. Surface Water Bodies

The Site is located on the southern shoreline of Guemes Channel. The western end of Guemes Channel connects to the Rosario Strait. Fidalgo Bay is connected to the eastern end of Guemes Channel and is adjacent to Padilla Bay. March Point separates the southern part of Fidalgo Bay and Padilla Bay east of the Site.

There are no significant freshwater streams that flow into Guemes Channel or the Fidalgo Bay area (Antrim et al. 2000). In the Guemes Channel area, the average difference in height between mean higher high water (MHHW) and MLLW is 8.2 feet. Currents in Guemes Channel are relatively strong (averaging 0.9 and 2.1 knots on flood and ebb tides respectively; Antrim et al. 2000). Tidal currents are affected to some



extent by winds. Much of the Site is protected from prevailing currents through Guemes Channel, and from northerly wind and waves by Pier 1 and 2, and Guemes Island located to the north.

1.4.5. Shoreline Features

The Port's Pier 1 facility located west of the Marine Area was originally constructed in the early 1900s and extends north from the historical shoreline (see historical aerial photographs presented in Appendix A). The northern portion of the Pier 1 facility is a pile supported wharf which operates as a deep-water berth. From the existing shoreline, the eastern portion of Pier 1 was infilled between the 1960s and 1970 to create the present-day marine terminal structure. Over-time, Pier 1 has undergone general improvements including paving, utility upgrades and the construction of warehouses that are utilized by Port, DCI and other tenants for marine-related operations.

The synchrolift system used by DCI to raise vessels for out of water hull maintenance is located in the western part of the Marine Area and was installed in the early 1980s. During installation, sediment was dredged to a depth of approximately -35 feet MLLW directly below the synchrolift and -15 feet MLLW in the area immediately east of the lift (Figure 3). To maintain the structural integrity of Pier 1, a sheet pile bulkhead was installed along the western side of the synchrolift berth area.

Along the southern shoreline, redevelopment activities were completed in 2008 to increase the capacity and efficiency of the DCI operations. The Project Pier 1 redevelopment project included the installation of a new bulkhead (i.e., Open Cell Bulkhead), pier and dredging of the Marine Area to approximately -35 MLLW to allow for more efficient dock-side work and dry-dock operations. Clean structural fill was placed in the area south (shoreward) of the new bulkhead alignment and the pre-existing marine railway structures along with some of the existing upland buildings were removed in order to allow for more efficient use of the Upland Area. The new bulkhead extends across the southern portion of the Site separating the Marine and Upland Areas. In addition, a new pier (Central Pier) was constructed as part of the redevelopment project. The Central Pier extends north from the new bulkhead bisecting the Marine Area of the Site. This structure is paved and is supported by concrete piling.

Pier 2 located east of the Upland and Marine Areas is an earthen fill pier with a pile supported wharf at the northern portion of the facility. Pier 2 operates as a deep-water berth and is primarily used for bulk product exports. Based on a review of aerial photographs, this facility was initially constructed in the early 1900 to support marine-related industry operating in this area.

The historical configuration of the shoreline and layout of the Site prior to redevelopment activities in 2008 is shown on Figure 3. The current configuration of the shoreline and layout of the Site is shown on Figure 4.

1.4.6. Stormwater/Wastewater Outfalls

The former Scott Paper Mill outfall historically discharged near the outer part of the Marine Area between 1963 and 1973 (Figure 3). After 1973, discharge from the former Scott Paper Mill was through a new outfall pipe that was constructed to take advantage of the dispersive effects and physical characteristics of the current in Guemes Channel by extending the discharge point 680 feet beyond the outer harbor line. Discharge continued until 1978 when the former Scott Paper Mill was closed.

Following the purchase of the Site by the Port in 1975, DCI's stormwater/wastewater was discharged under Western Washington Phase II Municipal Stormwater National Pollution Discharge Elimination System



(NPDES) General Permit WAR045711. Prior to the Project Pier 1 redevelopment in 2008, stormwater/wastewater at the Site either infiltrated into the ground (at this time, DCI maintained a compact gravel working surface for facility operations), sheet flowed into Guemes Channel, or passed through oil separating catch basins before joining the City's storm drain system. Currently, stormwater/wastewater collected from the Site is collected from one of three outfall locations (Outfall 001 through Outfall 003; Figure 5) and treated before being discharged.

In addition to DCI's stormwater system, the City maintains two outfalls in the vicinity of the Marine Area. Treated wastewater from the City's wastewater treatment plant located at intersection of 5th Street and T Avenue is discharged to Guemes Channel from an outfall (R Avenue Outfall) located at the northwest corner of the Port's Pier 2 Facility (Figure 5). The City also maintains a combined sewer outfall (CSO) that discharges into the Marine Area (Q Avenue Outfall; Figure 5) to manage the throughput of stormwater to the City's wastewater treatment plant. Stormwater and wastewater collected from the Port's Pier 2 facility is captured by a large detention pond located east of the Site. Collected stormwater/wastewater is recycled for use in their truck wash station. Excess water from this system is discharged to the City's sanitary sewer system (P2O 003; Figure 5).

The current stormwater collection, treatment and discharge for the Marine Area and surrounding area is shown on Figure 5 and further discussed in the following sections.

1.4.6.1. Outfall 001 - DCI Shipyard

In 2008, new storm drains were installed throughout the Site as part of the Project Pier 1 redevelopment to collect stormwater for treatment prior to discharge to Guemes Channel. Stormwater in the shipyard is collected from a network of catch basins and storm drainpipes and conveyed to an Aquip StormwateRx treatment system. The treatment system consists of a stormwater storage tank and enhanced media filtration tank with a buffering pre-treatment chamber and an inert and sorptive filtration media chamber. The treatment system is designed to reduce suspended solids, turbidity, heavy metals (including dissolved metals), and organics prior to discharge through a 36" diameter pipe into Guemes Channel at Outfall 001 (Figure 5).

The current system does not discharge bypass stormwater. Overflow from excessive storm events is routed back to an in-ground sump which then cycles back to the treatment system prior to discharge. DCI personnel conduct sampling of the stormwater/wastewater collection system in accordance with the DCI's Stormwater Pollution Prevention Plan (SWPPP; DCI 2017) to ensure compliance with the discharge requirements of the NPDES permit.

1.4.6.2. Outfall 002 - Drydock Floodwater

DCI uses a floating drydock that is moored at the northern end of Pier 1. The shipyard uses the drydock to clean and repair ships. Vessels hauled out periodically require pressure washing. Drydock floodwater is discharged when the drydock is flooded to dock or float a vessel onto or off the drydock floor (Outfall 002; Figure 5). To minimize the potential for pollutants to enter Guemes Channel when the drydock floor submerges, the drydock floor is cleaned of debris following vessel cleaning and repair activities. Prior to each lowering of the drydock, DCI personnel thoroughly sweep and clean the deck and stairwells to remove any visible debris. In addition, the surfaces of the drydock are pressure washed on an as needed basis to remove any oily substances that may be present. Pressure wash wastewater is collected in a trough located on the east side of the drydock which connects to a removable collection sump located on the southeast



corner of the drydock. The collection sump is removed and hoisted ashore prior to lowering of the drydock where it is routinely cleaned.

Before each lowering of the drydock, DCI inspects and photographs the cleanliness of the drydock deck. Sampling for Outfall 002 is completed from the catwalk above the southeast corner of the drydock in accordance with DCI's SWPPP to ensure compliance with the discharge requirements of the NPDES permit.

1.4.6.3. Outfall 003 - Wastewater

Pressure wash wastewater and other wastewater/liquids generated during vessel cleaning operations at the shipyard is collected by sumps that service the drydock, rails area and mechanics shop. Collected wastewater is transported by tanker truck to a wastewater treatment system where the wastewater is processed prior to discharge to the City's sanitary sewer system at Outfall 003 (Figure 5). Wastewater is processed using an Ultrasorb® and Electrocoagulation (ELCO) Treatment Systems prior to discharge. The Ultrasorb® System is comprised of an accumulation tank with oil skimmer, coalescers and filters to remove oil and volatile organic compounds from the wastewater. The ELCO System targets the removal of suspended solids and metals by inducing a charge in the partials causing them to bond together and fall out of solution.

DCI personnel conduct sampling of the wastewater treatment system in accordance with the DCI's SWPPP to ensure compliance with the discharge requirements of the NPDES permit.

1.4.6.4. R Avenue Outfall

The City's wastewater utility serves over 4,000 acres of residential and commercial customers within the City of Anacortes. The system is classified as a combined stormwater and wastewater system. The wastewater system is responsible for operating and maintaining 96.8 miles of gravity sewers, 23 pump stations, 9.4 miles of force mains ranging in size from 1.5 to 12 inches in diameter, and 175 septic tanks. The system collects and transports wastewater to the City's treatment plant for processing prior to discharge. Effluent discharge from the treatment plant to the Guemes Channel is through the R Avenue outfall (Figure 5).

1.4.6.5. Q Avenue Outfall

The Q Avenue CSO allows for the discharge of untreated wastewater to an outfall located beneath DCI's Syncrolift Pier (Figure 5). Overflow to this outfall is separated from the wastewater flows by a concrete overflow dam positioned in an upgradient manhole connected to the City's wastewater system. During elevated precipitation events, wastewater is allowed to overflow the dam and discharge directly to Guemes Channel to prevent capacity exceedance of the City's wastewater treatment plant. The City has permanent flow meters installed at each of the CSO structures to measure CSO activity. The flows are reported and discussed in annual CSO reports that are submitted to Ecology. The current average overflow rate per year for this CSO since 2000 is 0.42 overflow events per year (PARIS 2019).

1.4.6.6. Pier 2 Stormwater Discharge

Pier 2 is a 14-acre paved pier owned and operated by the Port. Prior to 2011, the facility operated under Industrial Stormwater General Permits (ISGP) WAR000849, WAR001004 and WA0020257. In the northern portion of the facility, collected stormwater was discharged directly to Guemes channel at outfall P20 001 under ISGP WAR000849 (Figure 3). In the central and southern portions of the facility, stormwater collected from the facility was conveyed to a settlement pond prior to discharge to the Marine Area at outfall P20 002 under ISGP WAR001004 (Figure 3). Wash water generated from a wheel wash station operating at the



facility during trucking operations is collected in a secondary settlement pond prior to discharge to the City's sanitary sewer system at outfall P20 003 under ISGP WA0020257 (Figure 5).

Following Pier 2 facility upgrades, stormwater and wash water is collected and discharged to the City's sanitary sewer system at outfall P2O 003 under State Waste Discharge Permit ST0045500. Facility upgrades for Pier 2 included construction of a new settling pond as well as the installation of two 15,000-gallon water tanks used to store recycled water for wheel wash operations and new manholes, pumps and piping to connect the system. Collected stormwater and wash water for the facility trucking operations is recycled. Excess water generated by the system is discharged to the City's sanitary sewer system at outfall P2O 003 for treatment prior to discharge to Guemes Channel (discussed above). The solid waste build-up in the facilities detention pond is removed on an annual basis for upland disposal to a permitted landfill.

Historical outfalls for the Port's Pier 2 facility are shown on Figure 3. Current outfalls for the Port's Pier 2 facility are shown on Figure 5.

1.4.7. Geological Setting

1.4.7.1. Local Geology

The United States Geological Survey (USGS) map of the Bellingham Quadrangle (Lapen 2000) was reviewed for geologic information in the vicinity of the Site. The geologic soil deposits in the vicinity of the Site are the result of both glacial and nonglacial processes that have occurred during the last 12,000 years.

Soil deposits at the Site consist of artificial fill overlying recessional marine (glaciomarine) drift from the Everson Interstade of the Fraser Glaciation. Artificial fill deposits are primarily characterized by silt, sand, and gravel that contain periodic wood debris, organic material, asphalt debris, concrete, and glass/tile debris resulting from historical land uses. Glacial marine deposits are primarily characterized by unsorted, unstratified silt and clay with varying amounts of sand, gravel, cobbles and occasional boulders deposited during the glacial advancement and retreat (melting). This material may contain shells, wood, and large erratics (boulders) as a result of sea level fluctuation relative to the land surface and present-day sea level.

East of the Site, bedrock is mapped at the ground surface and is part of the Lummi Formation which consists of marine metasedimentary rock. The Lummi Formation is a metamorphosed pebble conglomerate, sandstone, and/or mudstone that were deposited during the early Cretaceous to late Jurassic age (140 to 150 million years before present).

1.4.7.2. Geologic Hazards

The Site is located within the Puget Sound region, which is seismically active. Seismicity in this region is attributed primarily to the interaction between the Pacific, Juan de Fuca and North American plates. The Juan de Fuca Plate is subducting beneath the North American Plate. It is thought that the resulting deformation and breakup of the Juan de Fuca Plate might account for the deep focus earthquakes in the region.

Research has concluded that historical large magnitude subduction-related earthquake activity has occurred along the Washington and Oregon coasts. Evidence suggests several large magnitude earthquakes (Richter magnitude 8 to 9) have occurred in the last 1,500 years, the most recent of which occurred about 300 years ago. No earthquakes of this magnitude have been documented during the recorded history of the Pacific Northwest.



A review of geologic maps has identified a small fault line that runs in a northwesterly direction between Guemes Channel and Cap Sante Marina. Based on review of USGS and Department of Natural Resources (DNR) maps, the fault does not appear to be connected to any named fault system.

Other geologic hazards for the region include liquefaction based on the presence of artificial fill. Liquefaction refers to a condition where vibration or shaking of the ground, usually from earthquake forces, results in the development of excess pore pressures in saturated soils and subsequent loss of strength. This can result in vertical oscillations and/or lateral spreading of the affected soils, with accompanying surface subsidence (sinking) and/or heaving. In general, soils that are susceptible to liquefaction include loose to medium dense clean to silty sands that are saturated (i.e., below the water table).

Based on the topography of the Site and surrounding area, geologic hazards from landslides were not identified.

1.4.8. Natural Resources

1.4.8.1. Terrestrial Habitat

Typical of industrialized waterfronts, sections of the shoreline adjacent to the Site are armored with riprap or are separated from the Marine Area with sheet pile bulkheads to prevent erosion. In the Upland Area, the ground surface is mostly paved with asphalt or concrete. In limited portions of the Upland Area, the ground surface consists of a crushed gravel working surface that is maintained for fabrication layout and equipment storage. As a result, the Site contains little to no vegetation that would serve as riparian or terrestrial habitat.

During development of the RI/FS Work Plan, Ecology requested that the Port complete a Terrestrial Ecological Evaluation (TEE) to determine if ecological based soil cleanup levels were applicable to the Site. The goal of the TEE process is the protection of terrestrial ecological receptors from exposure to contaminated soil with the potential to cause significant adverse effects. For species protected under the Endangered Species Act (ESA) or other applicable laws that extend protection to individuals of a species, a significant adverse effect means an impact that would significantly disrupt normal behavior patterns that include, but are not limited to, breeding, feeding, or sheltering. For all other species, significant adverse effects are effects that impair reproduction, growth or survival.

In accordance with WAC 173-340-7491, a simplified TEE was completed for the Site. The results of the exposure analysis determined that the existing land surface (asphalt, concrete, compacted gravel, buildings, etc.) at the site and surrounding area make substantial wildlife exposure unlikely based on completion of Table 749-1. Ecology, during a visit in August 2008 to observe the condition of the Site confirmed that the working surface provided little to no habitat value. Additional paving of the previous gravel surfaces has occurred since that time, further reducing the potential for habitat at the Site.

The process specified under MTCA for identifying the requirements of a TEE (WAC 173-340-7491 and -7492) for the Site and results of the simplified TEE are presented in Appendix B.

1.4.8.2. Groundwater Potability

The City owns and operates a Class A water system which serves a much larger area than the sewer system boundary. The water system serves approximately 56,000 customers, with regional customers that include two refineries, the Skagit Public Utilities District, the town of La Conner, the Swinomish Tribal Community, and the City of Oak Harbor, including the Naval Air Station Whidbey Island. Because drinking water for the



Site and vicinity is supplied by the City, water supply wells are not known to exist at or near the Site, and groundwater beneath the Site is not used as drinking water. Groundwater at the Site is classified as non-potable.

Specific criteria used to evaluate groundwater potability (WAC 173-340-720(2)) and their applicability to the Site, are as follows:

- The ground water does not serve as a current source of drinking water WAC 173-340-720(2)(a).
 Applicability: Drinking water is currently supplied by the City. Water supply wells are not known to exist at or near the Site.
- 2. The Department (Ecology) determines it is unlikely that hazardous substances will be transported from the contaminated ground water to ground water that is a current or potential future source of drinking water, as defined in (a) and (b) of this subsection [i.e., -720(2)], at concentrations which exceed ground water quality criteria published in Chapter 173-200 WAC WAC 173-340-720(2)(c).
 - Applicability: Contaminated groundwater beneath the Site occurs in an unconfined shallow water-bearing zone contained within artificial fill. Shallow groundwater at the Site discharges directly to Guemes Channel and is not known to flow toward other aquifers that may be a current or potential future source of drinking water.
- 3. Even if ground water is classified as a potential future source of drinking water, the Department recognizes that there may be sites where there is an extremely low probability that the ground water will be used for that purpose because of the site's proximity to surface water that is not suitable as a domestic water supply. An example of this situation would be shallow ground waters in close proximity to marine waters such as on Harbor Island in Seattle. At such sites, the Department may allow ground water to be classified as non-potable if each of the following conditions can be demonstrated. These determinations must be for reasons other than that the ground water or surface water has been contaminated by a release of a hazardous substance at the site WAC 173-340-720(2)(d).
 - a. There are known or projected points of entry of the ground water into the surface water WAC 173-340-720(2)(d)(i).
 - Applicability: Groundwater at the Site is in close proximity to the Guemes Channel which is tidally influenced. This tidal influence results in the tidal exchange of saline surface water and upland groundwater within the Site as observed during groundwater monitoring activities (Section 5.3). Water quality parameters measured during monitoring activities show that total dissolved solids (TDS) in several wells located throughout the Site have elevated concentrations indicative of slightly saline (greater than 1,000 milligrams per liter [mg/L] TDS at wells MW-1 and MW-7) to highly saline (greater than 10,000 mg/L TDS at wells MW-2, MW-3 and MW-6).
 - b. The surface water is not classified as a suitable domestic water supply source under Chapter 173-201A WAC WAC 173-340-720(2)(d)(ii).
 - **Applicability**: Guemes Channel is a marine surface water body and is not suitable as a domestic water supply under Chapter 173-201A WAC.
 - c. The ground water is sufficiently hydraulically connected to the surface water that the ground water is not practicable to use as a drinking water source WAC 173-340-720(2)(d)(iii).



Applicability: The shallow water-bearing zone at the Site is directly connected with and discharges into the Guemes Channel. It is not practicable to utilize the shallow aquifer for water supply due to the potential for drawing saline water into the aquifer (i.e., saltwater intrusion).

1.4.9. Cultural Resources

Guemes Channel connects Rosario Strait with Fidalgo and Padilla Bays, which are high-priority, "early-action" cleanup areas under the Puget Sound Initiative. Ecology is working with stakeholders, including tribes, to keep them informed of the cleanup of contaminated sites and sediments in the vicinity of the Fidalgo/Padilla Bay areas. Tribes that are interested in engaging with Ecology under the Puget Sound Initiative at Fidalgo/Padilla Bays include the Swinomish, Samish, Upper Skagit, Suquamish, Skagit River System Cooperative, Tulalip and Lummi Tribes.

Cultural records (Lenz 2013) indicate that the Samish occupied the shoreline areas of Guemes Channel. Large historical middens representing winter villages and smaller sites related to camping and shellfish gathering are common in similar settings. Currently, no archaeological or culturally important sites are known to exist at the Site.

1.5. Ecological Setting

The Site is located on Fidalgo Island along the southern shoreline of Guemes Channel (Figure 1). Properties located to the west and south have industrial use and properties located to the east have commercial and residential uses. Guemes Channel to the north provides habitat for various marine fish, anadromous salmonids and invertebrate species of commercial and recreational value. The area also provides seasonal habitat for adult marine mammals, seabirds and other waterfowl.

The United States Fish and Wildlife Service (USFWS) Information for Planning and Consultation (IPaC) report (USFWS 2019) includes a total of nine threatened, endangered, or candidate species and one critical habitat on the species list known to occur, or potentially occur, within an approximate 5-mile radius Site including:

Mammals:

- Grey Wolf (Canis lupus) Proposed Endangered
- North American Wolverine (Gulo gulo luscus) Proposed Threatened

Birds:

- Marbled Murrelet (Brachyramphus marmoratus) Threatened
- Northern Spotted Owl (Strix occidentalis caurina) Threatened
- Streaked Horned Lark (Eremophila alpestris strigata) Threatened
- Yellow-billed Cuckoo (Coccyzus americanus) Threatened

Fishes:

- Bull trout (Salvelinus confluentus) Threatened
- Dolly Varden (Salvelinus malma) Proposed Similarity of Appearance (Threatened)

Flowering Plants:

■ Golden paintbrush (Castilleja levisecta) – Threatened



Critical Habitat:

Bull trout (Salvelinus confluentus) – Critical Habitat

The National Oceanic and Atmospheric Administration (NOAA) Fisheries Status of ESA Listings & Critical Habitat Designations for the West Coast region (NOAA 2019) includes a total of four threatened and three endangered species their critical habitat with potential to occur at the Site including:

- Puget Sound Steelhead (Oncorhynchus mykiss) Threatened & Critical Habitat
- Puget Sound Chinook Salmon (Oncorhynchus tshawytscha) Threatened & Critical Habitat
- Southern Resident DPS orcas (Orcinus orca) Endangered & Proposed Critical Habitat
- Western DPS Steller sea lion (Eumetopias jubatus) Endangered & Critical Habitat
- Puget Sound/Georgia Basin DPS bocaccio (Sebastes paucispinis) Endangered & Critical Habitat
- Puget Sound/Georgia Basin DPS yelloweye rockfish (Sebastes ruberrimus) Threatened & Critical Habitat
- Puget Sound/Georgia Basin DPS canary rockfish (Sebastes pinniger) Threatened & Critical

Within a mile of the assessment area, the Washington Department of Fish and Wildlife (WDFW) Priority Habitat and Species (PHS) reports document surf smelt and Pacific herring breeding areas, Dungeness crab presence, cliffs/bluffs and a biodiversity area/corridor (Cap Sante Park; WDFW 2018).

1.6. Regulatory Framework

Environmental studies completed at the Site since approximately 1991 have identified that historical uses including vessel moorage, bulk fuel and oil storage, and shipbuilding activities have resulted in the release of contaminants to soil, groundwater and sediment. Partial cleanup of the Site has been completed as voluntary actions by the Port whereas, the final cleanup will be completed under a formal order with Ecology.

1.6.1. Voluntary Cleanup Action

The Port has completed the following voluntary cleanup actions to address known soil contamination:

- 1991 UST Cleanup Action In 1991, two underground storage tanks (USTs) located near the south end of L dock were removed from the Site for permanent closure. During the removal of these tanks, approximately 20 cubic yards of petroleum impacted soil was removed from this area and transferred from the Site for landfill disposal. Verification samples at the final excavation limits were obtained to confirm the removal of the petroleum impacted soil observed during tank removal activities.
- 2001 Hydraulic Winch Cleanup Action In 2001, a hydraulic winch and its timber frame located near the south end of the east marine railway were removed from the Site. During removal of this structure and associate components, approximately 30 cubic yards of petroleum impacted soil were excavated and transferred from the Site for landfill disposal. Verification samples at the final excavation limits were obtained to confirm the removal of the petroleum impacted soil observed during removal of the hydraulic winch and associated timber frame.
- 2002 Petroleum and Marine Railways Cleanup Actions In 2002, the Port completed a voluntary cleanup action to address known soil contamination in the Petroleum Cleanup Action Area extending



from the aluminum shop (building formerly identified as the equipment maintenance shed) to the former bulk fuel storage ASTs; and the Marine Railway Cleanup Action Area located near the eastern marine railway structure. Cleanup actions were completed under Ecology's Voluntary Cleanup Program (VCP) to remove soil with contaminant concentrations exceeding cleanup levels developed for the Site (Landau 2002a). The extent of the VCP cleanup actions is shown on Figure 6. Upon completion of the remedial excavation activities, discrete confirmation samples from the excavation sidewalls and base were collected to verify the removal of soil contamination in these areas. Based on the verification sample results, previously identified contamination in these areas was successfully removed from the Site and the excavation areas were backfilled to the original grade with clean imported soil (Landau 2002b). Although the cleanup actions confirmed the removal of soil contamination in these areas, the work was never finalized under the VCP.

1.6.2. Ecology Agreed Order

On December 12, 2007, the Port entered Agreed Order No. DE-07TCPHQ-5080 with Ecology. Under the Agreed Order, the Port is required to complete an interim cleanup action in the Marine Area, evaluate the nature and extent of contamination in affected media on a Site-wide basis and develop and evaluate cleanup alternatives for addressing the identified contamination including:

- Preparation of a RI/FS Work Plan (completed);
- Field data collection to fill data gaps identified in the Work Plan (completed);
- Completion of an interim remedial action to remove contaminated media from the Marine Area prior to redevelopment activities (completed);
- Preparation of RI/FS document to present the results of the field data collection and to identify and evaluate cleanup alternatives for contaminated media at the Site (subject of this report); and,
- Preparation of a Draft Cleanup Action Plan (DCAP) provides a proposed remedial action to address the contamination present on the Site (*future task*).

Completion of the RI/FS and DCAP documents will fulfill the remaining work requirements required by the Agreed Order. Field data collection in accordance with the Ecology-approved RI/FS Work Plan is summarized in Section 3.5. Interim Action activities completed in accordance with the Ecology-approved RI/FS Work Plan and Interim Action Work Plan Addendum to remove previously identified sediment contamination as part of the Port's Project Pier 1 redevelopment is summarized in Section 4.0.

The final cleanup action at the Site as determined by the Cleanup Action Plan (CAP) is anticipated to be completed under a Consent Decree between the Port and Ecology.

2.0 CLEANUP STANDARDS

Cleanup standards consist of: 1) cleanup levels that are protective of human health and the environment; 2) the point of compliance at which the cleanup levels must be met; and 3) additional regulatory requirements, specified in applicable state and federal laws, that apply to a cleanup action because of the type of action and/or the location of the Site.



Preliminary cleanup levels) for Site media of concern including sediment, groundwater and soil were developed during preparation of the RI/FS Work Plan. These preliminary cleanup levels have been updated (since development of the RI/FS Work Plan) to meet the current MTCA standards and are the proposed cleanup levels (PCULs) for defining the nature and extent of Site contamination, for developing cleanup action objectives, and developing remedial alternatives for the Site. Sediment PCULs for protection of benthic organisms and protection of human health and higher trophic level ecological receptors are presented in Tables 1 and 2, respectively. PCULs for groundwater and soil are presented in Tables 3 and 4, respectively. Sediment, groundwater and soil PCULs, and points of compliance for each of these media are discussed below.

2.1. Proposed Sediment Cleanup Levels

The RI/FS Work Plan included preliminary cleanup levels for sediment protective of benthic organisms using Sediment Quality Standards (SQS) and Cleanup Screening Level (CSL) criteria established under the Sediment Management Standard (SMS; WAC 173-204) available at the time. In December 2019, Ecology issued the revised Sediment Cleanup User's Manual II (SCUM II, Ecology 2019) as a guidance document for implementing the cleanup provisions of the SMS under WAC 173-204 that included development of cleanup levels protective of benthic organisms, and cleanup levels protective of human health and higher trophic level ecological receptors. Preliminary cleanup levels developed for the RI/FS Work Plan revised to meet current standards are the PCULs for sediment and are presented in Tables 1 and 2 and are further discussed in the following sections.

2.1.1. Proposed Sediment Cleanup Levels for the Protection of Benthic Organisms

Sediment PCULs for benthic invertebrate community health are the numeric Sediment Cleanup Objectives (SCO) from SMS that correspond to sediment quality that will result in no adverse effects to the benthic community (WAC 173-204-562). PCULs for protection of benthic organisms are presented in Table 1.

The SMS benthic community health-based sediment cleanup objective of WAC 173-204-562 provide numeric criteria for a broad range of chemicals. The benthic community health-based criteria for specific chemicals are based on either dry-weight or organic carbon-normalized concentrations. The analytical results for nonpolar organics are organic carbon normalized when the total organic carbon (TOC) concentration for a sample range from 0.5 to 3.5 percent (inclusive). The carbon normalized analytical results are then compared to the organic carbon-normalized SCO. Analytical results for nonpolar organics that include samples with TOC concentrations outside of the 0.5 to 3.5 percent range are screened against Marine Sediment Apparent Effects Threshold (AET) values on a dry-weight basis (Table 8-1 of SCUM II). SMS and AET screening level criteria for benthic community health are presented in Table 1.

2.1.2. Proposed Sediment Cleanup Levels for the Protection of Human Health and Higher Trophic Level Ecological Receptors

Sediment PCULs for protection of human health and protection of higher trophic level ecological receptors are presented in Table 2. PCULs for human health exposure to sediment via ingestion and dermal contact were developed utilizing equations and parameter values from Ecology's SCUM II guidance.

The sediment PCULs based on sediment ingestion and dermal contact shown in Table 2 represent the values for an adult exposed during net fishing (subsistence harvesting). Based on current Site conditions (i.e., deep-water berth with no accessible intertidal beach zone), the exposure scenario is expected to only



apply to subtidal sediments that are below -3 feet MLLW. Therefore, exposure to sediment by children during beach play and adults to sediment during clam digging in the intertidal zone above -3 feet MLLW is not applicable to the Marine Area.

Tissue data do not exist for the Site and site-specific biota-sediment accumulation factors (BSAFs) are not available to back-calculate risk-based sediment PCULs. Therefore, a simplified approach (Option 1 within SCUM II – Section 9.2) where the SCO and CSL are established at background (natural and regional, respectively) or the practical quantitation limit (PQL) was selected to develop sediment PCULs based on bioaccumulation exposure for human health and higher trophic level organisms. For bioaccumulative chemicals such as dioxins/furans, total polychlorinated biphenyls (PCBs), polycyclic aromatic hydrocarbons (PAHs), arsenic, cadmium, lead and mercury, sediment screening levels based on bioaccumulation are based on either the 90/90 Upper Tolerance Limit (UTL) on the mean natural background concentrations derived from the entire Bold Plus dataset (DMMP 2009; see SCUM II, Table 10-1), or the Ecology-accepted PQL, whichever is higher. Sediment PCULs for human health and higher trophic level ecological receptors were chosen from the lowest of bioaccumulative and direct contact pathways. The PCULs for subtidal areas include marine areas at elevations below -3 feet MLLW and the applicable direct contact pathway is net fishing.

Consistent with the SCUM II guidance, where the risk-based value is lower than natural background or PQL, the PCUL defaults to the higher of natural background or PQL. Table 2 presents the natural background, regional background, PQL and the PCUL level selected for each chemical.

2.1.3. Point of Compliance in Sediment

In accordance with SMS requirements, the point of compliance for protection of benthic organisms and human health and higher trophic level species exposure in subtidal sediment is represented by the biologically active zone within the uppermost 10 centimeters (cm) below mudline.

2.2. Proposed Groundwater Cleanup Levels

The Site meets the definition of an industrial property under MTCA (WAC 173-340-200) as it is zoned for industrial use and has been and is being used for industrial purposes. The Site also meets the requirements for use of industrial cleanup levels for soil as hazardous substances remaining at the property do not pose a threat to human health and the environment in non-industrial areas (WAC 173-340-745[1][a][iii]). The Site is surrounded by properties that are also zoned for industrial land use. The location and use of the Site and land use in the surrounding area restricts access by the general public. Based on zoning, current and anticipated future land use, PCULs for groundwater were selected from the most conservative (lowest) published values from the following transport and exposure pathways:

- Acute and chronic effects to aquatic organisms resulting from exposure to contaminants in surface water and sediment where groundwater discharges to adjacent marine surface water;
- People consuming seafood exposed to contaminants in surface water and sediment where groundwater discharges to adjacent marine surface water; and
- People inhaling volatile organic compounds in enclosed spaces resulting from vapor intrusion.

As discussed in Section 1.4.8, groundwater at, or potentially affected by, the Site is not used for drinking water at this time and is not a reasonable future source of drinking water due to its proximity to marine



surface water and the availability of a municipal water supply. In accordance with WAC 173-340-720(2)(d), Site groundwater qualifies as a non-potable water source, therefore, people ingesting hazardous substances in groundwater is not a potential exposure pathway.

Groundwater PCULs that were selected are the lowest of the applicable numerical values from the regulatory criteria presented below. In accordance with WAC 173-340-705(6), the PCULs were adjusted as necessary based on Washington State groundwater background concentrations for metals (PTI 1989) and PQL to derive the groundwater PCULs such that groundwater PCULs for a given constituent shall not be set at a level below the natural background concentration or the PQL, whichever is higher. The PQLs were referenced from the Ecology-approved RI/FS Work Plan which were obtained from OnSite Environmental, Inc. (OnSite) of Redmond, Washington, a Washington-certified laboratory.

Groundwater PCULs for the Upland Area are presented in Table 3 and are further discussed in the following sections.

2.2.1. Proposed Groundwater Cleanup Levels for the Protection of Surface Water

Groundwater PCULs were selected from available state and federal surface water criteria according to WAC 173-340-730(3). The most conservative (lowest) published values were selected from the following regulatory criteria:

- Water Quality Standards for Surface Waters of the State of Washington. These marine surface water criteria for protection of aquatic life (acute and chronic exposures) and human health (fish consumption) are published in Chapter 173-201A WAC.
- Federal Marine Water Quality Criteria for Washington State. These criteria are from United States Environmental Protection Agency's (EPA's) Final Revision of Federal Human Health Criteria Applicable to Washington from 40 CFR 131.45 (EPA 2016).
- Federal National Recommended Water Quality Criteria. These marine surface water criteria for protection of aquatic life (acute and chronic exposures) and human health (fish consumption) are established under Section 304 of the Clean Water Act.
- MTCA Method B standard formula values (for carcinogens and non-carcinogens) protective of human health (consumption of aquatic organisms) (WAC 173-340-730[3]).
 - Surface water criteria are not currently available for gasoline-, diesel, and oil-range petroleum hydrocarbons. Therefore, as recommended in WAC 173-340-730(3)(b)(iii)(C), the MTCA Method A groundwater cleanup levels for gasoline-, diesel, and oil-range petroleum hydrocarbons were used as the MTCA Method B surface water cleanup levels for these analytes.

2.2.2. Proposed Groundwater Cleanup Levels for the Protection of Sediment

Groundwater concentrations protective of sediment were calculated assuming equilibrium partitioning between sediment and groundwater in sediment pore spaces. The following equation, from Ecology's Lower Duwamish Waterway Preliminary Cleanup Level Workbook Supplemental Information document dated December 2018, was used to calculate groundwater concentrations protective of dry weight SCO criteria:



Equation:

 $C_w = SCO/(CF \times DF [K_d + \theta_w/\rho_b])$

Where:

C_w = groundwater concentration protective of sediment (µg/L)

SCO = sediment cleanup objective (WAC 173-204-560[3]) (mg/kg dry weight)

CF = conversion factor (0.001 mg/µg)

DF = dilution factor (unitless) (default value of 1 for saturated sediment)

 K_d = soil-water distribution coefficient (L/kg). K_d for organic chemicals is calculated per MTCA Equation 747-2 below.

 $\theta_{\rm w}$ = water-filled porosity (0.615 ml/ml)

 ρ_b = dry sediment bulk density (1.02 kg/L)

Equation:

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

Where:

 K_{oc} = soil organic carbon-water partitioning coefficient (ml/g) (0.019 g/g)

 F_{oc} = sediment fraction organic carbon (g/g)

Values for Kd and Koc are from Ecology's "CLARC Master Spreadsheet.xlsx" dated May 2019.

2.2.3. Proposed Groundwater Cleanup Levels for the Protection of Vapor Intrusion

PCULs were developed for the groundwater to indoor air or vapor intrusion transport pathway. The groundwater to vapor intrusion transport pathway used in this RI are based on values for industrial land use. As described above, the Site meets the definition of an industrial property under MTCA (WAC 173-340-200) as it is zoned for industrial use and is being used for industrial purposes now and for the foreseeable future.

2.2.4. Point of Compliance for Groundwater

Groundwater at the Site does not meet the definition of potable water as outlined in WAC 173-340-720(2) based on the following factors: a) the groundwater does not serve as a current source of drinking water; and b) the groundwater is not a potential future source of drinking water given the Site's proximity to surface water that is not suitable as a domestic water supply (Section 1.4.8). Therefore, a conditional point of compliance located as close as technically possible to the point where groundwater discharges to surface water is proposed.

2.3. Proposed Soil Cleanup Levels

The Site meets the definition of an industrial property under MTCA (WAC 173-340-200) as it is zoned for industrial use and has been and is being used for industrial purposes (Section 1.1). The Site also meets the requirements for use of industrial cleanup levels for soil as hazardous substances remaining at the property do not pose a threat to human health and the environment in nearby non-industrial areas



(WAC 173-340-745[1][a][iii]). The Site is surrounded by properties that are also zoned for industrial land use. Residential areas are not located in proximity to the Site. The location and use of the Site and land use in the surrounding area restricts access to the Site by the general public. Based on zoning, current and anticipated future land use, soil PCULs were selected from the most conservative (lowest) published values from the following transport and exposure pathways:

- People ingesting soil through direct soil contact.
- Soil to groundwater transport pathway; including the protection of surface water via groundwater and protection of sediment via groundwater transport pathways.

As stated above (Section 1.4.8), groundwater at the Site is non-potable. Therefore, the soil PCULs used in this RI and discussed below include those derived for protection of non-potable groundwater. In addition, a TEE completed for the Site (Section 1.4.8) indicates that the Site is exempt under WAC 173-340-7491(1)(b) such that soil at the Site is covered by physical barriers (such as buildings or paved roads/working surfaces) that prevent exposure to plants and wildlife.

Soil PCULs that were selected are the lowest of the applicable numerical values from the regulatory criteria presented below. In accordance with WAC 173-340-705(6), the PCULs were adjusted as necessary based on background concentrations and PQLs to derive the soil PCULs such that soil PCULs for a given constituent shall not be set at a level below the natural background concentration or the PQL, whichever is higher. Natural background concentrations (except for arsenic) were referenced from Ecology Publication 94-115 "Natural Background Soil Metals Concentrations in Washington State" (Ecology 1994) using 90th percentile values published for the Puget Sound Basin. The arsenic background is established in regulation and published as the MTCA Method A value. The PQLs were obtained from OnSite of Redmond, Washington, a Washington-certified laboratory and presented in the Ecology-approved RI/FS Work Plan.

Soil PCULs for the Upland Area are presented in Table 4 are further discussed in the following sections.

2.3.1. Proposed Soil Cleanup Levels for the Protection of Human Health

Soil PCULs for the for protection of human health were identified from MTCA standard Method C soil cleanup levels for industrial land use – soil direct contact (WAC 173-340-745(5)(b)(iii)(B)). MTCA Method A soil cleanup levels for industrial land use (WAC 173-340-745[3]) are used for analytes without Method C soil cleanup levels, which include lead and petroleum hydrocarbons. Note that the Method A soil cleanup level for total PCBs is based on applicable federal law (40 CFR 761.61).

2.3.2. Proposed Soil Cleanup Levels for the Protection of Groundwater

Screening levels were developed for the soil to groundwater transport pathway using the MTCA fixed parameter three-phase partitioning model (WAC 173-340-747[4]). Default assumptions provided in WAC 173-340-747(4)(b) (Equation 747-1 and Equation 747-2) for saturated zone soils and Ecology default model input parameter values (soil organic carbon-water partition coefficient [K_{oc}] Henry's Law constants) were used in the calculations.



2.3.3. Point of Compliance for Soil

Soil PCULs developed for the Site are based on the protection of surface water via groundwater and protection of sediment via groundwater transport pathways. Therefore, the point of compliance for the soil will be throughout the soil column in accordance with WAC 173-340-740(6)(b).

3.0 SITE CHARACTERIZATION STUDIES

3.1. Historical Sediment Characterization

Historical environmental studies completed to assess sediment quality in and near the Marine Area included the following:

- Phase 2 Environmental Assessment (Otten Engineering 1997)
- Marine Area Surface Dioxin Study (Floyd | Snider 2007)
- Fidalgo Bay Sediment Investigation (SAIC 2008)

The sediment characterization studies listed above resulted in the collection of surface samples ranging between 0 and 20 cm below the mudline at 20 locations. The sediment samples were submitted for a combination of analyses including TOC, total solids (TS), and grain size, SMS metals, semi-volatile organic compound (SVOCs), PAHs, volatile organic compound (VOCs), PCBs, pesticides, tributyltin (TBT), and dioxins and furans. The schedule of laboratory analysis for sediment samples as part of these studies are summarized in Table 5. Historical sediment sample locations within the Marine Area are shown on Figure 7. Historical sediment sample results are presented in Tables C-1 and 2 (Appendix C) and are summarized in the following sections.

3.1.1. Phase 2 Environmental Assessment (1997)

In August 1997, Otten Engineering collected surface (0 to 10 cm) sediment samples at three locations (DC-SED-01 through DC-SED-03) from the intertidal portion of the Marine Area using hand tools during low tide. In addition, four surface sediment samples (DC-SED-05, DCI-SED-06, DCI-SED-08 and DC-SED-9) were collected from the subtidal portion of the Marine Area using a grab sampler deployed from a research vessel. Samples collected from these locations were submitted for a combination of metal, TOC, TS, TBT, PAH and PCB analysis.

The results of the chemical analysis identified detectable concentrations of metals including arsenic, cadmium, chromium, copper, lead, mercury, nickel, silver and zinc, TBT, low molecular weight PAHs (LPAHs), high molecular weight PAHs (HPAHs) and PCBs. Sediment sample results for this study are summarized in Tables C-1 and C-2 (Appendix C).

3.1.2. Marine Area Surface Dioxin Study (2006)

In 2006, Floyd Snider collected surface (0 to 20 cm) sediment samples at locations DCl06-1 through DCl06-9 within the Marine Area for dioxin and furan analysis to further evaluate potential discharges from former Scott Paper Mill operations. Total dioxin and furan toxic equivalency quotients (TEQs) in sediment samples were less than the natural background level of 5 nanograms per kilogram (ng/kg), except for surface sediment at locations DC-106-4 through DC-106-8 collected in the vicinity of the east marine railway. Dioxin and furan results for this study are summarized in Tables C-1 and C-2 (Appendix C).



3.1.3. Fidalgo Bay Sediment Investigation (2007)

In 2007, Science Applications International Corporation (SAIC) completed a sediment investigation of the aquatic areas of Fidalgo Bay for Ecology. The objectives of this investigation were to conduct a multi-faceted, tiered sediment characterization in order to help define the nature and extent of the sediment contamination in Fidalgo Bay. Investigation activities included sediment profile imaging (SPI), surface sediment chemistry, and sediment toxicity testing.

Within the study's sub area located nearest the Marine Area (DUA 4, decision area encompassing the Site), a total of 36 SPI images were analyzed from 30 stations. At sample stations FB-4A-14, FB-4A-15 and FB-4A-17 located in close proximity to the Site, fine to medium sand with occasional gravel was identified as the prominent shallow (0 to 10 cm) substrate consistent with an intermediate energy environment. The results of the chemical analysis identified detectable concentrations of metals including arsenic, cadmium, chromium, copper, lead, mercury, silver and zinc, SVOCs including LPAHs and HPAHs, phthalates, miscellaneous extractables and phenols. Sediment sample result for sample stations FB-4A-14, FB-4A-15 and FB-4A-17 are summarized in Tables C-1 and C-2 (Appendix C).

3.1.4. DCI Basin Dredged Material Characterization and Recency Determination (2000 and 2004)

Between April 2000 and July 2004, surface and subsurface sediment samples were also collected from the Marine Area for the purpose of DMMP dredge material characterization and suitability determination (Hart Crowser 2000 and Anchor 2004). For the purposes of the dredge material characterization, the Marine Area was divided into two Dredge Material Management Units (DMMUs). DMMU-D1 encompassed the outer half of the Marine Area while DMMU-D2 encompassed the nearshore half of the Marine Area (Figure 7).

In April 2000, Hart Crowser completed an initial dredged material characterization. During this study, discrete samples from 0 to 10 cm were collected throughout DMMU-D1 and DMMU-D2 that were later composited for laboratory analysis of DMMP parameters to evaluate open-water disposal suitability. Based on the results of this initial study and previous environmental studies completed, the DMMP determined that surface and native subsurface (i.e., hard glacially compacted native sediments) sediment within DMMU-D1 was suitable for unconfined open-water disposal. The DMMP also determined that native sediment from DMMU-D2 was also suitable for unconfined open-water disposal. However, surficial fill material from DMMU-D2 was determined to not be suitable for open-water disposal.

To evaluate potential dioxin contamination from the former Scott Paper Mill outfall located in the vicinity of the proposed dredge prism, Anchor Environmental completed a supplemental dredged material characterization in July 2004 within the Marine Area and vicinity. As part of this study, subsurface sediment ranging in depth between 0 and 5 feet below the mudline were collected for laboratory analysis of dioxins and furans. Total dioxin and furan TEQs for these samples were less than the natural background level of 5 ng/kg. Based on these supplemental results, the DMMP determined that the sediment characterized by these samples were suitable for unconfined open-water disposal under the existing DMMP evaluation framework at the time of the determination.

The recency date for the initial April 2000 dredged material characterization of the Marine Area was set to expire in April 2005. However, the supplemental data collected during the July 2004 dredge material characterization recency evaluation suggested that sediment quality had not changed since the initial characterization, and that the recency date could be extended to July 2009. This recency extension



maintained that surficial fill material above the hard glacially compacted native sediment in DMMU-D2 was not suitable for open-water disposal. Dredged Material suitability and recency determination by the DMMP is presented in Appendix D.

In 2008, the Port completed dredging of the Marine Area as part of their Project Pier 1 redevelopment activities. The contaminated material not suitable for open-water disposal was removed from the Site and transferred to an upland landfill as part of an Ecology-approved Interim Action. Following removal of the contaminated material, the remaining dredge prism approved by the DMMP for unconfined open-water disposal was dredged and transferred to the Rosario Strait disposal site. The Ecology-approved Interim Action is further discussed in Section 4.0.

3.2. Historical Groundwater Characterization

Historical environmental studies completed to assess groundwater quality in the Upland Area of the Site included the following:

- Remedial Investigation Study (Landau 2002b)
- Groundwater Characterization Study (Floyd | Snider 2007)

Groundwater characterization studies listed above included in the installation of four monitoring wells (MW-1 through MW-4) and collection of groundwater samples for chemical analysis to evaluate groundwater conditions at the Site. Groundwater samples were submitted for a combination of analysis including total/dissolved metals, gasoline-, diesel- and heavy oil-range petroleum hydrocarbons, VOCs, SVOCs including PAHs, pesticides and herbicides based on review of historical activities at the Site. The schedule of laboratory analysis for groundwater samples collected as part of these studies are summarized in Table 6. Groundwater sample locations are shown on Figure 8. Well completion logs for historical groundwater monitoring wells installed at the Site are presented in Appendix E. Historical groundwater sample results are presented in Table F-1 (Appendix F) and summarized in the following sections.

3.2.1. Remedial Investigation (2001)

Four groundwater monitoring wells (MW-1 through MW-4) were installed by Landau Associates (Landau) as part of the 2001 Remedial Investigation Study. Three of the monitoring wells (MW-1 through MW-3) were installed along the shoreline to evaluate downgradient groundwater conditions while the fourth monitoring well (MW-4) was installed at the south end of the Site near 3rd Street to evaluate upgradient conditions. Groundwater samples from these wells were collected during two separate monitoring events in September 2001 and October 2001 and were submitted for a combination of analyses including dissolved metals, gasoline-, diesel- and heavy oil-range petroleum hydrocarbons and VOCs based on a review of historical activities at the Site.

Results of the chemical analysis identified detectable concentrations of metals including arsenic, chromium, copper, nickel, mercury and zinc, and diesel- and heavy oil-range petroleum hydrocarbons. Other analytes evaluated including lead, gasoline-range petroleum hydrocarbons and VOCs were not detected in groundwater. Groundwater sample results for this study are presented in Table F-1 (Appendix F).



3.2.2. Groundwater Characterization Study (2006)

Floyd | Snider collected samples from monitoring well locations MW-1 through MW-4 on November 17, 2006 to further evaluate groundwater conditions at the Site. Petroleum hydrocarbons, the primary contaminant of concern for the 2002 independent cleanup action (summarized below), was not detected in groundwater samples collected from each four Site monitoring wells. In addition, other analytes evaluated either were not detected or were detected at concentrations less than 2002 VCP cleanup levels at monitoring well locations MW-1 through MW-4, except for arsenic at MW-4. Arsenic at monitoring well location MW-4 was detected at a concentration of 11.6 μ g/L which exceeded the Washington State background level of 8 μ g/L.

3.3. Historical Soil Characterization Studies

Historical environmental studies completed to assess soil quality in the Upland Area of the Site included the following:

- Phase 2 Environmental Site Assessment (Otten Engineering 1997)
- EPA Site Inspection (Weston 2001)
- Soil Characterization Study (Landau 2002b)

Soil characterization studies listed above included the collection of 17 shallow surface samples and completion of 25 subsurface explorations in the Upland Area of the Site. Soil samples were submitted for a combination of analyses including metals, gasoline-, diesel- and heavy oil-range petroleum hydrocarbons, organotins, VOCs, SVOCs including PAHs, pesticides based on review of historical Site uses and potential source areas. The schedule of laboratory analysis for soil samples collected as part of these studies are summarized in Table 7. Soil sample locations for these historical environmental studies are shown on Figure 9. Exploration logs for the historical soil explorations completed at the Site are presented in Appendix E. Historical soil sample results are presented in Tables G-1 and G-2 (Appendix G) and summarized in the following sections.

3.3.1. Phase 2 Environmental Site Assessment (1997)

In 1997, Otten Engineering completed a Phase 2 Environmental Assessment in the Upland Area of the Site to evaluate soil conditions. The Phase 2 Environmental Assessment included the collection of surficial samples from suspected source areas based on historical Site use and included areas near the synchrolift, in the vicinity of the marine railways and in the 1975 earth fill area (Figure 9). Samples were analyzed for metals, petroleum hydrocarbons, VOCs, SVOCs including PAHs, pesticides and PCBs.

Results of the chemical analysis identified detectable concentrations of metals including arsenic, cadmium, chromium, copper, lead, nickel, mercury, silver and zinc, gasoline-, diesel- and heavy oil-range petroleum hydrocarbons, VOCs, SVOCs including PAHs, pesticides and PCBs. Soil sample results collected for this study are summarized in Tables G-1 and G-2 (Appendix G).

3.3.2. Environmental Protection Agency (EPA) Site Inspection (2001)

Weston (on behalf of the EPA) collected 20 surface and subsurface soil samples at the Site during the 2001 EPA site inspection. During this study, five split samples were collected by Landau Associates (Landau) from near the marine railway and southwest of the former L dock.



Data results for samples collected by Weston are not available. Sample results from split samples collected by Landau identified detectable concentrations of metals, organotins, SVOCs, pesticides and PCBs. Soil sample results for split samples collected by Landau are summarized in Tables G-1 and G-2 (Appendix G).

3.3.3. Soil Characterization Study (2001)

In August 2001, soil sampling and analysis were completed to characterize soil conditions in the Upland Area of the Site. As part of this study, Landau completed 13 borings to evaluate up to six potential source areas based on historical land use and the results of previous environmental studies. Potential source areas evaluated as part of this study included:

- Former Welding Shop;
- Former Machine Shop;
- 1975 Earth Fill Area;
- Former AST and underground storage tanks (UST) Area (Petroleum Area);
- Paint House; and
- East Marine Railway.

Based on the results of the initial investigation, Landau collected soil samples from an additional three surface locations near the marine railway and 10 borings in the Petroleum Area on October 24, 2001. Borings completed at the Property were advanced to depths of 7 to 13 feet below ground surface (bgs). Results of the soil characterization study identified three areas of concern in which contaminants exceeded the 2002 VCP cleanup levels. The areas of concern included:

- Petroleum Area Metals including arsenic, copper, mercury, nickel, zinc, gasoline-, diesel-, and oil-range petroleum hydrocarbons, cPAHs, and PCBs were detected at concentrations greater than the 2002 VCP cleanup levels in the central portion of the Site south of the former East dock. In this area, identified soil contamination extended from approximately 1 to 7 feet bgs.
- Marine Railway Area Metals including arsenic, copper, mercury, nickel, zinc, gasoline- and diesel-range petroleum hydrocarbons were detected at concentrations greater than the 2002 VCP cleanup levels south of the former east marine railway. In this area, identified soil contamination extended from approximately 0 to 2 feet bgs.
- 1975 Earth Fill Area Metals including arsenic, copper, mercury, nickel, zinc, cPAHs, and methylene chloride were detected in surface and near-surface soil at concentrations greater than the 2002 VCP cleanup levels in the southwest portion of the Site. In this area, identified soil contamination extended from approximately 1 to 7 feet bgs.

3.4. Historical Cleanup Actions and Confirmation Sampling

Historical cleanup actions were completed following the removal and closure of three USTs; (A-1 1991), removal of a hydraulic hoist located near the east marine railway (Landau 2001) and to address soil contamination identified during Landau's 2002 Soil Characterization Study (Landau 2002c). Cleanup actions completed and confirmation soil sampling results are summarized in the following sections.



3.4.1. UST Removal and Closure (1991)

A-1 Pump Service oversaw the removal and closure of two USTs (one gasoline and one diesel) located near the south end of L dock in 1991 (Figure 3). The diesel UST, which was installed by DCI was in service less than six years before its removal. However, the age of the gasoline UST was not determined. As part of the UST removal and closure activities, approximately 20 cubic yards of soil was removed from this area and transferred from the Site for landfill disposal. At the limits of the UST removal excavation, verification base and sidewall samples were collected for petroleum hydrocarbon (gasoline and diesel) and benzene, ethylbenzene, toluene and total xylenes (BETX) analysis.

The UST removal and closure area and confirmation sample locations are shown on Figure 9. Confirmation soil sample results are summarized in Tables G-1 and G-2 (Appendix G). Based on a review of the confirmation sample results, gasoline-range petroleum hydrocarbon was detected in soil at concentrations ranging from 59 to 166 mg/kg and diesel-range petroleum hydrocarbon was detected in soil at concentrations ranging from 35 to 136 mg/kg. Other analytes evaluated including heavy oil-range petroleum hydrocarbons and BETX were not detected in base and sidewall samples collected from the UST removal excavation.

3.4.2. Marine Railway Hydraulic Winch Remedial Excavation (2001)

In July 2001, the hydraulic winch and its timber frame were removed from their former location south of the east marine railway (Figure 3). Petroleum hydrocarbon impacted soil resulting from historical releases to soil around the former winch and frame were excavated and transferred from the Site for landfill disposal. To verify the removal of the petroleum impacted soil, a total of six confirmation samples (VS-1 through VS-3 and VS-6 through VS-8) were collected by Landau from the remedial excavation limit.

The hydraulic winch remedial excavation area and confirmation sample locations are shown on Figure 9. Confirmation soil sample results are summarized in Tables G-1 and G-2 (Appendix G). Based on a review of the confirmation sample results, diesel- and heavy oil-range petroleum hydrocarbon were detected in soil at concentrations ranging from 6.6 to 1,900 mg/kg.

3.4.3. Independent Cleanup Action (2002)

Independent cleanup actions were completed to address soil contamination previously identified including the Petroleum Area, Marine Railway Area and 1975 Earth Fill Area. Cleanup of these areas were completed between August 19 and August 30, 2002. Based on the historical and future use of the Site and land zoning, MTCA Method A cleanup levels for petroleum hydrocarbons, lead, and PCBs and Method C cleanup levels industrial soil cleanup levels for all other analytes were established as the remediation levels for the independent cleanup action.

Cleanup actions for the Petroleum Area, Marine Railway Area, and 1975 Earth Fill Area are summarized in the following sections. Cleanup action areas and confirmation sample locations are shown on Figures 10 and 11. Confirmation soil sample results are summarized in Tables G-1 and G-2 (Appendix G).

3.4.3.1. Petroleum Cleanup Action Area

The Petroleum Area was defined as the area where soil with petroleum hydrocarbons (predominantly gasoline-range and diesel-range) concentrations exceeding the cleanup levels and known or suspected sources of releases had been identified. A remediation level of 2,000 mg/kg was established for diesel-and heavy oil-range petroleum hydrocarbons, and 100 mg/kg for gasoline-range petroleum hydrocarbons.



The cleanup action for the Petroleum Area extended from the location of a building formerly identified as an equipment maintenance shed to the former location of several ASTs (Figure 10).

Excavation depths ranged from 1.5 feet at the south end of the excavation, near the aluminum shop, to 8 feet in the area further north of the aluminum shop. Based on confirmation sample results, a total of approximately 1,300 cubic yards of contaminated soil was excavated and transferred from the Site for landfill disposal. An additional 1,300 cubic yards of excavated soil was temporarily stockpiled onsite, tested, and used as backfill in the completed excavation.

Forty-four confirmation samples were collected from the bottom and sidewalls of the excavation to verify the removal of the petroleum contaminated soil. Gasoline and/or diesel-range hydrocarbons exceeded remediation levels in five of these samples (CS-17, CS-19, CS-20, CS-26 and CS-38). Subsequently, soil represented by these samples was over-excavated and transferred from the Site for landfill disposal. To evaluate other contaminants of potential concern based on known or suspected sources, samples CS-30 and CS-33 were also submitted for chemical analysis of metals, PAHs, and PCBs. Metals, PAHs and PCBs were not detected except for lead, which was detected at a concentration of 8 mg/kg.

3.4.3.2. Marine Railway Cleanup Action Area

The Marine Railway Area was defined as the area near the east marine railway structure in which petroleum hydrocarbons and arsenic concentrations exceeded the 2002 VCP cleanup levels. A cleanup level of 2,000 mg/kg was established for diesel-range hydrocarbons and oil-range hydrocarbons and 100 mg/kg for gasoline-range hydrocarbons. For arsenic, a cleanup level of 88 mg/kg was established. The cleanup action for the Marine Railway Area included seven separate excavation areas (Figure 11).

Surface soil (0 to 1 feet bgs) was excavated to remove previously identified petroleum hydrocarbon contamination at four locations in the Marine Railway Area. The four approximately 10- by 10-foot excavations were centered on the sample locations from previous investigations with elevated petroleum hydrocarbons. Discrete confirmation samples were collected from base of each of these excavation areas to verify the removal of petroleum contamination. Petroleum hydrocarbons were not detected at concentrations greater than the 2002 VCP cleanup levels in each of completed excavation areas.

Two surface soil samples in the Marine Railway Area, that were collected during the 2001 EPA Site Inspection (described above), contained concentrations of PAHs that were elevated (with concentrations of individual PAHs up to 8.9 mg/kg), but below cleanup levels protective of the direct contact pathway. Even through the detected concentrations were below cleanup levels, the Port elected to excavate surface soil in this area and transport the soil offsite for disposal. No confirmation samples were collected from the excavation base at these locations to verify the removal of PAHs.

The main excavation in the Marine Railway Area (Figure 11) extended to depths ranging from 3.5 to 5 feet bgs. Approximately 300 cubic yards of soil were excavated from this area for landfill disposal. A total of seven confirmation samples were collected from the excavation bottom and sidewalls to verify the removal of petroleum contamination. Petroleum hydrocarbons were not detected in any of the confirmation samples collected from this area.

3.4.3.3. 1975 Earth Fill Area

Based on a review of historical Sanborn maps and aerial photographs, this area was used for residential purposes from before 1925 until after 1966. When DCI became a tenant on this parcel in 1975, this area



was topographically lower than the surrounding ground surface. This area is called the 1975 Earth Fill Area because fill material was used to bring it to grade around 1975.

The results of the Landau 2001 Soil Characterization Study identified petroleum hydrocarbons and PAHs in surface and near-surface soil in the 1975 Earth Fill Area at concentrations exceeding the 2002 VCP cleanup levels protective of surface water however, these concentrations did not exceed the MTCA Method C Cleanup Levels protective of direct contact with soil. Based on empirical evidence that groundwater was not being adversely affected by soil (i.e., groundwater results from MW-1) and that the results were less than the MTCA method C cleanup levels protective of direct contact with the soil, no cleanup action was completed for the 1975 Earth Fill Area.

3.4.4. Post-Independent Cleanup Action Groundwater Monitoring

Following completion of the 2002 Independent Cleanup Action, two rounds of groundwater monitoring (June 2002 and August 2002) were completed at monitoring well locations MW-1 through MW-4. The results of the sampling activities relative to the 2002 VCP cleanup levels are summarized below:

- Arsenic was detected in groundwater at concentrations greater than the Washington State background level of 8 μg/L at monitoring well location MW-4 during both the June 2002 and August 2002 monitoring events.
- Diesel-range petroleum hydrocarbons exceeded VCP cleanup levels at monitoring well locations MW-2 and MW-3 during the June 2002 monitoring event. However, during the August 2002 monitoring event, petroleum hydrocarbons were not detected in any of the wells sampled.

Other analytes evaluated at part of the post-independent cleanup action monitoring events either were not detected or were detected at concentrations less than the 2002 VCP cleanup levels.

3.5. 2007 Ecology Agreed Order Remedial Investigation Field Activities

As required by the 2007 Agreed Order, the Port completed a RI field investigation to fill data gaps in the characterization of Site sediment, groundwater and soil conditions and to define the nature and extent of contamination. The investigation activities completed to meet the objectives of the Ecology-approved RI/FS Work Plan are summarized in the following sections.

3.5.1. Contaminants of Potential Concern

As described in the Ecology-approved RI/FS Work Plan, Site contaminants of potential concern (COPCs) for the Marine Area sediment and Upland Area soil and groundwater were established for the RI based on a review of the previous environmental studies completed at the Site and historical/current land uses. The COPCs and rationale for their selection is summarized below.

MARINE AREA SEDIMENT CONTAMINANTS OF POTENTIAL CONCERN

Contaminants of Potential Concern	Rationale
Copper	Exceeded preliminary screening level protective of benthic organisms in surface (0-10cm) sediment at location DC-SED-03.



Contaminants of Potential Concern	Rationale
Lead	Exceeded preliminary screening level protective of human health and higher trophic level receptors in surface (0-10cm) sediment at location DC-SED-02 and DC-SED-03.
Mercury	Exceeded preliminary screening level protective of human health and higher trophic level receptors in surface (0-10cm) sediment in the composite DMMP sample for DMMU-D2.
Zinc	Exceeded preliminary screening level protective of benthic organisms in surface (0-10cm) sediment at location DC-SED-03.
Tributyltin (Bulk and Porewater)	Exceeded preliminary screening level protective of human health and higher trophic level ecological receptors in surface (0-10cm) at location DC-SED-05.
Low Molecular Weight Polycyclic Aromatic Hydrocarbons (LPAHs)	Exceeded preliminary screening level protective of benthic organisms and/or human health and higher trophic level ecological receptors in surface (0-10cm) sediment at locations DC-SED-02 and DC-SED-08, and in surface (0-10cm) sediment in the composite DMMP sample for DMMU-D2.
High Molecular Weight Polycyclic Aromatic Hydrocarbons (HPAHs)	Exceeded preliminary screening level protective of benthic organisms and/or human health and higher trophic level ecological receptors in surface (0-10cm) sediment at locations DC-SED-02, DC-SED-03 and DC-SED-08, and in surface (0-10cm) sediment in the composite DMMP sample for DMMU-D2.
Total cPAHs (TEQ)	Exceeded preliminary screening level protective of human health and higher trophic level ecological receptors in surface (0-10cm) at locations multiple locations within the Marine Area.
Total PCBs	Exceeded preliminary screening level protective of benthic organisms and/or human health and higher trophic level ecological receptors in surface (0-10cm) sediment at locations DC-SED-02, DC-SED-03 and DC-SED-08.
Dioxins/Furans	Exceeded preliminary screening level protective of human health and higher trophic level ecological receptors in surface (0-10cm) at locations DC-106-4, DC-106-5, DC-106-6 and DC-106-7.

The listed contaminants above were identified as COPCs based on preliminary cleanup level exceedances and were the focus of the sediment investigation completed for the RI. In addition, other metals (arsenic, cadmium, chromium and silver) and SVOCs (including polycyclic aromatic hydrocarbons [PAHs], phenols, phthalates, chlorinated organics and miscellaneous extractables) were also analyzed as part of the RI for consistency with SMS requirements.

UPLAND AREA SOIL AND GROUNDWATER CONTAMINANTS OF POTENTIAL CONCERN

Contaminants of Potential Concern (COPCs)	Rationale
Arsenic	Exceeded preliminary cleanup levels at multiple locations throughout the Upland Area in soil and groundwater.
Nickel	Exceeded preliminary cleanup levels at multiple locations throughout the Upland Area in soil and groundwater.
Gasoline-Range Hydrocarbons	Exceeded preliminary cleanup levels in upland soil in the 1975 Earth Fill and Petroleum Areas.



Contaminants of Potential Concern (COPCs)	Rationale
Diesel and Heavy Oil-Range Hydrocarbons	Exceeded preliminary cleanup levels in upland soil and groundwater in the Petroleum Area.
Total cPAHs (TEQ)	Exceeded preliminary cleanup levels at multiple locations throughout the Upland Area in soil.

The listed contaminants above were identified as COPCs based on preliminary cleanup level exceedances and were the focus of the upland soil and groundwater investigations completed for the RI. Additionally, other metals (including cadmium, chromium, copper, lead, mercury and zinc), SVOCs, VOCs (including BETX, methyl tertiary-butyl ether [MTBE], dibromoethane, 1-2 [EDB], dichloroethane, 1-2 [EDC]), and dioxins and furans were also analyzed as part of the RI to further evaluate subsurface conditions based on historical and current land use.

3.5.2. Sediment Remedial Investigation

The sediment RI was completed in general accordance with the Ecology-approved RI/FS Work Plan to characterize the vertical extent of sediment contamination in areas previously identified as exceeding the SMS criteria and to evaluate sediment quality in the Marine Area where no data previously existed. Sample locations are shown on Figure 12. Field procedures including sample handling, equipment decontamination and field screening are presented in Appendix H. Surface and subsurface sediment sampling activities are summarized in the following sections.

3.5.2.1. Sediment Sample Collection

The sediment samples were collected in from seven locations (G-1 through G-7) within the Marine Area (Figure 12). Sediment samples at locations G1, G2 and G7 were collected using a vibracore deployed from a research vessel. Sediment samples at locations G3 through G6 were collected from the upland area using a limited access direct-push drill rig during low tide. Surface and subsurface sediment samples were collected from cores that were advanced to depths ranging from approximately 4 to 7 feet below the mudline surface.

Upon collection, sediment samples were visually evaluated for the presence of wood debris, visually classified in accordance with ASTM International (ASTM) D 2488 methods and the Unified Soil Classification System (ASTM D 2487), homogenized in a stainless-steel bowl to a uniform in color and texture and placed into laboratory-prepared sample containers for analysis. Field screening results, observed wood content and a description of the material encountered during surface sediment sampling activities are summarized on the exploration logs presented in Appendix H.

3.5.2.2. Sediment Sample Laboratory Analysis

Samples collected for the sediment RI were submitted to Analytical Resources, Inc. (ARI) located in Tukwila, Washington. Selected samples were submitted for analysis of one or more of the COPCs identified above based on proximity to specific historical activities and previous sample results in accordance with the RI/FS Work Plan, including:

- Grain size by Puget Sound Estuary Program (PSEP) 1986 protocol;
- Total organic carbon (TOC) by PSEP protocol 1986;



- Total volatile solids (TVS) by PSEP protocol 1986;
- Total Solids (TS) by PSEP protocol 1986;
- Total Ammonia by EPA Method 350.1 M;
- Total Sulfides Standard Method (SM) 4500-S2;
- Tributyltin by EPA Method 8270D-SIM/KRONE;
- SMS metals by EPA Method 6000/7000 series;
- SMS SVOCs by EPA Method 8270/8270-SIM; and,
- PCBs by EPA Method 8082.

The schedule of analysis for sediment samples collected during the sediment RI are summarized in Table 5. Sediment sample results for this and previous environmental studies are presented in Table C-1 and C-2 (Appendix C) and were used as the basis for the 2008 Interim Action. Interim action activities including confirmation sampling are further discussed in Sections 4.0.

3.5.2.3. Deviations from RI/FS Work Plan

The surface and subsurface samples were collected in general accordance with the Ecology-approved RI/FS Work Plan. However, sediment sample location G-2 was moved approximately 15 feet to the west due to core refusal on the initial attempt. During the initial sampling attempt, the vibracore reached refusal at a depth of 2 feet below the mudline surface. At the new sample location, the vibracore was advanced to a depth of approximately 4 feet below mudline.

3.5.3. Groundwater Remedial Investigation

The groundwater RI was completed in general accordance with the Ecology-approved RI/FS Work Plan to characterize groundwater conditions at the shoreline where groundwater discharges to surface water, evaluate groundwater conditions within the Petroleum Area (independent cleanup action area), to estimate the hydraulic conductivity of the shallow aquifer, and to evaluate tidal influence on the shallow aquifer. Groundwater sampling activities are summarized in the following sections. Sample locations are shown on Figure 13. Field procedures including monitoring well installation and development, sample handling, equipment decontamination and field screening are presented in Appendix H. Groundwater RI field activities are summarized below.

3.5.3.1. Groundwater Sample Collection

The groundwater RI included the installation and development of new monitoring wells MW-2A, MW-2B, MW-3A, MW-5, MW-6, MW-7 and MW-8 and collection of groundwater samples from the four existing monitoring wells (MW-1 through MW-4) at the Site¹. Following installation, a GeoEngineers field representative completed monitoring well development and a field survey to obtain coordinates and top of casing rim elevation for new monitoring wells. Well installation, development, and surveying activities completed for MW-5 is summarized in Appendix H.

In accordance with the Ecology-approved RI/FS Work Plan, monitoring well MW-5 was initially installed in May 2008 for completion of the June 2008 monitoring event to evaluate groundwater conditions. Following

¹ Monitoring wells with the suffix "A" or "B" indicate a replacement well was installed in place of the original well.



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completion of the 2008 Interim Action in the Marine Area of the Site and reconfiguration of the shoreline by the Project Pier 2 redevelopment, Ecology required that four additional rounds of quarterly groundwater monitoring be completed to evaluate groundwater conditions. Quarterly groundwater monitoring events were completed in May 2012, August 2012, November 2012 and February 2013. In preparation for these monitoring activities, replacement wells MW-2A and MW-3A, and new monitoring wells MW-6 and MW-7 were installed in advance of the quarterly monitoring events to further evaluate groundwater conditions at the Site.

Due to inconclusive evidence linking contaminant exceedances identified in soil to contaminant exceedances in groundwater (further discussed in Section 5.5), Ecology required that four additional rounds of groundwater monitoring be completed on a semi-annual basis to further evaluate the potential source of soil contamination to groundwater (Ecology 2015). In addition, Ecology determined that the location of monitoring well MW-1 (Figure 13) may not adequately represent the conditional point of compliance in this area of the Site and that a new well (MW-8) be installed north of MW-1 to serve as the point of compliance. To avoid potential utilities, structural obstructions and minimize impacts to DCI's operations, MW-8 was positioned within Warehouse 9 located west of the Syncorlift Pier in an area which soil and groundwater conditions had not been previously evaluated.

During each monitoring event, groundwater samples were collected for chemical analytical testing using a peristaltic pump and disposable polyethylene tubing using low-flow/low-turbidity sampling techniques. A Horiba or YSI multi-probe field meter with a flow-through cell and/or Hach Turbidimeter were used to monitor water quality parameters during purging: electrical conductivity, dissolved oxygen, pH, salinity, TDS, turbidity, oxidation-reduction potential and temperature. Water samples were obtained once these parameters were measured to vary by less than 10 percent on three consecutive measurements. If water quality parameters did not stabilize, samples were collected after purging approximately three well-volumes.

3.5.3.2. Groundwater Sample Laboratory Analysis

Samples collected for the groundwater RI were submitted to CCI Analytical (CCI) located in Everett, Washington, ARI located in Tukwila, Washington, and OnSite located in Redmond, Washington. Samples were submitted for analysis of one or more of the COPCs identified based on proximity to specific historical activities and previous sample results in accordance with the RI/FS Work Plan and in discussions with Ecology following their review of the initial 2008 and 2012/2013 groundwater monitoring data. Groundwater samples were submitted for a combination of the following analysis:

- Total and dissolved metals (arsenic, cadmium, chromium, copper, lead, mercury, nickel, silver and zinc) by EPA Method 6000/7000 series;
- Gasoline-, diesel- and heavy oil-range petroleum hydrocarbons by NWTPH-Gx and NWTPH-Dx;
- VOCs by EPA Method 8260;
- SVOCs/PAHs by EPA Method 8270/SIM;
- Pesticides and herbicides by EPA Method 8081/8151; and
- Dioxins and Furans using EPA Method 8290 or EPA Method 1613B.

The schedule of laboratory analysis for samples collected during the groundwater RI are summarized in Table 6. Groundwater sample results for this and previous environmental studies are presented in Table F-1 (Appendix F) and discussed in Sections 5.4 and 5.5.



3.5.3.3. Hydraulic Conductivity Study

Hydraulic conductivity (K) testing was performed on June 16, 2008 on monitoring wells MW-1 through MW-5 within the shallow groundwater unit. The location of monitoring wells used to measure hydraulic conductivity are shown on Figure 13. Falling and rising head slug tests were performed on each well. Prior to conducting each slug test, an electronic water-level sensor consisting of a pressure transducer and automated datalogger was installed in the well. The hydraulic response was measured by the electronic water-level sensor, which was programmed to record the hydraulic pressure at 1-second intervals. Additionally, the depth to groundwater was measured manually using an electronic water level indicator to document the static groundwater level prior to initiating the slug tests as well as during and after each slug test.

The falling head slug test was performed by rapidly lowering a slug constructed of a sealed and weighted 5-foot-long section of polyvinyl chloride (PVC) pipe of known volume into the well causing the water level to rise rapidly above the initial, static water level. The groundwater level was then monitored until it returned (fell) to the approximate initial water level. Falling head tests were not evaluated for wells where the water table was within the well screen because the falling head response is affected by drainage into the unsaturated zone above the water table.

The rising head slug test was conducted following the falling head test in each well after recovery of the water table to the initial water level. Rising head tests were conducted by rapidly removing the slug from the well causing the water level to fall rapidly below the initial level. The groundwater level was monitored until it returned (rose) to the approximate initial water level. Wells where the water table was within the well screen interval were evaluated for hydraulic conductivity using the rising head data.

Data from the falling head or rising head tests were used to estimate hydraulic conductivity at each well using the Bouwer-Rice (1976) method. Field procedures for the slug tests are presented in Appendix H. The results of the slug tests are summarized in Section 5.3.

3.5.3.4. 72-hour Tidal Study

A 72-hour tidal study was completed between June 17 to June 20, 2008 using selected monitoring wells in the Upland Area to evaluate the influence of tidal variations in the level of surface water in the adjacent Guemes Channel on groundwater levels at the Site. Monitoring locations adjacent to the shoreline including MW-1 through MW-3 and at varying distances from the shoreline including MW-4 and MW-5 were selected to evaluate the lateral influence of tidal action on groundwater. The locations that were monitored as part of the tidal study are shown on Figure 13.

The tidal study recorded groundwater/potentiometric level response to tidal fluctuations using electronic water-level sensors consisting of a pressure transducer and automated datalogger installed in each monitoring location. Additionally, an electronic water-level sensor was attached to the East Dock (now removed) within the Marine Area to directly monitor and record the surface water level for comparison to water levels recorded in upland monitoring locations. The water-level sensors were removed from the monitoring locations and Marine Area after completion of the tidal study.

The data generated as part of the tidal study was analyzed using the Serfes (1987) method to identify the mean groundwater elevations and flow direction during the 72-hour tidal study and the Ferris (1951) method to evaluate hydraulic parameters of the shallow and deep groundwater units. Field procedures for the tidal study are presented in Appendix H. Results of the tidal study are summarized in Section 5.3.



3.5.3.5. Deviations from RI/FS Work Plan

The following deviations from the Ecology-approved RI/FS Work Plan occurred during the groundwater RI:

- During the February 2016 groundwater monitoring event, monitoring well MW-2A was observed to be damaged and as a result, no water samples were collected from this location for laboratory analysis. In August 2016, a replacement monitoring well (MW-2B) was installed for evaluating groundwater conditions.
- Prior to the 2012/2013 quarterly groundwater monitoring activities, the location of monitoring well MW-5 could not be identified. Ecology determined that a replacement well for MW-5 was not necessary and the existing monitoring well network was sufficient to evaluate groundwater conditions at the Site.

3.5.4. Soil Remedial Investigation

The soil RI was completed in general accordance with the Ecology-approved RI/FS Work Plan to characterize soil conditions in the Upland Area of the Site. The primary objective of the soil RI was to fill identified data gaps in the historical site characterization studies. Soil sample locations for the RI are shown on Figure 14. Field procedures including sample collection and handling, equipment decontamination and field screening are presented in Appendix H. The soil RI field activities are summarized below.

3.5.4.1. Soil Sample Collection

The soil RI included the collection of samples from the Site using a combination of hollow stem auger (HSA) drilling, direct push (DP) drilling, test pit (TP) and hand auger (HA) drilling technologies. In June 2008, subsurface soil samples were collected from eleven HSA explorations, ten test pit explorations and three hand auger explorations to meet the objectives of the RI/FS Work Plan.

As described in Section 3.5.3, the Port completed additional quarterly groundwater monitoring activities between May 2012 and February 2013 following completion of the 2008 Interim Action and reconfiguration of the shoreline during the Project Pier 1 redevelopment to evaluate whether previously identified contaminants in soil were adversely effecting groundwater. The results of the quarterly groundwater monitoring activities (further discussed in Sections 5.4 and 5.5) identified concentrations of arsenic, nickel and cPAHs exceeding groundwater PCULs during one or more quarterly monitoring events. In 2014, a supplemental soil investigation was completed to further characterize the nature and extent of these contaminants in soil. During this supplemental soil investigation, 43 DP explorations were completed to further evaluate subsurface soil conditions.

Three additional DP explorations were completed in July 2018 to evaluate soil conditions adjacent to and upgradient from monitoring well MW-8 based on the detected concentrations of arsenic and cPAHs in groundwater at this location.

Soil from each exploration was visually classified in general accordance with ASTM D-2488 and screened in the field for the presence of contamination. In addition, the presence of wood debris by type (i.e., saw dust, bark, chips, chunks, twigs, fibers, etc.) was also recorded when encountered. Field screening consisted of visual observation of contamination (i.e., staining, discoloration, etc.), water sheen testing, and organic vapor monitoring using a photo-ionization detector (PID). Exploration logs for soil investigation activities detailing observed soil conditions and field screening results are presented in Appendix H.



3.5.4.2. Laboratory Analyses

Samples collected as part of the soil RI were submitted to CCI located in Everett, Washington, and OnSite located in Redmond, Washington. Samples were submitted for analysis of one or more of the COPCs identified based on proximity to specific historical activities and previous sample results in accordance with the RI/FS Work Plan and discussions with Ecology following review of the initial 2008 and 2012/2013 groundwater monitoring data. Soil samples were submitted for a combination of the following analysis:

- Metals (arsenic, copper, nickel and zinc) by EPA Method 6000/7000 series;
- PAHs by SW-846 8270-SIM;
- Gasoline-range hydrocarbons by NWTPH-Gx;
- Diesel- and oil-range petroleum hydrocarbons by NWTPH-Dx with silica gel cleanup;
- VOCs including MTBE, EDB, and EDC by EPA Method 8260B and 8011; and
- Dioxins and furans using EPA Method 8290 or EPA Method 1613B.

3.5.4.3. Deviations from RI/FS Work Plan

The following deviations from the Work Plan were noted during the soil field investigation:

- Exploration locations for SB-1, SB-2, SB-4, SB-7 and SB-11 were adjusted in the field due to access restrictions resulting from shipyard operations (i.e., equipment storage and staged steel for vessel construction).
- Ten test pit explorations (TP-3 through TP-5 and TP-10 through TP-16) were completed in the east portion of the Site in October 2008. The purpose of the test pit explorations was to supplement the existing data in this area and to further evaluate the limits of arsenic, copper and zinc exceedances previously identified in this area. The test pits were completed during installation of subsurface infrastructure as part of the Project Pier 1 Redevelopment.

3.5.5. Evaluation of Site Outfalls and Catch Basins

During development of the RI/FS Work Plan, Ecology identified sediment in DCI catch basins as a potential source of contamination to the Marine Area sediments. However, the DCI stormwater system has been significantly modified since preparation of the RI/FS Work Plan. Over-time, the majority of the Site surfaces have been paved and a new stormwater collection system (described in detail in Section 1.4.6) was installed as part of the Port's Project Pier 1 redevelopment to capture the stormwater and wastewater from the Site for treatment prior to discharge. Stormwater and wastewater captured at the Site is treated prior to discharge to the City's sanitary sewer or to Guemes Channel under NPDES General Permit WAR045711. DCI performs regular monitoring of the water collection systems in accordance with DCI's SWPPP to ensure compliance with the discharge requirements of the NPDES permit.

Potential historical contaminant discharges to the Marine Area prior to the stormwater system upgrade activities have been addressed as part of the 2008 Interim Action (further discussed below), where the contaminated surficial and underlying clean native sediments were removed. The 2008 Interim Action dredging resulted in the removal of the known contaminated sediments from the Marine Area, therefore establishing a new baseline relative to potential contaminant sources.



As a result of the ongoing treatment and monitoring activities for stormwater and wastewater from the Site, discharges from the DCI stormwater/wastewater collection systems are no-longer a potential source of contamination to the Marine Area.

3.6. Environmental Data Used for the RI

Data sources for this RI report include data collected in general accordance with the Ecology-approved RI/FS Work Plan. Environmental data collected under the RI/FS Work Plan to fulfil the requirements of the Agreed Order were reviewed for technical quality. Based on this technical review, the data were determined to be of acceptable quality, as qualified and have been entered into Ecology's Environmental Information Management (EIM) System under Study ID FS2670 DCI. Laboratory data reports for sediment, groundwater and soil RI activities completed by GeoEngineers are presented in Appendix I. Validation reports for these data are presented in Appendix J.

