FOCUSED FEASIBILITY STUDY

Olympic Water & Sewer, Inc. Site 781 Walker Way Port Ludlow, Washington VCP Identification No. SW1311

Prepared for: Olympic Property Group and Pope Resources, LP

Project No. 130046-001-02 • September 24, 2013 Agency Review Draft





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Aspect Consulting, LLC

Brett Carp Sr. Environmental Scientist bcarp@aspectconsulting.com

earth + water

Eric J. Marhofer, PE Sr. Project Environmental Engineer emarhofer@aspectconsulting.com William V. Goodhue, LHG Sr. Associate Hydrogeologist cgoodhue@aspectconsulting.com

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Acronyms

AGI	Applied Geotechnology, Inc.
Aspect	Aspect Consulting, LLC
bgs	below ground surface
BTEX	benzene, toluene, ethylbenzene, and xylenes
COC	constituents of concern
DCA	disproportionate cost analysis
DPE	dual-phase extraction
Ecology	Washington State Department of Ecology
FS	feasibility study
FFS	Focused Feasibility Study
mg/m ³	milligrams per cubic meter
mg/kg	milligrams per kilogram
$\mu g/L$	micrograms per liter
MNA	monitored natural attenuation
MTCA	Model Toxics Control Act
NAPL	nonaqueous-phase liquid
NFA	No Further Action
NPV	net present value
OPG	Olympic Property Group and Pope Resources, LP
OWSI	Olympic Water & Sewer, Inc. Site
PID	photoionization detector
RAOs	Remedial action objectives
RI	Remedial Investigation
SLR	SLR International Corporation
SVE	soil vapor extraction

TEE	Terrestrial Ecological Evaluation
TPH	total petroleum hydrocarbons
UST	underground storage tank
VCP	Voluntary Cleanup Program

Washington Administrative Code

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1 Introduction

Aspect Consulting, LLC (Aspect) has prepared this Focused Feasibility Study (FFS) on behalf of Olympic Property Group and Pope Resources, LP (OPG) for the Olympic Water & Sewer, Inc. (OWSI) property located at 781 Walker Way in Port Ludlow, Washington (herein referred to as the Site) (Figure 1). This FFS incorporates the results of the previous investigations, the completed soil cleanup action, and soil vapor extraction (SVE) and SVE with groundwater pumping (DPE) pilot test activities completed by others, identifies and evaluates technically feasible cleanup action alternatives, and provides the basis for recommendation of the preferred final cleanup action for the Site. This FFS has been prepared in accordance with the Washington State Model Toxics Control Act Cleanup Regulation (MTCA), as established in Chapter 173-340 of the Washington Administrative Code (WAC) 173-340. The following subsections present the purpose and objectives of the FFS as well as an overview of this report organization.

1.1 Purpose and Objectives

Previous subsurface investigations and impacted soil excavation activities completed by others confirmed the release of gasoline-range total petroleum hydrocarbons to soil and groundwater at the Site from three former gasoline underground storage tanks (USTs) that were permanently decommissioned by removal in September 1990. Collectively, the completed UST decommissioning and soil removal activities, subsurface investigations, SVE and DPE pilot testing, and terrestrial ecological evaluation (TEE) meet the remedial investigation (RI) requirements of WAC 173-340-350. Details regarding the RI activities completed at the Site are provided in the following documents and are referenced throughout this report:

- Hydrocarbon Contamination Assessment and Underground Storage Tank Removal, Port Ludlow Water District, Port Ludlow, Washington, dated March 4, 1991, prepared by Applied Geotechnology, Inc. (AGI) (UST Removal Report; AGI, 1991).
- Well 17 Site Contamination, Initial Findings, and Recommendations, dated April 26, 2009, prepared by Robinson Noble & Saltbush, Inc. (Robinson Noble) (Initial Findings Report; Robinson Noble, 2009);
- Site Characterization Report, Olympic Water & Sewer, Inc. Property, 781 Walker Way, Port Ludlow, Washington, dated December 17, 2010, prepared by SLR International Corporation (SLR) (Site Characterization Report; SLR, 2010);
- Additional Investigation Report, Olympic Water & Sewer, Inc. Property, 781 Walker Way, Port Ludlow, Washington, dated August 2, 2011, prepared by SLR (Additional Investigation Report; SLR, 2011); and

• Soil Vapor Extraction Pilot Test Report, Olympic Water & Sewer, Inc. Facility, 781 Walker Way, Port Ludlow, Washington, dated May 8, 2012, prepared by SLR (SVE Pilot Test Report; SLR, 2012).

Based on the results of the RI activities, the Site has been sufficiently characterized to support the development and evaluation of technically feasible cleanup alternatives in accordance with WAC 173-340-360 through 173-340-390.

A traditional feasibility study (FS) typically includes an extensive development, screening, and evaluation process for numerous remedial alternatives. However, given the high frequency of sites with gasoline-impacted soil and groundwater, the range of applicable and effective remedial technologies is relatively well defined. In addition, Site-specific conditions preclude many potential remediation alternatives from application at the Site. Therefore, an FFS is considered sufficient for this Site.

The specific objectives of this FFS are to:

- Provide a summary of completed remedial investigation, cleanup activities and current site conditions, and present a concise Site conceptual model.
- Present a detailed analysis and feasibility evaluation of the completed SVE/groundwater pumping pilot test;
- Present the results of the completed soil vapor pathway evaluation;
- Identify and evaluate technically feasible cleanup action alternatives; and
- Present a recommendation for a final cleanup action for the Site in accordance with WAC 173-340-350(8).

The final cleanup action will be conducted independently under the Washington State Department of Ecology (Ecology) Voluntary Cleanup Program (VCP) with the objective to obtain a No Further Action (NFA) determination for the Site. The Site has been enrolled in the Ecology VCP and has been assigned VCP Identification No. SW1311.

1.2 Organization

This report has been organized into the following sections:

- Section 2 Summary of Site Conditions provides a summary of Site conditions including location and description, environmental setting, and geology and hydrogeology. This section also details recent soil vapor sampling work completed by Aspect.
- Section 3 Conceptual Site Model provides a summary of the conceptual site model including a discussion of the constituents of concern (COCs), affected media, sources and inferred extent of soil and groundwater impacts, potential receptors and exposure assessment, and contaminant fate and transport mechanisms for the Site. This section also presents the simplified Terrestrial Ecological Evaluation (TEE) completed for the Site.
- Section 4 Basis for Additional Remedial Action presents the objectives and standards by which evaluation of additional remedial action(s), beyond those already completed at the Site, will be measured.

- Section 5 Focused Feasibility Study presents a screening of potential remedial technologies, the retained cleanup action alternatives, and an evaluation of those alternatives.
- Section 6 Summary of Preferred Cleanup Action provides a summary of the recommended cleanup action alternative for implementation at the Site.
- Section 7 References provides a list of the source materials referenced in this report.

2 Summary of Site Conditions

This section provides a summary of Site conditions including Site location and description, environmental setting, geology and hydrogeology. It also details the recent soil vapor sampling work completed by Aspect.

2.1 Site Location and Description

The Site is located in Section 8, Township 28 North, Range 1 East in Port Ludlow, Washington (Figure 1). Identified as Jefferson County Parcel No. 821084004, the Site consists of an approximately 2.2-acre parcel of land located approximately ½-mile northwest of the Port Ludlow bay. The Site is located at the southwest corner of the intersection of Walker Way and Rainer Lane (Figure 2). Properties adjacent to the south, west, and east, beyond Rainier Lane are developed with single-family residences (Jefferson County, 2013). Properties to the north, across Walker Way, are developed with a mini-storage facility and single-family residences.

The parcel is partially developed with an OWSI operations and maintenance facility, which consists of an approximate ¹/₂-acre area that includes an office/shop/garage building (garage building), a public water supply well (i.e., Well #2) and associated pump house building, and a storage trailer (Figure 2). The ground surface within the developed portion of the Site is primarily unpaved, except for a narrow asphalt driveway that runs down the center of the OWSI facility from Walker Way to approximately the storage trailer. Areas surrounding the facility are undeveloped and covered with dense vegetation. The OWSI facility has been in operation since first development in 1968, following the installation of the water supply Well #2 (Figure 2) (SLR, 2011). Additional details regarding water supply Well #2 are provided in the following sections of this report.

2.2 Environmental Setting

This subsection provides a summary of the environmental setting of the Site. The information presented here has been obtained from a review of national, state, and local records and previous environmental work completed at the Site by others.

2.2.1 Land-Use

According to Jefferson County Assessor's Office, the property land use code is 4800–Utilities, non-public (Jefferson County, 2013). The land use code for properties adjacent to the south and west is 1100–Houses, single units, non-farm. According to the Jefferson County Assessor's Office, properties adjacent to the south, west, and east, beyond Rainier Lane are zoned MPR-Single Family. Properties to the north, across Walker Way, are zoned Rural Residential. However, the Site is used for commercial purposes by OWSI as an operations and maintenance facility, including water supply Well #2. The current and future land use for the Site is and will likely remain commercial.

2.2.2 Topography

The ground surface elevation proximate to the northern property boundary of the Site is approximately 290 feet above mean sea level (Google Earth, 2013). The ground surface of the OWSI facility slopes gently toward the southwest (Figure 2). Areas surrounding the facility are undeveloped and covered with dense vegetation. A densely vegetated gulley, containing an intermittent seasonal stream, is located to the west of the OWSI facility.

2.3 Geology, Hydrogeology, and Groundwater Use

2.3.1 Geology

Based on the results of multiple investigations completed to date at the Site, Site soils consist of dense glacial advance outwash (sand, gravel, and silt units) with interbedded lacustrine silts to the maximum depth drilled (approximately 60 feet [below ground surface] bgs). Specifically, thin surficial gravel fill is underlain by a sand (silty to gravelly) to gravel unit that is approximately 29 to 43 feet thick. Beneath the central and southern parts of the property, a 5- to 10-foot-thick sandy silt to silt is interbedded within the sand to gravel unit. The sand to gravel unit is underlain by clayey to gravelly silt that is 15 to more than 23 feet thick. At the northern and central parts of the OWSI facility, the clayey to gravelly silt unit is overlain by a silty sand that is up to 11 feet thick. In the central part of the OWSI facility (at MW-1 and MW-2), the top of the clayey to gravelly silt occurs at an elevation of approximately 251 feet above the NAVD 88 datum, while at the northern, southern, and eastern parts of the facility (at MW-4, MW-3, and MW-5, respectively), the clavey to gravelly silt occurs at higher elevations (approximately 260 to 263 feet above the NAVD 88 datum). At MW-2, MW-3, and MW-4, a gravelly sand to sand and gravel that is approximately 5 to 7.5 feet thick is interbedded within the clavey to gravelly silt unit (SLR, 2010). At MW-1, the clayey to gravelly silt unit is underlain by silty sand that extends beyond the bottom of the boring. According to the driller's log for the water supply well (Well #2) located in the northern part of the property, a thick sequence of clay and cemented sand occurs from approximately 49 to 215 feet bgs (SLR, 2010).

2.3.2 Hydrogeology

Shallow groundwater at the Site occurs under perched conditions within the glacial advance outwash and lacustrine deposits at depths above approximately 60 bgs. Deeper regional water-bearing units are present beneath a thick aquitard comprised of clay and cemented silty sand. These deep water-bearing units at Well #2 occur at depths of between 215 and 245 feet bgs, or over at least 150 feet below the top of the aquitard and base of the perched units.

In early April 2011, the depths to perched groundwater in the Site monitoring wells and SVE points ranged from 19.80 to 36.98 feet below the tops of the well casings, and the groundwater elevations ranged from 256.89 to 275.85 feet above the NAVD 88 datum (Table 1). At wells MW-1 through MW-4, the groundwater elevations in April 2011 were 3.32 to 5.68 feet higher than in October 2010, and from June 2010 to April 2011, the groundwater fluctuations in the wells ranged from 3.81 to 5.68 feet. The higher

groundwater elevations in April 2011 likely reflect seasonal recharge from infiltration of precipitation during the autumn and winter months.

Shallow groundwater beneath the Site occurs within the sand to gravel unit, and is perched on top of the underlying clayey to gravelly silt unit (see Figure 5 of the Additional Investigation Report; SLR, 2011). During periods of seasonal recharge, groundwater appears to collect above the silt and overlying silty sand units. In areas where the silty sands and silts are present at higher elevations, the groundwater elevations are higher. For example, groundwater elevations were 266.35 feet at MW-3, 273.19 feet at SVE-1, 273.38 at MW-4, and 274.07 feet at SVE-2 (see Figure 4 of the Additional Investigation Report; SLR, 2011). This groundwater is hydraulically continuous with the deeper perched groundwater intercepted by wells MW-1 and MW-2. The horizontal hydraulic conductivity of the sand to gravel unit is expected to be significantly (i.e., orders of magnitude) greater than the vertical hydraulic conductivity of the underlying silt and silty sand. Therefore, groundwater accumulating above the 265-foot elevation is expected to primarily flow laterally toward the gravels encountered at MW-2, or toward the intermittent stream in the gulley where stream sampling was completed in April 2011.

The groundwater flow within the perched zone appears to be controlled by the geometry of the clayey to gravelly silt, with flow converging into the low point of the top of the silt unit (SLR, 2010). As described above, the elevation of the silt unit is about 10 feet lower in the central part of the OWSI facility than at the northern, southern, and eastern parts of the facility. This interpretation is consistent with the high petroleum hydrocarbon concentrations occurring in the groundwater at wells MW-1 and MW-2 (SLR, 2011). Based on the known clayey to gravelly silt geometry and the area of petroleum hydrocarbon-impacted groundwater, there appears to be a flow component to the south-southwest (SLR, 2011). Perched groundwater appears to discharge to the intermittent stream at locations near the southern end of the property.

2.3.3 Groundwater Use

Well records obtained by SLR from Ecology and OWSI identified 12 water supply wells located within a ¹/₂-mile radius of the property (SLR, 2010). Approximate locations for the water supply wells are shown on Figure 7 of the Site Characterization Report (SLR, 2010). Copies of the water supply well completion logs and a table prepared by SLR that presents the well completion details are included as Appendix A. According to the well records, groundwater from the water supply wells, including Well #2 located at the Site, is used for domestic purposes. Shallow perched groundwater at the Site is not used for drinking purposes (SLR, 2011).

Water supply Well #2 is screened at depths ranging from 214 to 245 feet bgs. All of the identified water supply wells in the vicinity of the Site are completed (i.e., screened or open casing) at depths ranging between 157 and 377 feet bgs. The soil descriptions on the well logs consistently note that a thick sequence of clay and cemented silty sand aquitard units occur above the deep groundwater-bearing zones. Groundwater flow directions in the deeper regional aquifer have been inferred to flow from the upland areas toward Port Ludlow, indicating that the water supply wells are located hydraulically up- or cross-gradient of the Site (EES, 1994).

Based on the presence of the thick aquitard and the inferred deep groundwater flow direction, shallow impacted perched groundwater beneath the Site is not considered a risk to water quality in the deep groundwater-bearing zones (SLR, 2010). The lack of detectable petroleum hydrocarbons in water samples collected from Well #2 in the 1990, 2009, and 2010 further supports this conclusion (SLR, 2010).

2.4 Soil Vapor Pathway Evaluation

Aspect completed an evaluation of the soil vapor pathway at the Site in June 2013. The purpose of the evaluation was to assess if concentrations of gasoline constituents exceeding the screening levels provided in Ecology's *Draft Guidance for Evaluating Soil Vapor Intrusion in Washington State* (Ecology, 2009) were present in soil vapor beneath the slab of the garage building. Aspect installed soil vapor points SV-1 and SV-2 at the Site on June 20, 2013 (Figure 3). Soil vapor samples were collected from vapor points VP-1 and VP-2 on June 21, 2013, in accordance with the Standard Operating Procedure provided in Appendix B.

As noted in Section 2.2.1, the current and future land use for the Site is and will likely remain commercial. The vapor sample analytical results were therefore evaluated for a commercial land use scenario (Table 2). No exceedances of screening levels were recorded in soil vapor, and therefore no further evaluation of the soil vapor pathway is considered warranted given the current and future Site land use. A copy of the soil vapor laboratory analytical report is provided in Appendix C.

2.5 Sufficiency of Remedial Investigation Activities

As previously noted in Section 1.1, the results of the UST decommissioning activities, completed soil cleanup action, subsurface investigations, and pilot test activities completed at the Site constitute a complete RI in general accordance with WAC 173-340-350. The complete RI activities are considered sufficient to support the development and evaluation of technically feasible cleanup alternatives in accordance with WAC 173-340-360 through 173-340-390. Details regarding the RI activities completed at the Site are provided in documents referenced in Section 1.1. The conceptual site model, based on the results of the RI activities completed by others and the soil vapor pathway evaluation recently completed by Aspect, is presented in the following section.

3 Conceptual Site Model

This section provides a summary of the conceptual site model including a discussion of the COCs, affected media, sources and inferred extent of soil and groundwater impacts, potential receptors and exposure assessment, and contaminant fate and transport mechanisms for the Site. This section also presents the simplified TEE completed for the Site.

3.1 Constituents of Concern

The COCs identified for the Site are based on the historical use of gasoline USTs at the Site and the results of the RI activities. Based on the available data, the following COCs have been identified for the Site:

- Gasoline-range total petroleum hydrocarbons (TPH); and
- The gasoline constituents benzene, toluene, ethyl-benzene, and total xylenes (BTEX).

3.2 Affected Media

Concentrations of one or more of the COCs have been confirmed in soil and/or groundwater at the Site. Therefore, soil and groundwater have been identified as affected media of concern for the Site. Based on the lack of detectable concentrations of COCs in surface water samples collected from the intermittent seasonal stream located west of the OWSI facility, surface water will not be retained as a media of concern (SLR, 2011). In addition, based on the results of the soil vapor evaluation recently completed by Aspect, indoor air will not be retained as a media of concern. Potential receptors and exposure pathways are summarized in Section 3.4.

3.3 Sources and Extent of Impacts

A source area is the location where a release has occurred at the Site. Based on the available data, the sources of the COCs in the affected media are the 1,000-gallon UST formerly located beneath the floor of the garage building and the 2,000-gallon UST formerly located along the west side of the garage building (SLR, 2011). As previously noted, these USTs were permanently decommissioned by removal in September 1990. Given the similar contents of these former USTs (i.e., gasoline) and the proximity of these two source areas relative to each other, the two source areas will be treated as a single source area for the purposes of evaluating technically feasible remedial alternatives.

The extent of soil and groundwater impacts at the Site is identified as areas where COCs in the affected media have come to be located. A description of the extent of soil and groundwater impacts at the Site is presented below.

3.3.1 Soil

Following removal of the USTs, a cleanup action consisting of excavation of gasolineimpacted soil was completed to the extent practicable in 1990 (SLR, 2011). To prevent structural damage to the garage building, residual gasoline-impacted soil was left in-place at the base of the 1,000-gallon UST excavation (SLR, 2011). Residual gasoline-impacted soil at that location is expected to occur from below approximately 10 feet bgs to the perched groundwater table between approximately 20 to 41 feet bgs (SLR, 2011).

Concentrations of gasoline-range TPH exceeding the MTCA Method A cleanup level of 30 milligrams per kilogram (mg/kg) were detected in soil samples collected at depths greater than 20 feet bgs at SVE-2 and MW-1B (Table 3). Similarly, concentrations of benzene exceeding the MTCA Method A cleanup level of 0.03mg/kg were also detected in the soil sample collected at 24.5 to 25 feet bgs at MW-1B (Table 3).

Based on the results of the RI activities, the area of impacted soil is estimated to extend beyond the western, eastern, and southern ends of the garage building and covers an area of approximately 3,140 square feet (see Figure 7 of the Additional Investigation Report; SLR, 2011). The extent of soil impacts at the Site has been sufficiently characterized to support the development and evaluation of technically feasible cleanup alternatives in accordance with WAC 173-340-360 through 173-340-390.

3.3.2 Groundwater

Concentrations of gasoline-range TPH exceeding the MTCA Method A cleanup level of 800 micrograms per liter (μ g/L) have been detected during multiple sampling events at MW-1, MW-2, SVE-1 and SVE-2 (Table 4). In addition, concentrations of one or more BTEX constituents exceeding MTCA cleanup levels have historically been detected in shallow perched groundwater samples collected at MW-1, MW-2, MW-4, and SVE-1.

The estimated area of the hydrocarbon-impacted shallow perched groundwater is depicted on Figure 4 of the Additional Investigation Report (SLR, 2011). The impacted groundwater is inferred to extend beyond the western fence line of the OWSI facility, but not as far west as the intermittent stream. The area west of the fence line to the intermittent stream is inaccessible.

The groundwater flow within the shallow perched zone appears to be controlled by the geometry of the clayey to gravelly silt, with flow converging into the low point of the top of the silt unit (SLR, 2010). Based on the known clayey to gravelly silt geometry and the area of petroleum hydrocarbon-impacted groundwater, there appears to be a flow component to the south-southwest (SLR, 2011). It is likely that the perched groundwater discharges to the intermittent stream, at locations near the southern end of the property, during periods of seasonal recharge. In addition, based on the presence of petroleum hydrocarbons at MW-4 and MW-5, there is a limited component of impacted groundwater migration, likely seasonally, to the north and east (SLR, 2011).

As discussed in Section 2.3.3, based on the presence of the thick clay and cemented silty sand aquitard between the perched and regional water-bearing units, and the inferred deep groundwater flow direction, it is unlikely that hydrocarbon-impacted shallow perched groundwater beneath the property could affect water quality in the deep groundwater-bearing zones (SLR, 2010). This conclusion is supported by the lack of

detectable petroleum hydrocarbons in water samples collected from Well #2 in the 1990, 2009, and 2010 (SLR, 2010). The extent of shallow groundwater impacts at the Site has been sufficiently characterized to support the development and evaluation of technically feasible cleanup alternatives in accordance with WAC 173-340-360 through 173-340-390.

3.4 Potential Receptors and Exposure Assessment

The two primary exposures associated with the presence of COCs at the Site are human health and terrestrial ecological risk. The nature and extent of concentrations of COCs in soil and groundwater determines the potential exposure scenarios for human health and terrestrial ecological effects.

Potential exposure pathways that may affect human health include soil, groundwater, surface water, and vapor intrusion. The following subsections present a description of each potential exposure pathway.

3.4.1 Soil Exposure Pathways

Two potential soil exposure pathways, direct-contact and soil-leaching to groundwater, have been identified for the Site. A discussion of each of the soil exposure pathways is presented below:

- **Direct-contact pathway:** The direct-contact pathway considers both dermal contact with and ingestion of soil from beneath the Site, to a maximum depth of 15 feet bgs. As previously noted, following removal of the USTs in 1990, an cleanup action consisting of excavation of gasoline-impacted soil was completed to the extent practicable in 1990 (SLR, 2011). However, to prevent structural damage to the garage building, residual gasoline-impacted soil was left in-place beneath the former location of the 1,000-gallon UST (SLR, 2011). Residual gasoline-impacted soil at that location is expected to occur from below approximately 10 feet bgs to the perched groundwater table at approximately 20 to 41 feet bgs (SLR, 2011). Although direct-contact with this shallow soil is considered unlikely, this exposure pathway will be considered during evaluation of potential remedial technologies and development of potential cleanup action alternatives.
- Soil-leaching to groundwater pathway: The soil-leaching to groundwater pathway requires consideration of the highest beneficial use of groundwater at the Site in accordance with WAC 173-340-357(3)(d). As described in Section 2.3.3, Ecology and OWSI well records identified 12 water supply wells located within a ½-mile radius of the property (see Figure 7 of the Site Characterization Report; SLR, 2010). However, given the geology and hydrogeology of the Site (i.e., the presence of the thick clay and cemented silty sand aquitard and the inferred deep groundwater flow direction), it is considered highly unlikely that residual concentrations of COCs in soil could affect water quality in the deep groundwater-bearing zones. Only the soil-leaching to the shallow perched groundwater-bearing zone pathway will be considered during evaluation of potential remedial technologies and development of potential cleanup action alternatives.

3.4.2 Shallow Perched Groundwater Exposure Pathway

This pathway includes ingestion of and dermal contact with groundwater from the shallow perched aquifer at the Site. As described in Section 2.3.3, Ecology and OWSI well records identified 12 water supply wells located within a ½-mile radius of the property (See Figure 7 of the Site Characterization Report; SLR, 2010). Although the shallow perched groundwater has been impacted by COCs, it is not used for drinking purposes (SLR, 2011). In addition, given the geology and hydrogeology of the Site (i.e., the presence of a thick clay and cemented sand aquitard), it is considered unlikely that the COCs could migrate from the shallow perched aquifer to the deeper regional aquifer.

Given that the existing and any potential future water supply wells at the Site target production zones at depths ranging from 215 to 245 feet bgs, it is unlikely that the shallow perched groundwater would be used for drinking water purposes in the foreseeable future (SLR, 2011). However, for the purpose of this FFS, it is assumed that the perched groundwater may be used for drinking water purposes by future residents. Therefore, exposure via ingestion and dermal contact with groundwater from the shallow perched aquifer will be considered during evaluation of potential remedial technologies and development of potential cleanup action alternatives.

3.4.3 Surface Water Exposure Pathway

Shallow perched groundwater appears to migrate southwest towards the gulley with an intermittent stream (SLR, 2011). However, concentrations of COCs have not been detected in water samples collected from the stream. Since the shallow groundwater likely discharges to the stream during periods of seasonal recharge, it appears that natural attenuation processes are reducing the hydrocarbon concentrations before groundwater discharges to the stream. The lack of detectable hydrocarbon concentrations in the stream suggests that it is unlikely that human receptors, terrestrial receptors, or aquatic organisms could have significant exposure to COCs present in the shallow groundwater. As a result, this exposure pathway is considered incomplete.

3.4.4 Vapor Intrusion Pathway

As noted in Section 2.4, an evaluation of the soil vapor pathway was completed in June 2013. Based on the results of the evaluation, no further evaluation of this pathway is warranted given the current and expected future Site land use.

3.5 Contaminant Fate and Transport

Released gasoline constituents can exist in the environment in four different phases: adsorbed to soil, dissolved in water, as vapors, and as separate nonaqueous-phase liquid (NAPL) or residual product. To date, no evidence of NAPL has been detected in the subsurface at the Site (SLR, 2011). The primary processes influencing transport of petroleum constituents in the subsurface include:

- Migration as NAPL both vertically due to gravity and laterally along less permeable soil;
- Leaching from soil to groundwater;
- Volatilization from soil or groundwater to air;
- Advection and dispersion in groundwater; and
- Natural degradation.

These potential fate and transport processes are further discussed below.

After a release from a UST, NAPL flows into the shallow soils near the tank. After saturating the soil, a portion of the NAPL can migrate downward and laterally through the vadose zone, and may reach the groundwater table, where it would float on the fluctuating groundwater. As the groundwater table rises and falls, a smear zone of residual hydrocarbons can form in the soil within the zone of groundwater fluctuation. A portion of the product can migrate with groundwater flow and based on solubility, the product also dissolves in the groundwater. However, there is no evidence that NAPL has migrated to the groundwater beneath the Site, and there is no current evidence of NAPL in the soil (SLR, 2011).

Much of the developed portion of the OWSI property is covered with gravel, and as rain falls on the ground surface and infiltrates into the subsurface, residual COCs in soil can dissolve in the water and percolate through the soils. Some of the COCs eventually reach the groundwater. Partitioning from soil to water is determined, in part, by the solubility of a particular hydrocarbon. Once dissolved in groundwater, COCs may be transported by diffusion and advection away from the source area.