Historical sediment, groundwater and soil data were also reviewed for technical quality. Environmental data in which sample locations, sample depth, analytical methods and chemical analytical results (as qualified) could be verified are considered acceptable for use to identify Site COPCs. Chemical analytical data used for this RI to identify COPCs and to evaluate the nature and extent of contaminates exceeding PCULs are described below.

3.6.1. Sediment

The sediment data used for this RI consists of samples obtained by GeoEngineers in March 2008 in general accordance with the RI/FS Work Plan to evaluate near shore sediment conditions as well as data collected by others to support dredge material suitability determination. These data were used to identify COPCs in the Marine Area and form the basis for Interim Action dredging completed in 2008 (further discussed in Section 4.0).

Confirmation sample results obtained following completion of the 2008 Interim Action as well as sediment sample results collected as part of the 2007 Fidalgo Bay Sediment Investigation represent current (post Interim Action) sediment conditions for the Marine Area and north adjacent Guemes Channel.

3.6.2. Groundwater

A network of monitoring wells installed in the Upland Area of the Site were used to evaluate hydrogeologic conditions and the nature and extent of contamination in groundwater. The network of groundwater wells used to evaluate groundwater conditions is comprised of eight monitoring wells (MW-1 through MW-8) screened in the shallow unconfined aquifer. Well completion logs for the groundwater monitoring well network are presented in Appendix E and H.

Groundwater data collected during previous environmental studies were used to identify COPCs². In accordance with the Ecology-approved RI/FS Work Plan and as requested by Ecology, groundwater data

² Groundwater data collected prior to the 2002 Petroleum and Marine Area Cleanup Actions are not considered representative of Site conditions given the extent of soil contamination removal completed for these areas (see Section 3.4), and therefore were not used to identify COPCs.



was obtained by GeoEngineers between June 2008 and August 2017 to evaluate the nature and extent of COPCs. These groundwater sampling results represent the data set for the RI.

3.6.3. Soil

Soil data collected during previous environmental studies were used to identify COPCs for the RI. In addition, Groundwater data obtained by GeoEngineers were used to support identification of soil COPCs (i.e., if a COPC was not previously identified in soil and the results of RI groundwater sampling confirmed that a COPC was not present, then it was not retained as a soil COPC for further evaluation). In accordance with the Ecology-approved RI/FS Work Plan and as requested by Ecology, soil data was obtained by GeoEngineers between June 2008 and July 2018 to evaluate the nature and extent of COPCs. These soil sampling results represent the data set for the RI.

4.0 INTERIM ACTION

An Interim Action was completed within the Marine Area as part of the Project Pier 1 redevelopment in 2008. The purpose of the Interim Action was to remove contaminated sediment from the Marine Area identified by the sediment RI and other sediment characterization studies. In addition, contaminated soil from the Upland Area of the Site where new underground utility infrastructure was installed was also removed. The extent of contaminated sediment for removal was based on the sample results from previous environmental studies and sediment investigation activities completed as part of the RI (Sections 3.1 and 3.5). Contaminated soil was removed from new subsurface utility corridors that were constructed as part of the Site redevelopment.

A detailed description of the Ecology-approved 2008 Interim Action is presented in the Interim Action Report (GeoEngineers 2010; Appendix K). Characterization activities and results for dredged material management and Upland Area soil, interim action dredging and nearshore excavation activities, confirmation sampling and restoration are summarized in the following sections.

4.1. Dredged Material Characterization

The Marine Area sediments were characterized for the purposes of dredged material disposal in addition to the RI sediment characterization. For the purposes of the dredged material characterization, the Marine Area was divided into two DMMUs (DMMU-D1 and DMMU-D2). DMMU-D1 encompassed sediment in the north and outer approximate half of the Marine Area and was located farther offshore from known sources while DMMU-D2 encompassed the sediment in the south and inner approximate half of the Marine Area, adjacent to the uplands and known historical sources of contamination (Figure 7).

Based on the results of the dredged material characterization, the DMMP determined that surface and subsurface sediment in DMMU-D1 was suitable for open-water disposal. In DMMU-D2, the DMMP determined that recent sediments (above the native glaciomarine layer) were contaminated and therefore, not suitable for open-water disposal. However, the underlying native glaciomarine sediments were suitable for open-water disposal. A copy of the DMMP open-water suitability determination is presented in Appendix D.



4.2. Marine and Upland Area Characterization Results

4.2.1. Marine Area Sediment Characterization

Marine Area sediment characterization (including the RI; Section 3.1 and 3.5) identified contaminant concentrations exceeding SMS cleanup standards including metals (arsenic, copper, lead, mercury and zinc), HPAHs, LPAHs, phenols and PCBs in recent fill sediment within DMMU-D2. At the time that the Interim Action was completed, PCULs for dioxins and furans, and cPAHs were not yet established by Ecology under the SMS. The existing data on the Marine Area sediments show however, that the detected exceedances of dioxins and furans, and cPAHs (relative to current SMS standards) were only present in the surficial fill sediment above the native glaciomarine deposits and that PCUL exceedances of these contaminants were not observed in subsurface sediment based on historical core sample results (i.e., AN-P1-2, AN-DCI-1A/B and AN-DCI-2) and Interim Action confirmation sample results (i.e., SMA-1 through SMA-5).

Sediment characterization activities completed as part of the 2008 RI showed that that the vertical extent of sediment contamination extended from the mudline to approximately 1-foot below the mudline surface along the eastern edge of the basin (sample location G-2), to approximately 4 feet below the mudline surface along the southwest corner of the Marine Area (sample locations G-5 and G-6) and to the native glaciomarine contact in the southeast corner of the basin (sample locations G-3 and G-4). In accordance with the DMMP suitability determination, contaminated sediment deposits within DMMU-D2 were removed as part of the 2008 Interim Action and transferred from the Site for upland disposal.

Historical and RI sediment sample results are summarized in Tables C-1 and C-2 (Appendix C) and shown relative to the Property on Figures 7 and 12.

4.2.2. Upland Area Soil Characterization

In the Upland Area, environmental studies (including the RI; Section 3.2 through 3.5) identified PCUL exceedances of metals (arsenic and nickel) and cPAHs in soil and/or groundwater (further discussed in Section 5.4 and 5.5). Arsenic results in soil samples collected during the 2008 RI were used to evaluate the Upland Area component of the Interim Action. Based on the analytical results of the 2008 RI, the native deposits underlying upland fill soil was identified as the vertical limit of the upland soil contamination. The interim action in the Upland Area included the excavation and disposal of fill soil excavated to facilitate the installation of new subsurface utility infrastructure as part of the Project Pier 1 redevelopment project.

Historical and RI soil sample results are summarized in Tables G-1 and G-2 (Appendix G) and shown relative to the Property on Figures 9 and 14.

4.3. Summary of the 2008 Interim Action Activities

Interim action dredging and excavation activities were completed between July and November 2008 in general accordance with the Ecology-approved RI/FS Work Plan and Interim Action Work Plan Addendum, and in general accordance with the MTCA Cleanup regulation and applicable state and federal laws described in WAC 173-340-430.

The Port's general contractor for interim action construction was Pacific Pile and Marine (PPM) of Seattle, Washington. The extent of contaminated sediment dredging and soil excavation during the interim action was defined based on the results of previous environmental studies as described above and field observations and chemical analyses of confirmation samples completed during construction. A



GeoEngineers field representative was onsite during dredging and excavation activities to field screen dredged and excavated materials for evidence of contamination and to assist the contractor in identifying the limits of contamination. In general, the native glaciomarine layer underlying the recent fill deposits was used as the lower limit of contamination for sediment dredging in the Marine Area and soil excavation in the Upland Area. Confirmation samples were collected from the post-dredge surface to confirm the completeness of the contaminated sediment removal action.

As a result of the interim action dredging, approximately 26,000 cubic yards (approximately 38,000 tons) of contaminated sediment was dredged from the Marine Area and transported by truck for disposal at the Waste Management's Subtitle D landfill facility in Wenatchee, Washington. Following removal of the contaminated material, dredging was completed to deeper elevations to meet the planned redevelopment navigation depths. Deeper dredging completed following the interim action included removal of an additional approximately 230,000 cubic yards of sediment from the Marine Area that was transported by barge to the Rosario Strait dispersive site to meet the design grade of -35 feet MLLW within the Marine Area. Prior to completion of the deeper dredging activities, confirmation samples (SMA-1 through SMA-5) were collected to verify the removal of sediment contamination. Confirmation sample locations are shown on Figure 12. Confirmation sample results documenting the removal of previously identified sediment contamination are summarized in Tables C-1 and C-2. Included in Tables C-1 and C-2 are sediment sample results collected during the 2007 Fidalgo Bay Sediment Characterization Study. Confirmation sample results and sediment sample results for the 2007 Fidalgo Bay Sediment Characterization Study represent the current sediment conditions for the Marine Area and surrounding area.

In the Upland Area, approximately 570 cubic yards (approximately 860 tons) of arsenic contaminated soil was excavated from the Upland Area and transported by truck for disposal at the Waste Management's Subtitle D landfill facility in Wenatchee, Washington. Excavation activities in the upland area ranged in depth from approximately 3 feet to 9.5 feet bgs to facilitate installation of the new subsurface utility corridors.

4.4. Backfill and Restoration

Project Pier 1 redevelopment involved expansion of the Upland Area northward by filling the part of the remediated Marine Area. To facilitate the filling, a permanent sheet pile wall (open cell bulkhead) was installed in the Marine Area (Figure 13) and the area behind the sheet pile wall was backfilled with imported material in accordance with the redevelopment project requirements to match the surrounding upland grade. Along the eastern slope of the Marine Area, up to 1-foot of habitat mix was imported and placed within SMA-1 (Figure 13) to restore the subtidal slopes to design grades following dredging.

The current Site layout including the location of new bulkhead and subsurface utility corridors is shown on Figure 4. The current Marine Area is maintained to an approximate navigational depth of -35 feet MLLW. The ground surface in the upland portion of the DCI shipyard has a grade of approximately +13 feet MLLW south of the open cell bulkhead.



5.0 REMEDIAL INVESTIGATION RESULTS

5.1. Sediment Stratigraphy

The stratigraphy of sediment in the Marine Area was characterized based on observations of materials encountered in explorations completed as part of the RI and as part the previous sediment investigations completed at the Site (Section 3.0). Prior to dredging in 2008, subsurface sediment conditions in the Marine Area generally consisted of approximately 3 to 7 feet of recent deposits consisting of loose silt and sand with occasional shell fragments overlying glaciomarine deposits consisting of very dense sand and hard silt. Near the outer harbor line, glaciomarine deposits were encountered at depths ranging between -35 and -40 feet MLLW. In the nearshore portion of the Marine Area, glaciomarine deposits were encountered at depths ranging between -10 and -15 MLLW.

During the 2008 Project Pier 1 redevelopment activities, sediment in the Marine Area north of the installed open cell bulkhead (Figure 5) was dredged to an approximate depth of -35 feet MLLW – well into the clean glaciomarine layer. South of the open cell bulkhead, Marine Area dredging was completed to remove the contaminated recent silt and sand deposits to the surface of the clean native glaciomarine layer. At the completion of the Project Pier 1 redevelopment, the area south of the open cell bulkhead was filled with clean imported material to the current grade of approximately 15 feet MLLW.

The stratigraphy of the Marine Area prior to dredging is shown in cross-section on Figure 15. The stratigraphy of the Marine Area following dredging is shown in cross-section on Figure 16. Information from the sediment cores from the RI and previous environmental studies were used to prepare these cross-sections.

5.2. Soil Stratigraphy

The stratigraphy of soil in the Upland Area was characterized based on observations of materials encountered in soil explorations completed as part of the RI and as part of the previous soil investigations completed at the Site (Section 3.0). The information from observations of the soil explorations was used to prepare cross-sections illustrating soil stratigraphy in the Upland Area. Cross-section locations in the Upland Area are shown on Figure 14. Cross-sections illustrating soil stratigraphy are presented on Figures 17 and 18.

Development of the Site has included filling of the nearshore part of the Marine Area to expand the Upland Area of the Site. Based on observations from the soil explorations, the stratigraphy at the Site generally consists of artificial fill soil deposits overlying native sand and silt. The stratigraphy of Upland Area generally includes the following:

Area northward of the historical shoreline after completion of the interim action dredging. To facilitate the infilling of this area, an open cell bulkhead was installed and the area behind the wall was backfilled to match the surrounding upland grade with clean imported material to meet the project design requirements. In addition, subsurface utilities (electric power, water, sewer, etc.) located within the upland interim action area (Figure 14) that served historical and existing facilities at the Site were decommissioned and replaced with new utility infrastructure to support DCI operations. The fill placed as part of Project Pier 1 ranges in thickness from approximately 3 to 20 feet with the thickest deposits located immediately south of the open cell bulkhead.



- Historical Fill Deposits: The historical fill deposits are comprised of layers of sand, silty sand and silt with variable gravel content ranging from approximately 2 to 16 feet thick that were placed during initial shoreline development in the 1960s to extend the historical shoreline northward. Contained in the historical fill deposits are occasional debris including concrete asphalt, brick and wood fragments. Historical fill deposits generally increase in thickness north of 3rd Street
- 1975 Earth Fill Area Deposits: The "1975 Fill Area" located in the southwestern portion of the Site was used for residential purposes from before 1925 until after approximately 1966 based on a review of historical Sanborn maps and aerial photographs. This area was topographically lower than the surrounding ground surface and was filled around 1975. In this area, fill deposits consist of layered silt, clay and silty sand deposits with occasional wood debris that are approximately 6 to 7 feet in thickness suggesting that some fill was placed prior to 1975.
- Native Deposits: Native material underlying the fill deposits at the Site include beach sands overlying glacial deposits. The beach sand deposits are typically poorly sorted and loose in nature and vary in thickness from 2 to 4 feet. Glacial deposits consist of a medium dense glaciomarine drift with varying amounts of silt, sand, and gravel that extend to all depths explored. A layer of dark brown organic deposits is present below the fill layer in central and southwestern portions of the Site. The organic layer varies in thickness from several inches to 2 feet.

5.3. Hydrogeology

Based on the results of the RI, groundwater occurs at the Site in a single aquifer comprised of two separate hydrostratigraphic units that control groundwater elevations, flow directions and the degree of influence from tidal fluctuations in the adjacent Guemes Channel. The two hydrostratigraphic units include a shallow unconfined aquifer and underlying confining unit. The shallow water-bearing unit at the Site is comprised of sand and gravel fill and native sand deposits (i.e., native beach sands). The confining unit at the Site is comprised of the glacially consolidated deposits.

Within the shallow groundwater unit, the groundwater elevation varies seasonally, with observed wet season elevations being higher than dry seasons by up to approximately 1 foot. Asphalt and concrete pavement at the Site inhibit the infiltration of precipitation across the majority of the Upland Area making the primary recharge mechanism for the shallow groundwater unit precipitation falling onto and infiltrating into soil south of the Site where it subsequently flows toward the Guemes Channel. Precipitation falling on the asphalt and concrete pavement is captured in catch basins and is treated prior to discharge to Guemes Channel. Precipitation falling on the limited areas that are gravel surfaced at the Site infiltrates into the ground and recharges to some degree, shallow groundwater within these areas.

5.3.1. Tidal Study Results

A 72-hour tidal study was completed using existing monitoring well locations at varying distances from the shoreline to evaluate the lateral influence of tidal action in the Guemes Channel on groundwater. A surface water location in the Marine Area was also monitored to directly record the surface water level for comparison to water levels recorded in the upland monitoring locations. Where present, tidal fluctuations in groundwater levels in individual monitoring locations were analyzed to identify the following:

The magnitude of the tidal influence on the groundwater level in the well relative to distance from the shoreline, which is identified as the stage ratio and presented as a percent (%);



- The length of time it took for the tidal effect observed at the shoreline to reach an individual monitoring well location, which is identified as the time lag and presented in hours; and
- The effect of tidal fluctuations on groundwater gradients.

Based on the results of the tidal study, shallow groundwater at the shoreline was determined to be tidally influenced. Tidal influence on groundwater at other monitoring locations located further upland of the shoreline and within the interior portion of the Upland Area was indeterminate based on the Serfes (1987) analysis method (Appendix H). The magnitude of tidal influence on shallow groundwater, as indicated by the stage ratio, was greatest at monitoring wells MW-2 (16 percent) and MW-3 (5.1 percent) located adjacent to the shoreline. Limited tidal influence (stage ratio of 3 percent or less) was observed at the other monitoring wells evaluated as part of this study. In general, the results of the tidal study indicate that there is limited communication between tidally influenced marine water and shallow groundwater at the Site except within approximately 150 feet of the shoreline.

The monitoring well locations used for the tidal study and the observed tidal effects are shown on Figure 19. A detailed description of the methodology used to perform the tidal study as well as the tidal study results are presented in Appendix H. Please note that the 72-hour tidal study was completed prior to installation of the open cell bulkhead as part the 2008 Interim Action for the Site. Currently, the open cell bulkhead which now separates the Upland Area from the Marine Area likely provides a physical barrier restricting the direct discharge of groundwater north to Guemes Channel.

5.3.2. Groundwater Gradients

Characterization of the groundwater gradient and flow direction in the Upland Area is based on the results of the tidal study by averaging the groundwater elevations measured over the 72-hour tidal study. The results of the tidal study indicate that shallow groundwater generally flows north toward the Marine Area and Guemes Channel. Using the results of the tidal study, shallow groundwater gradients were also estimated for the Site. In the central portion of Upland Area, the shallow groundwater gradient is estimated to be 0.0134 feet per feet (ft/ft) between monitoring wells MW-4 (upgradient well location) and MW-2 (shoreline well location).

Groundwater flow based on average groundwater elevation measured over the 72-hour tidal study are shown on Figure 19. A detailed description of the methodology used to estimate groundwater gradients is presented in Appendix H.

5.3.3. Hydraulic Conductivity

Hydraulic conductivity for the shallow and deep groundwater units were evaluated using information from slug tests performed on selected monitoring wells (Figure 19). The values resulting from slug tests are considered estimates of the hydraulic conductivity at the test location or in a localized portion of the aquifer where the test was performed due to the radius of influence generated during slug testing. The methodology and procedures for evaluating and calculating hydraulic conductivities based on slug tests as well as the slug test results are presented in Appendix H.

Slug tests were performed on five monitoring wells (MW-1 through MW-5). Hydraulic conductivity (K) values calculated from slug test data for the shallow groundwater unit ranged from approximately 1.43 ft/day to



14.93 feet per day (ft/day) with an average K value of 4.29 ft/day. These calculated K values were used to estimate the average linear groundwater velocity, as discussed in the following section.

The K values calculated based on the slug tests generally correspond to the material in which the wells are screened that is identified in the exploration logs presented in Appendix E and H. The wide range of K values calculated for the shallow groundwater unit reflect the heterogeneous nature of the hydrogeologic unit, which includes fine to coarse sand, gravel, silt and clay.

5.3.4. Groundwater Velocity

Calculated hydraulic conductivity values and groundwater gradients were used to estimate linear groundwater velocities for the Site. The average linear groundwater velocity between monitoring well pair MW-4 and MW-2 was 0.06 ft/day and the average linear groundwater velocity between monitoring well pair MW-4 and MW-3 was 0.07 ft/day with a northerly flow direction.

Groundwater monitoring wells used to calculate groundwater velocities are shown on Figure 19. A detailed description of the methodology used to evaluate groundwater conditions is presented in Appendix H.

5.4. Determination of Contaminants of Concern

COPCs were screened to 1) identify which contaminants were not detected and which contaminants were detected, but at concentrations less than the PCUL; and 2) identify frequency at which a contaminant exceeds the PCUL for identifying Site contaminants of concern (COCs). Tables 8 through 11 summarize the frequency at which analytes were detected and identifies which analytes were detected at a concentration exceeding the PCULs. The PCULs developed for the Site are summarized in Section 2.0.

Analytes that were detected at concentrations greater than the PCUL were considered as a COC if they met either of the following criteria:

- 1. The analyte had an exceedance frequency of at least 10 percent; or
- 2. The analyte had an exceedance factor of 2 or more.

The frequency at which a contaminant exceeds the PCUL is termed the "exceedance frequency". The magnitude by which a contaminant exceeds the PCUL is termed the "exceedance factor". The exceedance factor is derived by dividing the detected contaminant concentration by the concentration of the PCUL.

Evaluation of COCs in each medium was completed in a step wise fashion where sediments were evaluated, then groundwater followed by soil. If an analyte was not detected in sediment above the PCUL, then the groundwater to sediment exposure pathway was determined to be incomplete and that the corresponding analyte in groundwater not need to be protective of sediment. Groundwater PCULs in Table 3 for the protection of sediment were then adjusted based on this evaluation. Similarly, if an analyte was not detected in groundwater above the PCUL, then the soil to groundwater exposure pathway was determined to be incomplete and that the corresponding analyte in soil not need to be protective of groundwater. Soil PCULs in Table 4 for the protection of groundwater were then adjusted based on this evaluation.

The selection of COCs for each medium of concern are summarized in Tables 8 through 11. These tables present frequency of exceedance and exceedance factor summary statistics for the analytes detected in



each medium as well as a description of other considerations that were evaluated as part of the COC selection process. COCs identified for each medium are summarized in the following Sections.

5.4.1. Sediment Contaminants of Concern

Tables 8 and 9 present the COC evaluation for Marine Area sediment. Table 8 presents the COC evaluation for the protection of benthic organisms. Table 9 presents the COC evaluation for the protection of human health and higher trophic level ecological receptors. Identified sediment COCs include the following:

- **Metals** Based on the frequency of detection and/or exceedance factor, metals including arsenic, copper, lead, mercury and zinc are identified as COCs for Marine Area sediment.
- **Tributyltins** Based on the frequency of detection and/or exceedance factor, tributyltin is identified as a COC for Marine Area sediment.
- LPAHs Based on the frequency of detection and/or exceedance factor, LPAH compounds are identified as COCs for Marine Area sediment.
- **HPAHs** Based on the frequency of detection and/or exceedance factor, LPAH compounds are identified as COCs for Marine Area sediment.
- Total cPAH TEQ Based on the frequency of detection and/or exceedance factor, total cPAH TEQ is identified as COCs for Marine Area sediment.
- PCBs Based on the frequency of detection and/or exceedance factor, PCBs are identified as COCs for Marine Area sediment.
- **Dioxins and Furans** Based on the frequency of detection and/or exceedance factor, dioxin and furans are identified as COCs for Marine Area sediment. However, dioxin and furan exceedances were only identified in surface (0 to 10 cm) sediment. Results of subsurface investigation activities indicate that dioxins and furan exceedances to do not extend below the surface sediment in the Marine Area.

Other analytes were not selected as COCs because they were infrequently detected, had a low exceedance frequency and/or only slightly exceeded the PCUL.

5.4.2. Groundwater Contaminants of Concern

Table 10 presents the COC evaluation for Upland Area groundwater. Identified groundwater COCs include the following:

- Metals Based on the frequency of detection and/or exceedance factor, metals including arsenic and nickel are identified as COCs for Upland Area groundwater.
- Total cPAH TEQ Based on the frequency of detection and/or exceedance factor, total cPAH TEQ is identified as COCs for Upland Area groundwater.

Other analytes were not selected as COCs because they were infrequently detected, had a low exceedance frequency and/or only slightly exceeded the PCUL.

5.4.3. Soil Contaminants of Concern

Table 11 presents the COC evaluation for Upland Area soil. Identified soil COCs include the following:



- Metals Based on the frequency of detection and/or exceedance factor, metals including arsenic and nickel are identified as COCs for Upland Area soil.
- Total cPAH TEQ Based on the frequency of detection and/or exceedance factor, total cPAH TEQ is identified as COCs for Upland Area soil.
- Petroleum Hydrocarbons Although verification sampling data confirmed the removal of petroleum contaminated soil from the 1991 UST Cleanup Action, 2001 Hydraulic Winch Cleanup Action and 2002 Petroleum and Marine Railway Cleanup Action areas, the technical quality of these data could not be independently validated. Therefore, gasoline-, diesel- and heavy oil-range petroleum hydrocarbons are retained as a COC for Upland Area soil.

Other analytes were not selected as COCs because they were infrequently detected, had a low exceedance frequency and/or only slightly exceeded the PCUL.

5.5. Nature and Extent of Contamination

The nature and extent of COCs in Site sediment, groundwater and soil are summarized in the following sections. As discussed in Section 3.5.5, catch basin solids are not considered a media of concern for further evaluation because the current stormwater collection system for DCI (described in detail in Section 1.4.6) installed as part of the Port's Project Pier 1 redevelopment captures stormwater and wastewater generated from the Site for treatment prior to discharge. Potential contaminant discharges from stormwater discharges prior to the Project Pier 1 redevelopment have been addressed as part of the 2008 interim action dredging in the Marine Area.

5.5.1. Sediment Contamination

Dredging within the Marine Area was completed as part of the 2008 Interim Action at the Site and is described in Section 4. The interim action dredging removed approximately 26,000 cubic yards (approximately 38,000 tons) of contaminated sediment, processed and transported by truck for disposal at the Waste Management's Subtitle D landfill facility in Wenatchee, Washington. Following removal of the contaminated near surface sediments, further dredging was completed into the clean native sediments to meet a design grade of -35 feet MLLW.

Confirmation sediment samples collected as part of the 2008 Interim Action dredging activities and sediment samples collected during previous environmental studies representing current conditions within the Marine Area are presented in Tables 12 and 13. Sample locations are shown relative to the Marine Area on Figure 13.

Based on the sediment sample results representative of the post-dredge condition, COC at concentrations exceeding PCULs were removed from the Marine Area. Due to the completeness of the interim action dredging and subsequent dredging of up to 30 additional feet of underlying native sediments, no sediment contamination is known to be present and sediment following the interim action is no longer considered a media of concern.

5.5.2. Groundwater Contamination

The groundwater data set for this RI consists of groundwater samples obtained by GeoEngineers between June 2008 and August 2017 from new and existing Upland Area monitoring wells. Groundwater results for



identified COCs (described above) are presented in Table 14. Groundwater monitoring locations are shown on Figures 20 through 22. Based on a review of the chemical analytical data, the nature and extent of groundwater contamination includes the following:

- MW-1 (Upland Well) Total and dissolved arsenic were detected at a concentration greater than the groundwater PCUL in monitoring well MW-1 which is located in the western portion of the Site during one or more monitoring events. However, dissolved arsenic concentrations at this location did not exceed the groundwater PCUL during semi-annual groundwater monitoring activities completed between February 2016 and August 2017.
- MW-2/2A/2B (Shoreline Well) Dissolved nickel was detected at a concentration greater than the groundwater PCUL in monitoring well MW-2A/2B which is located in the north central portion of the Site during one or more monitoring events. However, dissolved nickel concentrations at this location did not exceed the groundwater PCUL during semi-annual groundwater monitoring activities completed between February 2016 and August 2017 except for in February 2017 in which dissolved nickel detected at a concentration of 8.3 μg/L marginally exceeded the groundwater PCUL of 8.2 μg/L. In addition, the cPAH TEQ concentration at this location slightly exceeded the groundwater PCUL during the June 2008 monitoring event. In subsequent monitoring events, cPAHs were not detected in this monitoring well.
- MW-3/3A (Shoreline Well) Dissolved nickel was detected at a concentration greater than the groundwater PCUL in monitoring well MW-3A which is located in the north central portion of the Site during one or more monitoring events. However, dissolved nickel concentrations at this location did not exceed the groundwater PCUL during semi-annual groundwater monitoring activities completed between February 2016 and August 2017.
- MW-4 (Upland Well) Total arsenic and cPAHs were detected at concentrations greater than the groundwater PCUL in monitoring well MW-4 which is located in the south-central portion of the Site. However, total arsenic and cPAH TEQ concentrations at this location did not exceed the groundwater PCUL during semi-annual groundwater monitoring activities completed between February 2016 and August 2017.
- MW-5 (Upland Well) Total arsenic was detected at a concentration greater that the groundwater PCUL in monitoring well MW-5 which is located in the central portion of the Site. However, dissolved arsenic at this location and down gradient of this location (i.e., MW-2) was detected at a concentration less than the groundwater PCUL.
- MW-6 (Shoreline Well) Dissolved nickel was detected at a concentration greater than the groundwater PCUL in monitoring well MW-6 which is located in the north central portion of the Site during one or more monitoring events. However, dissolved nickel concentrations at this location did not exceed the groundwater PCUL during semi-annual groundwater monitoring activities completed between February 2016 and August 2017.
- MW-7 (Upland Well) Total and dissolved arsenic and nickel and cPAH TEQ concentrations exceeded groundwater PCLs in monitoring well MW-7 located in the southeastern portion of the Site during one or more monitoring events. Downgradient of MW-7, total/dissolved arsenic and nickel, and cPAHs did not exceed groundwater PCULs in MW-6 during semi-annual groundwater monitoring activities completed between February 2016 and August 2017.
- MW-8 (Shoreline Well) Total and dissolved arsenic and nickel, and cPAH TEQ concentrations exceeded groundwater PCULs in monitoring well MW-8 located in the northwestern portion of the Site



during one or more monitoring events. The concentration of dissolved nickel did not exceed PCUL levels in MW-8 during each of the semi-annual monitoring events. Although dissolved arsenic and cPAHs exceeded the groundwater PCUL, the concentration of dissolved arsenic appears to be stable and the concentration of cPAHs exceeding the PCUL was only observed during the August 2017 groundwater monitoring event with no detected concentrations of cPAHs during previous events.

Between 2015 and 2016, DCI replaced a significant portion of their gravel working surface with asphalt pavement that prevents stormwater infiltration through the soil column. A comparison of the initial (2008 to 2013) groundwater monitoring results to the recent semi-annual groundwater monitoring results (2016 to 2017) suggest that the paving activities have significantly reduced contaminant exceedances in groundwater and that in shoreline monitoring wells, exceedances of the groundwater PCULs are generally not observed. Trend plots for COCs including arsenic, nickel and cPAHs which were detected in monitoring wells MW-1 through MW-8 at concentrations exceeding groundwater PCULs during one or more monitoring event are shown on Figures 20 through 22.

5.5.3. Soil Contamination

The soil data set for this RI consists of soil samples obtained by GeoEngineers between June 2008 and August 2018 from subsurface explorations. In addition, soil sample results from previous environmental studies in which PCUL exceedances were observed are also being used to support the delineation of COC exceedances. Soil sample results for identified COCs (described above) are presented in Table 15. Soil sample locations and the distribution of soil COCs including arsenic, nickel, cPAHs and petroleum hydrocarbons are shown on Figures 23 through 26. Based on a review of the chemical analytical data, the nature and extend of identified soil contamination at the Site includes the following:

- Metals Arsenic and/or nickel was detected at concentrations exceeding PCULs in soil throughout the Site as follows:
 - In the eastern portion of the Site, arsenic and nickel exceeded PCULs in historical fill deposits from the ground surface up to a depth of approximately 8 feet bgs.
 - In the north central portion of the Site, arsenic and nickel exceeded PCULs in historical fill deposits from the ground surface up to a depth of approximately 10 feet bgs.
 - In the south-central portion of the Site, arsenic exceeded the PCUL in historical fill deposits from between approximately 5 and 8 feet bgs.
 - In the western portion of the Site, nickel exceeded the PCUL in the 1975 Earth fill area deposits from the ground surface up to a depth of approximately 10 feet bgs.

Results of soil/sediment samples collected at the Site from the underlying native surface suggest that the observed arsenic and nickel exceedance are contained within the overlying fill soil. Although soils at the Site contain concentrations of arsenic and nickel greater than the PCUL, groundwater monitoring data (summarized above) show that concentrations exist but are not exacerbating the current groundwater contamination at the Site. Decreasing concentrations due to paving indicates hydraulic connectivity and adverse effects from soil, but the modifications appeared to minimize contribution.

- Total cPAH TEQ Concentrations of cPAHs were detected at concentrations exceeding PCULs in soil as follows:
 - In the south and central portions of the Site, total cPAH TEQ concentrations exceeded the PCUL in historical fill deposits between approximately 5 and 13 feet bgs.



• In the western portion of the Site, total cPAH TEQ concentrations exceeded the PCUL in the 1975 Earth fill area deposits from the ground surface to a depth of approximately 8 feet bgs.

Results of soil/sediment samples collected at the Site from the underlying native surface suggest that the observed cPAHs exceedance are contained within the overlying fill soil. Although soil at the Site contain total cPAH TEQ concentrations greater than the PCUL, groundwater monitoring data (summarized above) indicate that the observed contamination is not adversely affecting groundwater at the Site and that the contaminant concentration in ground is stable or is decreasing over-time.

- Petroleum Hydrocarbons Results of previous environmental studies identified concentrations of gasoline-, diesel- and heavy oil-range petroleum hydrocarbons exceeding PCULs in historical fill deposits from the ground surface to a depth of approximately 8 feet bgs within the footprint of the 2002 Petroleum and Marine Railway Cleanup Action Areas. The nature and extent of petroleum hydrocarbons is as follows:
 - As previously discussed, the Port completed independent cleanup actions (i.e., 1991 UST, 2001 Hydraulic Winch and 2002 Petroleum and Marine Railway Cleanup Actions) to remove the previously identified petroleum contamination from the Site. Confirmation sample results obtained from the limits of these excavations indicated that the petroleum contamination was successfully removed from the Site. However, confirmation soil sample results for these areas could not be independently validated. Therefore, the absence or presence of petroleum hydrocarbons within the footprints of the previously completed cleanup action areas are not confirmed until subsequent sampling result confirm their removal³. In groundwater, petroleum hydrocarbons and petroleum-related constituents (i.e., BETX, EDB, EDC and MTBE) either were not detected or were detected at concentrations less than the PCULs within and downgradient of the previously completed cleanup action areas during each of the RI monitoring events providing further evidence for the completeness of the removal actions.
 - Petroleum hydrocarbons exceeding the PCULs were also identified in surficial soil (0-4 feet) in the western portion of the Site (i.e., Earth Fill Area). However, because petroleum hydrocarbons are not in contact with groundwater and groundwater monitoring results downgradient of the Earth Fill Area did not identify PCUL exceedances, PCULs for petroleum hydrocarbons in this area were adjusted for protection of direct contact only using Ecology's worksheet calculating soil cleanup levels for petroleum contaminated sites (MTCA TPH 11.1 Excel Workbook) in accordance with WAC 173-340-745. The maximum detected concentrations of petroleum hydrocarbons for this area were used as input parameters in this worksheet to calculate adjusted PCULs for gasoline- and heavy oil-range hydrocarbons for protection of direct contact. For this calculation, all petroleum hydrocarbons were assumed to be present in the most toxic form for each hydrocarbon range (i.e., gasoline-range hydrocarbons were assumed to be aliphatics with a 10- to 12-carbon chain and diesel range-hydrocarbons were assumed to be aliphatics with a 12- to 16-carbon chain). Adjusted PCUL calculations for the Earth Fill Area are presented in Appendix L. The resulting adjusted PCULs for this area were not exceeded, therefore do not require further cleanup evaluation.

³ To verify the completeness of the previously completed cleanup actions, additional soil investigation activities is being proposed within the footprints of the previously completed remedial excavations. Previsions for additional soil investigation activities to verify the completeness of the previous cleanup actions will be presented in the CAP to support cleanup action alternative refinement to address Site contamination.



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6.0 CONCEPTUAL SITE MODEL

For the RI, a conceptual site model (CSM) for potential contaminant sources, release mechanisms, transport processes, and exposure routes by which receptors may be affected was developed. The CSM incorporating the results of RI is shown on Figure 27 and is discussed below.

6.1. Media of Concern

6.1.1. Marine Area

Results of the RI identified COCs including metals, tributyltins, LPAHs, HPAHs, cPAHs, PCBs and dioxins and furans in the Marine Area. However, the interim action completed in the Marine Area followed by additional dredging resulted in the complete removal of known sediment contamination from the Site. As a result, sediment is not identified as a media of concern at the Site.

6.1.2. Upland Area

In the Upland Area, COCs including metals (arsenic and nickel) and cPAHs were identified in soil and groundwater at concentrations exceeding PCULs. Based on the results of the RI, Upland Area groundwater and soil are media of concern for the Site.

6.1.3. Stormwater Conveyance

As part of the Project Pier 1 redevelopment project, new systems were installed to capture and treat stormwater and wastewater at the Site prior to discharge. Because historical stormwater impacts (if any) to the Marine Area were removed by the interim action dredging and new infrastructure is in place to collect and treat stormwater prior to discharge, stormwater solids, are not identified as a media of concern for the Site.

6.2. Sources of Contamination to Media of Concern

The following have been identified as potential contaminant sources to the Site:

- Historical spills and releases onto Site soils;
- Placement of contaminated fill material; and
- Atmospheric deposition.

These potential contaminant sources are further described below.

6.2.1. Historical Spills and Releases onto Site Soils

The shoreline at the Site was historically located southwest of the former railway spur (Figure 4) based on a review of Sanborn maps and aerial photographs. Historical commercial and industrial activities at the Site since the late 1800s, including bulk fuel storage, shipping, shipbuilding, ship repairs and other maritime-related operations may have caused releases of contamination to the native and fill soils. The source of PAH contamination at the Site may include historical combustion of fossil fuels from residents, machinery (boiler, power, etc.), vehicles and marine vessels operating and wood burning. Metals contamination is likely the result of direct discharge of industrial wastes such as paint chips, grinding and blast grit residues to the ground from vessel maintenance and repair activities at the Site. These historical operations represent a potential source of contamination to the identified media of concern.



As previously discussed, independent cleanup actions of the Upland Area completed in 2002 resulted in the removal of identified petroleum contamination sourcing from bulk fuel storage and distribution operations. The results of confirmation sampling competed as part of these cleanup activities verified the removal of petroleum contaminated soil from the Site.

6.2.2. Placement of Contaminated Fill Material

Infilling of the historical Site shoreline has resulted in the placement of up to approximately 16 feet of fill material prior to the 1960s to expand the shoreline northward in support of maritime operations. These historical fill deposits are comprised of layers of sand, silty sand and silt with variable gravel content and contain debris including concrete asphalt, brick and wood fragments suggesting that they may have been re-used from other industrial sources and have the potential to contain contaminants of concern. Releases from historical commercial and industrial activities at the Site and surrounding area during infilling of the historical shoreline as well as contaminants contained within the fill material placed represent potential contaminant sources to the Site.

6.2.3. Atmospheric Deposition

Gases and particulates may be released to the atmosphere from current and historical combustion (vehicle emissions, burning, etc.) and/or industrial operations and may contain concentrations of metals and PAHs. Contaminants present in the atmosphere may be deposited by settling of particulates or precipitation directly to the land surfaces at the Site and surrounding area.

As described in previous sections, Project Pier 1 redevelopment in 2008 resulted in the installation of new infrastructure to collect and treat stormwater/wastewater generated at the Site prior to discharge. Since this time, DCI has completed facility upgrades that include paving much of the Upland area with asphalt that limits the infiltration of precipitation falling to the ground surface. Stormwater runoff is subsequently captured by a network of storm drains and is treated prior to discharge. However, atmospheric deposition prior to redevelopment and paving of the Site as well as atmospheric deposition onto currently unpaved areas activities represent a potential source of contamination to the Site.

6.3. Fate and Transport

The fate and transport of contaminants are affected by their chemical properties and the physical, chemical, and biological processes which they are exposed to at the Site. These properties and how they impact the fate and transport of the Site contaminants are discussed in the following sections.

6.3.1. Environmental Fate

6.3.1.1. Arsenic

Arsenic is a naturally occurring and persistent mineral that is not naturally degraded. Arsenic exists in the environment in four oxidation states that include +3 (arsenite), +5 (arsenate), 0 (arsenic metal) and -3 (arsine). The two most common oxidation states are +3 and +5. The mobility of arsenic in the environment depends strongly on the pH and redox conditions and can change oxidation states in response to environmental redox conditions. However, the change is generally slow. In the environment, arsenic bonds with a wide variety of other elements, producing inorganic and organic compounds with a range of mobilities.



Arsenic is capable of dissolution in pH ranges from 2.0 to 11.0 under suitable physical and chemical conditions. It is most soluble at both high (basic) pH and low (acidic) pH conditions, and less soluble at a neutral pH. Arsenic is also mobile under both oxidizing and reducing conditions. Arsenic transport could be characterized as cycling between sorbed and aqueous phases due to environmental changes that affect pH, redox, concentrations of other chemicals present in the water, and biologic activity (Panagiotaras and Nikolopoulos 2015).

6.3.1.2. Nickel

Nickel is a naturally occurring and persistent mineral present in soil is not naturally degraded. Nickel exists in several oxidation states, of which +2 is the most common. Nickel is generally, strongly adsorbed by soil. There are many factors that affect the extent to which nickel is adsorbed by soil, so the adsorption of nickel by soil is site specific. Amorphous oxides of iron and manganese, and to a lesser extent clay minerals are the most important adsorbents in soil (ASTDR 2017).

The solubility of nickel increases as pH decreases, causing it to become more mobile. Most nickel compounds are relatively soluble at pH values less than 6.5 and is relatively insoluble at high pH conditions.

6.3.1.3. PAHs

Transport and partitioning of PAHs in the environment are determined to a large extent by physicochemical properties such as water solubility, vapor pressure, Henry's law constant, octanol-water partition coefficient (Kow), and organic carbon partition coefficient (Koc). In general, PAHs have low water solubilities. Because of their low solubility and high affinity for organic carbon, PAHs in aquatic systems are primarily found sorbed to particles that either have settled to the bottom or are suspended in the water column. Volatilization and adsorption to suspended sediments with subsequent deposition are the primary removal processes for medium and high molecular weight PAHs, whereas volatilization and biodegradation are the major removal processes for low molecular weight compounds.

6.3.1.4. Petroleum Hydrocarbons

Gasoline is a mixture of relatively volatile hydrocarbons, including normal and branched chain alkanes, cycloalkanes, alkenes, and aromatics, that vary widely in their physical and chemical properties. Upon release to the environment, gasoline is not transported as a mixture; rather, the various components of the mixture selectively partition to the atmosphere, soil, or water according to their individual physical/chemical properties. Gasoline released to surface soils will differentially partition by volatilization, dissolution, or adsorption of individual constituents according to their physical/chemical properties. Gasoline exists in soil in four states: 1) as a free-moving liquid, 2) adsorbed to soil particulates, 3) dissolved in groundwater, and 4) as a vapor. Components of gasoline that are not volatilized or sorbed to non-colloidal soil particulates will migrate downward through the unsaturated zone by gravity or leaching to the water table. Liquid gasoline, as a result of its lower kinematic viscosity, is expected to move through the unsaturated zone of the soil at a velocity 2-3 times that of water. The amount of liquid product that reaches the water table is dependent upon the amount of product released and site-specific soil and hydrogeological conditions.

The transport and dispersion of diesel and heavy oil petroleum hydrocarbons are dependent on the water solubility and volatility of the aliphatic and aromatic hydrocarbon fractions. Lower molecular weight hydrocarbons such as n-alkanes may volatilize relatively quickly from both water and soil, while larger aliphatics (greater than C₉ chain length) may be sorbed to organic particles in water or soil. Aromatic hydrocarbons will be dissolved in the aqueous phase in both soil and water and may undergo some volatilization. Liquid diesel and heavy oils, as a result of their higher kinematic viscosity, are expected to



move through the unsaturated zone of the soil at a slower velocity as compared to gasoline. Similar to gasoline, the amount of liquid product that reaches the water table is dependent upon the amount of product released and site-specific soil and hydrogeological conditions.

Petroleum hydrocarbons immobilized in the unsaturated zone may be solubilized by downward moving soil water or fluctuating elevations of groundwater, and this residual material may serve as a source of contamination through leaching of solubilized components for long periods of time. Water-soluble components will dissolve in groundwater, whereas insoluble components will float as a separate phase on top of the water table. Water-soluble compounds, such as benzene, toluene, and xylene, show a greater potential for transport in groundwater aquifer than insoluble forms.

Overall, mass reduction of petroleum hydrocarbons occurs naturally through biodegradation. Biodegradation of petroleum is most efficient under aerobic conditions. Biodegradation of petroleum also results in reducing conditions as the available oxygen in the substrate is consumed during aerobic biodegradation. Anaerobic biodegradation of petroleum also occurs, but it is a slower process than aerobic biodegradation.

6.3.2. Environmental Transport

Release and transport mechanisms for Site contaminants to media of concern are presented on Figure 27. The specific release and transport mechanisms by which Site contaminants have been transported at the Site include:

- Infiltration and Leaching Infiltration of precipitation and leaching of contaminants in unpaved portions of the Site can transport COCs contained within the soil column. COCs that are soluble may enter solution and remain in solution until reaching groundwater or may be sorbed to soil in other portions of the soil column.
 - Since 2015, DCI has paved a significant portion of the Upland Area. Groundwater sampling results prior to this paving of the Site (i.e., 2008 to 2013 monitoring events) identified detected COC concentrations exceeding PCULs. However, groundwater monitoring completed since paving of the majority of the Site (i.e., 2016 to 2018 monitoring events) show a decrease in detected COC concentrations over-time. These findings indicate that infiltration and leaching of COCs from soil to groundwater is significantly reduced by the pavement and stormwater management practices at the Site.
- Shallow Unconfined Aquifer Solute Transport Solute transport in the shallow unconfined aquifer is governed by the environmental chemistry of the COCs described above and aquifer properties including hydraulic conductivity, groundwater velocity, gradient and tidal influence. As described in Section 5.3, the average shallow groundwater flow direction was measured to be to the north toward Guemes Channel with an estimated velocity of approximately 0.06 feet per day.
- Tidal Mixing Based on the results of the tidal study, there is a tidal mixing zone within approximately 150 feet of the shoreline. The results of the tidal study show that groundwater in shoreline wells are being influenced by fluctuations in sea level. However, monitoring wells located in the central and southern portions of the Site do not show a response to fluctuations in sea level and are therefore, less likely to be influenced by tidal mixing.



6.4. Exposure Pathways and Receptors

COCs released as the result of historic commercial and industrial operations at the Site have resulted in direct impacts to soil and secondary impacts to groundwater. Potential exposure pathways related to these media are discussed below.

6.4.1. Soil

The following potential exposure pathways and receptors existed for contaminants in Site soil:

- Contact (dermal, incidental ingestion or inhalation) by Site workers (including workers excavating soil);
 and
- Leaching of contaminants contained within the soil column to groundwater.

Visitors and terrestrial wildlife are not potential receptors of concern because access is limited to authorized personnel performing work at the Site (Section 1.1), and because pavement and gravel working surfaces limit the area of terrestrial habitat.

6.4.2. Groundwater

The following are potential exposure pathways and receptors for contaminants in Site groundwater:

- Contact (dermal or incidental ingestion) by Site workers (including workers excavating soil below the water table);
- Contact (dermal or incidental ingestion) by aquatic receptors to impacted groundwater that may discharge to the Marine Area resulting in acute or chronic effects; and
- Ingestion of aquatic organisms affected by the discharge of impacted groundwater to the Marine Area.

Human ingestion of contaminated groundwater from the Site is not a potential exposure pathway because groundwater at the Site is not a current or reasonable future source of drinking water (Section 1.4)

7.0 BASIS FOR CLEANUP ACTION

7.1. Cleanup Action Objectives

Cleanup action objectives (CAOs) are established to eliminate, reduce, or otherwise control to the maximum extent feasible and practicable, unacceptable risks to human health and the environment posed by Site-related hazardous substances in Upland Area soil and groundwater in accordance with the MTCA Cleanup Regulation (WAC 173-340) and other applicable regulatory requirements. CAOs form the basis for evaluating and selecting remedial technologies and cleanup actions that will be successful. CAOs consist of location-, chemical- and medium-specific goals for protecting human health and the environment. CAOs are dependent on the chemicals and pathways that represent a risk to people and natural resources associated with a site. Development of CAOs involves 1) identification of potentially applicable or relevant and appropriate requirements (ARARs) that set the framework and requirements for the development of cleanup standards and implementation of a cleanup action; 2) development of cleanup levels and points of compliance at which an acceptable risk level can be attained; and 3) identification of the locations and media requiring cleanup based on selected cleanup standards.



The following CAOs for the Site have been developed to mitigate risks associated with the Site COCs discussed in Section 5.5 and to address potential exposure routes and receptors discussed in Section 6.4 based on known subsurface conditions, and current and future land use:

- Contact (dermal, incidental ingestion or inhalation) by Site workers (including workers excavating soil below the water table);
- Leaching of contaminants contained within the soil column to groundwater;
- Contact (dermal or incidental ingestion) by aquatic receptors to impacted groundwater that may discharge to the Marine Area resulting in acute or chronic effects; and
- Ingestion of aquatic organisms affected by the discharge of impacted groundwater to the Marine Area by higher trophic level ecological receptors.

PCULs and points of compliance for sediment, groundwater and soil considered for the development of the CAOs are discussed in Section 2.0 and are expected to be adopted as final cleanup levels by Ecology for the Cleanup Action Plan (CAP). Areas and media requiring cleanup considered for the development of the CAOs are discussed in Section 5.5. ARARs considered for the development of the CAOs are further discussed below.

7.2. Potentially Applicable or Relevant and Appropriate Requirements

Under Washington Administrative Code (WAC) 173-340-710, MTCA requires that cleanup actions comply with all legally applicable local, state and federal laws, and requirements that are legally applicable and determined by Ecology to be relevant and appropriate requirements for the cleanup site. Legally "applicable" requirements under MTCA are those cleanup standards, standards of control, and other human health and environmental protection requirements, criteria, or limitations adopted under state or federal law that specifically address a hazardous substance, cleanup action, location, or other circumstance at a site (WAC 173-340-200). "Relevant and appropriate" requirements include those cleanup standards, standards of control, and other human health and environmental requirements, criteria, or limitations established under state or federal law that, while not legally applicable to the hazardous substance, cleanup action, location, or other circumstance at a site, address problems or situations sufficiently similar to those encountered at the site that their use is well suited to the particular site (WAC 173-340-200).

Potential ARARs and their descriptions/applicability are presented in Table 16. In accordance with WAC 173-340-710(9)(b), cleanup actions conducted by Ecology under MTCA are exempt from most procedural requirements of state and local laws, and related permitting requirements. Although exempt from procedural requirements of certain state and local laws and related permitting requirements, pertinent substantive compliance requirements remain applicable.

In addition, the Fidalgo Bay region is known to be archaeologically sensitive. Therefore, previsions of the National Historic Preservation Act of 1966, and the Archeological and Historical Preservation Act (16 USCA 469) are to remain applicable. The Samish Indian Nation, Swinomish Tribal Community, and other interested tribes, and the Washington State Department of Archaeology and Historic Preservation (DAHP) will be consulted on potential cultural resource and archaeological matters as they relate to the cleanup action. Ecology will take lead and carry out consultation in accordance with Executive Order 05-05 on locations not covered under Section 106.



7.3. Supplemental Soil Investigation

As previously discussed, environmental data collected during previous studies to confirm the completeness of the remedial actions could not be independently validated. As a result, a supplemental soil investigation is proposed as a pre-remedial design activity for the Site to verify the completeness of the 1991 UST removal excavation, 2001 marine railway winch remedial excavation, 2002 Petroleum and Marine Railway Cleanup Action areas. In general, borings will be advanced within the footprint of the previously completed remedial excavations which extended to depths greater than 1-foot (i.e., remedial excavations completed to remove previously identified soil contaminated located beneath the former gravel working surface). The borings will be advanced to a depth of below the base of the previous remedial excavations to collect soil samples that are representative of the previous verification sample interval. Samples will be analyzed for COCs previously identified for these areas including gasoline-, diesel- and heavy oil-range petroleum hydrocarbons and/or cPAHs.

The results of this supplemental soil investigation will be used to confirm the completeness of the previous removal actions. Sampling and analysis will be completed under an Ecology-approved addendum to the RI/FS Work Plan. The results of this investigation will be reported in the CAP and used to refine the preferred cleanup action alternative for the Site. Proposed sampling locations are shown relative to the previous removal action areas on Figure 28.

8.0 CLEANUP ACTION ALTERNATIVE DEVELOPMENT

8.1. Identification and Screening of Remedial Technologies

Potentially applicable remedial technologies for identified COCs in Upland Area media of concern (i.e., soil and groundwater) were screened and evaluated for developing cleanup action alternatives in accordance with MTCA requirements (WAC 173-340-350). Sources of information used to develop the list of remedial technologies include EPA publications and databases, vendor information, and professional experience gained at similar sites.

Under MTCA, remedial alternatives are developed from remedial technologies that are screened and identified as capable of meeting cleanup requirements to achieve the CAOs. Initial screening of remedial technologies allows development of a range of tools that can be used individually or in combination to address contamination at the Site. The screening process determines the most appropriate technologies and process options for addressing COCs in soil and groundwater based on their expected implementability, reliability, and relative cost as follows:

- Implementability This evaluation encompasses both technical and administrative feasibility of implementing a technology. Aspects of implementability include the ability to obtain permits, the availability of treatment methods, physical conditions of the site, and availability of required equipment and skilled workers.
- Effectiveness This evaluation focuses on 1) the potential effectiveness of a technology in handling the estimated areas or volumes of media and meeting CAOs; 2) the potential impacts to human health and the environment during the construction and implementation phase; and 3) how proven and reliable a technology is with respect to the contaminants and conditions at the site.



Cost - This evaluation takes into consideration relative capital, and operation and maintenance (0&M) cost rather than detailed estimates. During the screening process, the relative capital and 0&M cost between alternatives (based on engineering judgement) is evaluated as to whether costs are high, low, or moderate relative to the other technologies. Since remedial alternatives and associated quantities are not defined during technology screening stage, relative cost is presented qualitatively as a range rather than quantitatively.

Remedial technologies to address COCs in soil and groundwater in the Upland Area portion of the Site are discussed in the following sections. In general, remedial technologies that had limited implementability, low effectiveness, and/or high relative cost were screened out and the most appropriate technologies were retained for use in the development of remedial alternatives. Technologies retained through the screening process were selected as is or combined into remedial alternatives, as appropriate, for a detailed alternative evaluation.

8.1.1. Remedial Technologies for Soil

Descriptions and screening of applicable remedial technologies for soil are presented in Table 17. Based on the results of screening, the following are the remedial technologies for soil which were retained for development of remedial alternatives:

- Institutional controls including environmental covenants, land use restrictions and fencing and signage;
- Containment and capping including low permeability caps comprised of asphalt or concrete pavement with drainage controls;
- In situ treatment including stabilization; and
- Removal including excavation and offsite disposal to a permitted landfill.

8.1.2. Remedial Technologies for Groundwater

Descriptions and screening of applicable remedial technologies for groundwater are presented in Table 18. Based on the results of screening, the following are the remedial technologies for groundwater that are retained for development of remedial alternatives:

- Institutional controls including environmental covenants and groundwater use restrictions;
- Containment including low permeability sheet pile wall to restrict groundwater flow and contaminant migration; and
- Monitoring to assess attenuation of contaminants in groundwater via natural processes.

8.2. Description of Cleanup Action Alternatives

Cleanup action alternatives for the Upland Area were developed by combining technologies and process options retained through the remedial technology screening evaluation (Tables 17 and 18) to address soil and groundwater contamination and meet the CAOs. Cleanup action alternatives developed for the Site are summarized in Table 19 and described in the following sections. These alternatives represent a reasonable number and range of potentially applicable cleanup actions to provide a further basis for comparative evaluation. The cleanup action alternatives developed for the Site are based on a conceptual-level design for the implementation of the individual technologies described above. The design parameters used to



develop the alternatives are based on engineering judgment and the current knowledge of Site conditions. The final design for the preferred cleanup action alternative may require additional characterization and analysis of Site media and potential changes to specific plans for the future development of the Site to better define the cleanup action and associated costs. The comparative analysis for the cleanup action alternatives summarized below is presented in Section 9.2.

8.2.1. Cleanup Action Alternative 1 - Containment and Compliance Groundwater Monitoring

Cleanup Action Alternative 1 relies on containment technologies (i.e., capping and sheet pile wall) in conjunction with institutional controls to address Site contaminants. Capping in the form of asphalt paving will be completed in unpaved portions of the Site to further prevent stormwater infiltration through the soil column and mobility of contaminants in the subsurface. Existing stormwater infrastructure will be used to capture and treat stormwater runoff prior to discharge from the Site. Long-term monitoring would then be performed to evaluate groundwater conditions over time to assess contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants. Groundwater data collected prior to and following significant asphalt paving of the Site provide a line of evidence that suggests COCs that remain in place in saturated zone soils have stabilized and are not migrating downgradient toward the Guemes Channel since paving was completed.

Specific actions to be performed at the Site as part of Cleanup Action Alternative 1 include the following:

- Install asphalt pavement in the central and eastern portions of the Upland Area that currently consist of a gravel working surface to further prevent potential stormwater infiltration and contaminant leaching/migration through the soil column. This physical barrier will also further prevent direct contact with Site contaminants contained in the subsurface. Stormwater captured by the new paving will be directed to the stormwater treatment system at the Site. Note that the existing treatment system will require a capacity upgrade to handle the additional volume of stormwater that will result from the new paving.
- Maintain the existing concrete and asphalt caps in other portions of the Site to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site contaminants.
- Maintain the existing sheet pile wall that separates the Marine Area from the Upland Area to contain Site COCs identified in Upland Area soil as well as to provide a physical barrier to prevent direct contact. As previously discussed, tidal study results represent groundwater conditions prior to the 2008 Interim Action. As part of the interim action, an open cell bulkhead was installed in the central portion of the Site to extend the Upland Area northward and create additional land to facilitate DCI operations. As a result, the open cell bulkhead (which now separates the Upland Area from the Marine Area) provides a physical barrier that restricts the direct discharge of groundwater north to Guemes Channel.
- Maintain the existing fencing and security procedures to restrict public access to the Site.
- Install warning signs to inform Site workers and/or visitors to the Site regarding health risks and land use restrictions (as necessary).
- Implement a deed restriction (environmental covenant) compliant with the Uniform Environmental Covenants Act and with Ecology's model environmental covenant.



- Develop and implement a Compliance Monitoring Plan describing the groundwater confirmational and long-term monitoring to verify the effectiveness of the cleanup action including procedures for sample collection, sample frequency, data review, quality control and reporting.
- Perform compliance groundwater monitoring utilizing the existing network of wells to evaluate groundwater conditions over time to evaluate contaminant concentrations relative to the cleanup standards and natural attenuation performance of Site contaminants. It is assumed that groundwater monitoring would be completed on a semi-annual basis for up to five years targeting wet and dry season months. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Perform long-term groundwater monitoring utilizing the existing network of monitoring wells (MW-1 through MW-8) to evaluate groundwater conditions over time to ensure compliance with the cleanup standards and effectiveness of the cleanup action. It is assumed that groundwater monitoring would be completed once per Ecology Five Year Periodic Review period for up to 25 years following completion of the compliance monitoring period. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Develop and implement an Engineering Controls Monitoring and Maintenance Plan to identify the engineering and institutional controls that are being utilized at the Site, to provide guidelines for the monitoring and maintenance of the Site controls to ensure protection of human health and the environment, and to provide guidelines on the proper handling and disposal of soil and groundwater encountered during future Site maintenance and/or development activities.
- Perform annual inspection of the asphalt/concrete pavement cap and sheet pile wall that separates the Upland and Marine Areas to ensure the long-term performance of these containment barriers.

The estimated cost of Cleanup Action Alternative 1 is \$1,180,000 (Table M-1, Appendix M). The cost estimate is in 2019 dollars, include contingencies, and represent order-of-magnitude with a range of -30 percent to +50 percent based on EPA guidance (EPA 2000) and does not include the stormwater treatment system capacity upgrade. Existing Site features including paved portions of the Site, the location of sheet pile walls and existing monitoring well network that would be utilized to evaluate long-term groundwater conditions are shown on Figure 29.

8.2.2. Cleanup Action Alternative 2 - Partial Source Area Removal

Cleanup Action Alternative 2 includes partial removal of contaminant source areas located in the southeast portion of the Site which is generally centered around sampling location SB-12 followed by Site restoration. Contaminant source areas are defined as areas in which concentrations of metals (arsenic and nickel) in soil exceed three times (3x) the PCUL. Partial source area removal under Cleanup Action Alternative 2 will be used in conjunction with existing containment barriers as well as institutional controls to address remaining COCs in the central and western portions of the Site. Compliance monitoring would then be performed to verify the effectiveness of the removal action. Long-term monitoring would then be performed to evaluate groundwater conditions over time to assess contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants.

Specific actions to be performed at the Site as part of Cleanup Action Alternative 2 include the following:



- Develop and implement an Engineering Design Report describing the plans and procedures that will be used for cleanup of the Site.
- Removal followed by offsite disposal of an estimated 3,600 in-place cubic yards (bcy) of soil to a permitted landfill from the identified southeast contaminant source area which is defined as an area in which the concentration of metals (arsenic and nickel) in soil exceed three times (3x) the PCUL. Prior to construction, monitoring well MW-7 located in the removal area will be decommissioned. Verification sampling will be performed to confirm the vertical and lateral extent of remediation from this area during construction. Upon verification of the remedial excavation extent, backfill consisting of overburden material generated during construction that is determined to be both structurally and chemically suitable for reuse and/or imported structural fill will be placed.
- Restore portions of the asphalt pavement which are disturbed by the removal action.
- Install a replacement well for MW-7 decommissioned as part of the removal action.
- Maintain the existing concrete and asphalt pavement caps in other portions of the Site to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to remaining Site contaminants.
- Maintain the existing sheet pile wall that separates the Marine Area from the Upland Area to contain Site COCs identified in Upland Area soil as well as to provide a physical barrier to prevent direct contact. As previously discussed, the open cell bulkhead (which now separates the Upland Area from the Marine Area) physical barrier that restricts the direct discharge of groundwater north to Guemes Channel.
- Maintain the existing fencing and security procedures to restrict public access to the Site.
- Install warning signs to inform Site workers and/or visitors to the Site regarding health risks and land use restrictions (as necessary).
- Implement a deed restriction (environmental covenant) compliant with the Uniform Environmental Covenants Act and with Ecology's model environmental covenant.
- Develop and implement a Compliance Monitoring Plan describing the performance, confirmational and long-term monitoring to verify the effectiveness of the cleanup action including procedures for sample collection, sample frequency, data review, quality control and reporting.
- Perform compliance monitoring utilizing existing network of monitoring wells (MW-2B through MW-6) and new replacement monitoring well (MW-7A) following soil removal activities and restoration to evaluate groundwater conditions relative to the cleanup standards and effectiveness of the removal action. It is assumed that groundwater monitoring would be completed on a quarterly basis for a period of one year followed by semi-annual monitoring for up to an additional four years targeting wet and dry season months. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Perform long-term groundwater monitoring utilizing the existing network of monitoring wells (MW-1 through MW-6 and MW-8) and new replacement monitoring well (MW-7A) to evaluate groundwater conditions over time to ensure compliance with the cleanup standards and natural attenuation of Site contaminants remaining in place at the Site. It is assumed that groundwater monitoring would be completed once per Ecology Five Year Periodic Review period for up to 25 years following completion of the compliance monitoring period. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.



- Develop and implement an Engineering Controls Monitoring and Maintenance Plan to identify the engineering and institutional controls that are being utilized at the Site, to provide guidelines for the monitoring and maintenance of the Site controls to ensure protection of human health and the environment, and to provide guidelines on the proper handling and disposal of soil and groundwater encountered during future Site maintenance and/or development activities.
- Perform annual inspection of the asphalt/concrete pavement cap and sheet pile wall that separates the Upland and Marine Areas to ensure the long-term performance of these containment barriers.

The estimated cost of Cleanup Action Alternative 4 is \$2,120,000 (Table M-2, Appendix M). The cost estimate is in 2019 dollars, include contingencies, and represent order-of-magnitude with a range of -30 percent to +50 percent based on EPA guidance (EPA 2000). Existing Site features including paved portions of the Site, the location of sheet pile walls and existing monitoring well network that would be utilized to evaluate long-term groundwater conditions are shown on Figure 30.

8.2.3. Cleanup Action Alternative 3 - Source Area In Situ Treatment

Cleanup Action Alternative 3 includes in situ treatment of contaminant source areas (described above) to stabilize contaminants. In situ treatment of contaminant source areas under Cleanup Action Alternative 3 will be used in conjunction with containment and institutional controls as discussed under previous alternatives to prevent direct contact and the migration of contaminants contained in the subsurface. Compliance monitoring would then be performed to verify the effectiveness of the cleanup action. Long-term monitoring would then be performed to evaluate groundwater conditions over time to assess contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants.

Specific actions to be performed at the Site as part of Cleanup Action Alternative 3 include the following:

- Develop and implement an Engineering Design Report describing the plans and procedures that will be used for cleanup of the Site.
- Perform in situ soil stabilization through the injection of chemical reagents into the subsurface within contaminant source areas to immobilize (i.e., precipitate and/or bond to soil particles) Site contaminants. In situ stabilization would be completed using standard drilling and injection technologies and would result in minimal disturbance to the existing asphalt paved surfaces that are working to restrict stormwater infiltration through the soil column and thus preventing contaminant leaching/migration through the subsurface. Contaminant source areas are defined as areas in which concentrations of metals (arsenic and nickel) in soil exceed three times (3x) the PCUL. Groundwater data in the vicinity of MW-2B and MW-7 indicate that PCUL exceedances are only observed in areas in which metals concentrations in soil exceed three times the PCUL. In other portions of the Site, groundwater data indicate that concentrations of arsenic and nickel in soil are stable and are not adversely impacting groundwater. In addition, groundwater data indicate that concentrations of total cPAH TEQ in soil are stable and are not adversely impacting groundwater at the Site. Based on a review of the existing data, three (3) contaminant source areas have been identified and are located in the eastern, southern and central portions of the Site (Figure 31).
- Install asphalt pavement in the central and eastern portions of the Upland Area that currently consist of a gravel working surface to further prevent potential stormwater infiltration and contaminant leaching/migration through the soil column. This physical barrier will also further prevent direct contact



with Site contaminants contained in the subsurface. Stormwater captured by the new paving will be directed to the stormwater treatment system at the Site. Note that the existing treatment system will require a capacity upgrade to handle the additional volume of stormwater that will result from the new paving.

- Maintain the existing concrete and asphalt pavement caps in other portions of the Site to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site contaminants.
- Maintain the existing sheet pile wall that separates the Marine Area from the Upland Area to contain Site COCs identified in Upland Area soil as well as to provide a physical barrier to prevent direct contact. As previously discussed, the open cell bulkhead (which now separates the Upland Area from the Marine Area) physical barrier that restricts the direct discharge of groundwater north to Guemes Channel.
- Maintain the existing fencing and security procedures to restrict public access to the Site.
- Install warning signs to inform Site workers and/or visitors to the Site regarding health risks and land use restrictions (as necessary).
- Implement a deed restriction (environmental covenant) compliant with the Uniform Environmental Covenants Act and with Ecology's model environmental covenant.
- Develop and implement a Compliance Monitoring Plan describing the groundwater performance, confirmational and long-term monitoring to verify the effectiveness of the cleanup action including procedures for sample collection, sample frequency, data review, quality control and reporting.
- Install temporary monitoring wells upgradient and downgradient of the identified source areas and complete performance monitoring to verify the effectiveness of the cleanup action and whether the injection of additional chemical reagents is necessary to stabilize Site contaminants remaining in place. It is assumed that performance groundwater monitoring would be completed on quarterly basis for up to 2 years.
- Perform compliance monitoring following performance monitoring activities utilizing the existing network of monitoring wells (MW-1 through MW-8) to evaluate groundwater conditions over time to assess residual contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants. It is assumed that groundwater monitoring would be completed on a semi-annual basis for up to 5 years targeting wet and dry season months. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Perform long-term groundwater monitoring utilizing the existing network of monitoring wells (MW-1 through MW-8) to evaluate groundwater conditions over time to ensure compliance with the cleanup standards and natural attenuation of Site contaminants remaining in place at the Site. It is assumed that groundwater monitoring would be competed once per Ecology Five Year Periodic Review period for up to 25 years following completion of the compliance monitoring period. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Develop and implement an Engineering Controls Monitoring and Maintenance Plan to identify the engineering and institutional controls that are being utilized at the Site, to provide guidelines for the monitoring and maintenance of the Site controls to ensure protection of human health and the environment, and to provide guidelines on the proper handling and disposal of soil and groundwater encountered during future Site maintenance and/or development activities.



Perform annual inspection of the asphalt/concrete pavement cap and sheet pile wall that separates the Upland and Marine Areas to ensure the long-term performance of these containment barriers.

The estimated cost of Cleanup Action Alternative 3 is \$2,610,000 (Table M-3, Appendix M). The cost estimate is in 2019 dollars, include contingencies, and represent order-of-magnitude with a range of -30 percent to +50 percent based on EPA guidance (EPA 2000). Existing Site features including paved portions of the Site, the location of sheet pile walls and existing monitoring well network that would be utilized to evaluate long-term groundwater conditions are shown on Figure 31.

8.2.4. Cleanup Action Alternative 4 - Source Area Removal

Cleanup Action Alternative 4 includes removal of contaminant source areas (described above) followed by Site restoration. Source area removal under Cleanup Action Alternative 4 will be used in conjunction with containment and institutional controls as discussed in previous alternatives to prevent direct contact and the migration of remaining contaminants contained in the subsurface. Compliance monitoring would then be performed to verify the effectiveness of the cleanup action. Long-term monitoring would then be performed to evaluate groundwater conditions over time to assess contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants.

Specific actions to be performed at the Site as part of Cleanup Action Alternative 4 include the following:

- Develop and implement an Engineering Design Report describing the plans and procedures that will be used for cleanup of the Site.
- Removal followed by offsite disposal of an estimated 9,000 in-place cubic yards (bcy) of soil to a permitted landfill from the identified contaminant source areas in the central and eastern portions of the Site. Contaminant source areas are defined as areas in which concentrations of metals (arsenic and nickel) in soil exceed three times (3x) the PCUL. Prior to construction, monitoring well MW-7 located in the removal area will be decommissioned. Verification sampling will be performed to confirm the vertical and lateral extent of remediation from this area during construction. Upon verification of the remedial excavation extent, backfill consisting of overburden material generated during construction that is determined to be both structurally and chemically suitable for reuse and/or imported structural fill will be placed.
- Restore portions of the asphalt pavement which are disturbed by the removal action.
- Install a replacement well for MW-7 decommissioned as part of the removal action.
- Maintain the existing concrete and asphalt pavement caps in other portions of the Site to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to remaining Site contaminants.
- Maintain the existing sheet pile wall that separates the Marine Area from the Upland Area to contain Site COCs identified in Upland Area soil as well as to provide a physical barrier to prevent direct contact. As previously discussed, the open cell bulkhead (which now separates the Upland Area from the Marine Area) physical barrier that restricts the direct discharge of groundwater north to Guemes Channel.
- Maintain the existing fencing and security procedures to restrict public access to the Site.
- Install warning signs to inform Site workers and/or visitors to the Site regarding health risks and land use restrictions (as necessary).



- Implement a deed restriction (environmental covenant) compliant with the Uniform Environmental Covenants Act and with Ecology's model environmental covenant.
- Develop and implement a Compliance Monitoring Plan describing the performance, confirmational and long-term monitoring to verify the effectiveness of the cleanup action including procedures for sample collection, sample frequency, data review, quality control and reporting.
- Perform compliance monitoring utilizing existing network of monitoring wells (MW-1 through MW-8) following soil removal activities and restoration to evaluate groundwater conditions over time to assess residual contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants in other portions of the Site. It is assumed that groundwater monitoring would be completed on a quarterly basis for a period of one year followed by semi-annual basis for up to an additional four years four additional years targeting wet and dry season months. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Perform long-term groundwater monitoring utilizing the existing network of monitoring wells (MW-1 through MW-8) to evaluate groundwater conditions over time to ensure compliance with the cleanup standards and effectiveness of the cleanup action. It is assumed that groundwater monitoring would be completed once per Ecology Five Year Periodic Review period for up to 25 years following completion of the compliance monitoring period. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Develop and implement an Engineering Controls Monitoring and Maintenance Plan to identify the engineering and institutional controls that are being utilized at the Site, to provide guidelines for the monitoring and maintenance of the Site controls to ensure protection of human health and the environment, and to provide guidelines on the proper handling and disposal of soil and groundwater encountered during future Site maintenance and/or development activities.
- Perform annual inspection of the asphalt/concrete pavement cap and sheet pile wall that separates the Upland and Marine Areas to ensure the long-term performance of these containment barriers.

The estimated cost of Cleanup Action Alternative 4 is \$4,390,000 (Table M-4, Appendix M). The cost estimate is in 2019 dollars, include contingencies, and represent order-of-magnitude with a range of -30 percent to +50 percent based on EPA guidance (EPA 2000). Existing Site features including paved portions of the Site, the location of sheet pile walls and existing monitoring well network that would be utilized to evaluate long-term groundwater conditions are shown on Figure 32.

8.2.5. Cleanup Action Alternative 5 - Site-Wide In Situ Treatment

Cleanup Action Alternative 5 includes Site-wide in situ treatment to stabilize Site contaminants. In situ treatment under Cleanup Action Alternative 5 will be used in conjunction with institutional controls as discussed under previous alternatives to prevent direct contact and the migration of contaminants contained in the subsurface. Compliance monitoring would then be performed to verify the effectiveness of the cleanup action. Long-term monitoring would then be performed to evaluate groundwater conditions over time to assess contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants.

Specific actions to be performed at the Site as part of Cleanup Action Alternative 5 include the following:



- Develop and implement an Engineering Design Report describing the plans and procedures that will be used for cleanup of the Site.
- Perform in situ soil treatment through the injection of chemical reagents/oxidants into the subsurface throughout the Site to immobilize (i.e., precipitate and/or bond to soil particles) and/or degrade Site contaminants. In situ treatment would be completed using standard drilling and injection technologies and would result in minimal disturbance to the existing asphalt paved surfaces.
- Install asphalt pavement in the central and eastern portions of the Upland Area that currently consist of a gravel working surface to provide a physical barrier to prevent direct contact to remaining Site contaminants. Direct stormwater captured by the new paving to the stormwater treatment system at the Site. Note that the existing treatment system will require a capacity upgrade to handle the additional volume of stormwater that will result from the new paving.
- Maintain the existing concrete/asphalt/gravel pavement caps in other portions of the Site to provide a physical barrier to prevent direct contact to remaining Site contaminants.
- Maintain the existing fencing and security procedures to restrict public access to the Site.
- Install warning signs to inform Site workers and/or visitors to the Site regarding health risks and land use restrictions (as necessary).
- Implement a deed restriction (environmental covenant) compliant with the Uniform Environmental Covenants Act and with Ecology's model environmental covenant.
- Develop and implement a Compliance Monitoring Plan describing the performance, confirmational and long-term monitoring to verify the effectiveness of the cleanup action including procedures for sample collection, sample frequency, data review, quality control and reporting.
- Install temporary monitoring wells upgradient and downgradient of the identified contamination areas and complete performance monitoring to verify the effectiveness of the cleanup action and whether the injection of additional chemical reagents is necessary to stabilize Site contaminants remaining in place. It is assumed that performance groundwater monitoring would be competed on quarterly basis for up to 2 years.
- After this time period, compliance monitoring would be completed utilizing the existing network of monitoring wells (MW-1 through MW-8) to evaluate groundwater conditions over time to assess residual contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants. It is assumed that groundwater monitoring would be competed on a semi-annual basis for up to 3 years following performance monitoring activities targeting wet and dry season months. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Perform long-term groundwater monitoring utilizing the existing network of monitoring wells (MW-1 through MW-8) to evaluate groundwater conditions over time to ensure compliance with the cleanup standards and effectiveness of the cleanup action. It is assumed that groundwater monitoring would be competed once per Ecology Five Year Periodic Review period for up to 25 years following completion of the compliance monitoring period. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Develop and implement an Engineering Controls Monitoring and Maintenance Plan to identify the engineering and institutional controls that are being utilized at the Site, to provide guidelines for the



monitoring and maintenance of the Site controls to ensure protection of human health and the environment, and to provide guidelines on the proper handling and disposal of soil and groundwater encountered during future Site maintenance and/or development activities.

Perform annual inspection of the asphalt/concrete pavement cap and sheet pile wall that separates the Upland and Marine Areas to ensure the long-term performance of these containment barriers.

The estimated cost of Cleanup Action Alternative 5 is \$7,030,000 (Table M-5, Appendix M). The cost estimate is in 2019 dollars, include contingencies, and represent order-of-magnitude with a range of -30 percent to +50 percent based on EPA guidance (EPA 2000). Existing Site features including paved portions of the Site, the location of sheet pile walls and existing monitoring well network that would be utilized to evaluate long-term groundwater conditions are shown on Figure 33.

8.2.6. Cleanup Action Alternative 6 - Site-Wide Removal

Cleanup Action Alternative 6 includes removal of contaminated soil throughout the Site followed by restoration. Compliance monitoring would then be performed to verify the effectiveness of the cleanup action. Specific actions to be performed at the Site as part of Cleanup Action Alternative 6 include the following:

- Removal followed by offsite disposal of an estimated 46,500 bcy of soil to a permitted landfill. Prior to construction, monitoring well MW-2B, MW-4 and MW-7 located in the removal area will be decommissioned. The vertical and lateral extent of remedial will be verified through confirmation samples collected during construction. Remedial excavations will be backfilled with overburden material generated during construction that is determined to be both structurally and chemically suitable for reuse and/or imported structural fill.
- Restore portions of the asphalt pavement which are disturbed by the removal action.
- Install replacement wells at locations MW-2B, MW-4 and MW-7 and complete compliance groundwater monitoring utilizing the existing network of monitoring wells (MW-1 through MW-8) to verify the effectiveness of the cleanup action. It is assumed that groundwater monitoring would be completed on quarterly basis for up to two years. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.
- Complete long-term groundwater monitoring at MW-8 to evaluate groundwater conditions over time to assess contaminant concentrations relative to the cleanup standards and natural attenuation of Site contaminants. It is assumed that groundwater monitoring would be completed on a semi-annual basis for up to 10 years targeting wet and dry season months. After this time period, Ecology would be consulted to determine additional groundwater requirements (if any) for the Site.

The estimated cost of Cleanup Action Alternative 4 is \$15,060,000 (Table M-6, Appendix M). The cost estimate is in 2019 dollars, include contingencies, and represent order-of-magnitude with a range of -30 percent to +50 percent based on EPA guidance (EPA 2000). Existing Site features including paved portions of the Site, the location of sheet pile walls and existing monitoring well network that would be utilized to evaluate long-term groundwater conditions are shown on Figure 34.



9.0 EVALUATION OF CLEANUP ALTERNATIVES

Evaluation criteria and a comparative analysis of the cleanup action alternatives developed for the Site are summarized in the following sections. Each alternative is evaluated with respect to the MTCA evaluation criteria and are compared to each other relative to their expected performance under each criterion. The components of the cleanup action alternatives are described above in Section 3.2 and are summarized in Table 19. A detailed evaluation of the alternatives relative to the MTCA evaluation criteria is presented in Table 20, and the results of the evaluation are summarized in Table 21. Concept design level cleanup action cost estimates for each alternative are presented in Appendix M.

9.1. Cleanup Alternative Evaluation Criteria

Threshold requirements for cleanup actions under MTCA and the additional criteria used to evaluate the cleanup action alternatives are described in the following sections.

9.1.1. Threshold Requirements

Cleanup actions performed under MTCA must comply with basic threshold requirements. Cleanup action alternatives that do not comply with the threshold requirements are not considered suitable cleanup actions under MTCA. As provided in WAC 173-340-360(2)(a), the four threshold requirements for remedial actions are that they must:

- Protect human health and the environment;
- Comply with cleanup standards;
- Comply with applicable state and federal laws; and
- Provide for compliance monitoring.

The following sections further describe the threshold requirements.

9.1.1.1. Protection of Human Health and the Environment

The results of cleanup actions performed under MTCA must ensure that both human health and the environment are protected.

9.1.1.2. Compliance with Cleanup Standards

Compliance with cleanup standards requires, in part, that cleanup levels are met at the applicable points of compliance. If a cleanup action does not comply with cleanup standards, the cleanup action is an interim action, not a cleanup action. Where a cleanup action involves containment of hazardous substance concentrations exceeding cleanup levels at the point of compliance, the cleanup action may be determined to comply with cleanup standards, provided the requirements specified in WAC 173-340-740(6)(f) are met.

9.1.1.3. Compliance with Applicable State and Federal Laws

Cleanup actions conducted under MTCA must comply with applicable state and federal laws. The term "applicable state and federal laws" includes legally applicable requirements and those requirements that Ecology determines to be relevant and appropriate as described in WAC 173-340-710.

9.1.1.4. Provision for Compliance Monitoring

The cleanup action must allow for compliance monitoring in accordance with WAC 173-340-410. Compliance monitoring consists of protection monitoring, performance monitoring and confirmational



monitoring. Protection monitoring is conducted to confirm that human health and the environment are adequately protected during construction, and the operation and maintenance period of a cleanup action. Performance monitoring is conducted to confirm that the cleanup action has attained cleanup standards, remediation levels and/or other performance standards, as appropriate. Confirmational monitoring is conducted to confirm the long-term effectiveness of the cleanup action.

9.1.2. Other MTCA Requirements

Under MTCA, when selecting from the alternatives that meet the minimum requirements, the alternatives shall be further evaluated against the following additional criteria:

- Use permanent solutions to the maximum extent practicable [WAC 173-340-360(2)(b)(i)]. MTCA requires that when selecting from cleanup action alternatives that fulfill the threshold requirements, the selected action shall use permanent solutions to the maximum extent practicable [WAC 173-340-360(2)(b)(i)]. MTCA specifies that the permanence of these qualifying alternatives shall be evaluated by balancing the costs and benefits of each of the alternatives using a "disproportionate cost analysis" in accordance with WAC 173-340-360(3)(e). The criteria for conducting this analysis are described in Section 4.1.3 below.
- Provide a reasonable restoration time frame [WAC 173-340-360(2)(b)(ii)]. In accordance with WAC 173-340-360(2)(b)(ii), MTCA places a preference on those cleanup action alternatives that, while equivalent in other respects, can be implemented in a shorter period of time. MTCA includes a summary of factors to be considered in evaluating whether a remedial action provides for a reasonable restoration time frame [WAC 173-340-360(4)(b)].
- Consideration of Public Concerns [WAC 173-340-360(2)(b)(iii)]. Ecology will consider public comments submitted during the RI and FS process in making its preliminary selection of an appropriate remedial action alternative. This preliminary selection is subject to further public review and comment when the proposed remedy is published in the Draft Cleanup Action Plan (DCAP).

9.1.3. MTCA Disproportionate Cost Analysis

The MTCA disproportionate cost analysis (DCA) is used to further evaluate which of the alternatives that meet the threshold requirements are permanent to the maximum extent practicable. This analysis involves comparing the costs and benefits of alternatives and selecting the alternative whose incremental costs are not disproportionate to the incremental benefits. The evaluation criteria for the DCA are specified in WAC 173-340-360(2) and (3), and include protectiveness, permanence, cost, long-term effectiveness, management of short-term risks, implementability and consideration of public concerns.

As outlined in WAC 173-340-360(3)(e), MTCA provides a methodology that uses the criteria listed below to determine whether the costs associated with each cleanup alternative are disproportionate relative to the incremental benefit of the alternative above the next lowest-cost alternative. The comparison of benefits relative to costs may be quantitative but will often be qualitative. When possible for this FS, quantitative factors such as mass of contaminant removed or percentage of area of impacts remaining were compared to costs for the alternatives evaluated, but many of the benefits associated with the criteria described below were necessarily evaluated qualitatively. Costs are disproportionate to benefits if the incremental costs of the more permanent alternative exceed the incremental degree of benefits achieved by the other lower-cost alternative [WAC 173-340-360(e)(i)]. Where two or more alternatives are equal in benefits, the less costly alternative is retained as the preferred alternative [WAC 173-340-360(e)(ii)(c)].



MTCA criteria used in the DCA are described in the following sections.

9.1.3.1. Protectiveness

The overall protectiveness of a cleanup action alternative is evaluated based on several factors. First, the extent to which human health and the environment are protected and the degree to which overall risk at a Site is reduced are considered. Both on-site and off-site reduction in risk resulting from implementing the alternative are considered.

9.1.3.2. Permanence

MTCA specifies that when selecting a cleanup action alternative, preference shall be given to actions that are "permanent solutions to the maximum extent practicable." Evaluation criteria include the degree to which the alternative permanently reduces the toxicity, mobility or mass of hazardous substances, including the effectiveness of the alternative in destroying the hazardous substances, the reduction or elimination of hazardous substance releases and sources of releases, the degree of irreversibility of waste treatment processes, and the characteristics and quantity of treatment residuals generated.

9.1.3.3. Cost

The analysis of cleanup action alternative costs under MTCA includes the costs associated with implementing an alternative, including design, construction, long-term monitoring, and institutional controls. Costs are intended to be comparable among different alternatives to assist in the overall analysis of relative costs and benefits of the alternatives. The costs to implement an alternative include the cost of construction, the net present value of any long-term costs, and agency oversight costs. Long-term costs include operation and maintenance costs, monitoring costs, equipment replacement costs, and the cost of maintaining institutional controls. Unit costs used to develop overall remediation costs for this FS were derived using a combination of published engineering reference manuals (i.e., R.S. Means); construction cost estimates solicited from applicable vendors and contractors; review of actual costs incurred during similar, applicable projects; and professional judgment.

9.1.3.4. Long-Term Effectiveness

Long-term effectiveness is a parameter that expresses the degree of certainty that the alternative will be successful in maintaining compliance with cleanup standards over the long-term performance of the cleanup action. The MTCA regulations contain a specific preference ranking for different types of technologies that is to be considered as part of the comparative analysis. The ranking places the highest preference on technologies such as reuse/recycling, treatment, immobilization/solidification, and disposal in an engineered, lined, and monitored facility. Lower preference rankings are applied for technologies such as on-site isolation/containment with attendant engineered controls, and institutional controls and monitoring.

9.1.3.5. Management of Short-term Risks

Evaluation of this criterion considers the relative magnitude and complexity of actions required to maintain protection of human health and the environment during implementation of the cleanup action. Cleanup actions carry short-term risks, such as potential mobilization of contaminants during construction, or safety risks typical of large construction projects. In-water dredging activities carry a risk of temporary water quality degradation and potential sediment recontamination. Some short-term risks can be managed through the use of Best Management Practices (BMPs) during project design and construction, while other risks are inherent to project alternatives and can offset the long-term benefits of an alternative.



9.1.3.6. Implementability

Implementability is an overall metric expressing the relative difficulty and uncertainty of implementing the cleanup action. Evaluation of implementability includes consideration of technical factors such as the availability of mature technologies and experienced contractors to accomplish the cleanup work. It also includes administrative factors associated with permitting and completing the cleanup.

9.1.3.7. Consideration of Public Concerns

The public involvement process under MTCA is used to identify potential public concerns regarding remedial action alternatives. The extent to which an alternative can address public concerns is considered as part of the evaluation process. This includes concerns raised by individuals, community groups, local governments, tribes, federal and state agencies, and other organizations that may have an interest in or knowledge of the Site. In particular, the public concerns for this Site would generally be associated with environmental concerns and performance of the cleanup action, which are addressed under other criteria such as protectiveness and permanence.

9.2. Evaluation and Comparison of Cleanup Action Alternatives

Cleanup action alternatives developed for the Site were evaluated with respect to the MTCA threshold and other relevant requirements described above, then were compared to each other relative to the expected performance under each criterion. The following sections provide an evaluation and comparative analysis of the cleanup action alternatives developed to address Site contamination.

9.2.1. Threshold Requirements

Cleanup action alternatives developed for the Site incorporate varying combinations of capping and containment, in situ treatment and/or removal technologies in combination with institutional controls to meet the minimum threshold requirements of protecting human health and the environment, complying with cleanup standards, and complying with applicable state and federal laws within a reasonable time frame. Remediation technologies are intended to be protective of the ecological receptors, prevent direct contact with Site workers and prevent the offsite migration of contaminants. Performance and/or compliance monitoring would be completed for each cleanup action alternative to confirm compliance with the cleanup standards at the point of compliance. To ensure the effectiveness and compliance with the cleanup standards over time, Cleanup Action Alternatives 1 through 5 which leave residual contamination in place also have a provision for long-term monitoring of engineering controls to contain Site contaminants and groundwater monitoring to confirm compliance with the cleanup standards. Due to the complete removal of contaminated soil under Cleanup Action Alternative 6, long-term compliance monitoring would only be required at MW-8 to evaluate groundwater conditions at this location over time to assess contaminant concentrations relative to the cleanup standards and potential for natural attenuation.

Cleanup Action Alternative 1 achieves the lowest level of protectiveness by isolating residual contamination with a combination of institutional and engineering controls (i.e., protective caps and containment barriers to prevent direct contact). Cleanup Action Alternative 3 results in a slightly higher degree of protectiveness relative to Alternative 1 through in situ treatment/stabilization, however the full extent of contamination remains in place at the site. Cleanup Action Alternatives 2 and 4 achieves a higher degree of protectiveness than Alternative 3 as the result of removal of contaminants from the identified source areas. Cleanup Action Alternatives 5 and 6 have the highest degree of protectiveness through Site-wide stabilization of residual contamination (Cleanup Action Alternative 5) and through complete removal of Site contamination (Cleanup Action Alternative 6).



9.2.2. Other MTCA Requirements

9.2.2.1. Permanent to the Maximum Extent Practicable

Similar to threshold criteria, Cleanup Action Alternative 1 achieves the lowest level of permanence while Cleanup Action Alternative 3 achieves a higher degree of permanence over Cleanup Action Alternative 1 with the use of treatment technologies. Cleanup Action Alternatives 2 and 4 provide a higher degree of permanence through the use of permanent removal technologies to address source area material. Sitewide in situ treatment to stabilize Site contaminants under Cleanup Action Alternative 5 followed by complete removal of soil contamination under Cleanup Action Alternative 6 provide the highest degree of permanence.

9.2.2.2. Reasonable Restoration Time Frame

Each of the cleanup action alternatives evaluated are all expected to achieve remedial action objectives in reasonable restoration time frames. The time frame for design, permitting, contracting, and construction for each of the proposed cleanup action alternatives is expected to be on the order of 1 to 4 years. The restoration time frame for Cleanup Action Alternatives 1, 2 and 4 is expected to be the lowest; however, monitoring of the engineered caps and groundwater conditions to document compliance with cleanup objectives is expected to occur over time (minimum of 5 years). Cleanup Action Alternatives 3 and 5 are expected to occur over a 2- to 3-year time frame due to the potential of multiple rounds of in situ chemical reagent injection or complexities in soil removal to address source areas and/or Site-wide contamination followed by compliance monitoring to confirm the effectiveness of the cleanup action over a 5 year period. Cleanup Action Alternative 6 is expected to have the longest restoration time frame due to the complexities of remediating within an active shipyard including sequencing of the remedial action to limit potential impacts to DCI operations.

9.2.2.3. Considerations of Public Concerns

The cleanup alternatives proposed for the Site are generally expected to be acceptable to the public. Cleanup Action Alternative 6 achieves the greatest level of protection and certainty as a result of the greatest level of contaminated removal; however, is the most intrusive alternative requiring a high level of coordination with DCI to minimize disruption to shipyard operations and with the city and local residents for any concerns related to increased truck traffic during construction. Cleanup Action Alternatives 2 through 5 also result in a high level of protection and certainty through source removal/treatment and containment technologies; however, residual contamination will remain in place with each of these alternatives which may draw public concern for potential exposure. The public may be concerned for Cleanup Action Alternative 1 due to the level of residual contamination left in place; however, containment technologies combined with institutional controls will results in protection to human health and the environment and prevent contaminant exposure which may limit the potential for public concern.

9.2.3. MTCA Disproportionate Cost Analysis

The MTCA DCA is used to make a relative comparison the costs and benefits of the remedial alternatives under consideration for the Site. The comparison of benefits relative to costs are quantitative; however, quantitative factors such as mass of contaminant treated/removed, percentage of area of impacts remaining following implementation of the cleanup alternative and/or actual alternative cost when compared to the relative benefit score criteria described above may be more of a qualitative assessment in that a cleanup alternative with similar relative benefit to cost ratios. The remedial alternative with the highest ratio of benefit to cost is identified as the preferred alternative.



The evaluation of the level of achievement for how each individual criterion applies to each alternative, using a numeric scoring scale of 1 (lowest) to 10 (highest) and the methodology described above in Section 9.1.3. Table 20 presents an evaluation of the relative benefits ranking and numeric score for the individual criterion. Table 21 summarizes the results of the DCA and ranks each of the cleanup action alternatives based on relative cost and benefit. Preliminary planning level construction cost estimate for each cleanup action alternative incorporated into the DCA are presented in Appendix M and have an accuracy that is considered to be -30 to +50 percent based on EPA's Guide to Developing and Documenting Cost Estimates During the Feasibility Study (EPA 2000). The conclusions of this evaluation are shown on Figure 35.

9.3. Preferred Cleanup Action Alternative and Basis for Selection

Under MTCA, "costs are disproportionate to benefits if the incremental costs of the alternative over that of a lower cost alternative exceed the incremental degree of benefits achieved by the alternative over that of lower cost alternative" (WAC 173-340-360[3][e][i]). From the resulting benefit/cost ratio (Figure 35), the overall cost for Cleanup Action Alternatives 3 through 6 are disproportionate to the environmental benefit that they provide relative to Cleanup Action Alternatives 1 and 2. Furthermore, the environmental benefit for Cleanup Action Alternative 2 is greater than for Cleanup Action Alternative 1. As a result, Cleanup Action Alternative 2 emerges as the preferred alternative for the Site. This alternative may be refined during development of the Draft Cleanup Action Plan (DCAP).

Cleanup Action Alternative 2 addresses Site contamination through partial source area removal in the southeast portion of the Site and is not expected to significantly disrupt DCI operations or limit access to buildings and other infrastructure utilized by DCI. Remaining Site contamination will be address through containment technologies (i.e., isolation by existing paving and sheet pile wall) in conjunction with institutional controls. This alternative is permanent to the maximum extent practicable and reduces immediate risk to potential human and ecological receptors through:

- Partial removal of Site contaminants exceeding three times the PCUL in the southeast portion of the Site. Results of groundwater monitoring provide a line of evidence that contaminant levels exceeding three times the PCUL in soil in this area are adversely impacting groundwater. Additionally, groundwater data collected prior to and following significant asphalt paving of the Site also provides a line of evidence that contaminants remaining in place in other portions of the Site have stabilized and are not migrating downgradient toward the Guemes Channel since paving was completed.
- Maintenance of the existing concrete and asphalt caps in other portions of the Site to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site contaminants.
- Maintain the existing sheet pile wall that separates the Marine Area from the Upland Area to contain Site COCs identified in Upland Area soil as well as to provide a physical barrier to prevent direct contact.
- Maintenance of the existing fencing and security procedures to restrict public access to the Site.
- Installation of warning signs to inform Site workers and/or visitors to the Site regarding health risks and land use restrictions (as necessary).
- Implementation of a deed restriction (environmental covenant) compliant with the Uniform Environmental Covenants Act and with Ecology's model environmental covenant. It is expected that an



environmental covenant will not be required for the southeast portion of the DCI lease area following remedial excavation as part of this cleanup action.

Implementation of Cleanup Action Alternative 2 will result in contaminant mass reduction, containment and prevention of direct human contact with remaining COCs, and the prevention of stormwater infiltration/leaching and migration of COCs contained in other portions of the Site to the Marine Area. Although Cleanup Action Alternative 2 does not achieve complete mass removal, containment of the COCs in the fill body by way of the existing paving has been shown empirically to be effective at preventing the mobility and migration of COCs in groundwater at the Site. Therefore, the contaminant mass reduction acts to increase protectiveness and permanence of the alternative over containment focused alternatives.

10.0 CONCLUSIONS

10.1. Remedial Investigation and Interim Action

Pursuant to the Ecology Agreed Order DE DE-07TCPHQ -5080 and MTCA cleanup regulations (Chapter 173-340 WAC), remedial investigation activities were completed by the Port to characterize environmental conditions at the Anacortes Port of Dakota Creek site located along the shoreline of Guemes Channel at the northern terminus of Q Avenue in Anacortes, Washington.

Upland Area environmental data was collected in accordance with the Ecology-approved RI/FS Work Plan to supplement and fill identified data gaps in existing data for the Site, to determine the nature and extent of contamination in soil and groundwater. Marine Area environmental data was collected in accordance with the Ecology-approved RI/FS Work Plan to characterize the nature and extent of sediment contamination in areas previously identified as exceeding the PCULs. The data collected for the RI provided the basis for identification and evaluation of cleanup action alternatives for addressing Site contamination. In addition, the Marine Area data was used to support planning and design of the Interim Action that was completed in 2008.

10.2. Nature and Extent of Contamination

Based on a review of the Upland and Marine Area RI results as well as a review of previous Site characterization data, the following COCs were identified for sediment, groundwater and soil at the Site:

- Sediment Metals including arsenic, cadmium, copper, lead, mercury and zinc, TBT, LPAHs, HPAHs, cPAHs, PCBs, dioxin and furans were identified as COCs for Marine Area sediment. However, interim action dredging in the Marine Area in 2008 resulted in the complete removal of sediment contamination at the Site. As a result, sediment is no longer considered a media of concern.
- **Groundwater** Metals including arsenic and nickel, and cPAHs were identified as groundwater COCs for the Upland Area.
- Soil Metals including arsenic and nickel, and cPAHs were identified as soil COCs for the Upland Area. In addition, results of previous environmental studies identified concentrations of gasoline-, diesel-and/or heavy oil-range petroleum hydrocarbons exceeding PCULs. As indicated above, the Port completed independent cleanup actions (i.e., 1991 UST, 2001 Hydraulic Winch and 2002 Petroleum and Marine Railway Cleanup Actions) to remove the previously identified petroleum contamination from the Site. Confirmation sample results obtained from the limits of these excavations indicated that the petroleum contamination was successfully removed from the Site. However, confirmation soil sample



results for these areas could not be independently validated. Therefore, petroleum hydrocarbons within the footprints of the previously completed cleanup action areas are unverified until subsequent sampling result confirm their removal.

The nature and extent for COCs in Site media is further discussed below.

10.2.1. Marine Area

Interim action dredging and excavation activities completed between July and November 2008 in general accordance with the Ecology-approved RI/FS Work Plan and Interim Action Work Plan Addendum were completed to remove identified COCs from the Marine Area. The interim action resulted in the removal of approximately 26,000 cubic yards (approximately 38,000 tons) of contaminated sediment from the southern portion of the Marine Area for upland landfill disposal. An additional 230,000 cubic yards of sediment (approximate) determined to be suitable for open-water disposal by the DMMP was then dredged from the Marine Area as part of the Project Pier 1 redevelopment for disposal at the Rosario Strait dispersive site to meet the design grade of -35 feet MLLW. Due to the completeness of the interim action and subsequent Marine Area dredging to remove contaminated sediment deposits and up to 30 feet of native glaciomarine deposits, no identified sediment contamination remained in the Marine Area and sediment is no longer considered a media of concern for the Site.

10.2.2. Upland Area

In the Upland area, COCs including arsenic, nickel and cPAHs were detected at concentrations exceeding PCULs in soil throughout the Site. In the eastern portion of the Site, arsenic and nickel exceeded PCULs in fill deposits from the ground surface up to a depth of approximately 8 feet bgs. In the north central portion of the Site, arsenic and nickel exceeded PCULs in fill deposits from the ground surface up to a depth of approximately 10 feet bgs. In the southwestern portion of the Site, arsenic, nickel and cPAHs exceeded the PCUL in fill deposits from the ground surface up to a depth of approximately 13 feet bgs. Results of soil/sediment samples collected at the Site from the underlying native surface show that the Upland Area PCUL exceedances are limited to the overlying fill soil and do not extend into the native surface.

Between 2015 and 2016, DCI replaced a significant portion of their gravel working surface with asphalt pavement which prevents stormwater infiltration through the soil column. A comparison of the initial (2008 to 2013) groundwater monitoring results to the recent semi-annual groundwater monitoring results (2016 to 2017) show that the paved surfaces are limiting the infiltration, leaching and subsequent migration of contaminants through the soil column to groundwater.

At shoreline monitoring well locations, the following COCs were detected at concentrations greater than the groundwater PCUL since completion of the Upland Area paving activities:

- Dissolved nickel detected at a concentration of 8.3 μg/L marginally exceeded the groundwater PCUL of 8.2 μg/L at shoreline monitoring well location MW-2B during the February 2017 monitoring event. However, total nickel at this location was not detected at a concentration exceeding the groundwater PCUL during this event and dissolved nickel did not exceed the PCUL in subsequent monitoring events at this location.
- Total cPAH TEQ concentrations and/or dissolved arsenic exceeded the groundwater PCUL at shoreline monitoring well location MW-8 during one or more semi-annual monitoring events between February



2016 and August 2017. However, cPAHs and arsenic concentrations at monitoring well location MW-1 (upgradient monitoring well location) were less than the PCUL for each of these monitoring events. Supplemental sampling and analysis to further evaluate soil conditions in the vicinity of MW-8 did not identify potential source materials for cPAHs or arsenic in saturated soil adjacent to or upgradient of this location.

At upgradient monitoring well locations, the following COCs were detected at concentrations greater than the groundwater PCUL since completion of the Upland Area paving activities:

■ Total and dissolved arsenic and nickel, and total cPAH TEQ concentrations exceeded the groundwater PCULs at monitoring well location MW-7 during one or more semi-annual monitoring events. However, at downgradient monitoring well locations MW-3A and MW-6, total and dissolved arsenic and nickel, and cPAHs either were not previously detected or were detected at concentrations less than the groundwater PCUL during each of the four semi-annual groundwater monitoring events.

The distribution of COCs including arsenic, nickel and cPAHs in Upland Area soil and groundwater representing current conditions at the Site are summarized on Figures 20 through 25.

10.3. Supplemental Soil Investigation

Results of previous environmental studies identified concentrations of gasoline-, diesel- and/or heavy oil-range petroleum hydrocarbons exceeding PCULs in historical fill deposits from the ground surface to a depth of approximately 8 feet bgs in the central and eastern portions of the Site. Between 1991 and 2002, the Port completed independent cleanup actions in these areas to remove the previously identified petroleum contamination from the Site. However, confirmation soil sample results for these areas could not be independently validated and will require subsequent sampling to confirm the completeness of the removal.

A supplemental soil investigation is proposed as a pre-remedial design activity to verify the effectiveness of the previous cleanup actions. As part of this investigation, sampling will be performed within the footprint of the previously completed remedial excavations which extended to depths greater than 1-foot (i.e., remedial excavations extending beneath the former gravel working surface). Samples will be collected from a depth of below the base of the previous remedial excavations to represent the previous verification sample interval and analyzed for COCs previously identified for these areas including gasoline-, diesel- and heavy oil-range petroleum hydrocarbons and/or cPAHs. The results of this investigation will be reported in the CAP and used to refine the preferred cleanup action alternative for the Site.

10.4. Feasibility Study Development and Preferred Cleanup Action Alternative Selection

The nature and extent of contamination in sediment has been characterized in accordance with the Ecology-approved RI/FS Work Plan using PCULs for identified human and ecological receptors and exposure pathways based on current and future land use. Based on the results of the RI, sufficient data has been collected to identify and evaluate cleanup alternatives for contaminated media at the Site. A range of remedial technologies were screened considering the media requiring cleanup, COC present and current and future land use to develop a reasonable number and range of potentially applicable cleanup actions which were then evaluated relative to the following criteria:

Compliance with cleanup standards and applicable laws;



- Provision for a reasonable restoration time frame; and
- Use of permanent solutions to the maximum extent practicable by comparison of the following criteria:
 - Protectiveness;
 - Permanence:
 - Cost:
 - Effectiveness over the long term;
 - Short-term risk management;
 - Net environmental benefit;
 - Technical and administrative implementability; and
 - Consideration of Public Concerns.

As a result of this evaluation, Cleanup Action Alternative 2 emerged as the preferred alternative which meets the minimum threshold requirements, achieves a high level of environmental benefit and is not disproportionate in cost relative to the other alternatives evaluated. Implementation of Cleanup Action Alternative 2 will result in contaminant mass reduction in the southeast portion of the Site and will be used in conjunction with containment technologies and institutional controls to prevent direct human contact and reduce the potential for leaching and migration of residual COCs contained within the fill soil within a reasonable restoration time frame.

11.0 LIMITATIONS

This report has been prepared for the exclusive use of the Port of Anacortes, their authorized agents and regulatory agencies for the Dakota Creek Industries Site located at 115 Q Avenue in Anacortes, Washington. No other party may rely on the product of our services unless we agree in advance and in writing to such reliance. Within the limitations of scope, schedule and budget, our services have been executed in accordance with generally accepted environmental science practices in this area at the time this report was prepared. No warranty or other conditions, express or implied, should be understood.

Any electronic form, facsimile or hard copy of the original document (email, text, table, and/or figure), if provided, and any attachments are only a copy of the original document. The original document is stored by GeoEngineers, Inc. and will serve as the official document of record.

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Table 1

Proposed Sediment Cleanup Levels for Protection of Benthic Organisms

Dakota Creek Industries Anacortes, Washington

		Critorio f	or Protection		Organisms					
	Sadi	ment Manage			rent Effects T	hrochold		Pronosed	Sediment	
		ment Manago Standard ¹ (SM		Арра	rent Effects (i (AET) Criteria				p Level ³	
		Sediment	Cleanup		(AET) CITTETIO	Second		Oleanu	Level	
		Quality	Screening		Lowest	Lowest		Organic		Organic
		Objectives	Level	_	AET	AET		Carbon		Carbon
Analyte	Units	(SQ0)	(CSL)	Units	(LAET)	(2LAET)	Units	(0.5% to 3.5%)	Units	(<0.5% or >3.5%)
Metals				T		Т				
Arsenic	_	57	93		57	93		57		57
Cadmium		5.1	6.7		5.1	6.7		5.1		5.1
Chromium		260	270	=	260	270	i	260		260
Copper	ma/ka	390	390	ma/ka	390	390	mg/kg	390	ma /lea	390
Lead	mg/kg	450	530	mg/kg	450	530	mg/kg	450	mg/kg	450
Mercury		0.41	0.59		0.41	0.59		0.41		0.41
Nickel Silver		NE 6.1	NE 6.1		NE 6.1	NE 6.1		NE 6.1		NE 6.1
Zinc		410	960		410	960		410		410
Low Molecular Weight Polycyclic A	romatic Hydr				410	300		410		410
Total LPAH ⁴	I I I I I I I I I I I I I I I I I I I	370	780	I	5,200	5,200		370		5,200
2-Methylnaphthalene	1	38	64	1	670	670		38		670
Acenaphthene	1	16	57	1	500	500		16		500
Acenaphthylene		66	66		1,300	1,300		66		1,300
Anthracene	mg/kg OC	220	1,200	µg/kg	960	960	mg/kg OC	220	µg/kg	960
Fluorene		23	79		540	540		23		540
Naphthalene		99	170	=	2,100	2,100		99		2,100
Phenanthrene		100	480		1,500	1,500		100		1,500
High Molecular Weight Polycyclic A	romatic Hydr	ocarbons (HP	AHs)	•						
Total HPAH ⁵		960	5,300		12,000	17,000		960		12,000
Benzo(a)anthracene		110	270		1,300	1,600		110		1,300
Benzo(a)pyrene		99	210		1,600	1,600		99		1,600
Total Benzofluoranthenes		230	450		3,200	3,600		230		3,200
Benzo(ghi)perylene	mg/kg OC	31	78	µg/kg	670	720	mg/kg OC	31	μg/kg	670
Chrysene		110	460	MB/ 1/8	1,400	2,800	IND NO OO	110	MB/ 1/8	1,400
Dibenzo(a,h)anthracene		12	33		230	230		12		230
Fluoranthene		160	1,200		1,700	2,500		160		1,700
Indeno(1,2,3-cd)pyrene		34	88		600	690		34		600
Pyrene		1,000	1,400		2,600	3,300		1000		2,600
Chlorinated Organic Compounds						I .	ı			
1,2,4-Trichlorobenzene		0.81	1.8	'	31	51	i	0.81		31
1,2-Dichlorobenzene	mg/kg OC	2.3	2.3	µg/kg	35	50	mg/kg OC	2.3	µg/kg	35
1,4-Dichlorobenzene		3.1	9		110	110	:	3.1		110
Hexachlorobenzene		0.38	2.3		22	70		0.38		22
Phthalates		17	70		1 200	1 000		47		1 200
Bis(2-Ethylhexyl) Phthalate Butyl Benzyl Phthalate		47 5	78 64		1,300 63	1,900 900		47 4.9		1,300 63
Dibutyl Phthalate		220	1,700	-	1,400	1,400	:	220		1,400
Diethyl Phthalate	mg/kg OC	61	110	µg/kg	200	> 1,200	mg/kg OC	61	µg/kg	200
Dimethyl Phthalate	1	53	53		71	160		53		71
Di-N-Octyl Phthalate		58	4,500	-	6,200	6,200		58		6,200
Phenols	Į.		.,		0,200	3,233				5,255
2,4-Dimethylphenol		29	29		29	29		29		29
2-methylphenol		63	63	1	63	63		63		63
4-methylphenol	µg/kg	670	670	µg/kg	670	670	µg/kg	670	µg/kg	670
Pentachlorophenol		360	690		360	690		360		360
Phenol		420	1,200		420	1,200		420		420
Miscellaneous Extractables										
Dibenzofuran		15	58		540	540		15		540
Hexachlorobutadiene	mg/kg OC	3.9	6.2	µg/kg	11	120	mg/kg OC	3.9	µg/kg	11
N-Nitrosodiphenylamine		11	11	<u> </u>	28	40		11		28
Benzoic Acid	μg/kg	650	650	μg/kg	650	650	μg/kg	650	μg/kg	650
Benzyl Alcohol	με/ ng	57	73	µg/ ng	57	73	μg/ ng	57	μg/ ng	57
Polychlorinated Biphenyls (PCBs)										
Total PCBs (Sum of	mg/kg OC	12	65	mg/kg	0.13	1	mg/kg OC	12	mg/kg	0.13
Aroclors)	3 5		-				3 000		3 .6	

Notes:

mg/kg OC = milligram per kilogram normalized to organic carbon

μg/kg = microgram per kilogram

ng/kg = nanogram per kilogram

-- = Criteria not applicable or not available



 $^{^{\}rm 1}$ Sediment Management Standards (SMS) (Chapter 173-204 WAC).

² Apparent Effects Threshold (AET) Criteria from Ecology's SCUM II guidance (Table 8-1; Ecology 2019).

³ The organic carbon normalized SMS criteria are applicable to sediment with a total organic carbon (TOC) concentration ranging from 0.5 to 3.5 percent. Sediment with TOC concentrations outside of the 0.5 to 3.5 percent range are screened against the AET Screening Level on a dry weight basis (EPA 1988).

⁴ Total LPAHs are the total of naphthalene, acenaphthylene, acenaphthene, fluorene, phenanthrene and anthracene; 2-methylnapthalene is not included in the sum of LPAHs.

⁵ Total HPAHs are the total of fluoranthene, pyrene, benzo(a)anthracene, chrysene, benzofluoranthenes, benzo(a)pyrene, indeno(1,2,3-c-d)pyrene, dibenzo(a,h)anthracene and benzo(g,h,i)perylene. mg/kg = milligram per kilogram

Table 2

Proposed Sediment Cleanup Levels for Protection of Human Health and Higher Trophic Level Ecological Receptors Dakota Creek Industries Anacortes, Washington

			Anacon	es, Washingt	011			
		Criteria for Prote Bioaccumulation via Consumption of		n Health		Adjus	tment	Proposed Sediment Cleanup Level ⁵ (After Adjustment for
		Aquatic Organisms Natural		shing ²	Proposed	_	tors	Background and PQL)
Analyte	Units	Background or PQL ¹	Carc. (at 10 ⁻⁶ risk)	Non- Carc.	Sediment Cleanup Level	Natural Background ³	PQL⁴	Subtidal Sediment (below -3 ft MLLW)
Metals		1	T		I	1		
Arsenic		11	3.0	1,300	3.0	11	5	11
Cadmium		0.8		4,400	0.8	0.8	0.2	0.8
Chromium Copper				6,900,000 180,000	6,900,000 180,000	62 45	0.5 0.2	6,900,000 180,000
Lead	mg/kg	21			21	21	2	21
Mercury		0.2		460	0.2	0.2	0.05	0.2
Nickel			-	92,000	92,000	50	1	92,000
Silver		-		23,000	23,000	0.24	0.3	23,000
Zinc				1,400,000	1,400,000	93	1	1,400,000
Tributyltin								
Tributyltin, bulk	µg/kg	73 ⁶		1,200	73	-	3.86	73
Interstitial Tributyltin,	µg/L	0.15 ⁶			0.15		0.0052	0.15
porewater Low Molecular Weight Polycycli	c Aromatic F	I lydrocarbons (I PAHs)						
2-Methylnaphthalene	omatic i			16,000	16,000		0.005	16,000
Acenaphthene		-		240,000	240,000		0.005	240,000
Acenaphthylene				240,000	240,000		0.005	240,000
Anthracene	mg/kg			1,200,000	1,200,000		0.005	1,200,000
Fluorene				160,000	160,000	2	0.005	160,000
Naphthalene		-	-	79,000	79,000	-	0.005	79,000
Phenanthrene				1,200,000	1,200,000		0.005	1,200,000
High Molecular Weight Polycycl	ic Aromatic I	T				1		
Benzo(a)anthracene					NE DALLETO		0.005	NE BALL TEO
Benzo(a)pyrene			cPAH TEQ	cPAH TEQ	cPAH TEQ	-	0.005	cPAH TEQ
Total benzofluoranthenes			-	120,000	NE 120,000		0.005 0.005	NE 120,000
Benzo(ghi)perylene Chrysene	mg/kg			120,000	NE		0.005	120,000 NE
Dibenzo(a,h)anthracene			_	-	NE		0.005	NE NE
Fluoranthene		7	-	160,000	160,000		0.005	160,000
Indeno(1,2,3-cd)pyrene			-	_	NE		0.005	NE
Pyrene		-		120,000	120,000		0.005	120,000
Carcinogenic Polycyclic Aromat	ic Hydrocarb	ons (cPAHs)						
Total cPAHs - TEQ	µg/kg	21	680	120,000	21	21	5	21
Chlorinated Organic Compound	S				П	1		
1,2,4-Trichlorobenzene		-	140	41,000	140		0.2	140
1,2-Dichlorobenzene	mg/kg	-	700	370,000	370,000		0.2	370,000
1,4-Dichlorobenzene		-	760	290,000	760		0.2	760
Hexachlorobenzene Phthalates		-	2.5	3,300	2.5		0.001	2.5
Bis(2-Ethylhexyl) Phthalate			290	82,000	290		0.05	290
Butyl Benzyl Phthalate		_	2,100	820,000	2,100		0.03	2,100
Dibutyl Phthalate				410,000	410,000		0.02	410,000
Diethyl Phthalate	mg/kg	<u> </u>		3,300,000	3,300,000		0.02	3,300,000
Dimethyl phthalate					NE		0.02	NE
Di-N-Octyl Phthalate		-		41,000	41,000		0.02	41,000
Phenois		T	T		П	1		
2,4-Dimethylphenol				82,000	82,000		25	82,000
2-Methylphenol	ma/ka			200,000	200,000		20	200,000
4-Methylphenol Pentachlorophenol	mg/kg		 8.7	410,000 17,000	410,000 9		20 100	410,000 100
Pentacnioropnenoi Phenol			8.7	1,200,000	1,200,000		100	1,200,000
Miscellaneous Extractables	1	I.	<u>I</u>	1,200,000		<u>I</u>	100	1,200,000
Dibenzofuran				4,100	4,100		0.02	4,100
Hexachlorobutadiene	mg/kg		52	4,100	52		0.001	52
N-Nitrosodiphenylamine			830		830		0.02	830
Benzoic Acid	μg/kg			16,000,000	16,000,000		200	16,000,000
Benzyl Alcohol				410,000	410,000		20	410,000
Polychlorinated Biphenyls (PCB	s)							
Total PCBs (Sum of	mg/kg	0.0035	1.9		0.0035	0.0035	0.001	0.0035
Aroclors) Dioxins and Furans			<u> </u>		<u> </u>		<u> </u>	
Total Dioxins/Furans -		_		F 000	_	Α.	- 7	-
Human Health TEQ	ng/kg	5	55	5,000	5	4	5 ⁷	5



Notes:

- ¹ Bioaccumulative chemicals include arsenic, cadmium, lead, mercury, carcinogenic polycyclic aromatic hydrocarbons (cPAHs), dioxin-like polychlorinated biphenyls (PCBs), and dioxins/furans. Currently site-specific human health and ecological risk-based sediment screening levels have not been developed for bioaccumulative chemicals. Therefore, sediment screening levels for these chemicals are based on the natural background or the practical quantification limit (PQL), whichever is higher.
- ² Sediment cleanup levels for the protection of human health via direct contact are calculated using equations and input parameters provided by Ecology in the Draft Sediment Cleanup Users Manual (SCUM) II guidance (Ecology 2019).
- ³ Natural background concentrations are derived from the calculated values (90/90 UTL) from the Bold plus dataset and presented in Table 10-1 of Ecology's Draft SCUM II (Ecology 2019) guidance document.
- ⁴ PQL is from the Remedial Investigation/Feasibility Study Work Plan (GeoEngineers 2008) and is the lowest available value from Analytical Resources Inc. (ARI) of Tukwila, Washington.
- The proposed cleanup levels (PCULs) presented in this table are to provide an evaluation of human health and ecological risk for higher trophic level ecological receptors. Human health and higher trophic level ecological receptor PCULs are chosen from lowest of bioaccumulative and direct contact pathways. If the risk-based value is lower than natural background or practical quantitation limit (PQL), the screening level defaults to the higher of natural background or PQL. The human health PCULs for subtidal areas include marine areas at elevations below -3 feet MLLW and the applicable direct contact pathway is net fishing.
- ⁶ The bioaccumulative cleanup levels protective of higher trophic level ecological receptors is from the Dredged Material Management Program (DMMP) bioaccumulation triggers for bulk and porewater tributyltin. Measurement of tributyltin in interstitial water provides a more direct measure of potential bioavailability, and hence toxicity, than bulk sediment concentrations. Therefore porewater tributyltin
- 7 PQL for Dioxin TEQ is the Programmatic PQL values from Ecology's SCUM II guidance (Table 11-1; Ecology 2019).

mg/kg = milligram per kilogram

μg/kg = microgram per kilogram

ng/kg = nanogram per kilogram

-- = No criterion is currently available for this analyte

Total LPAHs are the total of naphthalene, acenaphthylene, acenaphthene, fluorene, phenanthrene and anthracene; 2-methylnaphthalene is not included in the sum of LPAHs.

Total HPAHs are the total of fluoranthene, pyrene, benzo(a)anthracene, chrysene, benzofluoranthenes, benzo(a)pyrene, indeno(1,2,3-cd)pyrene, dibenzo(a,h)anthracene and benzo(g,h,i)perylene. Blue shading identifies the basis for proposed sediment cleanup level.

Green shading identifies the proposed sediment cleanup level after adjustment for background and the PQL.



Table 3

Proposed Groundwater Cleanup Levels

Dakota Creek Industries Anacortes, Washington

										Anacorto	s, Washingt											
						Protection ce Water						or Protection or Intrusion		С	Criteria for Pro of Sedime							
		(Ch	ater Quality		Federal Marine Water Quality Criteria for Washington ² (40 CFR 131.45)		(CWA §30	ality Criteria ³ 94(a))	Surfac Cleanu	lethod B e Water ip Level		Method C		nt Cleanup	Equili Part	ition		Groundwater		Adjust Fact		Proposed Groundwater
		Protection Aquat	n of Marine tic Life	Protection of Human Health (Organisms	Protection of Human Health		n of Marine tic Life	Protection of Human Health (Organisms	`	d Formula lue) Non-		ndwater ning Level Non-		ive (SCO) ng Level ⁴	Coeffic (L/		Groundwater Concentration Protective of	to Sediment Pathway Complete ⁷	Proposed Groundwater	Background	Practical Quantitation Limit9	Cleanup Level ¹⁰ (After Adjustment for Background and
Analyte	Units	Acute	Chronic	Only)	(Organisms Only)	Acute	Chronic	Only)	Carc.	Carc.	Carc.	Carc.	C _{sed}	Units	(CLARC)	(metals)	Sediment ⁶	(Yes/No)	Cleanup Level	Concentration ⁸	(PQL)	PQL)
Metals																						
Arsenic	μg/L	69	36	10	0.14	69	36	0.14	0.1	18		-	11	mg/kg	-	29	3.7E+02	Yes	0.1	8	4.5	8
Cadmium	μg/L	42	9.3		-	33	7.9		-	-	-	-	0.8	mg/kg	-	6.7	1.1E+02	No	7.9	2	4.0	7.9
Total Chromium ¹¹	μg/L	1,100	50	-	-	1,100	50		-	490	-		260	mg/kg	-	19	1.3E+04	No	50	10	10.0	50
Copper	μg/L	4.8	3.1	-	_	4.8	3.1		-	2,900	1	-	390	mg/kg	-	22	1.7E+04	Yes	3.1	20	10.0	20
Lead	μg/L	210	8.1	-	-	210	8.1	-	-	-	-	-	21	mg/kg		10,000	2.1E+00	Yes	2.1	10	1.0	10
Mercury ¹²	μg/L	1.8	0.025	-	-	1.8	0.94				- '	1.9	0.2	mg/kg	-	52	3.8E+00	Yes	0.025	-	0.025	0.025
Nickel	μg/L	74	8.2	190	100	74	8.2	4,600	-	1,100	-	-	92,000	mg/kg	-	65	1.4E+06	No	8.2	-	5.6	8.2
Silver	μg/L	1.9	-			1.9		-	-	26,000	-		6.1	mg/kg	-	8	6.9E+02	No	1.9		10.0	10
Zinc	μg/L	90	81	2,900	1,000	90	81	26,000	-	17,000	-		410	mg/kg	-	62	6.5E+03	Yes	81	160	25.0	160
Petroleum Hydrocarbons ¹³																						
Gasoline-Range	μg/L	-	-				-	-	-	800	ï	-		-			-	No	800		250	800
Diesel-Range	μg/L	-	-			-			-	500	-	-		-			-	No	500		250	500
Heavy Oil-Range	μg/L		-				-		-	500	-	-		-		-	-	No	500	-	500	500
BETX Compounds				-														-	-		-	
Benzene	μg/L			1.6	-			16	23	2,000	24	230	-		62			No	1.6	_	1.0	1.6
Ethylbenzene	μg/L	-	-	270	31	-	-	130	-	6,900	45	6,100	_	-	204	-	-	No	31	_	1.0	31
Toluene	μg/L	-	-	410	130	-	-	520	-	19,000	-/	34,000	-	-	140		-	No	130	-	1.0	130
Xylenes	µg/L		-	_				-	-	-	T-	630	-		311		-	No	630		2.0	630
Volatile Organic Compounds (VOCs)		•	•	•	•	•				•		•	•		•	•		•	•	•		•
1,1,1,2-Tetrachloroethane	μg/L	_	-	_		_	-/		_	_	74	_				-	-	No	74		1.0	74
1,1,1-Trichloroethane	μg/L	-	-	160,000	50,000		-	200,000	7	930,000	-	12,000			135	-		No	12,000		1.0	12,000
1,1,2,2-Tetrachloroethane	μg/L	-	-	0.46	0.3		-	3	6.5	10,000	62				79	-	-	No	0.3		1.0	1
1,1,2-trichloro-1,2,2-	µg/L			_		_						2,400					_		2,400	_	1.0	2,400
trifluoroethane (CFC113)	μg/ L						-					2,400	-					No	2,400		1.0	2,400
1,1,2-Trichloroethane	μg/L	-	-	1.8	0.9	-	- '	8.9	25	2,300	79	10			75	-	-	No	0.9	-	1.0	1.0
1,1-Dichloroethane	μg/L	-				-		-	-	-	110	-			53		-	No	110		1.0	110
1,1-Dichloroethene	μg/L	-	-	4,100	4,000		-	20,000	-	23,000		280	-		65		-	No	280	-	1.0	280
1,1-Dichloropropene	μg/L		-	-	-		-	-				-			-	-	-	No	NE	-	1.00	NE
1,2,3-Trichlorobenzene	μg/L	-	-	-	-	-	-	-				-	-				-	No	NE		5.00	NE
1,2,3-Trichloropropane	μg/L		-	-	-		-	-	7			-		-	-	-		No	NE	-	2.0	NE
1,2,4-Trichlorobenzene	μg/L	-	-	0.14	0.04			0.076	2	230	-	84			1659			No	0.04	-	5.0	5.0
1,2,4-Trimethylbenzene	μg/L		-		-							61					-	No	61	-	1.0	61
1,2-Dibromo-3-chloropropane	μg/L	-	-	-	-	-	-	-	-	-	-	-			-		-	No	NE		5.0	NE
1,2-Dibromoethane (EDB)	μg/L	-	-			-	-				2.7	590	-	-	66	-	-	No	2.7	-	1.0	2.7
1,2-Dichlorobenzene	µg/L	-	-	2,500	800	-	-	3,000	-	4,200	-	5,700			379	-	-	No	800	-	1.0	800
1,2-Dichloroethane (EDC)	μg/L	-	-	120	73	-	-/	650	59	13,000	42	310	-	-	38	-	-	No	42	-	1.0	42
1,2-Dichloropropane	μg/L	_	-	3.1		-	-	31	43	25,000	39	62			47	-	-	No	3.1		1.0	3.1
1,3,5-Trimethylbenzene	μg/L	-	-	-	-		-	-	-	-	-	-			-		-	No	NE	-	1.0	NE
1,3-Dichlorobenzene	μg/L	-	-	16	2		-	10	-	-	-						-	No	2		1.0	2
1,3-Dichloropropane	µg/L		-	-	-		-		-		-						-	No	NE		1.0	NE
1,4-Dichlorobenzene	µg/L	-	-	580	200	-		900	22	3,300	49	17,000			616	-	-	No	22	-	1.0	22
2,2-Dichloropropane	μg/L	-	-	-					-	-	-	-					-	No	NE		1.00	NE
2-Butanone (MEK)	μg/L	-	-	-	-	-			-	-	-	3,800,000					-	No	3,800,000		5.0	3,800,000
2-Chloroethyl Vinyl Ether	μg/L		-	-		-	-		-	-	-	-	-	-	-	-	-	No	NE	_	1.0	NE
2-Chlorotoluene	μg/L	-	-	-		-	-	-	_	-	-	-				-	-	No	NE		1.0	NE



Application							tection	riteria for Prot	Cı		r Protection	Criteria fo					Protection	Criteria for					
Mart Quality Cities Peters Water			1]	TV	nt	of Sedimen			r Intrusion	of Vapo		1			ce Water	1					
Application	Proposed Groundwater				Groundwater		ition	Partit	•				e Water p Level	Surface Cleanu	•	(CWA §30	<u> </u>	Quality Criteria for Washington ²		apter 173-2	(Cha		
Management Man	Cleanup Level ¹⁰	Practical			to Sediment					_				1 '	Protection of				Protection of				
Manage M	on (After Adjustment	Quantitation		Proposed	Pathway	Concentration	· · · · · · · · · · · · · · · · · · ·	(L/k	ng Level*	Screeni	ing Level	Screen	ue)	Val	Human Health	tic Life	Aquat	Protection of	Human Health	tic Life	Aquat		
4 Abenduncer 1951	for Background and PQL)	Limit9 (PQL)				_	_	(CLARC)	Units	C _{sed}		Carc.		Carc.		Chronic	Acute		, ,	Chronic	Acute	Units	Analyte
Abstract Processor No.	NE	5.0		NE	No	-					-	-	-	-		-	-	-		-	-	µg/L	2-Hexanone
Marchenic Marc	NE	1.00		NE	No		-	-		-	-	1	-	-	_		-	-	-	-	-	µg/L	
Advantage Adva	1,000,000	5.0	_	1,000,000					_		1,000,000				_				_			µg/L	,
Arrisina	00.000.000			00 000 000							00 000 000												, , ,
Memoratement Memo	32,000,000	5.0				-		0.575					-								-		
Development 19/4	50	50			1					-						-	1				-		
Service processors	1.0	1.0			-		-			-	850				· ·	-					-		·
Secondary 1966 - - 77 19 - - 100 220 14,000 2,000 - - 100 - 10	NE	1.00			-		-				_				1	-		+			-		
Seminormentance 1961	NE 10	1.00														-							
Garbon Fearbristie Bigl.	12	1.0			1	-		-		-	200				1	-					-		
Gation friscriptions	28	1.0		_	1							_	970		10,000				,		-		
Characterise Mart -	870	1.0				1				-						1		+					
Choresthane	1.00	1.0			1										1	-							
Cheromorem	200	1.0			-	-		224		-			5,000		800	-					-		
Chiconstature	40,400	1.0	-	,		-	-				-					-				-	-		
Cist_12-Dicharcerehene	12	1.0	-					-		-	,	12	6,900	56	2,000			600	1,200	-	-		
Cite All Control proposed 1971	340	1.0	-			-			-	-	340				-	-	-	-	-	-	-		
Distribution Dist	NE	1.0			-		-	35.5	-		-	-		-		-	-	-	-	-	-		·
Dibromomethane 19/2 - - - - - - - - -	NE	1.0				-					-	-	-			-	-	-		-	-		
Dichlorothromethane 19/2 - - 3.6 2.8 - - 27 28 14.000 18 - - - 55 - - No 2.8 -	2.2	1.0				-		63.1	-	-		45	14,000	21	21	-	-	2.2	3	-	-	μg/L	Dibromochloromethane
Dichlorodiffuoromethane (CFC 12) Eg/L - - - - - - - - -	NE	1.0			1	-					-		-			-	-			-	-	μg/L	Dibromomethane
Hessachlorobutadiene	2.8	1.0				-		55				18	14,000	28	27	-	-	2.8	3.6	-	-	μg/L	
Sopropylbenzene	12	1.0	-		-	-	-		-	-	12			-		-	-		-				` ′
Methyl lodide	5.0	5.0			1	-		53700	-	-		8	930	30	0.01	-	-	0.01	4.1	-	-		Hexachlorobutadiene
Methyl Ether (MTBE) μ/L - - - - - - - - -	1,600	1.0		, , , , , , , , , , , , , , , , , , ,	1	-					1,600		-	_	-	-	-	-	-	-	-		
Methylene Chloride Mg/L	NE	1.0	-		-	-							-	-	-		-			-	-	μg/L	·
Naphthalene	6,100	5.0		,		-		-	-	-			-			-						μg/L	, , ,
n-Butylbenzene μg/L	100	2.0			1	-	-	1						3,600	1,000	-	-	100	250	-	-		•
n-Propylbenzene	89	5.0			No	-		1191	-		360	89	4,900			-	-			-	-	µg/L	Naphthalene
P-Isopropyltoluene	NE	1.0				-						-	-	-		-	-	-		-	-		·
sec-Butylbenzene μg/L <td>4,900</td> <td>1.0</td> <td>-</td> <td></td> <td>No</td> <td>-</td> <td>-</td> <td></td> <td>-</td> <td>-</td> <td>4,900</td> <td>-</td> <td>-</td> <td>-</td> <td>-</td> <td>-</td> <td></td> <td></td> <td></td> <td>-</td> <td>-</td> <td></td> <td>n-Propylbenzene</td>	4,900	1.0	-		No	-	-		-	-	4,900	-	-	-	-	-				-	-		n-Propylbenzene
Styrene	NE	1.0	-	NE	No	-						-	-	-	-	-		-	-		-	µg/L	p-lsopropyltoluene
tert-Butylbenzene µg/L No NE 1 Tetrachloroethene (PCE) µg/L 7.1 2.9 29 100 500 240 100 265 No 2.9 0 Trans-1,2-Dichloroethene µg/L 5,800 1,000 4,000 33,000 250 38 No 250 38 No No 250 38 No No 250 38 No No <t< td=""><td>NE</td><td>1.0</td><td>-</td><td></td><td>-</td><td>-</td><td></td><td></td><td></td><td></td><td></td><td>-</td><td></td><td>-</td><td>-</td><td>-</td><td>-</td><td>-</td><td></td><td></td><td></td><td></td><td>· · · · · · · · · · · · · · · · · · ·</td></t<>	NE	1.0	-		-	-						-		-	-	-	-	-					· · · · · · · · · · · · · · · · · · ·
Tetrachloroethene (PCE) μg/L 7.1 2.9 29 100 500 240 100 265 No 2.9 - No 2.9 - Νο Τrans-1,2-Dichloroethene μg/L 5,800 1,000 4,000 - 33,000 - 250 380 No 250 - Νο	18,000	1.0			1	-		912	-	-	18,000	-	-	-			-			-	-		•
Trans-1,2-Dichloroethene μg/L 5,800 1,000 4,000 - 33,000 - 250 380 No 250 - No NE NE - No NE NE - No NE	NE	1.0	-		-	-	-		-	-						-	-	-		-			•
Trans-1,3-Dichloropropene μg/L No NE NE Trans-1,4-Dichloro-2-butene μg/L No NE NE Trans-1,4-Dichloro-2-butene μg/L No NE NE Trichloroethene (TCE) μg/L No NE NE No NE NE No NE NE No NE	2.9	0.2			1	-		1	-	-		240		100		-	-			-	-		` ,
Trans-1,4-Dichloro-2-butene μg/L No NE !! Trichloroethene (TCE) μg/L 0.86 0.7 7 13 120 26 8 94 No 0.7 17 13 120 26 8 No 0.7 No 0.7 17 13 120 26 8 No 0.7 No 0.7 No 0.7 17 13 120 26 8 No 0.7	250	1.0				-		38					33,000	-	4,000	-		1,000	-		-		,
Trichloroethene (TCE) μg/L 0.86 0.7 7 13 120 26 8 94 No 0.7 - Τητικη στικη στικ	NE	1.0	-			-		-	-	-	-	-	-	-	-	-	-	-		-	-		, ,
Trichlorofluoromethane (CFC 11)	NE	5.0	-		1	-			-	-				+	-		-			-			·
Vinyl Acetate µg/L No 17,000 !	1.0	1.0				-		94				26	120	13	7	-	-	0.7	0.86	-	-	µg/L	` '
	260	1.0	-			-		-		-				-	-	-		-	-				, ,
	17,000	5.0	-			-		1									-						•
	1.0	1.0		0.18	No			18.6	-	-	120	3	6,600	3.7	1.6	-		0.18	0.26			μg/L	Vinyl Chloride
Semi-Volatile Organic Compounds (SVOCs)			T	·	<u>, </u>	T	T			T	1		,	ı	T								• • •
	1.0	1.0	-	0.04	No				mg/kg			-		2.0	1	-	-						1,2,4-Trichlorobenzene
1,2-Dichlorobenzene µg/L 2,500 800 3,000 - 4,200 - 5,700 0.035 mg/kg 379 - 4.5E+00 No 800 - 5	800	1.0		800	No	4.5E+00		379	mg/kg	0.035	5,700	-	4,200	-	3,000	-	-	800	2,500	-	-	μg/L	1,2-Dichlorobenzene
	2	1.0	-		No					-			-			-	-			-		μg/L	· ·
	22	1.0		22	No	8.9E+00	-	616	mg/kg	0.11	17,000	49	3,300	22	900	-	-	200	580	-	-	µg/L	
2,2'-Oxybis[1-chloropropane] µg/L 900 4,000 37 42,000 No 37 - :	37	1.0		37	No	-					-	-	42,000	37	4,000	-	-	900		-	-	μg/L	2,2'-0xybis[1-chloropropane]
2,4,5-Trichlorophenol µg/L 1597 No 600 !	600	5.0	-	600	No		-	1597	-	-		-			600	-	-	-		-	-	µg/L	2,4,5-Trichlorophenol
2,4,6-Trichlorophenol µg/L 0.28 2.8 3.9 17 381 No 0.3 - No 0.3 - !	5.0	5.0		0.3	No			381		-			17	3.9	2.8				0.28			μg/L	2,4,6-Trichlorophenol



					Criteria for of Surfac							or Protection r Intrusion		C	riteria for Pro							
		(Ch	ater Quality	•	Federal Marine Water Quality Criteria for Washington ² (40 CFR 131.45)	Federa	(CWA §30	ality Criteria ³)4(a))	MTCA N Surface Cleanu		мтса	Method C		nt Cleanup	Equili Parti	brium ition		Groundwater		Adjust Fact		Proposed Groundwater
		Protection Aquat	n of Marine tic Life	Protection of Human Health	Protection of	Protection Aquat	n of Marine tic Life	Protection of Human Health		l Formula lue)		ndwater ning Level	1	ive (SCO) ing Level ⁴	Coeffic (L/		Groundwater Concentration	to Sediment Pathway	Proposed		Practical Quantitation	Cleanup Level ¹⁰ (After Adjustment
Analyte	Units	Acute	Chronic	(Organisms Only)	Human Health (Organisms Only)	Acute	Chronic	(Organisms Only)	Carc.	Non- Carc.	Carc.	Non- Carc.	\mathbf{C}_{sed}	Units	K _{oc} (CLARC)	K _d (metals)	Protective of Sediment ⁶	Complete ⁷ (Yes/No)	Groundwater Cleanup Level	Background Concentration ⁸	Limit9 (PQL)	for Background and PQL)
2,4-Dichlorophenol	μg/L	-	-	34	10	-	-	60	-	190	-		-	-	147		-	No	10		5.0	10
2,4-Dimethylphenol	µg/L	-		97	-		-	3,000		550	-		0.029	mg/kg	209		6.3E+00	No	97		1.0	97
2,4-Dinitrophenol	μg/L	-	-	610	100	-	-	300	-	3,500		-		-	0.01	-	-	No	100	-	10	100
2,4-Dinitrotoluene	μg/L	-	-	0.18		-	-	1.7	5.5	1,400	-	-		-	95.5	-	-	No	0.18		5.0	5.0
2,6-Dinitrotoluene	μg/L	-	-				-	- 4 000	-	- 4 000	-	-		-	69.2	-	-	No	NE 100		5.0	NE 100
2-Chloronaphthalene	µg/L	-		180	100	-	-	1,000		1,000		-	-	-	200	-	-	No	100	-	1.0	100
2-Chlorophenol	μg/L	-	-	17	-	-	-	800		97	-	-		-	388	_	-	No	17 NE	-	1.0	17 NE
2-Nitroaniline 2-Nitrophenol	µg/L	-	_	-		-	-			-	-		-		-		-	No No	NE NE		5.0 5.0	NE NE
3,3'-Dichlorobenzidine	µg/L µg/L		-	0.0033	-		_	0.15	0.046		-			-	724		_	No	0.003		5.0	5.0
3-Nitroaniline	μg/L μg/L			0.0033		-	-	0.15		-	_							No	NE		5.0	NE
4,6-Dinitro-2-methylphenol	μg/L	_	_	25	7		-	30	_	_	_			-				No	7	_	10.0	10
4-Bromophenyl-phenylether	µg/L			-	_						_	-						No	NE	_	1.0	NE NE
4-Chloro-3-methylphenol	µg/L	_		36				2000				-	-	_	_	_	_	No	36		5.0	36
4-Chloroaniline	µg/L	_	-		_		-	_	_	_	_	-	_	_	66.1		_	No	NE	_	5.0	NE
4-Chlorophenyl-phenylether	µg/L	_	_	_	_	_	_	_		/ -		-		_	_	_	_	No	NE		1.0	NE
4-Nitroaniline	µg/L	_	_			_	-		_	-	_		-	-			_	No	NE	_	5.0	NE
4-Nitrophenol	µg/L	-	-			-			_		-	-	-	-	_	_	-	No	NE		5.0	NE
Benzoic acid	μg/L	-	-			-	-		-	-	-	-	0.65	mg/kg	0.6		1.1E+03	No	NE	_	10.0	NE
Benzyl alcohol	μg/L			-	-			-	-	7	-		0.057	mg/kg	-	-	-	No	NE	-	5.0	NE
bis(2-Chloroethoxy)methane	μg/L			-				-				-		-	-	-	-	No	NE	-	1.0	NE
bis(2-chloroethyl)ether	μg/L			0.06	-			2.2	0.85		259	-	_	-	76	-		No	0.06	-	1.0	1.0
bis(2-Ethylhexyl)phthalate	μg/L	-	-	0.25	0.05	-	-	0.37	3.6	400	-	-	0.3	mg/kg	111,123	-	1.4E-01	No	0.05	-	1.0	1.0
Butylbenzylphthalate	μg/L			0.58	0.013		-	0.10	8.2	1,300			0.063	mg/kg	13,746	-	2.4E-01	No	0.013	-	1.0	1.0
Carbazole	μg/L	-	-	-	-		-	-	_		-	-			3390		_	No	NE	-	1.0	NE
Dibenzofuran	µg/L	-	-			-	-	-				-	0.54	mg/kg			-	No	NE		1.0	NE
Diethylphthalate	μg/L	-	-	5,000	200	-	-	600	7	28,000		-	0.2	mg/kg	82	-	9.3E+01	No	200	-	1.0	200
Dimethylphthalate	μg/L	-	-	130,000	600	-	-	2,000		-	-	-	0.071	mg/kg	-	-	-	No	600	-	1.0	600
Di-n-butylphthalate	μg/L	-	-	510	8	-	-	30		2,900	_	-	1.4	mg/kg	1,567	-	4.6E+01	No	8	-	1.0	8
Di-n-octylphthalate	µg/L				-		-	_					6.2	mg/kg	83,200,000		3.9E-03	No	NE		1.0	NE
Hexachlorobenzene	µg/L	-	-	0.000052	0.000005		-	0.000079	0.0005	0.24	-	-	0.003	mg/kg	80,000	-	1.7E-03	No	0.000005		1.0	1.0
Hexachlorobutadiene	µg/L	-		4.1	0.01	-	-	0.01	30	930	8	-	0.011	mg/kg	53,700	-	1.1E-02	No	0.01	-	1.0	1.0
Hexachlorocyclopentadiene	μg/L	-	-	630	1	-	-	4		3,600		-			200000	-	-	No	1		5.0	5.0
Hexachloroethane	μg/L			0.13	0.02		-	0.1	1.90	21	31	414			1780		-	No	0.02	-	2.0	2.0
Isophorone	μg/L	-	-	110	422		-	1,800	1,600	120,000	- 4 000	-			46.8		-	No	110	-	1.0	110
Nitrobenzene	μg/L		-	320	100		-	600	- 0.00	1,800	1,600	23,000		-	119		-	No	100		1.0	100
n-Nitroso-di-n-propylamine	μg/L			0.058	-	-	-	0.51	0.82			-	- 0.000		24	-	1 15+00	No	0.06		5.0	5.0
n-Nitrosodiphenylamine	µg/L			0.69		-		6.0	9.7				0.028	mg/kg	1,290	-	1.1E+00 2.7E+01	No No	0.7		1.0	1.0
o-Cresol (2-Methylphenol)	µg/L		-		-	-		_					0.063	mg/kg	91.2	-	2.7E+01 -	No No	NE NE		1.0	NE NE
p-Cresol (4-Methylphenol) Pentachlorophenol	μg/L	13	7.9	- 0.1	0.002	13	-	0.04	2	1,200		-	0.67	mg/kg	- 502	-	8.4E+00	No No	NE 0.002	-	1.0 5.0	NE 5
Phenol	μg/L μg/L	13	7.9	0.1 200000	70000	-	8	300,000.0	-	560,000			1.0E-01 4.2E-01	mg/kg mg/kg	592 28.8	-	3.7E+02	No No	70,000	-	1.0	70,000
Non-Carcinogenic Polycyclic Aromat		hons (PAHs)	1	200000	10000			300,000.0		300,000		<u> </u>	7.2L-UI	ilig/ Ng	20.0		J.7 L±UZ	INU	7 0,000		±.∪	10,000
1-Methylnaphthalene	µg/L									l			_	Ι _		_	_	No	NE	_	0.01	NE
2-Methylnaphthalene	μg/L μg/L		-				-						0.67	mg/kg	_			Yes	NE NE	-	0.01	NE NE
Acenaphthene	μg/L μg/L		_	110	30	_		90		640		_	0.67	mg/kg	4,898		5.3E+00	Yes	5.3		0.01	5.3
Acenaphthylene	μg/L μg/L		-	-	-			-					1.3	mg/kg	4,090		3.3L100	Yes	NE		0.01	NE
Anthracene	μg/L μg/L			4,600	100			400		26,000	_	_	0.96	mg/kg	23,493		2.1E+00	Yes	2.1	-	0.01	2.1
Benzo[g,h,i]perylene	μg/L μg/L		_	4,600	-			400					0.96	mg/kg	20,430		2.1L+00 	Yes	NE		0.01	NE
Fluoranthene	μg/L μg/L			16	6			20		90			2.1	mg/kg	49,096		2.2E+00	Yes	2.2		0.01	2.2
Fluorene	μg/L μg/L			610	10			70		3,500	-		0.54	mg/kg	7,707		3.7E+00	Yes	3.7		0.01	3.7
i luorette	μg/ L			010	10			10		5,500			0.54	ilig/ Ng	1,101		J.7 L 100	162	3.1	_	0.01	3.1



					Criteria for of Surfac							r Protection r Intrusion		C	Criteria for Pro							
			ater Quality apter 173-2		Federal Marine Water Quality Criteria for Washington ² (40 CFR 131.45)	Feder	al Water Qu (CWA §30	ality Criteria ³ 14(a))	MTCA M Surface Cleanu	Water	MTCA I	Method C		t Cleanup	Equilii Parti			Groundwater		Adjust Fact		Proposed Groundwater
			n of Marine	Protection of			of Marine	Protection of	•	l Formula		ndwater	1	ve (SCO)	Coeffic		Groundwater	to Sediment			Practical	Cleanup Level ¹⁰
		Aqua	tic Life	Human Health	Protection of	Aqua	tic Life	Human Health	Val	ue)	Screen	ing Level	Screeni	ng Level ⁴	(L/		Concentration	Pathway	Proposed		Quantitation	(After Adjustment
k _{oc} = Soil organic carbon-water partitioning	Units	Acute	Chronic	(Organisms Only)	Human Health (Organisms Only)	Acute	Chronic	(Organisms Only)	Carc.	Non- Carc.	Carc.	Non- Carc.	C _{sed}	Units	(CLARC)	K _d (metals)	Protective of Sediment ⁶	Complete ⁷ (Yes/No)	Groundwater Cleanup Level	Background Concentration ⁸	Limit9 (PQL)	for Background and PQL)
Naphthalene	μg/L	-	-	-		-	-		-	4,900	89	360	2.1	mg/kg	1,191		9.0E+01	Yes	89		0.01	89
Phenanthrene	µg/L	-	-	-	-	-	-	-	-	-	-	-	1.5	mg/kg	-	-		Yes	NE		0.01	NE
Pyrene	μg/L	-	-	460	8	-		30	-	2,600	-		2.6	mg/kg	67,992		2.0E+00	Yes	2		0.01	2
Carcinogenic Polycyclic Aromatic Hyd		(cPAHs)	Ī	T	1	T	T	T	Ī										<u></u>			
Benzo[a]anthracene	μg/L			0.021	0.00016	-	-	0.0013		-	-		1.3	mg/kg	357,537		1.9E-01	Yes	0.00016		0.01	0.01
Benzo[a]pyrene	μg/L	-	-	0.0021	0.000016	-	-	0.00013	0.04	26	-	-	1.2	mg/kg	968,774	-	6.5E-02	Yes	0.000016		0.01	0.01
Benzo[b]fluoranthene	µg/L	-	-	0.021	0.00016	-	-	0.0013	-		-	-	3.2	mg/kg	1,230,000	_	1.4E-01	Yes	0.00016		0.01	0.01
Benzo[k]fluoranthene	µg/L		-	0.21	0.0016	-	-	0.013			-		3.2	mg/kg	1,230,000	-	1.4E-01	Yes	0.0016		0.01	0.01
Chrysene	µg/L		-	2.1	0.016	-	-	0.13			-		1.4	mg/kg	398,000		1.9E-01	Yes	0.016		0.01	0.016
Dibenz[a,h]anthracene Indeno[1,2,3-c,d]pyrene	µg/L		-	0.0021 0.021	0.000016 0.00016	-	-	0.00013 0.0013	-		-		0.012	mg/kg	1,789,101 3,470,000	*	3.5E-04 5.2E-04	Yes Yes	0.000016 0.00016		0.01 0.01	0.01 0.01
cPAHs TEQ (ND = 0.5RL)	µg/L		_	0.021	0.00016		-	0.0013	0.04	26		_	0.034	mg/kg	968,774		1.1E-03	Yes	0.00016		0.01	0.01
Pesticides and Herbicides	μg/L	-	-	0.0021	0.000016	_	_	0.00013	0.04	26	-	-	0.021	mg/kg	968,774	-	1.11-03	res	0.000016		0.01	0.01
4,4'-DDD	μg/L			0.000036	0.0000079			0.00012	0.0005	0.0015					45,800		l	No	0.0000079		0.10	0.1
4,4'-DDE		-	-	0.000051	0.0000079	-		0.00012	0.0003	0.0015	_	-	-		86,405			No	0.0000079		0.10	0.1
4,4'-DDT	μg/L μg/L	0.13	0.001	0.000031	0.0000008	0.13	0.001	0.000018	0.00036	0.013	-	_	-	_	677,934			No	0.00000088		0.10	0.1
Aldrin		0.13	0.001	0.000025	0.0000012	1.3	0.001	0.00003	0.00038	0.024	3		-	-	48,685			No	0.0000012		0.10	0.05
Alpha-BHC	µg/L	0.71	0.0019	0.000056	0.0000041	1.5		0.0000077	0.000082	160		-	_		1,762			No	0.0000041		0.05	0.05
Alpha-Chlordane (cis)	µg/L			0.00036	0.00048			0.00039		100	-	-	-	-	1,702			No	0.000048 NE		0.05	NE
Beta-BHC	μg/L μg/L		_	0.002	0.0014		_	0.014	0.028	-	-			_	2,139			No	0.0014		0.05	0.05
Delta-BHC	μg/L μg/L			0.002	0.0014			0.014	0.028		-	-			2,139			No	0.0014 NE		0.05	NE
Dieldrin	μg/L	0.71	0.0019	0.0000061	0.0000007	0.71	0.0019	0.0000012	0.000087	0.028					25,546	_		No	0.00000007		0.10	0.1
Endosulfan I	μg/L	0.71	-	10	7	0.034	0.0013	30	-	0.020					25,540			No	0.0087		0.10	0.05
Endosulfan II	μg/L											_						No	NE		0.10	NE
Endosulfan Sulfate	μg/L			10				40		_				_				No	10		0.10	10
Endrin	μg/L	0.037	0.0023	0.035	0.002	0.037	0.0023	0.03		0.20				_	10,811			No	0.002		0.10	0.1
Endrin Aldehyde	μg/L	-	-	0.035		-	- 0.0020	1		-				_	-			No	0.035		0.10	0.1
Endrin Ketone	μg/L	_				_			-					_				No	NE		0.10	NE
Gamma-Chlordane	µg/L	_		_				(_							No	NE		0.05	NE NE
Heptachlor	μg/L	0.053	0.0036	0.00001	0.0000034	0.053	0.0036	0.0000059	0.00013	0.12	_	_		_	9,528			No	0.00000034	_	0.05	0.05
Heptachlor Epoxide	µg/L		-	0.0000074	0.0000024	0.053	0.0036		0.000064		_	_			83,200		_	No	0.0000024		0.05	0.05
Lindane (Gamma-BHC)	µg/L	0.16	-	17	0.43	0.16	-	4.4	0.045	6.0	-	_	_	_	1,352	_		No	0.045		0.05	0.05
Herbicides		-	•	•									•				•	•	•			
2,4,5-T	μg/L	-			- /	-	-	-	-	-	_	-	_	_	-	_		No	NE		0.25	NE
2,4-D	μg/L				-			12,000	-	-	-		_	-	-			No	12,000		1.00	12,000
2,4-DB	μg/L				-	-		-	7-	-					_			No	NE		5.00	NE
Dalapon (DPA)	μg/L				-			-	_	-	-		_	-	-			No	NE		1.00	NE
Dicamba	μg/L				-		-	-	-	_	-		-	-	-		-	No	NE	-	0.50	NE
Dichlorprop	µg/L					-		-	-	-	-			-			-	No	NE	-	1.00	NE
Dinoseb	µg/L			-	-	-			-	-	-	-	-	-	-	-	-	No	NE	-	0.25	NE
МСРА	μg/L	-	-			-	-		-	-	-	-	-	-		-		No	NE		250	NE
Mecoprop (MCPP)	μg/L	-	-			-			-	-	-	-	-		-			No	NE		250	NE
Silvex (Fenoprop or 2,4,5-TP)	µg/L					-		400			-	_	-	-	_			No	400		0.25	400
Polychlorinated Biphenyls (PCBs)																						
Total PCBs (Sum of	μg/L	10	0.03	0.00017	0.000007	_	0.03	0.000064	0.00010				0.0035	mg/kg	309,000		6.0E-04	Yes	0.000007		0.01	0.01
Aroclors)	μg/ L	10	0.03	0.00011	0.00001		0.03	0.00004	0.00010	_			0.0000	1116/ Ng	303,000		0.01-04	163	0.000007		0.01	0.01
Dioxins and Furans	1			1	ī	1	1	1		1		1		_	1	1	ī	Ţ	I			
Total Dioxins/Furans - Human Health TEQ	pg/L			0.064	0.014			0.0051	0.010	0.36	-		5	ng/kg				Yes	0.0051		5	5



Notes:

¹ Water quality criteria for protection of aquatic life from WAC 173-201A-240 (Water Quality Standards for Surface Waters of the State of Washington).

² EPA's Final Revision of Federal Human Health Criteria Applicable to Washington from 40 CFR 131.45; effective date of December 28, 2016.

³ National Recommended Water Quality Criteria (http://water.epa.gov/scitech/swguidance/standards/criteria/current/index.cfm; accessed February 2019).

⁴ Proposed sediment cleanup objective (SCO) values are the lowest sediment risk-based concentration protective of benthic organisms, human health and higher trophic level receptors adjusted for natural background, if available, and PQL (see Tables 1 and 2).

 5 Values for K_{oc} and K_{d} are from Ecology's "CLARC Master Spreadsheet.xlsx" dated May 2019.

⁶ Proposed groundwater cleanup levels protective of sediment were calculated for analytes that were detected in sediment at concentrations greater than their respective proposed SCOs. See text for equation and assumptions used.

T Groundwater to sediment pathway is not complete if analyte was not detected in sediment at a concentration greater that its corresponding proposed sediment cleanup level (Tables 1 and 2). Groundwater to Sediment pathway evaluation is presented in Tables 8 and 9.

 8 PTI, 1989. Background Concentrations of Selected Chemicals in Water, Soil, Sediments, and Air of Washington State.

9 PQL is from the Remedial Investigation/Feasibility Study Work Plan (GeoEngineers 2008) and is the lowest available value from Analytical Resources Inc. (ARI) of Tukwila, Washington.

¹⁰ Screening level is based on lowest of Federal and State marine surface water concentrations protective of aquatic life and human health from consumption of aquatic life including MTCA Method B standard formula values for carcinogens and non-carcinogen, and adjusted for background and the practical quantification limit (PQL) for all analytes with available surface water criteria.

11 State Surface Water Quality Criteria, National Toxic Rule and Clean Water Act values are based on hexavalent chromium; trivalent chromium values are not available. MTCA Method B Surface Water Cleanup Levels are based on trivalent chromium.

 12 Clean Water Act [CWA 304(a)] for Protection of Human Health value for mercury is based on methylmercury.

12 MTCA Method A groundwater cleanup level. Value for gasoline-range petroleum hydrocarbons if benzene is present. If benzene is not present, screening level is 1,000 µg/L.

C_{sed} = Sediment cleanup level

C_w = Groundwater/Surface water screening level

f_{oc} = Sediment fraction of organic carbon

k_d = Distribution coefficient

k_{oc} = Soil organic carbon-water partitioning coefficient

θw = Water-filled porosity

pb = Dry sediment bulk density

MTCA = Model Toxics Control Act

μg/L = Microgram per liter

mg/kg = Milligrams per kilogram

ng/L = Nanogram per liter

NE = Not established

ND = Non-detect

RL = Reporting limit

TEQ = Toxic equivalent concentration

- = No screening criteria available.

Blue shading identifies the basis for proposed groundwater cleanup level.

Green shading identifies the proposed groundwater cleanup level after adjustment for background and the PQL.