Horizontal migration with groundwater (i.e., advection) is expected to be significantly more extensive at the OWSI property than vertical migration (SLR, 2011). The top of the impacted perched groundwater occurs at depths between approximately 20 to 41 feet bgs.

Dispersion, retardation, and biodegradation act to reduce the dissolved petroleum hydrocarbon concentrations as groundwater migrates from the source area. A growing body of evidence suggests that in most systems, biodecay is a significant loss mechanism for many petroleum constituents such as benzene. The intermittent stream is located over 200 feet southwest of the contaminant source area and based on topography and water levels, the perched groundwater discharges to the stream during periods of seasonal recharge. Based on the lack of detectable petroleum hydrocarbons in the stream sample, it appears that the concentrations in the groundwater have degraded sufficiently prior to discharge such that petroleum hydrocarbons are not detectable in the surface water (SLR, 2011).

The primary beneficial use aquifer in the vicinity of the Site occurs at depths of approximately 215 to 245 feet bgs beneath the Site. Given the geology and hydrogeology of the Site (i.e., the presence of the clay and cemented silty sand aquitard and the inferred deep groundwater flow direction), it is considered highly unlikely that residual concentrations of COCs in soil could affect water quality in the deep groundwater-bearing zones. None of the data collected during the RI suggests that COCs have or will migrate to deeper groundwater that is currently being used as a drinking water source.

3.6 Terrestrial Ecological Evaluation

A TEE is intended to assess potential risk to terrestrial plants and/or animals that live entirely or primarily on affected land. A simplified TEE is required under MTCA to assess potential ecological risk posed by the COC at the Site and to determine whether a more detailed investigation of potential ecological risk is required. Aspect completed a preliminary TEE for the Site in accordance with WAC 173-340-7491. A copy of the completed Ecology VCP TEE Form is provided as Appendix D. The Site qualifies for a TEE exclusion under WAC 173-340-7491(1)(a) (*All soil contamination is, or will be, at least 6 feet below the surface (or alternative depth if approved by Ecology), and institutional controls are used to manage remaining contamination.*). No further evaluation of potential threats to terrestrial plants or animals from the Site is considered warranted.

4 Basis for Additional Remedial Action

As previously noted, following removal of the USTs in 1990, a cleanup action consisting of excavation of gasoline-impacted soil was completed to the extent practicable in 1990 (SLR, 2011). This section presents the objectives and standards by which evaluation of additional remedial action(s), beyond those already completed at the Site, will be measured.

4.1 Remedial Action Objectives

Remedial action objectives (RAOs) established for the Site are intended to comply with applicable environmental regulations and protect human health and the environment. The Site-specific RAOs include the following:

- Protection from direct-contact and ingestion of petroleum-impacted soil;
- Protection from direct-contact and ingestion of petroleum-impacted shallow perched groundwater;
- Protection of drinking water in the deep groundwater-bearing zone; and
- Protection of surface water for beneficial use.

4.2 Cleanup Standards

As defined in WAC 173-340-700, cleanup standards for the Site include establishing cleanup levels and points of compliance at which those cleanup levels will be attained. The following presents a discussion of the preliminary cleanup levels and points of compliance for the Site.

4.2.1 Preliminary Cleanup Levels

Based on the exposure pathways described above (i.e., dermal contact with and/or ingestion of soil and/or shallow perched groundwater), recommended cleanup levels for the Site are MTCA Method A cleanup levels for soil and groundwater. Evaluation of additional remedial action(s), beyond the remedial actions completed to date, will address achievement of these recommended cleanup levels.

4.2.2 Points of Compliance

The points of compliance are defined in WAC 173-340-200 as the locations where cleanup levels established in accordance with WAC 173-340-720 through WAC 173-340-760 will be attained to meet the requirements of MTCA. Once the cleanup levels have been attained at the defined points of compliance, the Site is no longer considered to be a threat to human health or the environment. Standard points of compliance which address potential receptors via the exposure pathways that are complete are presented below:

- Soil for protection from direct-contact: Ground surface to a depth of 15 feet bgs; and
- Shallow perched groundwater for protection of drinking water and surface water: Within the perched aquifer extending vertically from the uppermost level of the saturated zone to the lowest depth potentially affected.

If it is not practicable to meet cleanup levels at the standard points of compliance discussed above within a reasonable restoration time frame, a conditional point of compliance for soil and/or groundwater may be established. Final points of compliance for the Site will be subject to Ecology approval.

5 Focused Feasibility Study

This section presents a screening of potential remedial technologies, the retained cleanup action alternatives, and an evaluation of those alternatives.

The purpose of the FFS is to screen cleanup alternatives and eliminate those that are not technically feasible, those whose costs are clearly disproportionate under WAC 173-340-360(3)(e), or those that will substantially affect the ability of the existing tenant to utilize the Site. In addition, the purpose of the FFS is to evaluate the most-advantageous remediation technologies using bench- and pilot-scale testing, where applicable, to recommend a final cleanup action for the Site in conformance with WAC 173-340-360 through 173-340-390. This FFS is intended to provide sufficient information to enable selection of a final cleanup action. As previously noted, the final cleanup action will be conducted independently under the Ecology VCP with the objective to obtain an NFA determination for the Site.

5.1 Potential Remedial Technologies

Aspect identified and evaluated potential remediation technologies for the Site with respect to the cleanup requirements set forth in MTCA and the RAOs and Cleanup Standards presented in Section 4. Potential remedial technologies for addressing the residual soil and groundwater impacts at the Site include the following:

- **Institutional Controls:** Measures to limit or prohibit activities that may interfere with the integrity of a cleanup action or result in exposure to hazardous substances (i.e., limitations on the use of the property or resources such as an environmental covenant or maintenance requirements for engineering controls).
- Engineering Controls: Containment and/or treatment systems that are designed and constructed to prevent or limit the movement of, or the exposure to, hazardous substances (i.e., asphalt or concrete paving/capping).
- **Monitored Natural Attenuation (MNA):** Monitoring the removal of contaminants by natural processes (i.e., biodegradation).
- Soil Vapor Extraction: Extracting and treating contaminated soil vapor. Pilot testing of this technology was completed.
- **Air Sparging:** Injecting air into contaminated groundwater to volatilize contaminants. This technology is often implemented in conjunction with SVE.
- Enhanced Aerobic Biodegradation: Injecting an oxygen source and, if necessary, bacteria to stimulate microbial biodegradation of contaminants.
- **In Situ Chemical Oxidation:** Injecting or mixing an oxidant, such as potassium permanganate or sodium persulfate, into the soil which reacts with and destroys contaminants.
- **Dual-Phase Extraction (DPE):** Extracting and treating impacted groundwater and vapor. Pilot testing of this technology was completed.

• Soil Excavation: Removal of impacted soil, followed by off-site disposal. This technology was implemented to the extent practical for cleanup of impacted soils during UST removal.

Each of these potential remedial technologies has been applied at sites with similar conditions and chemical occurrences. Appendix E provides a general description of each technology and their general applicability to comparable sites.

5.1.1 Screening of Potential Remedial Technologies

Preliminary screening of the potential remedial technologies based on effectiveness, implementability, and comparative costs is shown in Table 5. The following potential remedial technologies were retained for development as potential cleanup action alternatives:

- Institutional and Engineering Controls
- Monitored Natural Attenuation
- In Situ Chemical Oxidation
- Soil Excavation

These remediation technologies which passed the initial screening were combined into remedial alternatives and further evaluated in Section 5.3.

The following potential remedial technologies were not retained for development as potential cleanup action alternatives:

- Air Sparging Air sparging was not retained for further development as a remedial alternative. The perched aquifer complexity and low aquifer permeability, coupled with the transient nature of perched groundwater, would make the implementability of air sparging and the recovery of sparged vapors problematic at the Site.
- Enhanced Aerobic Biodegradation Enhanced Aerobic Biodegradation was not retained for further development as a remedial alternative primarily because similar to air sparging, the perched aquifer complexity and low aquifer permeability are considered critical factors that would make this technology difficult to implement, and of likely limited effectiveness.
- Soil Vapor Extraction An SVE pilot test was conducted at the Site between December 2011 and January 2012 (SLR, 2012). An evaluation of SVE pilot testing performance is included in Section 5.1.2 below. The evaluation confirms that SVE is not a viable technology for remediation of impacted soil, and therefore SVE was not retained for consideration.
- **Dual-Phase Extraction** A DPE test was completed in conjunction with the SVE pilot test between December 2011 and January 2012 (SLR, 2012). An evaluation of the DPE pilot testing performance is included in Section 5.1.2 below. The evaluation confirms that the addition of groundwater extraction did not significantly improve SVE mass recovery, and groundwater recovery rates were very low. As such, DPE is not considered a viable technology for

remediation of impacted soil and groundwater, and therefore was not retained for consideration.

5.1.2 Evaluation of SVE and DPE Pilot Test Data

A series of four SVE tests were conducted by SLR using a standard SVE system, with and without simultaneous removal of groundwater (DPE), between December 12, 2011 and January 5, 2012. A complete summary of the testing can be found in SLR's report included in Appendix F. These tests were designed to evaluate SVE and DPE as potential remedial options for removing gasoline constituents from soil and groundwater. Both laboratory samples and photoionization detector (PID) readings were collected over the course of the testing. PID readings are affected by many factors, and therefore cannot confidently be used as a surrogate for actual gasoline-range TPH vapor concentrations. The laboratory analytical results were therefore used in evaluating SVE performance and mass removal rates.

Soil vapor samples were collected during all four tests and were analyzed for a suite of hydrocarbons including gasoline-range TPH by Northwest Method NWTPH-Gx. Test 1 was conducted at well SVE-1 without the removal of groundwater, and resulted in a relatively low concentration of gasoline-range TPH in extracted vapor (i.e., 47 milligrams per cubic meter [mg/m³] after 6 hours of operation). Test 3 was conducted on a combination of MW-1 and MW-2, and confirmed that the MW-1 is not suitable for SVE.

The results from Tests 2 and 4 both indicate that pumping groundwater while extracting vapor (i.e., DPE) from SVE-1 modestly improved removal of gasoline-range TPH. DPE resulted in a lower water table, with a resulting increase in the removed volatile gasoline constituents. The concentration of gasoline-range TPH in the vapor sample collected from the blower influent line after 9 hours of DPE operation was 1,900 mg/m³.

During Test 4, two effluent samples were collected from the blower influent line. The first was collected after 7 days of continuous DPE and had a gasoline-range TPH concentration of 30 mg/m³. The second sample was collected 2 days after the DPE system was restarted following a 4-day inadvertent shut-down. This second sample had a measured gasoline-range TPH concentration of 180 mg/m³, which indicates that some "rebound" of volatile compounds likely occurred in the subsurface during the 4-day shutdown.

The SVE/DPE removal rate trend for the period from Test 2 through Test 4 was evaluated using the gasoline-range TPH concentrations from effluent samples and the respective airflows measured at the time of sampling. Due to the intermittent nature of the testing, mass recovery was evaluated relative to the duration of sampling time from individual test startups, as this provided the best assessment of how a continuously operating system would perform.

Figure 4 provides a plot of measured effluent removal rates, using the duration of sampling time from individual test startups (as opposed to a synchronous analysis). The data collected from the SVE/DPE pilot tests indicate a relatively low starting mass removal rate of approximately 3 pounds per day, with a subsequent logarithmic decay in mass removal rate to very low recovery (0.14 pounds per day) after 7 days of operation. This type of decay is typical of SVE systems, which is one reason why high initial removal rates are a key to successful implementation of SVE. Based on evaluation of the

SVE/DPE test performance, SVE and DPE were not retained as viable remedial technologies.

5.2 Potential Cleanup Action Alternatives

As detailed in the Site UST Removal Report (AGI, 1991), following removal of the USTs in 1990, a cleanup action consisting of excavation of gasoline-impacted soil was completed to the extent practicable. Each of the cleanup action alternatives developed for the Site include this soil cleanup action as the initial component.

Four retained cleanup action alternatives were fully evaluated for comparison with MTCA criteria (WAC 173-340-350(8)). The four cleanup action alternatives are as follows:

- Alternative 1 Completed Soil Removal and No Additional Action;
- Alternative 2 Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls;
- Alternative 3 Completed Soil Removal and *In Situ* Chemical Oxidation; and
- Alternative 4 Completed Soil Removal and Excavation and Off-Site Disposal.

A description of each of these cleanup action alternatives and corresponding evaluation, based on the cleanup requirements set forth in MTCA, are provided in the following subsection.

5.3 Evaluation of Potential Cleanup Action Alternatives

This FFS considers the requirements under WAC 173-340-350, Site-specific conditions, and the criteria defined in WAC 173-340-360 for screening of potentially feasible remedial alternatives for the Site. A cleanup action alternative must satisfy the following threshold criteria, as specified in WAC 173-340-360(2):

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

These criteria represent the minimum standards for an acceptable cleanup action alternative. In addition to meeting the threshold criteria, cleanup action alternatives under MTCA will also:

- Use permanent solutions to the maximum extent practicable;
- Provide for a reasonable restoration timeframe, and
- Consider public concerns.

Evaluation of each of the cleanup action alternatives is provided below. FFS-level cost estimates for each alternative were calculated in accordance with U.S. Environmental Protection Agency (EPA) cost estimating guidance and professional experience with

similar projects (EPA, 2000). The cost for Alternative 2 was calculated as net present value (NPV) assuming a discount rate of 4 percent for a 15-year period. If long-term monitoring were to extend past this period, the NPV costs for monitoring after 15 years would be negligible. Cost estimate details and assumptions are provided in Tables 6 through 8.

5.3.1 Alternative 1 – Completed Soil Removal and No Additional Action

This alternative includes no additional action beyond the soil cleanup action completed in 1990. Though not implementable from a regulatory perspective, this alternative has zero cost and provides a baseline against which to compare other alternatives.

5.3.2 Alternative 2 – Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls

This alternative includes MNA with an environmental covenant. Specifically, this alternative would include:

- The soil removal action completed in 1990 which significantly reduced the mass of COCs in the proximal source area.
- MNA to reduce concentrations of COCs in soil and shallow groundwater through biodegradation, volatilization, and other naturally occurring processes.
- An environmental covenant (filed with the property deed) incorporating institutional controls to prevent exposure to residual concentrations of COCs in soil or shallow perched groundwater, and a groundwater monitoring plan to document the progress on MNA in reducing COC concentrations.

This alternative highly implementable and economical, poses very little short-term risk, and is minimally disruptive to the operations of the OWSI facility. The potential for human exposure through direct-contact or ingestion of soil with residual concentrations of COCs under this alternative is prevented through institutional controls and restrictions on excavation or subsurface penetration established in the environmental covenant

The potential for human exposure through direct-contact or ingestion of COCs in shallow groundwater is prevented under this alternative through a restriction on shallow groundwater use in the environmental covenant, even though shallow perched waterbearing zone is not currently used for drinking water, and is a not expected to be a future source. The MNA component of this alternative provides for monitoring of the natural degradation of dissolved phase COCs in shallow perched groundwater.

This cleanup alternative would likely eventually achieve the proposed cleanup levels for the Site, complies with applicable State and Federal laws, provides for performance and compliance monitoring, and considers public concerns. This cleanup alternative would also eventually result in a permanent solution. Based on Aspect's previous experience on similar sites, groundwater monitoring to assess the progress of MNA on reducing dissolved concentrations of COCs in groundwater would be required over an extended period of time. At achievement of groundwater compliance, soil confirmation sampling would also likely be required. The estimated cost of this alternative is \$130,000 (Table 6). Restoration time frame is estimated at 15 years.

5.3.3 Alternative 3 – Completed Soil Removal and In Situ Chemical Oxidation

This alternative includes the following components:

• *In situ* chemical oxidation to reduce concentrations of COCs in shallow soil and perched groundwater to below MTCA Method A cleanup levels.

The *in situ* chemical oxidation component is estimated to include injection of a chemical oxidant at up to 20 permanent injection wells spaced approximately 15 feet apart to treat up to a 20-feet-thick zone of impacted soil within the mapped area of impacts. Seven separate injection events are scoped in this alternative. Injection-point spacing is an estimation based on Regenesis documentation.

The *in situ* chemical oxidation technology will require bench-scale and/or pilot testing to evaluate its potential effectiveness, select the appropriate oxidant, and design an injection program. Based on a preliminary estimate of the total mass of TPH in soil (approximately 880 pounds), approximately 28,000 pounds of RegenOx, a chemical oxidant supplied by Regenesis will be required over the seven injection events. Periodic post-injection protection groundwater monitoring would also be required to confirm that groundwater quality achieves compliance with MTCA Method A cleanup levels.

Assuming successful bench scale testing, this cleanup alternative would likely achieve the proposed cleanup levels for the Site. It complies with applicable State and Federal laws, provides for performance and compliance monitoring, and considers public concerns. This cleanup alternative would also eventually result in a permanent solution. Based on Aspect's previous experience on similar sites, groundwater monitoring to assess the progress of *in situ* chemical oxidation on reducing dissolved concentrations of COCs in groundwater would be required over an approximate 5-year period.

The estimated cost of this alternative is \$650,000 (Table 7). Restoration time frame is estimated at 5 years.

5.3.4 Alternative 4 – Completed Soil Removal and Excavation and Off-Site Disposal

This alternative includes soil excavation to address residual concentrations of COCs in soil above the shallow perched groundwater-bearing zone, and follow-up groundwater monitoring to confirm that MNA reduces residual concentrations of COCs in shallow groundwater to below MTCA Method A cleanup levels. The excavation component of this alternative would require demolition of the existing garage building, excavation and off-Site disposal of residual gasoline-impacted soil, and construction of a new garage building.

As noted in Section 3.3.1, residual gasoline-impacted soil is expected to occur beneath the garage building from below approximately 10 feet bgs to the perched groundwater table at depths ranging from approximately 20 to 41 feet bgs (SLR, 2011). Based on the results of the RI activities, the estimated area of impacted soil covers an area of

approximately 3,140 square feet (see Figure 7 of the Additional Investigation Report; SLR, 2011). This alternative includes excavation of approximately 12,000 bank cubic yards of soil, including overburden and gasoline-impacted soil. Scoping of this alternative assumes the excavation can be completed by sloping, and without shoring. Shoring would add significantly to both the complexity and cost of implementation.

During excavation, overburden would be stockpiled, tested, and reused as backfill. Gasoline-impacted soil with COC concentrations above MTCA Method A cleanup levels would be transported off-site for disposal at a permitted landfill. Periodic post-excavation groundwater monitoring would also be required to confirm that shallow perched groundwater quality achieves compliance with MTCA Method A cleanup levels.

This cleanup alternative would likely achieve the proposed cleanup levels for the Site. It complies with applicable State and Federal laws, provides for performance and compliance monitoring, and considers public concerns. This cleanup alternative would also eventually result in a permanent solution. Based on Aspect's previous experience on similar sites, groundwater monitoring to assess post-excavation attenuation of residual COCs in groundwater would be required over an approximate 3-year period.

The estimated cost of this alternative is \$1,250,000 (Table 8). Restoration time frame is estimated at 4 years.

5.4 Disproportionate Cost Analysis

A disproportionate cost analysis (DCA) was completed in accordance with WAC 173-340-360. The DCA provides a means to balance the cost to benefit associated with an alternative and allows for elimination of alternatives for which the incremental costs are disproportionate relative to the benefits. The DCA for the retained four alternatives is presented in Table 9. Figure 5 provides a graphical presentation of the cost to overall alternative ranking (e.g.: benefit) comparison for each of the four alternatives. The criteria used to qualitatively evaluate potentially applicable cleanup alternatives in the DCA were derived from WAC 173-340-360(3)(f). These criteria, which were assigned weighting factors in Table 9 in accordance with applicable Ecology guidance, include:

- **Protectiveness:** Overall protectiveness of human health and the environment, including the degree to which existing risks are reduced, the time required to reduce risk at the Site and attain cleanup standards, on-Site risks resulting from implementing the alternative, and the improvement of overall environmental quality.
- **Permanence:** The degree to which the alternative permanently reduces the toxicity, mobility, or volume of hazardous substances, including the adequacy 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 the waste treatment process, and the characteristics and quantity of treatment residuals generated.
- **Long-term effectiveness:** Long-term effectiveness includes the degree of certainty that the alternative will be successful, the reliability of the alternative during the period of time that hazardous substances are expected to remain on the

Site at concentrations that exceed cleanup levels, and the magnitude of residual risk with the alternative in place.

- **Management of short-term risks:** The risk to human health and the environment associated with the alternative during construction and implementation, and the effectiveness of measures that will be taken to manage such risks.
- **Technical and administrative implementability:** Ability to be implemented, including consideration of whether the alternative is technically feasible, administrative and regulatory requirements, permitting, scheduling, size, complexity, monitoring requirements, and access.
- **Consideration of public concerns:** Whether the community has concerns regarding the alternative and, if so, the extent to which the alternative addresses those concerns. This process involves concerns from individuals, community groups, local governments, federal and state agencies, or any other organization that may have an interest in or knowledge of the Site.
- **Cost:** The cost to implement the alternative, including the cost of construction, the net present value of any long-term costs, and Ecology oversight costs. Long-term costs include operation and maintenance, monitoring, and reporting costs.

The Site DCA documented in Table 9 and Figure 5 assigns each alternative an overall MTCA benefit ranking on a scale of 1 to 10, with 1 representing the lowest protectiveness, permanence, effectiveness, risk, implementability, and greatest level of public concern. The evaluated alternatives for the Site are ranked as follows:

- Alternative 1 Completed Soil Removal and No Additional Action: This alternative was assigned an overall MTCA benefit ranking value of 4.7. The estimated cost of implementation is \$0;
- Alternative 2 Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls: This alternative was assigned an overall MTCA benefit ranking value of 7.1. The estimated cost of implementation is \$130,000;
- Alternative 3 Completed Soil Removal and *In-Situ* Chemical Oxidation: This alternative was assigned an overall MTCA benefit ranking value of 7.2. The estimated cost of implementation is \$650,000; and
- Alternative 4 Completed Soil Removal and Excavation with Off-Site Disposal: This alternative was assigned an overall MTCA benefit ranking value of 7.5. The estimated cost of implementation is \$1,250,000.

As shown on Figure 5, Alternatives 2, 3 and 4 have essentially comparable benefit rankings. Alternative 3 provides a nominal net 1% incremental benefit over Alternative 2, and Alternative 4 provides a nominal net benefit of 4% over Alternative 2. Despite these limited incremental benefits, the costs to implement Alternatives 3 and 4 range from approximately 6 times (Alternative 3) to ten times (Alternative 4) the cost for implementation of Alternative 2. Based on the comparable protectiveness and effectiveness provided by Alternative 2, and disproportionate cost of the nominal incremental benefits provided by either Alternatives 3 or 4, Alternative 2 is identified as the preferred alternative.

6 Summary of Preferred Cleanup Action

The preferred cleanup action alternative for the Site is Cleanup Alternative 2 – Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls. This cleanup action would be conducted independently under the Ecology VCP with the long-term objective of obtaining an NFA determination for the Site.

Alternative 2, as the recommended cleanup action for this Site, would include the specific elements detailed below.

- **Institutional Controls:** These would be incorporated into an environmental covenant filed with the deed on the property. The covenant would restrict certain activities that could cause exposure to impacted soils or groundwater, or result in unacceptable mobilization of subsurface COCs. Non-commercial land uses would also be prohibited by the covenant unless and until a new analysis of remedial alternatives is prepared and Ecology approves additional cleanup actions designed to protect public health and the environmental under non-commercial land use scenarios.
- **COC Monitoring Program:** The covenant would include a groundwater sampling plan addressing implementation of an MNA groundwater sampling program to document the progress of natural attenuation of residual COCs. The groundwater sampling plan would include sampling of the selected existing wells on an annual basis, with analysis for TPH as gasoline and BTEX compounds. After groundwater compliance is achieved, the covenant would also likely include a requirement for confirmation of COC attenuation in soil as a prerequisite to removal of the environmental covenant and issuance of an NFA letter.

Based on the results of the DCA, the recommended cleanup action alternative for the Site is Cleanup Alternative 2 – Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls.

7 References

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Limitations

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TABLES

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Table 1 - Summary of Groundwater Elevation Data

Olympic Water and Sewer, Inc. Site 781 Walker Way, Port Ludlow, Washington

Well Number	Top of Casing Elevation ^a (feet)	Date Measured	Depth to Groundwater ^b (feet)	Groundwater Elevation (feet)
		06/14/10	41.33	252.69
MW-1	294.02	10/20/10	40.30	253.72
		04/08/11	36.98	257.04
		06/14/10	39.63	254.16
MW-2	293.79	10/20/10	40.71	253.08
		04/08/11	36.90	256.89
		06/14/10	25.19	264.18
MW-3	289.37	10/20/10	28.70	260.67
		04/08/11	23.02	266.35
		06/14/10	23.92	271.41
MW-4	295.33	10/20/10	26.67	268.66
		04/08/11	21.95	273.38
MW-5 ^c	299.40	04/08/11	23.55	275.85
SVE-1	294.41	04/08/11	21.22	273.19
SVE-2	293.87	04/08/11	19.80	274.07

Notes:

^aTop of casing elevations were surveyed relative to the NAVD 88 datum.

^bDepth to groundwater measured in feet below top of PVC casing.

^cThe top of the casing of MW-5 is 2.92 feet above the ground surface. All of the other wells and the soil vapor extraction points are flush-grade completions.

Table 1

Table 2 - Summary of June 21, 2013 Sub-Slab Soil Vapor Sampling Results

Olympic Water and Sewer, Inc. Site

781 Walker Way, Port Ludlow, Washington

Sample ID	Analyte Group	Analyte	Carcinogen(C) or Non- Carcinogen(NC)	Units	Result	Result with Leakage Correction	Hag	Reporting Limit	MTCA Method B Indoor Air CUL	Site-Specific Commercial Land Use Sub-Slab Soil Vapor Screening Level ^{4,5}
		C ₅ - C ₈ Aliphatic Hydrocarbons ^{1,2}	NC	µg/m3	110	110.02		28	2,700	59,063
	APH	C9 - C12 Aliphatic Hydrocarbons1.3	NC	µg/m3	2,100	2100.38		14	140	3,063
		C9 - C10 Aromatic Hydrocarbons	NC	µg/m3	42	42.01		3.5	180	3,938
		Methyl tert-Butyl Ether	С	µg/m3			ND	0.70	9.6	960
		n-Hexane	NC	µg/m3			ND	0.70	320	7,000
		1,2-Dichloroethane	С	µg/m3			ND	0.70	0.096	9.6
		Benzene	С	µg/m3			ND	0.70	0.32	32.0
VP-1-062113		Toluene	NC	µg/m3	9.8	9.80		0.70	2,200	48,125
VF-1-002113	TO-15	1,2-Dibromoethane (EDB)	С	µg/m3			ND	0.70	0.01	1.1
	10-13	Ethylbenzene	NC	µg/m3	3.2	3.20		0.70	460	10,063
		m,p-Xylenes	NC	µg/m3	15	15.00		1.4	46	1,006
		o-Xylene	NC	µg/m3	4.4	4.40		0.70	46	1,006
		1,3,5-Trimethylbenzene	NC	µg/m3	2.4	2.40		0.70	2.7	59
		1,2,4-Trimethylbenzene	NC	µg/m3	7.7	7.70		0.70	2.7	59
		Naphthalene	NC	µg/m3	1.2	1.20		0.70	1.4	31
	Tracer	Helium		ppmV	180			28	-	-
		C ₅ - C ₈ Aliphatic Hydrocarbons ^{1,2}	NC	µg/m3	100			27	2,700	59,063
	APH	C9 - C12 Aliphatic Hydrocarbons1.3	NC	µg/m3	790			14	140	3,063
		C9 - C10 Aromatic Hydrocarbons	NC	µg/m3	16			3.4	180	3,938
		Methyl tert-Butyl Ether	С	µg/m3			ND	0.68	9.6	960
		n-Hexane	NC	µg/m3			ND	0.68	320	7,000
		1,2-Dichloroethane	С	µg/m3			ND	0.68	0.10	9.6
		Benzene	С	µg/m3			ND	0.68	0.32	32.0
VP-2-062113		Toluene	NC	µg/m3	12			0.68	2,200	48,125
11 2 002110	TO-15	1,2-Dibromoethane (EDB)	С	µg/m3			ND	0.68	0.01	1.1
	10 10	Ethylbenzene	NC	µg/m3	10			0.68	460	10,063
		m,p-Xylenes	NC	µg/m3	45			1.4	46	1,006
		o-Xylene	NC	µg/m3	28			0.68	46	1,006
		1,3,5-Trimethylbenzene	NC	µg/m3	1.60			0.68	2.7	59
		1,2,4-Trimethylbenzene	NC	µg/m3	3.5			0.68	2.7	59
		Naphthalene	NC	µg/m3			ND	0.68	1.4	31
	Tracer	Helium		ppmV			ND	27	-	-

Notes:

Significant non-petroleum related peaks are subtracted from the APH hydrocarbon range areas when present.