Table 4

Proposed Soil Cleanup Levels

Dakota Creek Industries Anacortes, Washington

		Protection of Human	Human H	ealth Direct				Soil to Groundy	vater Protection	<u></u>						
		Health and the Environment ¹ (MTCA Method A Standard Table Value	Contact (MTCA Meth Formula Valu	Pathway od C Standard e for Industrial	-	ilibrium Part Coefficients (L/kg)		Using Groundv Screening Le per WAC 173-	vater Proposed evel (Table 3) 340-740(1)(d) -1/747-2	Soil to Groundwater Pathway	-	osed ng Level	Adjustm Factor		Cleanu	sed Soil p Level ⁶ ustment for nd and PQL)
		for Industrial Land	Carainagan	Non-	K _{oc}	K _d	H	Vadose	Saturated	Complete ³	Vadose	Saturated	Background	PQL ⁵	Vadose Zone	Saturated
Analyte	Units	Use)	Carcinogen	Carcinogen	(CLARC)	(metals)	(Unitless)	Zone	Zone	(Yes/No)	Zone	Zone	Concentration ⁴	PŲL	Zone	Zone
Metals	1		00	1 100		0.05.04	0.05.00	0.057	0.0000		0.057	0.0000	207	-	00	00
Arsenic	mg/kg		88	1,100		2.9E+01	0.0E+00	0.057	0.0029	Yes	0.057	0.0029	20 ⁷	5	20	20
Cadmium Total Chromium ⁸	mg/kg			3,500		1.0E+03	0.0E+00 0.0E+00	1 000	50	No	3,500	3,500.000 5,300,000	1 48	0.2	3,500	3,500
	mg/kg	-		5,300,000 140,000		2.2E+01	0.0E+00 0.0E+00	1,000 1.4	0.069	No	5,300,000 140,000	140,000	36	0.5 0.2	5,300,000 140,000	5,300,000 140,000
Copper	mg/kg	1 000		140,000				420	21	No		· · · · · · · · · · · · · · · · · · ·	24	2	,	
Lead	mg/kg	1,000		1.100 ⁹		1.0E+04 5.2E+01	0.0E+00 4.7E-01	0.026	0.0013	No No	1,000 1,100	1,000 1,100	0.07	0.05	1,000 1,100	1,000 1,100
Mercury	mg/kg			,	-				0.53		,	,	48		48	
Nickel	mg/kg		-	70,000		6.5E+01	0.0E+00	0.22		Yes	11	0.53		1 0.3		48 18,000
Silver Zinc	mg/kg	-		18,000 1,100,000		8.3E+00 6.2E+01	0.0E+00 0.0E+00	0.32 101	0.016 5.0	No No	18,000 1,100,000	18,000 1,100,000	 85	0.3	18,000 1,100,000	1,100,000
Petroleum Hydrocarbons	mg/kg	-		1,100,000	-	6.2E+01	0.0E+00	101	5.0	INO	1,100,000	1,100,000	85	1	1,100,000	1,100,000
Gasoline-Range	ma/ka	100 ¹⁰		_		Ī				No	100	100		5	100	100
Diesel-Range	mg/kg	2.000						_	-	No	2,000	2,000		5	2,000	2,000
Heavy Oil-Range	mg/kg mg/kg	2,000			-					No	2,000	2,000		10	2,000	2,000
BETX Compounds	IIIg/ kg	2,000								NO	2,000	2,000		10	2,000	2,000
Benzene	mg/kg		2,400	14,000	6.2E+01	Ī	1.3E-01	2.0	0.10	No	2,400	2.400		0.001	2,400	2.400
Ethylbenzene	mg/kg			350,000	2.0E+02		1.6E-01	127	6	No	350,000	350,000		0.001	350.000	350.000
Toluene	mg/kg	_		280,000	1.4E+02		1.5E-01	365	18	No	280,000	280,000		0.001	280,000	280,000
Xylenes	mg/kg			700,000	2.3E+02		1.4E-01	2,938	147	No	700,000	700,000		0.001	700,000	700.000
Volatile Organic Compounds (VOCs)	1118/118			700,000	2.5L102		1.72-01	2,338	177	110	700,000	700,000		0.001	700,000	100,000
1,1,1,2-Tetrachloroethane	mg/kg	_	5,000	110,000			4.6E-02		_	No	5,000	5.000		0.001	5,000	5,000
1.1.1-Trichloroethane	mg/kg			7,000,000	1.4E+02	-	4.2E-01	32,457	1,623	No	7,000,000	7,000,000		0.001	7,000,000	7,000,000
1,1,2,2-Tetrachloroethane	mg/kg		660	70.000	7.9E+01		7.0E-03	0.48	0.024	No	660	660		0.002	660	660
1,1,2-trichloro-1,2,2- trifluoroethane (CFC113)	mg/kg	-	-	110,000,000	-	-	1.3E+01	-		No	110,000,000	110,000,000		0.002	110,000,000	110,000,000
1,1,2-Trichloroethane	mg/kg	-	2,300	14,000	7.5E+01	-	2.0E-02	1.4	0.068	No	2,300	2,300		0.001	2,300	2,300
1,1-Dichloroethane	mg/kg	-	23,000	700,000	5.3E+01	-	1.4E-01	117	5.9	No	23,000	23,000		0.001	23,000	23,000
1,1-Dichloroethene	mg/kg	-	/-	180,000	6.5E+01		7.1E-01	365	18	No	180,000	180,000		0.001	180,000	180,000
1,1-Dichloropropene	mg/kg	-	_	-						No	NE	NE		0.001	NE	NE
1,2,3-Trichlorobenzene	mg/kg	-	-	_					-	No	NE	NE		0.005	NE	NE
1,2,3-Trichloropropane	mg/kg	-	4.4	14,000	-		7.9E-03		-	No	4.4	4.4		0.002	4.4	4.4
1,2,4-Trichlorobenzene	mg/kg	-	4,500	35,000	1.7E+03		2.4E-02	1.2	0.061	No	4,500	4,500		0.005	4,500	4,500
1,2,4-Trimethylbenzene	mg/kg	-	-	35,000	7-		1.1E-01			No	35,000	35,000		0.001	35,000	35,000
1,2-Dibromo-3-chloropropane	mg/kg		160	700	-					No	160	160		0.005	160	160
1,2-Dibromoethane (EDB)	mg/kg		66	32,000	6.6E+01		1.5E-02	3.6	0.18	No	66	66		0.001	66	66
1,2-Dichlorobenzene	mg/kg	-	-	320,000	3.8E+02		3.5E-02	6,067	303	No	320,000	320,000		0.001	320,000	320,000
1,2-Dichloroethane (EDC)	mg/kg	-	1,400	21,000	3.8E+01		2.3E-02	32	1.6	No	1,400	1,400		0.001	1,400	1,400
1,2-Dichloropropane	mg/kg	-	3,500	140,000	4.7E+01		6.5E-02	2.9	0.15	No	3,500	3,500		0.001	3,500	3,500
1,3,5-Trimethylbenzene	mg/kg			35,000			1.1E-01		_	No	35,000	35,000		0.001	35,000	35,000



		Protection of Human Health and the Environment ¹ (MTCA Method A Standard Table Value	Contact (MTCA Meth Formula Valu	ealth Direct t Pathway od C Standard ee for Industrial d Use)		ilibrium Part Coefficients (L/kg)	_	Using Ground Screening L per WAC 173	vater Protection water Proposed evel (Table 3) -340-740(1)(d) -1/747-2	Soil to Groundwater Pathway	Prop Screenii	osed ng Level	Adjustm Factor		Propos Cleanu _l (After adju backgroun	stment for
Analista	Heito	for Industrial Land	Carcinogen	Non- Carcinogen	K _{oc} (CLARC)	K _d (metals)	H (Unitless)	Vadose Zone	Saturated Zone	Complete ³	Vadose Zone	Saturated Zone	Background Concentration ⁴	PQL ⁵	Vadose Zone	Saturated Zone
Analyte 1,3-Dichlorobenzene	Units	Use) 			(OEAI(O)		6.0E-02			(Yes/No)	NE	NE		0.001	NE NE	NE NE
1,3-Dichloropropane	mg/kg mg/kg						0.0E-02			No	NE NE	NE NE		0.001	NE NE	NE NE
1,4-Dichlorobenzene	mg/kg		24,000	250,000	6.2E+02		4.6E-02	46.831.504	2,341,575	No	24,000	24,000		0.067	24,000	24,000
2,2-Dichloropropane	mg/kg		24,000		0.21.02		4.0L-02		2,541,575	No	NE	NE		0.001	NE	NE
2-Butanone (MEK)	mg/kg			2,100,000	_		1.3E-03		_	No	2,100,000	2,100,000		0.005	2,100,000	2,100,000
2-Chloroethyl Vinyl Ether	mg/kg						-			No	NE	NE		0.005	NE	NE
2-Chlorotoluene	mg/kg			70,000					-	No	70.000	70.000		0.001	70,000	70,000
2-Hexanone	mg/kg			18,000					-	No	18.000	18.000		0.001	18,000	18,000
4-Chlorotoluene	mg/kg								_	No	NE.	NE		0.001	NE	NE
4-Methyl-2-Pentanone (Methyl Isobutyl Ketone)	mg/kg			280,000		-	2.9E-03	-		No	280,000	280,000		0.005	280,000	280,000
Acetone	mg/kg	-		3,200,000	5.8E-01		9.7E-04	496,054	24,803	No	3,200,000	3,200,000		0.005	3,200,000	3,200,000
Acrolein	mg/kg				_				-	No	NE	NE		0.05	NE	NE
Acrylonitrile	mg/kg	-	240	140,000	-		2.3E-03	-	-	No	240	240		0.005	240	240
Bromobenzene	mg/kg	-		2.80E+04	2.3E+02		4.3E-02		-	No	28,000	28,000		0.001	28,000	28,000
Bromochloromethane	mg/kg				_			-	-	No	NE	NE		0.005	NE	NE
Bromoform	mg/kg	-	17,000	70,000	1.3E+02		1.2E-02	30	1.5	No	17,000	17,000		0.001	17,000	17,000
Bromomethane	mg/kg	-		4,900	9.0E+00		1.8E-01	5.2	0.26	No	4,900	4,900		0.001	4,900	4,900
Carbon Disulfide	mg/kg	-		350,000	4.6E+01		8.0E-01	800	40	No	350,000	350,000	-	0.001	350,000	350,000
Carbon Tetrachloride	mg/kg	-	1,900	14,000	1.5E+02		7.4E-01	1.1	0.053	No	1,900	1,900		0.001	1,900	1,900
Chlorobenzene	mg/kg			70,000	2.2E+02		7.9E-02	897	45	No	70,000	70,000		0.001	70,000	70,000
Chloroethane	mg/kg	-			-	-	2.5E-01	-		No	NE	NE		0.005	NE	NE
Chloroform	mg/kg	-	4,200	35,000	5.3E+01		9.2E-02	13	0.63	No	4,200	4,200		0.001	4,200	4,200
Chloromethane	mg/kg	-			6.0E+00		2.7E-01	42	2.1	No	NE	NE		0.001	NE	NE
Cis-1,2-Dichloroethene	mg/kg	-		7,000	3.6E+01		1.0E-01	-	-	No	7,000	7,000		0.001	7,000	7,000
Cis-1,3-Dichloropropene	mg/kg	-		-	_	4 /		-	-	No	NE	NE		0.001	NE	NE
Dibromochloromethane	mg/kg	-	1,600	70,000	6.3E+01		2.1E-02	2.8	0.14	No	1,600	1,600		0.001	1,600	1,600
Dibromomethane	mg/kg	-		35,000	-		2.0E-02		-	No	35,000	35,000		0.001	35,000	35,000
Dichlorobromomethane	mg/kg	-	2,100	70,000	5.5E+01		3.7E-02	3.1	0.15	No	2,100	2,100		0.001	2,100	2,100
Dichlorodifluoromethane (CFC 12)	mg/kg		-	700,000	-		8.1E+00			No	700,000	700,000		0.001	700,000	700,000
Hexachlorobutadiene	mg/kg	-	1,700	3,500	5.4E+04	-	1.4E-01	11	0.54	No	1,700	1,700		0.005	1,700	1,700
Isopropylbenzene	mg/kg	-	-	350,000	-		2.6E-01		-	No	350,000	350,000		0.001	350,000	350,000
Methyl lodide	mg/kg		70.000		4.45	\	4.05.55			No	NE 70.000	NE 70.000		0.001	NE 70.000	NE
Methyl t-Butyl Ether (MTBE)	mg/kg		73,000		1.1E+01		1.6E-02	1,354	68	No	73,000	73,000		0.001	73,000	73,000
Methylene Chloride	mg/kg		66,000	21,000	1.0E+01		5.7E-02	20	1.020	No	21,000	21,000		0.002	21,000	21,000
Naphthalene	mg/kg		-	70,000	1.2E+03		8.2E-03	2,125	106	No	70,000	70,000		0.005	70,000	70,000
n-Butylbenzene	mg/kg			180,000			2.4E-01			No	180,000	180,000		0.001	180,000	180,000
n-Propylbenzene	mg/kg			350,000	-		2.0E-01		-	No	350,000	350,000		0.001	350,000	350,000
p-Isopropyltoluene	mg/kg			350,000	-		0.05.04			No	NE 350,000	NE		0.001	NE 350,000	NE 350,000
sec-Butylbenzene	mg/kg		-	350,000	0.45+00		2.6E-01		16.400	No	350,000	350,000		0.001	350,000	350,000
Styrene	mg/kg			700,000	9.1E+02		5.6E-02	328,394	16,420	No	700,000	700,000		0.001	700,000	700,000
tert-Butylbenzene	mg/kg			350,000	0.75+00		2.6E-01			No	350,000	350,000		0.001	350,000	350,000
Tetrachloroethene (PCE)	mg/kg		63,000	21,000	2.7E+02		4.0E-01	15	0.77	No	21,000	21,000		0.001	21,000	21,000



		Protection of Human Health and the Environment ¹ (MTCA Method A Standard Table Value	Contact (MTCA Methor Formula Valu	ealth Direct : Pathway od C Standard e for Industrial d Use)		ilibrium Part Coefficients (L/kg)	_	Using Groundv Screening Lo per WAC 173	vater Protection vater Proposed evel (Table 3) -340-740(1)(d) -1/747-2	Soil to Groundwater Pathway	Prop Screenii		Adjustm Factoi		Propos Cleanup (After adju backgroun	stment for
		for Industrial Land	Oavelneden	Non-	K _{oc}	K _d	H	Vadose	Saturated	Complete ³	Vadose	Saturated	Background	PQL ⁵	Vadose	Saturated
Analyte	Units	Use)	Carcinogen	Carcinogen	(CLARC)	(metals)	(Unitless)	Zone	Zone	(Yes/No)	Zone	Zone	Concentration*	•	Zone	Zone
Trans-1,2-Dichloroethene	mg/kg	-		70,000	3.8E+01		2.4E-01	191	9.6	No	70,000	70,000		0.001	70,000	70,000
Trans-1,3-Dichloropropene	mg/kg				-					No	NE	NE		0.001	NE	NE
Trans-1,4-Dichloro-2-butene	mg/kg	-	-		-		-			No	NE	NE		0.005	NE 1 aaa	NE
Trichloroethene (TCE)	mg/kg		2,800	1,800	9.4E+01	-	2.4E-01	1.3	0.066	No	1,800	1,800	-	0.001	1,800	1,800
Trichlorofluoromethane (CFC 11)	mg/kg	-	-	1,100,000			2.7E+00		-	No	1,100,000	1,100,000		0.001	1,100,000	1,100,000
Vinyl Acetate	mg/kg	-		3,500,000	5.3E+00		1.2E-02	1,853	93	No	3,500,000	3,500,000		0.005	3,500,000	3,500,000
Vinyl Chloride	mg/kg	-	88	11,000	1.9E+01		8.1E-01	0.068	0.0034	No	88	88		0.001	88	88
Semi-Volatile Organic Compounds (SVO	1	I	4.500	25.000	4.75.00	I	0.45.00		2 224		4.500	4.500	1	0.007	4.500	4.500
1,2,4-Trichlorobenzene	mg/kg		4,500	35,000	1.7E+03		2.4E-02	1.2	0.061	No	4,500	4,500		0.067	4,500	4,500
1,2-Dichlorobenzene	mg/kg			320,000	3.8E+02		3.5E-02	6,067	303	No	320,000	320,000	-	0.067	320,000	320,000
1,3-Dichlorobenzene	mg/kg		- 04.000				6.0E-02	- 074	-	No	NE 04.000	NE 04.000		0.067	NE 04.000	NE 04.000
1,4-Dichlorobenzene	mg/kg		24,000	250,000	6.2E+02		4.6E-02	271	14	No	24,000	24,000	-	0.067	24,000	24,000
2,2'-0xybis[1-chloropropane]	mg/kg			250,000	4.05.00				-	No	NE 250,000	NE 250,000		0.067	NE 250,000	NE 250,000
2,4,5-Trichlorophenol	mg/kg		- 40.000	350,000	1.6E+03		6.5E-05	19,166	958	No	350,000	350,000		0.33	350,000	350,000
2,4,6-Trichlorophenol	mg/kg	-	12,000	3,500	3.8E+02		1.1E-04	2.1	0.11	No	3,500	3,500	-	0.33	3,500	3,500
2,4-Dichlorophenol	mg/kg	-		11,000	1.5E+02		3.4E-05	29	1.5	No	11,000	11,000		0.33	11,000	11,000
2,4-Dimethylphenol	mg/kg	-	-	70,000	2.1E+02		3.0E-05	406	20.3	No	70,000	70,000	-	0.067	70,000	70,000
2,4-Dinitrophenol	mg/kg	-		7,000	1.0E-02		1.5E-06	0.42	0.021	No	7,000	7,000		0.67	7,000	7,000
2,4-Dinitrotoluene	mg/kg	-	420	7,000	9.6E+01		1.0E-06	0.34	0.017	No	420	420		0.33	420	420
2,6-Dinitrotoluene	mg/kg		88	1,100	6.9E+01		8.8E-06		-	No	88	88		0.33	88	88
2-Chloronaphthalene	mg/kg	-	-	280,000		-		100	-	No	280,000	280,000		0.067	280,000	280,000
2-Chlorophenol	mg/kg	-		18,000	3.9E+02	-	7.3E-03	132	6.6	No	18,000	18,000	-	0.067	18,000	18,000
2-Nitroaniline	mg/kg			35,000	-				-	No	35,000	35,000		0.33	35,000	35,000
2-Nitrophenol	mg/kg			-	7.05+00				- 0.0004	No	NE	NE		0.33	NE 200	NE 200
3,3'-Dichlorobenzidine	mg/kg		290		7.2E+02		2.2E-08	0.048	0.0024	No	290	290		0.33	290	290
3-Nitroaniline	mg/kg				_		_			No	NE	NE		0.33	NE 000	NE 200
4,6-Dinitro-2-methylphenol	mg/kg	-		280	-			-		No	280 NE	280		0.67 0.067	280	280 NE
4-Bromophenyl-phenylether 4-Chloro-3-Methylphenol	mg/kg	_	-	350,000		-	-	-		No No	NE	NE	_	0.067	NE	NE 350,000
4-Chloroaniline	mg/kg		660	14,000	6.6E+01	_	4.9E-06			No	350,000 660	350,000 660	_	0.33	350,000 660	350,000 660
4-Chlorophenyl-phenylether	mg/kg		000		0.02+01		4.91-00			No	NE	NE		0.067	NE	NE
, ,, ,	mg/kg				-	-								0.067		
4-Nitrophonal	mg/kg			14,000	-	-				No No	14,000 NE	14,000 NE		0.33	14,000 NE	14,000 NE
4-Nitrophenol	mg/kg	-	_	14,000,000	C OF O1	_	4.05.06									
Benzoic acid	mg/kg			14,000,000 350,000	6.0E-01		4.9E-06		-	No No	14,000,000 350,000	14,000,000 350,000		0.67	14,000,000 350,000	14,000,000 350,000
Benzyl alcohol	mg/kg						-				i i	· ·			·	· · · · · · · · · · · · · · · · · · ·
bis(2-Chloroethoxy)methane	mg/kg		120	-	7.65104		2.05.04	0.001		No	NE 120	NE 120		0.067	NE 130	NE 120
bis(2-chloroethyl)ether	mg/kg		120	70,000	7.6E+01 1.1E+05		2.9E-04	0.091	0.0046 5.1	No	120	120 9,400		0.067 0.067	120	120
bis(2-Ethylhexyl)phthalate	mg/kg		9,400				6.6E-07	102		No	9,400	,			9,400	9,400
Butylbenzylphthalate	mg/kg		69,000	700,000	1.4E+04		1.1E-05	3.6	0.18	No	69,000	69,000		0.067	69,000	69,000
Carbazole	mg/kg			2.500	3.4E+03		1.6E-07			No	NE 2.500	NE 2.500		0.067	NE 2.500	NE 2.500
Dibenzofuran Diethylahtholete	mg/kg			3,500	0.05+04		1.7E-04			No	3,500	3,500		0.067	3,500	3,500
Diethylphthalate	mg/kg			2,800,000	8.2E+01		4.7E-06	329	16	No	2,800,000	2,800,000	-	0.067	2,800,000	2,800,000
Dimethylphthalate	mg/kg		-							No	NE	NE		0.067	NE	NE



		Protection of Human Health and the Environment ¹ (MTCA Method A Standard Table Value	Contact (MTCA Meth Formula Valu	ealth Direct : Pathway od C Standard e for Industrial d Use)	_	ilibrium Part Coefficients (L/kg)	_	Soil to Groundy Using Groundy Screening Lo per WAC 173- EQ. 747-	evel (Table 3) 340-740(1)(d)	Soil to Groundwater Pathway	(For Nonlo	nic & Ionizii	, Adjustm Factoi		Cleanu	sed Soil p Level ⁶ ustment for nd and PQL)
		for Industrial Land		Non-	K _{oc}	K _d	Н	Vadose	Saturated	Complete ³	Vadose	Saturated	Background	5 5	Vadose	Saturated
Analyte	Units	Use)	Carcinogen	Carcinogen	(CLARC)	(metals)	(Unitless)	Zone	Zone	(Yes/No)	Zone	Zone	Concentration ⁴	PQL ⁵	Zone	Zone
Di-n-butylphthalate	mg/kg	-		350,000	1.6E+03		8.2E-09	251	13	No	350,000	350,000		0.067	350,000	350,000
Di-n-octylphthalate	mg/kg	-		35,000	8.3E+07		5.2E-04			No	35,000	35,000		0.067	35,000	35,000
Hexachlorobenzene	mg/kg	-	82	2,800	8.0E+04	-	1.4E-02	0.0080	0.00040	No	82	82		0.067	82	82
Hexachlorobutadiene	mg/kg	-	1,700	3,500	5.4E+04	-	1.4E-01	11	0.54	No	1,700	1,700		0.067	1,700	1,700
Hexachlorocyclopentadiene	mg/kg	-		21,000	2.0E+05		4.2E-01	4,000	200	No	21,000	21,000		0.067	21,000	21,000
Hexachloroethane	mg/kg	-	3,300	2,500	1.8E+03		7.2E-02	0.71	0.04	No	2,500	2,500		0.067	2,500	2,500
Isophorone	mg/kg	-	140,000	700,000	4.7E+01		1.1E-04	103	5.2	No	140,000	140,000		0.067	140,000	140,000
Nitrobenzene	mg/kg			7,000	1.2E+02		4.0E-04	238	12	No	7,000	7,000		0.067	7,000	7,000
n-Nitroso-di-n-propylamine	mg/kg	-	19		2.4E+01		5.4E-05	0.028	0.0014	No	19	19		0.067	19	19
n-Nitrosodiphenylamine	mg/kg		27,000		1.3E+03		1.0E-04	18	0.89	No	27,000	27,000		0.067	27,000	27,000
o-Cresol (2-Methylphenol)	mg/kg			180,000	9.1E+01		2.0E-05		-	No	180,000	180,000		0.067	180,000	180,000
p-Cresol (4-Methylphenol)	mg/kg	-		350,000				-		No	350,000	350,000		0.067	350,000	350,000
Pentachlorophenol	mg/kg		330	18,000	5.9E+02		2.1E-07	0.024	1.2E-03	No	330	330		0.17	330	330
Phenol	mg/kg	-		1,100,000	2.9E+01		6.5E-06	40,600	2,030	No	1,100,000	1,100,000		0.033	1,100,000	1,100,000
Non-carcinogenic Polycyclic Aromat	tic Hydrocarbons	(PAHs)			ı					ī	I	ı	T			
1-Methylnaphthalene	mg/kg		4,500	250,000			1.6E-02	-	-	No	4,500	4,500		0.005	4,500	4,500
2-Methylnaphthalene	mg/kg			14,000			7.0E-03	-	-	No	14,000	14,000		0.005	14,000	14,000
Acenaphthene	mg/kg	-		210,000	4.9E+03		2.1E-03	523	26	No	210,000	210,000		0.005	210,000	210,000
Acenaphthylene	mg/kg		-					-	-	No	NE	NE		0.005	NE	NE
Anthracene	mg/kg			1,100,000	2.3E+04		7.6E-04	1009	50	No	1,100,000	1,100,000		0.005	1,100,000	1,100,000
Benzo[g,h,i]perylene	mg/kg		-		-			-		No	NE	NE		0.005	NE	NE
Fluoranthene	mg/kg			140,000	4.9E+04	-	1.7E-04	2209	110	No	140,000	140,000		0.005	140,000	140,000
Fluorene	mg/kg			140,000	7.7E+03	-	8.6E-04	566	28	No	140,000	140,000		0.005	140,000	140,000
Naphthalene	mg/kg			70,000	1.2E+03	-	8.2E-03	2,125	106	No	70,000	70,000	-	0.005	70,000	70,000
Phenanthrene	mg/kg		-	-	-	/		-	-	No	NE	NE		0.005	NE	NE
Pyrene	mg/kg	-		110,000	6.8E+04		1.1E-04	2,736	137	No	110,000	110,000	_	0.005	110,000	110,000
Carcinogenic Polycyclic Aromatic H	ydrocarbons (cPA	(Hs)									1					
Benzo[a]anthracene	mg/kg		-	-	3.6E+05		2.8E-05	1.1	0.057	Yes	1.1	0.06	-	0.01	1.1	0.06
Benzo[a]pyrene	mg/kg		130	1,100	9.7E+05	-	6.4E-06	0.31	0.016	Yes	0.31	0.016		0.01	0.31	0.016
Benzo[b]fluoranthene	mg/kg	-		-	1.2E+06	-	7.7E-04	3.9	0.20	Yes	3.9	0.2	-	0.01	3.9	0.2
Benzo[k]fluoranthene	mg/kg	-	-	-	1.2E+06	-	5.1E-06	39	1.97	Yes	39	2.0	-	0.01	39	2.0
Chrysene	mg/kg	-		-	4.0E+05		7.1E-04	127	6.4	Yes	127	6.4	-	0.01	127	6.4
Dibenz[a,h]anthracene	mg/kg	_	-	-	1.8E+06	-	6.0E-07	0.57	0.029	Yes	0.57	0.029		0.01	0.57	0.029
Indeno[1,2,3-c,d]pyrene	mg/kg	-	-	-	3.5E+06		8.4E-06	11	0.56	Yes	11	0.56		0.01	11	0.56
cPAHs TEQ (ND = 0.5RL)	mg/kg		130	1,100	9.7E+05	-	6.4E-06	0.31	0.016	Yes	0.31	0.016		0.01	0.31	0.016
Pesticides		,				•						.				
4,4'-DDD	mg/kg	-	550	110	4.6E+04		2.9E-05	0.0072	0.00036	No	110	110		0.003	110	110
4,4'-DDE	mg/kg	-	390	1,100	8.6E+04		1.9E-04	0.0015	0.000076	No	390	390	-	0.003	390	390
4,4'-DDT	mg/kg	4	390	1,800	6.8E+05		3.8E-05	0.016	0.00081	No	4	4		0.003	4	4
Aldrin	mg/kg		7.7	110	4.9E+04	_	1.6E-03	0.00004	0.000002	No	7.7	7.7		0.002	7.7	7.7
Alpha-BHC	mg/kg		21	28,000	1.8E+03	_	1.0E-04	0.0017	0.000085	No	21	21		0.002	21	21
Alpha-Chlordane (cis)	mg/kg		_	-		-	-			No	NE	NE		0.002	NE	NE
Beta-BHC	mg/kg	-	73		2.1E+03		4.8E-06	0.06	0.003	No	73	73		0.002	73	73



		Protection of Human Health and the Environment ¹ (MTCA Method A Standard Table Value	Contact (MTCA Meth Formula Valu	ealth Direct : Pathway od C Standard e for Industrial d Use)	•	ilibrium Part Coefficients (L/kg)		Using Ground Screening L per WAC 173	vater Protection vater Proposed evel (Table 3) :340-740(1)(d) :1/747-2	Soil to Groundwater Pathway	Prop Screenii		Adjustm Factor		Propos Cleanup (After adju backgroun	stment for
		for Industrial Land		Non-	K _{oc}	K _d	Н	Vadose -	Saturated _	Complete ³	Vadose	Saturated	Background	5	Vadose -	Saturated
Analyte	Units	Use)	Carcinogen	Carcinogen	(CLARC)	(metals)	(Unitless)	Zone	Zone	(Yes/No)	Zone	Zone	Concentration ⁴	PQL⁵	Zone	Zone
Delta-BHC	mg/kg	-		-			-		-	No	NE	NE		0.002	NE	NE
Dieldrin	mg/kg	-	8	180	2.6E+04		1.1E-04	0.000036	0.0000018	No	8.2	8.2	-	0.003	8.2	8.2
Endosulfan I	mg/kg			-					-	No	NE	NE		0.002	NE	NE
Endosulfan II	mg/kg				-		-		-	No	NE	NE		0.003	NE	NE
Endosulfan Sulfate	mg/kg	-	-	21,000				-	-	No	21,000	21,000		0.003	21,000	21,000
Endrin	mg/kg	-		1,100	1.1E+04		6.6E-05	0.43	0.022	No	1,100	1,100	-	0.003	1,100	1,100
Endrin Aldehyde	mg/kg	-	-	-					-	No	NE	NE		0.003	NE	NE
Endrin Ketone	mg/kg	-	_	-	-		-		_	No	NE	NE		0.003	NE	NE
Gamma-Chlordane	mg/kg	-	_	_	-		-		-	No	NE	NE		0.002	NE	NE
Heptachlor	mg/kg	-	29	1,800	9.5E+03		1.3E-02	0.000065	0.0000032	No	29	29	-	0.002	29	29
Heptachlor Epoxide	mg/kg	-	14	46	8.3E+04		8.0E-05	0.004	0.0002	No	14	14		0.002	14	14
Lindane (Gamma-BHC)	mg/kg	0.01	120	1,100	1.4E+03		1.3E-04	0.54	0.027	No	0.01	0.01		0.002	0.01	0.01
Methoxychlor	mg/kg	-		18,000	8.0E+04		1.2E-04	0.051	0.0026	No	18,000	18,000	-	0.002	18,000	18,000
Toxaphene	mg/kg	-	120	320	9.6E+04		5.2E-05	86	4.3	No	120	120		0.002	120	120
Polychlorinated Biphenyls (PCBs)	-															
Total PCBs (sum of Aroclors)	mg/kg	10	66.0		3.1E+05		1.2E-02	0.043	0.002	No	10	10		0.05	10	10
Dioxins and Furans																
Total Dioxins/Furans - Human Health TEQ	ng/kg		1,700	4,100					-	No	1,700	1,700	5.2	5	1,700	1,700

MTCA = Washington State Model Toxics Control Act

K_{oc} = Soil organic carbon-water partitioning coefficient (L/kg)

 K_d = Distribution coefficient for metals (L/kg)

H = Henrys Law constant (unitless)

PQL = Practical quantitation limit

mg/kg = Milligrams per kilogram

ng/kg = Nanogram per kilogram

-- = No screening criteria available.

NE = not established

ND = Non-detect

RL = Reporting limit

TEQ = toxic equivalent concentration (toxicity equivalency factor [TEF] values are presented in Table 4).

Blue shading identifies the basis for proposed soil cleanup level.

Green shading identifies the proposed soil cleanup level after adjustment for background and the PQL.

¹The MTCA A screening value is shown for those chemicals for which Method C values are not available (e.g., petroleum hydrocarbons and lead). The MTCA Method A value for total PCBs is also included in the table because it captures the chemical-specific level mandated in the Federal Toxic Substance Control Act.

 $^{^2}$ For ionizing and non-ionizing organics, $K_d = K_{oc} \times f_{oc}$ and uses the MTCA default f_{oc} of 0.1% in upland soil. Values for Kd and/or Koc and/or Henry's Law Constant are from Ecology's "CLARC Master Spreadsheet.xlsx" dated May 2019 where available.

³ Soil to groundwater pathway is not complete if analyte was not detected in groundwater at a concentration greater that its corresponding proposed groundwater cleanup level (Table 3). Soil to groundwater pathway evaluation is presented in Table 9.

⁴ Metals background values (Puget Sound Region 90th percentile values) are from Natural Background Soil Metals Concentrations in Washington State (Ecology Publication #94-115, 1994).

⁵ PQL is from the Remedial Investigation/Feasibility Study Work Plan (GeoEngineers 2008) and is the lowest available value from Analytical Resources Inc. (ARI) of Tukwila, Washington.

⁶ Screening level is based on lowest of soil concentrations protective of human health and the environment (MTCA Method A table value for industrial sites), human health - direct contact (MTCA Method C standard formula values for carcinogens and non-carcinogens), and protection of groundwater, adjusted for background and PQL. Calculated concentrations protective of groundwater as marine surface water assume unsaturated and saturated soil, and are calculated based on proposed groundwater cleanup levels before adjustment for background and PQLs.

 $^{^{7}}$ Background for arsenic as established in the MTCA A Table 745-1 (WAC 173-340-900).

⁸ Based on chromium (III).

⁹ Based on mercuric chloride.

¹⁰ Value for gasoline-range petroleum hydrocarbons if benzene is not present. If benzene is present, screening level is 30 mg/kg.

Schedule of Laboratory Analysis for Sediment Investigations

Dakota Creek Industries Anacortes, Washington

			Conventional Analyses ² Chemical Analyses ² Chemical Analyses ² Sauric Sauric																	
Sample Location ¹	Date Sampled	Sample Interval	Total Organic Carbon (TOC)	ø	Total Solids (TS)	Total Ammonia	Total Sulfides	Grain Size	TBT	Metals	LPAHs	нранs	сРАНѕ	Chlorinated Hydrocarbons	Phthalates	Miscellaneous	Phenois	Pesticides	PCB Aroclors	Dioxins and Furans
Phase 2 Environmental As	ssessment ³ (Ott	en Engineering	1997)																
DC-SED-01 ³	07/03/97	0-10 cm																		
DC-SED-02 ³	07/03/97	0-10 cm						,				•								
DC-SED-03 ³	07/03/97	0-10 cm								•										
DC-SED-05 ³	08/06/97	0-10 cm							•											
DC-SED-06 ³	08/06/97	0-10 cm							7	•										
DC-SED-08 ³	08/06/97	0-10 cm				\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \						•								
DC-SED-09	08/06/97	0-10 cm			1															
Dredged Material Charact	erization (Hart	Crowser 2000)																		
DMMU-D1-Comp-A ³	04/25/00	0-10 cm		•	•		-													
DMMU-D2-Comp-A ³	04/25/00	0-10 cm																		
Supplemental Dredged M	aterial Characte	erization (Ancho	or 2004	4)																
AN-P1-2	07/15/04	1-3 ft		K																
AN-DCI-1A/B ³	07/15/04	1-3 ft			-															
AN-DCI-2 ³	07/15/04	1-3 ft				ı														
DCI Basin Surface Sedime	ent Dioxin Study	(Floyd Snider	2007)					,			T		T			T				
DCI06-1A ³	N/A	0-10 cm	-																	
DCI06-2A	N/A	0-10 cm	•																	
DCI06-2D	N/A	0-10 cm	4																	
DCI06-3A ³	N/A	0-10 cm	-																	
DCI06-4A ³	N/A	0-10 cm																		
DCI06-4B ³	N/A	10-20 cm																		
DCI06-5A ³	N/A	0-10 cm																		
DCI06-5B ³	N/A	10-20 cm																		

File No. 5147-006-13 Table 5 | April 27, 2020



				Conv	ention	al Ana	lyses ²						Ch	emical	Analys	ses ²				
Sample Location ¹	Date Sampled	Sample Interval	Total Organic Carbon (TOC)	Total Volatile Solids (TVS)	Total Solids (TS)	Total Ammonia	Total Sulfides	Grain Size	твт	Metals	LPAHS	нранѕ	сРАНѕ	Chlorinated Hydrocarbons	Phthalates	Miscellaneous Extractables	Phenois	Pesticides	PCB Aroclors	Dioxins and Furans
DCI06-6A ³	N/A	0-10 cm																		
DCI06-7A ³	N/A	0-10 cm																		
DCI06-7B ³	N/A	10-20 cm																		
DCI06-8A ³	N/A	0-10 cm																		
DCI06-9A ³	N/A	0-10 cm																		
Fidalgo Bay Sediment Stu	dy (SAIC 2008)																			
FB-A4-14	09/06/07	0-10 CM																		
FB-A4-15	09/06/07	0-10 CM																		
FB-A4-17 ³	09/06/07	0-10 CM					•													
Sediment Remedial Inves	tigation (GeoEn	gineers 2008)																		
G-1(s) ³	03/13/08	0-20 CM					•				5									
G-2 (s) ³	03/13/08	0-20 CM				•				7										
G-3-0-1 ³	03/14/08	0-1 ft								7										
G-4-2-3 ³	03/14/08	2-3 ft	•		4		-													
G-5-0-1 ³	03/14/08	0-1 ft			-															
G-6-2-3 ³	03/14/08	2-3 ft	H				•													
G-7(s) ³	03/13/08	0-20 cm	Í																	
Interim Action Confirmation	on Sampling (G	eoEngineers 20	(80																	
SMA 1-1	09/30/08	0 - 10 cm	•		•															
SMA 2-1	09/30/08	0 - 10 cm	-																	
SMA 3-2	08/28/08	0 - 10 cm																		
DCI 4-1	10/10/08	0 - 10 cm																		
DCI 4-1A	10/10/08	0 - 10 cm																		
SMA 5-2	08/26/08	0 - 10 cm																		
SMA 5-3	08/26/08	0 - 10 cm											•	•	•					



- ¹ Sediment sampling locations are shown on Figure 12.
- ² Laboratory results are summarized in Appendix C.
- ³ Sediment represented by this sample was subsequently removed from the Marine Area during redevelopment of the Property in 2008. cm = centimeters

ft = feet

TBT = Tributyltin

LPAHs = Low Molecular Weight Polycyclic Aromatic Hydrocarbons

HPAHs = High Molecular Weight Polycyclic Aromatic Hydrocarbons

cPAHs = Carcinogenic Polycyclic Aromatic Hydrocarbons

PCBs = Polychlorinated Biphenyls

■ = Selected sample submitted for chemical analysis.





Schedule of Laboratory Analysis for Groundwater Investigations

								Chem	ical Ana	alyses ²					
Sample Location ¹	Date Sampled	Well Location	Total Metals	Dissolved Metals	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	BETX	Other VOCs	SVOCS	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
Remedial Investigation Study ³	(Landau 2002b)							·	•		•	•			
MW-1	09/04/01	Charalina													
IVIVV-T	10/24/01	Shoreline			•		•								
MW-2	09/04/01	Shoreline		•											
IVIVV-Z	10/24/01	Shoreline		•	-	•									
MW-3	09/04/01	Shoreline		•		-									
IVIVV-3	10/24/01	Silorenne		-			•								
MW-4	09/04/01	Upland				•	•								
10100-4	10/24/01	Оріана		-	•										
Independent Cleanup Action (L	andau 2002c)														
MW-1	06/05/02	Shoreline	-												
	08/19/02	Onorchine													
MW-2	06/05/02	Shoreline													
2	08/19/02	Silordinio													
MW-3	06/05/02	Shoreline													
	08/19/02	Gilordiiilo													
MW-4	06/05/02	Upland											•		
	08/19/02	S prior i d													
Groundwater Characterization	Study (Floyd Snider 20			•	•	•			T	•	T	T	·	•	
MW-1	11/17/06	Shoreline				•	•						•		
MW-2	11/17/06	Shoreline				•	•						•		
MW-3	11/17/06	Shoreline				•	•						•		
MW-4	11/17/06	Upland													



								Chem	nical Ana	alvses ²					
					eg "		ge ,								
Sample Location ¹	Date Sampled	Well Location	Total Metals	Dissolved Metals	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHS	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
Groundwater Remedial Investi	gation ⁴ (GeoEngineers	2010)													
MW-1	06/17/08	Shoreline													
MW-2	06/17/08	Shoreline					•			•					
MW-3	06/17/08	Shoreline			•		•			•					
MW-4	06/17/08	Upland			•		•		•						
MW-5	06/17/08	Upland				•									
Groundwater Remedial Investi	gation ⁵ (GeoEngineers	2013)													
	05/23/12				•		•								
MW-1	08/16/12	Shoreline													
IVIVV-T	11/13/12	Shoreline			-	•									
	02/13/13			•	6	•									
	05/23/12			-			•								
MW-2A	08/16/12	Shoreline				-	•								
IVIVV-ZA	11/13/12	Shorenie	•		•										
	02/13/13		•												
	05/23/12		-	-		-									
MW-3A	08/16/12	Shoreline													
WW-5A	11/13/12	Shoreline		•											
	02/13/13														
	05/23/12														
MW-4	08/16/12	Upland	•												
10100-4	11/13/12	Opiana													
	02/13/13														
	05/23/12														
MW-6	08/16/12	Shoreline													
IVIVV-O	11/13/12	Shoreline													
	02/13/13														



								Chem	ical Ana	lyses ²					
Sample Location ¹	Date Sampled	Well Location	Total Metals	Dissolved Metals	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SNOCs	PAHS	Pesticides	Herbicides	PCB Arociors	Dioxins and Furans
	05/23/12														
MW-7	08/16/12	Upland													
10100-7	11/13/12	оріапи					•								
	02/13/13														
Groundwater Remedial Investi	gation ⁶ (GeoEngineers	2017)													
	02/10/16							•							
MW-1	08/18/16	Shorolino													
IVIVV-1	02/15/17														
	08/23/17														
MW-2A	02/10/16		-7	-7											
	08/19/16	Shoreline	-												
MW-2B	02/15/17	Shoreline													
	08/23/17														
	02/11/16		•	-											
MW-3A	08/19/16	Charolina	•												
IVIW-SA	02/16/17	Shoreline	-												
	08/24/17														
	02/11/16														
MW-4	08/18/16	Unload													
IVI VV-4	02/15/17	upiand													
	08/24/17		•												
	02/11/16														
MW-6	08/19/16	Charolina													
IVIVV-O	02/16/17	Shoreline													
	08/24/17														
	02/10/16			•											
N // N/ 7	08/19/16	Holond													
MW-7	02/16/17	upiand													
	08/23/17	Shoreline Upland Shoreline Upland Upland Upland Upland													



								Chem	ical Ana	llyses ²					
Sample Location ¹	Date Sampled	Well Location	Total Metals	Dissolved Metals	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHS	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	02/10/16														
MW-8	08/18/16	Shoreline ⁸													
IVIVV-O	02/15/17	Shoreline													
	08/23/17														

BETX = Benzene, Ethylbenzene, Toluene and Xylenes

VOCs = Volatile Organic Compounds

SVOCs = Semi-Volatile Organic Compounds

cPAHs = Carcinogenic Polycyclic Aromatic Hydrocarbons

PCBs = Polychlorinated Biphenyls

■ = Selected sample submitted for chemical analysis.



¹ Groundwater sampling locations are shown on Figure 13.

² Laboratory results are summarized in Appendix F.

³ Groundwater monitoring activities completed prior to the 2002 independent cleanup action completed at the Site.

⁴ Remedial investigation activities were completed to evaluate groundwater conditions in accordance with the RI/FS Work Plan (GeoEngineers 2008).

⁵ Additional groundwater monitoring activities were completed on a quarterly basis between May 2012 and February 2013 as directed by Ecology to further evaluate groundwater conditions at the Site.

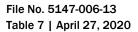
⁶ Due to inconclusive evidence linking contaminant exceedances in soil to contaminant exceedances in groundwater, Ecology required that four additional rounds of groundwater monitoring be completed on a semi-annual basis at the Site (Ecology 2015).

⁷ Monitoring well was damaged. In August 2016, a replacement monitoring well (MW-2B) was installed for evaluating groundwater conditions.

⁸ Ecology has determined that the location of MW-1 was not an appropriate location for evaluation of the conditional point of compliance and that a new monitoring well (MW-8) be installed north of MW-1 to serve as the point of compliance (Ecology 2015).

Laboratory Analysis for Soil Investigations

	1															
									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	ТВТ	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
UST Removal and C	Closure (A-1 1991)							·								
#2	North Wall	10/02/91	N/A													
#4	South Wall	10/02/91	N/A						•							
#6	West Wall	10/02/91	N/A													
#7	North Wall	10/02/91	N/A													
#8	East Wall	10/02/91	N/A					·								
#9	South Wall	10/02/91	N/A				•									
#10	Base Center	10/02/91	N/A													
#1A	Tank Hole	10/02/91	N/A													
#3A	Tank Hole	10/02/91	N/A													
#5A	Tank Hole	10/02/91	N/A													
Phase 2 Environme	ental Site Assessment (Otter	1997)		_												
S-1	DC-B-1	07/14/97	4.5 ft													
S-1	DC-B-1B	07/14/97	4.5 ft													
S-2	DC-B-2	07/14/97	4.5 ft													
S-2	DC-B-2A	07/14/97	2.5 ft													
SS-1A	DC-UPLD SS-1A	07/03/97	0-1 ft													
SS-1B	DC-UPLD SS-1B	07/03/97	0-1 ft													
SS-2A	DC-UPLD SS-2A	07/03/97	0-1 ft													
SS-2B	DC-UPLD SS-2B	07/03/97	0-1 ft													
SS-3	DC-UPLD SS-3	07/30/97	0-1 ft					•								
SS-4	DC-UPLD SS-4	07/30/97	0-1 ft					•	•							
SS-6	DC-UPLD SS-6	07/30/97	0-1 ft					•	•							
SS-9	DC-UPLD SS-9	07/30/97	0-1 ft													





									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHS	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
SS-11	DC-UPLD SS-11 ³	07/30/97	0-1 ft						•							
SS-13A	DC-UPLD SS-13A ³	07/30/97	0-1 ft													
SS-14A	DC-UPLD SS-14A ³	07/30/97	0-1 ft				•									
SS-14B	DC-UPLD SS-14B ³	07/30/97	0-1 ft													
EPA Site Inspection	(Landau 2001)				47											
DCI-SB-UL01	0020-LAI	07/17/01	2 ft					\								
DCI-SB-UL01	0040-LAI	07/17/01	4 ft		•											
DCI-SB-UL01	0070-LAI	07/17/01	7 ft													
DCI-SB-UL03	0020-LAI	07/17/01	2 ft													
DCI-SB-UL03	0060-LAI	07/17/01	6 ft	í	•											
Marine Railway Hyd	raulic Winch Soil Excavatior	n (Landau 2001)						,								
VS-1	VS-1 DH66A	06/28/01	5 ft				▶ ■									
VS-2	VS-2 DH66B	06/28/01	2.5 ft													
VS-3	VS-3 DH66C	06/28/01	2.5 ft													
VS-6	VS-6 DL 19A	07/03/01	2.5 ft													
VS-7	VS-7 DL 19B	07/03/01	2.5 ft													
VS-8	VS-8 DL 19C	07/03/01	2.5 ft													
Remedial Investigat	tion Study (Landau 2002b)															
	S-1-WS-0	08/22/01	0.5-1 ft													
S-1-WS	S-1-WS-1	08/22/01	1-4 ft													
3-1-443	S-1-WS-2	08/22/01	4-7 ft													
	S-1-WS-3	08/22/01	7-10 ft													
	S-2-MS-0	08/22/01	0.5-1 ft													
S-2-MS	S-2-MS-1	08/22/01	1-4 ft													
	S-2-MS-2	08/22/01	4-7 ft													
	S-3-EFA-0	08/22/01	0-1 ft													
S-3-EFA	S-3-EFA-1	08/22/01	1-4 ft													
3-3-LI A	S-3-EFA-2	08/22/01	4-7 ft													
	S-3-EFA-3	08/22/01	10-13 ft													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SOOAS	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	S-4-EFA-0	08/22/01	0-1 ft				4		•							
S-4-EFA	S-4-EFA-1	08/22/01	1-4 ft													
	S-4-EFA-2	08/22/01	4-7 ft				•									
	S-5-EFA-0	08/22/01	0-1 ft													
	S-5-EFA-1	08/22/01	1-4 ft			•	•									
S-5-EFA	S-5-EFA-2	08/22/01	4-7 ft													
	S-5-EFA-3	08/22/01	7-10 ft			T										
	S-5-EFA-4	08/22/01	10-13 ft													
	S-6-TPH-0	08/22/01	0-1 ft													
S-6-UST	S-6-TPH-1	08/22/01	1-4 ft	į												
3-0-031	Dup (S-6-TPH-1)	08/22/01	1-4 ft	-		-										
	S-6-TPH-2	08/22/01	4-7 ft				▶ ■									
	S-7-TPH-0 ³	08/22/01	0-1 ft			•										
S-7-UST	S-7-TPH-1 ³	08/22/01	1-4 ft													
3-7-031	S-7-TPH-2 ³	08/22/01	4-7 ft													
	S-7-TPH-3 ³	08/22/01	7-10 ft													
	S-8-TPH-0 ³	08/22/01	0-1 ft													
S-8-UST	S-8-TPH-1 ³	08/22/01	1-4 ft													
3-6-031	S-8-TPH-2 ³	08/22/01	4-7 ft													
	S-8-TPH-3 ³	08/22/01	7-10 ft													
	S-9-CPH-0	08/22/01	0-1 ft													
	Dup (S-9-CPH-0)	08/22/01	0-1 ft													
S-9-CPH	S-9-CPH-1	08/22/01	1-4 ft													
3-3-0FF	S-9-CPH-2	08/22/01	4-7 ft													
	S-9-CPH-3	08/22/01	7-9 ft													
	S-9-CPH-3A	08/22/01	9-10 ft													



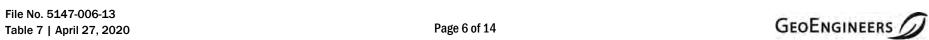
									Chemi	ical Ana	alyses ²	!				
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	181	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	BETX	Other VOCs	SOOAS	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	S-10-MR-0	08/22/01	0-1 ft													
S-10-MR	S-10-MR-1	08/22/01	1-4 ft													
3-10-WIK	S-10-MR-2	08/22/01	4-7 ft			4	•			•						
	S-10-MR-3	08/22/01	7-10 ft													
S-11-MR	S-11-MR ³	08/22/01	0-1 ft			•	•									
S-12-MR	S-12-MR-0 ³	08/22/01	0-1 ft													
3-12-WIK	Dup (S-12-MR-0) ³	08/22/01	0.7 ft			T										
S-13-MR	S-13-MR ³	08/22/01	0-0.5 ft				•									
	S-14-TPH-1	10/24/01	1-3.1 ft													
S-14-TPH	S-14-TPH-4	10/24/01	4-6.4 ft			•										
	S-14-TPH-7	10/24/01	7-10 ft			•										
	S-15-TPH-1	10/24/01	1-3.8 ft				•									
S-15-TPH	S-15-TPH-4	10/24/01	4-6.1 ft													
	S-15-TPH-7	10/24/01	7-9.9 ft													
	S-16-TPH-1 ³	10/24/01	1-3.7 ft													
S-16-TPH	S-16-TPH-4 ³	10/24/01	4-6.3 ft													
	S-16-TPH-7 ³	10/24/01	7-10 ft													
	S-17-TPH-1	10/24/01	1-3.7 ft													
S-17-TPH	S-17-TPH-4A	10/24/01	4-4.4 ft													
3-17-1111	S-17-TPH-4B	10/24/01	4.4-6.3 ft													
	S-17-TPH-7	10/24/01	7-9.8 ft													
	S-18-TPH-1	10/24/01	1-3.4 ft													
S-18-TPH	S-18-TPH-4	10/24/01	4-6.7 ft													
	S-18-TPH-7	10/24/01	7-9.9 ft													
	S-19-TPH-1	10/24/01	1-3.6 ft													
S-19-TPH	S-19-TPH-4	10/24/01	4-6.4 ft													
	S-19-TPH-7	10/24/01	7-9.9 ft													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	S-20-TPH-1 ³	10/24/01	1-3.9 ft													
S-20-TPH	S-20-TPH-4 ³	10/24/01	4-6.5 ft													
	S-20-TPH-7 ³	10/24/01	7-10 ft													
	S-21-TPH-1 ³	10/24/01	1-2.2 ft													
S-21-TPH	S-21-TPH-4 ³	10/24/01	4-4.1 ft			•	_									
	S-21-TPH-7 ³	10/24/01	7-9.4 ft													
	S-22-TPH-1A ³	10/24/01	1-2.5 ft			7										
S-22-TPH	S-22-TPH-1B ³	10/24/01	2.5-4 ft				•									
5-22-1PH	S-22-TPH-4 ³	10/24/01	4-5 ft													
	S-22-TPH-7 ³	10/24/01	7-9.5 ft													
	S-23-TPH-1 ³	10/24/01	1-3.4 ft			•										
S-23-TPH	S-23-TPH-4 ³	10/24/01	4-6.7 ft				▶ ■									
	S-23-TPH-7 ³	10/24/01	7-9.6 ft													
Independent Cleanu	up Action (Landau 2002c)							•								
CS-1	CS-1 8-20	08/20/02	N/A													
CS-2	CS-2 8-20	08/20/02	N/A		ŀ											
CS-3	CS-3 8-20	08/20/02	N/A													
CS-4	CS-4 8-20	08/20/02	N/A													
CS-5	CS-5 8-20	08/20/02	N/A													
CS-6	CS-6 8-20	08/20/02	N/A													
CS-7	CS-7 8-20	08/20/02	N/A													
CS-8	CS-8 8-20	08/20/02	N/A													
CS-9	CS-9 8-20	08/20/02	N/A													
CS-10	CS-10 8-20	08/20/02	N/A													
CS-11	CS-11 8-20	08/20/02	N/A													
CS-12	CS-12 8-20	08/20/02	N/A				-									
CS-13	CS-13 8-20	08/20/02	N/A				•									
CS-14	CS-14 8-20	08/20/02	N/A													
CS-15	CS-15 8-20	08/20/02	N/A													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
CS-16	CS-16 8-20	08/20/02	N/A													
CS-17	CS-17 8-20 ³	08/20/02	N/A													
CS-18	CS-18 8-20	08/20/02	N/A				•									
CS-19	CS-19 8-20 ³	08/20/02	N/A													
CS-20	CS-20 8-20 ³	08/20/02	N/A			•	_									
CS-21	CS-21 8-20	08/20/02	N/A													
CS-22	CS-22 8-20	08/20/02	N/A			T										
CS-23	CS-23 8-20	08/20/02	N/A				•									
CS-24	CS-24 8-20	08/20/02	N/A				•									
CS-25	CS-25 8-20	08/20/02	N/A													
CS-26	CS-26 8-20 ³	08/20/02	N/A			•										
CS-27	CS-27 8-20	08/20/02	N/A				■									
CS-28	CS-28 8-20	08/20/02	N/A	1												
CS-29	CS-29 8-20	08/20/02	N/A													
CS-30	CS-30 8-20	08/20/02	N/A													
CS-31	CS-31 8-20	08/20/02	N/A		ŀ											
CS-32	CS-32 8-20	08/20/02	N/A													
CS-33	CS-33 8-20	08/20/02	N/A													
CS-34	CS-34 8-20	08/20/02	N/A													
CS-35	CS-35 8-20	08/20/02	N/A													
CS-36	CS-36 8-20	08/20/02	N/A													
CS-37	CS-37 8-20	08/20/02	N/A													
CS-38	CS-38 8-20 ³	08/20/02	N/A													
CS-39	CS-39 8-20	08/20/02	N/A													
CS-40	CS-40 8-20	08/20/02	N/A													
CS-41	CS-41 8-20	08/20/02	N/A													
CS-42	CS-42 8-20	08/20/02	N/A													
CS-43	CS-43 8-20	08/20/02	N/A				•									
CS-44	CS-44 8-20	08/20/02	N/A													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
CS-45	CS-45 8-20	08/20/02	N/A													
CS-46	CS-46 8-20	08/20/02	N/A													
CS-47	CS-47 8-20	08/20/02	N/A				•									
CS-48	CS-48 8-20	08/20/02	N/A													
CS-49	CS-49 8-20	08/20/02	N/A			•	_									
CS-50	CS-50 8-20	08/20/02	N/A													
CS-51	CS-51 8-20	08/20/02	N/A			-										
CS-52	CS-52 8-20	08/20/02	N/A				•									
CS-53	CS-53 8-20	08/20/02	N/A			F	•									
CS-54	CS-54 8-20	08/20/02	N/A													
CS-55	CS-55 8-20	08/20/02	N/A			•										
CS-56	CS-56 8-20	08/20/02	N/A				-									
Remedial Investigat	tion ⁴ (GeoEngineers 2010)			V 7				•					•			
SS-1	SS-1-1	06/16/08	1-1.5 ft													
SS-2	SS-2-1	06/16/08	1-1.5 ft													
SS-3	SS-3-1	06/16/08	1-1.5 ft		ŀ											
SS-4	SS-4-0.5	06/16/08	0.5 -1 ft													
MW-5	MW-5-5.0	05/27/08	5-6.5 ft													
C-VVIVI	MW-5-10.0	05/27/08	10-11.5 ft													
SB-1	SB-1-2.0	06/16/08	2-3 ft													
2B-T	SB-1-4.0	06/16/08	4-5 ft													
SB-2	SB-2-2.0	06/16/08	2-3 ft													
3D-2	SB-2-4.0	06/16/08	4-5 ft													
SB-4	SB-4-3.0	06/16/08	3-4 ft				•	•								
SĎ-4	SB-4-9.0	06/16/08	9-10 ft													•
SB-5	SB-5-3.0	06/16/08	3-4 ft													
G-9C	SB-5-9.0	06/16/08	9-10 ft													-
SB-7	SB-7-3.0	06/16/08	3-4 ft													
3D-1	SB-7-9.0	06/16/08	9-10 ft													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
SB-8	SB-8-0.5	06/17/08	0.5-1.5 ft													
30-0	SB-8-4.0	06/17/08	4-5 ft													
SB-9	SB-9-0.5	06/16/08	0.5-1.5 ft													
36-9	SB-9-4.0	06/16/08	4-5 ft													
SB-10	SB-10-0.5	06/17/08	0.5-1.5 ft													
3B-10	SB-10-4.0	06/17/08	4-5 ft				1									
SB-11	SB-11-0.5	06/17/08	0.5-1.5 ft													
3D-11	SB-11-4.0	06/17/08	4-5 ft													
SB-12	SB-12-0.5	06/16/08	0.5-1.5 ft													
2D-12	SB-12-4.0	06/16/08	4-5 ft													
SB-13	SB-13-0.5	06/16/08	0.5-1.5 ft													
2B-T2	SB-13-4.0	06/16/08	4-5 ft				•									
SB-14	SB-14-0.5	06/16/08	0.5-1.5 ft													
3B-14	SB-14-4.0	06/16/08	4-5 ft													
SB-15	SB-15-0.5	06/16/08	0.5-1.5 ft													
3B-13	SB-15-4.0	06/16/08	4-5 ft													
TP-3	TP-3-6	09/05/08	6-6.5 ft													
TP-4	TP-4-6	09/08/08	6-6.5 ft													
TP-5	TP-5-2	09/08/08	2-2.5 ft													
11-5	TP-5-4	09/08/08	4-4.5 ft													
TP-10	TP-10-4	09/08/08	4-4.5 ft													
16-10	TP-10-6	09/08/08	6-6.5 ft													
TP-11	TP-11-6	09/08/08	6-6.5 ft													
TP-12	TP-12-3	09/08/08	3-3.5 ft													
TP-13	TP-13-2	09/08/08	2-2.5 ft													
11-13	TP-13-4	09/08/08	4-4.5 ft													
TP-14	TP-14-0-2	09/18/08	0-2 ft													
TP-15	TP-15-2-4	09/18/08	2-4 ft													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Arociors	Dioxins and Furans
TP-16	TP-16-0-2	09/18/08	0-2 ft													
	TP-16-4-6	09/18/08	4-6 ft													
Remedial Investiga	tion ⁵ (GeoEngineers 2014)						· ·									
GEI-1	GEI-01_3-4_092914	09/29/2014	3-4 ft													
GLI-I	GEI-02_1-2_092914	09/29/2014	1-2 ft													
GEI-2	GEI-02_4-5_092914	09/29/2014	4-5 ft													
GLI-2	GEI-02_7-8_092914	09/29/2014	7-8 ft													
GEI-3	GEI-03_2.5-3.5_092914	09/29/2014	2.5-3.5 ft													
GEI-3	GEI-03_7-8_092914	09/29/2014	7-8 ft													
	GEI-04_1-2_092914	09/29/2014	1-2 ft													
GEI-4	GEI-04_3-4_092914	09/29/2014	3-4 ft	-												
	GEI-04_6-7_092914	09/29/2014	6-7 ft													
GEI-5	GEI-05_7-8_092914	09/29/2014	7-8 ft													
	GEI-06_1.5-2.5_092914	09/29/2014	1.5-2.5 ft													
GEI-6	GEI-06_4-5_092914	09/29/2014	4-5 ft													
	GEI-06_7-8_092914	09/29/2014	7-8 ft	-												
GEI-7	GEI-07_1.5-2.5_092914	09/29/2014	1.5-2.5 ft													
GEI-7	GEI-07_7-8_092914	09/29/2014	7-8 ft													
	GEI-08_1.5-2.5_092914	09/29/2014	1.5-2.5 ft													
GEI-8	GEI-08_4-5_092914	09/29/2014	4-5 ft													
	GEI-08_7-8_092914	09/29/2014	7-8 ft													
	GEI-09_0.5-1.5_092914	09/29/2014	0.5-1.5 ft													
GEI-9	GEI-09_3-4_092914	09/29/2014	3-4 ft													
	GEI-09_6-7_092914	09/29/2014	6-7 ft													
GEI-10	GEI-10_2-3_092914	09/29/2014	2-3 ft													
GEI-10	GEI-10_7-8_092914	09/29/2014	7-8 ft													
	GEI-11_2-3_092914	09/29/2014	2-3 ft													
GEI-11	GEI-11_7-8_092914	09/29/2014	7-8 ft													
	GEI-11_9-10_092914	09/29/2014	9-10 ft													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	твт	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	GEI-12_2-3_092914	09/29/2014	2-3 ft													
GEI-12	GEI-12_4-5_092914	09/29/2014	4-5 ft													
	GEI-12_7-8_092914	09/29/2014	7-8 ft													
	GEI-13_2-3_093014	09/30/2014	2-3 ft													
GEI-13	GEI-13_5-6_093014	09/30/2014	5-6 ft													
	GEI-13_7-8_093014	09/30/2014	7-8 ft	-												
	GEI-14_2-3_093014	09/30/2014	2-3 ft													
GEI-14	GEI-14_3.5-4.5_093014	09/30/2014	3.5-4.5 ft													
GEI-14	GEI-14_7-8_093014	09/30/2014	7-8 ft	•												
	GEI-14_9-10_093014	09/30/2014	9-10 ft	7												
	GEI-15_10-11_093014	09/30/2014	10-11 ft	1												
GEI-15	GEI-15_2-3_093014	09/30/2014	2-3 ft													
	GEI-15_5.5-6.5_093014	09/30/2014	5.5-6.5 ft													
	GEI-16_2-3_093014	09/30/2014	2-3 ft													
GEI-16	GEI-16_6-7_093014	09/30/2014	6-7 ft													
	GEI-16_8-9_093014	09/30/2014	8-9 ft	-												
	GEI-17_1-2_093014	09/30/2014	1-2 ft													
GEI-17	GEI-17_4-5_093014	09/30/2014	4-5 ft													
GEI-17	GEI-17_7-8_093014	09/30/2014	7-8 ft													
	GEI-17_9-10_093014	09/30/2014	9-10 ft													
	GEI-18_1-2_093014	09/30/2014	1-2 ft													
GEI-18	GEI-18_4-5_093014	09/30/2014	4-5 ft													
GEI-10	GEI-18_8-9_093014	09/30/2014	8-9 ft													
	GEI-18_9-10_093014	09/30/2014	9-10 ft													
	GEI-19_2-3_093014	09/30/2014	2-3 ft													
GEI-19	GEI-19_4-5_093014	09/30/2014	4-5 ft													
GEI-19	GEI-19_7-8_093014	09/30/2014	7-8 ft													
	GEI-19_9-10_093014	09/30/2014	9-10 ft													



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHS	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	GEI-20_2-3_093014	09/30/2014	2-3 ft													
GEI-20	GEI-20_4-5_093014	09/30/2014	4-5 ft													
GLI-20	GEI-20_6-7_093014	09/30/2014	6-7 ft													
	GEI-20_8-9_093014	09/30/2014	8-9 ft													
	GEI-21_1-2_093014	09/30/2014	1-2 ft													
GEI-21	GEI-21_5-6_093014	09/30/2014	5-6 ft	-												
	GEI-21_7.5-8.5_093014	09/30/2014	7.5-8.5 ft													
	GEI-22_2-3_100114	10/01/2014	2-3 ft													
GEI-22	GEI-22_5-6_100114	10/01/2014	5-6 ft													
	GEI-22_7.5-8.5_100114	10/01/2014	7.5-8.5 ft													
	GEI-23_1-2_093014	09/30/2014	1-2 ft	-												
GEI-23	GEI-23_5-6_093014	09/30/2014	5-6 ft													
	GEI-23_7.5-8.5_093014	09/30/2014	7.5-8.5 ft													
	GEI-24_2-3_093014	09/30/2014	2-3 ft	•												
GEI-24	GEI-24_4-5_093014	09/30/2014	4-5 ft													
GLI-24	GEI-24_6-7_093014	09/30/2014	6-7 ft													
	GEI-24_9-10_093014	09/30/2014	9-10 ft													
	GEI-25_1-2_093014	09/30/2014	1-2 ft													
GEI-25	GEI-25_4-5_093014	09/30/2014	4-5 ft													
GLI 23	GEI-25_7-8_093014	09/30/2014	7-8 ft													
	GEI-25_9-10_093014	09/30/2014	9-10 ft													
GEI-26	GEI-26_2 -3_093014	09/30/2014	2-3 ft													
GLF20	GEI-26_6-7_093014	09/30/2014	6-8 ft													
	GEI-27_1-2_100114	10/01/2014	1-2 ft													
GEI-27	GEI-27_5-6_100114	10/01/2014	5-6 ft													
	GEI-27_9-10_100114	10/01/2014	9-10 ft													
	GEI-28_10-11_100114	10/01/2014	10-11 ft													
GEI-28	GEI-28_2-3_100114	10/01/2014	2-3 ft													
	GEI-28_5-6_100114	10/01/2014	5-6 ft													7



									Chemi	cal Ana	alyses ²					
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	TBT	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SVOCs	PAHs	Pesticides	Herbicides	PCB Arociors	Dioxins and Furans
	GEI-29_2-3_093014	09/30/2014	2-3 ft													
GEI-29	GEI-29_5-6_093014	09/30/2014	5-6 ft													
GEI 25	GEI-29_8-9_093014	09/30/2014	8-9 ft				ŀ									
	GEI-29_9-10_093014	09/30/2014	9-10 ft													
	GEI-30_3-4_093014	09/30/2014	3-4 ft													
GEI-30	GEI-30_7-8_093014	09/30/2014	7-8 ft													
	GEI-30_9-10_093014	09/30/2014	9-10 ft													
	GEI-31_1-2_100114	10/01/2014	1-2 ft													
GEI-31	GEI-31_4-5_100114	10/01/2014	4-5 ft													
GEI-SI	GEI-31_6-7_100114	10/01/2014	6-7 ft	į												
	GEI-31_9-10_100114	10/01/2014	9-10 ft													
GEI-32	GEI-32_1-2_100114	10/01/2014	1-2 ft													
GEI-32	GEI-32_6-7_100114	10/01/2014	6-7 ft													
GEI-33	GEI-33_1-2_100114	10/01/2014	1-2 ft													
	GEI-34_2.5-3.5_100114	10/01/2014	2.5-3.5 ft													
GEI-34	GEI-34_6-7_100114	10/01/2014	6-7 ft													
	GEI-34_9-10_100114	10/01/2014	9-10 ft													
	GEI-35_3-4_100114	10/01/2014	3-4 ft	•												
GEI-35	GEI-35_4-5_100114	10/01/2014	4-5 ft													
GLI-55	GEI-35_8-9_100114	10/01/2014	8-9 ft													
	GEI-35_9-10_100114	10/01/2014	9-10 ft													
GEI-36	GEI-36_1-2_100114	10/01/2014	1-2 ft													
GEI-30	GEI-36_5-6_100114	10/01/2014	5-6 ft													
GEI-37	GEI-37_1-2_100114	10/01/2014	1-2 ft													
GEI-31	GEI-37_6-7_100114	10/01/2014	6-7 ft													
GEI-38	GEI-38_1-2_100114	10/01/2014	1-2 ft													
GEI-30	GEI-38_6-7_100114	10/01/2014	6-7 ft													



									Chemi	cal Ana	alyses ²	!				
Sample Location ¹	Sample Identification	Date Sampled	Sample Interval	Metals	ТВТ	Gasoline-Range Hydrocarbons	Diesel-Range Hydrocarbons	Heavy Oil-Range Hydrocarbons	ВЕТХ	Other VOCs	SOOAS	PAHs	Pesticides	Herbicides	PCB Aroclors	Dioxins and Furans
	GEI-39_1.5-2.5_100114	10/01/2014	1.5-2.5 ft													
GEI-39	GEI-39_4-5_100114	10/01/2014	4-5 ft													
	GEI-39_6-7_100114	10/01/2014	6-7 ft													
GEI-40	GEI-40_2-3_100114	10/01/2014	2-3 ft													
	GEI-41_1-2_100114	10/01/2014	1-2 ft													
GEI-41	GEI-41_4-5_100114	10/01/2014	4-5 ft													
GEI 41	GEI-41_6-7_100114	10/01/2014	6-7 ft													
	GEI-41_8-9_100114	10/01/2014	8-9 ft													
	GEI-42_1-2_100114	10/01/2014	1-2 ft													
GEI-42	GEI-42_4-5_100114	10/01/2014	4-5 ft	Í												
	GEI-42_6-7_100114	10/01/2014	6-7 ft	-												
GEI-43	GEI-43_1-2_100114	10/01/2014	1-2 ft													
GEI 45	GEI-43_6-7_100114	10/01/2014	6-7 ft	1												
Remedial Investiga	tion ⁶ (GeoEngineers 2018)															
	GEI-44_1.5-2	07/23/18	1.5-2 ft													
GEI-44	GEI-44_7.5-10	07/23/18	7.5-10 ft													
	GEI-44_16-17.5	07/23/18	16-17.5 ft													
	GEI-45_1-3	07/23/18	1-3 ft													
GEI-45	GEI-45_9-10	07/23/18	9-10 ft													
	GEI-45_17-20	07/23/18	17-20 ft													
GEI-46	GEI-46_7-8.5	07/23/18	1-2 ft													
GLI-40	GEI-46_13.5-15	07/23/18	4-5 ft													



BETX = Benzene, Ethylbenzene, Toluene and Xylenes

VOCs = Volatile Organic Compounds

SVOCs = Semi-Volatile Organic Compounds

cPAHs = Carcinogenic Polycyclic Aromatic Hydrocarbons

PCBs = Polychlorinated Biphenyls

■ = Selected sample submitted for chemical analysis.





¹ Soil sampling locations are shown on Figure 14.

² Laboratory results are summarized in Appendix G.

³ Soil represented by this sample was subsequently removed from the Upland Area during the 2002 independent cleanup action completed at the Site.

⁴ Remedial investigation activities were completed to evaluate soil conditions in accordance with the RI/FS Work Plan (GeoEngineers 2008a).

⁵ Additional soil investigation activities were completed as directed by Ecology to fill identified data gaps. Soil investigation activities were completed in accordance with the Ecology-approved RI/FS Work Plan (GeoEngineers 2008a).

⁶ Additional soil investigation activities were completed as directed by Ecology based on semi-annual groundwater monitoring results at MW-8. Soil investigation activities were completed in accordance with the Ecology-approved RI/FS Work Plan Addendum (GeoEngineers, 2018).

Table 8

Summary Statistics and Evaluation of Sediment Contaminants of Concern - Protection of Benthic Organisms

	1		ı	1						wasningi					1			
		osed			1		1		Evaluation of RI	Data Results ³						Contaminant of	f Concern (COC)	Selection Considerations
		ment							RL Exceedance	Evaluation	SCO/AET Exceeda	nce Evaluation	CSL/2AET Exceeda	ance Evaluation				
		anup .2					Maximum	Maximum	Frequency	Maximum	Frequency	Maximum	Frequency	Maximum	Initial COC		Proposed	
Contaminant of		vel ²				Detection	Non-Detect	Detected	of RL	Exceedance	of PCUL	Exceedance	of PCUL	Exceedance	Selection	Groundwater	Sediment	
Potential Concern ¹	SCO/	CSL/		Number	Number of	Frequency	Concentration	Concentration	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	Criteria Met ⁶	COC ⁷	coc	
(COPC)	LAET	2LAET	Units	Samples	Detections	(%)	(µg/L)	(µg/L)	(%)	(ER)	(%)	(ER)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
Metals			1			Г	T			T	T			T	I	1	I	
Arsenic	57	73	mg/kg	18	16	89%	10	300	No Exceedance	0.2	11%	5.3	6%	4.1	Yes	Yes	Yes	Retained as a COC
Cadmium	5.1	6.7	mg/kg	18	13	72%	0.6	1.2	No Exceedance	0.1	No Exceedance	0.2	No Exceedance	0.2	No	No	No	None
Chromium	260	270	mg/kg	18	18	100%	-	55	No Exceedance	-	No Exceedance	0.2	No Exceedance	0.2	No	No	No	None
Copper	390	390	mg/kg	18	18	100%		3,870	No Exceedance		28%	9.9	28%	9.9	Yes	No	Yes	Retained as a COC
Lead	450	530	mg/kg	18	18	100%	-	939	No Exceedance		11%	2.1	11%	1.8	Yes	No	Yes	Retained as a COC
Mercury	0.41	0.59	mg/kg	18	11	61%	-	17.8	No Exceedance	-	28%	43.4	22%	30.2	Yes	No	Yes	Retained as a COC
Nickel	NE	NE	mg/kg	9	9	100%		35.5	No Exceedance	-	No Exceedance		No Exceedance	-	No	Yes	No	None
Silver	6.1	6.1	mg/kg	18	11	61%	0.9	0.5	No Exceedance	0.1	No Exceedance	0.1	No Exceedance	0.1	No	No	No	None
Zinc	410	960	mg/kg	18	18	100%	-	974	No Exceedance	-	17%	2.4	6%	1.0	Yes	No	Yes	Retained as a COC
Low Molecular Weight Polycyclic Aron	1	1	s) (Dry Weig	1	_		•	1		1					I		1	_
Sum of LPAHs ¹	5,200	5,200	µg/kg	3	3	100%	-	10,290	No Exceedance		33%	2.0	33%	2.0	Yes	No	Yes	
2-Methylnaphthalene	670	670	µg/kg	3	1	33%	59	4,100	No Exceedance	<0.1	33%	6.1	33%	6.1	Yes	No	Yes	
Acenaphthene	500	500	µg/kg	3	1	33%	20	230	No Exceedance	<0.1	No Exceedance	0.5	No Exceedance	0.5	No	No	Yes	LPAHs are retained as a COC based on
Acenaphthylene	1,300	1,300	µg/kg	3	0	0%	59	250	No Exceedance	<0.1	No Exceedance	0.2	No Exceedance	0.2	No	No	Yes	one or more analytes meeting the initial
Anthracene	960	960	μg/kg	3	3	100%	9	1,420	No Exceedance	-	33%	1.5	33%	1.5	Yes	No	Yes	selection criteria.
Fluorene	540	540	μg/kg	3	3	100%	20	742	No Exceedance	<0.1	33%	1.4	33%	1.4	Yes	No	Yes	-
Naphthalene	2,100	2,100	μg/kg 	3	1	33%	59	3,060	No Exceedance	<0.1	33%	1.5	33%	1.5	Yes	No	Yes	-
Phenanthrene	1,500	1,500	µg/kg	3	3	100%	9	5,070	No Exceedance	-	33%	3.4	33%	3.4	Yes	No	Yes	
Low Molecular Weight Polycyclic Aron		,	<u> </u>	1	1	Т	T	1					Г	Т	T	1	I	1
Sum of LPAHs ¹	370	780	mg/kg OC		10	100%		263.0	No Exceedance	4	No Exceedance	0.7	No Exceedance	0.3	No	No	Yes	
2-Methylnaphthalene	38	64	mg/kg OC	10	7	70%	0.36	6.4	No Exceedance	<0.1	No Exceedance	0.2	No Exceedance	0.1	No	No	Yes	
Acenaphthene	16	57	mg/kg OC	10	8	80%	0.36	16.5	No Exceedance	<0.1	10%	1.0	No Exceedance	0.3	No	No	Yes	LPAHs are retained as a COC based on
Acenaphthylene	66	66	mg/kg OC	10	9	90%	3.3	15.6	No Exceedance	0.1	No Exceedance	0.2	No Exceedance	0.2	No	No	Yes	one or more analytes meeting the initial
Anthracene	220	1,200	mg/kg OC	10	10	100%	-	118.8	No Exceedance	-	No Exceedance	0.5	No Exceedance	0.1	No	No	Yes	selection criteria for dry weight evaluation.
Fluorene	23	79	mg/kg OC	10	9	90%	3.3	19.4	No Exceedance	0.1	No Exceedance	0.8	No Exceedance	0.2	No	No	Yes	evaluatiOII.
Naphthalene	99	170	mg/kg OC	10	8	80%	3.3	11	No Exceedance	<0.1	No Exceedance	0.1	No Exceedance	0.1	No	No	Yes	
Phenanthrene	100	480	mg/kg OC	10	10	100%	-	122	No Exceedance	-	10%	1.2	No Exceedance	0.3	No	No	Yes	
High Molecular Weight Polycyclic Aro	1	· ·		 	1						T .		T .			1		Ţ
Sum of HPAHs ⁸	12,000	17,000	μg/kg	3	3	100%	_	28,020	No Exceedance		33%	2.3	33%	1.6	Yes	No	Yes	
Benzo(a)anthracene	1,300	1,600	µg/kg	3	3	100%	-	1,420	No Exceedance		33%	1.1	No Exceedance	0.9	Yes	Yes	Yes	
Benzo(a)pyrene	1,600	1,600	µg/kg	3	3	100%	-	4,100	No Exceedance		33%	2.6	33%	2.6	Yes	Yes	Yes	
Total Benzofluoranthenes ⁹	3,200	3,600	μg/kg	3	3	100%	-	2,990	No Exceedance		No Exceedance	0.9	No Exceedance	0.8	Yes	Yes	Yes	Retained as a COC based on the
Benzo(g,h,i)perylene	670	720	μg/kg	3	3	100%	-	4,850	No Exceedance		33%	7.2	33%	6.7	Yes	No	Yes	frequency of initial selection criteria
Chrysene	1,400	2,800	µg/kg	3	3	100%	-	4,150	No Exceedance		33%	3.0	33%	1.5	Yes	Yes	Yes	exceedances for HPAHs.
Dibenzo(a,h)anthracene	230	230	µg/kg	3	2	67%	6.1	3,060	No Exceedance	<0.1	33%	13.3	33%	13.3	Yes	No	Yes	
Fluoranthene	1,700	2,500	µg/kg	3	3	100%		2,100	No Exceedance		67%	1.2	No Exceedance	0.8	Yes	No	Yes	
Indeno(1,2,3-c,d)pyrene	600	690	μg/kg	3	3	100%		2,050	No Exceedance		33%	3.4	33%	3.0	Yes	Yes	Yes	
Pyrene	2,600	3,300	μg/kg	3	3	100%	-	3,300	No Exceedance		33%	1.3	No Exceedance	1.0	Yes	No	Yes	



	Prop	osed			1				Evaluation of RI	Data Results	1		1			Contaminant o	t Concern (COC)	Selection Considerations
	Sedi	ment							RL Exceedance	Evaluation	SCO/AET Exceeda	nce Evaluation	CSL/2AET Exceeda	nce Evaluation				
	Clea	nup					Maximum	Maximum	Frequency	Maximum	Frequency	Maximum	Frequency	Maximum	Initial COC		Proposed	
Contaminant of	Lev	/el²				Detection	Non-Detect	Detected	of RL	Exceedance	of PCUL	Exceedance	of PCUL	Exceedance	Selection	Groundwater	Sediment	
Potential Concern ¹	SCO/	CSL/		Number	Number of	Frequency	Concentration	Concentration	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	Criteria Met ⁶	COC ⁷	coc	
(COPC)	LAET	2LAET	Units	Samples	Detections	(%)	(µg/L)	(µg/L)	(%)	(ER)	(%)	(ER)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
ligh Molecular Weight Polycyclic Arc	omatic Hydroca	rbons (HPAI	ls) (OC Norm	alized)														
Sum of HPAHs ⁸	960	5,300	mg/kg OC	10	10	100%		1,888.1	No Exceedance		30%	2.0	No Exceedance	0.4	Yes	No	Yes	
Benzo(a)anthracene	110	270	mg/kg OC	10	10	100%		166.7	No Exceedance		20%	1.5	No Exceedance	0.6	Yes	Yes	Yes	
Benzo(a)pyrene	99	210	mg/kg OC	10	10	100%		135.9	No Exceedance		30%	1.4	No Exceedance	0.6	Yes	Yes	Yes	
Total Benzofluoranthenes ⁹	230	450	mg/kg OC	10	10	100%	-	281.6	No Exceedance	-	20%	1.2	No Exceedance	0.6	Yes	Yes	Yes	Detained as a COC based on the
Benzo(g,h,i)perylene	31	78	mg/kg OC	10	8	80%	0.56	70.9	No Exceedance	<0.1	30%	2.3	No Exceedance	0.9	Yes	No	Yes	 Retained as a COC based on the frequency of initial selection crite
Chrysene	110	460	mg/kg OC	10	10	100%	-	193.8	No Exceedance		40%	1.8	No Exceedance	0.4	Yes	Yes	Yes	exceedances for HPAHs.
Dibenzo(a,h)anthracene	12	33	mg/kg OC	10	8	80%	0.56	17.3	No Exceedance	<0.1	30%	1.4	No Exceedance	0.5	Yes	No	Yes	exceedances for the Aris.
Fluoranthene	160	1,200	mg/kg OC	10	10	100%		687.5	No Exceedance	_	50%	4.3	No Exceedance	0.6	Yes	No	Yes	
Indeno(1,2,3-c,d)pyrene	34	88	mg/kg OC	10	10	100%		70.9	No Exceedance	_	30%	2.1	No Exceedance	0.8	Yes	Yes	Yes	
Pyrene	1,000	1,400	mg/kg OC	10	10	100%		468.8	No Exceedance		No Exceedance	0.5	No Exceedance	0.3	No	No	Yes	
Chlorinated Hydrocarbons (Dry Weig	(ht)															-		-
1,2,4-Trichlorobenzene	31	51	µg/kg	1	0	0%	6.2	-	No Exceedance	0.2	No Exceedance	_	No Exceedance	-	No	No	No	None
1,2-Dichlorobenzene	35	50	µg/kg	1	0	0%	6.2		No Exceedance	0.2	No Exceedance		No Exceedance	-	No	No	No	None
1,4-Dichlorobenzene	110	110	µg/kg	1	0	0%	6.2	-	No Exceedance	0.1	No Exceedance	-	No Exceedance	-	No	No	No	None
Hexachlorobenzene	22	70	µg/kg	1	0	0%	6.2		No Exceedance	0.3	No Exceedance	-	No Exceedance	-	No	No	No	None
Chlorinated Hydrocarbons (OC Norm	alized)												•	•	•	•	•	•
1,2,4-Trichlorobenzene	0.81	1.8	mg/kg OC	9	0	0%	0.6		No Exceedance	0.7	No Exceedance		No Exceedance	-	No	No	No	None
1,2-Dichlorobenzene	2.3	2.3	mg/kg OC	9	3	33%	0.6	1.3	No Exceedance	0.3	No Exceedance	0.6	No Exceedance	0.6	No	No	No	None
1,4-Dichlorobenzene	3.1	9	mg/kg OC	9	2	22%	0.6	1	No Exceedance	0.2	No Exceedance	0.3	No Exceedance	0.1	No	No	No	None
Hexachlorobenzene	0.38	2.3	mg/kg OC	9	0	0%	0.6		No Exceedance	1.6	No Exceedance	-	No Exceedance	-	No	No	No	None
Phthalates (Dry Weight)	•			•	•								•	•	•		-	
Bis(2-Ethylhexyl) Phthalate	1,300	1,900	µg/kg	1	1	100%		400	No Exceedance	_	No Exceedance	0.3	No Exceedance	0.2	No	No	No	None
Butyl benzyl Phthalate	63	900	µg/kg	1	0	0%	15	_	No Exceedance	0.2	No Exceedance	-	No Exceedance		No	No	No	None
Dibutyl Phthalate	1,400	1,400	µg/kg	1	0	0%	59	_	No Exceedance	<0.1	No Exceedance	_	No Exceedance	_	No	No	No	None
Diethyl Phthalate	200	200	µg/kg	1	0	0%	59	_	No Exceedance	0.3	No Exceedance	_	No Exceedance		No	No	No	None
Dimethyl Phthalate	71	160	µg/kg	1	0	0%	59	_	No Exceedance	0.8	No Exceedance	_	No Exceedance	_	No	No	No	None
Di-N-Octyl Phthalate	6,200	6,200	µg/kg	1	0	0%	59		No Exceedance	<0.1	No Exceedance	_	No Exceedance		No	No	No	None
Phthalates (OC Normalized)	0,200	0,200	MB/ 1/8	-		• • • • • • • • • • • • • • • • • • • •	- 55		The Energianise		TTO Excoordance	1	TTO EXCOCUATION	1				
Bis(2-Ethylhexyl) Phthalate	47	78	mg/kg OC	9	7	78%	5.6	36.7	No Exceedance	0.1	No Exceedance	0.8	No Exceedance	0.5	No	No	No	None
Butyl benzyl Phthalate	5	64	mg/kg OC	9	1	11%	5.6	0.45	No Exceedance	1.1	No Exceedance	0.1	No Exceedance	<0.1	No	No	No	None
Dibutyl Phthalate	220	1,700	mg/kg OC	9	0	0%	4.2	-	No Exceedance	<0.1	No Exceedance		No Exceedance	-	No	No	No	None
Diethyl Phthalate	61	110	mg/kg OC	9	0	0%	4.2	-	No Exceedance	0.1	No Exceedance		No Exceedance		No	No	No	None
Dimethyl Phthalate	53	53	mg/kg OC	9	0	0%	4.2	-	No Exceedance	0.1	No Exceedance	_	No Exceedance		No	No	No	None
Di-N-Octyl Phthalate	58	4,500	mg/kg OC	-	0	0%	4.2		No Exceedance	0.1	No Exceedance	_	No Exceedance	-	No	No	No	None
,	38	4,500	ilig/ kg oc	9	U	076	4.2	-	NO Exceedance	0.1	NO Exceedance	_	NO Exceedance		NO	NO	INO	Notice
Phenols (Dry Weight)	20	20	ug/kg	10	2	20%	20	40	No Evocadance	0.7	10%	1.4	10%	1.4		No	No	None
2,4-Dimethylphenol 2-methylphenol (o-Cresol)	29 63	29 63	µg/kg µg/kg	10	0	20% 0%	60		No Exceedance No Exceedance	0.7 1.0	No Exceedance	1.4	No Exceedance	1.4	- No	No No	No	None
4-methylphenol (p-Cresol)	670	670		10	1	40%	58	59	No Exceedance	0.1	No Exceedance	0.1	No Exceedance	0.1	No	No	No	None
Pentachlorophenol	360	690	µg/kg	10	3	30%	31	70	No Exceedance	0.1	No Exceedance	0.1	No Exceedance	0.1	No	No	No	None
Phenol	420	1,200	µg/kg	10	4	40%	58	76	No Exceedance	0.1	No Exceedance	0.2	No Exceedance	0.1	No	No	No	None
		±,∠∪∪	µg/kg	10	4	40%	J0	10	NO Exceedance	0.1	INO EXCEPTION	∪.∠	INO EXCEPTION	0.1	INU	INU	INU	NOTIC
Miscellaneous Extractables (Dry Wei	540	540	pa/l-a	1	0	00/	59		No Exceedance	-O 1	No Evocadanas	ı	No Evocadanas	ı	No	No	No	None
Dibenzofuran			µg/kg	1		0%		_		<0.1	No Exceedance	-	No Exceedance	-	No	No	No	None
Hexachlorobutadiene	11	120	µg/kg	1	0	0%	6.2		No Exceedance	0.6	No Exceedance	-	No Exceedance	-	No	No	No	None
N-Nitrosodiphenylamine	28	40	μg/kg	1	0	0%	6.2	-	No Exceedance	0.2	No Exceedance	-	No Exceedance		No	No	No	None
Benzoic Acid	650	650	µg/kg	10	1	10%	600	200	No Exceedance	0.9	No Exceedance	0.3	No Exceedance	0.3	No	No	No	None
Benzyl Alcohol	57	73	µg/kg	10	1	10%	50	6.9	No Exceedance	0.9	No Exceedance	0.1	No Exceedance	0.1	No	No	No	None



	Prop	osed							Evaluation of RI	Data Results ³						Contaminant of	Concern (COC)	Selection Considerations
	Sedi	ment							RL Exceedance	Evaluation	SCO/AET Exceeda	nce Evaluation	CSL/2AET Exceeda	nce Evaluation				
Contaminant of	Clea Le	anup vel ²				Detection	Maximum Non-Detect	Maximum Detected	Frequency of RL	Maximum Exceedance	Frequency of PCUL	Maximum Exceedance	Frequency of PCUL	Maximum Exceedance	Initial COC Selection	Groundwater	Proposed Sediment	
Potential Concern ¹ (COPC)	SCO/ LAET	CSL/ 2LAET	Units	Number Samples	Number of Detections	Frequency (%)			Exceedance ⁴ (%)	Ratio ⁵ (ER)	Exceedance ⁴ (%)	Ratio ⁵ (ER)	Exceedance ⁴	Ratio ⁵ (ER)	Criteria Met ⁶ (Yes/No)	COC ⁷ (Yes/No)	COC (Yes/No)	Comments/Rationale
Miscellaneous Extractables (OC Norm	alized)		Cinto	Gampios	Dotootiono	(10)	(48/ -)	(P6/ =/	(%)	(211)	(70)	(=11)	(70)	(2.11)	(100) 110)	(100) 110)	(100) 110)	Commonto, Rationalo
Dibenzofuran	15	58	mg/kg OC	9	5	56%	5.6	9.4	No Exceedance	0.4	No Exceedance	0.6	No Exceedance	0.2	No	No	No	None
Hexachlorobutadiene	3.9	6.2	mg/kg OC	9	0	0%	0.6		No Exceedance	0.2	No Exceedance	-	No Exceedance		No	No	No	None
N-Nitrosodiphenylamine	11	11	mg/kg OC	9	1	11%	0.6	1	No Exceedance	0.1	No Exceedance	0.1	No Exceedance	0.1	No	No	No	None
Pesticides	•																	
4,4'-DDD	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	-	No Exceedance		No	No	No	None
4,4'-DDE	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	/	No Exceedance		No	No	No	None
4,4'-DDT	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	-	No Exceedance		No	No	No	None
Total DDT (4,4 isomers)	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	-	No Exceedance		No	No	No	None
Aldrin	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	-	No Exceedance		No	No	No	None
Total Chlordane ¹⁰	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	-	No Exceedance		No	No	No	None
Dieldrin	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	-	No Exceedance		No	No	No	None
Heptachlor	NE	NE	µg/kg	2	0	0%	1	-	No Exceedance	-	No Exceedance	_	No Exceedance		No	No	No	None
Polychlorinated Biphenyls (PCBs) (Dry	Weight)																	
Total PCBs (Sum of Aroclors)	0.13	1	mg/kg	3	2	67%	0.021	0.168	No Exceedance	0.2	33%	1.3	No Exceedance	0.2	Yes	-	Yes	Retained as a COC
Polychlorinated Biphenyls (PCBs) (OC	Normalized)	-	-	-	-					-			-	-				
Total PCBs (Sum of Aroclors)	12	65	mg/kg OC	10	5	50%	3	37	No Exceedance	0.3	40%	3.1	No Exceedance	0.6	Yes	-	Yes	Retained as a COC

RL = Reporting Limit

NE = Not Established

mg/kg = milligrams per kilogram

μg/kg = micrograms per kilogram

OC = organic carbon normalized
-- = not applicable

Bold indicate satisfaction of initial COC selection criteria or consideration of other selection criteria.