¹Hydrocarbon Range data from total ion chromatogram excluding any internal/tuning standards eluting in that range.

²C₅-C₈ Aliphatic Hydrocarbons exclude the concentration of Target APH analytes eluting in that range.

³C₉-C₁₂ Aliphatic Hydrocarbons exclude concentration of Target APH Analytes eluting in that range and concentration of C₉-C₁₀ Aromatic Hydrocarbons.

⁴Conservative cross-slab attenuation factor of 10, per Ecology's Draft Guidance for Evaluating Soil Vapor Intrusion in Washington State (2009).

⁵Site-specific correction for adult worker exposure scenario, calculated in accordance with WAC 173-340-750 and Ecology's Draft Guidance for Evaluating Soil Vapor Intrusion in Washington State (2009).

Table 2

Table 3 - Summary of Soil Analytical Data

Olympic Water and Sewer, Inc. Site

781 Walker Way, Port Ludlow, Washington

			Approximate			Analytic	al Results	(mg/kg)		
Soil Boring Number	Sample Name	Date Collected	Sample Depth (feet)	Benzene ^a	Toluene ^a	Ethylbenzene ^a	Total Xylenes ^a	Naphthalene ^a	Gasoline- range TPH [♭]	Lead ^c
MTCA Method A Cleanup Levels ^d			0.03	7	6	9	5	30	250	
SVE-1	SVE1-14	04/04/11	14 to 14.3	<0.02	0.04	<0.02	0.074	NA	<2.0	
SVE-2	SVE2-20	04/04/11	20 to 20.5	<0.02	0.64	0.55	1.50	NA	110	
SB-1	SB1-35	04/04/11	35 to 35.5	<0.02	< 0.02	<0.02	<0.06	NA	<2.0	
MW-5	MW5-25	04/06/11	25 to 25.5	<0.02	< 0.02	<0.02	<0.06	NA	<2.0	
MW-1B	MW1-24.5-25	04/14/10	24.5 to 25	0.49	5.70	1.20	6.70	0.58	140	1.11
MW-1	MW-1-40	06/08/10	40 to 40.3	<0.03	< 0.05	<0.05	<0.15	<0.05	<2	NA
MW-1	MW1-55	06/08/10	55 to 55.5	<0.03	<0.05	<0.05	<0.15	<0.05	<2	NA
MW-2	MW2-40	06/09/10	40 to 40.3	.21 ^e	0.062	0.11	0.066	<0.05	2.90	NA
MW-2	MW2-55.5	06/09/10	55.5 to 55.8	.21 ^e	<0.05	<0.05	<0.15	<0.05	<2	NA
MW-3	MW-3-30.5	06/09/10	30.5 to 30.9	< 0.03	<0.05	<0.05	<0.15	<0.05	<2	NA
MW-3	MW3-45.5	06/10/10	45.5 to 45.9	0.036 ^f	<0.05	<0.05	<0.15	<0.05	<2	NA
MW-4	MW4-31	06/10/10	30.5 to 31	<0.03	<0.05	<0.05	<0.15	<0.05	<2	NA
MW-4	MW4-55	06/11/10	55 to 55.5	<0.03	<0.05	<0.05	<0.15	<0.05	<2	NA

Notes:

mg/kg = milligrams per kilogram (ppm).

Values in **bold** exceed the soil cleanup levels.

NA = Not analyzed.

TPH = Total Petroleum Hydrocarbons

^aBenzene, toluene, ethylbenzene, total xylenes (BTEX), and naphthalene by EPA Method 8021B.

^bGasoline-range TPH by Northwest Method NWTPH-Gx or 8260c.

^cLead by EPA Method 200.8.

^dChapter 173-340 WAC, Model Toxics Control Act (MTCA) Cleanup Regulation, Method A Cleanup Levels. Amended 2007.

^eThe benzene concentration in this sample likely reflects dissolved benzene in pore water rather than benzene adsorbed to the soil.

All data from this table is from Site Characterization Report, Olympic Water & Sewer, Inc. Property, 781 Walker Way, Port Ludlow, Washington dated December 17, 2010 prepared by SLR and Additional Investigation Report, Olympic Water & Sewer, Inc. Property, 781 Walker Way, Port Ludlow, Washington dated August 2, 2011, prepared by SLR.

Table 4 - Summary of Groundwater Analytical Data

Olympic Water and Sewer, Inc. Site

781 Walker Way, Port Ludlow, Washington

	Dete		Analytical Results (µg/L)										
Well Number	Date Collected	Gasoline-range TPH ^a	Benzene ^b	Toluene ^b	Ethylbenzene ^b	Total Xylenes ^b	Naphthalene ^b	MTBE ^b	EDC [♭]	EDB℃	Total Lead ^d		
MTCA Method A	Cleanup Levels ^e	800	5	1,000	700	1,000	160 ^f	20	5	0.01	15		
	06/14/10	990	110	45	1.10	186	<1	<1	<1	<0.01	<1		
MW-1	10/20/10	1,900	520	140	110	221	15	NA	NA	NA	NA		
	04/07/11	3,000	530	82	160	120	NA	NA	NA	NA	NA		
	06/14/10	8,400	2,100	620	960	650	100	<1	<1	<0.01	<1		
MW-2	10/20/10	3,900	1,300	290	430	530	35	NA	NA	NA	NA		
	04/07/11	5,600	500	730	160	410	NA	NA	NA	NA	NA		
	06/14/10	<100	0.36	<1	<1	<3	<1	<1	<1	<0.01	<1		
MW-3	10/20/10	<100	<0.35	<1	<1	<3	<1	NA	NA	NA	NA		
	04/07/11	<100	<1	<1	<1	<3	NA	NA	NA	NA	NA		
	06/14/10	<100	<0.35	<1	<1	<3	<1	<1	<1	<0.01	<1		
MW-4	10/20/10	<100	<0.35	<1	<1	<3	<1	NA	NA	NA	NA		
	04/08/11	380	5.30	75	13	47	NA	NA	NA	NA	NA		
MW-5	04/08/11	220	3.40	43	7.80	25	NA	NA	NA	NA	NA		
SVE-1	04/07/11	34,000	550	5,700	850	3,300	NA	NA	NA	NA	NA		
SVE-2	04/07/11	2,000	5.0	14	18	35	NA	NA	NA	NA	NA		

Notes:

 μ g/L = micrograms per liter (ppb).

Values shaded and in **bold** exceed the groundwater cleanup levels.

NA = Not analyzed.

^aGasoline-range TPH by Northwest Method NWTPH-Gx or 8260c..

^bBenzene, toluene, ethylbenzene, and total xylenes (BTEX), naphthalene, methyl tertiary butyl ether (MTBE), and 1,2-dichloroethane (EDC) by EPA Method 8260C, or BTE

 $^{\rm c}$ 1,2-dibromoethane (EDB) by EPA Method 8011 Modified.

^dTotal lead by EPA Method 200.8.

^eChapter 173-340 WAC, Model Toxics Control Act (MTCA) Cleanup Regulation, Method A Cleanup Levels. Amended 2007.

^fThe cleanup level is the total value for naphthalene, 1-methyl naphthalene, and 2-methyl naphthalene.

Data from this table is from Site Characterization Report, Olympic Water & Sewer, Inc. Property, 781 Walker Way, Port Ludlow, Washington dated December 17, 2010 prepared by SLR and Additional Investigation Report, Olympic Water & Sewer, Inc. Property, 781 Walker Way, Port Ludlow, Washington dated August 2, 2011, prepared by SLR.

Table 5 - Preliminary Screening of Remedial Technologies

Olympic Water and Sewer, Inc. Site 781 Walker Way, Port Ludlow, Washington

Remedial Technology	Effectiveness	Implementability	Comparative Cost	Screening Result
Institutional Controls	low	high	low	Retained
Engineering Controls	low	high	low	Retained
Monitored Natural Attenuation	medium	high	low	Retained
Soil Vapor Extraction and Dual- Phase Extraction	low	low	medium	Not Retained
Air Sparging	low	medium	medium	Not Retained
Enhanced Aerobic Biodegradation	low	low	medium	Not Retained
In-Situ Chemical Oxidation	medium	medium	medium	Retained
Soil Excavation	high	low	high	Retained

Table 6 - Alternative 2 Cost Estimate - Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls

Olympic Water and Sewer, Inc. Site

781 Walker Way, Port Ludlow, Washington

Site:	Olympic Wate	r and Sewer, In	c.						
Remedial Action Description:	Completed So	Completed Soil Removal and MNA with Environmental Covenant with Institutional Controls							
Key Assumptions:	Annual groundwater and surface water monitoring required Future costs are adjusted to present value using a discount rate of 4 percent Environmental covenant required for soil and shallow groundwater								
Item	Quantity Ur	it Unit Cost	E	Est. Cost	Notes				
Professional Services									
project management	20%	\$ 75,592	\$	15,118	Planning and reporting				
remedial design	1 ls	\$ 3,500	\$	3,500	Develop monitoring plan				
environmental covenant	1 ls	\$ 10,000	\$	10,000	Covenant for soil and shallow groundwater				
Subtotal			\$	28,618					
Monitoring									
					5 shallow wells, Well #2, and 1 surface water				
Groundwater for 15 years	15 ea	\$ 5,000	\$	55,592	sample per event				
confirmation soil sampling	1 ls	\$ 20,000	\$	20,000	upon completion of active remediatior				
Subtotal			\$	75,592					
Contingency	25%		\$	26,053	25% scope and restoration time frame contingence				
Total Estimated Cost			\$	130,000	(rounded to the nearest \$10,000)				

Notes:

ea = each

ls = lump sum

Table 7 - Alternative 3 Cost Estimate - Completed Soil Removal and In Situ Chemical Oxidation

Olympic Water and Sewer, Inc. Site

781 Walker Way, Port Ludlow, Washington

Site:	Olympic Water and Sewer, Inc.									
Remedial Action Description:	Completed Soil F	Rem	noval and	In S	itu Chemica	al Oxidation				
Key Assumptions:	Average TPH-gas concentration in soil = 125 mg/kg Area of impacts = 3,200 sq ft Treatment thickness = 20 ft Average soil density = 110 lb/cu ft Oxidant requirements estimated using: <i>Principles of Chemical Oxidation Technology, Design and Application Manual, V.3.0</i> (Regenesis, 2010)									
ltem	Quantity Unit	U	nit Cost	E	st. Cost	Notes				
Professional Services										
project mgmt	6%	\$	387,400	\$	23 244	percentage of capital and monitoring costs				
remedial design	12%		295.400			percentage of capital and monitoring costs				
construction mgmt	8%		295,400			percentage of capital costs				
Subtotal	078	Ψ	233,400	\$	82,324	percentage of capital costs				
ISCO Pilot Test										
install injection wells	3 ea	\$	8,000	\$	24,000					
injection event	2 day	\$	2,500	\$	5,000					
Regen-Ox amendment	600 lb	\$	1.50	\$	900	assumes 200 lbs per injection well				
Subtotal				\$	29,900	-				
In Situ Chemical Oxidation										
install injection wells	17 ea	\$	8,000	\$	136,000					
injection events	7 week	\$	12,500	\$	87,500	7 events (20 wells total, inject 5 wells per day)				
Regen-Ox Amendment	28000 lb	\$	1.50		42,000	estimated using Regenesis design guidelines				
Subtotal				\$	265,500					
Monitoring										
quarterly groundwater sampling	5 yr	\$	12,000	\$	60,000	5 wells quarterly to monitor performance and confirmation				
confirmation soil sampling	1 ls	\$	20,000		20,000	upon completion of active remediation				
well abandonment	1 ls	\$	12,000		12,000					
Subtotal				\$	92,000					
Тах	9.5%			\$	28,063	Washington Sales Tax (applied to capital costs)				
Contingency	30%			\$	149,336	25% scope and restoration time frame contingency				
Total Estimated Cost				\$	650.000	(rounded to the nearest \$10,000)				

Notes:

ea = each lb = pound

ls = lump sum

yr = year

Table 8 - Alternative 4 Cost Estimate - Completed Soil Removal and Excavation and Off-Site Disposal

Olympic Water and Sewer, Inc. Site

781 Walker Way, Port Ludlow, Washington

Site:	Olympic Water	and S	Sewer, Inc.								
Remedial Action Description:	Completed Soil Removal and Excavation and Off-Site Disposal										
Cost Estimate Accuracy:		easibility Level (+50/-30 percent)									
Key Assumptions:	4,225 square foot area (65 feet by 65 feet) excavated to 40 foot depth 3/4:1 (H:V) sloped excavation, no shoring required Area of impacted soil = 3,200 sq ft Overburden average thickness = 15 feet Impacted zone thickness = 25 ft Average soil density = 110 lb/cu ft Clean overburden and soil from sloping excavation reused for backfill Costs for tree restoration are not included										
ltem	Quantity Unit	ιL	Init Cost	Т	otal Cost	Notes					
Professional Comisso											
Professional Services	6%	¢	773,738	¢	16 104	perceptage of capital and manitaring costs					
project mgmt remedial design	6%	ф Ф	705,738			percentage of capital and monitoring costs permitting, planning, geotechnical engineering					
•	4%		705,738			percentage of capital costs					
construction mgmt Subtotal	4%	φ	105,136	م \$	116,998						
ousional				Ψ	110,000						
Soil Excavation											
mobilization/demobilization	1 ls	\$	10,000	\$	10,000	recent project experience					
building demo and disposal	26680 cf	\$	0.40	\$	10,672	RSMeans for building demo/recent project experience ¹					
tree removal	0.1 acre	\$	10,430	\$	1,043	RSMeans for clearing and grubbing up to 12" trees ¹					
decommission/replace monitoring wells	3 ea	\$	7,500	\$	22,500	MW-1, MW-2, MW-5					
excavation/loading/stockpiling	12037 bcy	\$	4	\$	48,148	RSMeans for bulk excavation and loading ¹					
PCS hauling and disposal	6455 ton	\$	50	\$	322,743	Local current pricing					
dewatering during excavation	1 ls	\$	10,000	\$		estimate for pumps, storage, labor					
impacted water disposal	5000 gal	\$	0.40	\$	2,000	Local current pricing					
purchase and import clean backfill	7000 ton	\$	10	\$		Local current pricing					
place and compact clean backfill	15046 bcy	\$	5	\$	75,231	RSMeans for backfill plus compaction ¹					
replace building	1334 sf	\$	100	\$	133,400	engineer estimate					
Subtotal				\$	705,738						
Monitoring											
quarterly groundwater sampling	3 yr	\$	12,000	\$	36 000	5 wells guarterly to monitor performance and confirmation					
confirmation soil sampling	1 ls	\$	20,000		20,000	e were quartery to monitor performance and committation					
well abandonment	1 Is	\$	12,000		12,000						
Subtotal	1.10	Ŷ	.2,000	\$	68,000	-					
Tax	9.5%			\$	67,045	Washington Sales Tax (applied to capital costs)					
Contingency	30%			\$	287,334	Volume and implementation contingency					
Total Estimated Cost				\$	1,250,000	(rounded to the nearest \$10,000)					

Notes:

¹ Unit cost estimates from rsmeansonline.com acre = acres bcy = bank cubic yard ea = each gal = gallons ls = lump sum sf = square foot yr = year

Table 9 - Summary of Alternatives Evaluation

Olympic Water and Sewer, Inc. Site 781 Walker Way, Port Ludlow, Washington

Alternative Number	Alternative 1	Alternative 2	Alternative 3	Alternative 4
Description	No Additional Action	Completed Soil Removal and MNA with	Completed Soil Removal and In Situ Chemical	Completed Soil Removal and Excavation and Off-
		Environmental Covenant with Institutional	Oxidation	Site Disposal
		Controls		
overall Alternative Ranking	4.7	7.1	7.2	7.5
ompliance with MTCA Threshold C		·	·	
Protection of Human Health	Potentially not, since it includes no covenant to	Yes - Alternative will protect human health and	Yes – Alternative will protect human health and	Yes – Alternative will protect human health and
and the Environment	prevent exposure to soil or groundwater	the environment.	the environment.	the environment.
Compliance with Cleanup	Long restoration time since no active measures are	Yes – However, long restoration time since no	Yes – Active remedial measures are used for	Yes – Active remedial measures are used for
Standards	used for soil of groundwater not complying with cleanup standards.	active measures are used for soils not complying with cleanup standards.	soils not complying with cleanup standards.	soils not complying with cleanup standards.
Compliance with Applicable	No – Alternative includes no monitoring or	Yes – Alternative complies with applicable	Yes – Alternative complies with applicable laws.	Yes – Alternative complies with applicable laws.
State and Federal Laws	environmental covenant.	laws.		
Provision for Compliance Monitoring	None	Yes – Alternative includes provisions for compliance monitoring.	Yes – Alternative includes provisions for compliance monitoring.	Yes – Alternative includes provisions for compliance monitoring.
Restoration Time Frame	Potentially greater than 15 years	Potentially greater than 15 years	Minimum of 4 years	Minimum of 3 years
valuation Criteria				
Protectiveness	2	8	8	9
(30% Weighted Factor):	Potentially not protective of exposure pathways.	Protective of exposure pathways through use of institutional controls/environmental covenant.	Protective of exposure pathways, some risk since the technology untested at the Site.	Highest degree of assurance for protection of identified exposure pathways.
Permanence	5	5	7	9
(20% Weighted Factor):	Natural attenuation will reduce the volume and concentration of residual impacted soil and groundwater over an extended restoration time frame.	concentration of residual impacted soil and	Volume and concentration of residual impacted soil and groundwater is reduced; technology less certain than Alternative 4 because it is unproven at the Site.	Highest degree of assurance for reduction short term reduction in volume and concentration of residual impacted soil and groundwater.
Long-Term Effectiveness (20% Weighted Factor):	4 Relies on natural attenuation to reduce the volume and concentration of residual impacted soil and groundwater over an extended restoration time frame.	soil and groundwater over an extended restoration time frame. Exposures prevented through institutional controls/environmental	7 Destroys petroleum compounds in soil and groundwater. Some implementation risk since technology is untested at the Site.	9 Soil removal coupled with groundwater natural attenuation is a proven effective remedial approach at petroleum sites.
Short-Term Risk Management	Q	covenant.	7	5
(10% Weighted Factor):	No short term risk, since alternative involves no construction.	No short term risk, since alternative involves no construction.	This alternative poses a moderate amount of risk from drilling and construction activities.	° °
Implementability	10	9	6	3
(10% Weighted Factor):	High implementability.	High implementability. Requires recording of covenant, and implementation of monitoring program and institutional controls.	Requires drilling wells inside building, drilling is limited by very dense soils, application is untested at the Site.	Implementation is complicated by need to demolish building, management of deep unshored excavation, limitations on equipment, truck, and overburden staging, and construction of a new building.
Public Concerns	4	7	7	4
(10% Weighted Factor):	Likely concerns over lack of monitoring or institutional controls.	Potential concerns over extended restoration time frame.	Potential concerns over moderate disruption and inconvenience to local residents.	Likely significant concerns over disruption, noise, and inconvenience to local residents.
Cost	\$0	\$130,000	\$650,000	\$1,250,000
	ψυ	φ130,000	ψυου,υυυ	ψ1,200,000

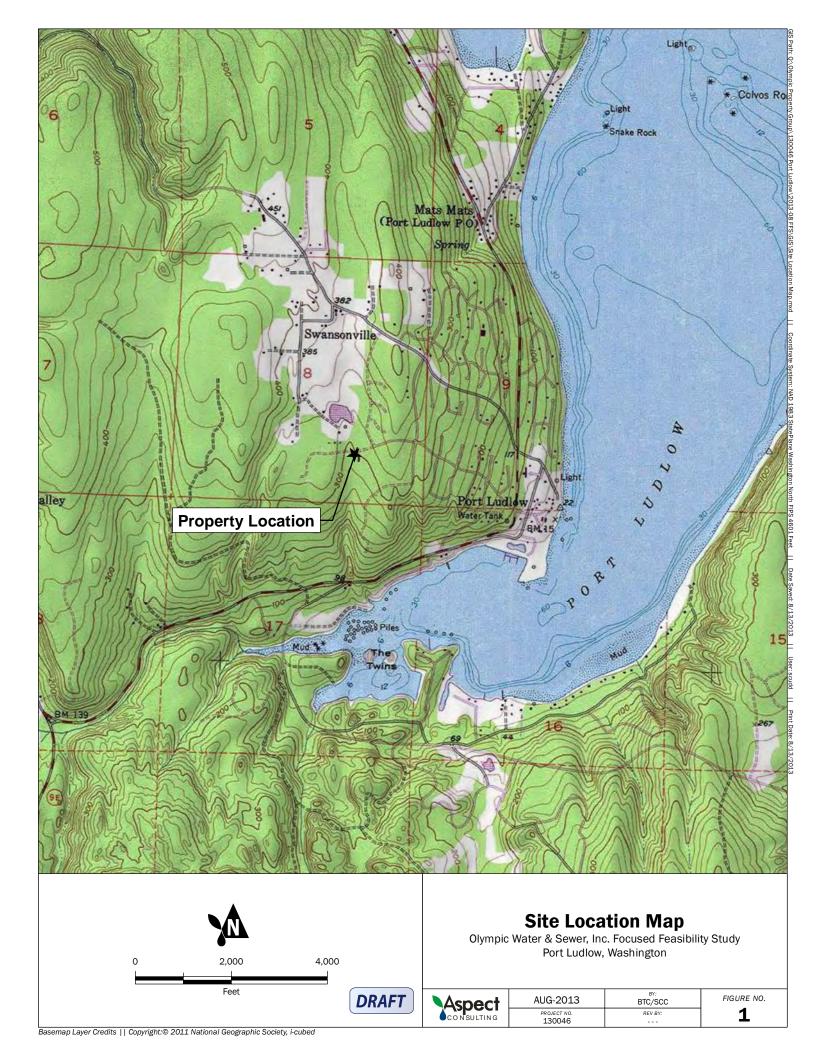
Notes:

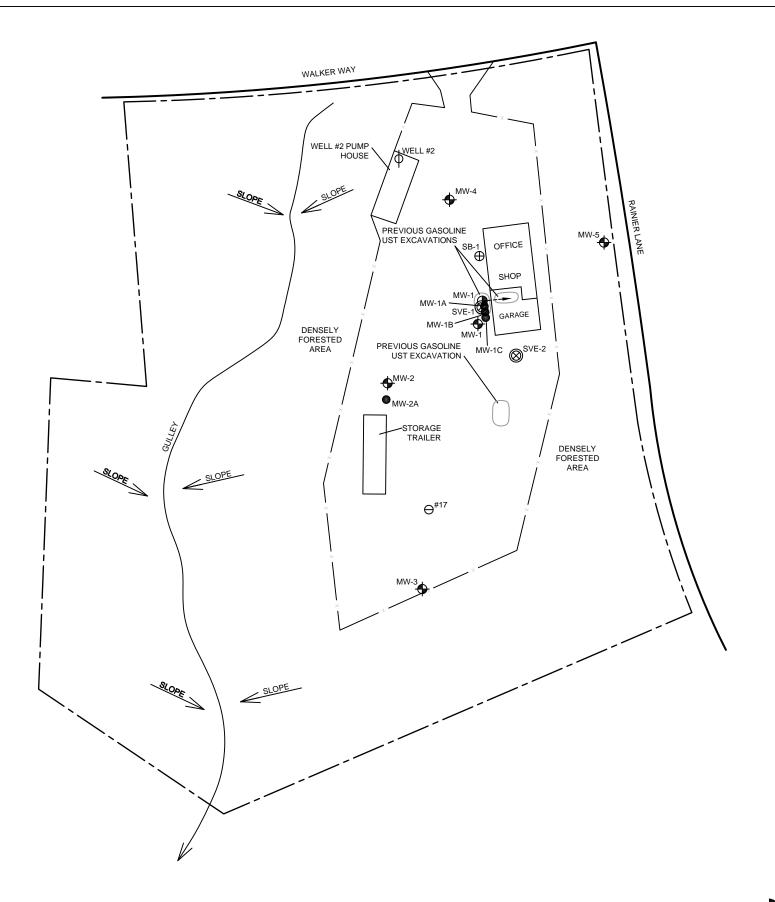
For each evaluation criterion, technologies are ranked on a scale of 1 to 10, with 1 representing the lowest protectiveness, permanence, effectiveness, risk and implementability, or greatest level of public concern. Aspect Consulting

9/24/2013

W:\130046 OPG Port Ludlow Property\Deliverables\FFS\Agency Draft\Tables\OSWI DCA Tables.xls

FIGURES





THIS DRAWING IS FOR CONCEPTUAL PURPOSES ONLY. ACTUAL LOCATIONS MAY VARY AND NOT ALL STRUCTURES ARE SHOWN.

BASE MAP PROVIDED BY SLR, FROM ADDITIONAL INVESTIGATION REPORT, DATED MAY 12, 2011.