Yellow shading indicates analyte is identified as a COC based on both satisfaction of initial selection criteria and consideration of other selection criteria, or on consideration of other selection criteria.



¹ Contaminants of potential concern (COPCs) were established for the RI based on a review of previous environmental studies. Previous sediment study results are summarized in Table C-1 (Appendix C).

 $^{^{2}}$ Proposed sediment cleanup levels for the protection of benthic organisms are referenced from Table 1.

³ The sediment data used for this RI consists of samples obtained by GeoEngineers in 2008 in general accordance with the RI/FS Work Plan (GeoEngineers 2008a) as well as data collected by others to support dredge material suitability determination. Sediment characterization results are summarized in Table C-1 (Appendix C).

⁴ Number of samples with analyte detected or non-detect at a concentration greater than PCUL / total number of samples analyzed for analyte.

⁵ Exceedance Ratio (max) = ratio of maximum detected or non-detect concentration divided by the Screening Level

⁶ Initial contaminant of concern (COC) selection criteria is met if exceedance frequency is greater or equal to 10 percent or if the exceedance ratio is greater than 2.

⁷ Total LPAHs are the total of naphthalene, acenaphthylene, acenaphthene, fluorene, phenanthrene and anthracene; 2-methylnapthalene is not included in the sum of LPAHs.

⁸ Total HPAHs are the total of fluoranthene, pyrene, benzo(a)anthracene, chrysene, benzofluoranthenes, benzo(a)pyrene, indeno(1,2,3-c-d)pyrene, dibenzo(a,h)anthracene and benzo(g,h,i)perylene.

 $^{^{9}}$ Total benzofluoranthenes represents the sum of concentrations of the b, j, and k isomers.

¹⁰ Total chlordane represents the sum of concentrations of alpha-Chlordane (cis), gamma-Chlordane (trans), cis-nonachlor, trans-nonachlor and oxychlordane.

Table 9
Summary Statistics and Evaluation of Sediment Contaminants of Concern - Protection of Human Health and Higher Trophic Level Ecological Receptors

Dakota Creek Industries

Anacortes, Washington

							Evaluation of R	I Data Baculta ³		_		Conton	ninant of Canaa	rn (COC) Selection Considerations
							Evaluation of R		Frankration	DOUB Francisco	- FI	Contain	illinant of Conce	(COC) Selection Considerations
								RL Exceedance	Evaluation Maximum	PCUL Exceedance	Maximum			
Ocutomiu out of	Proposed					Maximum	Maximum	Frequency of RL	Exceedance	Frequency of PCUL	Exceedance	Initial COC	Proposed	
Contaminant of	Sediment		l		Detection -	Non-Detect	Detected	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	Selection	Sediment	
Potential Concern ¹	Cleanup		Number	Number of	Frequency	Concentration	Concentration	(%)	(ER)	(%)	(ER)	Criteria Met ⁶	COC	O manage of the Continue of th
(COPC)	Level ²	Units	Samples	Detections	(%)	(µg/L)	(µg/L)	(70)	(LIV)	(70)	(LK)	(Yes/No)	(Yes/No)	Comments/Rationale
Metals	1 44		1 4-	1 4- 1	200/	1 40	222			2001			•	
Arsenic	11	mg/kg	17	15	88%	10	300	No Exceedance	0.9	29%	27.3	Yes	Yes	Retained as a COC
Cadmium	0.8	mg/kg	17	11	65%	0.6	1	No Exceedance	0.8	6%	1.5	No	No	None
Chromium	6,900,000	mg/kg	17	17	100%	-	55	No Exceedance		No Exceedance	<0.1	No	No	None
Copper	180,000	mg/kg	17	17	100%		3,870	No Exceedance	-	No Exceedance	<0.1	No	No	None
Lead	21	mg/kg	17	17	100%	_	939	No Exceedance	-	29%	44.7	Yes	Yes	Retained as a COC
Mercury	0.2	mg/kg	17	10	59%		17.8	No Exceedance	-	35%	89.0	Yes	Yes	Retained as a COC
Nickel	NE	mg/kg	9	9	100%		36	No Exceedance	_	No Exceedance		No	No	None
Silver	23,000	mg/kg	17	10	59%	0.9	1	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Zinc	1,400,000	mg/kg	17	17	100%		974	No Exceedance	-	No Exceedance	<0.1	No	No	None
Organometallic Compounds														
Tributyltin, bulk	73	μg/kg					+		-	-				None
Interstitial Tributyltin, porewater	0.15	μg/L	5	4	80%	0.08	0.45	No Exceedance	0.5	20%	3.0	Yes	Yes	Retained as a COC
Low Molecular Weight Polycyclic Aromat	tic Hydrocarbons (LF	PAHs)	•			•					•	•		
2-Methylnaphthalene	16,000,000	µg/kg	13	8	62%	20	4,100	No Exceedance	<0.1	No Exceedance	<0.1	No	Yes	
Acenaphthene	240,000,000	µg/kg	13	9	69%	20	230	No Exceedance	<0.1	No Exceedance	<0.1	No	Yes	
Acenaphthylene	240,000,000	µg/kg	13	9	69%	20	250	No Exceedance	<0.1	No Exceedance	<0.1	No	Yes	Retained as a COC based on the results
Anthracene	1,200,000,000	µg/kg	13	13	100%	-	1,900	No Exceedance	_	No Exceedance	<0.1	No	Yes	of the benthic organism evaluation (Table
Fluorene	160,000,000	µg/kg	13	12	92%	20	742	No Exceedance	<0.1	No Exceedance	<0.1	No	Yes	8).
Naphthalene	79,000,000	µg/kg	13	12	92%	20	3,060	No Exceedance	<0.1	No Exceedance	<0.1	No	Yes	
Phenanthrene	1,200,000,000	µg/kg	13	13	100%		5,070	No Exceedance		No Exceedance	<0.1	No	Yes	
High Molecular Weight Polycyclic Aroma							3,313				V		100	I
Benzo(a)anthracene	NE	µg/kg	13	13	100%		2,600	No Exceedance		No Exceedance		No	Yes	
Benzo(a)pyrene	cPAH TEQ	µg/kg	13	13	100%	-	4,100	No Exceedance	_	cPAH TEQ	cPAH TEQ	Yes	Yes	
Total Benzofluoranthenes ⁸	NE NE	µg/kg	13	13	100%		3,300	No Exceedance		No Exceedance	-	No	Yes	
Benzo(g,h,i)perylene	120,000,000	µg/kg	13	11	85%	20	4,850	No Exceedance	<0.1	No Exceedance	<0.1	No	Yes	Retained as a COC based on the results
Chrysene	NE	µg/kg	13	13	100%	-	4,150	No Exceedance		No Exceedance		No	Yes	of the benthic organism evaluation (Table
Dibenzo(a,h)anthracene	NE	µg/kg	13	10	77%	6.1	3,060	No Exceedance	<0.1	No Exceedance		No	Yes	8).
Fluoranthene	160,000,000	µg/kg	13	13	100%	-	11,000	No Exceedance	-	No Exceedance	<0.1	No	Yes	
Indeno(1,2,3-c,d)pyrene	NE	µg/kg	13	13	100%	-	2,050	No Exceedance		No Exceedance		No	Yes	
Pyrene	120,000,000	µg/kg	13	13	100%		7,500	No Exceedance		No Exceedance	<0.1	No	Yes	
Carcinogenic PAHs (cPAHs)										-				
Total cPAH TEQ ⁹ (ND=0.5 RL)	21	µg/kg	13	13	100%		5,094	No Exceedance		92%	242.5	Yes	Yes	Retained as a COC
Chlorinated Hydrocarbons	1		1	<u> </u>		1	, -			<u> </u>				
1,2,4-Trichlorobenzene	140,000	µg/kg	10	0	0%	9.9	_	No Exceedance	<0.1	No Exceedance		No	No	None
1.2-Dichlorobenzene	370,000,000	µg/kg	10	3	30%	9.9	18	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
1,4-Dichlorobenzene	780,000	μg/kg	10	2	20%	9.9	14	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
_,	. 55,555	~0/ '`0		1			_ · _ ·		- · -		, v.±			



							Evaluation of R	I Data Results ³				Contan	ninant of Conce	rn (COC) Selection Considerations
Contaminant of Potential Concern¹ (COPC) Phthalates Bis(2-Ethylhexyl) Phthalate	Proposed Sediment Cleanup Level ²	Units µg/kg	Number Samples	Number of Detections	Detection Frequency (%)	Maximum Non-Detect Concentration (µg/L)	Maximum Detected Concentration (μg/L)	RL Exceedance Frequency of RL Exceedance (%)	Evaluation Maximum Exceedance Ratio ⁵ (ER)	PCUL Exceedance Frequency of PCUL Exceedance (%)	Example Example Exceedance Ratio (ER)	Initial COC Selection Criteria Met ⁶ (Yes/No)	Proposed Sediment COC (Yes/No)	Comments/Rationale None
Butyl benzyl Phthalate	2,100,000	µg/kg	10	1	10%	20	12	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Dibutyl Phthalate	410,000,000	µg/kg	10	0	0%	59	-	No Exceedance	<0.1	No Exceedance		No	No	None
Diethyl Phthalate	3,100,000,000	µg/kg	10	0	0%	59	_	No Exceedance	<0.1	No Exceedance		No	No	None
Dimethyl Phthalate	NE NE	µg/kg	10	0	0%	59		No Exceedance		No Exceedance		No	No	None
Di-N-Octyl Phthalate	41,000,000	µg/kg	10	0	0%	59	_	No Exceedance	<0.1	No Exceedance		No	No	None
Phenols		10, 0				I.						1		
2,4-Dimethylphenol	82,000,000	μg/kg	10	2	20%	20	40	No Exceedance	<0.1	No Exceedance	<0.1		No	None
2-methylphenol (o-Cresol)	200,000,000	µg/kg	10	0	0%	60	-	No Exceedance	<0.1	No Exceedance	_	No	No	None
4-methylphenol (p-Cresol)	390,000,000	μg/kg	10	4	40%	58	59	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Pentachlorophenol	10,000,000	μg/kg	10	3	30%	31	70	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Phenol	1,200,000,000	μg/kg	10	4	40%	58	76	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Miscellaneous Extractables														
Dibenzofuran	4,100,000	μg/kg	10	5	50%	20	130	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Hexachlorobutadiene	52,000	μg/kg	10	0	0%	20	-	No Exceedance	<0.1	No Exceedance	-	No	No	None
N-Nitrosodiphenylamine	830,000	µg/kg	10	1	10%	20	6.1	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Benzoic Acid	16,000,000,000	µg/kg	10	1	10%	600	200	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Benzyl Alcohol	410,000,000	µg/kg	10	1	10%	50	6.9	No Exceedance	<0.1	No Exceedance	<0.1	No	No	None
Polychlorinated Biphenyls (PCBs)														
Total PCBs (Sum of Aroclors)	0.0035	mg/kg	13	7	54%	0.021	0.362	38%	6.0	54%	103.4	Yes	Yes	Retained as a COC
Dioxins and Furans														
Total Dioxin/Furan TEQ ¹⁰ (ND=0.5 RL)	5	mg/kg OC	18	18	100%	-	148.94	No Exceedance		33%	29.8	Yes	Yes	Retained as a COC

RL = Reporting Limit

NE = Not Established

mg/kg = milligrams per kilogram

μg/kg = micrograms per kilogram

OC = organic carbon normalized

-- = not applicable

Bold indicate satisfaction of initial COC selection criteria or consideration of other selection criteria.

Yellow shading indicates analyte is identified as a COC based on both satisfaction of initial selection criteria and consideration of other selection criteria, or on consideration of other selection criteria alone.



¹ Contaminants of potential concern (COPCs) were established for the RI based on a review of previous environmental studies. Previous sediment study results are summarized in Table C-2 (Appendix C).

² Proposed sediment cleanup levels for the protection of human health and higher trophic level ecological receptors are referenced from Table 2.

³ The sediment data used for this RI consists of samples obtained by GeoEngineers in 2008 in general accordance with the RI/FS Work Plan (GeoEngineers 2008a) as well as data collected by others to support dredge material suitability determination. Sediment characterization results are summarized in Table C-1 (Appendix C).

⁴ Number of samples with analyte detected or non-detect at a concentration greater than PCUL / total number of samples analyzed for analyte.

⁵ Exceedance Ratio (max) = ratio of maximum detected or non-detect concentration divided by the Screening Level

⁶ Initial contaminant of concern (COC) selection criteria is met if exceedance frequency is greater or equal to 10 percent or if the exceedance ratio is greater than 2.

⁷ Groundwater contaminants of concern (COCs) are presented in Table 10.

⁸Total benzofluoranthenes represents the sum of concentrations of the b, j, and k isomers.

⁹ Total cPAH Toxic Equivalency Quotients (TEQs) were calculated using Toxicity Equivalency Factors (TEFs) values referenced from MTCA Table 708.2 (WAC 173-340-900).

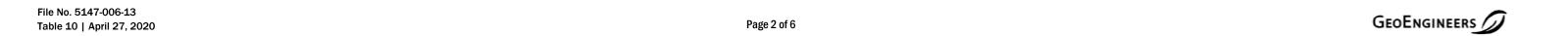
¹⁰ Total dioxin and furan TEQs were calculated using United States Environmental Protection Agency (USEPA) TEF values for human health (EPA, 2003).

Summary Statistics and Evaluation of Groundwater Contaminants of Concern

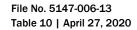
							Evaluation of R	I Data Results ³					Contar	ninant of Conce	rn (COC) Selection Considerations
								RL Exceedance	Evaluation	PCUL Exceedance	ce Evaluation				
Contaminant of Potential Concern ¹ (COPC)	Proposed Groundwater Cleanup Level ² (PCUL)	Units	Number Samples	Number of Detections	Detection Frequency (%)	Maximum Non-Detect Concentration (μg/L)	Maximum Detected Concentration (µg/L)	Frequency of RL Exceedance ⁴ (%)	Maximum Exceedance Ratio ⁵ (ER)	Frequency of PCUL Exceedance ⁴ (%)	Maximum Exceedance Ratio ⁵ (ER)	Proposed Sediment COC ⁶ (Yes/No)	Initial Groundwater COC Selection Criteria Met ⁷ (Yes/No)	Proposed Groundwater COC (Yes/No)	Comments/Rationale
Total Metals												•			
Arsenic	8	µg/L	53	28	53%	16	153	11 %	2.0	38%	19.1	Yes	Yes	Yes	Retained as a COC
Cadmium	7.9	µg/L	18	0	0%	8	-	22%	1.0	No Exceedance	-	No	No	No	RL for non-detect only slightly exceeded the PCUL and was not detected in other samples analyzed.
Total Chromium	50	μg/L	18	6	33%	20	16	No Exceedance	0.4	No Exceedance	0.3	No	No	No	None
Copper	20	µg/L	30	12	40%	20	20	No Exceedance	1.0	No Exceedance	1.0	No	No	No	None
Lead	10	μg/L	18	2	11%	5	2	No Exceedance	0.5	No Exceedance	0.2	Yes	No	No	None
Mercury	0.025	μg/L	30	14	47%	0.2	0.037	53%	8.0	7%	1.5	Yes	No	No	RL for non-detect only slightly exceeded the PCUL and was not detected in other samples analyzed.
Nickel	8.2	µg/L	53	28	53%	20	27	8%	2.4	25%	3.3	Yes	Yes	Yes	Retained as a COC
Silver	10	μg/L	24	0	0%	10	-	No Exceedance	1.0	No Exceedance	-	No	No	No	None
Zinc	160	μg/L	45	5	11%	50	30	No Exceedance	0.3	No Exceedance	0.2	No	No	No	None
Dissolved Metals															
Arsenic	8	μg/L	47	20	43%	16	143	15%	2.0	16%	17.9	Yes	Yes	Yes	Retained as a COC
Cadmium	7.9	µg/L	12	0	0%	8		25%	1.0	No Exceedance	-	No	No	No	RL for non-detect only slightly exceeded the PCUL and was not detected in other samples analyzed.
Total Chromium	50	μg/L	12	0	0%	20	12	No Exceedance	0.4	No Exceedance	0.2	No	No	No	None
Copper	20	μg/L	24	4	17%	20	17	No Exceedance	1.0	No Exceedance	0.9	No	No	No	None
Lead	10	μg/L	12	0	0%	5	-	No Exceedance	0.5	No Exceedance		Yes	No	No	None
Mercury	0.025	µg/L	24	0	0%	0.13		67%	5.2	No Exceedance		Yes	No	No	RL for non-detect only slightly exceeded the PCUL and was not detected in other samples analyzed.
Nickel	8.2	μg/L	47	18	38%	10	27	9%	1.2	14%	3.3	Yes	Yes	Yes	Retained as a COC
Silver	10	µg/L	24	0	0%	10	-	No Exceedance	1.0	No Exceedance		No	No	No	None
Zinc	160	μg/L	24	1	4%	50	28	No Exceedance	0.3	No Exceedance	0.2	No	No	No	None
Petroleum Hydrocarbons															
Gasoline-Range	800	µg/L	29	0	0%	100		No Exceedance	0.1	No Exceedance		No	No	No	None
Diesel-Range	500	μg/L	29	1	3%	256	180	No Exceedance	0.5	No Exceedance	0.4	No	No	No	None
Heavy Oil-Range	500	μg/L	29	0	0%	500	-	No Exceedance	1.0	No Exceedance		No	No	No	None
BETX Compounds															
Benzene	2	µg/L	6	0	0%	2	-	100%	1.3	No Exceedance	-	No	No	No	Benzene not historically detected and RL for previous events below the PCUL.
Ethylbenzene	31	μg/L	6	0	0%	2	-	No Exceedance	0.1	No Exceedance		No	No	No	None
Toluene	130	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Xylenes	630	μg/L	6	0	0%	4		No Exceedance	<0.1	No Exceedance		No	No	No	None



							Evaluation of R	I Data Results ³					Contan	ninant of Conce	rn (COC) Selection Considerations
								RL Exceedance	Evaluation	PCUL Exceedan	ce Evaluation				(
												1	Initial		
	Proposed					Maximum	Maximum	Frequency	Maximum	Frequency	Maximum	Proposed	Groundwater	Proposed	
Contaminant of	Groundwater				Detection	Non-Detect	Detected	of RL	Exceedance	of PCUL	Exceedance	Sediment	COC Selection	Groundwater	
Potential Concern ¹	Cleanup Level ²		Number	Number of	Frequency	Concentration	Concentration	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	COC ₆	Criteria Met ⁷	COC	
(COPC)	(PCUL)	Units	Samples	Detections	(%)	(μg/L)	(µg/L)	(%)	(ER)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
Volatile Organic Compounds (VOCs)															
1,1,1,2-Tetrachloroethane	74	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance	_	No	No	No	None
1,1,1-Trichloroethane	12,000	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
															Other VOCs either were not detected or
1,1,2,2-Tetrachloroethane	1	µg/L	6	0	0%	2		100 %	2.0	No Exceedance		No	No	No	detected at concentrations greater than
															the PCUL.
1,1,2-trichloro-1,2,2- trifluoroethane	2,400	ud/l	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No		No	None
(CFC113)	2,400	µg/L	0	U	0%	2		NO Exceedance	\U.1	No exceedance	_	INO	No	NO	Notie
															Other VOCs either were not detected or
1,1,2-Trichloroethane	1	μg/L	6	0	0%	2		100 %	2.0	No Exceedance		No	No	No	detected at concentrations greater than
															the PCUL.
1,1-Dichloroethane	110	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance	-	No	No	No	None
1,1-Dichloroethene	280	μg/L	6	0	0%	1		No Exceedance	<0.1	No Exceedance	_	No	No	No	None
1,1-Dichloropropene	NE	μg/L	6	0	0%	2		No Exceedance	-	No Exceedance	-	No	No	No	None
1,2,3-Trichlorobenzene	NE	μg/L	6	0	0%	5		No Exceedance	-	No Exceedance		No	No	No	None
1,2,3-Trichloropropane	NE	μg/L	6	0	0%	3		No Exceedance	-	No Exceedance		No	No	No	None
1,2,4-Trichlorobenzene	5.0	μg/L	6	0	0%	5		No Exceedance	1.0	No Exceedance		No	No	No	None
1,2,4-Trimethylbenzene	61.0	μg/L	6	0	0%	2	-	No Exceedance	<0.1	No Exceedance		No	No	No	None
1,2-Dibromo-3-chloropropane	NE	μg/L	6	0	0%	10		No Exceedance	-	No Exceedance		No	No	No	None
1,2-Dibromoethane (EDB)	2.7	μg/L	6	0	0%	2		No Exceedance	0.7	No Exceedance		No	No	No	None
1,2-Dichlorobenzene	800	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
1,2-Dichloroethane (EDC)	42	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
1,2-Dichloropropane	3.1	μg/L	6	0	0%	2		No Exceedance	0.6	No Exceedance	-	No	No	No	None
1,3,5-Trimethylbenzene	NE	μg/L	6	0	0%	2		No Exceedance	-	No Exceedance		No	No	No	None
1,3-Dichlorobenzene	2	μg/L	6	0	0%	2	-	No Exceedance	1.0	No Exceedance		No	No	No	None
1,3-Dichloropropane	NE	μg/L	6	0	0%	2	-	No Exceedance		No Exceedance		No	No	No	None
1,4-Dichlorobenzene	22	μg/L	6	0	0%	2		No Exceedance	0.1	No Exceedance		No	No	No	None
2,2-Dichloropropane	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
2-Butanone (MEK)	3,800,000	μg/L	6	0	0%	10		No Exceedance	<0.1	No Exceedance		No	No	No	None
2-Chloroethyl Vinyl Ether	NE	μg/L	6			-	-		-	_			-	No	None
2-Chlorotoluene	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
2-Hexanone	NE	μg/L	6	0	0%	10		No Exceedance		No Exceedance		No	No	No	None
4-Chlorotoluene	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
4-Methyl-2-Pentanone	1,000,000	ug/l	6	0	0%	10		No Exceedance	<0.1	No Exceedance		No	No	No	None
(Methyl Isobutyl Ketone)	1,000,000	μg/L	0	U	0%	10		NO Exceedance	\0.1	No exceedance		No	No	No	Notie
Acetone	32,000,000	μg/L	6	0	0%	25		No Exceedance	<0.1	No Exceedance		No	No	No	None
Acrolein	50	μg/L	6	0	0%	50	-	No Exceedance	1.0	No Exceedance		No	No	No	None
Acrylonitrile	1.0	μg/L	6	0	0%	1		No Exceedance	1.0	No Exceedance		No	No	No	None
Bromobenzene	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
Bromochloromethane	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
Bromoform	12	μg/L	6	0	0%	2		No Exceedance	0.2	No Exceedance		No	No	No	None
Bromomethane	28	μg/L	6	0	0%	2		No Exceedance	0.1	No Exceedance		No	No	No	None
Carbon Disulfide	870	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Carbon Tetrachloride	1.00	μg/L	6	0	0%	1		No Exceedance	1.4	No Exceedance		No	No	No	None
Chlorobenzene	200	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Chloroethane	40,400	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Chloroform	12	μg/L	6	0	0%	2		No Exceedance	0.2	No Exceedance		No	No	No	None
Chloromethane	340.0	μg/L	6	0	0%	2	-	No Exceedance	<0.1	No Exceedance		No	No	No	None
Cis-1,2-Dichloroethene	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None

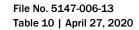


							Evaluation of F	I Data Results ³					Contar	ninant of Conce	rn (COC) Selection Considerations
								RL Exceedance	Evaluation	PCUL Exceedance	ce Evaluation				
								_		_			Initial		
	Proposed					Maximum	Maximum	Frequency	Maximum	Frequency	Maximum	Proposed	Groundwater	Proposed	
Contaminant of	Groundwater				Detection	Non-Detect	Detected	of RL	Exceedance	of PCUL	Exceedance	Sediment	COC Selection	Groundwater	
Potential Concern ¹	Cleanup Level ²		Number	Number of	Frequency	Concentration	Concentration	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	COC ⁶	Criteria Met	coc	
(COPC)	(PCUL)	Units	Samples	Detections	(%)	(μg/L)	(µg/L)	(%)	(ER)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
Cis-1,3-Dichloropropene	NE	µg/L	6	0	0%	2	-	No Exceedance	-	No Exceedance	-	No	No	No	None
Dibromochloromethane	2.2	µg/L	6	0	0%	2		No Exceedance	0.9	No Exceedance		No	No	No	None
Dibromomethane	NE	µg/L	6	0	0%	2		No Exceedance	-	No Exceedance		No	No	No	None
Dichlorobromomethane	2.8	µg/L	6	0	0%	2		No Exceedance	0.7	No Exceedance		No	No	No	None
Dichlorodifluoromethane															Not a COC. Other VOCs either were not
(CFC 12)	12	μg/L							_	-	_		-	No	detected or detected at concentrations
The cold cold to disc.	F 0	(I	0	0	20/			N. E	1.0	N. E d		NI -	N.	NI-	greater than the PCUL.
Hexachlorobutadiene	5.0	µg/L	6	0	0%	5		No Exceedance	1.0	No Exceedance		No	No	No	None
Isopropylbenzene	1,600	µg/L	6	0	0%	2	-	No Exceedance	<0.1	No Exceedance	-	No	No	No	None
Methyl lodide	NE C 100	µg/L	6	0	0%	2		No Exceedance	10.4	No Exceedance		No	No	No	None
Methyl t-Butyl Ether (MTBE)	6,100	µg/L	6	0	0%	2	-	No Exceedance	<0.1	No Exceedance	'	No	No	No	None
Methylene Chloride	100	µg/L	6	0	0%	5		No Exceedance	0.1	No Exceedance		No	No	No	None
Naphthalene	89	µg/L	6	0	0%	5		No Exceedance	0.1	No Exceedance		No	No	No	None
n-Butylbenzene	NE	μg/L	6	0	0%	2		No Exceedance	-	No Exceedance		No	No	No	None
n-Propylbenzene	4,900	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
p-Isopropyltoluene	NE	μg/L	6	0	0%	2	-	No Exceedance	-	No Exceedance		No	No	No	None
sec-Butylbenzene	NE	µg/L	6	0	0%	2	'	No Exceedance	-	No Exceedance		No	No	No	None
Styrene	18,000	μg/L	6	0	0%	2	-	No Exceedance	<0.1	No Exceedance		No	No	No	None
tert-Butylbenzene	NE	μg/L	6	0	0%	2	-	No Exceedance	-	No Exceedance		No	No	No	None
Tetrachloroethene (PCE)	2.9	μg/L	6	0	0%	1		No Exceedance	0.3	No Exceedance		No	No	No	None
Trans-1,2-Dichloroethene	250	μg/L	6	0	0%	2	-	No Exceedance	<0.1	No Exceedance		No	No	No	None
Trans-1,3-Dichloropropene	NE	μg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
Trans-1,4-Dichloro-2-butene	NE	μg/L	6	0	0%	5	-	No Exceedance		No Exceedance		No	No	No	None
Trichloroethene (TCE)	1.0	μg/L	6	0	0%	1		No Exceedance	1.0	No Exceedance		No	No	No	None
Trichlorofluoromethane (CFC 11)	260	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Vinyl Acetate	17,000	μg/L	6	0	0%	5	-	No Exceedance	<0.1	No Exceedance	-	No	No	No	None
Vinyl Chloride	1.0	μg/L	6	0	0%	1	_	No Exceedance	1.0	No Exceedance		No	No	No	None
Semi-Volatile Organic Compounds (S\	/OCs)								•						
1,2,4-Trichlorobenzene	1.0	μg/L	6	0	0%	2		No Exceedance	2.0	No Exceedance	-	No	No	No	None
1,2-Dichlorobenzene	800	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance	-	No	No	No	None
1,3-Dichlorobenzene	2	μg/L	6	0	0%	2	-	No Exceedance	1.0	No Exceedance	-	No	No	No	None
1,4-Dichlorobenzene	22	μg/L	6	0	0%	2		No Exceedance	0.1	No Exceedance		No	No	No	None
2,2'-0xybis[1-chloropropane]	37	μg/L	6	0	0%	2	_	No Exceedance	0.1	No Exceedance		No	No	No	None
2,4,5-Trichlorophenol	600	μg/L	6	0	0%	5		No Exceedance	<0.1	No Exceedance		No	No	No	None
2,4,6-Trichlorophenol	5.0	μg/L	6	0	0%	5		No Exceedance	1.0	No Exceedance	_	No	No	No	None
2,4-Dichlorophenol	10	μg/L	6	0	0%	5		No Exceedance	0.5	No Exceedance	-	No	No	No	None
2,4-Dimethylphenol	97	μg/L	6	0	0%	3		No Exceedance	<0.1	No Exceedance	-	No	No	No	None
2,4-Dinitrophenol	100	μg/L	6	0	0%	25		No Exceedance	0.3	No Exceedance		No	No	No	None
2,4-Dinitrotoluene	5.0	μg/L	6	0	0%	5		No Exceedance	1.0	No Exceedance		No	No	No	None
2,6-Dinitrotoluene	NE	μg/L	6	0	0%	5		No Exceedance	-	No Exceedance		No	No	No	None
2-Chloronaphthalene	100	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
2-Chlorophenol	17	µg/L	6	0	0%	2		No Exceedance	0.1	No Exceedance	-	No	No	No	None
2-Nitroaniline	NE	µg/L	6	0	0%	5		No Exceedance		No Exceedance	_	No	No	No	None
2-Nitrophenol	NE NE	μg/L	6	0	0%	5		No Exceedance		No Exceedance		No	No	No	None
3,3'-Dichlorobenzidine	5.0	μg/L	6	0	0%	5		No Exceedance	1.0	No Exceedance		No	No	No	None
S,S DIGINGIONOLIZIGNIO	0.0	M8/ ₽		Ŭ	U /0		-	110 Exocodanos	1.0	THE EXOCOGRAPHOC		110	110	110	None



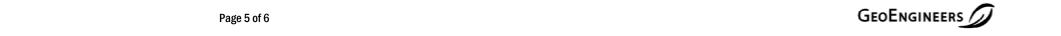


							Evaluation of R	I Data Results ³					Contar	ninant of Conce	rn (COC) Selection Considerations
								RL Exceedance	Evaluation	PCUL Exceedance	ce Evaluation				
Contaminant of Potential Concern ¹ (COPC)	Proposed Groundwater Cleanup Level ² (PCUL)	Units	Number Samples	Number of Detections	Detection Frequency (%)	Maximum Non-Detect Concentration (μg/L)	Maximum Detected Concentration (μg/L)	Frequency of RL Exceedance ⁴ (%)	Maximum Exceedance Ratio ⁵ (ER)	Frequency of PCUL Exceedance ⁴ (%)	Maximum Exceedance Ratio ⁵ (ER)	Proposed Sediment COC ⁶ (Yes/No)	Initial Groundwater COC Selection Criteria Met ⁷ (Yes/No)	Proposed Groundwater COC (Yes/No)	Comments/Rationale
4,6-Dinitro-2-methylphenol	10	µg/L	6	0	0%	2	(P8/ =/	No Exceedance	0.2	No Exceedance	-	No	No	No	None
4-Bromophenyl-phenylether	NE	µg/L	6	0	0%	2		No Exceedance	_	No Exceedance	_	No	No	No	None
4-Chloro-3-methylphenol	36	µg/L	6	0	0%	5	_	No Exceedance	0.1	No Exceedance		No	No	No	None
4-Chloroaniline	NE	µg/L	6	0	0%	5		No Exceedance	_	No Exceedance		No	No	No	None
4-Chlorophenyl-phenylether	NE	µg/L	6	0	0%	2		No Exceedance	_	No Exceedance		No	No	No	None
4-Nitroaniline	NE	µg/L	6	0	0%	5		No Exceedance	-	No Exceedance	-	No	No	No	None
4-Nitrophenol	NE	µg/L	6	0	0%	5		No Exceedance	- /	No Exceedance	-	No	No	No	None
Benzoic acid	NE	µg/L	6	0	0%	50		No Exceedance	-	No Exceedance	-	No	No	No	None
Benzyl alcohol	NE	µg/L	6	0	0%	5	_	No Exceedance	-	No Exceedance	-	No	No	No	None
bis(2-Chloroethoxy)methane	NE	µg/L	6	0	0%	2		No Exceedance	-	No Exceedance		No	No	No	None
bis(2-chloroethyl)ether	1.0	μg/L	6	0	0%	0.54		No Exceedance	0.5	No Exceedance	-	No	No	No	None
bis(2-Ethylhexyl)phthalate	1.0	µg/L	6	0	0%	2	_	100%	2.0	No Exceedance	_	No	No	No	Other VOCs either were not detected or
Butylbenzylphthalate	1.0		6	0	0%	2		100%	2.0	No Exceedance		No	No	No	detected at concentrations greater than
		µg/L		-					2.0						the PCUL.
Carbazole	NE	µg/L	6	0	0%	2		No Exceedance	-	No Exceedance		No	No	No	None
Dibenzofuran	NE	µg/L	6	0	0%	2		No Exceedance	-	No Exceedance		No	No	No	None
Diethylphthalate	200	µg/L	6	0	0%	2	1	No Exceedance	<0.1	No Exceedance	-	No	No	No	None
Dimethylphthalate	600	µg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance	-	No	No	No	None
Di-n-butylphthalate	8	µg/L	6	0	0%	2		No Exceedance	0.3	No Exceedance	-	No	No	No	None
Di-n-octylphthalate	NE	µg/L	6	0	0%	2		No Exceedance		No Exceedance	-	No	No	No	None Other VOCs either were not detected or
Hexachlorobenzene	1.0	μg/L	6	0	0%	2	-	100%	2.0	No Exceedance	-	No	No	No	detected at concentrations greater than
Hexachlorobutadiene	1.0	μg/L	6	0	0%	2	-	100%	2.0	No Exceedance		No	No	No	the PCUL.
Hexachlorocyclopentadiene	5.0	μg/L	6	0	0%	5	_	No Exceedance	1.0	No Exceedance	_	No	No	No	None
Hexachloroethane	2.0	μg/L	6	0	0%	2	-	No Exceedance	1.0	No Exceedance	-	No	No	No	None
Isophorone	110	μg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Nitrobenzene	100	μg/L	6	0	0%	2	-/	No Exceedance	<0.1	No Exceedance		No	No	No	None
n-Nitroso-di-n-propylamine	5.0	μg/L	6	0	0%	5		No Exceedance	1.0	No Exceedance		No	No	No	None
n-Nitrosodiphenylamine	1.0	µg/L	6	0	0%	2	-	100%	2.0	No Exceedance	_	No	No	No	Other VOCs either were not detected or detected at concentrations greater than the PCUL.
o-Cresol (2-Methylphenol)	NE	µg/L	6	0	0%	2		No Exceedance		No Exceedance		No	No	No	None
p-Cresol (4-Methylphenol)	NE NE	μg/L	6	0	0%	2	_	No Exceedance	_	No Exceedance		No	No	No	None
Pentachlorophenol	5.0	μg/L	6	0	0%	5	-	No Exceedance	1.0	No Exceedance		No	No	No	None
Phenol	70,000	µg/L	6	0	0%	2		No Exceedance	<0.1	No Exceedance		No	No	No	None
Non-Carcinogenic Polycyclic Aromati							1				1		1		1
1-Methylnaphthalene	NE NE	µg/L	30	0	0%	0.095	_	No Exceedance		No Exceedance	-	No	No	No	None
2-Methylnaphthalene	NE	µg/L	30	0	0%	0.095		No Exceedance		No Exceedance	-	No	No	No	None
Acenaphthene	5	µg/L	30	2	7%	0.095	0.03	No Exceedance	<0.1	No Exceedance	<0.1	No	No	No	None
Acenaphthylene	NE	µg/L	30	2	7%	0.095	0.3	No Exceedance		No Exceedance	-	No	No	No	None
Anthracene	2	µg/L	30	3	10%	0.095	0.05	No Exceedance	<0.1	No Exceedance	<0.1	No	No	No	None
Benzo[g,h,i]perylene	NE	µg/L	30	4	13%	0.095	0.02	No Exceedance		No Exceedance		No	No	No	None
Fluoranthene	2.2	µg/L	30	2	7%	0.095	0.07	No Exceedance	<0.1	No Exceedance	<0.1	No	No	No	None
Fluorene	4	μg/L	30	2	7%	0.095	0.09	No Exceedance	<0.1	No Exceedance	<0.1	No	No	No	None
Naphthalene	89	μg/L	30	2	7%	0.100	0.03	No Exceedance	<0.1	No Exceedance	<0.1	No	No	No	None
Phenanthrene	NE NE	μg/L	30	3	10%	0.095	0.14	No Exceedance	-	No Exceedance	-	No	No	No	None
Pyrene	2	μg/L	30	4	13%	0.095	0.07	No Exceedance	<0.1	No Exceedance	<0.1	No	No	No	None





							Evaluation of R	I Data Results ³					Contar	ninant of Concer	n (COC) Selection Considerations
								RL Exceedance	e Evaluation	PCUL Exceedance	ce Evaluation				(000)
							-	NE EXOCOGUNIO	Evaluation	1 OOL EXOCCULIN	L L L L L L L L L L L L L L L L L L L		Initial		
	Proposed					Maximum	Maximum	Frequency	Maximum	Frequency	Maximum	Proposed	Groundwater	Proposed	
Contaminant of	Groundwater				Detection	Non-Detect	Detected	of RL	Exceedance	of PCUL	Exceedance	Sediment	COC Selection	Groundwater	
Potential Concern ¹	Cleanup Level ²		Number	Number of	Frequency	Concentration	Concentration	Exceedance ⁴	Ratio ⁵	Exceedance ⁴	Ratio ⁵	COC ⁶	Criteria Met ⁷	coc	
(COPC)	(PCUL)	Units	Samples	Detections	(%)	(µg/L)	(μg/L)	(%)	(ER)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
Carcinogenic Polycyclic Aromatic Hy	drocarbons (cPAHs)														
Benzo[a]anthracene	0.01	μg/L	42	8	19%	0.018	0.10	10%	1.8	8%	10	Yes	Yes	Yes	
Benzo[a]pyrene	0.01	μg/L	42	4	10%	0.018	0.12	10%	1.8	3%	12	Yes	Yes	Yes	
Benzo[b]fluoranthene	0.01	μg/L	42	4	10%	0.018	0.18	10%	1.8	5%	18	Yes	Yes	Yes	
Benzo[k]fluoranthene	0.01	μg/L	42	4	10%	0.018	0.07	10%	1.8	5%	6.8	Yes	Yes	Yes	Retained as a COC
Chrysene	0.016	μg/L	42	5	12%	0.018	0.14	10%	1.1	3%	8.8	Yes	Yes	Yes	Notaliled as a coo
Dibenz[a,h]anthracene	0.01	μg/L	42	1	2%	0.018	0.02	10%	1.8	1%	1.8	Yes	No	Yes	
Indeno[1,2,3-c,d]pyrene	0.01	μg/L	42	3	7%	0.018	0.09	10%	1.8	3%	9.4	Yes	Yes	Yes	
cPAHs TEQ 8 (ND = 0.5RL)	0.01	μg/L	42	9	21%	0.01	0.17	10%	1.3	7%	16.7	Yes	Yes	Yes	
Pesticides and Herbicides															
4,4'-DDD	0.1	μg/L	18	0	0%	0.1		No Exceedance	1.0	No Exceedance		No	No	No	None
4,4'-DDE	0.1	μg/L	18	0	0%	0.1		No Exceedance	1.0	No Exceedance		No	No	No	None
4,4'-DDT	0.1	μg/L	18	0	0%	0.1		No Exceedance	1.0	No Exceedance		No	No	No	None
Aldrin	0.05	μg/L	18	0	0%	0.052		No Exceedance	1.0	No Exceedance	-	No	No	No	None
Alpha-BHC	0.05	μg/L	18	0	0%	0.052		No Exceedance	1.0	No Exceedance	-	No	No	No	None
Alpha-Chlordane (cis)	NE	μg/L	18	0	0%	0.052	-	No Exceedance	-	No Exceedance		No	No	No	None
Beta-BHC	0.05	μg/L	18	0	0%	0.052	-	No Exceedance	1.0	No Exceedance		No	No	No	None
Delta-BHC	NE	μg/L	18	0	0%	0.052		No Exceedance	-	No Exceedance		No	No	No	None
Dieldrin	0.1	μg/L	18	0	0%	0.1		No Exceedance	1.0	No Exceedance		No	No	No	None
Endosulfan I	0.05	μg/L	18	0	0%	0.052		No Exceedance	1.0	No Exceedance		No	No	No	None
Endosulfan II	NE	µg/L	18	0	0%	0.1		No Exceedance	-	No Exceedance		No	No	No	None
Endosulfan Sulfate	10	μg/L	18	0	0%	0.1		No Exceedance	0.0	No Exceedance	-	No	No	No	None
Endrin	0.1	μg/L	18	0	0%	0.1	-	No Exceedance	1.0	No Exceedance		No	No	No	None
Endrin Aldehyde	0.1	μg/L	18	0	0%	0.1	-	No Exceedance	1.0	No Exceedance	-	No	No	No	None
Endrin Ketone	NE	μg/L	18	0	0%	0.1		No Exceedance		No Exceedance	_	No	No	No	None
Gamma-Chlordane	NE	μg/L	18	0	0%	0.052		No Exceedance		No Exceedance		No	No	No	None
Heptachlor	0.05	µg/L	18	0	0%	0.052	_	No Exceedance	1.0	No Exceedance	-	No	No	No	None
Heptachlor Epoxide	0.05	μg/L	18	0	0%	0.052		No Exceedance	1.0	No Exceedance		No	No	No	None
Lindane (Gamma-BHC)	0.05	μg/L	18	0	0%	0.052		No Exceedance	1.0	No Exceedance		No	No	No	None
Herbicides			1						Ţ		Ţ	1			
2,4,5-T	NE	μg/L	18	0	0%	1	-	No Exceedance		No Exceedance			No	No	None
2,4-D	12,000	μg/L	18	0	0%	1.5		No Exceedance	<0.1	No Exceedance			No	No	None
2,4-DB	NE	μg/L	18	0	0%	10		No Exceedance		No Exceedance			No	No	None
Dalapon (DPA)	NE	μg/L	18	0	0%	2		No Exceedance		No Exceedance			No	No	None
Dicamba	NE	μg/L	18	0	0%	1	-	No Exceedance		No Exceedance		-	No	No	None
Dichlorprop	NE	μg/L	18	0	0%	3	-	No Exceedance		No Exceedance		_	No	No	None
Dinoseb	NE	μg/L	18	0	0%	2		No Exceedance		No Exceedance		-	No	No	None
MCPA	NE	μg/L	18	0	0%	260		No Exceedance		No Exceedance		-	No	No	None
Mecoprop (MCPP)	NE	μg/L	18	0	0%	250		No Exceedance		No Exceedance		-	No	No	None
Silvex (Fenoprop or 2,4,5-TP)	400	μg/L	18	0	0%	1		No Exceedance	<0.1	No Exceedance		-	No	No	None
Polychlorinated Biphenyls (PCBs)			1	1		<u> </u>	,		T		T	1		1	
Total PCBs (Sum of	0.01	μg/L	_							-		No	-	No	None
Aroclors or Congeners) Dioxins and Furans									<u> </u>		<u> </u>				
Total Dioxins/Furans -	_						I		1		1				
Human Health TEQ ⁹	5	pg/L	13	8	62%	2	2.18	No Exceedance	0.39	No Exceedance	0.4	No	No	No	None



1 Contaminants of potential concern (COPCs) were established for the RI based on a review of previous environmental studies. Previous sediment study results are summarized in Table F-1 (Appendix F).

 $^{\rm 2}$ Proposed groundwater cleanup levels are referenced from Table 3.

³ The groundwater data used for this RI consists of samples obtained by GeoEngineers between June 2008 and August 2017 in general accordance with the RI/FS Work Plan (GeoEngineers 2008a) and in coordination with Ecology. Groundwater results are summarized in Table F-1 (Appendix F).

⁴ Number of samples with analyte detected or non-detect at a concentration greater than PCUL / total number of samples analyzed for analyte.

⁵ Exceedance Ratio (max) = ratio of maximum detected or non-detect concentration divided by the Screening Level

 $^{\rm 6}$ Sediment contaminants of concern (COCs) are presented in Tables 8 and 9.

⁷ Initial contaminant of concern (COC) selection criteria is met if exceedance frequency is greater or equal to 10 percent or if the exceedance ratio is greater than 2.

⁸ Total cPAH Toxic Equivalency Quotients (TEQs) were calculated using Toxicity Equivalency Factors (TEFs) values referenced from MTCA Table 708.2 (WAC 173-340-900).

⁹ Total dioxin and furan TEQs were calculated using United States Environmental Protection Agency (USEPA) TEF values for human health (EPA, 2003).

NE = Not Established

μg/L = micrograms per liter

ng/L = nanograms per liter

ND = Non-detect

RL = reporting limit

TEQ = toxicity equivalency concentration

Bold indicated satisfaction of initial COC or consideration of other selection criteria.

Yellow shading indicates analyte is identified as a COC based on both satisfaction of initial selection criteria and consideration of other selection criteria, or on consideration of other selection criteria alone.



Summary Statistics and Evaluation of Soil Contaminants of Concern

Dakota Creek Industries

	1		1	1						Anacortes, Wa							1		
									Evaluat	on of RI Data Res			1					COC Sele	ection Considerations
		sed Soil Ip Level ¹		Total	Total	Detection	RL Maximum Non-Detect	Exceedance Evalua Frequency of RL	tion Maximum Non-Detect	Vadose 2 Maximum Detected	Zone Exceedance E Frequency of PCUL	Waluation Maximum Exceedance	Saturate Maximum Detected	Frequency of PCUL	Evaluation Maximum Exceedance	Proposed Groundwater	Initial Soil COC Selection	Proposed	
Analyte	Vadose Zone	Saturated Zone	Units	Number Samples	Number of Detections	Frequency (%)	Concentration (µg/L)	Exceedance ³ (%)	Exceedance Ratio ⁴	Concentration (µg/L)	Exceedance ³ (%)	Ratio ⁴ (ER)	Concentration (µg/L)	Exceedance ³ (%)	Ratio ⁴ (ER)	COC ⁵ (Yes/No)	Criteria Met ⁶ (Yes/No)	Soil COC (Yes/No)	Comments/Rationale
Metals																			
Arsenic	20	20	µg/L	133	33	25%	6.9	No Exceedance	0.3	910	27%	45.5	92	10%	4.6	Yes	Yes	Yes	Retained as a COC
Cadmium	3,500	3,500	μg/L	-	-	-			-		-	-/-	-		-	No	-	No	Not identified as a COPC
Total Chromium	5,300,000	5,300,000	μg/L	-	-	-			-		-	-	-	-	-	No	-	No	Not identified as a COPC
Copper	140,000	140,000	µg/L	22	22	100%	-	No Exceedance	-	1,100	No Exceedance		2,000.0	No Exceedance	-	No	-	No	None
Lead	1,000	1,000	μg/L	-	-	-			-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
Mercury	1,100	1,100	µg/L	-	-			-	-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
Nickel	48	48	μg/L	85	25	29%	48	No Exceedance	1.0	150	60%	3.1	200	21%	4.2	Yes	Yes	Yes	Retained as a COC
Silver	18,000	18,000	µg/L		-	-	-	-	-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
Zinc	1,100,000	1,100,000	µg/L	33	33	100%	0	No Exceedance	<0.1	2,800	No Exceedance	-	720	No Exceedance	-	No	-	No	None
etroleum Hydrocarbons	r	1	1	1	1		•	T	•	•				_	_	•	_		
Gasoline-Range	100	100	μg/L	2	0	0%	3	No Exceedance	-	-	No Exceedance		-	No Exceedance	-	No	No	Yes	Cleanup actions were previously completed to remove petroleum hydrocarbons in soil exceeding PCULs However, the data used to confirm the
Diesel-Range	2,000	2,000	μg/L	5	3	60%	25	No Exceedance	<0.1		No Exceedance	-	420	No Exceedance	0.2	No	No	Yes	removal of soil in these areas cannot verified. Additional soil investigation activities will be performed within the footprints of the previously completed
Heavy Oil-Range	2,000	2,000	μg/L	5	3	60%	50	No Exceedance	<0.1	-	No Exceedance	-	330	No Exceedance	0.2	No	No	Yes	remedial excavations to verify the completeness of the previously comp cleanup actions,
BETX Compounds	•			•			1					•	•	•	•	•	•		
Benzene	2,400	2,400	µg/L	2	0	0%	0.030	No Exceedance	<0.1	-	No Exceedance	-	-	No Exceedance	-	No	No	No	None
Ethylbenzene	350,000	350,000	µg/L	2	0	0%	0.050	No Exceedance	<0.1	-	No Exceedance	-		No Exceedance		No	No	No	None
Toluene	280,000	280,000	μg/L	2	0	0%	0.050	No Exceedance	<0.1	-	No Exceedance	-	_	No Exceedance	-	No	No	No	None
Xylenes	700,000	700,000	μg/L	2	0	0%	0.02	No Exceedance	<0.1	-	No Exceedance	-	_	No Exceedance	-	No	No	No	None
olatile Organic Compounds (VOCs)	s)		•	•								•	•			*			•
1,1,1,2-Tetrachloroethane	5,000	5,000	μg/L		-	_	-	-	<0.1	-	_					No	-	No	Not identified as a COPC
1,1,1-Trichloroethane	7,000,000	7,000,000	µg/L	-	_	_			-		_	_			_	No	_	No	Not identified as a COPC
1,1,2,2-Tetrachloroethane	660	660	μg/L	-	-	-		-	-	-	-				-	No	-	No	Not identified as a COPC
1,1,2-trichloro-1,2,2- trifluoroethar (CFC113)	110,000,000	110,000,000	µg/L	-	-	-	-	-			-	-	-	-		No	-	No	Not identified as a COPC
1,1,2-Trichloroethane	2,300	2,300	µg/L	-	-	-		-	-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
1,1-Dichloroethane	23,000	23,000	μg/L	-	-	7	-	-	_	-	-	-	-	-	-	No	-	No	Not identified as a COPC
1,1-Dichloroethene	180,000	180,000	μg/L	-	- 4	-	-	-		-	-		-	-	-	No	-	No	Not identified as a COPC
1,1-Dichloropropene	NE	NE	μg/L		-(1	-	-			-	-	-	-	-	-	No	-	No	Not identified as a COPC
1,2,3-Trichlorobenzene	NE	NE	µg/L	-	-	-	-	_	-	-	-	-	-	-	-	No	_	No	Not identified as a COPC
1,2,3-Trichloropropane	4.4	4.4	μg/L	-	-						-	-			-	No	-	No	Not identified as a COPC
1,2,4-Trichlorobenzene	4,500	4,500	µg/L		-	-	-		-		_	_	_	-	_	No	_	No	Not identified as a COPC
1,2,4-Trimethylbenzene	35,000	35,000	µg/L	_	-		-	/	-		_	_	_	-	_	No	_	No	Not identified as a COPC
1,2-Dibromo-3-chloropropane	160	160	µg/L		_	-	-	-		-	_	_	_	-	_	No	_	No	Not identified as a COPC
1,2-Dibromoethane (EDB)	66	66	μg/L	5	0	0%	0.005	No Exceedance	<0.1	_	No Exceedance	_	_	No Exceedance	_	No	No	No	None
1,2-Dichlorobenzene	320,000	320,000	μg/L	_	-		-	-		_	-	_		-		No	-	No	Not identified as a COPC
1,2-Dichloroethane (EDC)	1400	1400	μg/L	5	0	0%	0.01	No Exceedance	<0.1		No Exceedance			No Exceedance		No	No	No	None
1,2-Dichloropropane	3,500	3,500	μg/L		-	-					-		_	-	_	No	-	No	Not identified as a COPC
1,3,5-Trimethylbenzene	35,000	35,000	μg/L			_			_		_	_	_	_		No	_	No	Not identified as a COPC
1,3-Dichlorobenzene	NE	33,000 NE	μg/L		_		-		_	_	_	_	_	-		No	_	No	Not identified as a COPC
1.3-Dichloropropage	NE	NE	μ8/ ∟	+	-	-	1	-		1 -	_	-	-	_	_	INU	_	INU	Not identified as a COPC



Not identified as a COPC

No

No

No

No

No

No

No

1,3-Dichloropropane

1,4-Dichlorobenzene

2,2-Dichloropropane

2-Butanone (MEK)

NE

24000

NE

NE

24000

NE

2,100,000 2,100,000

μg/L

μg/L

μg/L

μg/L

									Evaluati	ion of RI Data Res	ults ²							COC Sel	ection Considerations
	Propo	sed Soil					RL E	xceedance Evalua	tion	Vadose Z	one Exceedance E	Evaluation	Saturated	Zone Exceedance					
	Cleanu						Maximum	Frequency	Maximum	Maximum	Frequency	Maximum	Maximum	Frequency	Maximum	Proposed	Initial Soil COC		
				Total	Total	Detection	Non-Detect	of RL	Non-Detect	Detected	of PCUL	Exceedance	Detected	of PCUL	Exceedance	Groundwater	Selection	Proposed	
	Vadose	Saturated		Number	Number of	Frequency	Concentration	Exceedance	Exceedance		Exceedance	Ratio ⁴	Concentration	Exceedance ³	Ratio ⁴	COC ⁵	Criteria Met ⁶	Soil COC	
Analyte	Zone	Zone	Units	Samples	Detections	(%)	(µg/L)	(%)	Ratio⁴	(µg/L)	(%)	(ER)	(µg/L)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
2-Chloroethyl Vinyl Ether	NE	NE	µg/L	-			-			-	-	-		_	-	No	-	No	Not identified as a COPC
2-Chlorotoluene	70,000	70,000	µg/L	-			-			-	-	-	-	-	-	No	-	No	Not identified as a COPC
2-Hexanone	18,000	18,000	µg/L		-	-	-	-		-		-	-	_	-	No	-	No	Not identified as a COPC
4-Chlorotoluene 4-Methyl-2-Pentanone	NE	NE	μg/L	-	-					-		-				No	-	No	Not identified as a COPC
(Methyl Isobutyl Ketone)	280,000	280,000	μg/L	-	-	-	-			-	-	-	-	-		No	-	No	Not identified as a COPC
Acetone	3,200,000	3,200,000	µg/L	-	-			-		-		-	1 -			No	_	No	Not identified as a COPC
Acrolein	NE	NE	μg/L		-	-		-	-	-			-	-		No	_	No	Not identified as a COPC
Acrylonitrile	240	240	μg/L	-				-	-	-	-	4	-	-		No	-	No	Not identified as a COPC
Bromobenzene	28,000	28,000	µg/L		-				-	-	-	_	-	-	-	No	-	No	Not identified as a COPC
Bromochloromethane	NE	NE	µg/L		-		-	-	-	-	-			-		No	_	No	Not identified as a COPC
Bromoform	17,000	17,000	μg/L		-	-	_		_	_	-	-		-	_	No	-	No	Not identified as a COPC
Bromomethane	4,900	4,900	μg/L	_	-			-	-	_		-		-		No	-	No	Not identified as a COPC
Carbon Disulfide	350,000	350,000	μg/L	-		-	-	-		-	4	-	-	-	-	No	-	No	Not identified as a COPC
Carbon Tetrachloride	1,900	1,900	µg/L	-	-	_	-	-	-	_	-	-	-	-	-	No	-	No	Not identified as a COPC
Chlorobenzene	70,000	70,000	µg/L		-				-	-	-	-	-		-	No	_	No	Not identified as a COPC
Chloroethane	NE	NE	μg/L	-	-		-	-	-	-	-	-	-			No	-	No	Not identified as a COPC
Chloroform	4200	4200	μg/L	-	-		-	_		-	-	-			-	No	-	No	Not identified as a COPC
Chloromethane	NE	NE	µg/L	-	-		_			_	-	-			_	No	-	No	Not identified as a COPC
Cis-1,2-Dichloroethene	7,000	7,000	µg/L		-		-			_	_	-		_	_	No	-	No	Not identified as a COPC
Cis-1,3-Dichloropropene	NE	NE	µg/L	_	_	_	_	_	_	_	_	-	_	_	_	No	_	No	Not identified as a COPC
Dibromochloromethane	1,600	1,600	µg/L	_	_	_	_	_			_	-		_	_	No	_	No	Not identified as a COPC
Dibromomethane	35,000	35,000	µg/L	_	_	_	_	_	_	-	-		-	_	_	No	_	No	Not identified as a COPC
Dichlorobromomethane	2,100	2,100	µg/L	_	_	_		-	_		-	-	-		_	No		No	Not identified as a COPC
Dichlorodifluoromethane (CFC 12)	700,000	700,000	µg/L	-		-	_				-	-			_	No	-	No	Not identified as a COPC
Hexachlorobutadiene	1,700	1,700	µg/L	_	_	-	_			-	-/	-		-	_	No	-	No	Not identified as a COPC
Isopropylbenzene	350,000	350,000	µg/L	-		_	_			-	-				_	No	-	No	Not identified as a COPC
Methyl lodide	NE	NE	µg/L			-	_	-	-	1		-		_	-	No	_	No	Not identified as a COPC
Methyl t-Butyl Ether (MTBE)	73,000	73,000	µg/L	5	0	0%	0	No Exceedance	<0.1	-	No Exceedance	-		No Exceedance		No	No	No	None
Methylene Chloride	21,000	21,000	µg/L	_	_	_	_	-		-	_			_	_	No		No	Not identified as a COPC
Naphthalene	70,000	70,000	µg/L	_	_	_	_	_	_		-				_	No		No	Not identified as a COPC
n-Butylbenzene	180,000	180,000	µg/L	_	_	_	- 4	-	-		-	_		_	_	No		No	Not identified as a COPC
n-Propylbenzene	350,000	350,000	µg/L	_	_	_		-	7		_				_	No		No	Not identified as a COPC
p-lsopropyltoluene	NE	NE	µg/L			_	-	-					_	_	_	No	_	No	Not identified as a COPC
sec-Butylbenzene	350,000	350,000	μg/L	_		_	_	-					_	_	_	No	_	No	Not identified as a COPC
Styrene	700,000	700,000	µg/L	_	_	_					_		_	_	_	No	_	No	Not identified as a COPC
tert-Butylbenzene	350,000	350,000	μg/L	_					-				-		_	No		No	Not identified as a COPC
Tetrachloroethene (PCE)	21,000	21,000	μg/L	_	_		-					_		_	_	No		No	Not identified as a COPC
Trans-1,2-Dichloroethene	70,000	70,000	μg/L	_		-			-		-	_				No		No	Not identified as a COPC
Trans-1,3-Dichloropropene	70,000 NE	70,000 NE	μg/L		_		_	-					-			No		No	Not identified as a COPC
Trans-1,4-Dichloro-2-butene	NE NE	NE NE	μg/L μg/L		-		-	-				-				No	_	No	Not identified as a COPC Not identified as a COPC
Trichloroethene (TCE)	1,800	1,800	μg/L μg/L			-		_				-				No		No	Not identified as a COPC Not identified as a COPC
Trichlorofluoromethane (CFC 11)	1,100,000	1,100,000	μg/L μg/L			-						-			-	No	_	No	Not identified as a COPC Not identified as a COPC
` ,	3,500,000					_													Not identified as a COPC Not identified as a COPC
Vinyl Acetate		3,500,000	µg/L	-	- '						-				-	No	-	No	
Vinyl Chloride	88 (00 a)	88	μg/L	-	-	-					-	-	-			No	_	No	Not identified as a COPC
Semi-Volatile Organic Compounds (SV	· ·	4.500		1			1		I			T	T		I	Al-	1	NI-	Not identified as a COPO
1,2,4-Trichlorobenzene	4,500	4,500	μg/L	-	-	-	-		-	-	-	-	-			No	-	No	Not identified as a COPC
1,2-Dichlorobenzene	320,000	320,000	μg/L	-	-											No	_	No	Not identified as a COPC



					,		,		Evaluatio	on of RI Data Res	ults ²							COC Sele	ection Considerations
	Propo	sed Soil					RL I	Exceedance Evalua		Vadose Z	one Exceedance E		Saturated	Zone Exceedance					
		ıp Level ¹					Maximum	Frequency	Maximum	Maximum	Frequency	Maximum	Maximum	Frequency	Maximum	Proposed	Initial Soil COC		
	Vadose	Saturated		Total	Total	Detection	Non-Detect	of RL	Non-Detect	Detected	of PCUL	Exceedance	Detected	of PCUL	Exceedance	Groundwater	Selection	Proposed	
Analyte	Zone	Zone	Units	Number Samples	Number of Detections	Frequency	Concentration	Exceedance	Exceedance Ratio ⁴	Concentration (µg/L)	Exceedance	Ratio ⁴ (ER)	Concentration (µg/L)	Exceedance ³ (%)	Ratio ⁴ (ER)	COC ⁵ (Yes/No)	Criteria Met ⁶	Soil COC (Yes/No)	Comments/Rationale
1.3-Dichlorobenzene	NE	NE NE	µg/L	Samples	Detections	(%) 	(μg/L) 	(%) 	Ratio	(μg/ L) 	(%) 	(ER) 	(μg/ L) 	(%)	(ER) 	No	(Yes/No)	No	Not identified as a COPC
.,4-Dichlorobenzene	24,000	24,000		-	-				-				-			No		No	Not identified as a COPC
,	24,000 NE	24,000 NE	µg/L	_					-			-				No		No	Not identified as a COPC
2,2'-Oxybis[1-chloropropane]	350,000	350,000	µg/L		-				-	-		-	-			No		No	Not identified as a COPC
2,4,5-Trichlorophenol		1	µg/L				-				_	-	_	-					
.4-Dichlorophenol	3,500	3,500	µg/L		-		-		-	-	-	-				No		No	Not identified as a COPC
2,4-Dichlorophenol	11,000	11,000 70,000	µg/L	-	-							-				No No		No No	Not identified as a COPC Not identified as a COPC
• • • • • • • • • • • • • • • • • • • •	70,000		µg/L	_	-		-		-			-		-		No		No	Not identified as a COPC Not identified as a COPC
2,4-Dinitrophenol	7,000	7,000	µg/L	-	-				-	-	-	_	-	-		-			
2,4-Dinitrotoluene	420	420	µg/L		-	-			-	-	-	-	-	-		No	-	No	Not identified as a COPC
2,6-Dinitrotoluene	88	88	µg/L	-			-		-	-	-				-	No	_	No	Not identified as a COPC
-Chloronaphthalene	280,000	280,000	µg/L	-	-		-		-	-	_	-		-		No	-	No	Not identified as a COPC
-Chlorophenol	18,000	18,000	µg/L	-	-		-	-	-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
-Nitroaniline	35,000	35,000	μg/L		-				-	-	-			-	-	No	-	No	Not identified as a COPC
-Nitrophenol	NE	NE	μg/L	-	-		-	-	-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
,3'-Dichlorobenzidine	290	290	μg/L	-	-		_	-	-	-			-			No	-	No	Not identified as a COPC
3-Nitroaniline	NE	NE	µg/L	-	-	-	-	-	-	-	-	-	-	-		No	-	No	Not identified as a COPC
1,6-Dinitro-2-methylphenol	280	280	μg/L		-		-	-	-	-	-	-		-		No	-	No	Not identified as a COPC
4-Bromophenyl-phenylether	NE	NE	μg/L	-		-	-	-	-		-	7		-		No	-	No	Not identified as a COPC
1-Chloro-3-methylphenol	350,000	350,000	μg/L	-	-	-	-	-	-	-	-	-	-	-	-	No		No	Not identified as a COPC
I-Chloroaniline	660	660	μg/L	-	-	-	_	-	-	-	-	_	_	-	_	No	-	No	Not identified as a COPC
-Chlorophenyl-phenylether	NE	NE	μg/L			-	-	-	-	-	-	-	-	-	-	No		No	Not identified as a COPC
1-Nitroaniline	14,000	14,000	μg/L			-	-	-	- '	-	-	-	-	-	-	No	-	No	Not identified as a COPC
1-Nitrophenol	NE	NE	μg/L	-	-	-	-	_		-	Á	-		-	-	No	-	No	Not identified as a COPC
Benzoic acid	14,000,000	14,000,000	μg/L	-		-	-	-	-	-	-	-		-	-	No	-	No	Not identified as a COPC
Benzyl alcohol	350,000	350,000	μg/L	-		-	-	-	-	-	-	-		-	-	No	-	No	Not identified as a COPC
ois(2-Chloroethoxy)methane	NE	NE	μg/L	-	-	-	-	-	-	-	-	_	-	-	-	No	-	No	Not identified as a COPC
ois(2-chloroethyl)ether	120	120	μg/L	-	-	-	_	-	-	-	-	-	_	-	-	No	_	No	Not identified as a COPC
ois(2-Ethylhexyl)phthalate	9,400	9,400	μg/L	_		-		-	-	-	-			-	_	No	-	No	Not identified as a COPC
Butylbenzylphthalate	69,000	69,000	μg/L	-		-	_	-		-	-	_	_	-	-	No	-	No	Not identified as a COPC
Carbazole	NE	NE	μg/L	-	-	_	_		-	-	-	_	-	-		No		No	Not identified as a COPC
Dibenzofuran	3,500	3,500	μg/L	-	-	_	-	-		- 1	-		_	-	-	No		No	Not identified as a COPC
Diethylphthalate	2,800,000	2,800,000	μg/L	-	-	-	-			- 1	-		-	_	-	No		No	Not identified as a COPC
Dimethylphthalate	NE	NE	μg/L	-	-	-	-	-		-	-	_	-	_	_	No		No	Not identified as a COPC
Di-n-butylphthalate	350,000	350,000	µg/L	-		_	-	-	_	-	-			_		No		No	Not identified as a COPC
Di-n-octylphthalate	35,000	35,000	μg/L	_	_	_	_	-			_		_	_	_	No		No	Not identified as a COPC
Hexachlorobenzene	82	82	µg/L		_	_	_	-	-	_	_	_	_		_	No	_	No	Not identified as a COPC
Hexachlorobutadiene	1,700	1,700	µg/L		_	-	-		_	_	_	_			_	No	_	No	Not identified as a COPC
Hexachlorocyclopentadiene	21,000	21,000	µg/L	_	_	-	_	_	_	_	_	_				No	_	No	Not identified as a COPC
Hexachloroethane	2,500	2,500	μg/L		_	-	-	-	_	_	_	_				No	_	No	Not identified as a COPC
sophorone	140,000	140,000	μg/L μg/L			-	-						_			No	-	No	Not identified as a COPC
Vitrobenzene	7,000	7,000				_	-									No	-	No	Not identified as a COPC
n-Nitroso-di-n-propylamine	19	19	µg/L µg/L						_							No	-	No	Not identified as a COPC
	27,000	27,000			-	_										No	-	No	
-Nitrosodiphenylamine -Orosol (2 Methylphonel)			µg/L		-			_		-	_					+			Not identified as a COPC
-Cresol (2-Methylphenol)	180,000	180,000	µg/L		-	-			-	-	-				-	No		No	Not identified as a COPC
o-Cresol (4-Methylphenol)	350,000	350,000	µg/L		-	-	-		-	_	_	_	-		-	No	-	No	Not identified as a COPC
Pentachlorophenol	330	330	µg/L		-		-		-	-		-	-		-	No	-	No	Not identified as a COPC
Phenol	1,100,000	1,100,000	μg/L	-		-	-	-	_	-	-	-	-	-	-	No	-	No	Not identified as a COPC
n-Carcinogenic Polycyclic Aromat					1 .								0.55		1		1 1		L
L-Methylnaphthalene	4,500	4,500	μg/L	11	4	36%	0.06	No Exceedance	<0.1	0.06	No Exceedance	-	0.28	No Exceedance	-	No	No	No	None
2-Methylnaphthalene	14,000	14,000	µg/L	11	4	36%	0.03	No Exceedance	<0.1	0.05	No Exceedance	-	0.032	No Exceedance		No	No	No	None
Acenaphthene	210,000	210,000	μg/L	11	2	18%	0.02	No Exceedance	<0.1	0.015	No Exceedance	-	0.041	No Exceedance	_	No	No	No	None



									Evaluati	on of RI Data Res	sults ²							COC Sele	ection Considerations
	Brono	sed Soil					RLI	Exceedance Evalua	tion	Vadose 2	Zone Exceedance E		Saturated	d Zone Exceedance	Evaluation				
		seu Soli ip Level ¹					Maximum	Frequency	Maximum	Maximum	Frequency	Maximum	Maximum	Frequency	Maximum	Proposed	Initial Soil COC		
				Total	Total	Detection	Non-Detect	of RL	Non-Detect	Detected	of PCUL	Exceedance	Detected	of PCUL	Exceedance	Groundwater	Selection	Proposed	
Amelida	Vadose Zone	Saturated Zone		Number	Number of	Frequency	Concentration	Exceedance	Exceedance	Concentration	Exceedance	Ratio ⁴	Concentration	Exceedance	Ratio ⁴	COC ⁵	Criteria Met ⁶	Soil COC	O
Analyte Acenaphthylene			Units	Samples	Detections	(%)	(μg/L)	(%)	Ratio ⁴	(μg/L)	(%)	(ER)	(μg/L)	(%)	(ER)	(Yes/No)	(Yes/No)	(Yes/No)	Comments/Rationale
	NE 4.400.000	NE 1 100 000	µg/L	11	2	18%	0.02	No Exceedance		0.12	No Exceedance	-	0.02	No Exceedance	-	No	No	No	None
Anthracene	1,100,000	1,100,000	µg/L	11	4	36%	0.02	No Exceedance	<0.1	0.12	No Exceedance	_	0.04	No Exceedance		No	No	No	None
Benzo[g,h,i]perylene	NE 110.000	NE 440.000	µg/L	11	6	55%	0.02	No Exceedance		0.56	No Exceedance	-	0.11	No Exceedance	-	No	No	No	None
Fluoranthene	140,000	140,000	μg/L	11	6	55%	0.05	No Exceedance	<0.1	2.0	No Exceedance	-	0.29	No Exceedance	-	No	No	No	None
Fluorene	140,000	140,000	μg/L	11	4	36%	0.03	No Exceedance	<0.1	0.56	No Exceedance	-	0.04	No Exceedance		No	No	No	None
Naphthalene	70,000	70,000	μg/L	11	4	36%	0.08	No Exceedance	<0.1	0.56	No Exceedance	-	0.08	No Exceedance	-	No	No	No	None
Phenanthrene	NE	NE	μg/L	11	6	55%	0.07	No Exceedance	-	0.62	No Exceedance	-	0.16	No Exceedance	-	No	No	No	None
Pyrene	110,000	110,000	μg/L	11	6	55%	0.05	No Exceedance	<0.1	1.80	No Exceedance	-	0.27	No Exceedance	-	No	No	No	None
arcinogenic Polycyclic Aromatic Hy	1	1				1		1	1	1	I		1 212						
Benzo[a]anthracene	1.1	0.06	μg/L	19	7	37%	0.02	26%	0.3	0.94	No Exceedance	16	0.13	11%	2.3	Yes	Yes	Yes	_
Benzo[a]pyrene	0.31	0.016	μg/L	19	7	37%	0.02	No Exceedance	1	1.00	60%	65	0.13	22%	9.2	Yes	Yes	Yes	_
Benzo[b]fluoranthene	3.9	0.20	μg/L	19	7	37%	0.02	No Exceedance	0.1	0.92	No Exceedance	4.7	0.12	No Exceedance	0.7	Yes	Yes	Yes	_
Benzo[k]fluoranthene	39	2.0	μg/L	19	6	32%	0.02	No Exceedance	0.0	1.10	No Exceedance	0.6	0.12	No Exceedance	<0.1	Yes	Yes	Yes	Retained as a COC
Chrysene	127	6.4	μg/L	19	7	37%	0.02	No Exceedance	0.0	1.10	No Exceedance	0.2	0.15	No Exceedance	<0.1	Yes	Yes	Yes	_
Dibenz[a,h]anthracene	0.6	0.029	μg/L	19	3	16%	0.02	No Exceedance	1	0.56	No Exceedance	20	0.04	11%	6.5	Yes	Yes	Yes	
Indeno[1,2,3-c,d]pyrene	11	0.56	μg/L	19	6	32%	0.02	No Exceedance	0.0	0.64	No Exceedance	1.2	0.09	No Exceedance	0.2	Yes	Yes	Yes	
cPAHs TEQ (ND = 0.5RL)	0.31	0.016	μg/L	19	7	37%	0.02	11%	2	1.38	40%	89	0.18	33%	12.7	Yes	Yes	Yes	
Pesticides and Herbicides		1	ı	1	1	ı	I	ī	1	1			1			1	Г		T
4,4'-DDD	110	110	μg/L	-		-	-		-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
4,4'-DDE	390	390	μg/L						-		-			-	-	No		No	Not identified as a COPC
4,4'-DDT	4	4	µg/L		-		-	-			-	-			-	No	_	No	Not identified as a COPC
Aldrin	7.7	7.7	µg/L	-	-	-	-			-	-		-		-	No		No	Not identified as a COPC
Alpha-BHC	21	21	µg/L	-	-		-		-	-		-	-			No		No	Not identified as a COPC
Alpha-Chlordane (cis)	NE 70	NE 70	µg/L	-	-	-	-		-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
Beta-BHC	73	73	µg/L	-	-	-	-		-	-		-	-			No	-	No	Not identified as a COPC
Delta-BHC	NE	NE	μg/L	-	-	-	-		-	-	_	_	-	-	-	No	-	No	Not identified as a COPC
Dieldrin	8.2	8.2	μg/L	-	-	-	-		-	-	-	-		-	-	No	-	No	Not identified as a COPC
Endosulfan I	NE	NE	μg/L	-	-	-	-	-	-	-	-	-	-			No	-	No	Not identified as a COPC
Endosulfan II	NE	NE	μg/L	-	-	-	-	-		-	-			-	-	No	-	No	Not identified as a COPC
Endosulfan Sulfate	21,000	21,000	μg/L	-	-	-	-		-	-	-	-	-			No	-	No	Not identified as a COPC
Endrin	1,100	1,100	μg/L	-	-	-	-	-	-	-	-			-	-	No	-	No	Not identified as a COPC
Endrin Aldehyde	NE	NE	µg/L	-		-		-		- \	-				-	No	-	No	Not identified as a COPC
Endrin Ketone	NE	NE	μg/L	-		-	-	-	-	-	-				_	No	-	No	Not identified as a COPC
Gamma-Chlordane	NE	NE	μg/L	-	-	-	-	- 4	-	-	-	-	_	-	-	No	-	No	Not identified as a COPC
Heptachlor	29	29	μg/L	-	-	-	-		-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
Heptachlor Epoxide	14	14	µg/L	-	-	-	-	-	-	-	-	-	-	-	-	No	-	No	Not identified as a COPC
Lindane (Gamma-BHC) Methoxychlor	0.01 18,000	0.01 18,000	µg/L	-			-	-	-		-	-			-	No No		No No	Not identified as a COPC Not identified as a COPC
Toxaphene	120	120	µg/L µg/L	-	_	-		-	-	_	_	_			_	No	_	No	Not identified as a COPC Not identified as a COPC
olychlorinated Biphenyls (PCBs)	120	120	µg/∟					-								140		INU	110t lucitation as a cor c
Total PCBs (Sum of	10	10	μg/L	-	-	-	_	-		_		-	-	-	_	No	-	No	Not identified as a COPC
Aroclors or Congeners) Dioxins and Furans	1	1	ļ	1			ļ		_	1	l .	ļ	1	ļ	ļ.	<u> </u>	ļ ļ		
Total Dioxins/Furans -				1													ļ		
Human Health TEQ	1,700	1,700	ng/L	6	6	100%		No Exceedance		4.27	No Exceedance	<1	0.22	No Exceedance	<1	No	No	No	None



- $^{\mbox{\scriptsize 1}}$ Proposed soil cleanup levels are referenced from Table 4.
- 2 Data used to evaluate Site soil conditions are presented in Table G-1 (Appendix G).
- ³ Number of samples with analyte detected or non-detect at a concentration greater than PCUL / total number of samples analyzed for analyte.
- ⁴ Exceedance Ratio (max) = ratio of maximum detected or non-detect concentration divided by the Screening Level
- $^{\rm 5}$ Groundwater contaminants of concern (COCs) are presented in Table 10.
- ⁶ Initial contaminant of concern (COC) selection criteria is met if exceedance frequency is greater or equal to 10 percent or if the exceedance ratio is greater than 2.

NE = Not Established

μg/L = micrograms per liter

ng/L = nanograms per liter

ND = Non-detect

RL = reporting limit

TEQ = toxicity equivalency concentration

Bold indicated satisfaction of initial COC or consideration of other selection criteria.

Yellow shading indicates analyte is identified as a COC based on both satisfaction of initial selection criteria and consideration of other selection criteria, or on consideration of other selection criteria alone.