SCALE: 1" = 50' WHEN PLOTTED AT 11" x 17" PAGE SIZE 50 100

LEGEND

	PROPERTY BOUNDARY
SB-1 🕀	APPROXIMATE 2011 SOIL BORING LOCATION AND DESIGNATION
SVE-1	PROPOSED SOIL VAPOR EXTRACTION POINT LOCATION AND DESIGNATION
MW-1 -	GROUNDWATER MONITORING WELL LOCATION AND DESIGNATION
MW-2A ●	APPROXIMATE SOIL BORING LOCATION AND DESIGNATION
#17 🕁	EXISTING CASING LOCATION AND DESIGNATION
WELL #2 🔿	EXISTING WATER SUPPLY WELL LOCATION AND DESIGNATION
MW-1	PREVIOUS ANGLE BORING LOCATION, DESIGNATION, AND DIRECTION FROM VERTICAL
×	FENCE

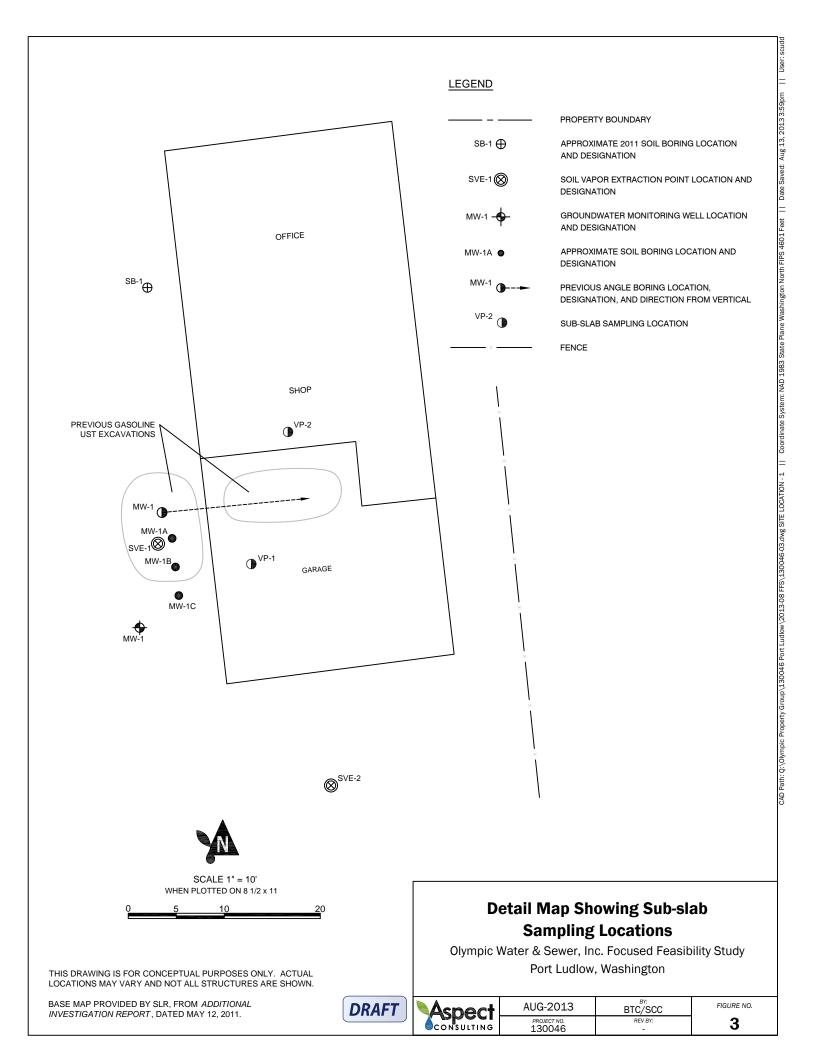


Site Plan

Olympic Water & Sewer, Inc. Focused Feasibility Study Port Ludlow, Washington

	AUG-2013	BTC/SCC	FIGURE NO.
CONSULTING	PROJECT NO. 130046	REV BY: -	2

CAD



	Testing Data Summary									
Sample		Days of continuous SVE operation with groundwater removal		Laboratory Result for GRO in mg/meter ³	GRO Removal Rate (Ibs/day)					
Test2-Sample2	12/13/2011	0.375	18	1900	3.07					
Test4-Sample2	1/5/2012	2	75	180	1.21					
Test4-Sample1	12/29/2011	7	53	30	0.14					

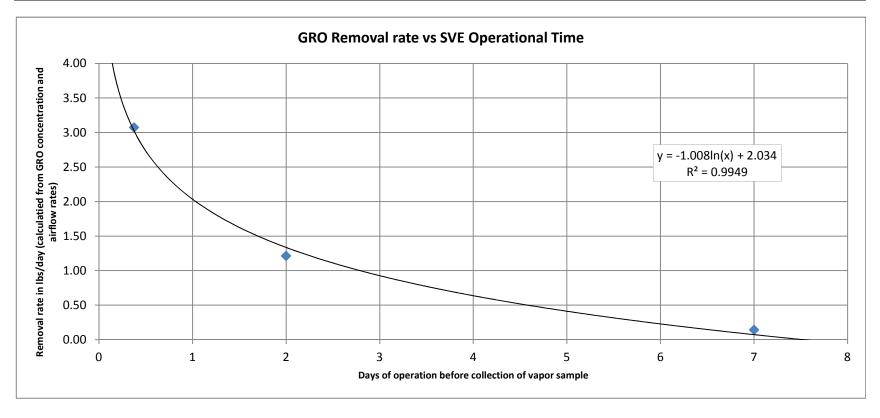


Figure 4 - Analysis of Mass Removal Rate versus Time -SVE Pilot Testing with Groundwater Extraction (DPE, Test 4)

DRAFT

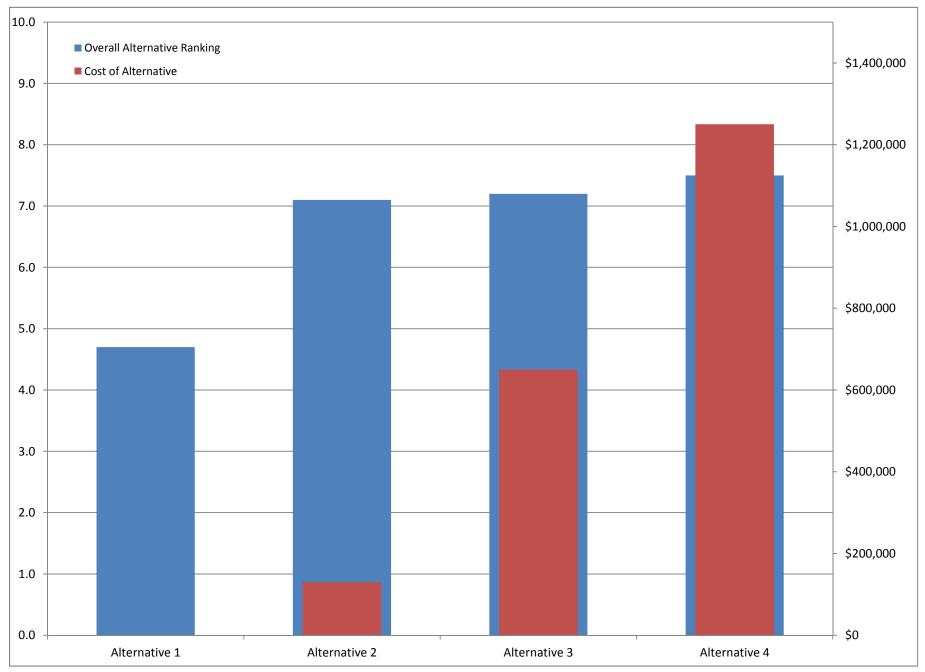


Figure 5 - Disproportionate Cost Analysis Summary

Olympic Water and Sewer, Inc. Site

APPENDIX A

Water Production Well Logs

Table C-1 Water Well Supply Well Construction Details Olympic Water and Sewer, Inc. Property Port Ludlow, Washington

SLR								Total	Casing				
Designated						Well Owner's Name		Depth	Depth	Screen Depth	Open	StaticWater Level	Producing
Number	Quarter		Section	Township	Range	(at time of installation)	Well Use	(ft. bgs)	(ft. bgs)	(ft. bgs)	(ft. bgs)	(ft. bgs)	Formation
1	NW	SE	8	28N	1E	Blaine Shaffer	Domestic	208	0-203	203-208	NA	108	Sand and Gravel
						Pope & Talbot			0-214;	214-224;			
2	SW	SE	8	28N	1E	Development, Inc.	Domestic	245	224-240	240-245	-	69.5	Sand and Gravel
						Pope & Talbot							
3	SE	NE	8	28N	1E	Development, Inc.	Domestic	257	241-257	0-241	-	144.5	Sand and Gravel
									0-315.5;				
						Pope & Talbot			329.7-	315.5-329.7;			Sand and Gravel;
4	SE	SW	8	28N	1E	Development, Inc.	Domestic	546	361.3	361.3-377.1	-	158.9	Pebble Conglomerate
5	NW	SE	8	28N	1E	Richard Werner	Domestic	157	0-157	None	157	118	Gravel
6	NW	SE	8	28N	1E	Ross Witter	Domestic	176	0-176	None	176	136	Sand and Gravel
7	NW	SE	8	28N	1E	Ross Witter	Domestic	178	0-178	None	178	133	Sand and Gravel
8	NE	SW	8	28N	1E	Ruth Altis	Domestic	211.5	0-211.5	None	211.5	191	Sandy Clay and Gravel
9	NE	SW	8	28N	1E	Chris Baschab	Domestic	276	0-270	270-276	-	226.5	Sand and Gravel
10	NE	SW	8	28N	1E	Frank Woodruff	Domestic	193	0-193	None	193	114	Gravel
11	S 1/2	NE	8	28N	1E	R.T. Moran	Domestic	290	0-285	285-290	-	201	Sand and Gravel
12	NW	SE	8	28N	1E	John Werner	Domestic	205	NA	NA	NA	142.65	NA
Notes: The well con NA= Informa	struction tion not a	details ir wailable.	n this table	e are basec	d on Wasi	hington Department of E	cology or C)lympic W	/ater & Sev	wer, Inc. record	5.		

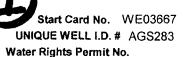
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File original	and first copy with
Department	of Ecology

Second Copy - Owner's Copy

WATER WELL REPORT

STATE OF WASHINGTON



Third Copy - Driller's Copy		Water Rights F	'erm
(1) OWNER: Name: BLAINE SHAFFER	Addr	ess: 425 SHINE RD., PORT LUDLOW, WA 94	8365
(2) LOCATION OF WELL: County: JEFFERSON		W 1/4 of the SE 1/4, Sec. 8, Twnsp 28 N,	
(2a)STREET ADDRESS OF WELL (or nearest address			
(3) PROPOSED USE: X Domestic Test well			
Irrigation		(10) WELL LOG or	
Industrial Dewater Other		ABANDONMENT PROCEDURE D	ESC
(4) TYPE OF WORK: Owner's number of well(if more than	1 one): 1		
Abandoned X New well Method: Dug	Bored	Formation: Describe by color, character, size	
	Driven	structure, and show thickness of aquifers and of the material in each stratum penetrated, wit	
Reconditioned Rotary	Jetted	each change of formation.	
(5) DIMENSIONS: Diameter of well: 6 inches	1 00000	MATERIAL	
	208 feet		ا
(6) CONSTRUCTION DETAILS:	.00 .001	BROWN HARDPAN	1
Casing Installed: 6 in. diam. from 0 ft. to	203 ft.	BROWN SAND & GRAVEL	
Welded in diam. from ft. to	200 ft.	BROWN SAND & GRAVEL	
Liner installed in. diam. from ft. to	ft.	GRAY HARDPAN	
Threaded in diam. from ft. to	ft.		
Perforations: Were perforations made?: Yes	16.	BROWN CLAY	
Perforator type:		BLUE CLAY	
Size of perforation: in. by; in.		BROWN CLAY	1
Perforations from: ft. to ft.			÷
Perforations from: ft. to ft.		GRAY SAND W/B	
Screens: X Screens installed		BROWN CLAY	
Manufacturer's name: Johnson		GRAY SAND & GRAVEL W/B	
Type: Telescoping	000 #		
Diam. 6 in. slot size .016 from 203 ft. to Diam. in. slot size from ft. to	208 ft. ft.		1
Gravel: Yes Size of gravel: in. Gravel Placed from: ft. to ft.			
Surface seal: XYes To what depth?:	18 ft.		
	18 ft	RA	
Material used in seal: Bentonite Did any strata contain unusable water 1 Yes			
Type of water: Depth of strata:	ft.	A11-	1
Method of sealing strata:	ц.	nug ne	
		RECE AUG 05 DEPT OF ECC	201
		UFFO	0
Type: Submersible HP: 1 (8) WATER LEVELS Land-surface elevation			えつ
(8) WATER LEVELS Land-surface elevation above mean sea-level: 310	ft.		
	/21/2005		
Artestian pressure: PSI Date:	21/2005		
Artesian water is controlled by:			
(9) WELL TESTS Drawdown is amount water level is lowered below	v static level.		
Was a pump test made? X Yes, by: Don Lofall		Work started: 6/13/2005 Work com	nplet
Yield: 15 GPM with 15 ft. drawdown after	1.5 hrs		
Yield: GPM with 15 ft. drawdown after	hrs	WELL CONSTRUCTOR CERTIFICATION: I constructed and / or accept responsibility fo	or cor
Recovery data (time taken as zero when pump turned off) (water level	1113	well, and to the compliance with all Washingt	
measured from well top to water level)		standards. Materials used and the information	on re
Water Water	Water	true to my best knowledge and belief.	
Time level Time level Time	level	Contractor: LOFALL WELL DRI	111
10 min. 108'		Contractor: LOFALL WELL DRI	LL
		Address: 180 NW Lofall Rd, Poulsbo	э, W
Date of test: 6/21/2005			
Bailer test: 20 GPM with 28 ft. drawdown after		Signed: Kong of Douly	Licer
Air test: GPM with stem set at ft. for	hrs.	(Well Driller)	
Artesian flow: GPM Date:		Contractorie	

16868

Was a chemical analysis made? 🕱 Yes Temp. of water: deg.

0) WELL LOG or BANDONMENT PROCEDURE DESCRIPTION

of the SE 1/4, Sec. 8, Twnsp 28 N, R. 1 E W.M.

nation: Describe by color, character, size of material and cture, and show thickness of aquifers and the kind and nature e material in each stratum penetrated, with at least on entry for h change of formation.

MATERIAL	FROM	то
BROWN HARDPAN	0'	67'
BROWN SAND & GRAVEL	67'	102'
GRAY HARDPAN	102'	108'
BROWN CLAY	108'	125'
BLUE CLAY	125'	144'
BROWN CLAY	144'	189'
GRAY SAND W/B	189'	192'
BROWN CLAY	192'	199'
GRAY SAND & GRAVEL W/B	199'	?



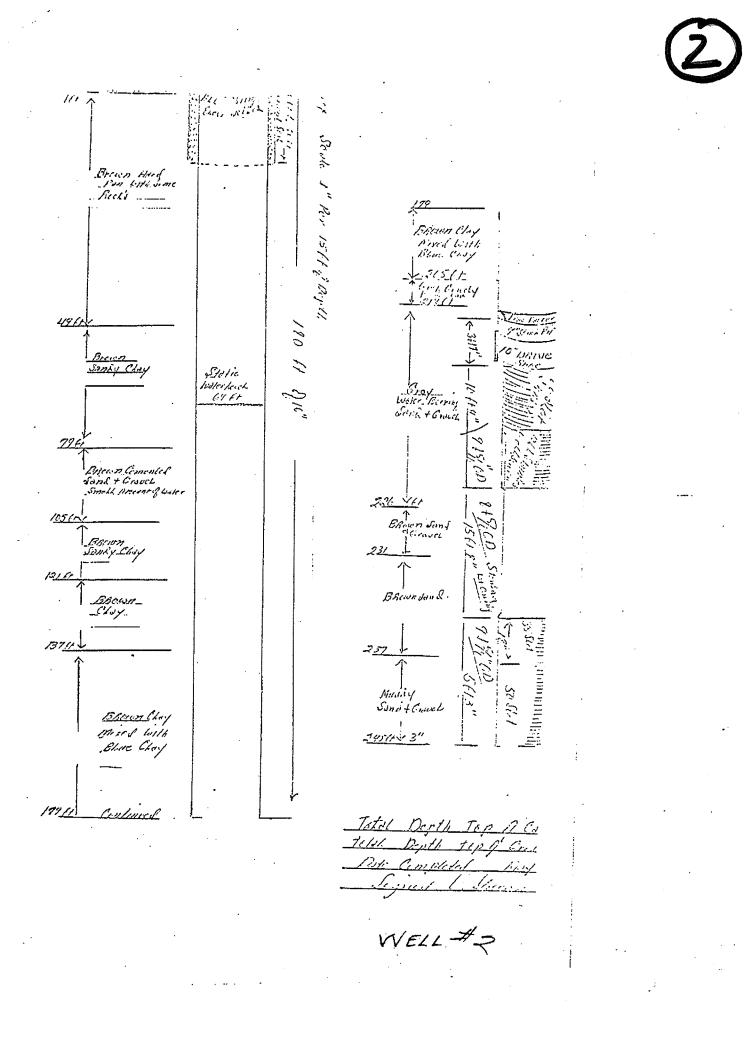
Nork started: 6/13/2005 Work completed: 6/20/2005

onstructed and / or accept responsibility for construction of this I, and to the compliance with all Washington well construction ndards. Materials used and the information reported above are to my best knowledge and belief.

Contractor: LOFALL WELL D	RILLING
Address: 180 NW Lofall Rd, Poul	sbo, WA 98370
Signed: Kaymond (Well Driller)	License # 1463
Contractor's Contr	Date: 7/12/2005

((ATE OF WASHINGTON (DEPARTMENT OF CONSERVATION DIVISION OF WATER RESOURCES つん WELL SCHEDULE No 8-2 Date Record by Source 1. Location: State of WASHINGTON Jef County_ Агез Map. DIAGRAM OF SECTION <u>SV SE</u> **1**4 14 Details - 14 4 Offia No. 2. Owner or Te Address 3. Driller γĆ ibor Land-surface datums Topography: thi ... Rept 5. Type: Dug : Drilled -De Jetted Mei Bored Data_ ጵ" 6. Casing: Diam Depth... ft. Finish ff. to 7. Chief aquifér(s): 8. Water levels Rept abova batov Maai bove which datum Capacity mi 9. Pump: Type Size horsepowe Driven by_ gal. min. Pump 160 gal. min. Mess (Rept) Est 10. Yield: Flow_ 2.5 ft. after. mi. mh hours pumping Drawdown. Adequacy, permanence 11. Use: Dom., Stock. PS. Ind. Irr. Obs. • . • 12. Quality: Sample No..... Temp. 19. . - • -• Taste, color, hardness, sanitation, etc. 13. Other datas Log Water levels Draft Pump test Analyses. Тит ир · . : : ----

LOCATION: $T 28$ R 1E BeC. B 1/4 1/4 <u>KQ</u> 13 LAND SUPFACE <u>300</u> <u>1/3</u> DEPTH <u>245</u> DATE <u>6000</u> ELEVATION <u>300</u> <u>1/3</u> DEPTH <u>245</u> DATE <u>6000</u> WELL OWNER: <u>PORE TRANSOF DEVEL OPMENT</u> <u>1/NC</u> . OWNERS DESIGNATION <u>WELL 2</u> , <u>N</u> UBE DRILLING DRILLING <u>BOURCE <u>R+N</u> 01-B2 METHOD <u>CACLE</u> DRILLING DRILLING <u>314-22.45</u> Sec. COMPLETION MODE <u>SCREEN</u> COMPLETION ZONE (B) <u>214-22.45</u> YIELD <u>158</u> <u>A</u> SPECIFIC CAPACITY <u>3.0</u> <u>A</u> SWL <u>67.2</u> <u>1/5</u> DATE <u>11/1968</u> SWL <u>67.2</u> <u>15,000</u> <u>B</u> STORAGE COEFFICIENT <u>5,000</u> <u>B</u> TOTHER WATER BEARING ZONES PENETRATED MAJOR AQUICLUDES PENETRATED <u>1/4 1/4 KQ</u> OTHER WATER BEARING ZONES PENETRATED <u>VER.CODE</u> <u>1/4 1/4 KQ</u> POST CONSTRUCTION SWL MEASUREMENTS WITH DATES <u>67.0</u> <u>B</u> <u>11/71</u> <u>47.2</u> <u>1/8</u> <u>0/20/75</u> <u>VER.CODE</u> <u>VER.CODE</u> PUNP TEST DATA FILE <u>9000001C LOG <u>SEE CACK</u> WATER CHEMISTRY REMARKS <u>11/71</u> <u>Q/A</u> = 4.0 <u>C</u> <u>1 MAS AND 160 gpm</u> <u>11/71</u> <u>TEMP</u> <u>49.1/2</u> <u>0H = 7.4</u> <u>MARD</u> <u>70</u> <u>FC = 0.3</u> <u>H.S.5 5C/0</u> <u>781</u> <u>MAIKEN</u> <u>481</u> <u>781</u> <u>WAIKEN</u> <u>WAR</u> <u>781</u> <u>WAIKEN</u> <u>WAR</u> <u>781</u> <u>WAIKEN</u> <u>WAR</u> 11</u></u>	,	PORT LUDLOW REGION
AND BUPFACE 300 13 DEPTH 245 DATE 6000 1940. ELEVATION		REFERENCE NO. 10 Well Data Summary
WELL OWNER: <u>POPE TRUGGE</u> <u>DEVELOPMENT</u> , <u>INC</u> OWNER'S DESIGNATION <u>INELL 2, N</u> UBE DRILLING <u>DRILLING</u> <u>DRILING</u> <u>DRILLING</u> <u>DRILING</u> <u>DRILNG</u> <u>DRILLI</u>		LOCATION: T 28 R 1 8 800. 8 1/4 1/4 RQ DATE YEA
OWNER'S DESIGNATION $(NELL 2, N)$ UBE		ELEVATION 300 BEPTH 245 DRILLED 1968
INFORMATION BOURCE $R+N$ $01-B2$ DRILLING $METHOD$ DRILLED BY $STOICAN$ CASING BIZE (B) $1/2$ $342 - 224$ COMPLETION MODE $SCREEN$ COMPLETION ZONE (B) $240 - 245$ VIELD 15.8 M SPECIFIC CAPACITY 3.0 M WIELD 15.8 M SPECIFIC CAPACITY 3.0 M aduifer 47.2 17.6 DATE $M/96.8$ aduifer TANBMISSIVITY 15.000 M BWL 67.2 17.6 M aduifer TANBMISSIVITY 15.000 $VER.CODE$ 114.114 $K.Q$ aduifer TAND TAND 114.114 $K.Q$ 114.114 $K.Q$ aduifer TAND TAND 114.114		WELL OWNER: POPE TALBOT DEVELOPMENT, INC.
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GEOLOGIC LOG <u>SEE CACK</u> WATER CHEMIBTRY REMARKS <u>11/71: Q/A = 4.0 @ 1 HRS AND 160 gpm</u> <u>11/71 TEMP 49.1/2 pH=7:4 , HARO = 70 , Fe = 0.3 , H.S = SLIG</u> <u>Ma = .16</u> <u>181 Walker Way</u>		
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REWARKS <u>$11/71$ Q/A = 4.0 @ 1 HRS AND 160 gpm</u> . <u>$11/71$ TEMP 49.1/2</u> <u>$DH = 7:4$</u> <u>$HARO = 70$</u> , <u>$Fe = 0.3$</u> <u>$H_2S = SLIG$ <u>$M_{R} = .16$</u> <u>781 Walker</u> Way</u>		
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$\frac{11/71}{M_{R} = .16} = \frac{16}{781} = \frac{16}{181} = \frac{16}{181} = \frac{16}{181}$	1	EMARKS 11/71. Q/A = 4.0 @ 1 HRS AND 160 gpm.
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	-	HII TEMP 49.12, H=7:4, HARO = 70, Fe = 0.3, H2S=SLIG
	-	$M_{R} = .16$
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Add	Alternation of the second static level if report Autor describe driller's terminology literally but faraphense as no al water-bearing, so state and record static level if report d-surface dutum unless otherwise indicated. Correlate with Following log of materials, list all casings, perforations,	From (feet)	To (feet) parentheses. 		· · · · · · · · · · · · · · · · · · ·
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Add and su WL: Corre- LATION (Train T materia	Attended in the second static level if report Autorial of the second static level if report d-surface dutum unless otherwise indicated. Correlate with Following log of materials, list all casings, perforations, Community domestic supply hardpan clay, sand, gravel hardpan	From (feet) ccssury, in cd. Give de h stratigram s.rcons, etc 0 34 39	To (teet) parenthuses. pths in feet white column, 34 39 48		
Add and su WL: Corre- LATION (Train T materia	Andrews and, gravel, some clay binder	From (feet) c.ssury, in cd. Give de h stratigram s.rcons, etc. 0 34 39 48 57	To (teet) parentheses. pths in feet whic column, 34 39 48 57 80		
Add and su WL: CORAB- ATION (Train materia	Artenation of water 80-82.	From (feet) c. ssury, in cd. Give do h stratigram screens, etc. 0 34 39 48 57 80	To (Teet) parentheses pths in feet thic column, 34 39 48 57 80 128		
Add and su WL: Corre- LATION (Train T materia	Artenation of water 80-82. clay, silt & sandy wet peat	From (feet) c.ssury, in ed. Give de h stratigra or cons, etc. 0 34 39 48 57 80 128	To (teet) parentheses pths in feet white column, 34 39 48 57 80 128 142		
Add and su WL: Corre- LATION (Train T materia	rface, datum	ims : 81 From (feet) c	To (Teet) parenthoses. pths in feet shic column, 34 34 39 48 57 80 128 142 162		
Add and su WL: Corre- LATION (Train T materia	Artenation of water 80-82. clay, silt & sandy wet peat clay, silt & sandy wet peat clay, silt & some sand	ims	To (feet) parentheses poths in feet poths in feet 34 39 48 57 80 128 142 162 186		
Add and su WL: Corre- LATION (Train T materia	Artenation of water 80-82. clay, sand hardpan clay, sand clay, sand clay, sand clay, sand clay, sand clay, sand clay, sand clay, sand hardpan clay, sand clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, some clay binder small amount of water 80-82. clay, silt & sandy wet peat clay, some gravel & peat clay, some gravel & peat	ims : 8 ¹¹ From (feet) c. 3 sury. in cd. Give de h stratigra s. r cons, etc. 0 34 39 48 57 80 128 142 162 186	To (teet) parentheses. pths in feet whic column. 34 39 48 57 80 128 142 162 186 214		
Add and su WL: Corra- LATION (Trai (Trai	rface, datum	ims : 8 ¹¹ From (feet) c	To (teet) parentheses. pths in feet shic column, 34 34 39 48 57 80 128 142 162 186 214 223		
Add and su WL: Corra- LATION (Trai (Trai	Artenation of water 80-82. clay, sand hardpan clay, sand clay, sand clay, sand clay, sand clay, sand clay, sand clay, sand clay, sand hardpan clay, sand clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, sand hardpan clay, some clay binder small amount of water 80-82. clay, silt & sandy wet peat clay, some gravel & peat clay, some gravel & peat	ims	To (feet) parentheses poths in feet poths in feet 34 39 48 57 80 128 128 142 162 186 214 223 235		
Add and su WL: Corra- LATION (Trai f materi	rface, datum	ims : 8 ¹¹ From (feet) c	To (teet) parentheses. pths in feet shic column, 34 34 39 48 57 80 128 142 162 186 214 223		