Table 12

Sediment Contaminant of Concern Chemical Analytical Data – Protection of Benthic Organisms Dakota Creek Industries

Anacortes, Washington

Samp	le Location ID ¹	DC-SED-09	P1-2	DC-106-2	DC-106-2	FB-A4-14	FB-A4-15	SMA-1		
Sample	Identification	DC-SED-09	AN-P1-2	DCI06-2A	DCI06-2-D	FB-A4-14	FB-A4-15	SMA 1-1		
	Sample Date	08/06/97	07/15/04	N/A	N/A	09/06/07	09/06/07	09/30/08	Prop	osed
Sample	Interval (dbm)	0-10 cm	1-3 ft	0-10 cm	0-10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	Sedi	ment
	Sample Study	1997	2004	2007	2007	2007	2007	2007		ening
		Phase II ESA	Sediment Study	Interim Action	Le	vel ²				
	Sample Type	Surface	Subsurface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	Otten Engineering	Anchor Env.	Floyd Snider	Floyd Snider	Ecology	Ecology	GeoEngineers	LAET	2LAET
Conventionals	•								•	_
Total Organic Carbon (TOC)	%	0.167	0.64	0.641	1.15	0.95 J	0.26	0.12	NE	NE
Total Volatile Solids (TVS)	%				-	2.46 J	2.35		NE	NE
Total Solids (TS)	%	N/A	78.2	78.3	78.2	79.1 J	76.8	78.1	NE	NE
Total Ammonia	mg-N/kg				-	13.6 J	5.9		NE	NE
Total Sulfide	mg/kg				_	0.07 J	6.7 UJ		NE	NE
Grain Size										
Gravel (>2,000 μm)	%	-		-	-	-	59.9		NE	NE
μm)	%		-	-	-		7.29	-	NE	NE
Coarse Sand (1,000 to 500 μm)	%		-		-			-	NE	NE
Medium Sand (500 to 250 μm)	%	_	-					-	NE	NE
Fine Sand (250 to 125 μm)	%		-	-	-			-	NE	NE
Very Fine Sand (125 to 62.5 μm)	%		-	-	-			-	NE	NE
Coarse Silt (62.5 to 31 µm)	%		-	-	-		3.52	-	NE	NE
Medium Silt (31 to 15.6 μm)	%		-	-	-			-	NE	NE
Fine Silt (15.6 to 7.8 μm)	%	_	-	-	-			-	NE	NE
Very Fine Silt (7.8 to 3.9 μm)	%		_		-				NE	NE
Clay (3.9 to <1 μm)	%	-	-	-	-		1.25	-	NE	NE
Total Fines (<62.5 μm)	%	-					4.77		NE	NE
Metals	•							•	•	l .
Arsenic	mg/kg	2.1 J	- 1			2.37	2.1		57	73
Cadmium	mg/kg	ND				0.09	0.08 U	0.054	5.1	6.7
Chromium	mg/kg	13.4				12.3	13.5	26.2	260	270
Copper	mg/kg	15.3				10.4	9.49	27.4	390	390
Lead	mg/kg	6.23 J				2.23 J	1.63	4.08	450	530
Mercury	mg/kg	ND				0.01 J	0.01	0.033	0.41	0.59
Silver	mg/kg	0.0582				0.02 U	0.03	0.07	6.1	6.1
Zinc	mg/kg	21.3	-	_		20.9	19.1	43.9	410	960



Sam	ple Location ID ¹	DC-SED-09	P1-2	DC-106-2	DC-106-2	FB-A4-14	FB-A4-15	SMA-1		
	le Identification	DC-SED-09	AN-P1-2	DCI06-2A	DCI06-2-D	FB-A4-14	FB-A4-15	SMA 1-1		
	Sample Date	08/06/97	07/15/04	N/A	N/A	09/06/07	09/06/07	09/30/08	Prop	osed
Samp	le Interval (dbm)	0-10 cm	1-3 ft	0-10 cm	0-10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	Sedi	iment
-	Sample Study	1997	2004	2007	2007	2007	2007	2007		ening
	Sample Study	Phase II ESA	Sediment Study	Interim Action	Le	vel ²				
	Sample Type	Surface	Subsurface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	Otten Engineering	Anchor Env.	Floyd Snider	Floyd Snider	Ecology	Ecology	GeoEngineers	LAET	2LAET
Low Molecular Weight Polycyclic A	1			<u> </u>	Τ		1	T	T	T
Sum of LPAHs ³	μg/kg	-				9.6	9.1 U	4.40	5,200	5,200
2-Methylnaphthalene	μg/kg 				-	9.70	9.1 U	0.85 J	670	670
Acenaphthene	µg/kg	-			-	9.7 U	9.1 U	0.91 J	500	500
Acenaphthylene	µg/kg	-				1.4 J	9.1 U	0.24 U	1,300	1,300
Anthracene	µg/kg	-		-	-	9.7 U	9.1 U	0.5 J	960	960
Fluorene	µg/kg				-	9.7 U	9.1 U	1.1 J	540	540
Naphthalene	µg/kg					9.7 U	9.1 U	2.5	2,100	2,100
Phenanthrene	µg/kg				-	8.2 J	9.1 U	2.9	1,500	1,500
Low Molecular Weight Polycyclic A	omatic Hydrocarbo	ns (LPAHs) (OC Normalize	ed)				1	1		
Sum of LPAHs ³	mg/kg OC	-				1.01 U	3.5 U	6.03 J	370	780
2-Methylnaphthalene	mg/kg OC					1.01 U	3.5 U	0.71 J	38	64
Acenaphthene	mg/kg OC				-	1.01 U	3.5 U	0.76 J	16	57
Acenaphthylene	mg/kg OC		-	-	-	0.15 J	3.5 U	0.2 U	66	66
Anthracene	mg/kg OC				-	1.01 U	3.5 U	0.42 J	220	1,200
Fluorene	mg/kg OC					1.01 U	3.5 U	0.92 J	23	79
Naphthalene	mg/kg OC				-	1.01 U	3.5 U	2.1 J	99	170
Phenanthrene	mg/kg OC		-	-	-	0.86 J	3.5 U	2.4 J	100	480
High Molecular Weight Polycyclic A	romatic Hydrocarbo	ons (HPAHs) (Dry Weight)								
Sum of HPAHs ⁴	µg/kg	-			-	121 J	2.2 J	9.35 J	12,000	17,000
Benzo(a)anthracene	µg/kg	_				4.1 J	9.1 U	1.3 J	1,300	1,600
Benzo(a)pyrene	µg/kg	-	-	_	-	2.6 J	9.1 U	0.86 J	1,600	1,600
Total Benzofluoranthenes ⁵	µg/kg	-	-	_	-	9.9 J	9.1 U	1.87 J	3,200	3,600
Benzo(g,h,i)perylene	µg/kg	-	-	-	-	2 J	9.1 U	0.94 J	670	720
Chrysene	μg/kg	-	-			18	9.1 U	1.1 J	1,400	2,800
Dibenzo(a,h)anthracene	µg/kg	-	_			9.7 U	9.1 U	0.28 U	230	230
Fluoranthene	µg/kg					50	2.2 J	3	1,700	2,500
Indeno(1,2,3-c,d)pyrene	µg/kg	-	-	-		1.8 J	9.1 U	0.83 J	600	690
Pyrene	µg/kg	_	-		-	33	9.1 U	4.0	2,600	3,300
High Molecular Weight Polycyclic A		ons (HPAHs) (OC Normaliz	ed)		ı		1	1	=,	-,
Sum of HPAHs ⁴	mg/kg OC	-		-		12.7 J	0.85 J	7.79 J	960	5,300
Benzo(a)anthracene	mg/kg OC	-		_	-	0.43 J	3.5 U	1.08 J	110	270
Benzo(a)pyrene	mg/kg OC		-	_		0.27 J	3.5 U	0.72 J	99	210
Total Benzofluoranthenes ⁵	mg/kg OC	-				1.04 J	3.5 U	1.56 J	230	450
Benzo(g,h,i)perylene	mg/kg OC		-			0.21 J	3.5 U	0.78 J	31	78
Chrysene	mg/kg OC					1.89 J	3.5 U	0.92 J	110	460
Dibenzo(a,h)anthracene	mg/kg OC					1.01 U	3.5 U	0.23 U	12	33
Fluoranthene			-			5.26 J	0.85 J	2.5	160	1,200
	mg/kg OC		<u></u>				3.5 U			+
Indeno(1,2,3-c,d)pyrene	mg/kg OC					0.19 J		0.69 J	34	88
Pyrene	mg/kg OC					3.47 J	3.5 U	3.3	1,000	1,400



Samr	ole Location ID ¹	DC-SED-09	P1-2	DC-106-2	DC-106-2	FB-A4-14	FB-A4-15	SMA-1		
Sample	e Identification	DC-SED-09	AN-P1-2	DCI06-2A	DCI06-2-D	FB-A4-14	FB-A4-15	SMA 1-1		
	Sample Date	08/06/97	07/15/04	N/A	N/A	09/06/07	09/06/07	09/30/08	Prop	osed
Sample	e Interval (dbm)	0-10 cm	1-3 ft	0-10 cm	0-10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm		ment
	Sample Study	1997	2004	2007	2007	2007	2007	2007		ening
		Phase II ESA	Sediment Study	Sediment Study	Sediment Study	Sediment Study	Sediment Study	Interim Action		vel ²
	Sample Type	Surface	Subsurface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	Otten Engineering	Anchor Env.	Floyd Snider	Floyd Snider	Ecology	Ecology	GeoEngineers	LAET	2LAET
Chlorinated Hydrocarbons (Dry Weig	1			T		9.7 U	9.1 U	0.611	24	
1,2,4-Trichlorobenzene 1,2-Dichlorobenzene	µg/kg	-		-		9.7 U	9.1 U	2.6 U 2.9 U	31 35	51 50
1,4-Dichlorobenzene	µg/kg	-	-	-	-	9.7 U	9.1 U	2.9 U	110	110
	µg/kg	-		-	-					
Hexachlorobenzene	µg/kg	-		-	-	9.7 U	9.1 U	1.2 U	22	70
Chlorinated Hydrocarbons (OC Norm				T		1.04.11	2511	0.011	0.01	1.0
1,2,4-Trichlorobenzene 1,2-Dichlorobenzene	mg/kg OC	-	-		-	1.01 U	3.5 U	2.2 U 2.4 U	0.81 2.3	1.8 2.3
1,4-Dichlorobenzene	mg/kg OC		-	-		1.01 U	3.5 U	2.4 U	3.1	9
•	mg/kg OC		-	-		1.01 U	3.5 U		0.38	2.3
Hexachlorobenzene	mg/kg OC		<u></u>	-		1.01 U	3.5 U	1 U	0.38	2.3
Phthalates (Dry Weight)						97	91	7 U	1,300	1,900
Bis(2-Ethylhexyl) Phthalate	µg/kg	-				9.7 U	9.1 U		63	900
Butyl benzyl Phthalate	µg/kg	-	-	-	-	9.7 U	9.1 0 16	3.2 U 7.9 U	1,400	1,400
Dibutyl Phthalate	µg/kg	-	-	-	-	1.9 J	9.1 U		1	
Diethyl Phthalate	µg/kg	-	-	-	-	1.9 J	9.1 U	1.3 U	200	> 1,200
Dimethyl Phthalate	µg/kg			-	-	9.7 U	9.1 U	1 U	71 6,200	160
Di-N-Octyl Phthalate Phthalates (OC Normalized)	µg/kg		-	-		9.7 0	9.10	1.7 U	6,200	6,200
· · · · · · · · · · · · · · · · · · ·	mg/kg 00			-	-	10.21	35	E 02 II	17	70
Bis(2-Ethylhexyl) Phthalate	mg/kg OC		-	-		1.01 U	3.5 U	5.83 U 2.67 U	47 5	78 64
Butyl benzyl Phthalate	mg/kg OC			-		0.97 J	6.15	6.58 U	220	1,700
Dibutyl Phthalate	mg/kg OC		-	-	-	0.97 J	3.50	1.08 U	61	110
Diethyl Phthalate Dimethyl Phthalate	mg/kg OC					0.2 J 0.11 J	3.5 U	0.83 U	53	53
Di-N-Octyl Phthalate	mg/kg OC mg/kg OC			-		1.01 U	3.5 U	1.42 U	58	4,500
Phenols (Dry Weight)	ilig/ kg OC					1.010	3.5 0	1.42 0	56	4,500
2,4-Dimethylphenol	ug/kg	-					46 R	5.5 U	29	29
2-methylphenol (o-Cresol)	µg/kg			-			9.1 U	1.5 U	63	63
4-methylphenol (p-Cresol)	µg/kg	-					9.1 U	3.4 J	670	670
Pentachlorophenol	µg/kg µg/kg	-	-				91 U	20 U	360	690
Phenol	μg/kg μg/kg						28 U	2 U	420	1,200
Miscellaneous Extractables (Dry Wei							200	2.0	420	1,200
Dibenzofuran	µg/kg					9.7 U	9.1 U	0.87 J	540	540
Hexachlorobutadiene	µg/kg µg/kg					9.7 U	9.1 U	2.5 U	11	120
N-Nitrosodiphenylamine	µg/kg µg/kg					9.7 U	9.1 U	2.5 U	28	40
Benzoic Acid	μg/kg					200	190 R	96 U	650	650
Benzyl Alcohol	µg/kg µg/kg			-		6.2 J	19 U	2.1 U	57	73
Miscellaneous Extractables (OC Norr				<u> </u>	<u> </u>	U.Z J	1 100	2.10	JI	13
Dibenzofuran	mg/kg OC					1.01 U	3.5 U	0.73 J	15	58
Hexachlorobutadiene	mg/kg OC					1.01 U	3.5 U	2.08 U	3.9	6.2
N-Nitrosodiphenylamine	mg/kg OC					1.01 U	3.5 U	1.33 U	3.9	11



Sample	e Location ID ¹	DC-SED-09	P1-2	DC-106-2	DC-106-2	FB-A4-14	FB-A4-15	SMA-1		
Sample	Identification	DC-SED-09	AN-P1-2	DCI06-2A	DCI06-2-D	FB-A4-14	FB-A4-15	SMA 1-1		
	Sample Date	08/06/97	07/15/04	N/A	N/A	09/06/07	09/06/07	09/30/08	Prop	osed
Sample	Interval (dbm)	0-10 cm	1-3 ft	0-10 cm	0-10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	Sedi	ment
	Sample Study	1997	2004	2007	2007	2007	2007	2007	Scree	ening
	Sample Study	Phase II ESA	Sediment Study	Interim Action	Lev	el2				
	Sample Type	Surface	Subsurface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	Otten Engineering	Anchor Env.	Floyd Snider	Floyd Snider	Ecology	Ecology	GeoEngineers	LAET	2LAET
Pesticides										
4,4'-DDD	μg/kg	-	<u></u>	-	-	0.99 U			NE	NE
4,4'-DDE	µg/kg	-	<u></u>	-	-	0.99 U			NE	NE
4,4'-DDT	µg/kg	-	<u></u>	-	-	0.99 U			NE	NE
Total DDT (4,4 isomers)	μg/kg	-	<u></u>		-	0.99 U			NE	NE
Aldrin	μg/kg	-	<u></u>		-	0.99 U			NE	NE
Total Chlordane	μg/kg	-	<u></u>		-	9.9 U	-		NE	NE
Dieldrin	µg/kg	-			_	0.99 U	-		NE	NE
Heptachlor	µg/kg	-			_	0.99 U			NE	NE
Polychlorinated Biphenyls (PCBs) (Dry	Weight)									
Total PCBs (Sum of Aroclors)	mg/kg	-	-	+	-	0.02 U	0.013 U	0.0013 U	0.13	1
Polychlorinated Biphenyls (PCBs) (OC	Normalized)									
Total PCBs (Sum of Aroclors)	mg/kg OC	-	-	-	-	2.11 U	5 U	1.08 U	12	65

SCO = Sediment Cleanup Objective

CSL = Cleanup Screening Level

LAET = Lowest Apparent Effects Threshold

2LAET = Second Lowest Apparent Effects Threshold

mg-N/kg = milligrams of nitrogen per kilogram

mg-N/L = milligrams of nitrogen per liter

mg/kg = milligram per kilogram

mg/kg OC = milligram per kilogram normalized to organic carbon

μg/kg = microgram per kilogram

-- = not analyzed

NE = not established

U = The analyte was not detected at a concentration greater than the value identified.

J = The analyte was detected and the detected concentration is considered an estimate.

cm = centimeters

Bold font type indicates the analyte was detected at the reported concentration.

Yellow shading indicates exceedance of the SCO/LAET screening level.

Orange shading indicates exceedance of the CSL/2LAET screening level.

Grey text indicates that the reported value is not compared to the screening levels because the TOC concentration of the sample is outside the specified range for application of the screening level.



¹ Sediment sample locations are shown on Figure 12.

 $^{^2}$ Proposed sediment cleanup levels for the protection of benthic organisms are referenced from Table 1.

³ Total LPAH represents the sum of the detected concentrations of the following LPAH compounds: acenaphthene, acenaphthylene, anthracene, fluorene, naphthalene, and phenanthrene. When all compounds are undetected, only the single highest individual chemical quantitation limit is reported. The result for 2-Methylnaphthalene is not included in the LPAH sum.

⁴ Total HPAH represents the sum of the detected concentrations of the following HPAH compounds: benz[a]anthracene, benzo[a,h,i]perylene, chrysene, dibenzo[a,h]anthracene, fluoranthene, indeno[1,2,3-c,d]pyrene, pyrene, and total benzofluoranthenes. When all compounds are undetected, only the single highest individual chemical quantitation limit is reported.

⁵ Total benzofluoranthenes represents the sum of concentrations of the b, j, and k isomers.

Table 12Sediment Contaminant of Concern Chemical Analytical Data – Protection of Benthic Organisms
Dakota Creek Industries
Anacortes, Washington

Sample	e Location ID ¹	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5		
Sample I	dentification	SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3		
	Sample Date	09/30/08	08/28/08	10/10/08	10/10/08	08/26/08	08/26/08	Prop	osed
Sample	Interval (dbm)	0 - 10 cm	Sedi	ment					
	Sample Study	2007	2007	2007	2007	2007	2007		ening
	. ,	Interim Action	Lev	/el ²					
	Sample Type	Surface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	LAET	2LAET
Conventionals									
Total Organic Carbon (TOC)	%	0.09	0.09	0.1	0.09	0.32	0.44	NE	NE
Total Volatile Solids (TVS)	%		-	-		-		NE	NE
Total Solids (TS)	%	73.2	73.2	86.7	85.7	83.6	85	NE	NE
Total Ammonia	mg-N/kg	-	-		-			NE	NE
Total Sulfide	mg/kg							NE	NE
Grain Size									
Gravel (>2,000 μm)	%	-	-	_	-		-	NE	NE
μm)	%	-		-	-			NE	NE
Coarse Sand (1,000 to 500 µm)	%	-		-	-			NE	NE
Medium Sand (500 to 250 μm)	%	-			-			NE	NE
Fine Sand (250 to 125 μ m)	%	-		-				NE	NE
Very Fine Sand (125 to 62.5 μm)	%	-		-				NE	NE
Coarse Silt (62.5 to 31 µm)	%	-		-				NE	NE
Medium Silt (31 to 15.6 μm)	%	-		-				NE	NE
Fine Silt (15.6 to 7.8 μm)	%	-		-				NE	NE
Very Fine Silt (7.8 to 3.9 μm)	%	-	-/_	-				NE	NE
Clay (3.9 to <1 μm)	%	-	-	-				NE	NE
Total Fines (<62.5 μm)	%	-	-	-				NE	NE
Metals									
Arsenic	mg/kg	-	-					57	73
Cadmium	mg/kg	0.091	0.078	0.077	0.071	0.3 U	0.3 U	5.1	6.7
Chromium	mg/kg	12.7	51.1	244	96.9	35.3	33.3	260	270
Copper	mg/kg	16.1	23.6	27.8	25.7	29.1	25.9	390	390
Lead	mg/kg	3.73	4.21	2.45	3	3 U	2.9 U	450	530
Mercury	mg/kg	0.0453	0.0221	0.032	0.036	0.041	0.036	0.41	0.59
Silver	mg/kg	0.05	0.07	0.07	0.06	0.8 U	0.8	6.1	6.1
Zinc	mg/kg	25.4	42	43.7	44.2	53	41.7	410	960



	mple Location ID ¹	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5		
Sam	ple Identification	SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3	Duan	
Com	Sample Date	09/30/08	08/28/08	10/10/08	10/10/08	08/26/08	08/26/08		osed ment
Sam	ple Interval (dbm)	0 - 10 cm 2007	0 - 10 cm 2007	0 - 10 cm 2007	0 - 10 cm 2007	0 - 10 cm 2007	0 - 10 cm 2007		ening
	Sample Study	Interim Action	Interim Action	Interim Action	Interim Action	Interim Action	Interim Action		vel ²
	Sample Type	Surface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	LAET	2LAET
Low Molecular Weight Polycyclic A	Aromatic Hydrocarbo	ons (LPAHs) (Dry Weight)						•	•
Sum of LPAHs ³	μg/kg	1.71	2.11	3.89	2.98	4.2	1.9	5,200	5,200
2-Methylnaphthalene	μg/kg	6.8	0.61 J	0.5 J	0.42 J	0.79 J	0.53 J	670	670
Acenaphthene	µg/kg	6.8	0.78 J	0.23 U	0.23 U	5.1 J	0.91 J	500	500
Acenaphthylene	µg/kg	0.45 J	0.44 J	0.24 U	0.24 U	0.39 J	0.24 U	1,300	1,300
Anthracene	μg/kg	8.3	2.5	0.47 U	0.47 U	2.8 J	0.82 J	960	960
Fluorene	μg/kg	8.6	1.3 J	0.5 U	0.5 U	4.5 J	1.2 J	540	540
Naphthalene	μg/kg	10	1.3 J	0.91 J	0.82 J	5 J	0.81 J	2,100	2,100
Phenanthrene	μg/kg	33	6.3	0.81 J	0.9 J	15 J	3.8	1,500	1,500
Low Molecular Weight Polycyclic A	Aromatic Hydrocarbo	ons (LPAHs) (OC Normalize	ed)					_	
Sum of LPAHs ³	mg/kg OC	67.1 J	13.2 J	1.72 J	1 .91 J	10.49 J	1.86 J	370	780
2-Methylnaphthalene	mg/kg OC	7.6	0.68 J	0.5 J	0.47 J	0.25 J	0.12 J	38	64
Acenaphthene	mg/kg OC	7.6	0.87 J	0.23 U	0.26 U	1.59 J	0.21 J	16	57
Acenaphthylene	mg/kg OC	0.5 J	0.49 J	0.24 U	0.27 U	0.12 J	0.05 U	66	66
Anthracene	mg/kg OC	9.2	2.8	0.47 U	0.52 U	0.88 J	0.19 J	220	1,200
Fluorene	mg/kg OC	9.6	1.4 J	0.5 U	0. 56 U	1.41 J	0.27 J	23	79
Naphthalene	mg/kg OC	11	1.4 J	0.91 J	0.91 J	1.56 J	0.18 J	99	170
Phenanthrene	mg/kg OC	37	7	0.81 J	1 J	4.69 J	0.86 J	100	480
High Molecular Weight Polycyclic	Aromatic Hydrocarbo	ons (HPAHs) (Dry Weight)							
Sum of HPAHs ⁴	µg/kg	114	40.98 J	1.82 J	1.26 J	58 J	16.97 J	12,000	17,000
Benzo(a)anthracene	μg/kg	15	5	0.48 U	0.48 U	3.2	0.98 J	1,300	1,600
Benzo(a)pyrene	µg/kg	13	5.3	0.14 U	0.14 U	2.9	0.61 J	1,600	1,600
Total Benzofluoranthenes ⁵	µg/kg	20.6	9.1	0.25 U	0.25 U	6.7	1.72	3,200	3,600
Benzo(g,h,i)perylene	µg/kg	9.9	3.5	0.64 U	0.64 U	2.3	0.64 U	670	720
Chrysene	µg/kg	16.0	7.1	0.47 J	0.25 U	5.4	1.8	1,400	2,800
Dibenzo(a,h)anthracene	µg/kg	3.5	0.98 J	0.28 U	0.28 U	0.5 J	0.28 U	230	230
Fluoranthene	µg/kg	36	10	0.63 J	0.61 J	14	4.4	1,700	2,500
Indeno(1,2,3-c,d)pyrene	μg/kg	8.9	4	0.16 U	0.16 U	2.3	0.54 J	600	690
Pyrene	μg/kg	40	11	0.72 J	0.65 J	14	3.7	2,600	3,300
High Molecular Weight Polycyclic	· ·				T	T		1	T
Sum of HPAHs ⁴	mg/kg OC	126.7	45.5 J	1.83 J	1.4 J	18.1 J	3.86 J	960	5,300
Benzo(a)anthracene	mg/kg OC	16.7	5.56	0.48 U	0.53 U	1.0	0.22 J	110	270
Benzo(a)pyrene	mg/kg OC	14.4	5.89	0.14 U	0.16 U	0.91	0.14 J	99	210
Total Benzofluoranthenes ⁵	mg/kg OC	22.9	10.1	0.25 U	0.28 U	2.09	0.39	230	450
Benzo(g,h,i)perylene	mg/kg OC	11	3.89	0.64 U	0.71 U	0.72	0.15 U	31	78
Chrysene	mg/kg OC	17.8	7.89	0.47 J	0.28 U	1.69	0.41	110	460
Dibenzo(a,h)anthracene	mg/kg OC	3.89	1.09 J	0.28 U	0.31 U	0.16 J	0.06 U	12	33
Fluoranthene	mg/kg OC	40	11.1	0.63 J	0.68 J	4.38	1.0	160	1,200
Indeno(1,2,3-c,d)pyrene	mg/kg OC	9.89	4.44	0.16 U	0.18 U	0.72	0.12 J	34	88
Pyrene	mg/kg OC	44.4	12.2	0.72 J	0.72 J	4.38	0.84	1,000	1,400



Samp	le Location ID ¹	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5		
Sample	Identification	SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3		
	Sample Date	09/30/08	08/28/08	10/10/08	10/10/08	08/26/08	08/26/08	Prop	osed
Sample	Interval (dbm)	0 - 10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	Sedi	ment
	Sample Study	2007	2007	2007	2007	2007	2007		ening
		Interim Action	Interim Action	Interim Action	Interim Action	Interim Action	Interim Action	Le	vel ²
	Sample Type	Surface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	LAET	2LAET
Chlorinated Hydrocarbons (Dry Weigl	i i	0.011	0.011	1 0011	0.011	0.011	0.011	24	
1,2,4-Trichlorobenzene 1,2-Dichlorobenzene	µg/kg	2.6 U 2.9 U	2.6 U 2.9 U	2.6 U 2.9 U	2.6 U 2.9 U	2.6 U 2.9 U	2.6 U 5.9 U	31 35	51 50
1,2-Dichlorobenzene	µg/kg	2.9 U	2.9 U	2.9 U	2.9 U	2.9 U	2.9 U	110	110
,	µg/kg								
Hexachlorobenzene	µg/kg	1.2 U	1.2 U	1.2 U	1.2 U	1.2 U	1.2 U	22	70
Chlorinated Hydrocarbons (OC Norma		2.9 U	2.9 U	2.6 U	2011	0.8 0	0.6 U	0.81	1.0
1,2,4-Trichlorobenzene 1,2-Dichlorobenzene	mg/kg OC	3.2 U	3.2 U	2.6 U	2.9 U 3.2 U	0.8 U	1.3 U	2.3	1.8 2.3
1,4-Dichlorobenzene	mg/kg OC	3.2 U	3.2 U	2.9 U	3.20	0.9 U	0.7 U	3.1	9
,	mg/kg OC				1.3 U				2.3
Hexachlorobenzene	mg/kg OC	1.3 U	1.3 U	1.2 U	1.30	0.4 U	0.3 U	0.38	2.3
Phthalates (Dry Weight)	ug/kg	9.4 J	7 U	7 U	18	7 U	7 U	1,300	1,900
Bis(2-Ethylhexyl) Phthalate Butyl benzyl Phthalate	µg/kg	3.2 U	3.2 U	3.2 U	3.2 U	3.2 U	3.2 U	63	900
,	µg/kg	7.9 U	7.9 U	7.9 U	7.9 U	8.4 J	8.9 J	1,400	1.400
Dibutyl Phthalate	µg/kg				1.4			200	,
Diethyl Phthalate	µg/kg	1.3 U	1.3 U	1.5 U		1.4 J	1.4 J	71	> 1,200
Dimethyl Phthalate	µg/kg	1 U	1 U	10	1 U 1.7 U	1 U	1 U		160
Di-N-Octyl Phthalate	μg/kg	1.7 U	1.7 U	1.7 U	1.70	1.7 U	1.7 U	6,200	6,200
Phthalates (OC Normalized)	m 4 /14 00	40.41	7,78 U	7.11	20.00	2.10.11	1 50 11	1 47	70
Bis(2-Ethylhexyl) Phthalate	mg/kg OC	10.4 J 3.56 U	3.56 U	3.2 U	20.00 3.56 U	2.19 U 1 U	1.59 U 0.73 U	47 5	78 64
Butyl benzyl Phthalate	mg/kg OC	8.78 U	8.78 U	7.9	8.78 U	2.63	2.02	220	1,700
Dibutyl Phthalate Diethyl Phthalate	mg/kg OC	1.44 U	1.44 U	1.5 U	1.56	0.44	0.32	61	110
Dimethyl Phthalate	mg/kg OC	1.11 U	1.14 U		1.11 U	0.44 0.31 U	0.23 U	53	53
Di-N-Octyl Phthalate	mg/kg OC	1.89 U	1.89 //	1.7 U	1.11 U	0.53 U	0.23 U	58	4,500
Phenols (Dry Weight)	mg/kg OC	1.09 U	1.09.0	1.70	1.09 0	0.55 0	0.59 0	56	4,500
2,4-Dimethylphenol	ug/kg	5.5 U	5.5 U	5.5 U	5.5 U	5.5 U	5.5 U	29	29
2-methylphenol (o-Cresol)	µg/kg µg/kg	1.5 U	1.5 U	1.5 U	1.5 U	1.5 U	1.5 U	63	63
4-methylphenol (p-Cresol)	μg/kg μg/kg	1.5 U	1.5 U	1.5 U	1.5 U	1.5 U	1.5 U	670	670
Pentachlorophenol	µg/kg	20 U	20 U	20 U	20 U	20 U	20 U	360	690
Phenol	μg/kg	2 U	2 U	2 U	2 U	2 U	2 U	420	1,200
Miscellaneous Extractables (Dry Weig		20	20	2.0	2 0	2 0	20	720	1,200
Dibenzofuran	µg/kg	6.3	0.85 J	0.59 U	0.59 U	2.8 J	1.2 U	540	540
Hexachlorobutadiene	µg/kg µg/kg	2.5 U	2.5 U	2.5 U	2.5 U	2.5 U	2.5 U	11	120
N-Nitrosodiphenylamine	µg/kg µg/kg	1.6 U	1.6 U	1.6 U	1.6 U	1.6 U	1.6 U	28	40
Benzoic Acid	μg/kg	96 U	96 U	96 U	96 U	96 U	96 U	650	650
Benzyl Alcohol	µg/kg	2.1 U	2.1 U	2.1 U	2.1 U	2.1 U	2.1 U	57	73
Miscellaneous Extractables (OC Norn		2.1 0	2.10	2.10	2.10	1 2.10	2.10	1 01	1 .5
Dibenzofuran	mg/kg OC	7.00	0.94 J	0.59 U	0.66 U	0.88 J	0.27 U	15	58
Hexachlorobutadiene	mg/kg OC	2.78 U	2.78 U	2.5 U	2.78 U	0.78 U	0.57 U	3.9	6.2
N-Nitrosodiphenylamine	mg/kg OC	1.78 U	1.78 U	1.6 U	1.78 U	0.5 U	0.36 U	11	11



Sample	e Location ID ¹	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5		
Sample I	Identification	SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3		
	Sample Date	09/30/08	08/28/08	10/10/08	10/10/08	08/26/08	08/26/08	Prop	osed
Sample 1	Interval (dbm)	0 - 1 0 cm	0 - 10 cm	0 - 1 0 cm	0 - 1 0 cm	0 - 1 0 cm	0 - 10 cm	Sedi	ment
	Sample Study	2007	2007	2007	2007	2007	2007	Scree	ening
	oumpio otuay	Interim Action	Interim Action	Interim Action	Interim Action	Interim Action	Interim Action	Lev	el2
	Sample Type	Surface	Surface	Surface	Surface	Surface	Surface	SCO/	CSL/
	Sampled By	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	LAET	2LAET
Pesticides									
4,4'-DDD	μg/kg	-		-	-/	-		NE	NE
4,4'-DDE	µg/kg	-			-	-		NE	NE
4,4'-DDT	µg/kg	-			_	-		NE	NE
Total DDT (4,4 isomers)	μg/kg	1		-		-		NE	NE
Aldrin	µg/kg	-	-	-		-		NE	NE
Total Chlordane	µg/kg	-		-				NE	NE
Dieldrin	µg/kg	-		-	-	-		NE	NE
Heptachlor	μg/kg	-		-	_	_		NE	NE
Polychlorinated Biphenyls (PCBs) (Dry	Weight)								
Total PCBs (Sum of Aroclors)	mg/kg	0.0031	0.0013 U	0.0013 U	0.0013 U	0.0013 U	0.0013 U	0.13	1
Polychlorinated Biphenyls (PCBs) (OC	Normalized)								
Total PCBs (Sum of Aroclors)	mg/kg OC	3.44	1.44 U	1.3 U	1.44 U	0.41 U	0.3 U	12	65

SCO = Sediment Cleanup Objective

CSL = Cleanup Screening Level

LAET = Lowest Apparent Effects Threshold

2LAET = Second Lowest Apparent Effects Threshold

mg-N/kg = milligrams of nitrogen per kilogram

mg-N/L = milligrams of nitrogen per liter

mg/kg = milligram per kilogram

mg/kg OC = milligram per kilogram normalized to organic carbon

μg/kg = microgram per kilogram

-- = not analyzed

NE = not established

U = The analyte was not detected at a concentration greater than the value identified.

J = The analyte was detected and the detected concentration is considered an estimate.

cm = centimeters

Bold font type indicates the analyte was detected at the reported concentration.

Yellow shading indicates exceedance of the SCO/LAET screening level.

Orange shading indicates exceedance of the CSL/2LAET screening level.

Grey text indicates that the reported value is not compared to the screening levels because the TOC concentration of the sample is outside the specified range for application of the screening level.



¹ Sediment sample locations are shown on Figure 12.

 $^{^2}$ Proposed sediment cleanup levels for the protection of benthic organisms are referenced from Table 1.

³ Total LPAH represents the sum of the detected concentrations of the following LPAH compounds: acenaphthene, acenaphthylene, anthracene, fluorene, naphthalene, and phenanthrene. When all compounds are undetected, only the single highest individual chemical quantitation limit is reported. The result for 2-Methylnaphthalene is not included in the LPAH sum.

⁴ Total HPAH represents the sum of the detected concentrations of the following HPAH compounds: benz[a]anthracene, benzo[a]pyrene, benzo[a,h,i]perylene, chrysene, dibenzo[a,h]anthracene, fluoranthene, indeno[1,2,3-c,d]pyrene, pyrene, and total benzofluoranthenes. When all compounds are undetected, only the single highest individual chemical quantitation limit is reported.

⁵ Total benzofluoranthenes represents the sum of concentrations of the b, j, and k isomers.

Table 13Sediment Contaminant of Concern Chemical Analytical Data – Protection of Human Health and Higher Trophic Level Ecological Receptors

Dakota Creek Industries

Anacortes, Washington

Sample Identification Sample Date Sample Interval (dbm) Sample Study Sample Type Sampled By Metals Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg Silver mg/kg	DC-SED-09 08/06/97 0-10 cm 1997 Phase II ESA Surface Otten Engineering 2.1 J ND 13.4 15.3 6.23 J	AN-P1-2 07/15/04 1-3 ft 2004 Sediment Study Subsurface Anchor Env.	DCI06-2A N/A 0-10 cm 2007 Sediment Study Surface Floyd Snider	DCI06-2-D N/A 0-10 cm 2007 Sediment Study Surface Floyd Snider	FB-A4-14 09/06/07 0 - 10 cm 2007 Sediment Study Surface Ecology	FB-A4-15 09/06/07 0 - 10 cm 2007 Sediment Study Surface Ecology	SMA 1-1 09/30/08 0 - 10 cm 2007 Interim Action Surface GeoEngineers	Proposed Sediment Screening
Sample Interval (dbm) Sample Study Sample Type Sampled By Metals Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	0-10 cm 1997 Phase II ESA Surface Otten Engineering 2.1 J ND 13.4 15.3 6.23 J	1-3 ft 2004 Sediment Study Subsurface Anchor Env.	0-10 cm 2007 Sediment Study Surface Floyd Snider	0-10 cm 2007 Sediment Study Surface	0 - 10 cm 2007 Sediment Study Surface	0 - 10 cm 2007 Sediment Study Surface	0 - 10 cm 2007 Interim Action Surface	Sediment Screening
Sample Study Sample Type Sampled By Metals Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	1997 Phase II ESA Surface Otten Engineering 2.1 J ND 13.4 15.3 6.23 J	2004 Sediment Study Subsurface Anchor Env.	2007 Sediment Study Surface Floyd Snider	2007 Sediment Study Surface	2007 Sediment Study Surface	2007 Sediment Study Surface	2007 Interim Action Surface	Sediment Screening
Study Sample Type Sampled By Metals Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	Phase II ESA Surface Otten Engineering 2.1 J ND 13.4 15.3 6.23 J	Sediment Study Subsurface Anchor Env.	Sediment Study Surface Floyd Snider	Sediment Study Surface	Sediment Study Surface	Sediment Study Surface	Interim Action Surface	Sediment Screening
Sample Type Sampled By Metals Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	Surface Otten Engineering 2.1 J ND 13.4 15.3 6.23 J	Subsurface Anchor Env.	Surface Floyd Snider	Surface	Surface	Surface	Surface	Sediment Screening
Metals Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	2.1 J ND 13.4 15.3 6.23 J	Anchor Env.	Floyd Snider 					_
MetalsArsenicmg/kgCadmiummg/kgChromiummg/kgCoppermg/kgLeadmg/kgMercurymg/kg	2.1 J ND 13.4 15.3 6.23 J		-	Floyd Snider	Ecology	Ecology	GeoEngineers	2
Arsenic mg/kg Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	ND 13.4 15.3 6.23 J	-						Level ²
Cadmium mg/kg Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	ND 13.4 15.3 6.23 J	-		_				
Chromium mg/kg Copper mg/kg Lead mg/kg Mercury mg/kg	13.4 15.3 6.23 J		-		2.37	2.1	4.40	11
Copper mg/kg Lead mg/kg Mercury mg/kg	15.3 6.23 J			-	0.09	0.08 U	0.054	0.8
Lead mg/kg Mercury mg/kg	6.23 J		1	-	12.3	13.5	26.2	6,900,000
Mercury mg/kg			1	-	10.4	9.49	27.4	180,000
			1	1	2.23 J	1.63	4.08	21
Silver mg/kg	ND		-		0.01 J	0.01	0.033	0.2
	0.0582				0.02 U	0.03	0.07	23,000
Zinc mg/kg	21.3		-	-	20.9	19.1	43.9	1,400,000
Organometallic Compounds								
Tributyltin, bulk µg/kg	-		-	-	-		7.24 J	73
Interstitial Tributyltin, porewater µg/L			-	1	-			0.15
Low Molecular Weight Polycyclic Aromatic Hydroca	arbons (LPAHs)							
2-Methylnaphthalene μg/kg		-	-		9.7 U	9.1 U	0.85 J	16,000,000
Acenaphthene µg/kg	-				9.7 U	9.1 U	0.91 J	240,000,000
Acenaphthylene µg/kg	-		-	-	1.4 J	9.1 U	0.24 U	240,000,000
Anthracene µg/kg	-	_	-		9.7 U	9.1 U	0.5 J	1,200,000,000
Fluorene µg/kg	-	-	-	_	9.7 U	9.1 U	1.1 J	160,000,000
Naphthalene µg/kg	-	-	-	-	9.7 U	9.1 U	2.5	79,000,000
Phenanthrene µg/kg	-	-	-	_	8.2 J	9.1 U	2.9	1,200,000,000
High Molecular Weight Polycyclic Aromatic Hydroc	carbons (HPAHs)							
Benzo(a)anthracene µg/kg	-	-		-	4.1 J	9.1 U	1.3 J	NE
Benzo(a)pyrene µg/kg	-	-	-	_	2.6 J	9.1 U	0.86 J	cPAH TEQ
Total Benzofluoranthenes ³ µg/kg		-	-	-	9.9 J	9.1 U	1.87 J	NE
Benzo(g,h,i)perylene µg/kg	-			_	2 J	9.1 U	0.94 J	120,000,000
Chrysene µg/kg	-	-	-		18	9.1 U	1.1 J	NE
Dibenzo(a,h)anthracene µg/kg	-	/	-		9.7 U	9.1 U	0.28 U	NE
Fluoranthene µg/kg	-	-	-	-	50	2.2 J	3	160,000,000
Indeno(1,2,3-c,d)pyrene µg/kg	-	-	_	-	1.8 J	9.1 U	0.83 J	NE
Pyrene µg/kg		-	-	-	33	9.1 U	4.0	120,000,000
Carcinogenic PAHs (cPAHs)	•						-	
Total cPAH TEQ ⁴ (ND=0.5 RL) µg/kg			1					



Sampl	le Location ¹	DC-SED-09	P1-2	DC-106-2	DC-106-2	FB-A4-14	FB-A4-15	SMA-1	
Sample Ide	entification	DC-SED-09	AN-P1-2	DCI06-2A ²	DCI06-2-D ²	FB-A4-14	FB-A4-15	SMA 1-1	
S	ample Date	08/06/97	07/15/04	N/A	N/A	09/06/07	09/06/07	09/30/08	
Sample In	terval (dbm)	0-10 cm	1-3 ft	0-10 cm	0-10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	
	Sample	1997	2004	2007	2007	2007	2007	2007	Proposed
	Study	Phase II ESA	Sediment Study	Sediment Study	Sediment Study	Sediment Study	Sediment Study	Interim Action	Sediment
	ample Type	Surface	Subsurface	Surface	Surface	Surface	Surface	Surface	Screening
	Sampled By	Otten Engineering	Anchor Env.	Floyd Snider	Floyd Snider	Ecology	Ecology	GeoEngineers	Level ²
Chlorinated Hydrocarbons	1		T	T				T	
1,2,4-Trichlorobenzene	µg/kg	-		-	-	9.7 U	9.1 U	2.6 U	140,000
1,2-Dichlorobenzene	µg/kg			-	_	9.7 U	9.1 U	2.9 U	370,000,000
1,4-Dichlorobenzene	µg/kg	<u></u>			-	9.7 U	9.1 U	2.9 U	780,000
Hexachlorobenzene	µg/kg				-	9.7 U	9.1 U	1.2 U	2,500
Phthalates			-	-				-	
Bis(2-Ethylhexyl) Phthalate	µg/kg				-	97	91	7 U	290,000
Butyl benzyl Phthalate	µg/kg		-	-		9.7 U	9.1 U	3.2 U	2,100,000
Dibutyl Phthalate	μg/kg			-	-	9.2 J	16	7.9 U	410,000,000
Diethyl Phthalate	µg/kg				-	1 .9 J	9.1 U	1.3 U	3,100,000,000
Dimethyl Phthalate	µg/kg					1 J	9.1 U	1 U	NE
Di-N-Octyl Phthalate	µg/kg			-	-	9.7 U	9.1 U	1.7 U	41,000,000
Phenols									
2,4-Dimethylphenol	µg/kg			-	-		46 R	5.5 U	82,000,000
2-methylphenol (o-Cresol)	µg/kg		-	-	-	-	9.1 U	1.5 U	200,000,000
4-methylphenol (p-Cresol)	μg/kg			-	_		9.1 U	3.4 J	390,000,000
Pentachlorophenol	μg/kg	_		-	-		91 U	20 U	10,000,000
Phenol	µg/kg		-	-	-	-	28 U	2 U	1,200,000,000
Miscellaneous Extractables									
Dibenzofuran	μg/kg		-/	-	-	9.7 U	9.1 U	0.87 J	4,100,000
Hexachlorobutadiene	µg/kg	-	-	-	-	9.7 U	9.1 U	2.5 U	52,000
N-Nitrosodiphenylamine	µg/kg	-	_	-	-	9.7 U	9.1 U	1.6 U	830,000
Benzoic Acid	µg/kg	_	-	-		200	190 R	96 U	16,000,000,000
Benzyl Alcohol	µg/kg	_				6.2 J	19 U	2.1 U	410,000,000
Polychlorinated Biphenyls (PCBs)	100			-	L			L	, ,
Total PCBs (Sum of Aroclors or Congeners)	mg/kg	-		-	-	0.02 U	0.013 U	0.0013 U	0.0035
Dioxins and Furans					<u> </u>			•	<u> </u>
2,3,7,8-TCDD	ng/kg		1 U	0.21 U	0.18 U				NE
1,2,3,7,8-PeCDD	ng/kg	-	2.5 U	1.1 U	0.91 U				NE
1,2,3,4,7,8-HxCDD	ng/kg	-	2.5 U	1.1 U	0.91 U				NE
1,2,3,6,7,8-HxCDD	ng/kg	-	2.5 U	1.1 U	0.91 U				NE
1,2,3,7,8,9-HxCDD	ng/kg	-	2.5 U	1.1 U	0.91 U				NE
1,2,3,4,6,7,8-HpCDD	ng/kg	_	2.5 U	2 J	5	_			NE NE
OCDD	ng/kg		5 U	14	35	_			NE NE
2,3,7,8-TCDF	ng/kg		1U	0.21 U	0.18 U	_		_	NE NE
1,2,3,7,8-PeCDF	ng/kg	<u></u>	2.5 U	1.1 U	0.91 U				NE NE
2,3,4,7,8-PeCDF	ng/kg		2.5 U	1.1 U	0.91 U			_	NE NE
1,2,3,4,7,8-HxCDF	ng/kg		2.5 U	1.1 U	0.91 U				NE NE
1,2,3,6,7,8-HxCDF	ng/kg		2.5 U	1.1 U	0.91 U				NE NE
2,3,4,6,7,8-HxCDF	ng/kg		2.5 U	1.1 U	0.91 U			-	NE NE



Samp	le Location ¹	DC-SED-09	P1-2	DC-106-2	DC-106-2	FB-A4-14	FB-A4-15	SMA-1	
Sample Id	entification	DC-SED-09	AN-P1-2	DCI06-2A ²	DCI06-2-D ²	FB-A4-14	FB-A4-15	SMA 1-1	
	Sample Date	08/06/97	07/15/04	N/A	N/A	09/06/07	09/06/07	09/30/08	
Sample In	terval (dbm)	0-10 cm	1-3 ft	0-10 cm	0-10 cm	0 - 10 cm	0 - 10 cm	0 - 10 cm	
	Sample	1997	2004	2007	2007	2007	2007	2007	Proposed
	Study	Phase II ESA	Sediment Study	Sediment Study	Sediment Study	Sediment Study	Sediment Study	Interim Action	Sediment
!	Sample Type	Surface	Subsurface	Surface	Surface	Surface	Surface	Surface	Screening
	Sampled By	Otten Engineering	Anchor Env.	Floyd Snider	Floyd Snider	Ecology	Ecology	GeoEngineers	Level ²
1,2,3,7,8,9-HxCDF	ng/kg		2.5 U	1.1 U	0.91 U	-	-		NE
1,2,3,4,6,7,8-HpCDF	ng/kg		2.5 U	1.1 U	0.91 U	-	-		NE
1,2,3,4,7,8,9-HpCDF	ng/kg		2.5 U	1.1 U	0.91 U		-		NE
OCDF	ng/kg	-	9.1	2.1 U	2.2 J				NE
Total Dioxin/Furan TEQ ⁵	ng/kg		3.13	1.32 J	1.09 J		_		_

ng/kg = nanogram per kilogram

mg/kg = milligram per kilogram

μg/kg = microgram per kilogram

- = not analyzed

NE = not established

ND = Not detected

RL = Reporting limit

ft = feet

U = The analyte was not detected at a concentration greater than the value identified.

J = The analyte was detected and the detected concentration is considered an estimate.

Bold font type indicates the analyte was detected at the reported concentration.

Yellow shading indicates that the identified concentration is greater than the sediment screening level for protection of human health (HH) and higher trophic level ecological receptors (HTLER).



¹ Sediment sample locations are shown on Figure 12.

² Proposed sediment cleanup levels for the protection of human health and higher trophic level ecological receptors are referenced from Table 2.

 $^{^{\}rm 3}\,\text{Total}$ benzofluoranthenes represents the sum of concentrations of the b, j, and k isomers.

⁴ Total cPAH Toxic Equivalency Quotients (TEQs) were calculated using Toxicity Equivalency Factors (TEFs) values referenced from MTCA Table 708.2 (WAC 173-340-900).

⁵ Total dioxin and furan TEQs were calculated using United States Environmental Protection Agency (USEPA) TEF values for human health (EPA, 2003).

Table 13Sediment Contaminant of Concern Chemical Analytical Data – Protection of Human Health and Higher Trophic Level Ecological Receptors

Dakota Creek Industries

Anacortes, Washington

Sample	e Location ¹	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5	
Sample Ide	entification	SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3	
Si	ample Date	09/30/08	08/28/08	10/10/08	10/10/08	08/26/08	08/26/08	
Sample Into	erval (dbm)	0 - 10 cm						
	Sample	2007	2007	2007	2007	2007	2007	Proposed
	Study	Interim Action	Sediment					
Si	ample Type	Surface	Surface	Surface	Surface	Surface	Surface	Screening
	Sampled By	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	Level ²
Metals								
Arsenic	mg/kg	1.71	2.11	3.89	2.98	4.2	1.9	11
Cadmium	mg/kg	0.091	0.078	0.077	0.071	0.3 U	0.3 U	0.8
Chromium	mg/kg	12.7	51.1	244	96.9	35.3	33.3	6,900,000
Copper	mg/kg	16.1	23.6	27.8	25.7	29.1	25.9	180,000
Lead	mg/kg	3.73	4.21	2.45	3	3 U	2.9 U	21
Mercury	mg/kg	0.0453	0.0221	0.032	0.036	0.041	0.036	0.2
Silver	mg/kg	0.05	0.07	0.07	0.06	0.8 U	0.8	23,000
Zinc	mg/kg	25.4	42	43.7	44.2	53	41.7	1,400,000
Organometallic Compounds								
Tributyltin, bulk	µg/kg	60.35 J	11.84 J	1.72 J	1.72 J	33.58 J	8.19 J	73
Interstitial Tributyltin, porewater	μg/L	-		-	-		-	0.15
Low Molecular Weight Polycyclic Arom	natic Hydrocar	bons (LPAHs)						
2-Methylnaphthalene	µg/kg	6.8	0.61 J	0.5 J	0.42 J	0.79 J	0.53 J	16,000,000
Acenaphthene	µg/kg	6.8	0.78 J	0.23 U	0.23 U	5.1 J	0.91 J	240,000,000
Acenaphthylene	µg/kg	0.45 J	0.44 J	0.24 U	0.24 U	0.39 J	0.24 U	240,000,000
Anthracene	µg/kg	8.3	2.5	0.47 U	0.47 U	2.8 J	0.82 J	1,200,000,000
Fluorene	µg/kg	8.6	1.3 J	0.5 U	0.5 U	4.5 J	1.2 J	160,000,000
Naphthalene	µg/kg	10	1.3 J	0.91 J	0.82 J	5 J	0.81 J	79,000,000
Phenanthrene	µg/kg	33	6.3	0.81 J	0.9 J	15 J	3.8	1,200,000,000
High Molecular Weight Polycyclic Aron	natic Hydroca	rbons (HPAHs)						
Benzo(a)anthracene	µg/kg	15	5	0.48 U	0.48 U	3.2	0.98 J	NE
Benzo(a)pyrene	µg/kg	13	5.3	0.14 U	0.14 U	2.9	0.61 J	cPAH TEQ
Total Benzofluoranthenes ³	µg/kg	20.6	9.1	0.25 U	0.25 U	6.7	1.72	NE
Benzo(g,h,i)perylene	µg/kg	9.9	3.5	0.64 U	0.64 U	2.3	0.64 U	120,000,000
Chrysene	µg/kg	16.0	7.1	0.47 J	0.25 U	5.4	1.8	NE
Dibenzo(a,h)anthracene	µg/kg	3.5	0.98 J	0.28 U	0.28 U	0.5 J	0.28 U	NE
Fluoranthene	µg/kg	36	10	0.63 J	0.61 J	14	4.4	160,000,000
Indeno(1,2,3-c,d)pyrene	µg/kg	8.9	4	0.16 U	0.16 U	2.3	0.54 J	NE
Pyrene	µg/kg	40	11	0.72 J	0.65 J	14	3.7	120,000,000
Carcinogenic PAHs (cPAHs)								
Total cPAH TEQ ⁴ (ND=0.5 RL)	µg/kg	17.96	7.28 J	0.005 J	0.003 J	4.22 J	0.95 J	21



Sample Identification	Sample Location	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5	
Sample Date 09/30/08 08/28/08 10/10/08 10/10/08 08/26		SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3	1
Sample Interval (dbm) O-10 cm Sample D-10 cm 2007 20	•			10/10/08	10/10/08			
Study	Sample Interval (di							
Sample Type Surface	Sam	2007	2007	2007	2007	2007	2007	Proposed
Chlorinated Hydrocarbons	Str	Interim Action	Sediment					
Chlorinated Hydrocarbons 1,2,4 Trichlorobenzene	Sample Ty							Screening
1.2.4-Trichlorobenzene	Sampled	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	Level ²
1.2-Dichlorobenzene	· · · · · · · · · · · · · · · · · · ·			T			T	
1.4-Dichlorobenzene								·
Hexachlorobenzene	13							370,000,000
Phthalates Bis(2-Ethylhexyl) Phthalate μg/kg 9.4 J 7 U 7 U 18 7 U 7 U 290,000	4-Dichlorobenzene μg/	2.9 U	2.9 U	2.9 U		2.9 U	2.9 U	
Bis(2-Ethylhexyl) Phthalate	exachlorobenzene µg/	1.2 U	2,500					
Butyl benzyl Phthalate	nalates							
Dibutyl Phthalate	s(2-Ethylhexyl) Phthalate µg/	9.4 J	7 U	7 U	18	7 U	7 U	290,000
Diethyl Phthalate	ıtyl benzyl Phthalate μg/	3.2 U				3.2 U	3.2 U	2,100,000
Dimethyl Phthalate μg/kg 1 U 1 U 1 U 1 U 1 U 1 U 1 U 1 U NE	butyl Phthalate μg/	7.9 U	7.9 U	7.9 U	7.9 U	8.4 J	8.9 J	410,000,000
Di-N-Octyl Phthalate	ethyl Phthalate µg/	1.3 U	1.3 U	1.5 U	1,4	1.4 J	1.4 J	3,100,000,000
Phenois 2,4-Dimethylphenol μg/kg 5.5 U 5.5 U 5.5 U 5.5 U 5.5 U 82,000,000 2-methylphenol (o-Cresol) μg/kg 1.5 U 2.0 U <td>methyl Phthalate µg/</td> <td>1 U</td> <td>1 U</td> <td>1 U</td> <td>10</td> <td>1 U</td> <td>1 U</td> <td>NE</td>	methyl Phthalate µg/	1 U	1 U	1 U	10	1 U	1 U	NE
2,4-Dimethylphenol μg/kg 5.5 U 82,000,000 2-methylphenol (o-Cresol) μg/kg 1.5 U 2.0	-N-Octyl Phthalate µg/	1.7 U	41,000,000					
2-methylphenol (o-Cresol) μg/kg 1.5 U 1.5 U 1.5 U 1.5 U 1.5 U 200,000,000 4-methylphenol (p-Cresol) μg/kg 1.5 U 1.5 U 1.5 U 1.5 U 1.5 U 1.5 U 390,000,000 Pentachlorophenol μg/kg 20 U	nols							
4-methylphenol (p-Cresol) μg/kg 1.5 U 1.5 U 1.5 U 1.5 U 1.5 U 390,000,000 Pentachlorophenol μg/kg 20 U <	4-Dimethylphenol μg/	5.5 U	82,000,000					
Pentachlorophenol μg/kg 20 U 20 U </td <td>methylphenol (o-Cresol) µg/</td> <td>1.5 U</td> <td>1.5 U</td> <td>1.5 U</td> <td>1.5 U</td> <td>1.5 U</td> <td>1.5 U</td> <td>200,000,000</td>	methylphenol (o-Cresol) µg/	1.5 U	200,000,000					
Phenol μg/kg 2 U 2 U 2 U 2 U 2 U 2 U 2 U 2 U 2 U 1,200,000,00 Miscellaneous Extractables Dibenzofuran μg/kg 6.3 0.85 J 0.59 U 0.59 U 2.8 J 1.2 U 4,100,000 Hexachlorobutadiene μg/kg 2.5 U 52,000 N-Nitrosodiphenylamine μg/kg 1.6 U 1.6 U 1.6 U 1.6 U 1.6 U 1.6 U 96 U 410,000,000 2.1 U <	methylphenol (p-Cresol) µg/	1.5 U	1.5 U	1.5 U	1 .5 U	1.5 U	1.5 U	390,000,000
Miscellaneous Extractables Dibenzofuran μg/kg 6.3 0.85 J 0.59 U 0.59 U 2.8 J 1.2 U 4,100,000 Hexachlorobutadiene μg/kg 2.5 U 2.5 U 2.5 U 2.5 U 2.5 U 2.5 U 52,000 N-Nitrosodiphenylamine μg/kg 1.6 U 96 U 2.1 U 2.1 U 2.1 U 2.1 U 410,000,000	entachlorophenol µg/	20 U	10,000,000					
Dibenzofuran μg/kg 6.3 0.85 J 0.59 U 0.59 U 2.8 J 1.2 U 4,100,000 Hexachlorobutadiene μg/kg 2.5 U 52,000 N-Nitrosodiphenylamine μg/kg 1.6 U 1.6 U 1.6 U 1.6 U 1.6 U 1.6 U 830,000 Benzoic Acid μg/kg 96 U 2.1 U 2.1 U 410,000,000 <td>nenol µg/</td> <td>2 U</td> <td>2 U</td> <td>2 U</td> <td>2 U</td> <td>2 U</td> <td>2 U</td> <td>1,200,000,000</td>	nenol µg/	2 U	2 U	2 U	2 U	2 U	2 U	1,200,000,000
Hexachlorobutadiene μg/kg 2.5 U 2.5 U 2.5 U 2.5 U 2.5 U 52,000 N-Nitrosodiphenylamine μg/kg 1.6 U 96 U 2.1 U 410,000,000	cellaneous Extractables							
N-Nitrosodiphenylamine μg/kg 1.6 U 1.6	benzofuran µg/	6.3	0.85 J	0.59 U	0.59 U	2.8 J	1.2 U	4,100,000
Benzoic Acid μg/kg 96 U 16,000,000,00 96 U 96 U 96 U 96 U 96 U 16,000,000,00 96 U 96 U 96 U 96 U 96 U 410,000,000,00 96 U 96 U 96 U 96 U 96 U 410,000,000,00 96 U 96 U 96 U 96 U 96 U 410,000,000,00 96 U 96 U 96 U 96 U 96 U 410,000,000,00 96 U <	exachlorobutadiene µg/	2.5 U	52,000					
Benzyl Alcohol μg/kg 2.1 U 2.1 U 2.1 U 2.1 U 2.1 U 2.1 U 410,000,00	Nitrosodiphenylamine µg/	1.6 U	830,000					
	enzoic Acid µg/	96 U	16,000,000,000					
Polychlarinated Rinhanyle (PCRs)	enzyl Alcohol µg/	2.1 U	410,000,000					
	ychlorinated Biphenyls (PCBs)							
Total PCBs (Sum of Aroclors or Congeners) mg/kg 0.0031 0.0013 U	roclors or Congeners) mg/	0.0031	0.0013 U	0.0035				
Dioxins and Furans				_	,			
2,3,7,8-TCDD ng/kg NE		-	-					
1,2,3,7,8-PeCDD ng/kg NE	2,3,7,8-PeCDD ng/	-						NE
1,2,3,4,7,8-HxCDD	2,3,4,7,8-HxCDD ng/	-		-		-		NE
1,2,3,6,7,8-HxCDD	2,3,6,7,8-HxCDD ng/	-	-	-		-		NE
1,2,3,7,8,9-HxCDD ng/kg NE	2,3,7,8,9-HxCDD ng/	-	-	-		-		NE
1,2,3,4,6,7,8-HpCDD ng/kg NE	2,3,4,6,7,8-HpCDD ng/	-	-	-	-	-	-	NE
OCDD ng/kg NE	CDD ng/	-	-	-	-	-	-	NE
2,3,7,8-TCDF ng/kg NE	3,7,8-TCDF ng/	-	-	-	-	-	-	NE
1,2,3,7,8-PeCDF ng/kg NE	2,3,7,8-PeCDF ng/	-		-		<u></u>		NE
2,3,4,7,8-PeCDF ng/kg NE	3,4,7,8-PeCDF ng/	-	<u></u>					NE
1,2,3,4,7,8-HxCDF ng/kg NE	2,3,4,7,8-HxCDF ng/	-	<u>-</u>	-	-		-	NE
1,2,3,6,7,8-HxCDF ng/kg NE	2,3,6,7,8-HxCDF ng/	-		-			-	NE
2,3,4,6,7,8-HxCDF ng/kg NE	3,4,6,7,8-HxCDF ng/	-		-				NE



Sample	Location ¹	SMA-2	SMA-3	SMA-4	SMA-4	SMA-5	SMA-5	
Sample Ider	ntification	SMA 2-1	SMA 3-2	DCI 4-1	DCI 4-1A	SMA 5-2	SMA 5-3	
Sa	mple Date	09/30/08	08/28/08	10/10/08	10/10/08	08/26/08	08/26/08	
Sample Inte	erval (dbm)	0 - 10 cm						
	Sample	2007	2007	2007	2007	2007	2007	Proposed
Study		Interim Action	Sediment					
Sample Type		Surface	Surface	Surface	Surface	Surface	Surface	Screening
Sampled By		GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	GeoEngineers	Level ²
1,2,3,7,8,9-HxCDF	ng/kg	-		-	-			NE
1,2,3,4,6,7,8-HpCDF	ng/kg	-		-	-	-		NE
1,2,3,4,7,8,9-HpCDF	ng/kg	-		-		-		NE
OCDF	ng/kg	-				-		NE
Total Dioxin/Furan TEQ ⁵ (ND=0.5 RL)	ng/kg			-		-		5

ng/kg = nanogram per kilogram

mg/kg = milligram per kilogram

µg/kg = microgram per kilogram

-- = not analyzed

NE = not established

ND = Not detected

RL = Reporting limit

ft = feet

 $\mbox{\bf U}$ = The analyte was not detected at a concentration greater than the value identified.

J = The analyte was detected and the detected concentration is considered an estimate.

Bold font type indicates the analyte was detected at the reported concentration.

Yellow shading indicates that the identified concentration is greater than the sediment screening level for protection of human health (HH) and higher trophic level ecological receptors (HTLER).



¹ Sediment sample locations are shown on Figure 12.

² Proposed sediment cleanup levels for the protection of human health and higher trophic level ecological receptors are referenced from Table 2.

 $^{^{\}rm 3}\,\text{Total}$ benzofluoranthenes represents the sum of concentrations of the b, j, and k isomers.

⁴ Total cPAH Toxic Equivalency Quotients (TEQs) were calculated using Toxicity Equivalency Factors (TEFs) values referenced from MTCA Table 708.2 (WAC 173-340-900).

⁵ Total dioxin and furan TEQs were calculated using United States Environmental Protection Agency (USEPA) TEF values for human health (EPA, 2003).

Groundwater Contaminant of Concern Chemical Analytical Data

Monitoring	Sample		Total N	/letals ²	Dissolve	d Metals ²	Total cPAH
Well ¹	Date	Units	Arsenic	Nickel	Arsenic	Nickel	TEQ ^{3,4}
-	06/05/02	μg/L	5	3.8	4	2.2	1.41 U
	08/19/02	µg/L	0.6	4.2	_		0.07 U
	11/17/06	µg/L	3.3	2.1	_	-	0.07 U
	06/17/08	µg/L	4.8	3.2	-		0.013 U
	06/17/08 ⁶	μg/L	4.9	3.3	- /		0.013 U
	05/23/12	μg/L	17 U	8 U	15 U	8 U	0.007 U
MW-1	08/16/12	μg/L	15	8 U	15	8 U	0.007 U
(Upland Well)	11/13/12	μg/L	16 U	5.4	16 U	5 U	0.007 U
	02/13/13	μg/L	15	8 U	14	8 U	0.007 U
	02/10/16	μg/L	1.3	-	1.2 J	-	
	08/18/16	μg/L	9.2		7.8 U	_	
	02/15/17	μg/L	11	-	5.6	_	
	08/23/17	μg/L	5.6 U	-	5.6 U	_	
	06/05/02	μg/L	3	7.5	3	7.5	1.41 U
	06/05/02 ⁶	μg/L	3	7.5	3	7.5	1.41 U
	08/19/02	μg/L	4	9.9	-		0.07 U
MW-2	08/19/02 ⁶	μg/L	2	8.2	-		0.07 U
(Shoreline Well)	11/17/06	µg/L	4	3.9	-		0.007
	11/17/06 ⁶	µg/L	3.8	3.9			0.007 U
	06/17/08	μg/L	3.4	2.4			0.015
	05/23/12	μg/L	12 U	20 U	11 U	20 U	0.007 U
	08/16/12	μg/L	7.5 U	17	7.5 U	16	0.007 U
MW-2A	11/13/12	µg/L	10 U	13	10 U	13	0.007 U
(Shoreline Well)	02/13/13	µg/L	8 U	7	8 U	8 U	0.007 U
	02/10/16	μg/L	_6 	_6 	_6	_6 	_
1411/05	08/19/16	μg/L	7.8 U	7.8 U	7 U	7 U	
MW-2B (Shoreline Well)	02/15/17	µg/L	5.6 U	8.2	5 U	8.3	_
(Shoreline Well)	08/23/17	µg/L	6	6.9	6.5	6	-
	06/05/02	µg/L	1 U	3.4	0.1 U	0.33	1.41 U
MW-3	08/19/02	µg/L	1	3.7	-	-	0.07 U
(Shoreline Well)	11/17/06	µg/L	0.9	1.5	-	-	0.007 J
	06/17/08	µg/L	0.8	2.2	-	-	0.013 U
	05/23/12	µg/L	4 U	8 U	4.5 U	8 U	0.007 U
	08/16/12	µg/L	7.5 U	19	7.5 U	18	0.007 U
	11/13/12	µg/L	8 U	18	8 U	17	0.007 U
MW-3A	02/13/13	μg/L	8 U	16	8 U	18	0.007 U
(Shoreline Well)	02/11/16	μg/L	-	0.5 J	_	0.5 U	
	08/19/16	μg/L	-	7.8 U	_	7 U	
	02/16/17	μg/L	_	5.6 U	-	5 U	_
	08/24/17	µg/L		5.6 U		5 U	-
Pronosed Gr	oundwater Cleanup	_	8	8.2	8	8.2	0.01



Monitoring	Sample		Total N	/letals ²	Dissolved	d Metals ²	Total cPAH -
Well ¹	Date	Units	Arsenic	Nickel	Arsenic	Nickel	TEQ ^{3,4}
	06/05/02	μg/L	8	3.4	9	1.1	_
	08/19/02	μg/L	12	3.3		-	0.07 U
	11/17/06	μg/L	11.6	2		_	0.007 U
	06/17/08	μg/L	8.1	1.1		_	0.013 U
	05/23/12	μg/L	9 U	20 U	9.5 U	20 U	0.007 U
MW-4	08/16/12	μg/L	11	8 U	10	8 U	0.007 U
(Upland Well)	11/13/12	μg/L	10 U	5 U	10 U	5 U	0.031
	02/13/13	μg/L	8 U	8 U	8 U	8 U	0.015
	02/11/16	μg/L	3.5		3.5	-	0.008
	08/18/16	μg/L	7.8 U	-	7.8 U		0.008
	02/15/17	μg/L	5.6 U		5 U	-	0.007 U
	08/24/17	μg/L	5.6 U		5 U	-	0.007 U
MW-5 (Upland Well)	06/17/08	μg/L	10	5.2		-	0.013 U
	05/23/12	μg/L	3.5 U	20 U	3 U	20 U	0.008
	08/16/12	μg/L	7.5 U	18	7.5 U	19	0.007 U
	11/13/12	μg/L	8 U	18	8 U	18	0.007 U
MW-6	02/13/13	μg/L	8 U	8 U	8 U	8 U	0.007 U
(Shoreline Well)	02/11/16	μg/L	_	0.4 U	-	0.4 U	-
	08/19/16	μg/L	-	7.8 U	-	7 U	-
	02/16/17	μg/L	1	5.6 U	1	5 U	ı
	08/24/17	μg/L	-	5.6 U	-	5 U	1
	05/23/12	μg/L	11 U	20 U	9.8 U	20 U	0.007 U
	08/16/12	µg/L	10	27	7.5 U	27	0.0104
	11/13/12	µg/L	8 U	18	8 U	19	0.013
MW-7	02/13/13	μg/L	9	18	8	18	0.007 U
(Upland Well)	02/10/16	μg/L	12.9	9.9	13	5.2 J	-
	08/19/16	μg/L	12	11	11	10	-
	02/16/17	µg/L	9.2	6.7	7.1	6.6	-
	08/23/17	µg/L	14	8.8	12	9.3	
	02/10/16	μg/L	16.6	9.1	16.1	7.4	0.008 U
	02/10/16 ⁶	μg/L	16.1	8.4	16.3	7.8	0.008 U
	08/18/16	µg/L	16	7.8 U	15	7.8 U	0.008 U
MW-8	08/18/16 ⁶	µg/L	14	7.8 U	14	7.8 U	0.008 U
(Shoreline Well)	02/15/17	µg/L	12	5.6 U	12	5 U	0.007 U
	02/15/17 ⁶	μg/L	5.6 U	5.6 U	5 U	5 U	0.007 U
	08/23/17	μg/L	17	6.7	17	5.6 U	0.167
	08/23/17	μg/L	17	6	15	5.6 U	0.040
Proposed Gr	oundwater Cleanup I	Level ⁷	8	8.2	8	8.2	0.01



- ¹Monitoring well locations are shown on Figure 13. Groundwater results are summarized on Figures 20 through 22.
- 2 Metals analyzed using EPA Method 6010/6020.
- ³ Carcinogenic Polycyclic Aromatic Hydrocarbons (cPAHs) analyzed by EPA method 8270D/SIM.
- ⁴ Total cPAH Toxic Equivalency Quotients (TEQs) were calculated using Toxicity Equivalency Factors (TEFs) values referenced from MTCA Table 708.2 (WAC 173-340-900).
- ⁵ Monitoring well not accessible at the time of sampling.
- $^{\rm 6}$ Duplicate sample collected for laboratory analysis.
- $^{\rm 7}$ Proposed groundwater cleanup level is referenced from Table 4.
- J = Estimated value
- T = qualifier indicating total concentration
- U = qualifier indicating analyte not detected at level above listed practical quantitation limit
- µg/L = microgram per liter
- -- = not analyzed

Bold indicates analyte was detected.



Yellow shading indicates analyte was detected at a concentration above proposed groundwater cleanup level.



Table 15Soil Contaminant of Concern Chemical Analytical Data
Dakota Creek Industries