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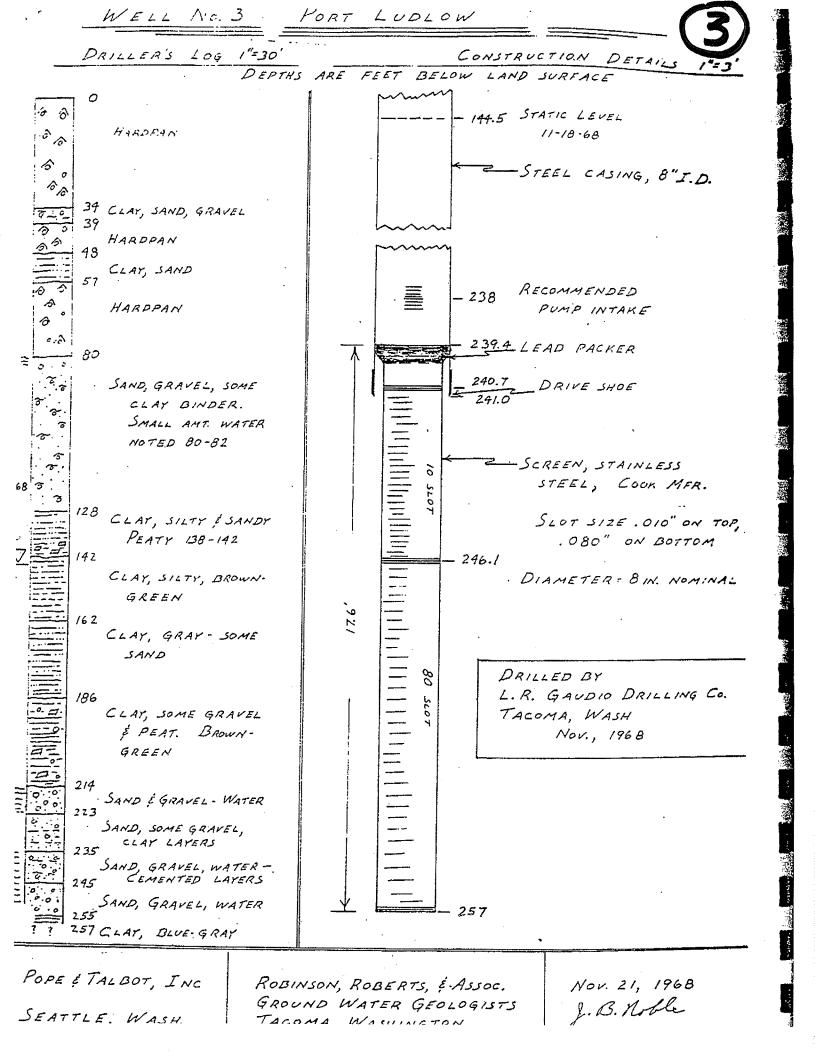
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C	ORE.		·····/ ·····	
	ATION	MATERIAL	From (feet)	To (feet)
:		Deptb forward		
		sand, gravel water		
		clay, blue grey	245	255
		Casing: 8" from Ol	255	257
) —		COOK granini.		
·		o slot size IO from 241		
				46
				57
· · · · · · · · · · · · · · · · · · ·				
		Pump: Gould, submersible 20 H	after]	hrs
.		20 H		[
				F
				ft
	+			
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· · · · · · · · · · · · · · · · · · ·	╞╧╧			f
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	· · · ·		4.2	
	<u> </u>			
S. F. No. 744	9-09	5-12-65		
- 195 - E				

PORT LUDLOW REGION	3
REFERENCE NO. 9 Well Data Summary	
LOCATION: T28 R 1F Boc. 8 1/4 1/4 HI B	
LAND SURFACE 380 B DEPTH 257 DATE CODE	5-
WELL OWNER: POPE AND TALBOT DEVELOPMENT INC	
OWNER'S DESIGNATION WELL'S, N USE	
INFORMATION BOURCE R+N 01-82 METHOD CABLE	
DRILLED BY GRUDIO CASING SIZE (8) 8	
COMPLETION MODE SCREEN COMPLETION ZONE (6) 241-257	•
VIELD 88 A SPECIFIC CAPACITY 2.3 A	
YIELD 88 A SPECIFIC CAPACITY 2.3 A SWL 144.5 15 DATE 11/18/68 Code SWL 144.5 15 DATE 11/18/68 Code	
AQUIFER TRANSMISSIVITY 6000 11/1971	
BTORAGE COEFFICIENT 1/4 1/4 // VER.CODE 1/4 1/4 /// OTHER WATER BEARING ZONES PENETRATED 1/4 1/4 ///	
MAJOR AQUICLUDES PENETRATED	
POST CONSTRUCTION SWL MEASUREMENTS WITH DATES 152 3 11/71	•
VEN. CODE	
156.5 <u>B</u> 8/21/75 VER. CODE VER. CODE VER. CODE	
DETAILED SUPPLEMENTARY FILES	
PUMP TEBT DATA FILE R+N FILE	
$a_{2}a_{3}a_{3}a_{3}a_{3}a_{3}a_{3}a_{3}a_{3$	
GEOLOGIC LOG <u>SEE BACK</u>	
WATER CHEMISTRY TEMP = 51 F, pl-1 = 7.5, HARD = 102, Cl = 6 Fe =	0.4 , H2S . 000R
REMARKS 11/71 Q/2 = 2.0 @ 107 gpm @ 1 HK.	STRONG
11/71 Fe = 0.1, H2S= SLIGHT, Mn = .13	
11/08 CL = 6	
-tali i u	
Talbot Way	
RECORD BY: MGS ROBINSON & NOBL	F INC



PORT LUDLOW REGION	<u>a</u>
EFERENCE NO. 8 Well Data Summary	
EFERENCE IC: 0 ALE BAC & 1/4 1/4 P B	-1 1
AND SURFACE 340 TOPO DEPTH 546 DRILLED 1980	l 1
IELL OWNER: POPE AND TALBOT DEVELOPMENT, INC.	L L
NFORMATION SOURCE R+N 79-81 METHOD CABLE	
	0 5607
COMPLETION MODE SCREEN COMPLETION ZONE (6) 361.3-377.1 10 + 30	SLOT
TIELD 94 A SPECIFIC CAPACITY 1.7 A VER CODE CODE CODE CODE CODE CODE CODE CODE	
QUIFER TRANSMISSIVITY 3030 + 4280	
	C
TORAGE COEFFICIENT VER.CODE	
THER WATER BEARING ZONES PENETRATED	
	. •
AAJOR AQUICLUDES PENETRATED	<u> </u>
POST CONSTRUCTION SWL MEASUREMENTS WITH DATES 154 A 5/8/80	• .
VER. CODE	
VER. CODE VER. CODE VER. CODE	
VER. GUDL	
DETAILED SUPPLEMENTARY FILES	
<u>DETAILED SUPPLEMENTARY FILES</u> PUMP TEST DATA FILE <u>R+N 79-81</u> W/ <u>RECOVERY HYDRGRAPH</u>	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-B1 W/ RECOVERY HYDRGRAPH</u> GEOLOGIC LOG <u>SEE BACK</u>	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-B1 W/ RECOVERY HYDRGRAPH</u> GEOLOGIC LOG <u>SEE BACK</u>	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-81 W/ RECOVERY HYDRERAPH</u> GEOLOGIC LOG <u>SEE BACK</u> WATER CHEMISTRY <u>R-N 79-81</u> OK	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-81</u> <u>W/ RECOVERY HYDRGRAPH</u> GEOLOGIC LOG <u>SEE BACK</u> WATER CHEMISTRY <u>R-N 79-81</u> , <u>OK</u> REMARKS <u>A A DANN</u> (Westurg "Extension"	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-81</u> <u>W/ RECOVERY HYDRGRAPH</u> GEOLOGIC LOG <u>SEE BACK</u> WATER CHEMISTRY <u>R-N 79-81</u> , <u>OK</u> REMARKS <u>A A DANN</u> (Westurg "Extension"	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-81</u> <u>W/ RECOVERY HYDRERAPH</u> GEOLOGIC LOG <u>SEE BACK</u> WATER CHEMISTRY <u>R-N 79-81</u> <u>OK</u>	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-81 W/ RECOVERY HYDRGRAPH</u> GEOLOGIC LOG <u>SEE BACK</u> WATER CHEMISTRY <u>R-N 79-81</u> , OK REMARKS <u>A A DANN</u> (Westury "Extension"	
DETAILED SUPPLEMENTARY FILES PUMP TEST DATA FILE <u>R+N 79-81</u> <u>W/ RECOVERY HYDRGRAPH</u> GEOLOGIC LOG <u>SEE BACK</u> WATER CHEMISTRY <u>R-N 79-81</u> , <u>OK</u> REMARKS <u>A A DANN</u> (Westurg "Extension"	

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spartment of Ecology word Copy - Owner's Copy	WATER WE	IT RELAKI.	whereare	1101	
wond Copy - Owner's Copy urd Copy - Driller's Copy	STATE OF W	ASHINGTON	Permit No.	• • • • • • • • • • • • • • • • • • •	[{
1) OWNER: Name Pope & Talbot Dev	elopment.Inc.	Address P. 0. Box 75.	Port Ludlow, WA	98365	
	Jefferson		_SW 14 Sec. 8 - 72	and the second second second second	and the second
ng and distance from section or subdivision corn	er				
3) PROPOSED USE: Damestic [] Industr	ial 🗋 Municipal	(10) WELL LOG:			835
Irrigation 🔲 Test W		Formation: Describe by color show thickness of aquifers or	, character, size of materi	al and struc	ture, and
4) TYPE OF WORK: Owner's number of w	eil as	show thickness of aquifers an stratum penetrated, with at	la the kind and nature of least one entry for each	the materia change of fi	i in each prinction.
(if more than one) New well Mathod: D	C SAMPLE CONTRACTOR AND	MATE		FROM	TO .
	able Driven	brown to gray til		0	52
Reconditioned C R	otary [] Jeited []	brown, sandy, peb	bly silt	52	131
5) DIMENSIONS: Dismeter of well	12 inches	sand and gravel		131	151
Drilled 546 ft. Depth of completed	0001	hrown silty sand	and gravel	151	
		gray sandy silt		157	171
6) CONSTRUCTION DETAILS:	•	gray provide sorte	d silty sand	+1/1	267
	1_ tt. to <u>3154</u> m.	and dravel	<u>a su cy saua _</u>	267	269
Threaded [] <u>10</u>	125. n. to 361 ft.	gray pebbly silt	••••	269	314
		brown, noorly sor	ted medium to:		
Perforations: Yes Ci No X		coarse sand and		314	332
Type of perforstor used	*****	aray pebbly silt		332	360
SIZE of perforations in, b;		gray sand and gra	vel	360	366
perforations from		grav silty sand w		366	380:
*	ft, to ft,	gray clayey silt		380	536
		weathered nebble	conglomerate ·	536	544
Screens: Yes X1 No D Manufacturer's Name UOP Johnson		gray pebble congl	omerate (bedroc	k) 544	546
Type PiDe Size	N No 30455				
Type Pipe Size Diam 10 Slot size .080- 100 31	5 H to 3293 H	· · · · · · · · · · · · · · · · · · ·	,		-
Diem. 10" Slot size .080- 1101 36	1_ st. to 378_ st.	****	• •,• •. • • • •		
Gravel packed: Yes O No XI Size of g					
Gravel placed from ft. to	ravel: ft.				·*
		PFCEIV	1 ····		
Surface seal: Yes & No C To what de	pth? _20 ft.				
Material used in seal	¥	ITIN 0.9.19	80		
Did any strata contain unusable water? Type of water?	Yes [] No []	<u>JUN 09 19</u>	00		
Method of sealing strate off		DEPARTMENT OF E	COLUGY	· _	
		SOUTHWEST REGION	AL OFFICEIN 5 19	80	
7) PUMP: Manufacturer's Name		- SOUTHWEST REGION			· · · ·
Тура:			UEPI. OF COU	- Har	•
8) WATER LEVELS: Land-surface elevation	ion 372 +		nill i ni	den	·
tatic level158.9ft. below top of well	Date 4/28/80	11 jaton F	STATIS LIDIT	کل	
rtesian pressure			Mer err		
Artesian water is controlled by	p. valve. etc.)				
				•	
9) WELL TESTS: Drawdown is amount lowered below staric	level Ropinson,	Work started 3/18	19.80 Completed	/29	1980
Vas a pump test made? Yes [Å. No [] If yes, by wh right: 94 gal./min. with 50.6 ft. drawdown	om: Noble Carr	WELL DRILLER'S S			
reid: 94 gal/min. with 50.6 ft. drawdowr	a after [/, 0/hrs.				
14 IV		This well was drilled true to the best of my l	under my jurisdiction	n and this	report is
anuen data (time taken as tern when numn tum	and AAL (mater Jassa)				
Recovery data (time taken as zero when pump turn measured from well top to water level)		NAME Story/Arms	trong Drilling		
Time Water Level Time Water Level Ti 10 min. 172.6 50 168.35 15	ime Water Level 35 164.8	(Person. 5	in, or corporation)	(Type or p	rint)
20 " 170.95 80 166.85 22		10711-1074	5 66th Ave. E.		
³ Ω " 169.8 100 166.1	A.T	Address Puyal Tup',	WA 98371 7.	1 2 2 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
Date of test4/28/80 - 4/29/80	******	Low Low	- / - Stan	- 5	
.er testgal/min. withft. drawdow	wn after	[Signed]	(Well Driller)		
ricsian flowg.p.m. Date		1		10 30	
Camperature of water	made? Yes 💭 No 🗍	License No	Date	Z.J	, 1992
	•	• •			
(USE ADDITIONAL SE	HEETS IF NECESSARY)			

Second Copy — Owner's Copy	CLL REPORT Application VASHINGTON Permit No.		
(1) OWNEB: Name Sichard Werner	11,1,1, - 10,1		
	- Adares 2 NUL 1/2 SE	8	Ew M
(3) PROPOSED USE: Domestic & Industrial I Municipal I Irrigation I Test Well I Other I	(10) WELL LOG: Formation: Describe by color, character, size of materia show thickness of aquifers and the kind and nature of stratum penetraied, with at legat one entry for each of	the materi	at in each
(4) TYPE OF WORK: Owner's number of well (if mgre Man one)	MATERIAL	FROM	TO
New well 🔄 Method: Dug 🔲 Bored 🗋	Bon landal Class	0	8
Despend Cable Driven Reconditioned Reconditioned	the state of the s		
	Bron Sandy land pan	8	20
(5) DIMENSIONS: Diameter of well inches. Drilled 57 tt. Depth of completed well 57 tt.	The second second		
Drifted J. J. J. Depth of completed water jugar	Bandandy lay	20	<u> 45</u>
(6) CONSTRUCTION DETAILS: Casing installed: <u>Diam. trom</u> <u>One</u> tt. to <u>157</u> tt.	graf sandy Clay	25	35
Threaded Diam. from	grof landy leftgrond HP.	35	45
Perforations: Yes 🗆 No 🖅	may and Clay	45	55
Type of perforator used	pt July		10 0
perforations from	Andendy Clay	32	100
perforations from	gray saddy lay	100	135
Screens: Yes D No C	UB. portant lay	135	155
Manufacturer's Name Model No	st & A. A.		1.000
Diam	Carrier Manuel	12-	12/
Diam			
Gravel packed: Yes D Not Size of gravel:	3		
Gravel placed from ft. to ft.	X		
Surface seal: Yes p No T Athat septin 2 Q			
Material used in sea full al loff Sintomitte.			
Did any strata contain unusable water? Yes in No is Type of water?			+
Method of sealing strate of			
(7) PUMP: Manufacturer's Name			ļ
Type:			
(8) WATER LEVELS: Land-surface elevation #30 #		+	+
(a) WAIED LEVELS. above mean sea level. 42.			
Artesian pressure			
Artesian water is controlled by (Cap, vaive, etc.)			
(9) WELL TESTS: Drawdown is amount water level is lowered below static level	Work started 2-13 1985. Completed 2	-27	1084
Was a pump test made? Yes [] No [] If yes, by whom?			
Yield: gal./min. with ft. drawdown after hrs.	WELL DRILLER'S STATEMENT:		
	This well was drilled under my jurisdiction true to the best of my knowledge and belief.	and this	report is
Recovery data (time taken as zero when pump turned off) (water level measured from well top to water level)	NAME RICHAND BEKKE	VD)	
Time Water Level Time Water Level Time Water Level	(Person, firm, or corporation)	(Type or	print)
	Address 2722 # Wy 10/85	egu	(m. 6-
	p.14p.		
Date of test	[Signed] flchand to kt	ena	u
Bailer test /2. f. gal/min. with /0. tt. drawdown after / 2. hrs. Artesian flow		\$ 1	0.
Temperature of water	License Nof		, 19 0

hird Copy—Driller's Copy		STAT	TE OF 1	NASHINGTON						
		_				ht Permit I				
1) OWNER: Name_Ko	ss witter	R		Address []] En	1 bodi	, RD.	Part	hudl	OLA, La
2) LOCATION OF WELL	· County JUFF	P.P.S. D.N.			NI	1.55	X Sec.	- + !	24	PIFWN
(a) STREET ADDDRESS O	-					N A.C () I 6	A.	
							UENT DO			
3) PROPOSED USE:	Irrigation	itriel 🗌 Munici Well 🗌 Other		(10) WELL LOG or ABANDONMENT PROCEDURE DESCRIPTIO						
				thickness of aquife with at least one en	e and the k	ind and nati	ure of the ma			
() (I)						ATERIAL			FROM	то
Abandoned 🗌 New we Deepen			red 🗌	CAY, SANC	L.G.K	avely	/		0	10
Record	tioned 🗌		led 🗋	HARDDAN	<u>/'</u>	· /			10	30
5) DIMENSIONS: Diamet	er of well		inches.	clay, SAN	- / /	Brely			30	60
Drilled 196 test.	Depth of complete	ed well17	<u>16.</u> n.	clay, san	-7	eaveli			130	11.5
) CONSTRUCTION DET				Sand GR		L H H	6		165	126
Casing Installed;	L · Diam. from	<u>- /17</u>	<u>6</u> n.		<u></u>				100	
Liner Installed		It. to								ļ
Threaded			ft.						<u> </u>	<u> </u>
	No						. <u> </u>	_1		
Type of perforator used		Y		•						
SIZE of perforations				· · · · · · · · · · · · · · · · · · ·						1
						8	Ð			
	ns from		łt.		s Ì:		'n		<u> </u>	
Screens: Yes Not					<u></u>			-		<u> </u>
Manufacturer's Name					in in z		1771			
Туре	from	Model No		-		\sim				+
Diam,		ft. to	ft.		<u> </u>					
Gravel packed: Yes						:5				
Gravel placed from)				2				
Surface seal: Yes	To what depti	h2 / {	<u>н.</u>							
Material used in seal	ATTALITE + US	W DRillings								
Did any strate contain unusab		No					<u></u>			1
Type of water?		Depth of strats		·····	^ 		·····		1	1
Method of sealing strats off							· · · · · · · · · · · · · · · · · · ·			
) PUMP: Manufacturer's N	IM#	<u></u>								
Туре:	<u></u>	H.P			_			<u></u>		
WATER LEVELS:	Land-surface elevation above mean sea level .		2-4 n.							
Statio ievel 2 6			-70							+
•										1
		(Cep, velve, etc.))		Work started	1-5		Rompiete	1_/=	16-	192
WELL TESTS: Drawd Was a pump test made? Yes				WELL CONS	TRUCTO	D CERTI		4-		
Yield:				I constructe					etruction of	of this wel
	**	31		and its cor Materials u	noliance v	vith all W	ashington.	well co	nstruction	standard
Recovery data (time taken ar		· · · · · · · · · · · · · · · · · · ·	eured	knowledge a						,
from well top to water level) Time Water Level	Time Water Level	Time Wate	wLavai	NAME MA	berry	1 Wel	1 DR.	llin	4	
			<u></u>	2	(PERSON	FIRM. OR CO		1		OR PRINT)
				Address A	BOX	255	Alui	Lleni	1. C. (J. (2,903
Date of test				1 1.5	1. 1	$2 \cdot l$	m.l.		11	10
Bailer test gai./r	nin, with _15 n	drawdown atter	<u> <u>Yr</u> hre.</u>	(Signed)	(WELL	DRILLER)	(All and a start of the second start of the se	License	No.	
				Contractor's		-				
Airtest	. with etem set at	fl. for	hre.	Redetration Not BEE	N		· · /	1.		9

(USE ADDITIONAL SHEETS IF NECESSARY)

File Original and First Copy with Department of Ecology Second Copy — Owner's Copy Third Copy — Driller's Copy			
A Marthalette	At Dave Al		9821
(1) OWNER: Name, County County (2) LOCATION OF WELL: County Count	- Address ICT 1 Dof 3 92 Clima - With vs S. E. V. Sec. B. 72		<u>Г</u> Ен.м.
(3) PROPOSED USE: Domestic [] Industrial [] Municipal [] Irrigation [] Test Well [] Other []	(10) WELL LOG: Formation: Describe by color, character, size of materi thow thickness of aquifers and the kind and nature of	the mater	al in each
(4) TYPE OF WORK: Owner's number of well (1) more than one) New well (1) Method: Dug [] Bored [] Deepened [] Cable [] Driven [] Reconditioned [] Rotary [] Jetted []	stratum penetroled, with at least one entry for each MATERIAL Brn Sandy Clay	FROM O	TO TO
(5) DIMENSIONS: Diameter of well 6 inches. Drilled 78 ft. Depth of completed well 78 ft.	grafsands las	8	2-5 5-3
(6) CONSTRUCTION DETAILS: Casing installed: <u>O Diam. from</u> <u>O R. to</u> 78 r.	Bin Sandy Clay	33	80
Threaded	group sound flor	80	130
Perforations: yes No E Type of perforator used	gray gravely landfan	130	137
perforations from ft. to ft. perforations from ft. to ft. to ft.	groysandy Clay	137	150
Screens: Yes D No Manufacturer's Name	grafhand sendy Cley	150	170
Diam. Slot size from ft. to ft. Diam. Slot size from ft. to ft.	lin sandy thank	170	128
Gravel packed: Yes No Size of gravel: Gravel placed from ft. to ft. to ft. Surface seal: Yes No To what depth? 20 ft. Material used in seal Did any strata contain unusable water Yes No F Type of water? Depth of strata Method of sealing strata off.			
(7) PUMP: Manufacturer's Name			
(8) WATER LEVELS: Land-surface elevation 4.30 above mean sea level			
(9) WELL TESTS: Drawdown is amount water level is lowered below static level Was a pump test made? Yes □ No □ If yes, by whom? Yield: gal./min. with ft. drawdown after hrs. """"""""""""""""""""""""""""""""""""	Work started /- 22	n and this	
Recovery data (time taken as zero when pump turned off) (water level measured from well top to water level) Time Water Level Time Water Level Time Water Level	NAME RICH Drd Det KC (Person, firm, or corporation) Address. 2722 Ruy 10/6	Va)	print)
Date of test, 1-29-85 Bailer test 27 gal/min. with 3 ft, drawdown after hrs. Artesian flow g.p.m. Date Temperature of water. Was a chemical analysis made? Yes I No	[Signed] J. Chand Files (Well Driller) License No	9-	

Second Copy — Owner's Copy	LL REPORT Application 1 Ashington Permit No.		
B. J. M.L.	FRUID FRUID I	1	1100
(1) OWNER: Nameffuth (114) (2) LOCATION OF WELL: County / Serson	Address Rt 1 BOX 3 94 A Pt Jud 5 E. N.E. V.S.We V. Sec. 8 The	Y N D	Ewy
(2) LOCATION OF WELLE: County (County County	J. M	······································	
(3) PROPOSED USE: Domestic Municipal	(10) WELL LOG:		
Irrigation [] Test Well [] Other []	Formation: Describe by color, character, size of materia show thickness of aquifers and the kind and nature of	the materu	ai in each
(4) TYPE OF WORK: Owner's number of well (if more than one)	stratum penetrated, with at least one entry for each c MATERIAL	FROM	TO
New well Method: Dug Dered	Rea Jandy Clay	0	20
Deepened D Cable Driven C Reconditioned Rotary Jetted D	The trans		1.0
(5) DIMENSIONS: Diameter of well inches.	lite gray sandy city	20	40
Drilled 2 4 2 it. Depth of completed well 2 4 tt.	UBillin Bran sould Cley had	40	50
(6) CONSTRUCTION DETAILS:	1 Rough All Charl	m	105
Casing installed: 6 " Diam. from O to 21 10-	fine for carage of	50	10,1
Threaded Diam. from	Conce Pro Sandy Clay	105	135
Perforations: Yes D Note	Ren In du Clas + oracel	135	14.5
Type of perforator used	the first first		150
SIZE of perforations	Bra Sandy Clay	143	13 8
perforations fromft. toft. toft. toft. toft. toft. toft.	Bin sandy laid ban	158	170
	- tiplet	170	200
Screens: Yes D No Manufacturer's Name	Ben Saray Cloy	110	100
Type	UB. Bron Sondy lay growt	200	2114
Diam. Slot size from			
Gravel packed: Yes D No & Size of gravel:			
Gravel placed from ft. to ft.		<u>10</u>	
Surface seal: Yes No D To what depth? the Material used in seal state the former Clay		6	
Did any strata contain unusable water? Yes Ves No		<u> </u>	
Type of water?			
(7) PUMP: Manufacturer's Name			
(/) FUMIL: Manufacturers Name	7		
(8) WATER LEVELS: Land-surface elevation 480 ft.			
Static level 19 ft. below top of well Date 12-6-84 Artesian pressurelbs, per square inch Date			+
Artesian pressure			
(a) THEFT TECTS. Drawdown is amount water level is		6	
(9) WELL IESIS: lowered below static level Was a pump test made? Yes No I If yes, by whom?	Work started 11 - 21, 19.84. Completed		1
Yield: gal./min. with ft. drawdown after hrs.	WELL DRILLER'S STATEMENT: This well was drilled under my jurisdiction	and this	report is
	true to the best of my knowledge and belief.		
Recovery data (time taken as zero when pump turned off) (water level measured from well top to water level)	NAME RICHArd Berke	rar	· · · · · · · · · · · · · · · · · · ·
Time Water Level Time Water Level Time Water Level	(Person, firm, or corporation)	(Type or	print)
	Address 27 22 Huy 10/6 00	19017	ηuA
	Richard Bor	5 ber	m
Date of test gal/min. with 15 ft. drawdown after 4 hrs.	[Signed] (Weh Driller)		
Artesian flow	-779 - 12		- 18
Temperature of water manage was a themata analysis mader its [] no L			

	a	WATER WELL REPORT		START CARD NO			
1.	04454	STATE O	F WASHINGTON	WATER RIGHT PERM	UNIQUE WELL I D #	AFC 96	0
(1)	OWNER: Name Chris Bascha	ab	Address	P.O. Box 650		8365	
(2)	LOCATION OF WELL: County	Jefferson		NE 1/4 S	W 1/4 Sect 8 T	28N R	1F W M
	STREET ADDRESS OF WELL (or nearest address)						
(3)	PROPOSED USE: Domestic Irrigation O Industrial DeWater	Dther	(10) WELL LOG	or ABANDONME	NT PROCEDURE DES	CRIPTION	
(4)	Municipal Test Well TYPE OF WORK: Owner's number of well (If more than	one) 1	and the kind and nature o		aterial and structure, and show atum penetrated, with at least of		
、·,	🛛 New Well 🔋 Method 🗌 Dug 📋 Bored		of information	MATERIAL		FROM	то
	□ Abandoned ⊠ Cable □ Driven □ Deepened □ Rotary □ Jetted		Brown_hard_clay,_gr	avel		0_	_35
			Gray_cemented_grav	vel		35	45
(5)	DIMENSIONS: Diameter of well	6" inches	Brown_packed_sand	I_&_clay		45	56
(-)	Drilled275 feet Depth of completed well		Brown_very_fine_san	nd, packed, dry		56	69
(6)	CONSTRUCTION DETAILS:		Brown_sandy_clay			69	75
(•)	Casing installed _6" diam From0 ft T	ro <u>271-10</u> tt	Brown_silty_clay			75	102
	⊗ Welded5" diam From269-1 ft 7		Brown_very_fine_san	•		102	137
	Threaded diam From ft T	fo ft	Brown_packed_sand			137	
	Perforations Yes No		Brown_tight_sand,_si		-	151	162
	Type perforator used		Brown_sandy_clay,_c	•		162 177	<u>177</u> 189
	Size of perforations in by		Brown_sandy_clay,_f			1.89	201
	perforations From ft T	fo ft				201	212
	perforations From ft T		· · ·			212	217
_	perforations From ft T	oft	Blue_sandy_firm_clay			217	224
	Screens 🛛 Yes 🗋 No		Blue_sandy_clay			224	230
	Manufacture's NameJohnsonJohnson		Blue_silty_clay			230	246
	TypeStainless V-SlotMo		Blue_fine_sand,_strat	tified,_dirty		246	250
	Diam _5" Slot size 010_ From _270-10_ ft T Diam		Light_brown_sandy_c	slay		250	260
	Diam		Light_brown_sandy,_	gravely_clay		260	266
	Gravel packed ☐ Yes ⊠ No Size of gravel		Brown_med_&_coars	•		266	27_1
			Brown_med_&_coars	e_sand, pea_grave	I, W/B	27_1	275
	Gravel placed from ft Tr Surface seal ⊠ Yes □ No To what depth	?25ft	Brown_tight_gravel,_(clay,_no_flow		275	
	Material used in seal Med_Chip_Benton						
	Did any strata contain unusable water? 📋 Yes 👩 No				yields down to 3 gpm_		
	Type of water Depth of strata	ft			o to high pumping rate 900 feet from this well.	· · ·	
	Method of sealing strata off		ol_Pope_Resource_w				
(7)	PUMP: Manufacture's Name			KEt	EIVED	·····	
	Туре Н Р	· · · · · · · · · · · · · · · · · · ·		DE	C 1 4 2001		
(8)	WATER LEVELS: Land surface elevation above sea level				······		
		May_15,_2001_		Was	nington State		
	Artesian pressure lbs per square inch Date Artesian water is controlled by	,			nent of Ecology		
	(Cap, valve, etc)	Worked started		_ Completed	May 15, 20	01
(9)	WELL TEST: Drawdown is amount water level is lowered by			IMai_20,_200_i		_iviay_1.0,_2.0	
	Was a pump test made? □ Yes ⊠ No if yes,by whom? Yield		WELL CONSTRUC	CTOR CERTIFICA	TION:		
	Yield gal /min with ft drawdown after				ibility for construction of this we construction standards Mater		
	Yield gal /min with ft drawdown after				are true to my best knowledge		
	Recovery data (time taken as zero when pump turned off) (water from well top to water level)	level measured	NAME Charlie	's Drilling			
	Time Water Level Time Water Level Time	e Water Level		,	rson, Firm, or Corporation)		
			Address P.O. Box		, VVA. 90939		
			(Signed)	(hali)	Shik Lice	nse No	0458
	Date of test			(Well Driller)	•		
	Bailer test gal /min_with ft drawdown after		Contractor's Regi	stration No.	CHARLD*06	<u> </u>	
	Airtest gal /min with stem set at ft for			D-4-			
		May 11, 2001		Date	May 16, 20	01	
	Temp of water Was a chemical analysis made	⇒. ∏tes⊠ivo					