Anacortes, Washington

		Sample			Total N	/letals ²	Petro	oleum Hydroca	rbons	Total cP	AH TEQ ^{3,4}
Sample Location ¹	Sample Identification	Depth (bgs)	Sample Date	Units	Arsenic	Nickel	Gasoline- Range	Diesel- Range	Heavy Oil- Range	Vadose Zone	Saturated Zone
S-1	DC-B-1	4.5 ft	07/14/97	mg/kg	5.24	32.7					
3-1	DC-B-1B	4.5 ft	07/14/97	mg/kg	8.85	25.6					
S-2	DC-B-2	4.5 ft	07/14/97	mg/kg	2.11	22.9					
3-2	DC-B-2A	2.5 ft	07/14/97	mg/kg	1.0	5.75				ND	
SS-1A/1B	DC-UPLD SS-1A	0-1 ft	07/03/97	mg/kg	32.1 J	20.9 J	-				
33-1A/ 1B	DC-UPLD SS-1B	0-1 ft	07/03/97	mg/kg	1.74 J	45.7 J					
SS-2A/2B	DC-UPLD SS-2A	0-1 ft	07/30/97	mg/kg	15 J	27.9 J					
30-2 <i>H</i> y 2B	DC-UPLD SS-2B	0-1 ft	07/30/97	mg/kg	1.44 J	52.6 J					
SS-3	DC-UPLD SS-3	0-1 ft	07/30/97	mg/kg	3	35.7 J	ND	10.9	63.9		
SS-4	DC-UPLD SS-4	0-1 ft	07/30/97	mg/kg	7.26	21.7 J	ND	203	2,220		
SS-6	DC-UPLD SS-6	0-1 ft	07/30/97	mg/kg			ND	492	2,100		
SS-9	DC-UPLD SS-9	0-1 ft	07/30/97	mg/kg			233	8,360	4,470	ND	
SS-11	DC-UPLD SS-11 ²	0-1 ft	07/30/97	mg/kg			126	16,300	1,980	ND	
SS-13A	DC-UPLD SS-13A ²	0-1 ft	07/30/97	mg/kg	22.6	15.1 J	26.7	421	843		
SS-14A	DC-UPLD SS-14A ²	0-1 ft	07/30/97	mg/kg	27	16.5 J	22.9	1,590	18,500	ND	
SS-14B	DC-UPLD SS-14B ²	0-1 ft	07/30/97	mg/kg	1.97	23.3 J	23.1	2,900	2,820	ND	
DCI-SB-UL01	0020-LAI	2 ft	07/17/01	mg/kg	5	46.7				0.266 J	
DCI-SB-UL01	0040-LAI	4 ft	07/17/01	mg/kg	6.5	35.4				0.131 J	
DCI-SB-UL01	0070-LAI	7 ft	07/17/01	mg/kg	5.3	42.9				0.44 U	
DCI-SB-UL03	0020-LAI	2 ft	07/17/01	mg/kg	2.7	17.7				0.285 J	
DCI-SB-UL03	0060-LAI	6 ft	07/17/01	mg/kg	3.0	21.8	-	-		0.315 J	
	S-1-WS-0	0.5-1 ft	08/22/01	mg/kg	3.4	58		-			
S-1-WS	S-1-WS-1	1-4 ft	08/22/01	mg/kg	3.8	52		-			
O-T-44O	S-1-WS-2	4-7 ft	08/22/01	mg/kg	3.1	59		-			
	S-1-WS-3	7-10 ft	08/22/01	mg/kg	2.2	16	-				
	Proposed So	il Cleanup Lev	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	Metals ²	Petro	oleum Hydrocai	bons	Total cP	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	S-2-MS-0	0.5-1 ft	08/22/01	mg/kg	3.8	17	5.8 U	8.1 J	18 J		-
S-2-MS	S-2-MS-1	1-4 ft	08/22/01	mg/kg	5.3	20	6 U	5.9 J	10 U		
	S-2-MS-2	4-7 ft	08/22/01	mg/kg	1.5	12	7.3 U	5.4 J	10 U		
	S-3-EFA-0	0-1 ft	08/22/01	mg/kg	25	63	200	990	620	0.28	
S-3-EFA	S-3-EFA-1	1-4 ft	08/22/01	mg/kg	4.3	32	250	370	50 U	0.056 U	
3-3-LI A	S-3-EFA-2	4-7 ft	08/22/01	mg/kg	3.6	30	7.2 U	19	22	0.06 U	
	S-3-EFA-3	10-13 ft	08/22/01	mg/kg	6.5	38	17 U	19 J	55 J	-	0.08 U
	S-4-EFA-0	0-1 ft	08/22/01	mg/kg	13.6	52	5.5 U	97	340	0.061	-
S-4-EFA	S-4-EFA-1	1-4 ft	08/22/01	mg/kg	5.4	42	6.4 U	6.6 J	24 J	0.061 U	
	S-4-EFA-2	4-7 ft	08/22/01	mg/kg	6.3	72	7.8 U	130 J	220	1.38	-
	S-5-EFA-0	0-1 ft	08/22/01	mg/kg	21	63	6.3 U	68	220	0.26	
	S-5-EFA-1	1-4 ft	08/22/01	mg/kg	6.1	39	6.7 U	9	15	0.061 U	
S-5-EFA	S-5-EFA-2	4-7 ft	08/22/01	mg/kg	6.5	37	8.4 U	10	23	0.12	-
	S-5-EFA-3	7-10 ft	08/22/01	mg/kg	4.5	54	7.8 U	8.2 J	35 J		0.075 U
	S-5-EFA-4	10-13 ft	08/22/01	mg/kg	3.1	43	6.8 U	5 U	10 U	_	0.05 U
	S-6-TPH-0	0-1 ft	08/22/01	mg/kg	8.0	105	5.7 U	46 J	230	0.2	-
S-6-UST	S-6-TPH-1	1-4 ft	08/22/01	mg/kg	3.9	36	6.6 U	65 J	42 J	0.05 U	-
3-0-031	Dup (S-6-TPH-1)	1-4 ft	08/22/01	mg/kg	3.8	41	7.1 U	330 J	100 J	0.0069	-
	S-6-TPH-2	4-7 ft	08/22/01	mg/kg	4.1	22	9.3 U	68 J	91	0.79	
	S-7-TPH-0 ³	0-1 ft	08/22/01	mg/kg	74	66	5.4 U	48	76	0.16	
S-7-UST	S-7-TPH-1 ³	1-4 ft	08/22/01	mg/kg	45	39	68	4,400	500 U	0.17 U	
3-7-031	S-7-TPH-2 ³	4-7 ft	08/22/01	mg/kg	6.2	28	560	7,600 J	500 U	0.10	
	S-7-TPH-3 ³	7-10 ft	08/22/01	mg/kg	1.8	12	7.8 U	360 J	40 U		0.0061 U
	S-8-TPH-0 ³	0-1 ft	08/22/01	mg/kg			130	970 J	4,100	-	-
S-8-UST	S-8-TPH-1 ³	1-4 ft	08/22/01	mg/kg			310	1,100 J	780	-	-
3-0-031	S-8-TPH-2 ³	4-7 ft	08/22/01	mg/kg			50	74 J	76		-
	S-8-TPH-3 ³	7-10 ft	08/22/01	mg/kg			35	13 J	26		
	Proposed Sc	oil Cleanup Lev	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	Metals ²	Petro	oleum Hydroca	rbons	Total cP	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	S-9-CPH-0	0-1 ft	08/22/01	mg/kg	5.3	59	5.6 U	14 J	52 J		-
	Dup (S-9-CPH-0)	0-1 ft	08/22/01	mg/kg	5.9	70	5.6 U	18 J	60 J		-
S-9-CPH	S-9-CPH-1	1-4 ft	08/22/01	mg/kg	2.3	28	6 U	5.5 J	10 U		-
5-9-CPH	S-9-CPH-2	4-7 ft	08/22/01	mg/kg	2.6	31	6.5 U	25 J	23 J		-
	S-9-CPH-3	7-9 ft	08/22/01	mg/kg	7.2	24	47	420 J	330 J		-
	S-9-CPH-3A	9-10 ft	08/22/01	mg/kg	4.7	24	9 U	94 J	82 J		-
	S-10-MR-0	0-1 ft	08/22/01	mg/kg	10.5	173	5.6 U	35 J	200 J		
S-10-MR	S-10-MR-1	1-4 ft	08/22/01	mg/kg	5.1	26	6.1 U	22 J	70 J		
3-10-WK	S-10-MR-2	4-7 ft	08/22/01	mg/kg	4.5	42	6.7 U	8.3 J	25 J		-
	S-10-MR-3	7-10 ft	08/22/01	mg/kg	4.2	28	8	8.0 J	29 J		-
S-11-MR	S-11-MR ³	0-1 ft	08/22/01	mg/kg	39	67 J	470	2,600	1,300		-
S-12-MR	S-12-MR-0 ³	0-1 ft	08/22/01	mg/kg	124 J	56 J	5.9 U	1,900	790		
3-12-WIK	Dup (S-12-MR-0) ³	0.7 ft	08/22/01	mg/kg	240 J	65 J	6 U	1,900	720		-
S-13-MR	S-13-MR ³	0-0.5 ft	08/22/01	mg/kg	270	22 J	7	120	340		
	S-14-TPH-1	1-3.1 ft	10/24/01	mg/kg			5.8 U	72	100		
S-14-TPH	S-14-TPH-4	4-6.4 ft	10/24/01	mg/kg			6.4 U	5 U	10 U		
	S-14-TPH-7	7-10 ft	10/24/01	mg/kg			7.9 U	5 U	10 U		
	S-15-TPH-1	1-3.8 ft	10/24/01	mg/kg			5.5 U	15	32		
S-15-TPH	S-15-TPH-4	4-6.1 ft	10/24/01	mg/kg		-	6.8 U	7	10 U		
	S-15-TPH-7	7-9.9 ft	10/24/01	mg/kg	-		7.6 U	5 U	10 U		-
	S-16-TPH-1 ³	1-3.7 ft	10/24/01	mg/kg	-		120	730	730		-
S-16-TPH	S-16-TPH-4 ³	4-6.3 ft	10/24/01	mg/kg			2,000	40,000	1,300		
	S-16-TPH-7 ³	7-10 ft	10/24/01	mg/kg			7.2 U	21	10 U		-
	S-17-TPH-1	1-3.7 ft	10/24/01	mg/kg			6.6 U	51	130		
S-17-TPH	S-17-TPH-4A	4-4.4 ft	10/24/01	mg/kg			9.3 U	500	100 U		
0-11-1111	S-17-TPH-4B	4.4-6.3 ft	10/24/01	mg/kg			7.3 U	6	10 U		
	S-17-TPH-7	7-9.8 ft	10/24/01	mg/kg			7.6 U	5 U	10 U		
	S-18-TPH-1	1-3.4 ft	10/24/01	mg/kg			5.9 UJ	48 J	150		
S-18-TPH	S-18-TPH-4	4-6.7 ft	10/24/01	mg/kg			6.2 U	9	10 U		
	S-18-TPH-7	7-9.9 ft	10/24/01	mg/kg			6.8 U	5 U	10 U		
	Proposed So	il Cleanup Lev	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	/letals ²	Petro	leum Hydroca	rbons	Total cP/	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	S-19-TPH-1	1-3.6 ft	10/24/01	mg/kg			6.1 U	350	100 U		-
S-19-TPH	S-19-TPH-4	4-6.4 ft	10/24/01	mg/kg			69	1700 J	36 J		-
	S-19-TPH-7	7-9.9 ft	10/24/01	mg/kg			7.5 ป	190	10 U		-
	S-20-TPH-1 ³	1-3.9 ft	10/24/01	mg/kg			5.8 U	9	15		-
S-20-TPH	S-20-TPH-4 ³	4-6.5 ft	10/24/01	mg/kg			210	2,600	140 J		-
	S-20-TPH-7 ³	7-10 ft	10/24/01	mg/kg			9 U	12	18		
	S-21-TPH-1 ³	1-2.2 ft	10/24/01	mg/kg		/	5.5 U	12	10 U		-
S-21-TPH	S-21-TPH-4 ³	4-4.1 ft	10/24/01	mg/kg			5.7 U	140	35		-
	S-21-TPH-7 ³	7-9.4 ft	10/24/01	mg/kg			8 U	8	10U		-
	S-22-TPH-1A ³	1-2.5 ft	10/24/01	mg/kg			6.9 U	1,600	960		-
S-22-TPH	S-22-TPH-1B ³	2.5-4 ft	10/24/01	mg/kg			700	6,700	110 J		-
5-22-1PH	S-22-TPH-4 ³	4-5 ft	10/24/01	mg/kg	-		360	380	39		-
	S-22-TPH-7 ³	7-9.5 ft	10/24/01	mg/kg			34	10	11		-
	S-23-TPH-1 ³	1-3.4 ft	10/24/01	mg/kg			5.9 U	5 U	10 U		-
S-23-TPH	S-23-TPH-4 ³	4-6.7 ft	10/24/01	mg/kg			6.9 U	3,800	210 J		-
	S-23-TPH-7 ³	7-9.6 ft	10/24/01	mg/kg			8.9 U	9	10 U		-
SS-1	SS-1-1	1-1.5 ft	06/16/08	mg/kg							
SS-2	SS-2-1	1-1.5 ft	06/16/08	mg/kg							
SS-3	SS-3-1	1-1.5 ft	06/16/08	mg/kg	-	-					
SS-4	SS-4-0.5	0.5 -1 ft	06/16/08	mg/kg	1		-	-			
MW-5	MW-5-5.0	5-6.5 ft	05/27/08	mg/kg	-		3 U			0.19	
10100-5	MW-5-10.0	10-11.5 ft	05/27/08	mg/kg	1		3 U				0.18
SB-1	SB-1-2.0	2-3 ft	06/16/08	mg/kg	8.7						-
30-1	SB-1-4.0	4-5 ft	06/16/08	mg/kg	5 U				-		ı
SB-2	SB-2-2.0	2-3 ft	06/16/08	mg/kg	5 U						
JD-2	SB-2-4.0	4-5 ft	06/16/08	mg/kg	5 U						
SB-4	SB-4-3.0	3-4 ft	06/16/08	mg/kg	-					0.02 U	
3b-4	SB-4-9.0	9-10 ft	06/16/08	mg/kg	-						-
SB-5	SB-5-3.0	3-4 ft	06/16/08	mg/kg	-			-	-	0.07	-
ა ნ -ა	SB-5-9.0	9-10 ft	06/16/08	mg/kg	-						
	Proposed Soil Cleanup Level ⁵					48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	/letals ²	Petro	oleum Hydroca	rbons	Total cP	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
SB-7	SB-7-3.0	3-4 ft	06/16/08	mg/kg	-	-				0.038	
3B-1	SB-7-9.0	9-10 ft	06/16/08	mg/kg	-	-					
SB-8	SB-8-0.5	0.5-1.5 ft	06/17/08	mg/kg	-		/				
30-8	SB-8-4.0	4-5 ft	06/17/08	mg/kg	1			-			-
SB-9	SB-9-0.5	0.5-1.5 ft	06/16/08	mg/kg			-	_			
30-9	SB-9-4.0	4-5 ft	06/16/08	mg/kg	1		-	-			1
SB-10	SB-10-0.5	0.5-1.5 ft	06/17/08	mg/kg	1		-	-	-		-
36-10	SB-10-4.0	4-5 ft	06/17/08	mg/kg	-		1		-		1
SB-11	SB-11-0.5	0.5-1.5 ft	06/17/08	mg/kg							
36-11	SB-11-4.0	4-5 ft	06/17/08	mg/kg	-		-				-
SB-12	SB-12-0.5	0.5-1.5 ft	06/16/08	mg/kg	910						
3D-12	SB-12-4.0	4-5 ft	06/16/08	mg/kg	48	-	-				-
SB-13	SB-13-0.5	0.5-1.5 ft	06/16/08	mg/kg	5.2	1	-				
30-13	SB-13-4.0	4-5 ft	06/16/08	mg/kg	5 U	-	1				-
SB-14	SB-14-0.5	0.5-1.5 ft	06/16/08	mg/kg	73	_	-	-	-		
3b-14	SB-14-4.0	4-5 ft	06/16/08	mg/kg	5 U		1	-	-		1
SB-15	SB-15-0.5	0.5-1.5 ft	06/16/08	mg/kg	180						
36-13	SB-15-4.0	4-5 ft	06/16/08	mg/kg	5 U		1				-
TP-3	TP-3-6	6-6.5 ft	09/05/08	mg/kg	5 U		-				
TP-4	TP-4-6	6-6.5 ft	09/08/08	mg/kg	5 U						-
TP-5	TP-5-2	2-2.5 ft	09/08/08	mg/kg	15						
11-5	TP-5-4	4-4.5 ft	09/08/08	mg/kg	9.6						
TP-10	TP-10-4	4-4.5 ft	09/08/08	mg/kg	5 U		-				-
17-10	TP-10-6	6-6.5 ft	09/08/08	mg/kg	1		-	-	-		-
TP-11	TP-11-6	6-6.5 ft	09/08/08	mg/kg	5 U				-		
TP-12	TP-12-3	3-3.5 ft	09/08/08	mg/kg	5 U				-		
TP-13	TP-13-2	2-2.5 ft	09/08/08	mg/kg	24						
11-13	TP-13-4	4-4.5 ft	09/08/08	mg/kg	34						
TP-14	TP-14-0-2	0-2 ft	09/18/08	mg/kg	5 U				-		
TP-15	TP-15-2-4	2-4 ft	09/18/08	mg/kg	5 U				-		
	Proposed Se	oil Cleanup Lev	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	Metals ²	Petro	oleum Hydroca	rbons	Total cP	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
TD 16	TP-16-0-2	0-2 ft	09/18/08	mg/kg	5 U			_			-
TP-16	TP-16-4-6	4-6 ft	09/18/08	mg/kg	5 U		-	-			-
GEI-01	GEI-01_3-4	3-4 ft	09/29/14	mg/kg	6 U		-/	-			
	GEI-02_1-2	1-2 ft	09/29/14	mg/kg	5.2 U	34					
GEI-02	GEI-02_4-5	4-5 ft	09/29/14	mg/kg	5.8 U		_				
	GEI-02_7-8	7-8 ft	09/29/14	mg/kg	5.3 U	8		-			
GEI-03	GEI-03_2.5-3.5	2.5-3.5 ft	09/29/14	mg/kg	7.8	25					
GEI-03	GEI-03_7-8	7-8 ft	09/29/14	mg/kg	5.7 U	21 J					
	GEI-04_1-2	1-2 ft	09/29/14	mg/kg	5.2 U	31					
GEI-04	GEI-04_3-4	3-4 ft	09/29/14	mg/kg	13						
	GEI-04_6-7	6-7 ft	09/29/14	mg/kg	33	43					
GEI-05	GEI-05_7-8	7-8 ft	09/29/14	mg/kg	6.0 U	88	-				-
	GEI-06_1.5-2.5	1.5-2.5 ft	09/29/14	mg/kg	23	58	-				
GEI-06	GEI-06_4-5	4-5 ft	09/29/14	mg/kg	5.2 U	-					
	GEI-06_7-8	7-8 ft	09/29/14	mg/kg	5.6 U	29					
GEI-07	GEI-07_1.5-2.5	1.5-2.5 ft	09/29/14	mg/kg	27	52					
GLI-01	GEI-07_7-8	7-8 ft	09/29/14	mg/kg	4.4	38					
	GEI-08_1.5-2.5	1.5-2.5 ft	09/29/14	mg/kg	5.2 U	50					
GEI-08	GEI-08_4-5	4-5 ft	09/29/14	mg/kg	5.3 U	-					-
	GEI-08_7-8	7-8 ft	09/29/14	mg/kg	6.3 U	27	-				1
	GEI-09_0.5-1.5	0.5-1.5 ft	09/29/14	mg/kg	62	57	1		-		1
GEI-09	GEI-09_3-4	3-4 ft	09/29/14	mg/kg	5.4 U						-
	GEI-09_6-7	6-7 ft	09/29/14	mg/kg	4.6	40					
GEI-10	GEI-10_2-3	2-3 ft	09/29/14	mg/kg	33	150					-
GLI-10	GEI-10_7-8	7-8 ft	09/29/14	mg/kg	6.1 U	8.3					-
	GEI-11_2-3	2-3 ft	09/29/14	mg/kg	5.9	26					
GEI-11	GEI-11_7-8	7-8 ft	09/29/14	mg/kg	5.5 U	38					
	GEI-11_9-10	9-10 ft	09/29/14	mg/kg	5.6 U	34					-
	GEI-12_2-3	2-3 ft	09/29/14	mg/kg	5.2 U	-	-		-		-
GEI-12	GEI-12_4-5	4-5 ft	09/29/14	mg/kg	19		-		-		-
	GEI-12_7-8	7-8 ft	09/29/14	mg/kg	5.8 U						
	Proposed So	oil Cleanup Leve	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	/letals ²	Petro	oleum Hydroca	rbons	Total cP/	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	GEI-13_2-3	2-3 ft	09/30/14	mg/kg	85	110					-
GEI-13	GEI-13_5-6	5-6 ft	09/30/14	mg/kg	5.2 U						-
	GEI-13_7-8	7-8 ft	09/30/14	mg/kg	5.3 U	12	/				-
	GEI-14_2-3	2-3 ft	09/30/14	mg/kg	91	43					
GEI-14	GEI-14_3.5-4.5	3.5-4.5 ft	09/30/14	mg/kg	6.5			1			
GLI-14	GEI-14_7-8	7-8 ft	09/30/14	mg/kg	7.4	13		-			
	GEI-14_9-10	9-10 ft	09/30/14	mg/kg	6.4 U	-		-	-		
	GEI-15_2-3	2-3 ft	09/30/14	mg/kg	5.2 U		-4	-	-		
GEI-15	GEI-15_5.5-6.5	5.5-6.5 ft	09/30/14	mg/kg	6.0 U	41	-			0.0061 U	
	GEI-15_10-11	10-11 ft	09/30/14	mg/kg	ı	39		-			
	GEI-16_2-3	2-3 ft	09/30/14	mg/kg	6.6	36					
GEI-16	GEI-16_6-7	6-7 ft	09/30/14	mg/kg	6.6 U	6.2					
	GEI-16_8-9	8-9 ft	09/30/14	mg/kg	5.8 U	15	-	-			
	GEI-17_1-2	1-2 ft	09/30/14	mg/kg	-	41					
GEI-17	GEI-17_4-5	4-5 ft	09/30/14	mg/kg	8	_			-		
GLI-I1	GEI-17_7-8	7-8 ft	09/30/14	mg/kg	31	39					
	GEI-17_9-10	9-10 ft	09/30/14	mg/kg	8.1	58					-
	GEI-18_1-2	1-2 ft	09/30/14	mg/kg	5.2	36	-	1	-		-
GEI-18	GEI-18_4-5	4-5 ft	09/30/14	mg/kg	18	37		-	-		-
GLI-10	GEI-18_8-9	8-9 ft	09/30/14	mg/kg	11	200		-	-		-
	GEI-18_9-10	9-10 ft	09/30/14	mg/kg	6.9	49					
	GEI-19_2-3	2-3 ft	09/30/14	mg/kg	-	43					_
GEI-19	GEI-19_4-5	4-5 ft	09/30/14	mg/kg	5.2 U						
GLI 10	GEI-19_7-8	7-8 ft	09/30/14	mg/kg	2.6						
	GEI-19_9-10	9-10 ft	09/30/14	mg/kg		29					-
	GEI-20_2-3	2-3 ft	09/30/14	mg/kg	-	37		-			-
GEI-20	GEI-20_4-5	4-5 ft	09/30/14	mg/kg	6.5						
GLI-20	GEI-20_6-7	6-7 ft	09/30/14	mg/kg						0.06	
	GEI-20_8-9	8-9 ft	09/30/14	mg/kg	5.7						<u></u>
	Proposed So	oil Cleanup Leve	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	/letals ²	Petro	oleum Hydrocai	rbons	Total cP	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	GEI-21_1-2	1-2 ft	09/30/14	mg/kg		33		- -			
GEI-21	GEI-21_5-6	5-6 ft	09/30/14	mg/kg	5.9 U	28	-			0.015	-
	GEI-21_7.5-8.5	7.5-8.5 ft	09/30/14	mg/kg	3.1	1	-				0.024
	GEI-22_2-3	2-3 ft	10/01/14	mg/kg	5.3 U	35		-			
GEI-22	GEI-22_5-6	5-6 ft	10/01/14	mg/kg	5.4 U	30		_		0.022	
	GEI-22_7.5-8.5	7.5-8.5 ft	10/01/14	mg/kg	92	-					-
	GEI-23_1-2	1-2 ft	09/30/14	mg/kg	5.2 U	27					
GEI-23	GEI-23_5-6	5-6 ft	09/30/14	mg/kg	12		-4				
	GEI-23_7.5-8.5	7.5-8.5 ft	09/30/14	mg/kg	6.1 U		/				0.0061 U
	GEI-24_2-3	2-3 ft	09/30/14	mg/kg	5.2 U	40					
GEI-24	GEI-24_4-5	4-5 ft	09/30/14	mg/kg	5.2 U						
GEI-24	GEI-24_6-7	6-7 ft	09/30/14	mg/kg	13						
	GEI-24_9-10	9-10 ft	09/30/14	mg/kg	6.3 U	-	-				
	GEI-25_1-2	1-2 ft	09/30/14	mg/kg	7.5	36					
GEI-25	GEI-25_4-5	4-5 ft	09/30/14	mg/kg	6.0 U	80					
GEI-25	GEI-25_7-8	7-8 ft	09/30/14	mg/kg	6.1 U	130					
	GEI-25_9-10	9-10 ft	09/30/14	mg/kg		130					
GEI-26	GEI-26_2 -3	2-3 ft	09/30/14	mg/kg	5.4	40					
GEI-20	GEI-26_6-7	6-7 ft	09/30/14	mg/kg	5.8 U	33					
	GEI-27_1-2	1-2 ft	10/01/14	mg/kg		28					
GEI-27	GEI-27_5-6	5-6 ft	10/01/14	mg/kg	22	20	-				
	GEI-27_9-10	9-10 ft	10/01/14	mg/kg	6.6 U	-					-
	GEI-28_2-3	2-3 ft	10/01/14	mg/kg	5.9 U	34					
GEI-28	GEI-28_5-6	5-6 ft	10/01/14	mg/kg	5.8 U	50				0.0058 U	
	GEI-28_10-11	10-11 ft	10/01/14	mg/kg	6.7	37					
_	GEI-29_2-3	2-3 ft	09/30/14	mg/kg	5.5 U	42					
GEI-29	GEI-29_5-6	5-6 ft	09/30/14	mg/kg	6.4	-	-				-
GLI-25	GEI-29_8-9	8-9 ft	09/30/14	mg/kg	3.6 U	1	-		-		1
	GEI-29_9-10	9-10 ft	09/30/14	mg/kg	6.0 U		-				
	Proposed Sc	oil Cleanup Leve	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total N	/letals ²	Petro	oleum Hydrocai	bons	Total cP	AH TEQ ^{3,4}
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	GEI-30_3-4	3-4 ft	09/30/14	mg/kg	5.4 U	7.3					
GEI-30	GEI-30_7-8	7-8 ft	09/30/14	mg/kg	5.7 U	8.9					-
	GEI-30_9-10	9-10 ft	09/30/14	mg/kg	6.0 U	11	/				
	GEI-31_1-2	1-2 ft	10/01/14	mg/kg	7.6	110					
GEI-31	GEI-31_4-5	4-5 ft	10/01/14	mg/kg		40		_			
GEI-SI	GEI-31_6-7	6-7 ft	10/01/14	mg/kg		86					
	GEI-31_9-10	9-10 ft	10/01/14	mg/kg	_	48					-
GEI-32	GEI-32_1-2	1-2 ft	10/01/14	mg/kg	5.3 U	64					
GEI-32	GEI-32_6-7	6-7 ft	10/01/14	mg/kg		14					-
GEI-33	GEI-33_1-2	1-2 ft	10/01/14	mg/kg	5.2 U	-					
	GEI-34_2.5-3.5	2.5-3.5 ft	10/01/14	mg/kg	5.3 U						
GEI-34	GEI-34_6-7	6-7 ft	10/01/14	mg/kg		38					-
	GEI-34_9-10	9-10 ft	10/01/14	mg/kg	1-	38					-
	GEI-35_3-4	3-4 ft	10/01/14	mg/kg	6.2 U	35				-	
GEI-35	GEI-35_4-5	4-5 ft	10/01/14	mg/kg	-	22					-
GEI-35	GEI-35_8-9	8-9 ft	10/01/14	mg/kg		40					
	GEI-35_9-10	9-10 ft	10/01/14	mg/kg		28					-
GEI-36	GEI-36_1-2	1-2 ft	10/01/14	mg/kg	5.7						
GEI-30	GEI-36_5-6	5-6 ft	10/01/14	mg/kg	6.9	50					
GEI-37	GEI-37_1-2	1-2 ft	10/01/14	mg/kg	5.2 U	-					
GEI-57	GEI-37_6-7	6-7 ft	10/01/14	mg/kg	5.7 U	8.1			-		-
GEI-38	GEI-38_1-2	1-2 ft	10/01/14	mg/kg	5.6 U				-		-
GLI-38	GEI-38_6-7	6-7 ft	10/01/14	mg/kg	5.8 U		-		-		-
	GEI-39_1.5-2.5	1.5-2.5 ft	10/01/14	mg/kg	6.0 U	38					-
GEI-39	GEI-39_4-5	4-5 ft	10/01/14	mg/kg		45					-
	GEI-39_6-7	6-7 ft	10/01/14	mg/kg	6.5 U	40			-		-
GEI-40	GEI-40_2-3	2-3 ft	10/01/14	mg/kg	6.2				-		-
	GEI-41_1-2	1-2 ft	10/01/14	mg/kg	6.2 U	46			-	-	-
GEI-41	GEI-41_4-5	4-5 ft	10/01/14	mg/kg	5.8 U	-			-		-
ULI-41	GEI-41_6-7	6-7 ft	10/01/14	mg/kg	-	43			-		-
	GEI-41_8-9	8-9 ft	10/01/14	mg/kg		30					
	Proposed So	oil Cleanup Leve	el ⁵		20	48	100	2,000	2,000	0.31	0.016



Sample	Sample	Sample Depth	Sample		Total Metals ² Petrole		oleum Hydrocai	rbons	Total cPAH TEQ ^{3,4}		
Location ¹	Identification	(bgs)	Date	Units	Arsenic	Nickel	Range	Range	Range	Zone	Zone
	GEI-42_1-2	1-2 ft	10/01/14	mg/kg	5.5	34		-			
GEI-42	GEI-42_4-5	4-5 ft	10/01/14	mg/kg	-	24					
	GEI-42_6-7	6-7 ft	10/01/14	mg/kg	6.0 U	31		-			
GEI-43	GEI-43_1-2	1-2 ft	10/01/14	mg/kg	6.6	52		-			
GLI-43	GEI-43_6-7	6-7 ft	10/01/14	mg/kg	-	37		1			
	GEI-44_1.5-2	1.5-2 ft	07/23/18	mg/kg	9.5	-		ı		0.049	
GEI-44	GEI-44_7.5-10	7.5-10 ft	07/23/18	mg/kg	5.6 U	-		-	-		0.006 U
	GEI-44_16-17.5	16-17.5 ft	07/23/18	mg/kg	5.4 U		-4	-	-		0.006 U
	GEI-45_1-3	1-3 ft	07/23/18	mg/kg	5.3 U	-	-			0.005 U	
GEI-45	GEI-45_9-10	9-10 ft	07/23/18	mg/kg	5.5 U	-	-				0.006 U
	GEI-45_17-20	17-20 ft	07/23/18	mg/kg	5.6 U	-		1			0.006 U
GEI-46	GEI-46_7-8.5	7-8.5 ft	07/23/18	mg/kg	5.7 U	1	-	1	-	0.006 U	
GLI-40	GEI-46_13.5-15	13.5-15 ft	07/23/18	mg/kg	6.0 U	-					0.006 U
	Proposed Soil Cleanup Level ⁵					48	100	2,000	2,000	0.31	0.016

bgs = below the ground surface

mg/kg = milligram per kilogram

J = Estimated value

U = qualifier indicating analyte not detected at level above listed practical quantitation limit

-- = not analyzed

Bold indicates analyte was detected.

Blue shading indicates that the reporting limit exceeds the proposed groundwater cleanup level.

Yellow shading indicates analyte was detected at a concentration above proposed groundwater cleanup level.



¹ Soil sample locations are shown on Figure 14. Soil sample results are summarized on Figures s 23 through 26.

² Metals analyzed using United States Environmental Protection Agency (EPA) Method 6010/6020.

³ Carcinogenic Polycyclic Aromatic Hydrocarbons (cPAHs) analyzed by EPA method 8270D/SIM.

⁴ Total cPAHs calculated using toxic equivalent (TEQ) methodology relative to benzo(a)pyrene. Non-detect compounds were assigned a value of one half the reporting limit for the calculation.

⁵ Proposed soil cleanup level is referenced from the Remedial Investigation/Feasibility Study Work Plan (GeoEngineers 2008).

Applicable or Relevant and Appropriate Requirements

Authorizing Statute and Implementing Regulation	Citation	Procedural/Substantive Requirements	Description and Applicability
Federal ARARs			
Archaeological Resources Protection Act	16 USC § 470aa et seq. 43 CFR Part 7	Prohibits the unauthorized disturbance of archaeological resources on public or Indian lands. Archaeological resources are "any material remains of past human life and activities which are of archaeological interest," including pottery, baskets, tools, and human skeletal remains. The unauthorized removal of archaeological resources from public or Indian lands is prohibited without a permit, and any archaeological investigations at a site must be conducted by a professional archeologist.	Potentially applicable to a site where response actions involve disturbance/ alteration of
American Indian Religious Freedom Act	42 USC § 1996 et seq	The American Indian Religious Freedom Act and implementing regulations are intended to protect Native American religious, ceremonial, and burial sites, and the free practice of religions by Native American groups. The requirements of this Act must be followed if sacred sites graves are discovered in the course of ground-disturbing activities.	the ground and/or site terrain. Appropriate measures will be taken to evaluate the presence of cultural resources. If a potential for an existence of cultural resources exists then appropriate measures will be taken during excavation activities and appropriate tribal members will be contacted in the event that an artifact is encountered.
Native American Graves Protection and Repatriation Act	25 USC § 3001 et seq 43 CFR Part 10 25 USC 3001 et seq. 43 CFR 10	Intended to protect Native American graves from desecration through the removal and trafficking of human remains and "cultural items" including funerary and sacred objects. The requirements of this Act must be followed when graves are discovered or ground-disturbing activities encounter Native American burial sites.	
Clean Air Act (CAA), National Ambient Air Quality Standards	42 USC 7401 et seq. 40 CFR 50	Provides air quality standards for six criteria pollutants, including particulate matter, to protect public health and welfare.	Applicable.
Endangered Species Act (ESA)	16 U.S.C. § 1531 – 1544 50 CFR Parts 17, 402	Provides for the protection of species of fish, wildlife, and plants that are listed as threatened or endangered with extinction. It also protects designated critical habitat for listed species. The Act outlines procedures for federal agencies to follow when taking actions that may jeopardize listed species, including consultation with resource agencies.	Potentially applicable to the site for listed and proposed to be listed threatened or endangered species and their habitat areas which will, or could, be impacted by cleanup action.
Federal Coastal Zone Management Act (CZMA)	16 USC 1451-1464 15 CFR 923-930	The CZMA requires that federal agency action that is reasonably likely to affect use of shorelines be consistent with the approved coastal zone management plan to the maximum extent practicable, subject to limitations set forth in the CZMA and requires that construction activities near the shoreline must be consistent with the State's Coastal Zone Management Program.	Applicable if construction is completed within 200 feet of the shoreline. The requirements will be met by preparing a CMZA form for Washington State Department of Ecology's review. Ecology reviews the proposed project for consistency with state environmental requirements, including shoreline permitting requirements.
Fish and Wildlife Conservation Act (FWCA)	16 U.S.C. § 661 et seq 50 CFA 83	Requires that adequate provision must be made for the conservation, maintenance, and management of wildlife resources and habitat and requires consultation with the U.S. Fish and Wildlife service and appropriate state agencies.	Applicable to the site if listed threatened or endangered species habitat areas will, or could, be impacted by cleanup action.



Authorizing Statute and Implementing Regulation	Citation	Procedural/Substantive Requirements	Description and Applicability
Hazardous Materials Transportation Act	49 USC 1801-1813 49 CFR 107, 171-177	Regulates the transportation of hazardous waste.	Applicable to the site if offsite disposal is included in cleanup action.
Migratory Bird Treaty Act (MBTA)	16 USC § 703 et seq	Makes it unlawful to "hunt, take, capture, kill" or take various other actions adversely affecting a broad range of migratory birds, including tundra swans, hawks, falcons, songbirds, without prior approval by the U.S. Fish and Wildlife Service.	Applicable for protecting migratory bird species if identified. The selected response action must be carried out in a manner that avoids the taking of protected migratory bird species, including individual birds or their nests or eggs.
Occupational Safety and Health Act (OSHA)	29 CFR 1904 29 CFR 1910 29 CFR 1926	Specifies minimum requirements to maintain worker health and safety during hazardous waste operations, including training and construction safety requirements.	Applicable to construction phases of a cleanup. Construction activities will be conducted in accordance with the requirements of OSHA.
Resource Conservation and Recovery Act (RCRA), Identification and Management of Hazardous Wastes	40 CFR 261 et seq.	Specifies how to determine whether a solid waste is considered hazardous (whether listed or based on characteristic) and how to manage hazardous wastes.	Relevant and appropriate to the site. Washington State is authorized for RCRA.
State ARARs			
Model Toxics Control Act (MTCA) Cleanup Regulation	RCW 70.105D Chapter 173-340 WAC	MTCA is the primary regulation governing cleanup actions.	Cleanup actions conducted by Ecology under MTCA are exempt from the procedural requirements of most state and local laws/permits; however, must meet substantive requirements of the laws/permits.
State Environmental Policy Act (SEPA)	Chapter 43.21C RCW Chapter 173-802 WAC Chapter 197-11 WAC	Prior to taking any action on a proposal, agencies must follow specific procedures to ensure that appropriate consideration has been given to the environment. The severity of potential environmental impacts associated with a project determines whether an Environmental Impact Statement is required.	Applicable. A SEPA checklist is required prior remedial construction activities.
Shoreline Management Act	Chapter 90.58 RCW Chapter 173-27-060 WAC	The Shoreline Management Act and its implementing regulations establish requirements for substantial developments occurring within waters of the state or within 200 feet of the shoreline. Local shoreline management programs are adopted under state regulations, creating an enforceable state law.	Applicable to upland cleanup action alternatives that include activities within 200 feet of the shoreline. Cleanup actions under MTCA are exempt from shoreline management act permitting; however, will need to meet substantive requirements.
MTCA, Site Cleanup and Monitoring	WAC 173-340-400 through 173-340-440	Provides requirements for implementation of the cleanup action, compliance monitoring, periodic review, interim action and institutional controls.	Applicable.
Washington Clean Air Act	Chapter 70.94 RCW Chapter 43.21A RCW Chapter 173-400 WAC	Requires all sources of air contaminants to meet emission standards for visible, particulate, fugitive, odors, and hazardous air emissions. Requires use of reasonably available control technology.	Substantive requirements are applicable for any response actions in the project area that may create fugitive dust or other regulated air emissions.
Puget Sound Clean Air Agency (PSCAA)	Regulation 1, Section 9.15.	Provides regulation for the visible emissions of fugitive dust and reasonable precautions that should be employed to minimize these emissions.	Substantive requirements are applicable for any response actions in the project area that may create fugitive dust or other regulated air emissions.



Authorizing Statute and Implementing Regulation	Citation	Procedural/Substantive Requirements	Description and Applicability
Hazardous Waste Management Act, Dangerous Waste Regulations	Chapter 70.105 RCW Chapter 173-303 WAC	Governs handling and disposition of dangerous waste, including identification, accumulation, storage, transport, treatment, and disposal.	Substantive requirements are applicable handling, storage, and disposal of hazardous material. These requirements will apply if dangerous wastes are generated during the cleanup action.
Solid Waste Handling Standards	Chapter 70.95 RCW Chapter 173-350 WAC.	The solid waste requirements are applicable to remedial alternatives that consist of off-site disposal of solid non-hazardous wastes and contaminated media.	For off site disposal activities, waste materials will be sent to facilities licensed and permitted to accept the specific waste material and documentation will be obtained of such disposition.
Washington Industrial Safety and Health Act (WISHA)	Chapter 49.17 RCW Chapter 296-62 WAC Chapter 296-843 WAC	Specifies minimum requirements to maintain worker health and safety during hazardous waste operations, including training and construction safety requirements.	Applicable to construction phases of a cleanup. Construction activities will be conducted in accordance with the requirements of WISHA.
City ARARs			
City of Anacortes Noise Ordinance	Chapter 17.54.010 Ordinance 2316 (part), 1994	Establishes noise levels and standards.	Applicable.
City of Anacortes Publicly Owned Treatment Water (POTW) Discharge Authorization	Chapter 13.40.060	Establishes the requirements and limitations for discharges to the POTW.	Potentially applicable if collected water from construction activities will be treated and permitted for disposal under a discharge authorization by the City.
City of Anacortes Stormwater Management Program	Chapter 13.36 Chapter 17.54.050	Provides the necessary measures to control the quantity and quality of stormwater produced by new development and redevelopment such that they comply with water quality standards and contribute to the protection of beneficial uses of the receiving waters.	Potentially applicable if dredged material is processed upland of the Site prior to permitted landfill disposal.

ARAR = Applicable or Relevant and Appropriate Requirement

CFR = Code of Federal Regulations

RCW = Revised Code of Washington

WAC = Washington Administrative Code

USC = United States Code



Soil Remedial Technologies Screening

Remedial Technology Identification			Description of	Implementability of	Effectiveness of	Relative Remedial T		Summary of	Technology Retained
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	(Yes/No)
No Action	No Action	None	No institutional controls or treatment.	Not effective for protecting human health and environment.	Implementable but not acceptable to the general public or government agencies.	None	None	Used as a baseline for comparison.	No
	Governmental/ Property Controls	Environmental Covenant	Legal restrictions associated with future land use and activities (e.g., development, construction, etc.); may also be used to specify long-term maintenance requirements of remediation systems.	Technically implementable. Specific legal requirements and authority would need to be met.	Not effective for remediating contaminants. Can be effective at reducing risks and maintaining integrity of a remedy.	Low	Low	Applicable and/or required in combination with other technologies.	Yes
		Land Use Restrictions, Soil Management Plans/ Requirements	Restrictions on activities such as excavation to prevent physical damage to in situ remedies (e.g., caps) and/or exposure to hazardous substances that remain in-place. Implement soil management plans/requirements so that contaminated soils are managed properly in an event that it is necessary to disturb/excavate (e.g., utility work, etc.).	Technically implementable but administratively more difficult. Requires an implementing agency.	Not effective for remediating contaminants. Enforcement would be required for restrictions to be effective.	Low	Low	Applicable and/or required in combination with other technologies.	Yes
	Access Restrictions and Information Devices	Fencing and Warning Signage	Placement of fencing and warning signs to prevent access and inform the public regarding health risks. Fencing currently exists at the Site to restrict public access. Signage would be used to inform site workers of potential health risks.	Technically implementable. Implementability and applicability depends on current and future site uses.	Not effective for remediating contaminants. Effective in minimizing human exposure to contaminated media by preventing access.	Low	Low	Applicable and/or required in combination with other technologies.	Yes
Containment/ Lo	Low-Permeability Cap with Drainage Controls	Asphalt and/or Concrete Cap	Maintain existing asphalt or concrete cap over contaminated soil as well as existing stormwater collection and treatment system. Installation of additional asphalt and/or concrete cap in unpaved areas of the Site. Primary function of the cap is to prevent/minimize stormwater infiltration, contaminant migration and exposure to hazardous substances that remain in-place.	Technically implementable. A majority of the Site is currently paved and stormwater collection systems are in place to manage and treat water prior to discharge.	Effective for preventing exposure to hazardous substances that remain in-place, erosion of source material, and reducing stormwater infiltration and contaminant migration.	Low to Moderate	Low	Applicable and/or required in combination with other technologies.	Yes
Capping		A minimum of 1-Foot of Soil Cover with Underlying Low- Permeability Barrier	Install soil cover (a minimum of 1-foot thick) with underlying barrier (plastic or similar) over contaminated soil in unpaved areas. Surface/storm water collection and discharge would be designed to reduce infiltration of stormwater at the site. Primary function of the cover is to prevent/minimize contaminant migration and exposure to hazardous substances that remain in-place.	Technically implementable. Implementability and applicability depends on current and future site uses. Additional considerations for stormwater collection, treatment, and discharge will be needed. May require off-site disposal of material removed to facilitate cap placement.	Effective for preventing exposure to hazardous substances that remain in-place, and reducing stormwater infiltration and contaminant migration. However, the use of heavy equipment on site may compromise the integrity of underlying barrier causing this technology to be less effective.	Low to Moderate	Low	May not be effective due to current site use.	No



	Remedial Technology Identification		Description of	Implementability of	Effectiveness of		e Cost of Technology	Summary of	Technology Retained
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	(Yes/No)
Containment/ Capping	Permeable Cap	Permeable Soil Cover	Install and/or maintain existing 6-feet thick (conditional point of compliance) soil cover over contaminated soil. Can be vegetated at the surface based on current/future site use. Primary function of the cover is to prevent/minimize exposure to hazardous substances that remain in-place. Not effective at reducing stormwater infiltration.	Technically implementable. Implementability and applicability depends on current and future site uses. Requires disposal of material removed to facilitate placement of cover.	Effective for preventing exposure to hazardous substances that remain in-place and erosion of source material. Not effective at reducing stormwater infiltration and contaminant migration.	Moderate to High	Low	Ineffective at reducing stormwater infiltration/contaminant migration and high cost relative to other remedial technologies.	No
	Monitored Natural Attenuation (MNA)	Natural Attenuation	Natural biotransformation processes such as volatilization, biodegradation, adsorption, and chemical reactions with soil materials are used to reduce contaminant concentrations.	Technically implementable. Monitoring may be required to ensure adequate reduction rate. Require institutional controls during treatment period.	Generally not effective for reducing risk to human health and ongoing threats to groundwater in a reasonable time frame. Effectiveness is highest in combination with other technologies as a final step to achieve cleanup levels when risks to human health and the environment are low.	Low	Moderate	Remediation time frame would not be reasonable.	No
	Physical Treatment	Soil Flushing	The extraction of contaminants from soil with aqueous solution accomplished by passing fluid through in-place soils using an injection or infiltration process. Extraction fluids must be recovered from underlying groundwater.	Technically implementable, but would require significant safety components to prevent exacerbating groundwater contamination. Regulatory concerns over potential to wash contaminants beyond fluid capture zones and introduction of surfactants in to the subsurface would make permitting difficult.	Effective for more soluble chemicals. Presence of fine-grained soils and debris limits effectiveness.	High	Moderate	High cost and uncertainty relative to other remedial technologies.	No
In Situ Treatment	Chemical	In Situ Stabilization (ISS)	ISS is accomplished by injecting solutions of chemical reagents with contaminated media. The reagents reduce the mobility/leachability of contaminants and stabilizes it.	Technically implementable. Buried debris or subsurface obstruction such as foundation/utilities may interfere and would require prior removal. Solidification and stabilization processes can result in an increase in volume. Treatability testing is required.	Effective for reducing mobility of metals. Most common in situ source control technology for metals used at CERCLA sites.	Moderate to High	Low	Applicable for site conditions and contaminants (metals) but requires treatability testing.	Yes
	In Situ Chemical chemical oxidation (ISCO) media. The re-		ISCO is accomplished by injecting solutions of chemical oxidation reagents with contaminated media. The reagents chemically oxidizes and destroys contaminants.	Technically implementable. Buried debris or subsurface obstruction such as foundation/utilities may interfere and would require prior removal. Treatability testing is required.	Effective for treating PAHs and other organics. Implemented widely on CERCLA sites.	Moderate to High	Low	Applicable for site conditions and contaminants (PAHs) but requires treatability testing.	Yes
	Biological Treatment	Phytoextraction/ Phytodegradation	Plants, called "Hyperaccumulators" (e.g. Chinese brake fern) have the capacity to extract and store large amounts of contaminants (metals, hydrocarbons etc.) from soil and use them as nutrients during metabolism. Phytoremediation typically involves interaction of plant roots and microorganisms associated with them to remediate soil.	Technically implementable. However, industrial site use would limit implementation. Disposal of accumulated waste materials or plant materials may be necessary. Pilot testing that would be required will significantly delay implementation of full-scale remediation.	Use for Chinese brake fern for remediating soil contaminated with metals were evaluated as part of Tacoma Smelter Plume and the study concluded that phytoremediation is not a good cleanup option due to following reasons: Phytoremediating plants (Chinese brake fern) did not grow well in climatic conditions of Puget Sound, arsenic level in the fronds pose a health risk, fronds hyperaccumulated arsenic and became dangerous waste and fern did not take up other metals.	Moderate	Moderate	Not applicable to all contaminants on site, industrial site use would limit implementation, and climatic conditions limit effectiveness.	No



	Remedial Technology Identification		Description of	Implementability of	Effectiveness of	Relative Cost of Remedial Technology		Summary of	Technology Retained
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	(Yes/No)
	Soil Excavation and Off-Site Disposal	Excavation and Landfill	Removal of impacted soil using common excavation techniques. Disposal of impacted soil at an off-site, permitted landfill. May include treatment of contaminated soil by off-site landfill prior to disposal.	Technically implementable where accessibility allows for excavation.	Effective for all site soil contaminants.	High	Low	Commonly used established technology effective for all site soil contaminants.	Yes
	Soil Excavation, Ex Situ Treatment and Off-Site Disposal/On Site Reuse	Solidification/ Stabilization (S/S)	Removal of impacted soil using common excavation techniques. Contaminants are physically bound or enclosed within a stabilized mass using cementitious reagents (cement, lime, etc.) or surface adsorption/chemical reagents.	Requires sufficient space on site to set up temporary treatment plant and treat/process excavated material prior to disposal. S/S processes may result in an increase in the overall volume of material for off-site disposal/on-site reuse. Additionally S/S processes increases density which increases disposal costs.	Stabilization is a common and effective technology for reducing the leachability of metals in soil.	High	Low	Requires sufficient space on site to set up ex-situ treatment. High capital cost and does not provide specific advantage over in situ S/S.	No
Removal		Soil Washing	Removal of impacted soil using common excavation techniques. Wash soil with water-based surfactants, detergents, acids, etc., to remove chemicals from soil particles. Treat or dispose of high chemical concentration residuals fluids.	Technically implementable. Require sufficient space on site to set up temporary treatment plant and treat/process excavated material prior to disposal/reuse. Require treatment of residual fluids.	Effective for more soluble chemicals. Presence of fine- grained soils and debris limits effectiveness.	High	Moderate	High cost and uncertainty relative to other remedial technologies.	No
	Soil Excavation, Ex Situ	Incineration	Removed soil is heated above approximately 1,600 degrees Fahrenheit to volatilize and combust organic contaminants. Incinerator off-gas is treated in an air pollution control system.	Potentially difficult to implement. Limited space for on- site treatment system and staging. Specific feed size and material handling requirements may impact implement ability. Suitable off-site facility not currently identified.	Proven effective treatment for organics, however, ineffective for inorganic hazardous substances.	High	High	High cost and uncertainty relative to other remedial technologies. Not applicable to all contaminants on site.	No
T	Treatment and Off-Site Disposal/On Site Reuse	Bioremediation	Biodegradation of contaminants in removed soil is enhanced through modification of the material for microbial growth. Treatment is conducted in landfarm arrangement, aboveground reactor, or in treatment cells (biopiles).	Difficult to implement. Landfarming option may require use of a large amount of space, depending on quantity of excavated material. Slurry and biopile treatment require reactor or treatment cell construction. Leachate and off-gas require collection and treatment. Addition of additives may increase total bulk volume of treated material.	While bioremediation cannot degrade inorganic contaminants, bioremediation can be used to change the valence state of inorganics and cause adsorption, immobilization onto soil particulates, precipitation, uptake, accumulation, and concentration of inorganics in micro or macroorganisms.	Moderate to High	Moderate to High	Likely not effective and difficult to implement.	No

O&M = Operations and Maintenance

MNR = Monitored Natural Recovery

ISS = In Situ Stabilization

ISCO = In Situ Chemical Oxidation

S/S = Solidification and Stabilization

Shading indicates remedial technology retained for cleanup action evaluation.



Groundwater Remedial Technologies Screening

	Remedial Technology Identification		Description of	Implementability of	Effectiveness of	Relative Cost of Remedial Technology		Summary of	Technology Retained
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	(Yes/No)
No Action	No Action	None	No institutional controls or treatment.	Not effective for protecting human health and environment.	Implementable but not acceptable to the general public or government agencies.	None	None	Used as a baseline for comparison.	No
		Environmental Covenant	Legal restrictions associated with future land use and activities (e.g., development, construction, etc.); may also be used to specify long-term maintenance requirements of remediation systems.	Technically implementable. Specific legal requirements and authority would need to be met.	Not effective for remediating contaminants. Can be effective at reducing risks and maintaining integrity of a remedy.	Low	Low	Applicable and/or required in combination with other technologies.	Yes
	Governmental/ Property Controls	Groundwater Use Restrictions, Groundwater Management Plans/ Requirements	Restrictions on groundwater extraction and use and/or exposure of humans and environment to hazardous substances present in groundwater. Implement groundwater management plans/requirements so that contaminated groundwater is managed properly in an event that it is necessary to remove groundwater (e.g., utility work, etc.).	Technically implementable but administratively more difficult. Requires an implementing agency.	Not effective for remediating contaminants. Enforcement would be required for restrictions to be effective.	Low	Low	Applicable and/or required in combination with other technologies.	Yes
Containment	Physical Groundwater Barrier	Low-Permeability Vertical Barrier	Construction of a low-permeability vertical barrier such as driven steel sheet piles, soil-bentonite or cement-bentonite wall to restrict groundwater flow and contaminant migration in the downgradient direction. Barrier can be installed down to the nearest aquitard to provide full containment, or installed at a partial depth to direct groundwater deeper. Groundwater extraction may be required to achieve containment under some scenarios. Longterm monitoring of containment structure required.	Technically implementable. A sheet pile wall is currently present which separates a majority of the Marine Area from the Upland Area.	Established technology effective for reducing mobility of contaminants. Effective for containing impacted groundwater or directing groundwater away from a source. However, does not provide treatment of contaminants. Effectiveness likely to increase if implemented to encapsulate the entire source area such that upgradient groundwater flows around the source area thereby minimizing contaminant mobility.	Low	Low	Applicable and/or required in combination with other technologies. A sheet pile wall barrier already exists at Site which separates a majority of the Marine and Upland Areas.	Yes
	Hydraulic Groundwater Barrier	Groundwater Pumping	Groundwater pumping to establish a hydraulic capture zone and restrict groundwater flow and contaminant migration in the downgradient direction. May be used in conjunction with a physical barrier to achieve full containment.	Technically implementable using standard groundwater extraction methods. The need to treat extracted groundwater to acceptable levels to allow discharge will reduce the implementability.	Potentially effective for hydraulic control of impacted groundwater. May be implemented to increase effectiveness of physical barrier technologies. Requires continuous long-term operation to achieve effective containment and maintenance of treatment components to prevent discharge of contaminated groundwater.	Moderate	High	Potentially applicable in combination with other technologies, but at high cost. Not expected to be cost effective if applied as sole containment method.	No



Remedial Technology Identification		Description of	Implementability of	Effectiveness of	Relative Cost of Remedial Technology		Summary of	Technology	
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	Retained (Yes/No)
	Monitored Natural Attenuation (MNA)	Natural Attenuation	Monitoring of naturally occurring physical, chemical and biological processes that reduce the mass, toxicity, mobility, volume, or concentration of contaminants in groundwater. Involves monitoring over time to confirm that natural processes are occurring to reduce risk associated with contaminant concentrations. A contingency plan is needed if the expected processes do not occur.	Technically implementable but requires long-term monitoring. Cleanup time frame may be longer than other remedial technologies. Source to groundwater generally requires treatment such as removal, containment or stabilization.	Effectiveness is dependent on site conditions and time frame for implementation. Not effective in preventing contaminant migration and/or exposure.	Low	Low	Applicable in combination with other technologies.	Yes
In Situ Treatment	Chemical Treatment	Chemical Oxidation	ENA is the use of low-energy, long-acting (sustainable) technologies to augment the natural attenuation processes. Oxygen releasing material (additive) is injected into ground to oxidize metals to a higher valence state which are more stable and less mobile in groundwater. Enhanced attenuation is based on a mass balance between contaminant loading into the system and the attenuation capacity of the system that will result in contaminants meeting the cleanup action objectives.	Technically implementable but requires long-term monitoring. Cleanup time frame longer than other remedial technologies but shorter compared to MNA. Source to groundwater generally requires treatment such as removal, containment or stabilization.	Not anticipated to be effective long term because additive injected into ground to alter geochemical parameters of groundwater can be consumed resulting in the release contaminants that were previously stabilized.	Low to Moderate	Low to Moderate	Not a permanent solution. Pre-treatment groundwater geochemical parameters may return after the additives are consumed resulting in the release contaminants. COCs in groundwater are below screening levels except for cPAHs at MW-8 which there is no identified source. Therefore the likelihood of implementing/maintaining remedy is uncertain/indefinite.	No
		Permeable Reactive Barrier (PRB)	PRBs are walls containing reactive media that are installed across the path of contaminated groundwater flow to intercept and treat contaminated groundwater. The barrier allows water to pass through while the media remove the contaminants by precipitation, degradation, adsorption, or ion exchange. PRB wall can be installed by excavating a trench (continuous or funnel/gate) or by injection method.	Technically implementable where accessibility allows for placement of reactive barrier.	Effective treatment configuration under proper hydrogeologic conditions that direct Site groundwater through PRB. Effectiveness relies on selecting an effective reactive treatment component. Reactive media used are chemical reagent such as zero valent iron (ZVI) or combination of ZVI, iron sulfide, iron oxide and/or calcium carbonate (pH adjusting agent).	Moderate	Moderate	PRB is most effective when combined with other technologies to reduce source area (i.e., removal or treatment). COCs in groundwater are below screening levels except for cPAHs at MW-8 which there is no identified source. Therefore the likelihood of implementing/maintaining PRB is uncertain/indefinite.	No



Remedial Technology Identification		Description of	Implementability of	Effectiveness of	Relative Cost of Remedial Technology		Summary of	Technology Retained	
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	(Yes/No)
In Situ Treatment	Physical Treatment	Electrokinetics Remediation (ER)	ER includes passing a low density current between electrodes to mobilize contaminants through soil and water in form of charged species. Positively-charged metal or metalloid cations migrate to the negatively-charged electrode (cathode), while metal or metalloid anions migrate to the positively charged electrode (anode). Contaminants arriving at the electrodes can be removed by means of electroplating/electrodeposition, precipitation/coprecipitation, adsorption, complexing with ion exchange resins, or by pumping of water (or other fluid) near the electrode.	Difficult implementability. ER is an emerging technology with relatively few applications for arsenic treatment.	Effectiveness may be limited by a variety of contaminants and soil and water characteristics. Treatment depth is limited by the depth to which the electrodes can be placed. ER is most applicable to saturated soil and soil with small particle sizes, such as clay.	High	High	Emerging technologies with limited case studies. Difficult Implementability. High capital and O&M cost.	No
Ex-Situ Treatment	Pump and Treat	Precipitation/ Coprecipitation	Extracted groundwater is treated by either mixing treatment chemicals into groundwater or by passing extracted groundwater through a fixed bed of media (e.g. ferric salts, alum) to form solid matrix through precipitation/coprecipitation. Usually involves pretreatment of pH adjustment and addition of chemical oxidant to create oxidizing environment to increase effectiveness. The precipitated/coprecipitated solid is then removed from the liquid phase by clarification or filtration.	Technically implementable. Long treatment time frame. Permitting may be required for discharge of treated water. May need to be combined with pre- and post-treatment steps. Treatment byproducts (e.g., settled solids) require management. Systems using this technology generally require skilled operators.	The effectiveness of this technology is less likely to be reduced by characteristics and contaminants other than arsenic, compared to other pump and treat water treatment technologies. It is also capable of treating heavy metals.	High	High	Potential physical constraints in relation to current/future site use. The nature of the groundwater contamination source at the site makes actively pumping and treating groundwater expensive and timeframe for running an active system uncertain if source to groundwater contamination is left inplace.	No
Treatment	Pump and Treat	Adsorption	Extracted groundwater is treated by passing extracted groundwater through a fixed bed of adsorption media (e.g. activated alumina, activated carbon). As contaminated water is passed through the adsorption media, contaminants are adsorbed. When adsorption sites become filled, the column must be regenerated or disposed of and replaced with new media. Like precipitation/coprecipitation, this technology requires pretreatment of pH adjustment and addition of chemical oxidant to create oxidizing environment to increase effectiveness.		Effectiveness of adsorption treatment process is sensitive to a variety of untreated water contaminants and characteristics. Competition for adsorption sites could reduce the effectiveness of adsorption because other constituents may be preferentially adsorbed, resulting in a need for more frequent bed regeneration or replacement. It is used less frequently than precipitation/coprecipitation, and is most commonly used as a polishing step for other water treatment processes.	High	High	Less effective in treating contaminants as compared to pump and treat with precipitation/coprecipitation.	No



	Remedial Technology Identification		Description of	Implementability of	Effectiveness of	Relative Cost of Remedial Technology		Summary of	Technology Retained	
General Response Action	Type of Remedial Technology	Process Option	Remedial Technology	Remedial Technology	Remedial Technology	Capital	0&M	Screening	(Yes/No)	
Ex-Situ Treatment	Pump and Treat	Ion Exchange: Xanthate Treatment	lon exchange removes ions from the aqueous phase by the exchange of cations or anions between the contaminants and the exchange medium. Ion exchange materials may consist of resins made from synthetic organic materials that contain ionic functional groups to which exchangeable ions are attached. They also may be inorganic and natural polymeric materials. After the resin capacity has been exhausted, resins can be regenerated for re-use.	Technically implementable. Long treatment time frame. Permitting may be required for discharge of treated water. May need to be combined with pre- and post-treatment steps. Treatment byproducts (e.g., treatment chemicals) require management. Systems using this technology generally require skilled operators.	Effectiveness of Ion Exchange treatment process is sensitive to a variety of untreated water contaminants and characteristics.	High	High	Less effective in treating contaminants as compared to pump and treat with precipitation/coprecipitation.	No	

Notes:

O&M = Operations and Maintenance

IC = Institutional Controls

MNR = Monitored Natural Recovery

ENA = Enhanced Natural Attenuation

ZVI = Zero Valent Iron

PRB = Permeable Reactive Barrier

ER = Electrokinetics Remediation

cm = centimeters





Cleanup Action Alternative Descriptions

Dakota Creek Industries Anacortes, Washington

					Cleanup Action Alte	rnative Components		
Matrix	Contaminants of Concern (COCs)	Cleanup Action Objectives (CAOs)	Alternative 1 - Containment and Compliance Monitoring	Alternative 2 - Partial Source Area Removal	Alternative 3 - Source Area In Situ Treatment	Alternative 4 - Source Area Removal	Alternative 5 - Site-Wide In Situ Treatment	Alternative 6 - Site-Wide Removal
Soil and Groundwater	■ Arsenic ■ Nickel ■ Total cPAH TEQ	 ■ Prevent contact (dermal or incidental ingestion) by workers, visitors and other Site users with hazardous substances in soil and groundwater. ■ Prevent leaching of hazardous substances through the soil column to groundwater. ■ Prevent contact (dermal or incidental ingestion) by aquatic receptors to impacted groundwater that may discharge to the Marine Area resulting in acute or chronic effects. ■ Prevent the ingestion of aquatic organisms affected by the discharge of impacted groundwater to the Marine Area by higher trophic level ecological receptors. 	■ Maintenance of existing physical containment barriers including surface pavement and sheet pile bulkhead to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ Installation of new physical containment barrier (i.e., asphalt/concrete pavement) to further prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ Compliance Groundwater Monitoring ■ Institutional Controls ■ Annual Cap Inspection	■ Asphalt demolition, soil removal and offsite disposal of COCs in the southwest Source Area generally centered around location SB-12. ■ Verification Soil Sampling ■ Site Restoration ■ Maintenance of existing physical containment barriers including surface pavement and sheet pile bulkhead to prevent stormwater infiltration and contaminant leaching/ migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ Compliance Groundwater Monitoring ■ Institutional Controls ■ Annual Cap Inspection	■ Installation of new physical containment barrier (i.e., asphalt/concrete pavement) to further prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ Maintenance of existing physical containment barriers including surface pavement and sheet pile bulkhead to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ In situ soil treatment through injection of chemical reagents to immobilize/treat COCs in Source Areas generally centered around locations SB-12, GEI-17 and GEI-22. ■ Institutional Controls ■ Performance/Compliance Groundwater Monitoring ■ Institutional Controls ■ Annual Cap Inspection	■ Asphalt demolition, soil removal and offsite disposal of COCs in Source Areas generally centered around SB-12, GEI-17 and GEI-22. ■ Verification Soil Sampling ■ Site Restoration ■ Maintenance of existing physical containment barriers including surface pavement and sheet pile bulkhead to prevent stormwater infiltration and contaminant leaching/ migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ Compliance Groundwater Monitoring ■ Institutional Controls ■ Annual Cap Inspection	■ Installation of new physical containment barrier (i.e., asphalt/concrete pavement) to further prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ Maintenance of existing physical containment barriers including surface pavement and sheet pile bulkhead to prevent stormwater infiltration and contaminant leaching/migration through the soil column as well as to provide a physical barrier to prevent direct contact to Site COCs. ■ In situ soil treatment through injection of chemical reagents to immobilize/treat COCs throughout the Site. ■ Performance/Compliance Groundwater Monitoring ■ Institutional Controls ■ Annual Cap Inspection	■ Asphalt demolition, soil removal and offsite disposal of COCs throughout the Site. ■ Verification Soil Sampling ■ Site Restoration ■ Compliance Groundwater Monitoring
	E	stimated Alternative Cost (+50%/-30%) ¹	\$1,180,000	\$2,120,000	\$2,610,000	\$4,390,000	\$7,030,000	\$15,060,000
	Estimated Volume	of Contaminated Soil Removed/Treated	N/A	3,600 bcy	9,000 bcy	9,000 bcy	46,500 bcy	46,500 bcy
		Estimated Restoration Time frame	1-2 Years ²	1-2 Years ²	2-3 Years ²	2-3 Years ²	3-4 Years ²	3-4 Years ²

Notes:

TEQ = Toxicity Equivalence

COC = Contaminant of Concern

bcy = bulk (in-place) cubic yards

% = percent

N/A = Not Applicable



¹ Alternative cost estimates are presented in Appendix A.

² Compliance groundwater monitoring is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.

cPAH = Carcinogenic Polycyclic Aromatic Hydrocarbons

Evaluation of Cleanup Action Alternatives Dakota Creek Industries Feasibility Report

Anacortes, Washington

Evaluation Criteria	Alternative 1 - Containment and Compliance Monitoring	Alternative 2 - Partial Source Area Removal	Alternative 3 - Source Area In Situ Treatment
Compliance with MTCA Threshold Crite	ria		
Protection of Human Health and the Environment	Yes - Alternative would protect human health and the environment through a combination of containment technologies and institutional controls.	Yes - Alternative would protect human health and the environment through a combination of source area removal, containment technologies, and institutional controls.	Yes - Alternative would protect human health and the environment through a combination of source area in situ treatment, containment technologies, and institutional controls.
Compliance With Cleanup Standards	Yes - Alternative is expected to comply with cleanup standards. This alternative utilizes containment technologies and institutional controls to prevent exposure to contaminants in the subsurface. Compliance would rely on long-term monitoring and maintenance of institutional controls. Future development of property could potentially require additional environmental cleanup or special provisions.	Yes - Alternative is expected to comply with cleanup standards. This alternative utilizes partial source area removal, containment technologies, and institutional controls to prevent exposure to contaminants in the subsurface. Compliance would rely on long-term monitoring and maintenance of institutional controls. Future development of property could potentially require additional environmental cleanup or special provisions.	Yes - Alternative is expected to comply with cleanup standards. This alternative utilizes in situ treatment and containment technologies, and institutional controls to prevent exposure to contaminants in the subsurface. Compliance would rely on long-term monitoring and maintenance of institutional controls. Future development of property could potentially require additional environmental cleanup or special provisions.
Compliance With Applicable State and Federal Regulations	Yes - Alternative complies with applicable state and federal regulations.	Yes - Alternative complies with applicable state and federal regulations.	Yes - Alternative complies with applicable state and federal regulations.
Provision for Compliance Monitoring	Yes - Alternative includes provisions for compliance monitoring.	Yes - Alternative includes provisions for compliance monitoring.	Yes - Alternative includes provisions for compliance monitoring.
Restoration Time Frame			
Restoration Time Frame	A significant portion of the containment barriers are currently in place. Additional containment in the form of asphalt paving of existing gravel surfaces is expected to occur over a 1-2 year period. Monitoring of containment elements (i.e., asphalt/concrete pavement and sheet pile wall) and groundwater conditions to document compliance with cleanup objectives. Compliance groundwater monitoring is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.	A significant portion of the containment barriers are currently in place. The removal of COCs in southeast source area followed by restoration is expected to occur over a 1-2 year period. Monitoring of containment elements (i.e., existing asphalt/concrete pavement and sheet pile wall) and groundwater conditions Site to document compliance with cleanup objectives. Compliance groundwater monitoring in portions of the Site containing residual contamination is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.	A significant portion of the containment barriers are currently in place. Additional containment in the form of asphalt paving of existing gravel surfaces and the injection of chemical reagents to treat source areas COCs are expected to occur over a 2-3 year period. More than one injection event may be necessary. Monitoring of containment elements (i.e., asphalt/concrete pavement and sheet pile wall) and groundwater conditions to document compliance with cleanup objectives. Compliance groundwater monitoring is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.
Relative Benefits Ranking (Scored fron	Score = 2	Score = 6	Score = 5
Protectiveness	Achieves a moderate-low level of protectiveness as all portions of the Site containing COCs receive a protective containment barriers under this alternative to prevent potential human exposure and/or stormwater infiltration. However, contamination will be left onsite throughout the uplands in a heavy industrial and active site.	Achieves a moderate-high level of protectiveness s this alternative improves overall environmental quality onsite by removing the source area with elevated contaminant levels in soil and groundwater greater than the PCULs in the eastern portion of the Site. The remaining residual contamination will utilize existing asphalt and sheetpile wall barriers to prevent worker exposure under this alternative. Short-term on-site and off-site risk of exposure are slightly increased due to removal action and off-site disposal of contaminated soil over	Achieves a moderate level of protectiveness as all portions of the Site containing COCs receive a protective containment barrier under this alternative. Achieves a higher score then Alternative 1 since this alternative improves overall environmental quality through in situ treatment of COCs in identified Source Areas. However, there is no contaminant mass removal under this alternative, therefore receives a slightly lower score than Alternative 2.



Evaluation Alternative 1 - Containment and Criteria Compliance Monitoring		Alternative 2 - Partial Source Area Removal	Alternative 3 - Source Area In Situ Treatment		
	Score = 2	Score = 6	Score = 5		
Permanence	Achieves a low level of permanence since COCs remain in-place and/or untreated. Alternative 1 relies on the installation of additional pavement combined with other technologies to reduce the mobility of COCs.	Achieves a moderate level of permanence. The alternative receives a higher score as compared to Alternative 1 and 3 due to the removal and off-site disposal of COCs which provides a relatively higher level reduction in the toxicity, mobility or volume of COCs.	Achieves a moderate level of permanence since under this alternative due to the treatment/stabilization of COCs in Source Areas through in situ technologies combined with other technologies to reduce the toxicity, mobility or volume of COCs. However the site is located along the shoreline of a marine system and it is relying on effective monitoring of the cap to remain in place and undamaged in a heavy industrial site. Therefore receives a lower score than Alternative 2.		
	Score = 2	Score = 6	Score = 6		
Long-Term Effectiveness	Provides a level of certainty in long-term effectiveness as all areas containing COCs receive a protective impermeable cap, which reduces exposure risk and contaminant leaching from vadose to saturated zone. However, relies on diligence of entity where history of leadership changes, frequent leasee modifications, and heavy industrial activity damage has been shown to alter priorities and increase risk of exposure. This puts a heavy and unidentified cost on PLP related to approval under the EC, reporting and repair costs along with potential contaminant release while exposed.	Provides a higher level of certainty in long-term effectiveness over Alternative 1 due to the permanent removal of COCs in the southeastern Source Area. However, it requires the central and western portions of the upland to remain under institutional controls which needs to be monitored in a heavy industrial and active site.	Provides a moderate level of certainty in long-term effectiveness. Slightly higher score than Alternative 2 is achieved due to in situ treatment of COCs within each of the Source Areas. However, it received a lower score than Alternative 4 as decreasing metals mobility through in situ reduction can be reversed under certain conditions.		
	Score = 8	Score = 7	Score = 8		
Management of Short-Term Risks	Achieves a high level of confidence in managing short-term risk to human health and environment since this alternative involves construction of pavement. Exposure risk to Site COCs during pavement construction is low to negligible.	alth and environment since this alternative involves construction of vement. Exposure risk to Site COCs during pavement construction is low to management, transport and disposal of contaminated material. Receives a			
	Score = 7	Score = 6	Score = 6		
Technical and Administrative Implementability	Achieves a high level of implementability since this alternative involves construction of an asphalt cap, which is a proven remedial technology.	Achieves a moderate level of implementability due to the design and coordination associated with implementation of soil removal. Implementation will be challenging since it will likely impact current site use at the property.	Achieves a moderate level of implementability due to the design and coordination associated with implementation of in situ treatment technologies. Implementation will be challenging since it may impact current site use at the Property.		
	Score = 3	Score = 7	Score = 6		
Consideration of Public Concerns	Residual contamination remaining in place below containment features could result in concerns by the public and nearby property owners and potentially affect the future development and Site use. However, the further addition of asphalt pavement to reduce the potential for contaminant migration and exposure would slightly reduce public concerns.	Residual contamination remaining in place below containment features could result in concerns by the public and nearby property owners and potentially affect the future development and Site use. However, the removal of source material to reduce the potential for contaminant migration and exposure would reduce public concerns.	result in concerns by the public and nearby property owners and potential affect the future development and Site use. However, the further addition		

Notes:

COC = Contaminant of Concern



Evaluation of Cleanup Action Alternatives Dakota Creek Industries Feasibility Report Anacortes, Washington

Evaluation Criteria	Alternative 4 - Source Area Removal	Alternative 5 - Site-Wide In Situ Treatment	Alternative 6 - Site-Wide Removal
Compliance with MTCA Threshold Crite	ria		
Protection of Human Health and the Environment	Yes - Alternative would protect human health and the environment through a combination of source area removal, containment technologies, and institutional controls.	Yes - Alternative would protect human health and the environment through a combination of site-wide in situ treatment, containment technologies, and institutional controls.	Yes - Alternative would protect human health and the environment through complete source removal.
Compliance With Cleanup Standards	Yes - Alternative is expected to comply with cleanup standards. This alternative utilizes source area removal, containment technologies, and institutional controls to prevent exposure to contaminants in the subsurface. Compliance would rely on long-term monitoring and maintenance of institutional controls. Future development of property could potentially require additional environmental cleanup or special provisions.	Yes - Alternative is expected to comply with cleanup standards. This alternative utilizes site-wide in situ treatment, containment technologies, and institutional controls to prevent exposure to contaminants in the subsurface. Compliance would rely on long-term monitoring and maintenance of institutional controls. Future development of property could potentially require additional environmental cleanup or special provisions.	Yes - Alternative is expected to comply with cleanup standards to the greatest extent practicable. All contaminant exceedance will be removed for offsite disposal.
Compliance With Applicable State and Federal Regulations	Yes - Alternative complies with applicable state and federal regulations.	Yes - Alternative complies with applicable state and federal regulations.	Yes - Alternative complies with applicable state and federal regulations.
Provision for Compliance Monitoring	Yes - Alternative includes provisions for compliance monitoring.	Yes - Alternative includes provisions for compliance monitoring.	Yes - Alternative includes provisions for compliance monitoring.
Restoration Time Frame			
Restoration Time Frame	A significant portion of the containment barriers are currently in place. Additional containment in the form of asphalt paving of existing gravel surfaces and the removal of COCs in identified source areas followed by restoration are expected to occur over a 2-3 year period. Monitoring of containment elements (i.e., asphalt/concrete pavement and sheet pile wall) and groundwater conditions to document compliance with cleanup objectives. Compliance groundwater monitoring to evaluate residual contamination in other portions of the Site is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.	A significant portion of the containment barriers are currently in place. The injection of chemical reagents to treat COCs Site-wide are expected to occur over a 3-4 year period. In situ treatment activities may require phasing during implementation to reduce disturbances to the DCl operations as well as more than one injection event if necessary. Monitoring of containment elements (i.e., asphalt/ concrete pavement and sheet pile wall) and groundwater conditions to document compliance with cleanup objectives. Compliance groundwater monitoring is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.	Complete removal of COCs Site-wide followed by restoration are expected to occur over a 3-4 year period. Removal activities may require phasing during implementation to reduce disturbances to the DCI operations. Compliance groundwater monitoring to verify the effectiveness of the cleanup action is expected to occur over a 1-2 year period following removal.
Relative Benefits Ranking (Scored from			
	Score = 7	Score = 8	Score = 9
Protectiveness	Achieves a moderate-high level of protectiveness as all portion of the Site containing COCs receive a protective containment barrier under this alternative. Therefore, this alternative receives a slightly higher score than Alternative 2. This alternative improves overall environmental quality onsite by removing COCs in Source Areas through removal. Similar to Alternative 2, short-term on-site and off-site risk of exposure are increased due to removal action and off-site disposal.	Achieves a high level of protectiveness as all COCs are treated/stabilized through in-situ treatment. Overall environmental quality on Site is increased as well as exposure risk to contamination are reduced to high degree under this alternative. No risk of exposure off-site as contamination is not removed.	Achieves a high level of protectiveness as all COCs are removed from the site thereby increasing the overall environmental quality on site to the highest degree. However, short-term on-site and off-site risk of exposure are increased due to removal action and off-site disposal. Therefore gets a slightly higher score than Alternative 5.



Evaluation Criteria	Alternative 4 - Source Area Removal	Alternative 5 - Site-Wide In Situ Treatment	Alternative 6 - Site-Wide Removal
	Score = 7	Score = 8	Score = 10
Permanence	Achieves a moderate-high level of permanence. The alternative receives a higher score due to the removal and off-site disposal of Source Area COCs which provides a relatively higher level reduction in the toxicity, mobility or volume of COCs than Alternative 3.	Achieves a high level of permanence by reducing toxicity, mobility and volume of COCs through Site-wide in-situ treatment of all COCs.	Achieves highest level of permanent reduction of mass, toxicity, and mobility of hazardous substances throughout the Site through removal and off-site permitted disposal. This alternative would eliminate/minimize to the need to perform additional cleanup actions.
	Score = 7	Score = 8	Score = 10
Long-Term Effectiveness	Provides a high level of certainty in long-term effectiveness due to the permanent removal of COCs in the Source Areas in addition to other technologies implemented similar to Alternative 2. However, it requires the entire upland to remain under institutional controls which needs to be monitored in a heavy industrial and active site.	Provides a high level of certainty in long-term effectiveness due to the site-wide treatment/stabilization of COCs. However, it requires the entire upland to remain under institutional controls which needs to be monitored in a heavy industrial and active site.	Achieves highest level of long-term effectiveness through removal of hazardous substances from the Site to the greatest degree feasible and utilizes approved off-site disposal facilities for final disposition
	Score = 5	Score = 4	Score = 3
Management of Short-Term Risks	Achieves a moderate level of confidence in managing short-term risk due to degree of health and safety risks associated with heavy earthwork construction, and potential for exposure to COCs during removal, on-site management, transport and disposal of contaminated material.	Achieves a moderate-high level of confidence in managing short-term risk as there is some potential for exposure to contamination during in-situ injection of reagents as well as other construction related risks. Since in situ treatment is to be performed site-wide under this alternative, the short-term risk are higher than the risk associated with in situ treatment of Source Areas under Alternative 3. Therefore is scored slightly lower than Alternative 3.	Achieves a low level of confidence in managing short-term risk due to degree of health and safety risks associated with heavy earthwork construction, and potential for exposure to COCs during removal, on-site management, transport and disposal of contaminated material. Achieves a lower score than Alternative 4 due to higher volume of contaminated material that will be removed under this alternative.
	Score = 6	Score = 5	Score = 4
Technical and Administrative Implementability	Achieves a moderate level of implementability due to the design and coordination associated with implementation of soil removal. Implementation will be challenging since it will likely impact current site use at the property.	Achieves a low-moderate level of implementability due to the design and coordination associated with implementation of in situ treatment technologies. Receives a lower score than Alternative 3 since the extent of in situ treatment is larger making implementation more challenging.	Achieves a low-moderate level of implementability due to the design and coordination associated with implementation of soil removal. Receives a lower score than Alternative 4 since the extent of soil removal is larger making implementation more challenging.
	Score = 7	Score = 9	Score = 8
Consideration of Public Concerns	Residual contamination remaining in place below containment features could result in concerns by the public and nearby property owners and potentially affect the future development and Site use. However, the further addition of asphalt pavement and removal of source areas to reduce the potential for contaminant migration and exposure would reduce public concerns.	Site-wide in situ treatment to reduce the potential for contaminant migration and exposure would produce minimum public concerns. However, there may be public concern for the temporary disruptions to Site operations and increased traffic resulting from construction activities. However, long-term public concerns are expected to be low.	Soil contamination would be removed to the extent practical under this alterative. However, there may be public concern for the temporary disruptions to Site operations, increased traffic resulting from construction activities and potential spills during transport of contaminated soil to the landfill. However, long-term public concerns are expected to be low.

Notes:

COC = Contaminant of Concern



Cleanup Action Alternative Evaluation Summary and Ranking

Dakota Creek Industries Anacortes, Washington

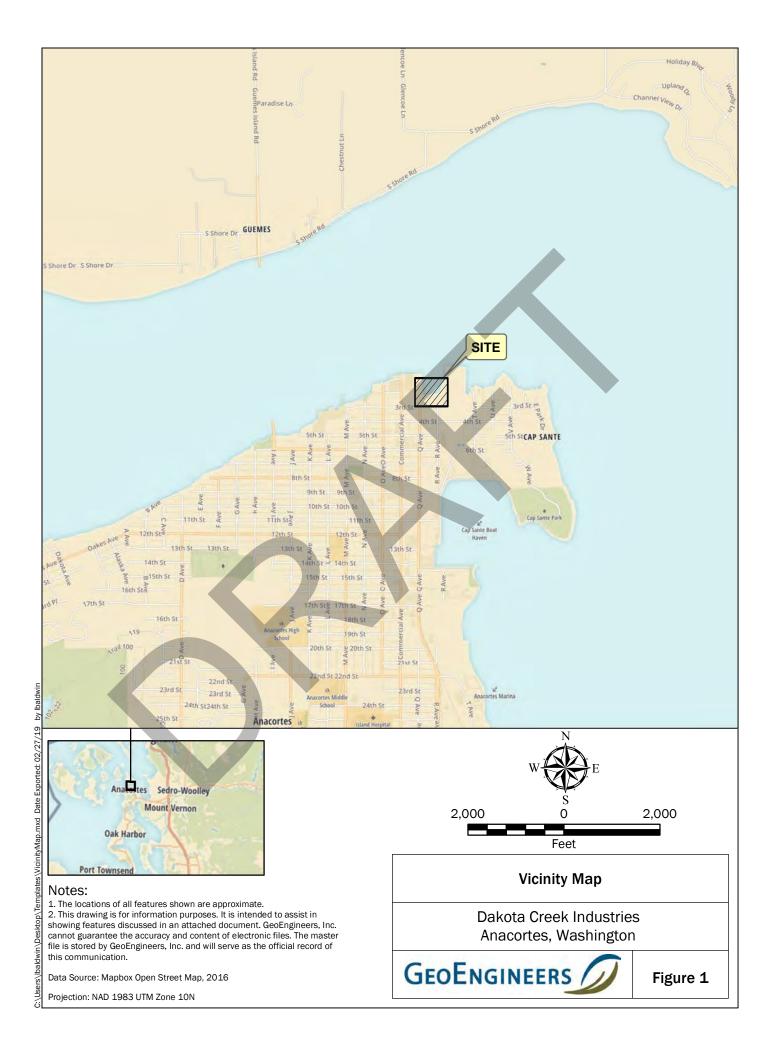
Remedial Alternative	Alternative 1 - Containment and Compliance Monitoring	Alternative 2 - Partial Source Area Removal	Alternative 3 - Source Area In Situ Treatment	Alternative 4 - Source Area Removal	Alternative 5 - Site-Wide In Situ Treatment	Alternative 6 - Site-Wide Removal
Evaluation						
Compliance with MTCA Threshold Criteria	Yes	Yes	Yes	Yes	Yes	Yes
Restoration Time Frame	1-2 Years ¹	1-2 Years ¹	2-3 Years ¹	2-3 Years ¹	3-4 Years ¹	3-4 Years
Estimated Volume of Contaminated Soil Removed/Treated	N/A	3,600 bcy	9,000 bcy	9,000 bcy	46,500 bcy	46,500 bcy
Relative Benefits Ranking ²						
Protectiveness (weighted as 30%)	0.6	1.8	1.5	2.1	2.4	2.7
Permanence (weighted as 20%)	0.4	1.2	1	1.4	1.6	2
Long-Term Effectiveness (weighted as 20%)	0.4	1.2	1.2	1.4	1.6	2
Management of Short-Term Risks (weighted as 10%)	0.8	0.7	0.8	0.5	0.4	0.3
Technical and Administrative Implementability (weighted as 10%)	0.7	0.6	0.6	0.6	0.5	0.4
Consideration of Public Concerns (weighted as 10%)	0.3	0.7	0.6	0.7	0.9	0.8
Overall Weighted Benefit Score	3.20	6,20	5.70	6.70	7.40	8.20
Disproportionate Cost Analysis						
Probable Remedy Cost (+50%/-30%, rounded)	\$1,180,000	\$2,120,000	\$2,610,000	\$4,390,000	\$7,030,000	\$15,060,000
Practicability of Remedy	Practicable	Practicable	Practicable	Practicable	Practicable	Practicable
Remedy Permanent to Maximum Extent Practicable	Yes	Yes	Yes	Yes	Yes	Yes
Relative Benefit Ranking to Remedial Cost (Benefit/\$1M)	2.71	2.92	2.18	1.53	1.05	0.54
Costs Disproportionate to Incremental Benefits	No	No	Yes	Yes	Yes	Yes
Overall Alternative Ranking	2 nd	1 st	3 rd	4 th	5 th	6 th

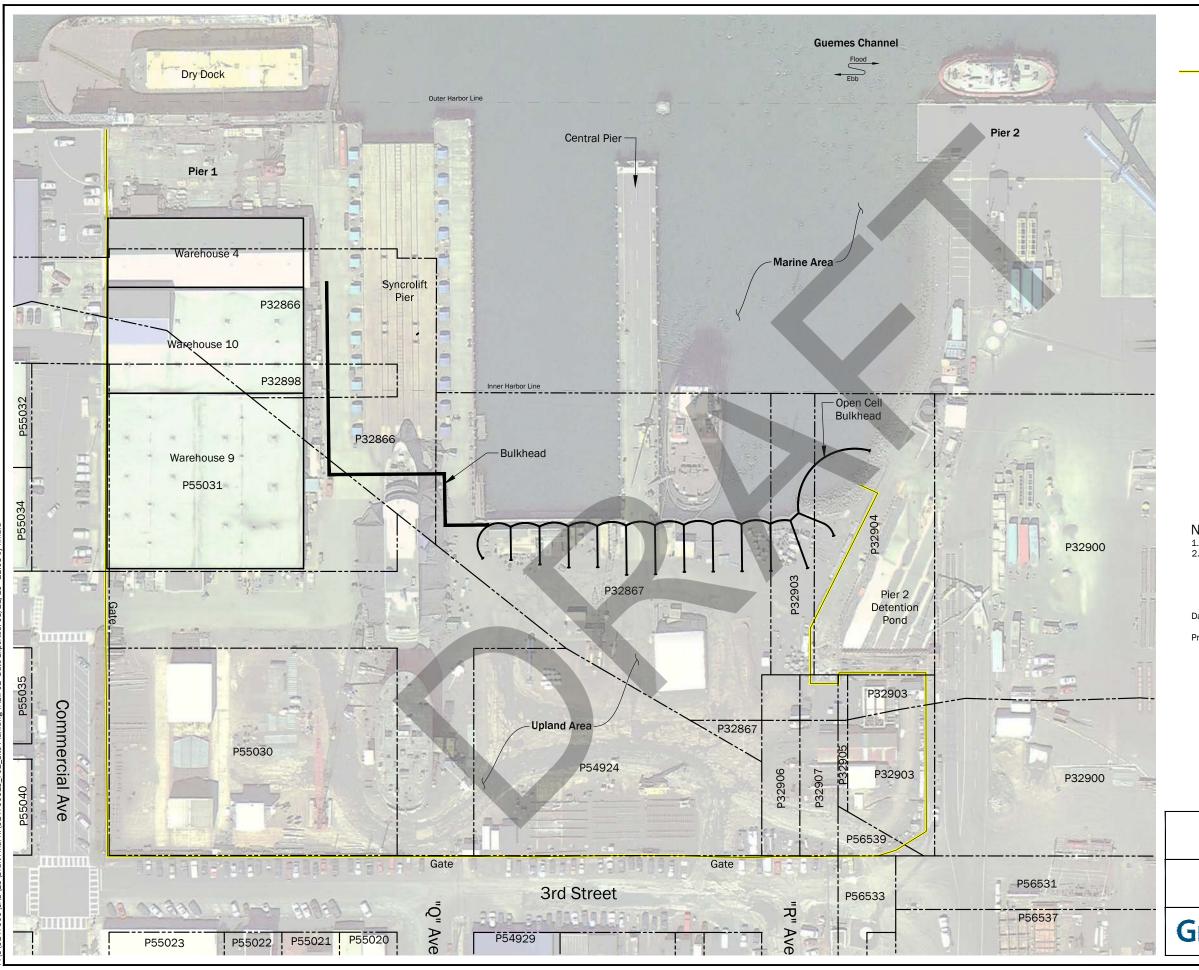
Note:



¹Compliance groundwater monitoring is expected to occur over a 5 year time frame (minimum). Additional long-term monitoring may be required to verify compliance with cleanup standards.

² Weightings were established by Ecology as referenced in their Opinion Letter dated December 28, 2009.





Legend

Dakota Creek Industries (DCI) Property Boundary

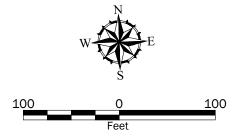
P32904 Skagit County Parcel Boundary and Number

Notes:

- The locations of all features shown are approximate.
 This drawing is for information purposes. It is intended to assist in showing features discussed in an attached document. GeoEngineers, Inc. cannot guarantee the accuracy and content of electronic files. The master file is stored by GeoEngineers, lead with large of the official record of this communication. Inc. and will serve as the official record of this communication.

Data Source: Aerial from Google Earth Pro dated 8/2011.