Second Copy — Owner's Copy		1 No	
(1) OWNER: Name FRANK L WOOD RUFF	1412 SE Mir LAW. LAAA	A. A. A. A. A.	90.00
(2) LOCATION OF WELL: County JeF/. Bearing and distance from section or subdivision corner			
(3) PROPOSED USE: Domestic Municipal	(10) WELL LOG:		
Irrigation 🗍 Test Well 🗌 Other 🗌	Formation: Describe by color, character, size of mater show thickness of aquifers and the kind and nature of stratum penetrated, with at least one entry for each	f the materia	it in each
(4) TYPE OF WORK: Owner's number of well (if more than one)	MATERIAL	FROM	TO
New well 🖉 Method: Dug 🔲 Bored 🗌	RACKY H.P.	0	14
Despened [] Cable [] Driven [] Reconditioned [] Rotary 💇 Jetted []	SANdy H.C.	19	50
	BROWN SANAT CLAY	50	18
(5) DIMENSIONS: Diameter of well inches. Drilled 122 ft. Depth of completed well 124 ft.	ETAT GYARLA HAP.		110
Drittet	LAREC AYTRA HARD RH	K/10	115
(6) CONSTRUCTION DETAILS:	BYOWN SAND - NSO		
Casing installed: Diam. from + L., ft. to + 9.8 ft.	PLA BYAKEL HEO	197	1. 8. 4
Threaded []	H.P.	112	2
Welded 🕱			
Perforations: Yes 🗆 No 🚔			
Type of perforator used in. by in.			
perforations from ft. to ft.			
perforations from	······································		f
perforations from			
Screens: Yes 🗆 No 💋			
Manufacturer's Name			
Type Model No			
Diam. Slot size trom tt. to tt.			
Gravel packed: Yes D No D Size of gravel:			
Gravel placed from			
Surface seal: Yes B No D To what depth? 20. rt.			
Material used in seal BANTANITE + Priklin 62			
Did any strata contain unusable water? Yes 🗋 No 🐙			
Type of water?			
(7) PUMP: Manufacturer's Name			
(8) WATER LEVELS: Land-surface elevation above mean sea level			
Static levelft. below top of well Date= TAM & 19 Artesian pressurelbs. per square inch Date	<u>/</u> /		
Artesian pressure Artesian water is controlled by (Cap, valve, etc.)			
(Cap, valve, etc.)			
(9) WELL TESTS: Drawdown is amount water level is lowered below static level	Work started		
Was a pump test made? Yes 🗋 No 🗊 If yes, by whom?	WELL DRILLER'S STATEMENT:		
Yield: gal./min. with ft. drawdown after hrs.			ancet !-
	This well was drilled under my jurisdiction true to the best of my knowledge and belief.	I ADG (DIS)	report is
Recovery data (time taken as zero when pump turned off) (water level	Si an Ala	4.	
measured from well top to water level) Time Water Level Time Water Level Time Water Level	NAME (Person, firm, or corporation)	the CO	(
		•	
	Address R.T. 1. B. 693.A. Q.	ulu	(
	here here here and		
Date of test	[Signed]		
Artesian flow	License No. D. 2. 8 5 Dates TA		
Temperature of water M.Q Was a chemical analysis made? Yes 🔲 No 🗌	License No. M. A. J. Datest. A.	₹.,₡₽	

- ()

File Original and First Copy with Department of Ecology Second Copy — Owner's Copy Third Copy — Driller's Copy	VATER WE	LL REPORT Application I		•
(1) OWNER: Name R. T. MO.TEN				
(2) LOCATION OF WELL: County Jes	fferson	TAX IS NE Sec. 8 T. 2	8 N. R.	
Bearing and distance from section or subdivision corner				
(3) PROPOSED USE: Domestic I Industrial	Municipal []	(10) WELL LOG:		<u></u>
(3) FROFUSED USE: Domestic (1) Industrial Irrigation [] Test Weil		Formation: Describe by color, character, size of materia show thickness of aquifers and the kind and nature of stratum penstrated, with at least one entry for each c	l and struc the materic hange of f	cture, and al in each formation.
(4) TYPE OF WORK: Owner's number of well (if more than one)		MATERIAL	TROM	то
New well 🖾 Method: Dug	Bored []	Brown top soil	0	
	e 💆 Driven 🗋	Brown sandy clay		11
		Brown sandy gravely clay &		
(5) DIMENSIONS: Diameter of well	0 inches.	large rocks	25	25
Drilled 290 10"rt. Depth of completed well		Brown clay	83	83_
(6) CONSTRUCTION DETAILS:		Brown gritty clay	128	138
Casing installed: 6 Diam. from	n. to 288 n.	Brown sandy clay Brown sand & gravel	138	142
Threaded []	5rt. to 205. 4.	Brown dry gravel	142	169
Weided 🕼	. ft. to ft.	Brown sandy clay	169	182
Perforations: Yes I No I		Brown fine dry sand	182	197
Type of perforator used		Brown dry large gravel &		
SIZE of perforations	in.	coarse sand	197	199
perforations from	to ft.	Greenish brown clay & wood		
perforations from ft.	to ft.	<u>partickes</u>	218	218
		Gray clay		
Screens: Yes A No D		Greenish silty clay	241	256
Manufacturer's NameJohnson Type_Stainless_StaelModel	No	clay (some seepage)	256	270
Diam 5 5/9810t size 020 from 285	811. to 29010	<u>clay (some seepage)</u> . Greenish brown sandy clay	270	287
Diam	ft. to ft.	Blue coarse sand & large		
Gravel packed: Yes D No I Size of grav	vel ·	gravel W/B Static approx.		
Gravel placed from ft. to		215	287	289
		Greenish brown clap	289	
Surface seal: Yes of No D To what depth Material used in seal Bentonite &	clav			<u></u>
Did any strata contain unusable water?	Yes 🚺 🛛 No 💾	I€H		
Type of water? Depth of sti	rata		κ_{Λ}	
Method of scaling strata off		SOUTHWEST OF	NU -	
(7) PUMP: Manufacturer's Name		SOUTHWEST REGIONAL OFF		
Туре:		ST Pr OF r		
(8) WATER LEVELS: Land-surface elevation	n #	REGIONAL OG		
(8) WATER LEVELS: above mean sea level. Static level	Date 6/30/78	OFFIC		
Artesian pressure	Date		<u>[]</u>	
Artesian water is controlled by	valve, etc.)	· · · · · · · · · · · · · · · · · · ·		
			6/70	
(3) WELL LEGIS. lowered below static le	ivel	Work started	0/ 20	
Was a pump test made? Yes No (I if yes, by whor Violation gal/min, with ft, drawdown		WELL DRILLER'S STATEMENT:		
Yield: gal./min. with ft. drawdown		This well was drilled under my jurisdiction	n and this	s report is
	F1	true to the best of my knowledge and belief.		
Recovery data (time taken as zero when pump turne measured from well top to water level) Time Water Level Time Water Level Tim			(Type or	print)
Time Water Level Time Water Level Time		Address P.O. Box 161 Sequim,	Wash.	
		1 1 1 to the		
		[Signed] Values Alucan Pro	esiden	1t_047
Bailer test 5 gel/min. with	n after	(Well Driller) Charlie Sherk 0458 Date		
Artesian flow		License No. 0458 Date	(/10	, 197.8
Temperature of watermanage was a creation menter		Sub Contractor-Bill Weber (Webers Well Drilling	1000	

(USE ADDITIONAL SHEETS IF N

	E
DEPARTMENT OF CONSERVATION DIVISION OF WATER ANDURCES	
WELL SCHEDULE Date 8-21 To75 No 28/1E 3K1	
Record by A Williamson	
Storres	
L Location State of WASHINGTON	
NNIASEN - R THE PL COMMENT OF SECTION	
	An and a second se
2. Owner or Temper	
Address.	
4. Land anthree datante	
5. Type Dug Drilled Driven Bught Rept	
Bored Jetted Seet	
6. Casing: Diam 6 to to In Type	
7. Chief aquifer(s)	
below above	
10. Yields Flow grif sile . Press. gal min. Mass. Rept. Est.	
Adequacy permanence	a contraction of the second
12. Quality Sample No.	
13. Other data: Log Water levels Draft Putup test. Analysee	
	NAN MENTAL AND AN

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APPENDIX B

Field Procedures Addendum

B.1 Field Procedures Addendum

B.1.1 Sub-Slab Soil Vapor Sampling Procedures

The purpose of this Addendum is to provide field personnel with an outline of the specific information needed to collect and document representative sub-slab soil vapor samples. The recommended sub-slab soil vapor sampling technique, as presented in this Addendum, is based on the assumption that soil vapor samples should be representative of chemicals that may volatilize from the uppermost aquifer into the vadose zone.

B.1.1.1 Sampling Equipment and Materials

The following equipment and materials are necessary to properly conduct sub-slab soil vapor sampling (see Figure B-1):

- Rotary hammer drill with a 2-inch and a 1-inch carbide tipped bit.
- Extension cord and generator (if no power outlets are available).
- 3-inch (length) stainless steel (SS) screen assembly with locking cap (AMS GVP probe assembly or equivalent).
- Hose barb, stainless steel (1/4-inch).
- Teflon® tape.
- 100% Beeswax, to seal vapor port borehole annulus.
- Quick Set Concrete Patch, to seal vapor port borehole annulus.
- Air pump and appropriate connection tubing, tee fittings, valves, and flow metering device for purging and sampling vapor ports.
- 1-liter Tedlar® bags to collect purged vapors.
- Sufficient number of Summa canisters and appropriate flow controllers to collect soil vapor samples.
- Equipment required for collection of samples using Summa canisters, including appropriate wrenches and pressure gauges.
- An accurate and reliable watch that has been properly set.
- A calculator.
- Field notebook, applicable sampling analysis plan, and Chain of Custody.
- Health-and-safety equipment and supplies (e.g., personal protective equipment [PPE]) as described in the relevant site health-and-safety plan (HSP).
- Shipping package for the Summa canisters.

When leak testing is required, additional equipment and materials include:

- Leak test shroud of sufficient size to cover soil gas vapor probe and sampling train (including Summa canister).
- A soft gasket to seal the leak test shroud to the floor.
- Tracer gas (helium), supplied in a 20 cubic foot gas cylinder with flow regulator.
- Flow regulator with 1/8-inch barbed outlet and tubing to connect the helium gas cylinder to the shroud.
- MGD-2002 helium meter or equivalent.

B.1.1.2 Sampling Procedure

Preparation

- Prior to beginning, clear sampling locations for utilities, verify access agreements are in place, and obtain required permits, as appropriate.
- Install sub-slab soil vapor sampling ports as follows:
 - Drill a 2-inch borehole to a depth of approximately 3 inches.
 - Drill a 1-inch borehole through the center of the 2-inch borehole through the floor slab of the building foundation to a depth of approximately 12 inches below the surface.
 - Construct the vapor point as shown in Figure B-1 and insert such that the top
 of the assembly is set approximately 1/8-inch below the top of the slab.
 - Seal the vapor port by melting the beeswax with a small butane torch. Pour the beeswax from the rubber plug up to the bottom ¹/₂-inch of the 2-inch borehole.
 - Allow beeswax to solidify and harden.
 - Mix Quick Set concrete patch and apply from top of beeswax seal to within ¹/₄-inch of the top of the slab.
- Assemble sampling train. The sampling train will be set up so that the Summa canister is in-line between the vapor port and the air pump, with a valve between the canister and the pump (see Figure B-1):
 - Verify the Summa canister number engraved on the canister matches the number listed on the certified clean tag to ensure proper decontamination of the canister was completed. Fill out the sample tag.
 - Verify the canister valve is closed tightly and remove the threaded cap at the inlet of the canister.
 - Attach the flow controller to the inlet of the canister; the flow controller will have a built in pressure gauge.
 - Connect the tubing from vapor port to inlet of a ¹/₄-inch tee fitting.

- Connect the Summa canister/flow controller to one outlet of the tee fitting.
- Connect air pump to the other outlet of the tee fitting, insert a ¹/₄-inch shutoff valve between the tee fitting and the air pump.
- Where leak testing is required, a shroud will be placed over the vapor port and the Summa canister to keep tracer gas in contact with the vapor port and fittings. The shroud consists of a plastic bin of a known volume. Two holes will be drilled near the top of the shroud; one for connection of the helium gas cylinder and one for connection of the air pump located outside the shroud. A third hole will be drilled near the base of the shroud to monitor the helium concentration inside during sampling (see Figure B-1).

B.1.1.3 Sampling Methodology

Sample Collection

• Purge the vapor port and sampling train at approximately 100 ml/min using the air pump to ensure the sample is representative of subsurface conditions. Capture purged vapor in 1-liter Tedlar® bags at the outlet of the air pump and release the vapor outdoors. Three-to-five tubing volumes should be removed. Use the following equation to calculate volume to be purged:

$$\mathbf{V} = \boldsymbol{\pi} \mathbf{x} \mathbf{r}^2 \mathbf{x} \mathbf{l}$$

Where:

V = Volume of tubing

r = the inner radius of the tubing being used [inches]

l = the length of the tubing being used [inches]

 $\pi = 3.14$

(Convert to ml using 1-inch³ = 16.387 ml to determine how long to purge port)

- If the sampling and analysis plan calls for Tedlar® bag samples to be collected for analysis, these samples will be collected at the outlet of the air pump following purging of the vapor port.
- Begin sample collection by closing the ¹/₄-inch shutoff valve between the Summa canister and the air pump and opening the valve on the Summa canister. Immediately record the pressure on the gauge as the "initial pressure" on the tag attached to the canister.
- After sampling begins and the apparatus is verified to be operating correctly, leave the canister to fill.
- Record all sample information in the field book and/or applicable field forms including the following:
 - Canister number and sample identification,
 - Sample start date and times,
 - Location of sample (distance from walls shown on building floor plan),

- Initial and final pressure of canister, and
- Notes regarding leak test, if applicable.
- Return to check canisters periodically (depending on length of sample period), to ensure proper operation. It is necessary to check the canister prior to completion because the accuracy of the flow regulators can vary, causing the canisters to fill faster than expected. The final pressure at the end of sampling should be approximately -5 to -6 inches mercury (Hg). If the canister has already reached this point, sampling is complete, the canister valve should be closed, and the pressure recorded as the "final pressure" on the sample tag, the field book, and applicable field forms. Sample collection will be considered complete, regardless of final pressure, after the stated sample period has elapsed.
- Record the exact pressure of the canister and time at the end of sampling on the sample tag for that canister, in the field book, and on the applicable field forms.
- Verify that the canister valve is closed tightly, remove the flow controller, and replace the threaded cap at the top of the canister. Discard all sample tubing.
- Abandon vapor port by removing vapor screen and tubing, backfilling with glass bead, and patching with concrete.

Leak Testing

- Before purging or sampling begins, place the leak test shroud over the vapor port/Summa canister sampling apparatus. The tubing from the tee connection above the canister will pass through the wall of the shroud to connect with the air pump outside.
- Connect the helium cylinder to the leak test shroud using tubing from the flow regulator on the cylinder, through a hole in the wall of the shroud. Be sure to keep the cylinder in an upright position at all times.
- Connect the helium meter to the leak test shroud using the hole near the base.
- Use the flow regulator to slowly release helium into the leak test shroud until a predetermined concentration of helium is contained within the enclosed area. The helium concentration will be measured using the helium meter. Maintain helium concentrations throughout the sampling period by continuously bleeding cylinder gas into the shroud as needed.
- Prior to collecting the canister sample, the vapor port will be purged as described above. Purged vapor contained in the Tedlar® bags will be field screened using the helium meter to ensure that the concentration of helium inside the bags is less than 5 percent of the shroud concentration. If leakage is detected, the vapor port seal will be enhanced and connections will be inspected and tightened. This process will be repeated until no significant leakage has been demonstrated.
- After confirming no significant leakage, the ¹/₄-inch shutoff valve between the Summa canister and the air pump will be closed and the canister valve will be opened to begin collecting the sample.

B.1.1.4 Post-Sample Collection Procedures

Label all sample containers with the following information: sample identification, date and time sample was collected, the starting and ending canister pressure, the site name, and the company name.

Include all this information in the field book plus the ending time of sample collection, and transfer pertinent information to the Chain-of-Custody record. Pack all Summa canisters in the original shipping containers, sealed with a custody seal, and send to the lab for analysis. The official holding time for this analysis is 30 days. However, attempt to get samples to the lab as soon as possible to allow lab time to conduct re-runs, dilutions, and low-level analyses, as necessary prior to sample expiration.

B.1.1.5 Analysis

The soil gas samples should be analyzed using EPA Methods TO-14 or TO-15, and when necessary/possible, low-level analysis or Selective Ion Mode (SIM) analysis to obtain the lowest achievable detection and reporting limits. Note the desired analytical methods are on the Chain-of-Custody form, and be sure analysis for helium is specified for leak-tested samples.

B.1.1.6 Decontamination

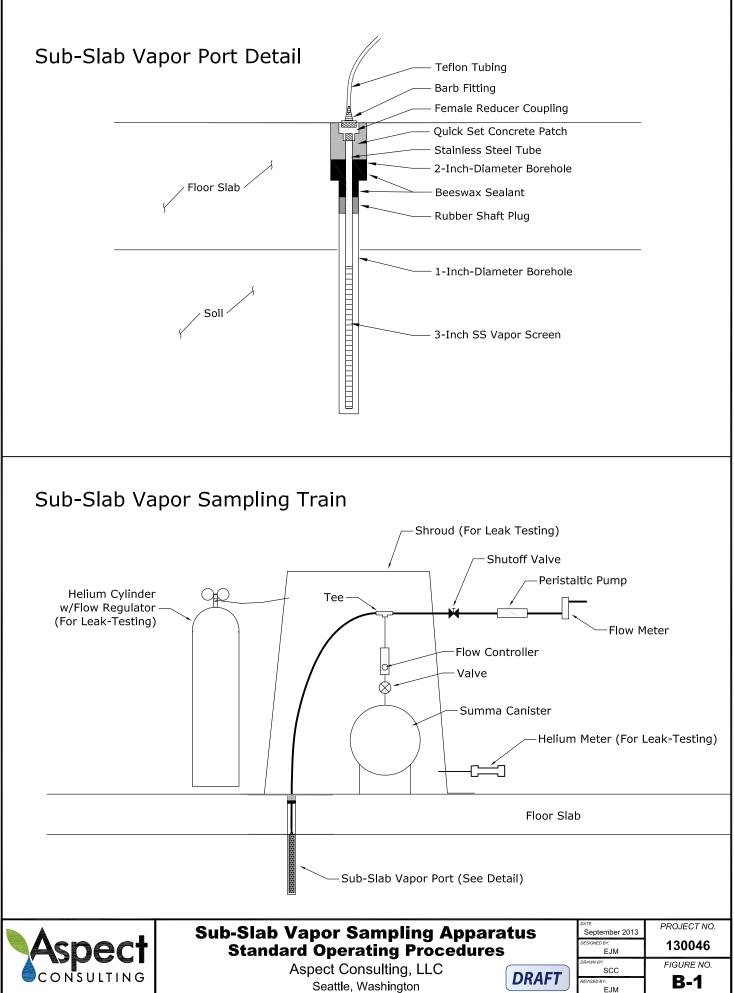
The equipment used for soil gas sampling does not require decontamination in the field. The Summa canisters will be individually cleaned and certified to 0.02 ppbv THC for the project-specific analyte list by the contract laboratory prior to shipment. Ensure that documentation of this certification is included on a tag attached to the canister and in the paperwork that accompanies the canister shipment from the lab.

B.1.1.7 Documentation

Record all field activities, environmental and building conditions, and sample documentation on the appropriate field forms and field notebook.

B.2 References

- Department of Environmental Protection, 2002, Commonwealth of Massachusetts, Indoor Air Sampling and Evaluation Guide, WSC Policy #02-430, Boston, Massachusetts, April 2002.
- EPRI, 2005, Reference Handbook for Site Specific Assessment of Sub-Surface Vapor Intrusion to Indoor Air, March 2005.
- New Jersey Department of Environmental Protection, 2005, Vapor Intrusion Guidance, October 2005.
- New York State Department of Health, 2006, Guidance for Evaluation Soil Vapor Intrusion in the State of New York, October 2006.
- USEPA, 1999, Center for Environmental Research Information, Office of Research and Development, Compendium of Methods for Determination of Toxic Organic Compounds in Ambient Air, Second Edition, Compendium Method To-14A, Determination of Volatile Organic Compounds (VOCs) in Ambient Air Using Specially Prepared Canisters with Subsequent Analysis by Gas Chromatography, January 1999.
- USEPA, 2002, Draft Guidance for Evaluating the Vapor Intrusion to Indoor Air Pathway Form Groundwater and Soils, EPA530-F-02-052, November 2002.



C:\Users\emarhofer\Desktop\Figure B-1.dwg 8.5x11 Portrait

sep 24, 2013 3:53pm emarhofer

APPENDIX C

Laboratory Analytical Report



LABORATORY REPORT

July 11, 2013

Greg Ferris Aspect Consulting 401 2nd Ave. S, Suite 201 Seattle, WA 98104-3814

RE: Port Ludlow OWSI / 130046

Dear Greg:

Enclosed are the results of the samples submitted to our laboratory on June 26, 2013. For your reference, these analyses have been assigned our service request number P1302737.

All analyses were performed according to our laboratory's NELAP and DoD-ELAP-approved quality assurance program. The test results meet requirements of the current NELAP and DoD-ELAP standards, where applicable, and except as noted in the laboratory case narrative provided. For a specific list of NELAP and DoD-ELAP-accredited analytes, refer to the certifications section at <u>www.alsglobal.com</u>. Results are intended to be considered in their entirety and apply only to the samples analyzed and reported herein.

If you have any questions, please call me at (805) 526-7161.

Respectfully submitted,

ALS | Environmental

Jadeca

By Kate Aguilera at 10:58 am, Jul 11, 2013 Kate Aguilera Project Manager



Client:Aspect ConsultingProject:Port Ludlow OWSI / 130046

Service Request No: P1302737

CASE NARRATIVE

The samples were received intact under chain of custody on June 26, 2013 and were stored in accordance with the analytical method requirements. Please refer to the sample acceptance check form for additional information. The results reported herein are applicable only to the condition of the samples at the time of sample receipt.

Fixed Gases Analysis

The samples were analyzed for fixed gases (oxygen/argon, nitrogen, methane and carbon dioxide) according to modified EPA Method 3C (single injection) using a gas chromatograph equipped with a thermal conductivity detector (TCD). This method is not included on the laboratory's NELAP or AIHA-LAP scope of accreditation.

<u>Helium Analysis</u>

The samples were also analyzed for helium according to modified EPA Method 3C (single injection) using a gas chromatograph equipped with a thermal conductivity detector (TCD). This method is not included on the laboratory's NELAP, DoD-ELAP, or AIHA-LAP scope of accreditation.

<u>Air-Phase Petroleum Hydrocarbons (APH) Analysis</u>

The samples were also analyzed for total aliphatic and aromatic gasoline range hydrocarbons by gas chromatography/mass spectrometry according to the Method for the Determination of Air-Phase Petroleum Hydrocarbons (APH), Massachusetts Department of Environmental Protection, Revision 1, December, 2009. This method is not included on the laboratory's DoD-ELAP or AIHA-LAP scope of accreditation.

Significant non-petroleum related peaks (i.e. halogenated, oxygenated, terpenes, etc.) are subtracted from the hydrocarbon range areas when present. Any internal/tuning standards and target APH analytes eluting in the hydrocarbon ranges are also subtracted. Additionally, $C_{g}-C_{10}$ Aromatic Hydrocarbons are excluded from the $C_{g}-C_{12}$ Aliphatic Hydrocarbon range.



Client: Aspect Consulting Project: Port Ludlow OWSI / 130046 Service Request No: P1302737

CASE NARRATIVE

Volatile Organic Compound Analysis

The samples were also analyzed for volatile organic compounds in accordance with EPA Method TO-15 from the Compendium of Methods for the Determination of Toxic Organic Compounds in Ambient Air, Second Edition (EPA/625/R-96/010b), January, 1999. The analytical system was comprised of a gas chromatograph / mass spectrometer (GC/MS) interfaced to a whole-air preconcentrator. The method was modified to include the use of helium as a diluent gas in place of zero-grade air for canister pressurization. When necessary, analytical sample volumes were adjusted by a correction factor for canisters pressurized with helium. A summary sheet has been included listing the affected samples. This method is not included on the laboratory's AIHA-LAP scope of accreditation. Any analytes flagged with an X are not included on the laboratory's NELAP or DoD-ELAP scope of accreditation.

The Summa canisters were cleaned, prior to sampling, down to the method reporting limit (MRL) reported for this project. Please note, projects which require reporting below the MRL could have results between the MRL and method detection limit (MDL) that are biased high.

The results of analyses are given in the attached laboratory report. All results are intended to be considered in their entirety, and ALS Environmental (ALS) is not responsible for utilization of less than the complete report.

Use of ALS Environmental (ALS)'s Name. Client shall not use ALS's name or trademark in any marketing or reporting materials, press releases or in any other manner ("Materials") whatsoever and shall not attribute to ALS any test result, tolerance or specification derived from ALS's data ("Attribution") without ALS's prior written consent, which may be withheld by ALS for any reason in its sole discretion. To request ALS's consent, Client shall provide copies of the proposed Materials or Attribution and describe in writing Client's proposed use of such Materials or Attribution. If ALS has not provided written approval of the Materials or Attribution shall be deemed denied. ALS may, in its discretion, reasonably charge Client for its time in reviewing Materials or Attribution requests. Client acknowledges and agrees that the unauthorized use of ALS's name or trademark may cause ALS to incur irreparable harm for which the recovery of money damages will be inadequate. Accordingly, Client acknowledges and agrees that a violation shall justify preliminary injunctive relief. For questions contact the laboratory.



ALS Environmental - Simi Valley

Certifications, Accreditations, and Registrations

Agency	Web Site	Number
АІНА	http://www.aihaaccreditedlabs.org	101661
Arizona DHS	http://www.azdhs.gov/lab/license/env.htm	AZ0694
DoD ELAP	http://www.pjlabs.com/search-accredited-labs	L11-203
Florida DOH (NELAP)	http://www.doh.state.fl.us/lab/EnvLabCert/WaterCert.htm	E871020
Maine DHHS	http://www.maine.gov/dhhs/mecdc/environmental-health/water/dwp- services/labcert/labcert.htm	2012039
Minnesota DOH (NELAP)	http://www.health.state.mn.us/accreditation	494864
New Jersey DEP (NELAP)	http://www.nj.gov/dep/oqa/	CA009
New York DOH (NELAP)	http://www.wadsworth.org/labcert/elap/elap.html	11221
Oregon PHD (NELAP)	http://public.health.oregon.gov/LaboratoryServices/EnvironmentalLaborat oryAccreditation/Pages/index.aspx	CA200007
Pennsylvania DEP	http://www.depweb.state.pa.us/labs	68-03307 (Registration)
Texas CEQ (NELAP)	http://www.tceq.texas.gov/field/qa/env_lab_accreditation.html	T104704413- 12-3
Utah DOH (NELAP)	http://www.health.utah.gov/lab/labimp/certification/index.html	CA01527201 2-2
Washington DOE	http://www.ecy.wa.gov/programs/eap/labs/lab-accreditation.html	C946

Analyses were performed according to our laboratory's NELAP and DoD-ELAP approved quality assurance program. A complete listing of specific NELAP and DoD-ELAP certified analytes can be found in the certifications section at <u>www.alsglobal.com</u>, or at the accreditation body's website.

Each of the certifications listed above have an explicit Scope of Accreditation that applies to specific matrices/methods/analytes; therefore, please contact the laboratory for information corresponding to a particular certification.

DETAIL SUMMARY REPORT

Client:	Aspect Consult	ng						Service Req	uest:	P130	0273	7	
Project ID:	Port Ludlow OV	WSI / 1300	046										
Date Received: Time Received:	6/26/2013 09:30								ed - Fxd Gases Can	ied - Helium Can	0 - MA VOC PH Can	OC Cans	1
Client Sample ID	Lab Code	Matrix	Date Collected	Time Collected	Container ID	Pi1 (psig)	Pf1 (psig)		3C Modified	3C Modified	MA APH 1.0	TO-15 - V	
VP-1-062113	P1302737-001	Air	6/21/2013	10:00	SC00683	-1.47	3.65		Х	Х	Х	Х	
VP-2-062113	P1302737-002	Air	6/21/2013	11:00	SC01531	-1.10	3.69		Х	Х	Х	Х	

ALS ENVIRONMENTAL Sample Volume Correction for Helium Pressurization for SCAN Analysis

			Sample	Adjusted
Sample ID	<u>Pi</u>	<u>Pf</u>	<u>Volume (L)</u>	<u>Volume (L)</u>
P1302737-001	-1.47	3.65	0.901	1.00
P1302737-002	-1.10	3.69	0.905	1.00

Flax (805) 526-7270		Requested Turnar 1 Day (100%) 2 Da	Requested Turnaround Time In Business Days (Surcharges) please circle 1 Day (100%) 2 Day (75%) 3 Day (50%) 4 Day (35%) 5 Day (25%) 10 Day-Standard	sss Days (Surch 4 Day (35%)	a rges) please (5 Day (25%) 10	circle Day-Standar		CAS Project	CAS Project No.
Compary Name & Address (Reporting Information)		Project Name				0	CAS Contact:		
Aspect Consulting		Port	r Ludlaw	ISMO	ZT Z		Analysis	Method	
		Project Number	074001				1		
Project Manager		P.O. # / Billing Information	mation				5.1 (51-	~~~~ ZN Way	
×		_	130046-001-01	0-100	1	0-1	- Guya	1 have	Comments e.g. Actual Preservative or
Email Address for Result Reporting offerris @ aspectconsulting, com		Sampler (Print & Sign)	Eric Geissing	acr 6	of G		n (m odzi Hd t	PD.4.	S
Client Sample ID Client Sample ID Collected	te Time cted Collected	Canister ID (Bar code # - AC, SC, etc.)	Flow Controller ID (Bar code #- FC #)		Canister End Pressure "Hg/psig	Sample Volume	95) 95) WW	snjd 7'20 100N	
10-17-1-062113 0-137 6/21/13	13 1000	S	FLOODTH	87-	-5.5	er er	×	\times	Plance not
	13 1100	5001531		- 29	- 5: 5	ر و ا	×	×	ી જ
	_								He 45 0
# C5-CB Aliphatic Hyddoca Son S									tracer gas
C9-C12 Aliphatic Hydrocarbons									
09-010 promatic Hydrocarty ns									
BTEX	/								
Naphthalene (MA APH ITO 15	-15 Recording	list (call	w/ questions	(4)	-			
MTBE	-			-					
n-herane									
1,2-dibromorathane									
1,2- dichlorethance					• .				
113, 5- trimetry beneenco									
1,2,4 -trimethylbeneere									
Report Tier Levels - please select Tier I - Results (Default if not specified)	Tier III (Results	Tier III (Results + OC & Calibration Summaries)	mmaries)				>		
Tier II (Results + QC Summaries)	Tier IV (Data Va	Tier IV (Data Validation Package) 10% Surcharge	Surcharge		- -	Type:	0NI / 621		Project Kequirements (MRLs, QAPP)
Relinquished by: (Sloggture)	Date: 0/24/13	Time: 13.90	Received by (SignAure) 00 0.0	200000	8	and a state	4772	Time:	
Relinquished by: (Signature)	Date:	Time:	Received by: (Signature)			Date:		Time:	Cooler / Blank

ot ____ Page___

4

Air - Chain of Custody Record & Analytical Service Request

Columbia Analytical Services*

ALS Environmental Sample Acceptance Check Form

	: Aspect Consult	ě.			-	Work order:	P1302737			
5	: Port Ludlow O									
-	(s) received on:			-	Date opened:	N	by:	MZAM		
		l samples received by ALS.		-	-	-			dication	of
compliance	or nonconformity.	Thermal preservation and p	H will only be ev	valuated either at t	he request or the	e client and/or as rec	juired by the memo	iod/SOP. <u>Yes</u>	<u>No</u>	N/A
1	Were sample (containers properly m	narked with cli	ient sample ID	?			X		
2	-	upplied by ALS?		-				X		
3	Did sample co	ontainers arrive in goo	od condition?					X		
4	Were chain-of	f-custody papers used	and filled out	.?				X		
5	Did sample co	ontainer labels and/or	tags agree wit	th custody pape	ers?			X		
6	Was sample v	olume received adequa	ate for analysi	is?				X		
7	Are samples w	vithin specified holding	g times?					X		
8	Was proper ter	emperature (thermal pr	reservation) o	f cooler at rece	pipt adhered t	ю?				X
								_	_	_
9	Was a trip bla								\mathbf{X}	
10	Were custody	seals on outside of coo							\mathbf{X}	
		Location of seal(s)?					Sealing Lid?			\mathbf{X}
	•	e and date included?								X
	Were seals inta									
	Were custody s	seals on outside of sam	-						\boxtimes	
	. .	Location of seal(s)?					Sealing Lid?			
	•	e and date included?								
	Were seals inta									X
11		rs have appropriate pr		-		Client specified	information?			X
		nt indication that the su	-		served?					X
		ials checked for presen								\mathbf{X}
		t/method/SOP require	•		mple pH and	if necessary alte	er it?			X
12	Tubes:	Are the tubes cappe	ed and intact?	?						X
		Do they contain me								X
13	Badges:	Are the badges pro								X
		Are dual bed badge	es separated <i>a</i>	und individually	y capped and	intact?				
Lab	Sample ID	Container	Required	Received	Adjusted	VOA Headspac	e Recei	ipt / Prese	ervatior	1
		Description	nH *	рН	рН	(Presence/Absence		Commen		

Lab Sample ID	Description	pH *	pH	Adjusted pH	(Presence/Absence)	-
P1302737-001.01	6.0 L Source Can					
P1302737-002.01	6.0 L Source Can					

Explain any discrepancies: (include lab sample ID numbers):

RSK - MEEPP, HCL (pH<2); RSK - CO2, (pH 5-8); Sulfur (pH>4)

RESULTS OF ANALYSIS

Page 1 of 1

Client: Client Sample ID: Client Project ID:	Aspect Consulting VP-1-062113 Port Ludlow OWSI / 130046			Project ID: P Sample ID: P	
Test Code: Instrument ID:	EPA Method 3C Modified HP5890 II/GC1/TCD			e Collected: 6/ e Received: 6/	
Analyst:	Jennifer Young			e Analyzed: 7/	
Sample Type: Test Notes:	6.0 L Summa Canister			s) Analyzed:	0.10 ml(s)
Container ID:	SC00683				
	Initial Pressure (psig): -1.47	Final Pressure (psig):	3.65		
			Ca	anister Dilution	Factor: 1.39
CAS #	Compound		Result %, v/v	MRL %, v/v	Data Qualifier
7782-44-7	Oxygen +		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	, , , , , , , , , , , , , , , , , , ,	Quantier
7440-37-1	Argon		15.0	0.14	
7727-37-9	Nitrogen		77.4	0.14	
74-82-8	Methane		ND	0.14	
124-38-9	Carbon Dioxide		7.54	0.14	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

RESULTS OF ANALYSIS

Page 1 of 1

Client: Client Sample ID: Client Project ID:	Aspect Consulting VP-2-062113 Port Ludlow OWSI / 130046			Project ID: P Sample ID: P	
Test Code:	EPA Method 3C Modified		Dut	e Collected: 6/	=1,10
Instrument ID:	HP5890 II/GC1/TCD			e Received: 6/	
Analyst:	Jennifer Young 6.0 L Summa Canister			e Analyzed: 7/	
Sample Type: Test Notes:	0.0 L Summa Canister		v olume(s) Analyzed:	0.10 ml(s)
Container ID:	SC01531				
	Initial Pressure (psig): -1.10	Final Pressure (psig):	3.69		
			Ca	nister Dilution	Factor: 1.35
CAS #	Compound		Result	MRL	Data
			%, v/v	%, v/v	Qualifier
7782-44-7	Oxygen +				
7440-37-1	Argon		12.9	0.14	
7727-37-9	Nitrogen		79.3	0.14	
74-82-8	Methane		ND	0.14	
124-38-9	Carbon Dioxide		7.78	0.14	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

RESULTS OF ANALYSIS

Page 1 of 1

Client: Client Sample ID: Client Project ID:	Aspect Consulting Method Blank Port Ludlow OWSI / 130046	ALS Project ID: P1302737 ALS Sample ID: P130701-MB
Test Code:	EPA Method 3C Modified	Date Collected: NA
Instrument ID:	HP5890 II/GC1/TCD	Date Received: NA
Analyst:	Jennifer Young	Date Analyzed: 7/01/13
Sample Type:	6.0 L Summa Canister	Volume(s) Analyzed: 0.10 ml(s)
Test Notes:		

CAS #	Compound	Result %, v/v	MRL %, v/v	Data Qualifier
7782-44-7	Oxygen +			
7440-37-1	Argon	ND	0.10	
7727-37-9	Nitrogen	ND	0.10	
74-82-8	Methane	ND	0.10	
124-38-9	Carbon Dioxide	ND	0.10	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

LABORATORY CONTROL SAMPLE SUMMARY

Page 1 of 1

Client:	Aspect Consulting	
Client Sample ID:	Lab Control Sample	ALS Project ID: P1302737
Client Project ID:	Port Ludlow OWSI / 130046	ALS Sample ID: P130701-LCS

Test Code:	EPA Method 3C Modified	Date Collected: NA	
Instrument ID:	HP5890 II/GC1/TCD	Date Received: NA	
Analyst:	Jennifer Young	Date Analyzed: 7/01/1	3
Sample Type:	6.0 L Summa Canister	Volume(s) Analyzed:	NA ml(s)
Test Notes:			

CAS #	Compound	Spike Amount ppmV	Result ppmV	% Recovery	ALS Acceptance Limits	Data Qualifier
7782-44-7	Oxygen +					
7440-37-1	Argon	50,000	55,200	110	85-111	
7727-37-9	Nitrogen	49,400	54,500	110	85-114	
74-82-8	Methane	39,500	42,700	108	90-114	
124-38-9	Carbon Dioxide	49,300	53,000	108	84-113	

RESULTS OF ANALYSIS

Page 1 of 1

Client:Aspect ConsultingClient Project ID:Port Ludlow OWSI / 130046

ALS Project ID: P1302737

Helium

Test Code:	EPA 3C Modified	
Instrument ID:	HP5890 II/GC8/TCD	Date(s) Collected: 6/21/13
Analyst:	Jennifer Young	Date Received: 6/26/13
Sample Type:	6.0 L Summa Canister(s)	Date Analyzed: 6/28/13
Test Notes:		

Client Sample ID	ALS Sample ID	Injection Volume ml(s)	Canister Dilution Factor	Result ppmV	MRL ppmV	Data Qualifier
VP-1-062113	P1302737-001	1.00	1.10	180	28	
VP-2-062113	P1302737-002	1.00	1.06	ND	27	
Method Blank	P130628-MB	1.00	1.00	ND	25	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

LABORATORY CONTROL SAMPLE SUMMARY

Page 1 of 1

-	Aspect Consulting Lab Control Sample Port Ludlow OWSI / 130046	ALS Project ID: P1302737 ALS Sample ID: P130628-LCS
Test Code: Instrument ID:	EPA 3C Modified HP5890 II/GC8/TCD	Date Collected: NA Date Received: NA
Analyst:	Jennifer Young	Date Analyzed: 6/28/13

					ALS	
CAS #	Compound	Spike Amount	Result	% Recovery	Acceptance	Data
		ppmV	ppmV		Limits	Qualifier
7440-59-7	Helium	10,000	9,620	96	70-127	

Volume(s) Analyzed:

NA ml(s)

Sample Type:

Test Notes:

6.0 L Summa Canister

RESULTS OF ANALYSIS

Page 1 of 1

Client:	Aspect Consulting			
Client Sample ID:	VP-1-062113	ALS Project ID: 1	P1302737	
Client Project ID:	Port Ludlow OWSI / 130046	ALS Sample ID:	P1302737-001	
Test Code:	Massachusetts APH, Revision 1, December 2009	Date Collected:	5/21/13	
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received:	6/26/13	
Analyst:	Chris Cornett	Date Analyzed:	7/4/13	
Sample Type:	6.0 L Summa Canister Vo	lume(s) Analyzed:	1.00 Liter(s	5)
Test Notes:				
Container ID:	SC00683			
	Initial Pressure (psig): -1.47 Final Pressure (psig):	3.65		
		Canister	Dilution Factor	: 1.39
Compound		Result	MRL	Data
		μg/m ³	µg/m³	Qualifier
C ₅ - C ₈ Aliphatic H	ydrocarbons ^{1,2}	110	28	

Significant non-petroleum related peaks (i.e. halogenated, oxygenated, terpenes, etc.) are subtracted from the hydrocarbon range areas when present.

¹Hydrocarbon Range data from total ion chromatogram excluding any internal/tuning standards eluting in that range.

 $^{2}C_{5}-C_{8}$ Aliphatic Hydrocarbons exclude the concentration of Target APH analytes eluting in that range.

 ${}^{3}C_{9}-C_{12}$ Aliphatic Hydrocarbons exclude concentration of Target APH Analytes eluting in that range and concentration of C₉-C₁₀ Aromatic Hydrocarbons. ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

2,100

42

14

3.5

MRL = Method Reporting Limit - The minimum quantity of a target analyte that can be confidently determined by the referenced method.

C₉ - C₁₂ Aliphatic Hydrocarbons^{1,3}

C₉ - C₁₀ Aromatic Hydrocarbons

RESULTS OF ANALYSIS

Page 1 of 1

Client: Client Sample ID: Client Project ID:	Aspect Consulting VP-2-062113 Port Ludlow OWSI / 130046	ALS Project ID: ALS Sample ID:	
Test Code: Instrument ID:	Massachusetts APH, Revision 1, December 2009 Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Collected: Date Received:	
Analyst:	Chris Cornett	Date Analyzed:	7/4/13
Sample Type: Test Notes:		Volume(s) Analyzed:	1.00 Liter(s)
Container ID:	SC01531 Initial Pressure (psig): -1.10 Final Pressure (psi	ig): 3.69	
		Canister	Dilution Factor: 1.35
Compound		Result µg/m³	MRL Data µg/m ³ Qualifier

$C_9 - C_{12}$ Aliphatic Hydrocarbons ^{1,3} 790 14 $C_9 - C_{10}$ Aromatic Hydrocarbons 16 3.4	$C_5 - C_8$ Aliphatic Hydrocarbons ^{1,2}	100	27	
$C_0 - C_{10}$ Aromatic Hydrocarbons 16 3 4	C ₉ - C ₁₂ Aliphatic Hydrocarbons ^{1,3}	790	14	
	C ₉ - C ₁₀ Aromatic Hydrocarbons	16	3.4	

Significant non-petroleum related peaks (i.e. halogenated, oxygenated, terpenes, etc.) are subtracted from the hydrocarbon range areas when present.

¹Hydrocarbon Range data from total ion chromatogram excluding any internal/tuning standards eluting in that range.

 $^{2}C_{5}-C_{8}$ Aliphatic Hydrocarbons exclude the concentration of Target APH analytes eluting in that range.

 ${}^{3}C_{9}-C_{12}$ Aliphatic Hydrocarbons exclude concentration of Target APH Analytes eluting in that range and concentration of C₉-C₁₀ Aromatic Hydrocarbons. ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

RESULTS OF ANALYSIS

Page 1 of 1

Client: Client Sample ID: Client Project ID:	Aspect Consulting Method Blank Port Ludlow OWSI / 130046	ALS Project ID: P1302737 ALS Sample ID: P130703-MB
Test Code:	Massachusetts APH, Revision 1, December 2009	Date Collected: NA
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received: NA
Analyst:	Chris Cornett	Date Analyzed: 7/3/13
Sample Type: Test Notes:	6.0 L Summa Canister	Volume(s) Analyzed: 1.00 Liter(s)

Compound	Result	MRL	Data
	μg/m³	µg/m³	Qualifier
C ₅ - C ₈ Aliphatic Hydrocarbons ^{1,2}	ND	20	
C ₉ - C ₁₂ Aliphatic Hydrocarbons ^{1,3}	ND	10	
C ₉ - C ₁₀ Aromatic Hydrocarbons	ND	2.5	

Significant non-petroleum related peaks (i.e. halogenated, oxygenated, terpenes, etc.) are subtracted from the hydrocarbon range areas when present.

¹Hydrocarbon Range data from total ion chromatogram excluding any internal/tuning standards eluting in that range.

 $^{2}C_{5}-C_{8}$ Aliphatic Hydrocarbons exclude the concentration of Target APH analytes eluting in that range.

 ${}^{3}C_{9}-C_{12}$ Aliphatic Hydrocarbons exclude concentration of Target APH Analytes eluting in that range and concentration of C₉-C₁₀ Aromatic Hydrocarbons. ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

LABORATORY CONTROL SAMPLE SUMMARY

Page 1 of 1

-	Aspect Consulting Lab Control Sample Port Ludlow OWSI / 130046	ALS Project ID: P1302737 ALS Sample ID: P130703-LCS
Test Code:	Massachusetts APH, Revision 1, December 2009	Date Collected: NA
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received: NA
Analyst:	Chris Cornett	Date Analyzed: 7/3/13
Sample Type: Test Notes:	6.0 L Summa Canister	Volume(s) Analyzed: 0.125 Liter(s)

				ALS	
Compound	Spike Amount	Result	% Recovery	Acceptance	Data
	$\mu g/m^3$	μg/m³		Limits	Qualifier
C5 - C8 Aliphatic Hydrocarbons	202	167	83	70-130	
C9 - C12 Aliphatic Hydrocarbons	204	184	90	70-130	
C9 - C10 Aromatic Hydrocarbons	402	366	91	70-130	

RESULTS OF ANALYSIS

Page 1 of 1

Client:	Aspect Consulting	
Client Sample ID:		ALS Project ID:
Client Project ID:	Port Ludlow OWSI / 130046	ALS Sample ID:
Test Code:	EPA TO-15 Modified	Date Collected:
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received:
Analyst:	Chris Cornett	Date Analyzed:
Sample Type:	6.0 L Summa Canister	Volume(s) Analyzed:
Test Notes:		
Container ID:	SC00683	

Initial Pressure (psig): -1.47

Final Pressure (psig): 3.65

Canister Dilution Factor: 1.39

1.00 Liter(s)

P1302737 P1302737-001

6/21/13 6/26/13 7/4/13

CAS #	Compound	Result	MRL	Result	MRL	Data
		μg/m ³	$\mu g/m^3$	ppbV	ppbV	Qualifier
1634-04-4	Methyl tert-Butyl Ether	ND	0.70	ND	0.19	
110-54-3	n-Hexane	ND	0.70	ND	0.20	
107-06-2	1,2-Dichloroethane	ND	0.70	ND	0.17	
71-43-2	Benzene	ND	0.70	ND	0.22	
108-88-3	Toluene	9.8	0.70	2.6	0.18	
106-93-4	1,2-Dibromoethane	ND	0.70	ND	0.090	
100-41-4	Ethylbenzene	3.2	0.70	0.73	0.16	
179601-23-1	m,p-Xylenes	15	1.4	3.5	0.32	
95-47-6	o-Xylene	4.4	0.70	1.0	0.16	
108-67-8	1,3,5-Trimethylbenzene	2.4	0.70	0.50	0.14	
95-63-6	1,2,4-Trimethylbenzene	7.7	0.70	1.6	0.14	
91-20-3	Naphthalene	1.2	0.70	0.23	0.13	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

RESULTS OF ANALYSIS

Page 1 of 1

Client:	Aspect Consulting		
Client Sample ID:	VP-2-062113	ALS Project ID: P1	302737
Client Project ID:	Port Ludlow OWSI / 130046	ALS Sample ID: P1	302737-002
Test Code:	EPA TO-15 Modified	Date Collected: 6/2	21/13
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received: 6/2	26/13
Analyst:	Chris Cornett	Date Analyzed: 7/4	4/13
Sample Type:	6.0 L Summa Canister	Volume(s) Analyzed:	1.00 Liter(s)
Test Notes:			
Container ID:	SC01531		

Initial Pressure (psig): -1.10

Final Pressure (psig): 3.69

Canister Dilution Factor: 1.35

CAS #	Compound	Result µg/m³	MRL µg/m³	Result ppbV	MRL ppbV	Data Qualifier
1634-04-4	Methyl tert-Butyl Ether	ND	0.68	ND	0.19	
110-54-3	n-Hexane	ND	0.68	ND	0.19	
107-06-2	1,2-Dichloroethane	ND	0.68	ND	0.17	
71-43-2	Benzene	ND	0.68	ND	0.21	
108-88-3	Toluene	12	0.68	3.2	0.18	
106-93-4	1,2-Dibromoethane	ND	0.68	ND	0.088	
100-41-4	Ethylbenzene	10	0.68	2.3	0.16	
179601-23-1	m,p-Xylenes	45	1.4	10	0.31	
95-47-6	o-Xylene	28	0.68	6.5	0.16	
108-67-8	1,3,5-Trimethylbenzene	1.6	0.68	0.33	0.14	
95-63-6	1,2,4-Trimethylbenzene	3.5	0.68	0.71	0.14	
91-20-3	Naphthalene	ND	0.68	ND	0.13	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

RESULTS OF ANALYSIS

Page 1 of 1

Client:Aspect ConsultingClient Sample ID:Method BlankClient Project ID:Port Ludlow OWSI / 130046

ALS Project ID: P1302737 ALS Sample ID: P130703-MB

Test Code:	EPA TO-15 Modified	Date Collected: N	A
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received: N	A
Analyst:	Chris Cornett	Date Analyzed: 7/	3/13
Sample Type:	6.0 L Summa Canister	Volume(s) Analyzed:	1.00 Liter(s)
Test Notes:			

Canister Dilution Factor: 1.00

CAS #	Compound	Result µg/m³	MRL µg/m³	Result ppbV	MRL ppbV	Data Qualifier
1634-04-4	Methyl tert-Butyl Ether	ND	0.50	ND	0.14	
110-54-3	n-Hexane	ND	0.50	ND	0.14	
107-06-2	1,2-Dichloroethane	ND	0.50	ND	0.12	
71-43-2	Benzene	ND	0.50	ND	0.16	
108-88-3	Toluene	ND	0.50	ND	0.13	
106-93-4	1,2-Dibromoethane	ND	0.50	ND	0.065	
100-41-4	Ethylbenzene	ND	0.50	ND	0.12	
179601-23-1	m,p-Xylenes	ND	1.0	ND	0.23	
95-47-6	o-Xylene	ND	0.50	ND	0.12	
108-67-8	1,3,5-Trimethylbenzene	ND	0.50	ND	0.10	
95-63-6	1,2,4-Trimethylbenzene	ND	0.50	ND	0.10	
91-20-3	Naphthalene	ND	0.50	ND	0.095	

ND = Compound was analyzed for, but not detected above the laboratory reporting limit.

SURROGATE SPIKE RECOVERY RESULTS

Page 1 of 1

Client:Aspect ConsultingClient Project ID:Port Ludlow OWSI / 130046

ALS Project ID: P1302737

Test Code:	EPA TO-15 Modified	
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date(s) Collected: 6/21/13
Analyst:	Chris Cornett	Date(s) Received: 6/26/13
Sample Type:	6.0 L Summa Canister(s)	Date(s) Analyzed: 7/3 - 7/4/13
Test Notes:		

		1,2-Dichloroethane-d4	Toluene-d8	Bromofluorobenzene		
Client Sample ID	ALS Sample ID	Percent	Percent	Percent	Acceptance	Data
		Recovered	Recovered	Recovered	Limits	Qualifier
Method Blank	P130703-MB	86	98	105	70-130	
Lab Control Sample	P130703-LCS	83	95	106	70-130	
VP-1-062113	P1302737-001	85	96	109	70-130	
VP-2-062113	P1302737-002	85	97	109	70-130	

Surrogate percent recovery is verified and accepted based on the on-column result.

Reported results are shown in concentration units and as a result of the calculation, may vary slightly from the on-column percent recovery.