Projection: WA State Plane, North Zone, NAD83, US Foot

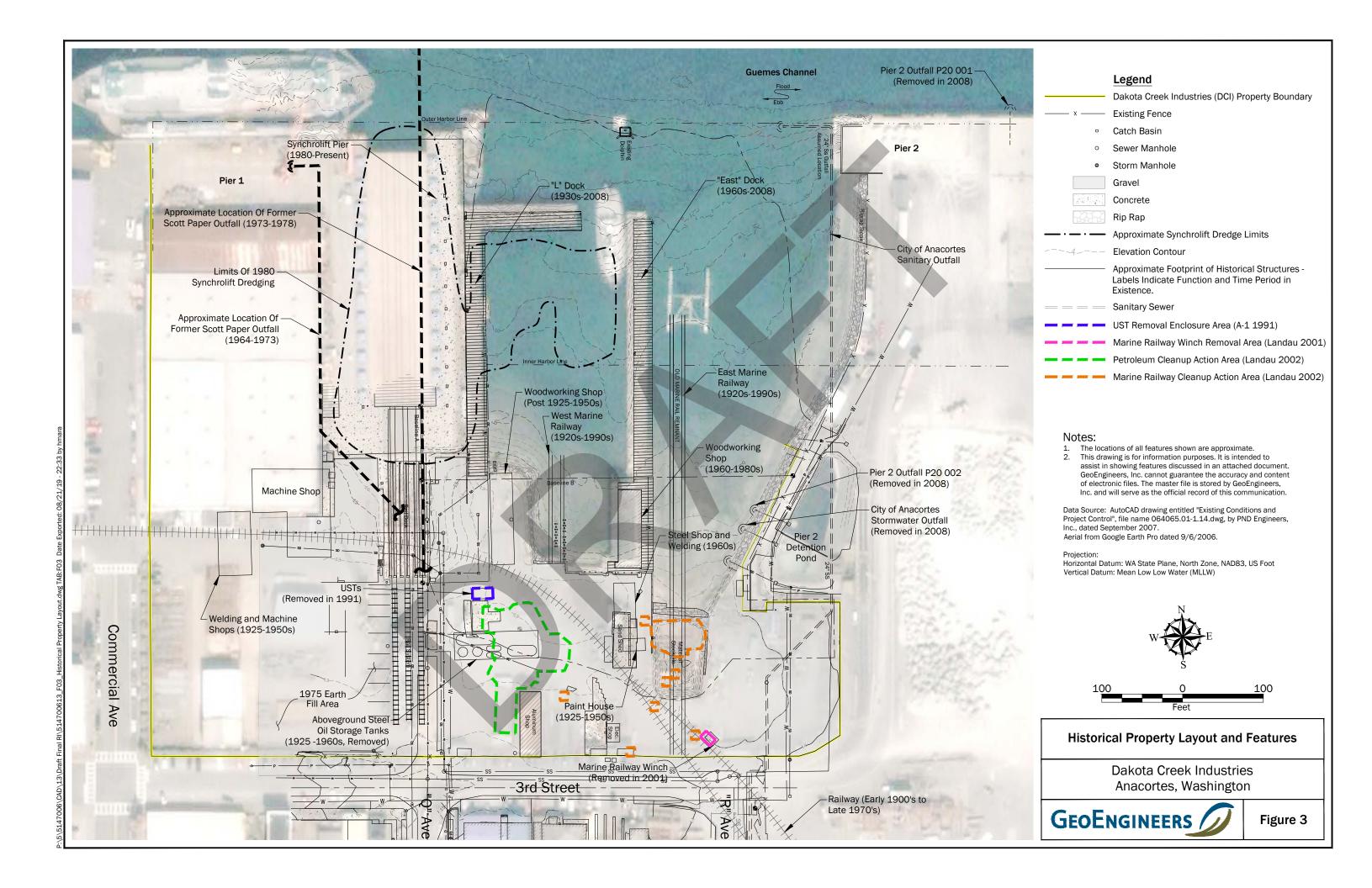


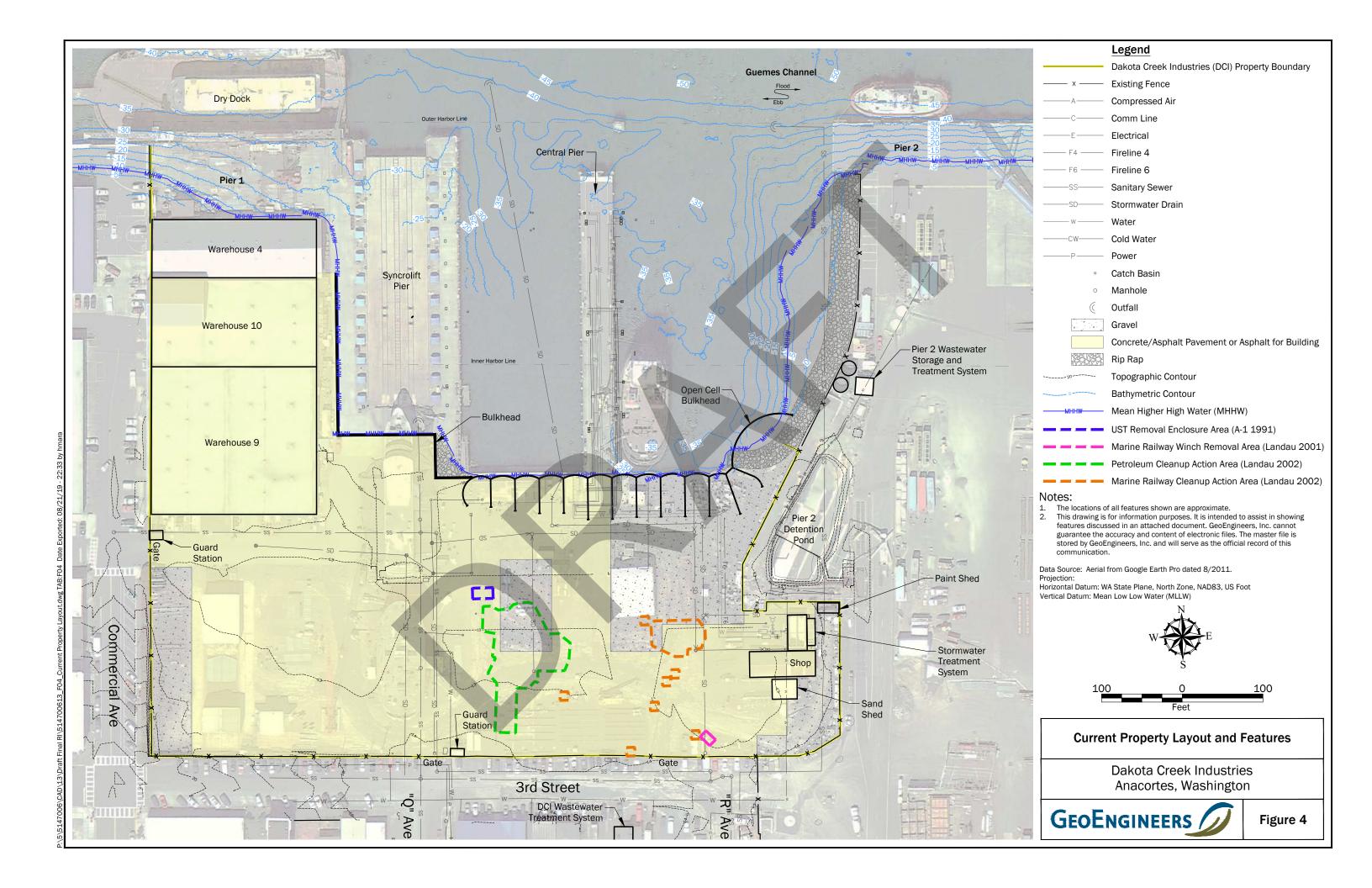
Site Plan

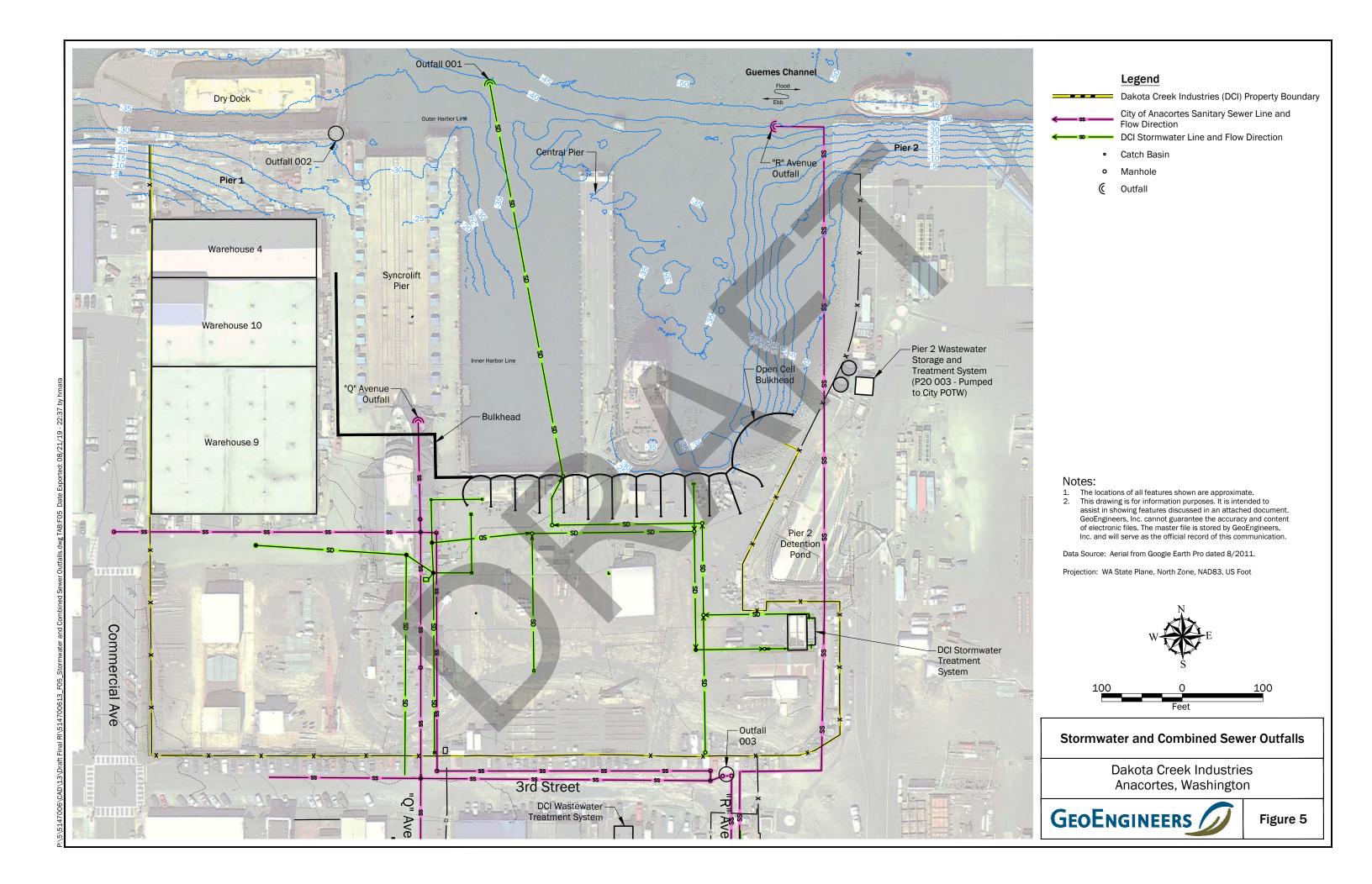
Dakota Creek Industries Anacortes, Washington

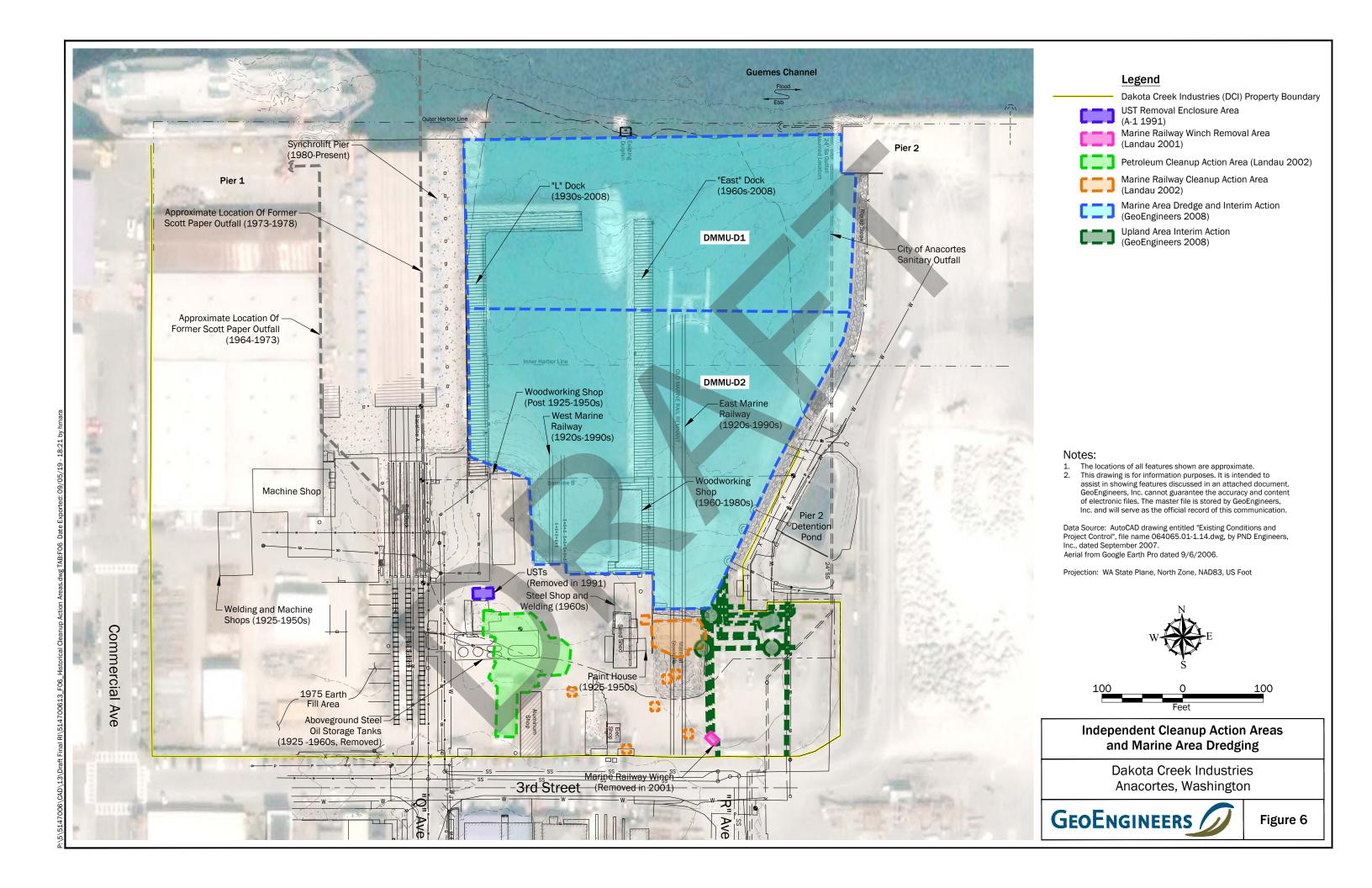


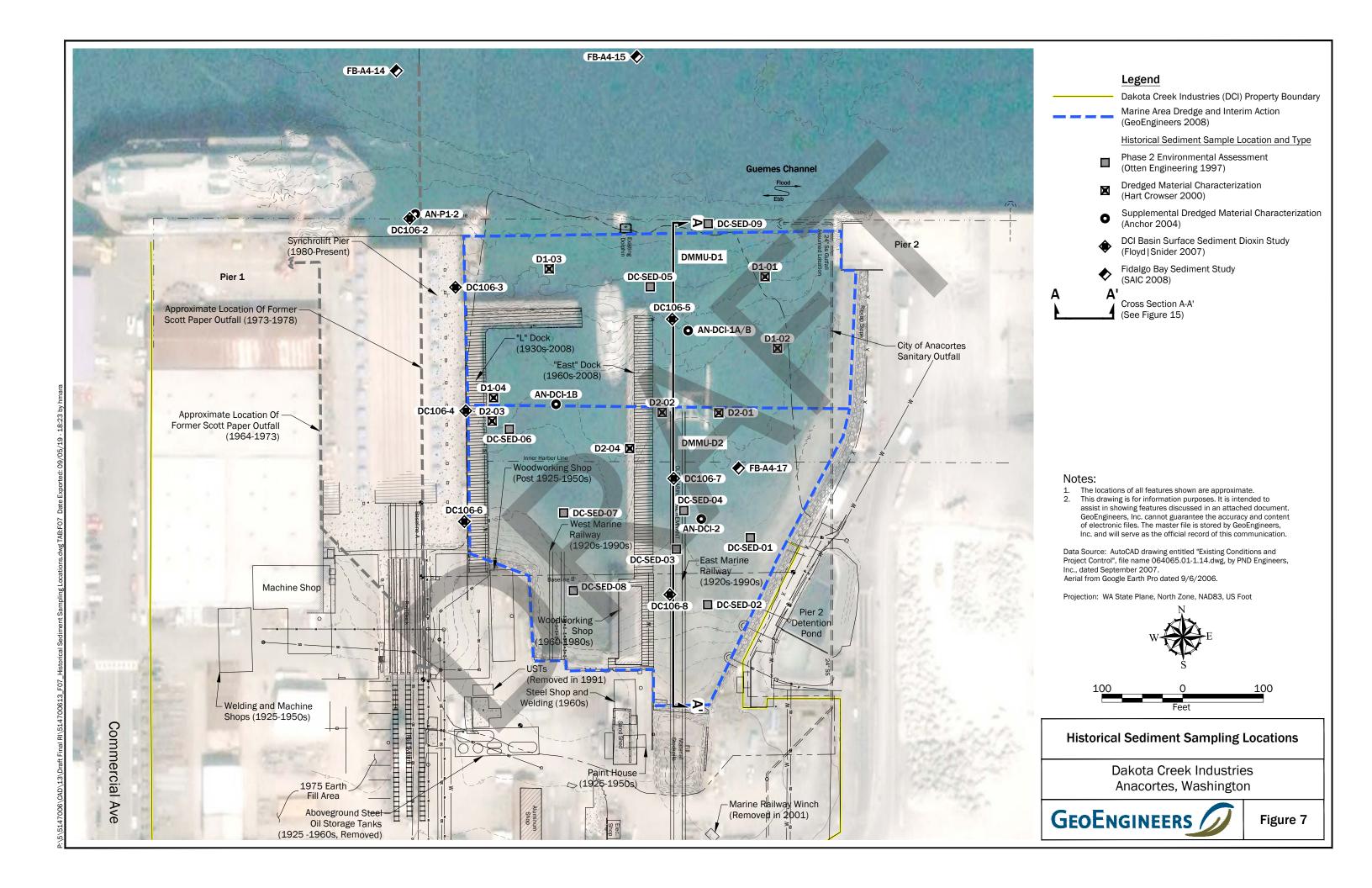
Figure 2

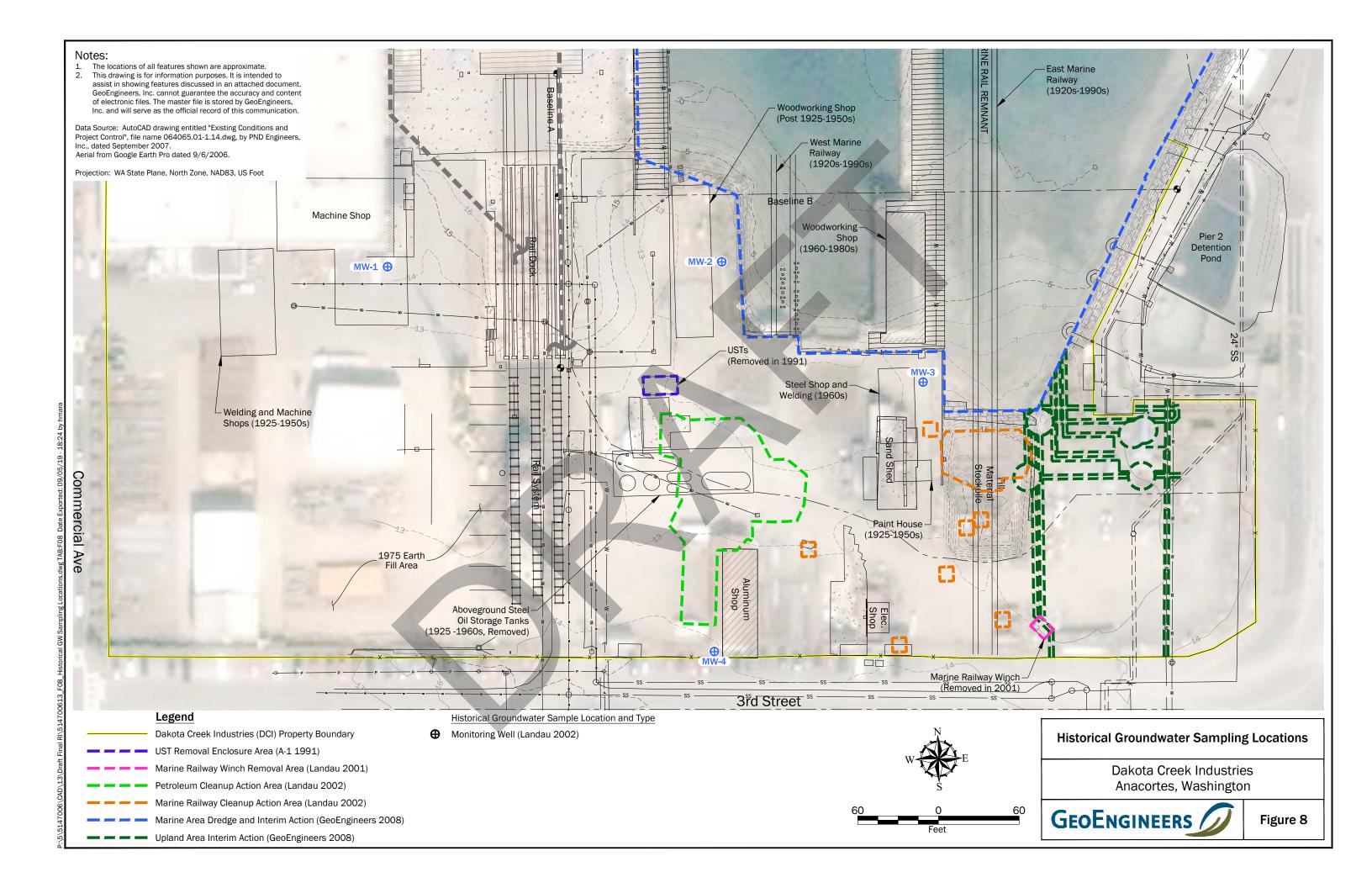


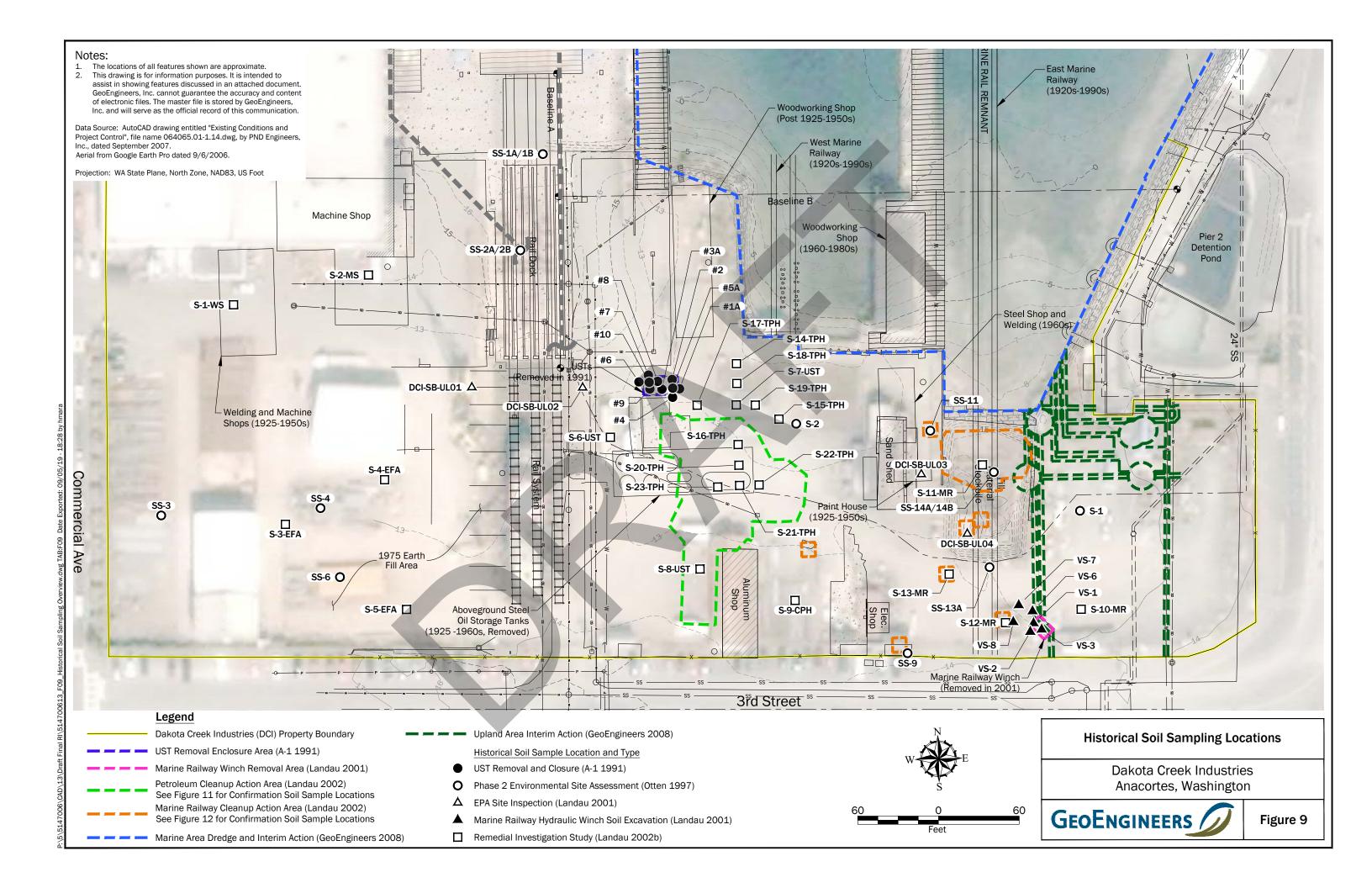


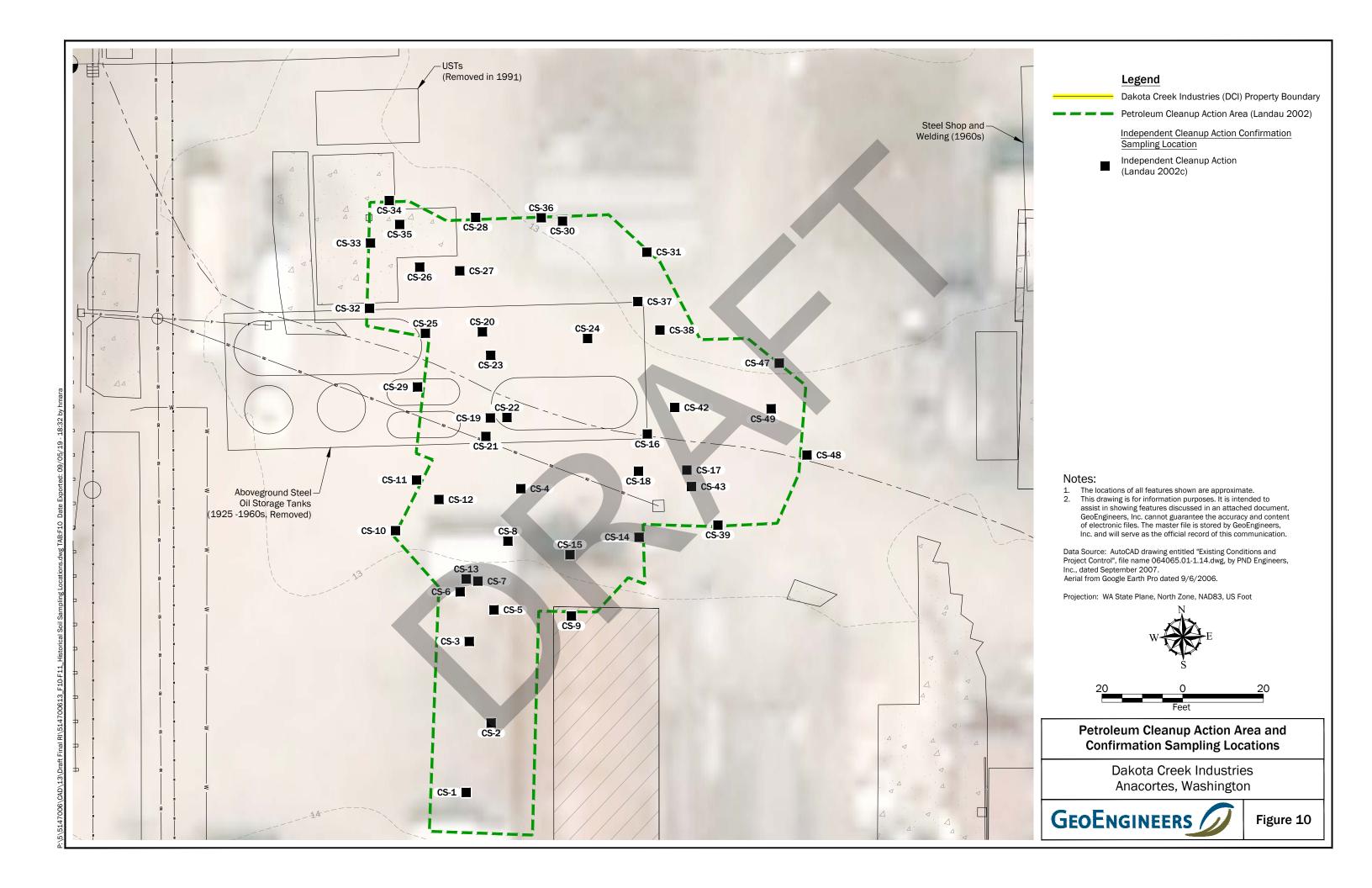


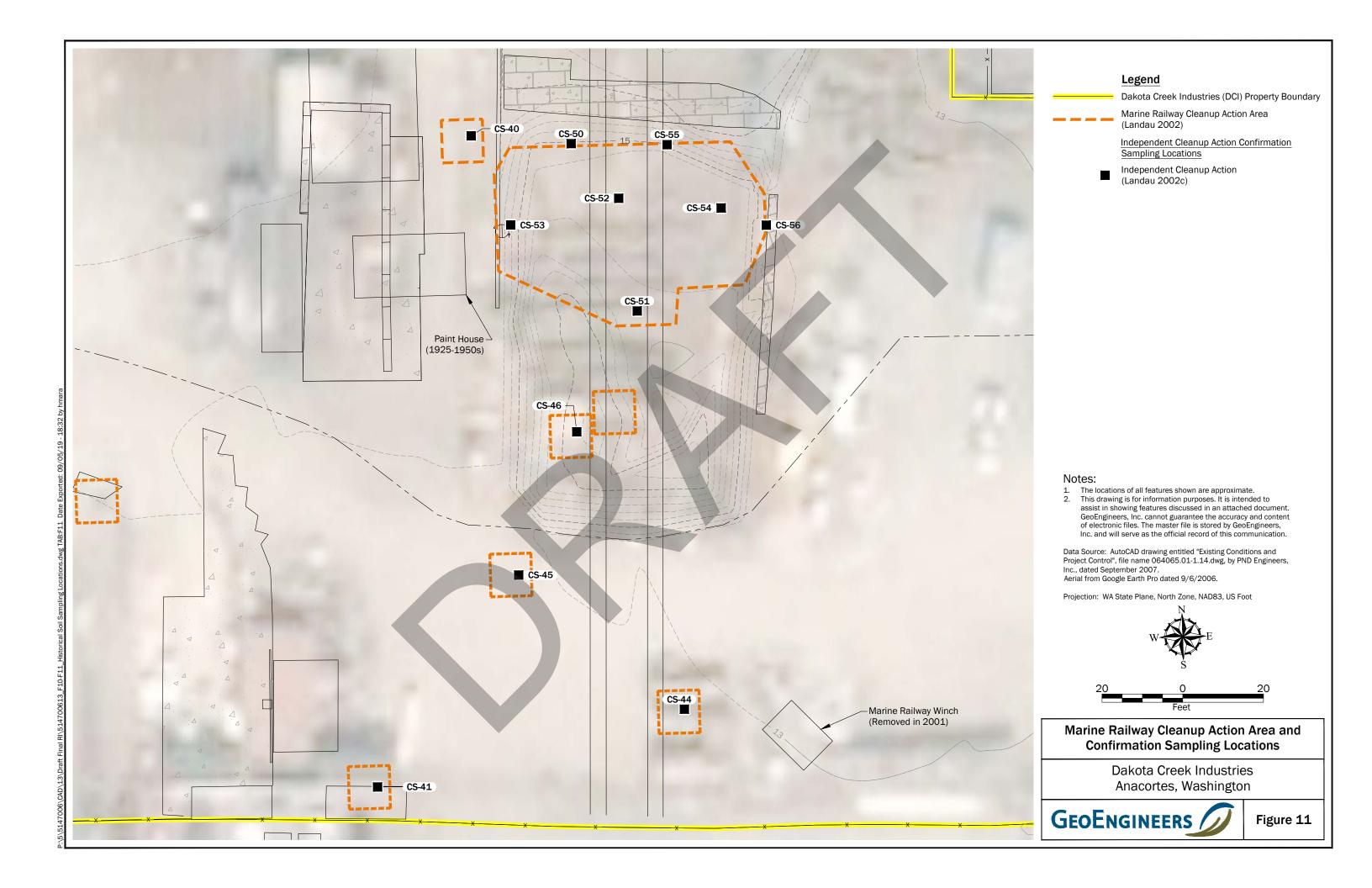


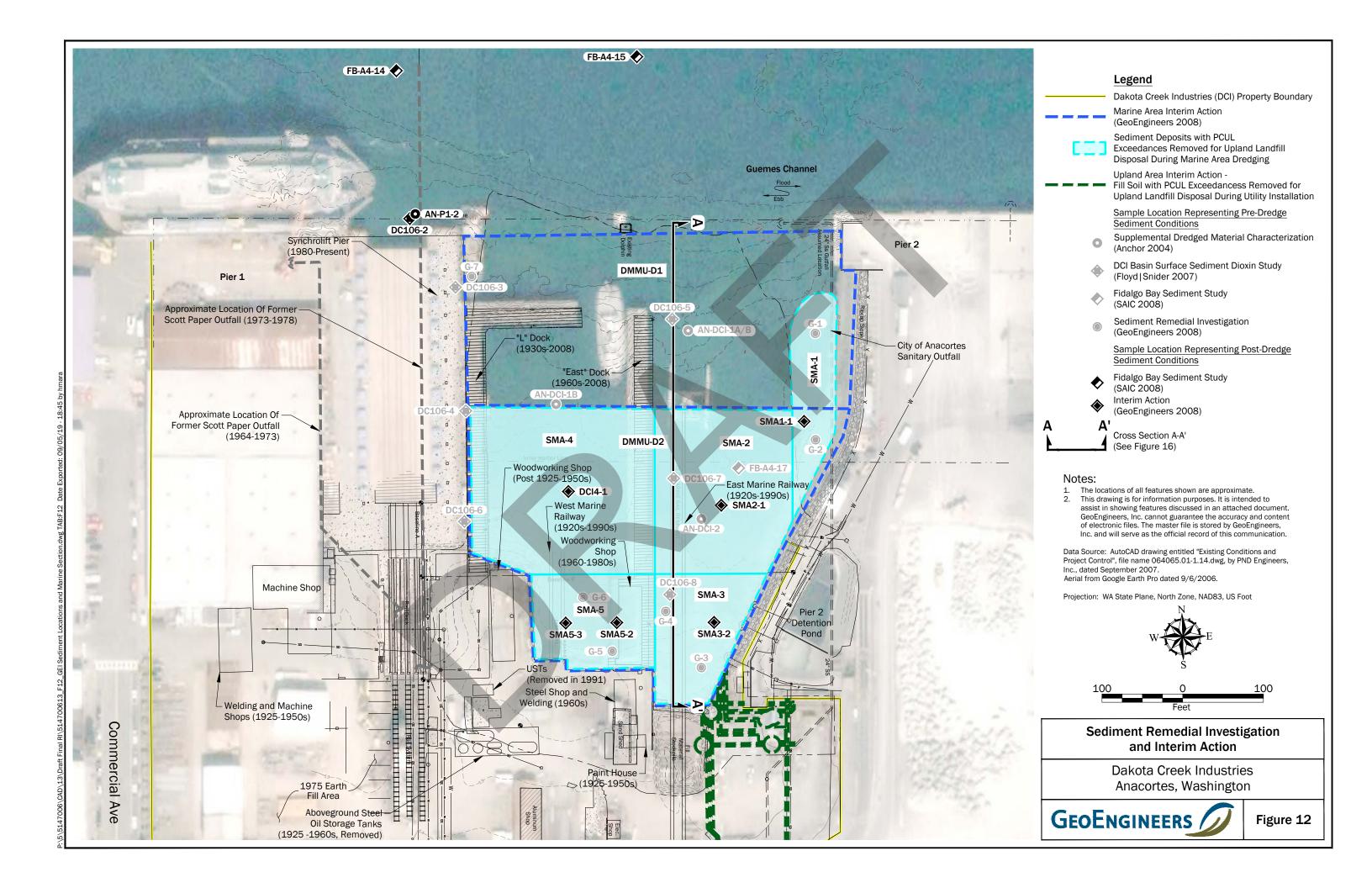


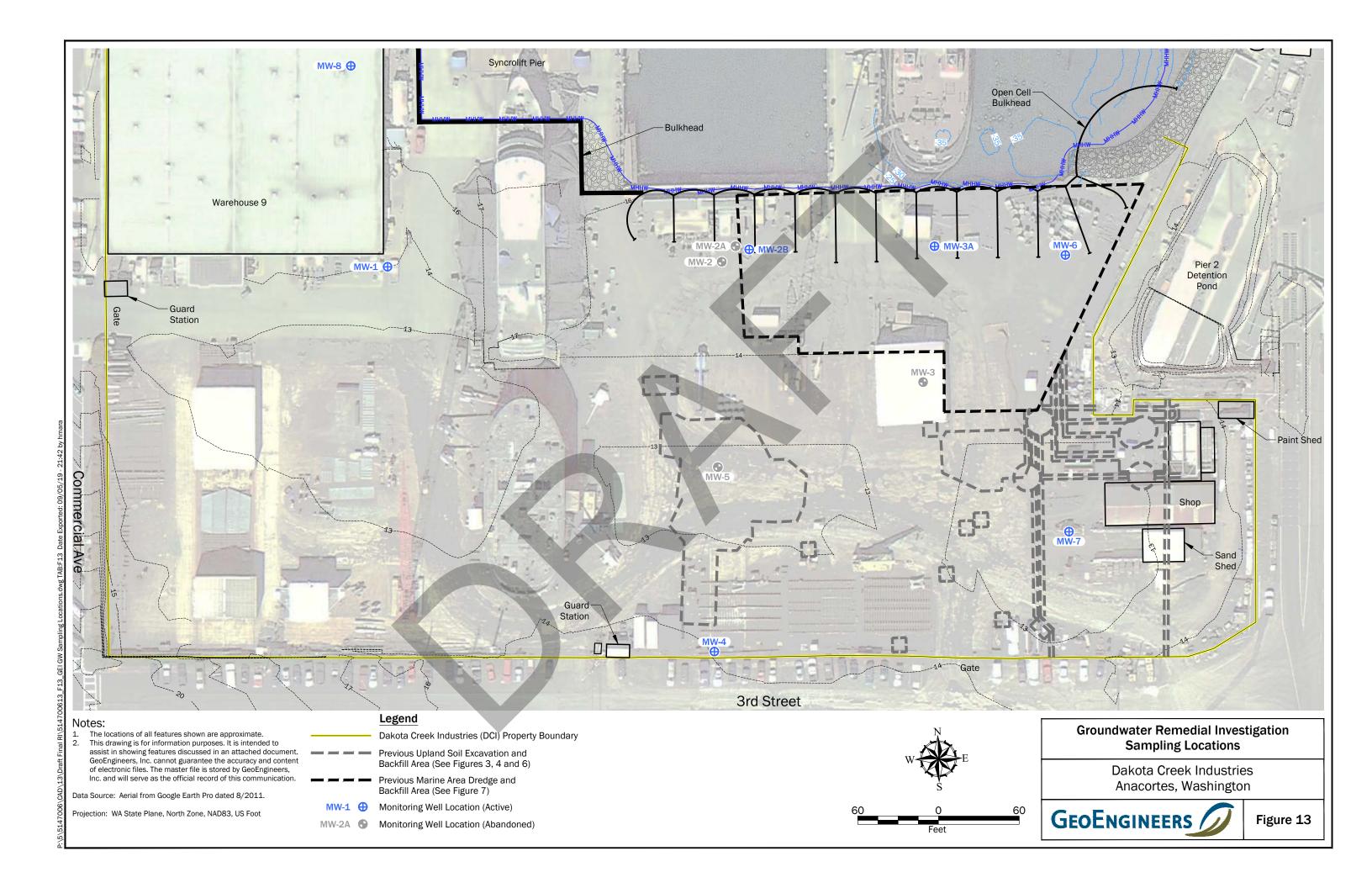


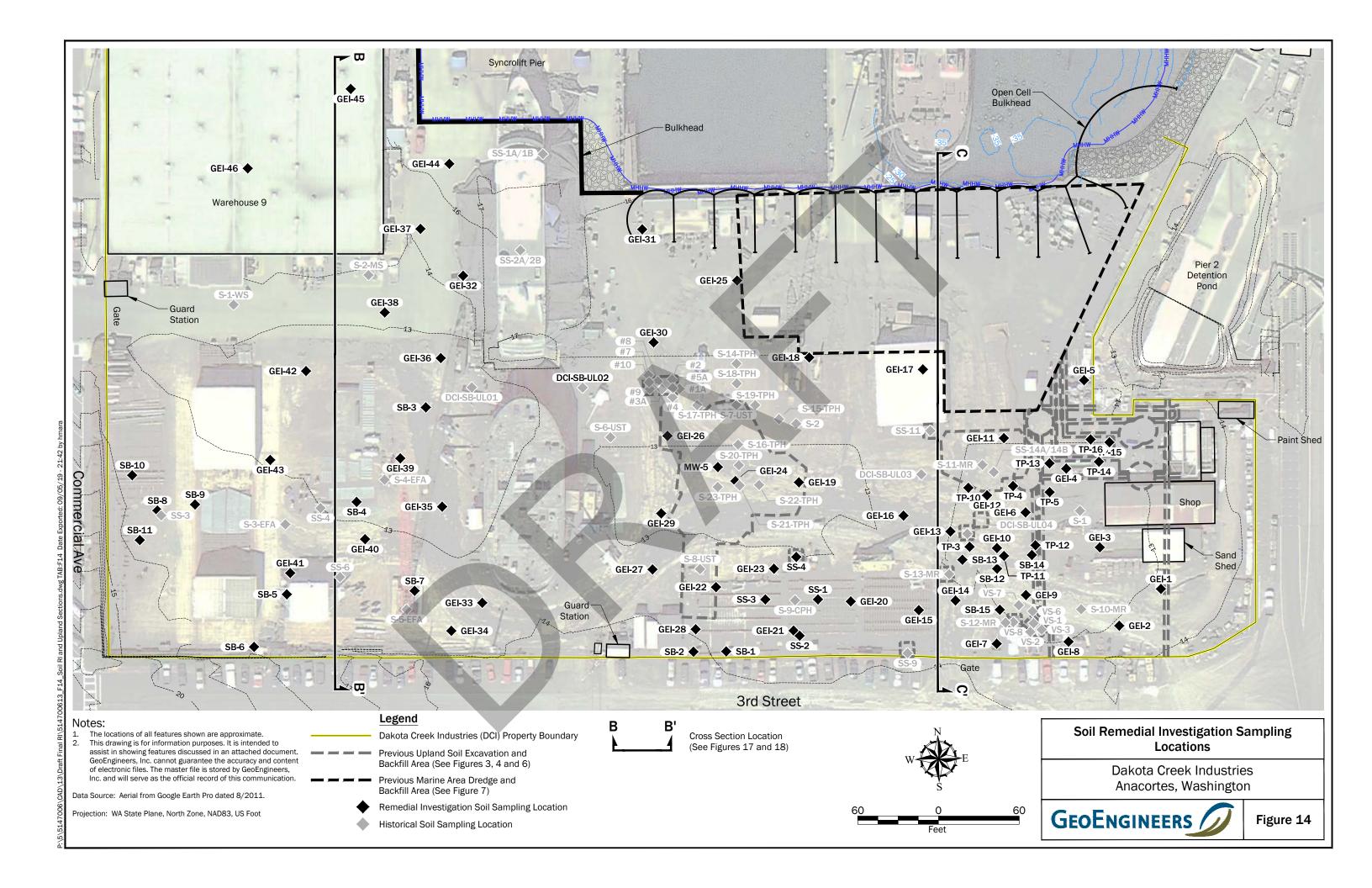


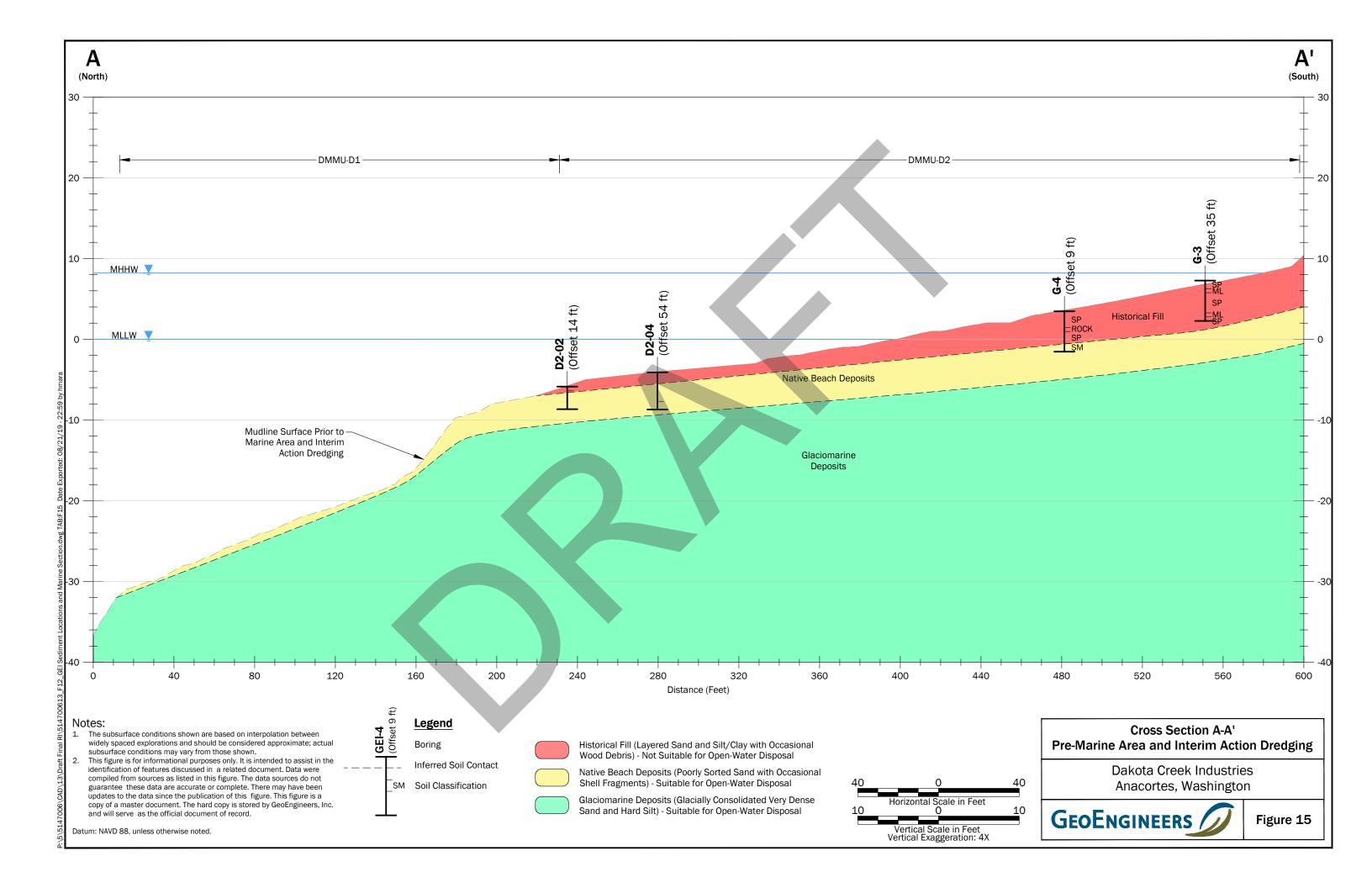


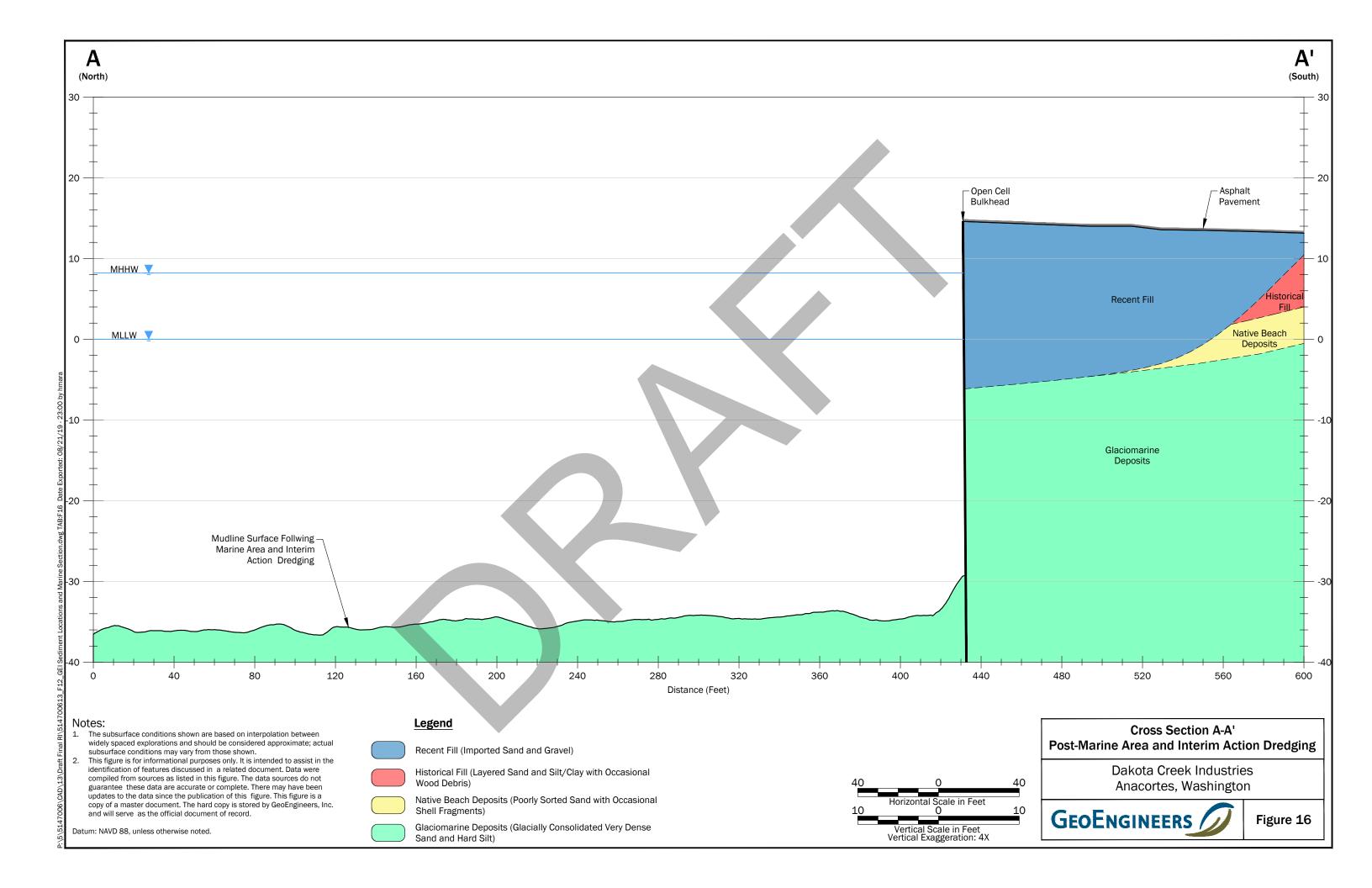


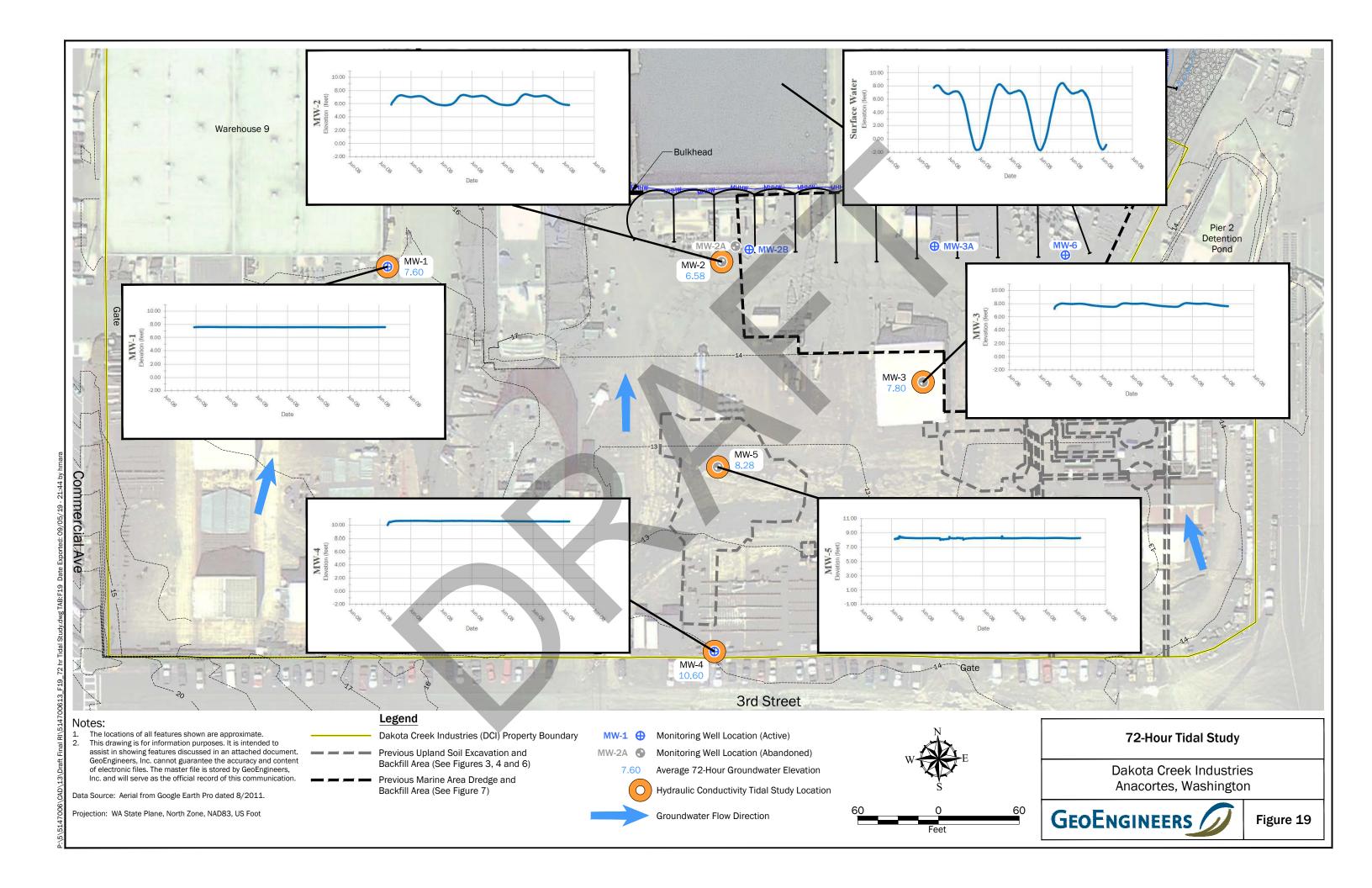


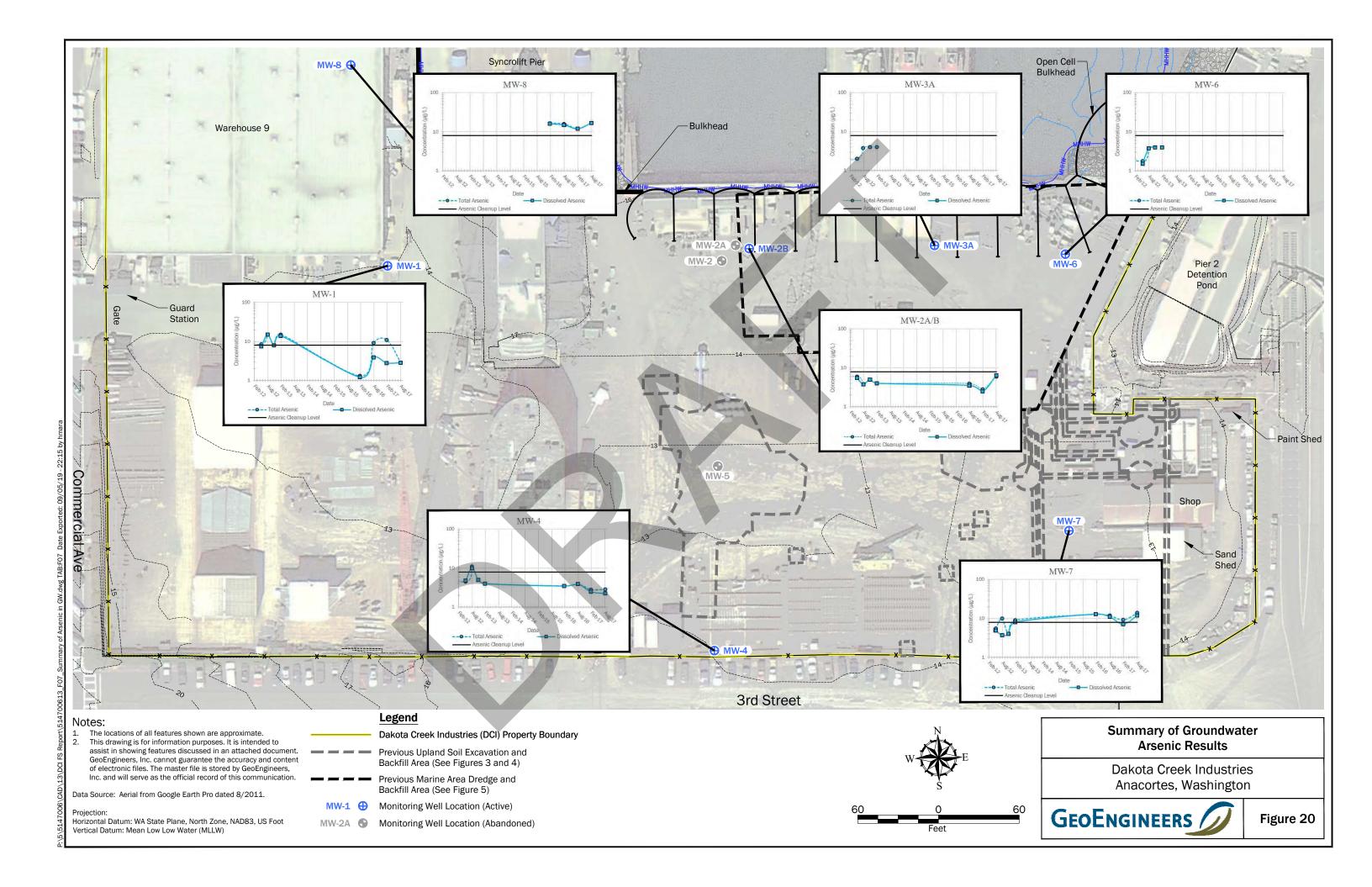


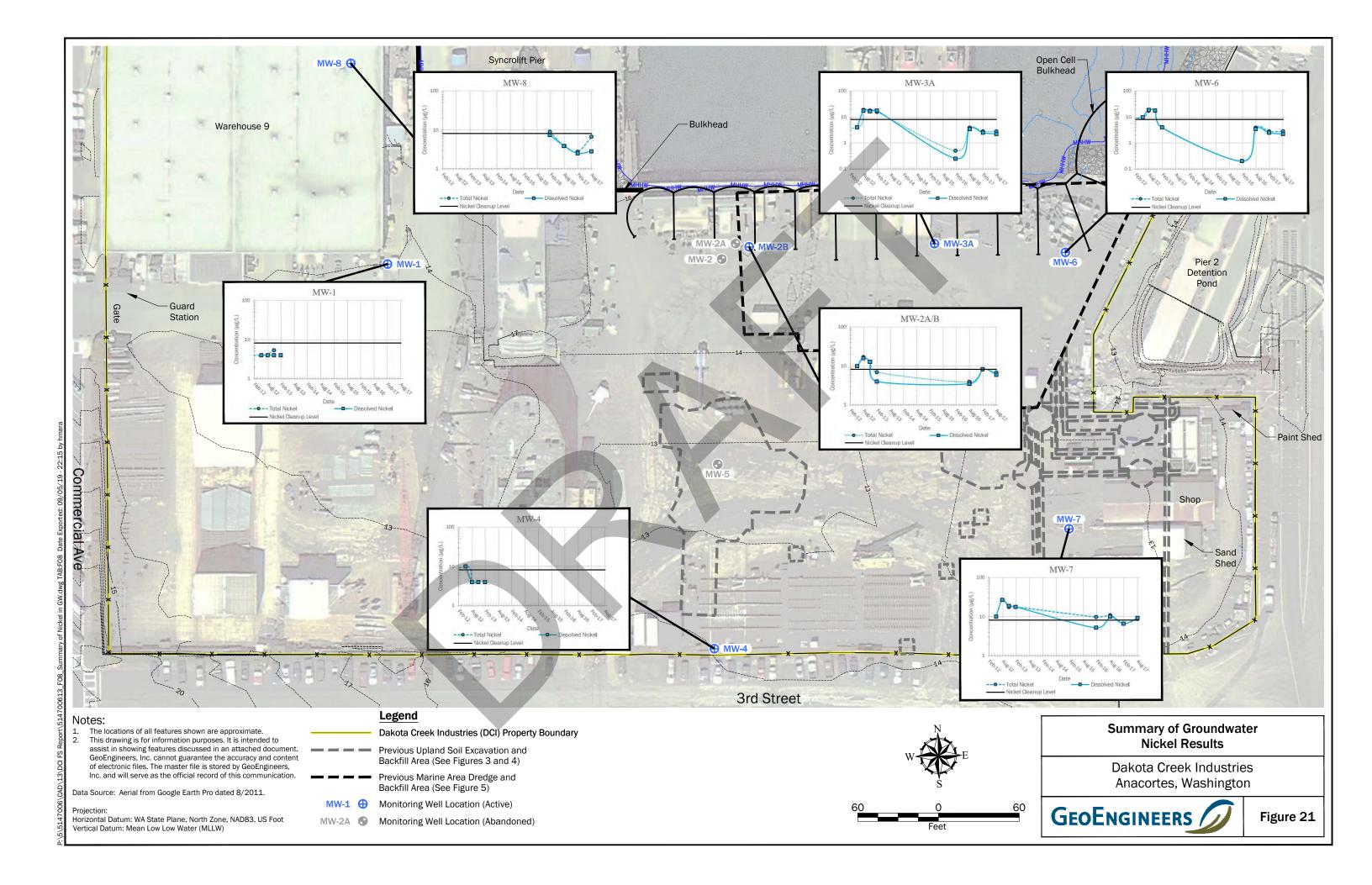


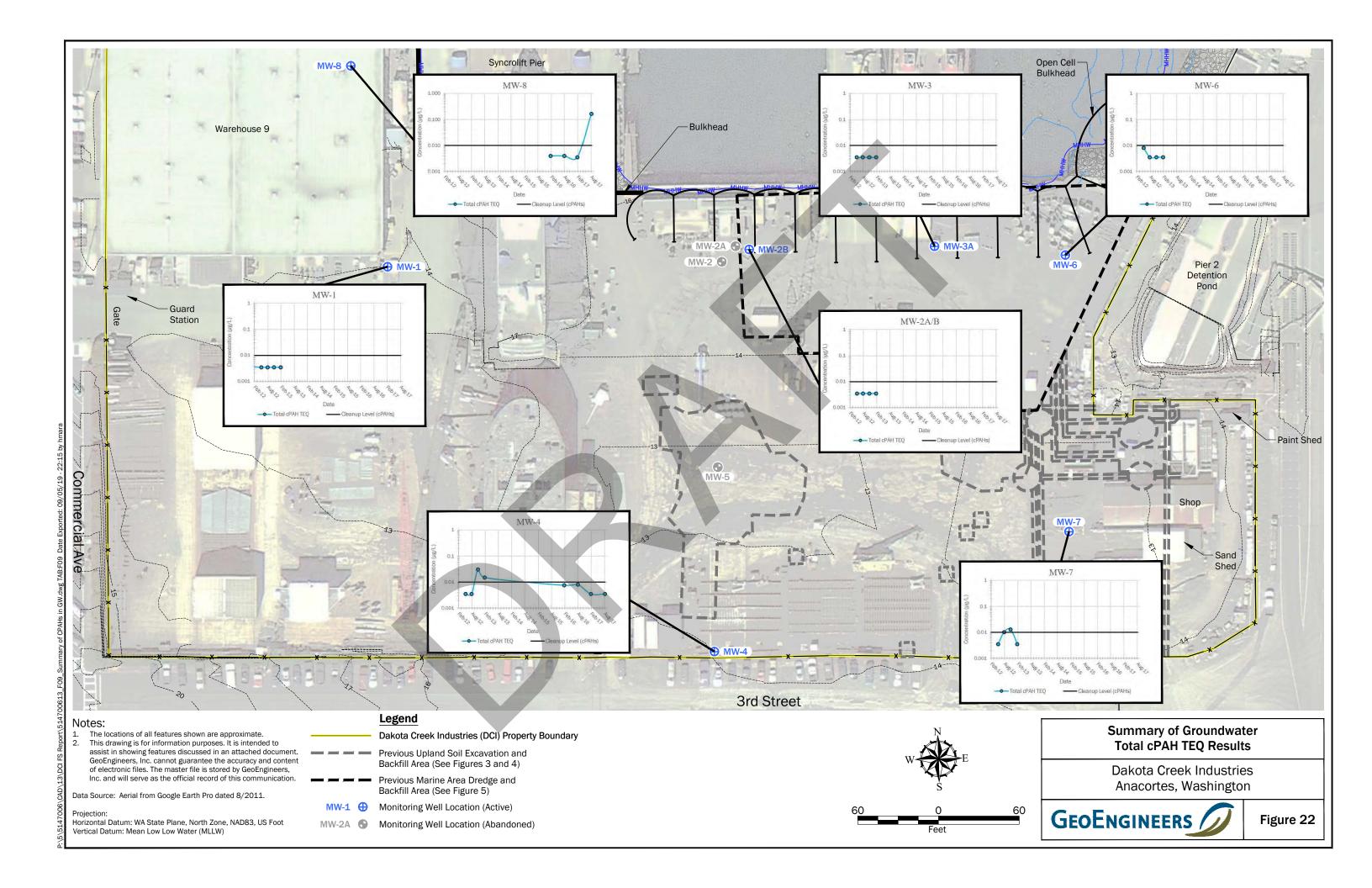


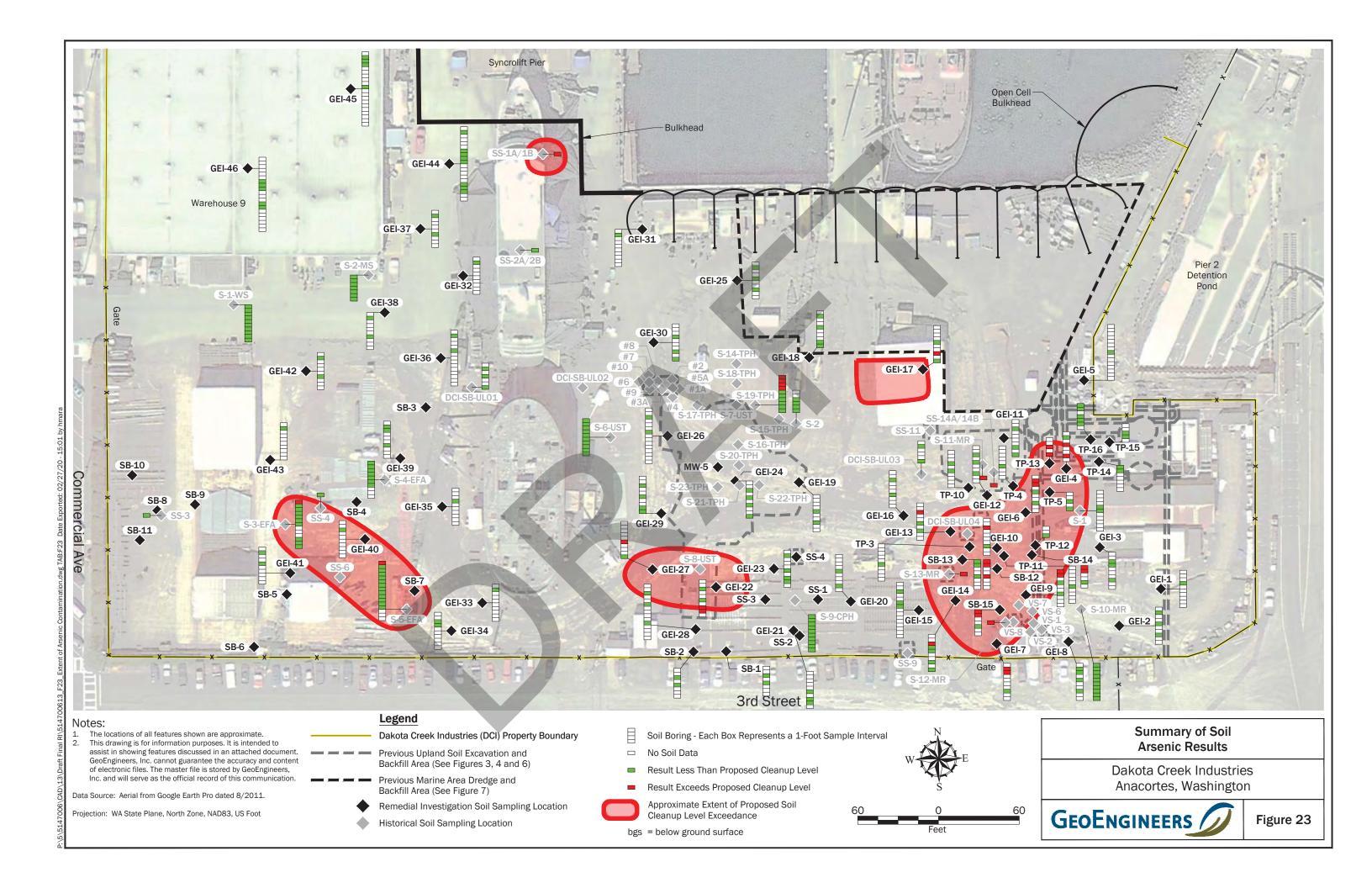


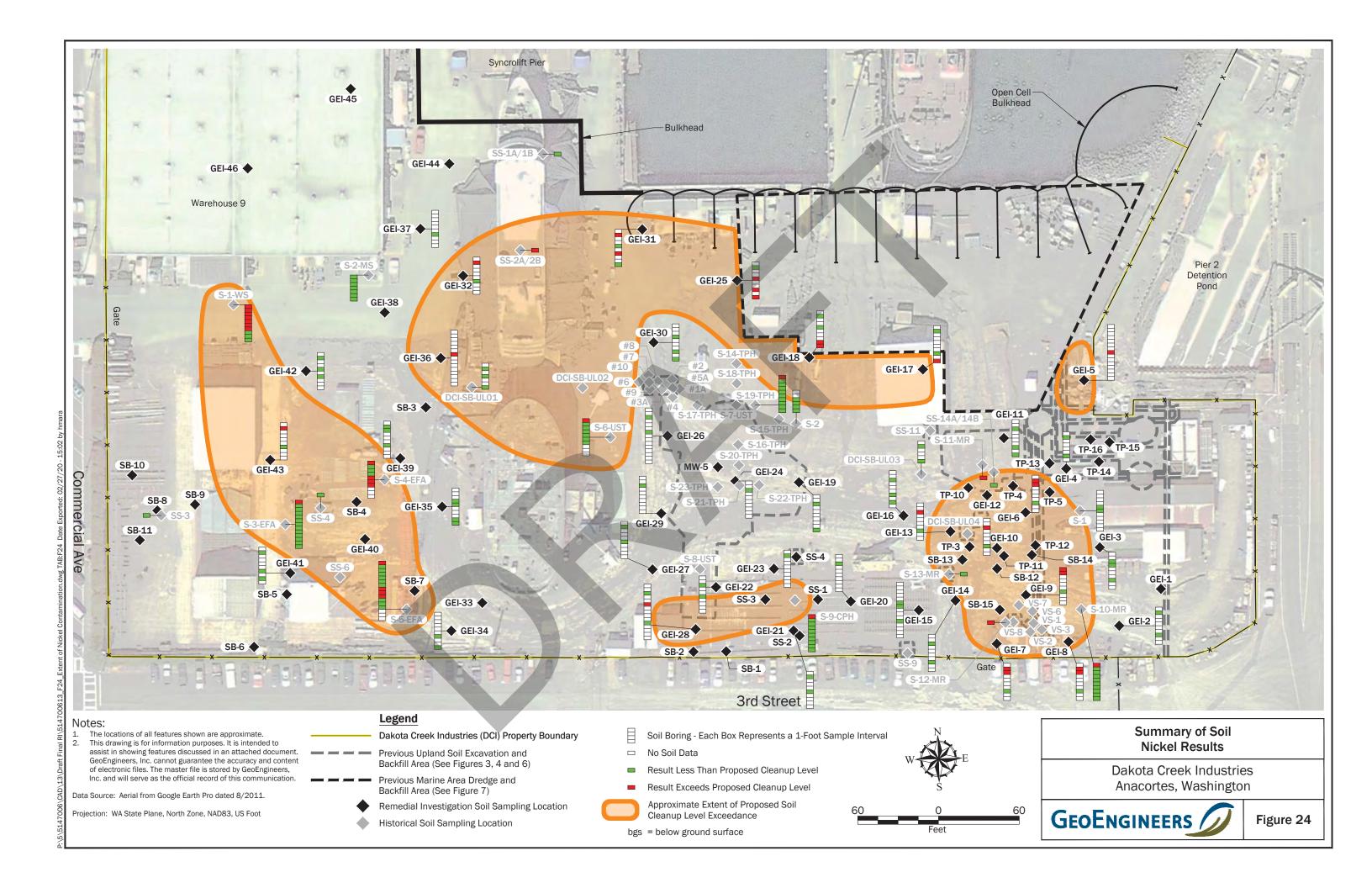


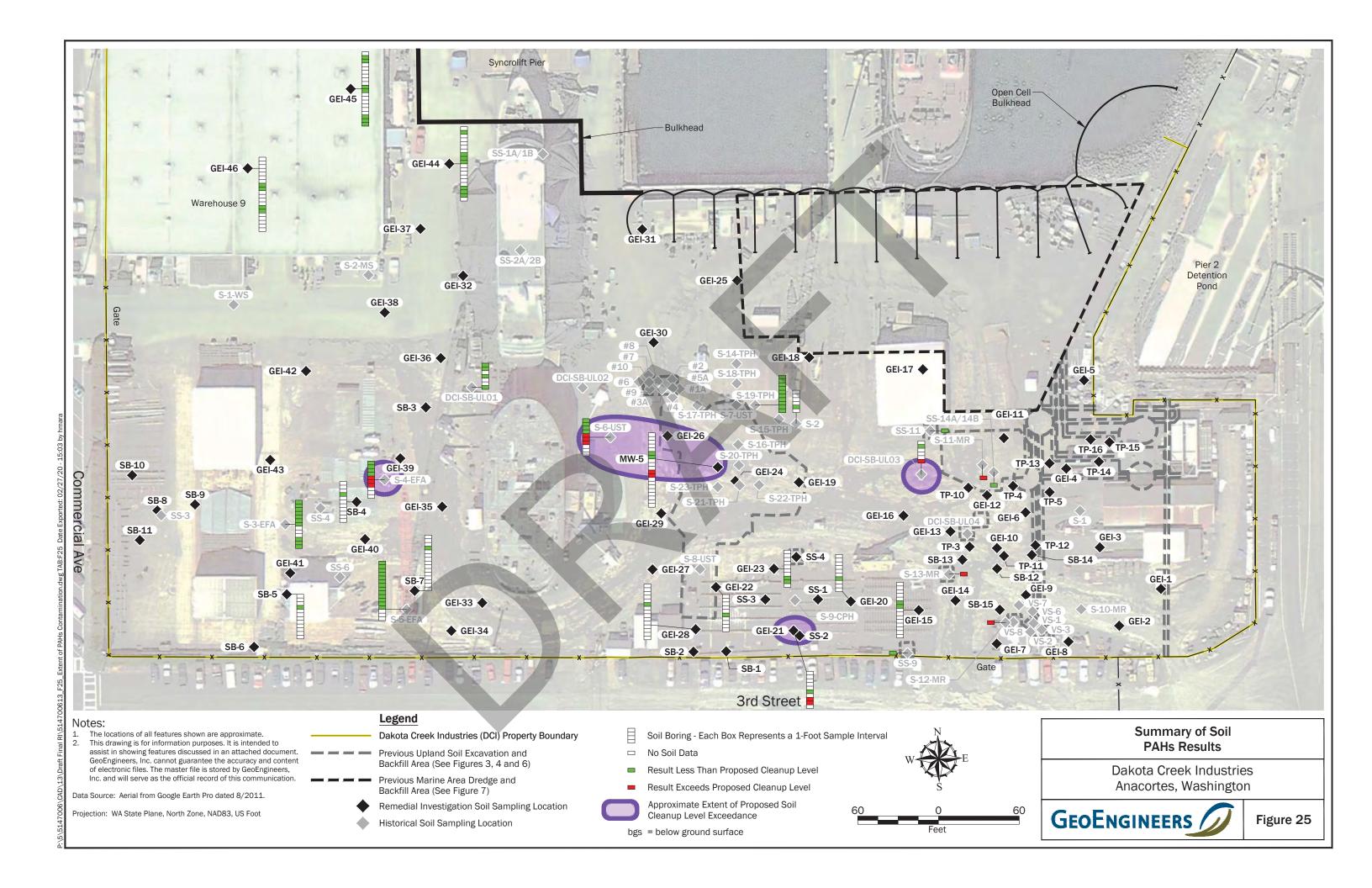


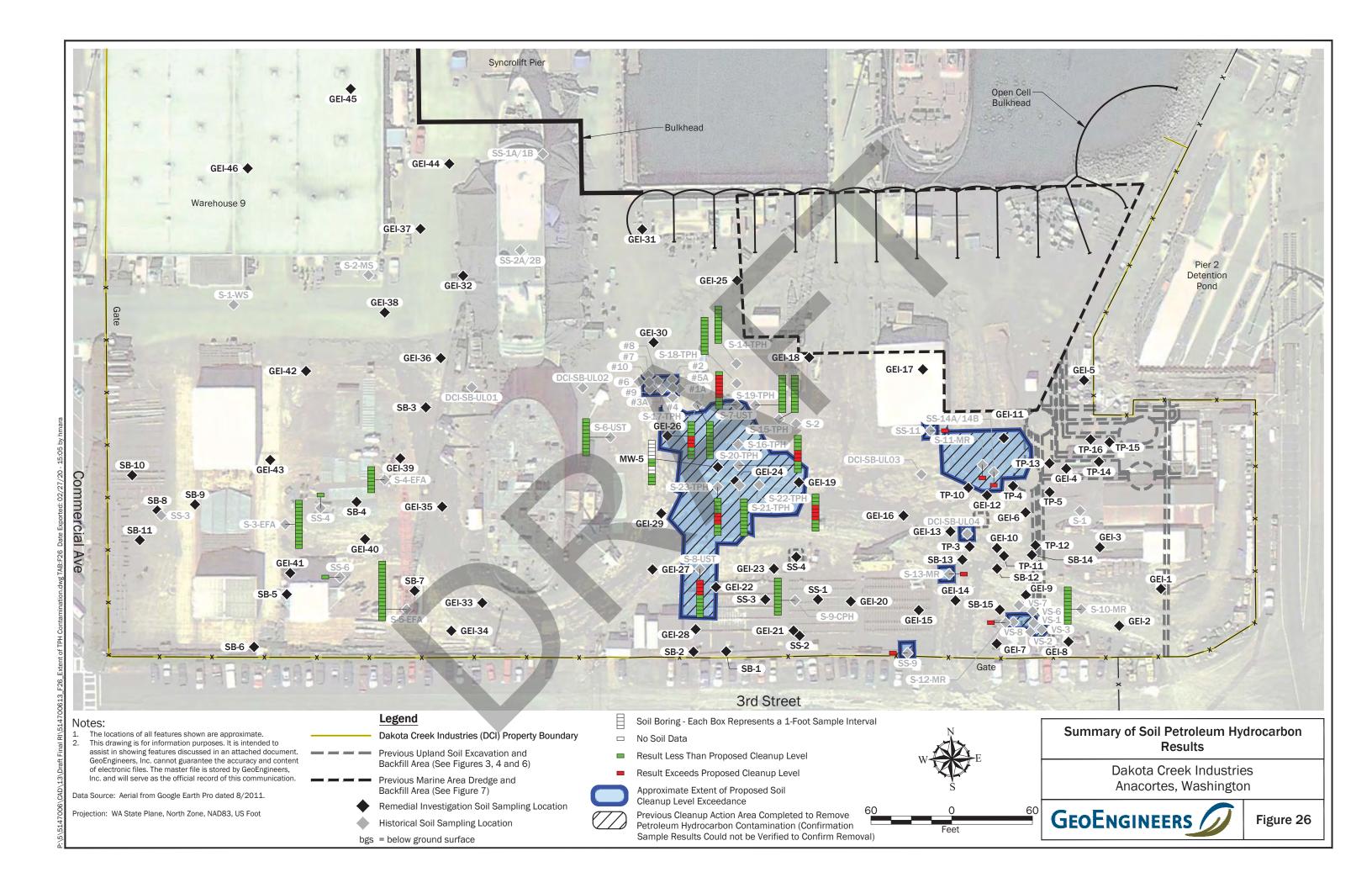












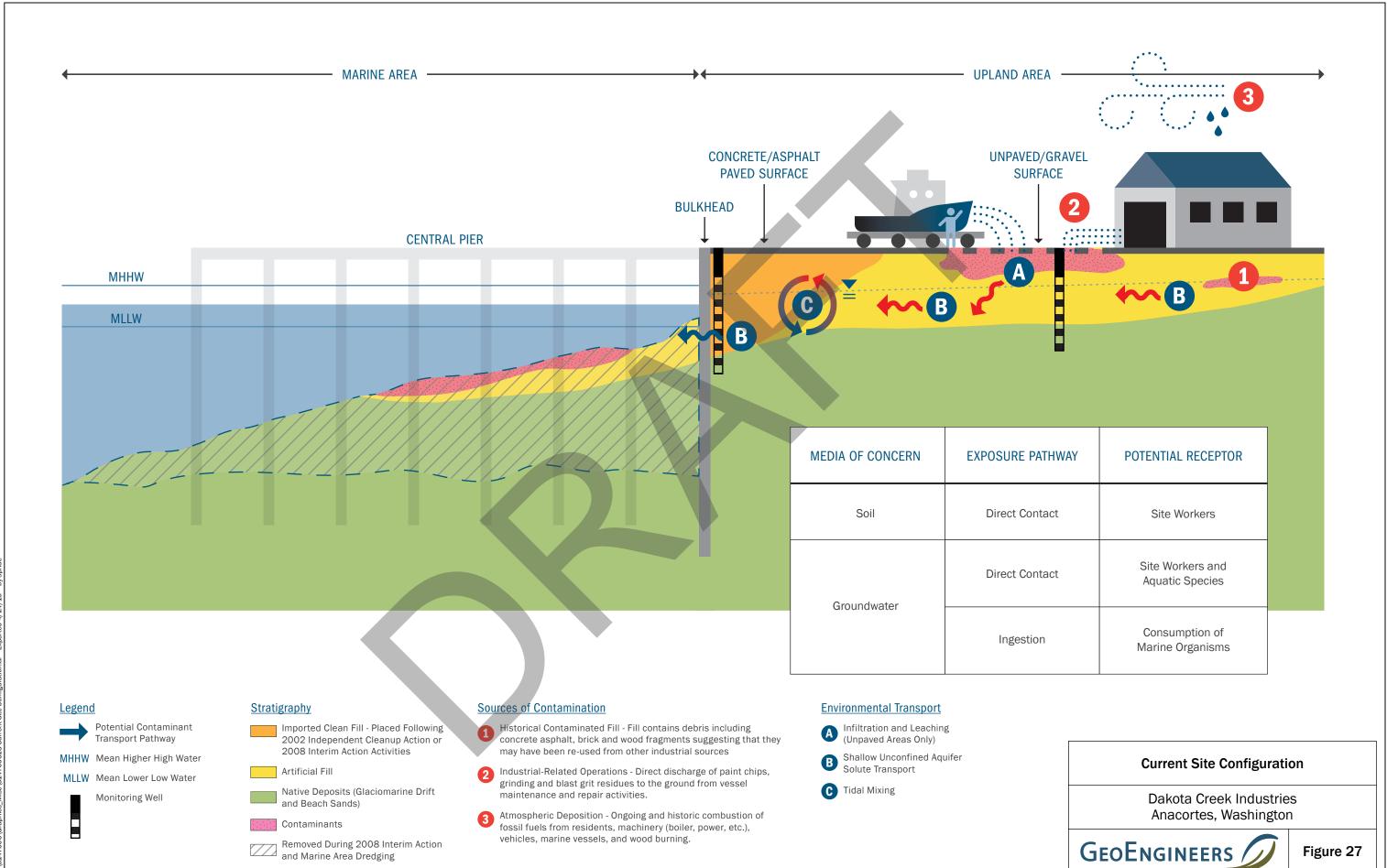


Figure 27

and Marine Area Dredging