LABORATORY CONTROL SAMPLE SUMMARY

Page 1 of 1

-	Aspect Consulting Lab Control Sample Port Ludlow OWSI / 130046	ALS Project ID: P1302737 ALS Sample ID: P130703-LCS
Test Code:	EPA TO-15 Modified	Date Collected: NA
Instrument ID:	Tekmar AUTOCAN/Agilent 5975Binert/6890N/MS13	Date Received: NA
Analyst:	Chris Cornett	Date Analyzed: 7/3/13
Sample Type: Test Notes:	6.0 L Summa Canister	Volume(s) Analyzed: 0.125 Liter(s)

					ALS	
CAS #	Compound	Spike Amount	Result	% Recovery	Acceptance	Data
		$\mu g/m^3$	μg/m³		Limits	Qualifier
1634-04-4	Methyl tert-Butyl Ether	204	173	85	69-120	
110-54-3	n-Hexane	206	157	76	63-115	
107-06-2	1,2-Dichloroethane	208	161	77	69-118	
71-43-2	Benzene	208	182	88	69-117	
108-88-3	Toluene	208	195	94	65-116	
106-93-4	1,2-Dibromoethane	208	204	98	69-130	
100-41-4	Ethylbenzene	206	200	97	66-119	
179601-23-1	m,p-Xylenes	412	388	94	64-118	
95-47-6	o-Xylene	200	191	96	65-120	
108-67-8	1,3,5-Trimethylbenzene	208	203	98	64-125	
95-63-6	1,2,4-Trimethylbenzene	200	201	101	64-131	
91-20-3	Naphthalene	178	185	104	56-143	

Laboratory Control Sample percent recovery is verified and accepted based on the on-column result. Reported results are shown in concentration units and as a result of the calculation, may vary slightly.

APPENDIX D

Terrestrial Ecological Evaluation Form



Voluntary Cleanup Program

Washington State Department of Ecology Toxics Cleanup Program

TERRESTRIAL ECOLOGICAL EVALUATION FORM

Under the Model Toxics Control Act (MTCA), a terrestrial ecological evaluation is necessary if hazardous substances are released into the soils at a Site. In the event of such a release, you must take one of the following three actions as part of your investigation and cleanup of the Site:

- 1. Document an exclusion from further evaluation using the criteria in WAC 173-340-7491.
- 2. Conduct a simplified evaluation as set forth in WAC 173-340-7492.
- 3. Conduct a site-specific evaluation as set forth in WAC 173-340-7493.

When requesting a written opinion under the Voluntary Cleanup Program (VCP), you must complete this form and submit it to the Department of Ecology (Ecology). The form documents the type and results of your evaluation.

Completion of this form is not sufficient to document your evaluation. You still need to document your analysis and the basis for your conclusion in your cleanup plan or report.

If you have questions about how to conduct a terrestrial ecological evaluation, please contact the Ecology site manager assigned to your Site. For additional guidance, please refer to www.ecy.wa.gov/programs/tcp/policies/terrestrial/TEEHome.htm.

Step 1: IDENTIFY HAZARDOUS WASTE SITE

Please identify below the hazardous waste site for which you are documenting an evaluation.

Facility/Site Name: 781 Walker Way, Port Ludlow

Facility/Site Address: 781 Walker Way, Port Ludlow, WA 98365

Facility/Site No:

VCP Project No.: SW1311

Title: Sr. Environmental Scientist

Step 2: IDENTIFY EVALUATOR

Please identify below the person who conducted the evaluation and their contact information.

Name:	Brett	Carn
iname.	DIEU	Carp

Organization: Aspect Consulting

Mailing address: 401 2nd Avenue South, #201

City: Seattle			te: WA	Zip code: 98104
Phone: 206-838-5836	Fax: 206-838-5853		E-mail: bcarp	@aspectconsulting.com

Step 3: DOCUMENT EVALUATION TYPE AND RESULTS					
A. Exclusion from further evaluation.					
1. Does the Site qualify for an exclusion from further evaluation?					
Yes If you answered " YES ," then answer Question 2 .					
□ No or If you answered " NO" or "UKNOWN," then skip to Step 3B of this form.					
2. What is the basis for the exclusion? Check all that apply. Then skip to Step 4 of this form.					
Point of Compliance: WAC 173-340-7491(1)(a)					
All soil contamination is, or will be,* at least 15 feet below the surface.					
All soil contamination is, or will be,* at least 6 feet below the surface (or alternative depth if approved by Ecology), and institutional controls are used to manage remaining contamination.					
Barriers to Exposure: WAC 173-340-7491(1)(b)					
All contaminated soil, is or will be,* covered by physical barriers (such as buildings or paved roads) that prevent exposure to plants and wildlife, and institutional controls are used to manage remaining contamination.					
Undeveloped Land: WAC 173-340-7491(1)(c)					
 There is less than 0.25 acres of contiguous[#] undeveloped[±] land on or within 500 feet of any area of the Site and any of the following chemicals is present: chlorinated dioxins or furans, PCB mixtures, DDT, DDE, DDD, aldrin, chlordane, dieldrin, endosulfan, endrin, heptachlor, heptachlor epoxide, benzene hexachloride, toxaphene, hexachlorobenzene, pentachlorophenol, or pentachlorobenzene. 					
For sites not containing any of the chemicals mentioned above, there is less than 1.5 acres of contiguous [#] undeveloped [±] land on or within 500 feet of any area of the Site.					
Background Concentrations: WAC 173-340-7491(1)(d)					
Concentrations of hazardous substances in soil do not exceed natural background levels as described in WAC 173-340-200 and 173-340-709.					
 * An exclusion based on future land use must have a completion date for future development that is acceptable to Ecology. [±] "Undeveloped land" is land that is not covered by building, roads, paved areas, or other barriers that would prevent wildlife from feeding on plants, earthworms, insects, or other food in or on the soil. 					
* "Contiguous" undeveloped land is an area of undeveloped land that is not divided into smaller areas of highways, extensive paving, or similar structures that are likely to reduce the potential use of the overall area by wildlife.					

в	B. Simplified evaluation.					
1. Does the Site qualify for a simplified evaluation?						
	□ Y	If you answered "YES," then answer Question 2 below.				
	☐ N Unkn	If you answered "NO" or "UNKNOWN," then skip to Step 3C of this form.				
2. Did you conduct a simplified evaluation?						
	🗌 Y	If you answered "YES," then answer Question 3 below.				
	□ N	If you answered "NO," then skip to Step 3C of this form.				
3. Was further evaluation necessary?						
	□ Y	If you answered "YES," then answer Question 4 below.				
		lo If you answered " NO ," then answer Question 5 below.				
4. If further evaluation was necessary, what did you do?						
	Used the concentrations listed in Table 749-2 as cleanup levels. <i>If so, then skip to</i> Step 4 of this form.					
		Conducted a site-specific evaluation. If so, then skip to Step 3C of this form.				
5. If no further evaluation was necessary, what was the reason? Check all that apply. Then skip to Step 4 of this form.						
	Exposure Analysis: WAC 173-340-7492(2)(a)					
		Area of soil contamination at the Site is not more than 350 square feet.				
	Current or planned land use makes wildlife exposure unlikely. Used Table 749-1.					
	Pathway Analysis: WAC 173-340-7492(2)(b)					
	No potential exposure pathways from soil contamination to ecological rece					
	Contaminant Analysis: WAC 173-340-7492(2)(c)					
		No contaminant listed in Table 749-2 is, or will be, present in the upper 15 feet at concentrations that exceed the values listed in Table 749-2.				
		No contaminant listed in Table 749-2 is, or will be, present in the upper 6 feet (or alternative depth if approved by Ecology) at concentrations that exceed the values listed in Table 749-2, and institutional controls are used to manage remaining contamination.				
		No contaminant listed in Table 749-2 is, or will be, present in the upper 15 feet at concentrations likely to be toxic or have the potential to bioaccumulate as determined using Ecology-approved bioassays.				
alternative depth if approved by Ecology) at the potential to bioaccumulate as determined		No contaminant listed in Table 749-2 is, or will be, present in the upper 6 feet (or alternative depth if approved by Ecology) at concentrations likely to be toxic or have the potential to bioaccumulate as determined using Ecology-approved bioassays, and institutional controls are used to manage remaining contamination.				

_						
C. Site-specific evaluation. A site-specific evaluation process consists of two parts: (1) formulating the problem, and (2) selecting the methods for addressing the identified problem. Both steps require consultation with and approval by Ecology. <i>See</i> WAC 173-340-7493(1)(c).						
1.	1. Was there a problem? See WAC 173-340-7493(2).					
	□ Y	Yes If you answered "YES," then answer Question 2 below.				
	□ N	o <i>lf you ans</i> below:	If you answered " NO," then identify the reason here and then skip to Questi below:			
			No issues were identified during the problem formulation step.			
			While issues were identified, those issues were addressed by the cleanup actions for protecting human health.			
2.	2. What did you do to resolve the problem? See WAC 173-340-7493(3).					
	Used the conc Question 5 be		entrations listed in Table 749-3 as cleanup levels. If so, then skip to elow.			
			nore of the methods listed in WAC 173-340-7493(3) to evaluate and entified problem. <i>If so, then answer Questions 3 and 4 below.</i>			
3.	3. If you conducted further site-specific evaluations, what methods did you use? <i>Check all that apply.</i> See WAC 173-340-7493(3).					
		Literature surveys.				
		Soil bioassays.				
		Wildlife exposure model.				
		Biomarkers.				
		Site-specific field studies.				
		Weight of evid	ence.			
		Other methods approved by Ecology. If so, please specify:				
4.	4. What was the result of those evaluations?					
		Confirmed the	re was no problem.			
		Confirmed the	re was a problem and established site-specific cleanup levels.			
5. Have you already obtained Ecology's approval of both your problem formulation and problem resolution steps?						
	Yes If so, please identify the Ecology staff who approved those steps:					
	□ No					

Step 4: SUBMITTAL

Please mail your completed form to the Ecology site manager assigned to your Site. If a site manager has not yet been assigned, please mail your completed form to the Ecology regional office for the County in which your Site is located.



If you need this publication in an alternate format, please call the Toxics Cleanup Program at 360-407-7170. Persons with hearing loss can call 711 for Washington Relay Service. Persons with a speech disability can call 877-833-6341.

APPENDIX E

Review of Potential Remediation Technologies for Petroleum-Impacted Sites

E.1 Institutional Controls

Institutional controls are measures to limit or prohibit activities that may interfere with the integrity of a cleanup action or result in exposure to hazardous substances. Examples of institutional controls are limitations on the use of the property or resources such as an environmental covenant or maintenance requirements for engineering controls.

Advantages

- Can be easy to implement without disrupting operations.
- Relatively low cost.

Limitations

- Does not address the destruction or remediation of COCs.
- Depending on site-specific conditions, may not be sufficient to prevent offproperty migration of COCs.
- May result in restrictions on the property use.

Summary Evaluation

Because of its low cost and ease of implementation, institutional controls can be a valuable component of a remediation strategy at sites impacted with petroleum hydrocarbons.

E.2 Engineering Controls

Engineering controls means containment and/or treatment systems that are designed and constructed to prevent or limit the movement of, or the exposure to, hazardous substances. An example of an engineering control would be a physical barrier such as asphalt or concrete paving/capping.

Advantages

- Can be easy to implement without disrupting operations.
- Can be relatively low cost.

Limitations

- Does not address the destruction or remediation of COCs.
- Depending on site-specific conditions, may not be sufficient to prevent offproperty migration of COCs.
- May result in restrictions on the property use.

Summary Evaluation

Because of its low cost and ease of implementation, engineering controls can be a valuable component of a remediation strategy for petroleum-impacted sites.

E.3 Monitored Natural Attenuation

Monitored natural attenuation involves the destruction of COCs in site soil and groundwater by *in situ* by natural processes, including as biodegradation by native bacteria. This technology typically involves periodic monitoring of soil, groundwater, and/or air to evaluate remediation progress and ensure continued protectiveness.

Advantages

- COCs are permanently destroyed.
- Easy to implement without disrupting operations.
- Relatively low cost.

Limitations

- Depending on site-specific conditions, remediation may take an extended time period.
- Depending on site-specific conditions, may not be sufficient to prevent offproperty migration of COCs.

Summary Evaluation

Because of its low cost and ease of implementation, monitored natural attenuation can be a valuable component of a remediation strategy at petroleum-impacted sites.

E.4 Soil Vapor Extraction

Soil vapor extraction involves removal of COCs in site soils above the water table by applying a vacuum to wells and treating constituents removed in the extracted soil gas. Equipment required with this technology includes wells, piping, a vacuum blower, moisture knockout pot, and treatment equipment (e.g., activated carbon vessels). Operation requirements include electricity for the vacuum blower, natural gas for the catalytic oxidizer, disposal of generated wastes (condensate water), equipment maintenance, and air monitoring.

Advantages

- COCs are permanently destroyed.
- Relatively non-disruptive technology (will require temporary disturbance to install wells and piping).
- Area of treatment can extend underneath otherwise inaccessible facility areas.

Limitations

- Removal of COCs from low-permeability soils can be limited by the rate of diffusion through these soils.
- Not effective in groundwater or soil below the water table.

Summary Evaluation

Because of its moderate cost and ease of implementation, soil vapor extraction can be a valuable component of a remediation strategy at petroleum-impacted sites. However, this technology was pilot tested at the Site between December 2011 and January 2012 and overall performance was poor. Initial soil vapor extraction mass removal rates were relatively low, and removal rates declined rapidly over the testing period, even with the addition of groundwater extraction (dual-phase extraction).

E.5 Air Sparging

Air sparging involves removal of COCs from groundwater and saturated soil and by injecting air in wells screened below the water table. Volatile contaminants evaporate into the injected air, which is typically collected and treated by through soil vapor extraction (see E.4 above). Equipment required with this technology includes wells, piping, and an air compressor. Operation requirements include electricity for the air compressor, equipment maintenance, and air monitoring.

Advantages

- COCs are permanently removed and destroyed (if collected/treated with soil vapor extraction).
- Relatively non-disruptive technology (will require temporary disturbances to install wells and piping).
- Area of treatment can extend underneath otherwise inaccessible facility areas.

Limitations

- Heterogeneous geology may limit the rate of diffusion through low-permeability soil layers and reduce treatment effectiveness.
- Preferential pathways for subsurface air movement may still result in incomplete treatment in some areas.
- Heterogeneous soils, especially low permeability zones above the air injection level, can make recovery of sparged air problematic and can result in unpredictable subsurface migration of contaminated soil vapor.

Summary Evaluation

Because of its moderate cost and ease of implementation, air sparging can be a valuable component of a remediation strategy at petroleum-impacted sites. However, given the heterogeneous subsurface soils present at the Site, and the transient nature of perched groundwater, both the implementability of air sparging, and the recovery of sparged vapors, would be problematic at the Site.

E.6 Enhanced Aerobic Biodegradation

Enhanced aerobic biodegradation is the practice of adding oxygen (an electron acceptor) to groundwater and/or soil to increase the number and vitality of indigenous microorganisms already naturally performing biodegradation of COCs at the Site. Application is typically accomplished via injection of a liquid compound to provide oxygen to the subsurface. This process is performed in several discrete injection events and does not require continuously-operating equipment on site.

Advantages

- COCs are permanently destroyed *in situ*.
- Easy to implement without significantly disrupting operations.
- Can enhance remediation in otherwise inaccessible areas by altering groundwater conditions over a localized area.

Limitations

- Although faster than natural attenuation, remediation will likely be limited by the rate at which COCs desorb from soil. Therefore, remediation time with this technology may be a decade or more.
- Generally not effective in soil above the water table.

Summary Evaluation

Enhanced aerobic biodegradation is not typically cost-effective for source removal, but is most applicable as a polishing technology. Although faster than unassisted natural attenuation, this is typically still a slow process that can take a number of years to destroy COCs, may require multiple injection events, and is generally employed following treatment of saturated soil and groundwater by more aggressive technologies.

E.7 In Situ Chemical Oxidation

For chemical oxidization, a strong oxidizing chemical (e.g., ozone, Fenton's reagent, activated persulfate, permanganate) is injected into groundwater or mixed into soil to react and mineralize (i.e., convert to carbon dioxide and water) organic contaminants. Ozone is typically applied in gas form as part of air sparging; Fenton's reagent and activated persulfate are typically injected as liquid solutions into groundwater.

Advantages

- COCs are permanently destroyed in situ.
- May not require installation of permanent wells, piping, or equipment which may help minimize disruption to business operations.
- Potential area of influence could extend underneath inaccessible areas of the Site.

Limitations

- Generally not effective in soil above the water table.
- Low-permeability soils may not be adequately addressed.

Summary Evaluation

The success of this technology is highly dependent on the chemical oxidant physically coming into contact and reacting with COCs in soil and groundwater. The presence of heterogeneous soils and low-permeability zones can limit the ability of this technology to completely treat impacted soil and groundwater.

E.8 Dual-Phase Extraction

This approach uses soil vapor extraction in conjunction with groundwater pumping to depress the water table, which exposes shallow saturated soils to treatment by soil vapor extraction, and provides hydraulic containment and removal of COCs in site groundwater. To increase effectiveness, this technology can be applied in conjunction with air sparging to provide additional groundwater treatment. In addition to equipment required by soil vapor extraction, this technology requires either submersible pumps or a high-vacuum blower to remove water, and additional treatment equipment. Water disposal can require obtaining a sewer discharge authorization and possibly treatment prior to discharge.

Advantages

- COCs are permanently removed and destroyed.
- Provides hydraulic control of chemical migration as well as on-site treatment.
- Area of influence from pumping can extend underneath inaccessible areas.

Limitations

- Heterogeneous soils and low-permeability zones can complicate application, increase costs, and result in incomplete treatment.
- High soil permeability can result in the need to remove and treat large volumes of water to adequately depress the water table.
- Requires significant above-ground space for required equipment.
- Can have relatively high cost for water disposal.

Summary Evaluation

The presence of heterogeneous soils and low-permeability zones can limit the ability of this technology to completely treat impacted soil and groundwater. This technology was pilot tested at the Site between December 2011 and January 2012 and overall performance was poor. Water removal rates were low, consistent with the dense, low permeability glacial soil of the shallow perched aquifer. Initial mass removal rates were relatively low, and mass removal rates declined rapidly over the testing period.

E.9 Soil Excavation

This technology involves removing contaminated soils and transporting the soil to a permitted disposal facility (e.g., landfill or soil recycler). Soil can be removed by a variety of techniques; shallow soil is typically removed with an excavator, while deeper soil may be removed using overlapping augers or a shored excavation.

Advantages

- For soil that can be accessed, this is the most certain method of removing COCs from the Site.
- For shallow impacted soils, excavation coupled with off-site disposal is typically the most cost-effective active remedial measure.

Limitations

- Excavation costs increase significantly with depth and proximity to load bearing structures and buildings, particularly when shoring is required.
- Impacted soil beneath buildings and other facilities typically requires the demolition of those structures to access soil.
- Excavation is potentially disruptive, particularly when the removal is not consistent with site development plans.

Summary Evaluation

Shallow impacted soil was removed the extent practical at the Site in 1990. Removal of additional deeper impacted soil at the Site would require a deep, sloped excavation. Demolition and replacement of the existing shop/garage building would also be required.

APPENDIX F

SLR Soil Vapor Extraction Pilot Test Report (on CD)



May 8, 2012 Project 101.00433.00003

Mr. Larry Smith Olympic Water & Sewer, Inc. 70 Breaker Lane Port Ludlow, Washington 98635

Re: Soil Vapor Extraction Pilot Test Report, Olympic Water & Sewer, Inc. Facility, 781 Walker Way, Port Ludlow, Washington

Dear Larry:

SLR International Corporation (SLR) has prepared this report to present the results of the soil vapor extraction (SVE) pilot tests that were recently conducted at the Olympic Water & Sewer, Inc. (OWSI) facility in Port Ludlow, Washington. The purposes of the pilot tests were to: 1) assess the potential effectiveness of SVE, with and without groundwater de-watering, at remediating the volatile petroleum hydrocarbon-impacted soil, 2) evaluate if groundwater extraction can effectively de-water the shallower portion of saturated zone (above the Unit B silty sand), expose the capillary fringe above the deeper water table (above the Unit D silts), and allow SVE to remove additional petroleum hydrocarbon vapors from the shallower saturated zone and the capillary fringe, and 3) obtain the information necessary for potential future design and implementation of an SVE system, with or without groundwater recovery/treatment.

PROPERTY DESCRIPTION

The subject property, which is owned by OWSI, is located at 781 Walker Way in Port Ludlow, Washington. The location of the property is shown on Figure 1. The OWSI property is an approximate 2.2-acre parcel that is partially developed with an OWSI operation and maintenance facility. The facility consists of an approximate ½-acre area that includes an office/shop/garage building (garage building), a public water supply well (Well #2) and associated pump house building, and a storage trailer (see Figure 2). The ground surface within the facility is primarily unpaved, except for a narrow asphalt driveway that runs down the center of the facility from Walker Way to approximately the storage trailer. Within the OWSI property, the facility is surrounded on all sides by dense forest, and a gulley containing an intermittent stream is located within the forest to the west of the facility. The facility has been operating since 1968, after the installation of the water supply well. Prior to 1968, the property was undeveloped.

SUMMARY OF ENVIRONMENTAL CONDITIONS

In 1990, the three gasoline underground storage tanks (USTs) at the property were removed. A 1,000-gallon UST was located beneath the floor of the northern garage bay, a 2,000-gallon UST was located outside (west) of the northern garage bay, and a 2,000-gallon UST was located approximately 40 feet south of the garage. The approximate locations of the UST excavations are shown on Figure 2. Volatile petroleum hydrocarbon-impacted soil was only present in the 1,000-gallon tank excavation and the northern 2,000-gallon tank excavation [Applied Geotechnology, Inc. (AGI), 1991]. Excavation activities were conducted to remove the impacted soil near the tanks; however, due to structural concerns for the building, the excavation at the former 1,000-gallon tank could only be extended to a depth of approximately 10 feet below ground surface (bgs).

During subsequent environmental investigations at the property, a total of 13 soil borings were drilled and sampled. The approximate locations of the borings are shown on Figure 2. The soil sample analytical results from the 1990 gasoline UST removals (including the subsequent soil excavations) (AGI, 1991), and the field screening and soil sample analytical results from the subsurface investigations at the property (SLR, 2010; and SLR, 2011) indicate that the former northern gasoline UST area is the source of the remaining volatile petroleum hydrocarbon-impacted soil and groundwater beneath the property. The hydrocarbon-impacted soil occurs at the base of the former 1,000-gallon UST excavation (at approximately 10 feet bgs) and extends to the saturated zone above the Unit B silty sands (approximately 20 feet bgs) near the garage building. The estimated area of impacted soil extends beyond the western, eastern, and southern ends of the garage building and covers an area of approximately 3,140 square feet. The estimated area of the hydrocarbon-impacted soil is shown on Figure 3.

Well #2, which is located at the northern portion of the property, is screened at depths ranging from 214 to 245 feet bgs. After 40 years of operation, the yield of Well #2 was decreasing, and in 2009, OWSI decided to install a replacement well (designated Well #17) at the southern part of the facility. The planned construction of Well #17 was similar to the construction of Well #2. On April 21, 2009, during the drilling of Well #17, the driller noticed a gasoline odor emanating from the well casing at a depth of approximately 50 feet [Robinson Noble & Saltbush, Inc. (Robinson Noble), 2009]. The drilling was discontinued, and soil and groundwater samples were collected from the bottom of the casing for laboratory analysis. The analytical results showed that the groundwater sample contained gasoline-range organics (GRO) and benzene concentrations [5,530 and 948 micrograms per liter (μ g/L), respectively] that exceeded the Model Toxics Control Act (MTCA) Method A groundwater cleanup levels (800 and 5 μ g/L, respectively). Due to the presence of the

gasoline-impacted groundwater, the well was not completed and the casing was capped. The location of the casing for Well #17 is shown on Figure 2.

During the 2010 and 2011 subsurface investigations, five of the soil borings were completed as groundwater monitoring wells (MW-1, MW-2, MW-3, MW-4, and MW-5), and groundwater sampling events were conducted in June 2010, October 2010, and April 2011. In addition, the soil borings that were completed as SVE points (SVE-1 and SVE-2) contained shallow groundwater, and groundwater samples were collected from the SVE points in April 2011. The locations of the monitoring wells and SVE points are shown on Figure 4. The groundwater sample analytical results showed that all of the samples from MW-1 and MW-2 and the April 2011 sample from SVE-1 contained benzene and GRO concentrations (up to 2,100 and 34,000 μ g/L, respectively) that exceeded the MTCA Method A cleanup levels (SLR, 2010; and SLR, 2011). The April 2011 samples from MW-4 and SVE-2 contained benzene and GRO concentrations, respectively, that exceeded the Mtthe Mtthe A cleanup levels.

Based on the groundwater sample analytical results from the 2009 drilling of Well #17 (Robinson Noble, 2009) and the 2010 and 2011 subsurface investigations (SLR, 2010; and SLR, 2011), petroleum hydrocarbon concentrations greater than the MTCA Method A groundwater cleanup levels occur near the source area (former northern gasoline UST area) and primarily extend to the south and southwest. The estimated area of the hydrocarbon-impacted groundwater is shown on Figure 4. The greatest petroleum hydrocarbon concentrations occurred in the shallow saturated zone above the Unit B silty sands beneath the western edge of the garage building (at SVE-1). Based on the presence of petroleum hydrocarbons at wells MW-4 and MW-5, there is also a limited component of impacted groundwater migration (likely above Unit B) to the north and east.

PROPERTY GEOLOGY AND HYDROGEOLOGY

The subsurface investigations indicate that the soils beneath the subject property consist of surficial gravel fill underlain by dense glacial advance outwash (sand, gravel, and silt units) with interbedded lacustrine silts to the maximum depth drilled (approximately 60 feet bgs). The distributions of subsurface soils (grouped by lithologies into Units A through E) are shown on Figure 5.

Unit A extends beneath all of the investigated portions of the property and varies from approximately 29 to 43 feet in thickness. Unit A primarily consists of very fine to fine sand or gravelly sand, and includes silty sands and a 5- to 10-foot-thick sandy silt to silt interbed beneath some portions of the property. Unit A is underlain by Unit B beneath the northeastern and eastern parts of the property (at MW-1, MW-4, MW-5, SB-1, SVE-1, and SVE-2), by Unit C beneath the central part of the property (at MW-2), and by Unit D beneath

the southern part of the property (at MW-3). Unit B consists of silty sand that is up to 11 feet thick where encountered, and is underlain by Unit D. Unit C was only encountered at MW-2, and consists of gravel with cobbles. Unit C has a maximum thickness of 12 feet and is underlain by Unit D. Unit D extends beneath all of the investigated portions of the property, consists of hard silt, clayey silt, and gravelly silt, and ranges from approximately 15 to more than 23 feet in thickness. In the central part of the OWSI facility (at MW-1 and MW-2), the top of Unit D occurs at an elevation of approximately 251 feet above the NAVD 88 datum, while at the northern and southern parts of the facility and to the east of the facility (at MW-4, MW-3, and MW-5, respectively), Unit D occurs at higher elevations (approximately 260 to 263 feet above the NAVD 88 datum). At MW-2, MW-3, MW-4, and SVE-2, Unit D contains an interbed of sand with gravel and cobbles that is approximately 5 to 7.5 feet thick. A very dense, very fine to fine silty sand that underlies Unit D at the bottom of the deepest boring at the facility (MW-1) is identified as Unit E.

According to the driller's log for the water supply well (Well #2) located in the northern part of the property (see Figure 2), a thick sequence of clays with cemented gravels extends from approximately 49 to 215 feet bgs (SLR, 2010). The deep water-bearing units at Well #2 occur at depths between 215 and 245 feet bgs.

In general, sands and gravels are significantly more permeable than silty sands, silts, sandy silts, clayey silts, and gravelly silts. Therefore, Unit A and Unit C are interpreted as being relatively permeable soils that provide preferential pathways for groundwater flow. Units B, D, and E are interpreted as being relatively impermeable soils that act as local aquitards. The uppermost clay identified in the driller's log for Well #2 is a "sandy clay" that extends from about 49 to 79 feet bgs, which is generally consistent with the depths of Units D and E. This suggests that an aquitard consisting of Unit B, Unit D, Unit E, and associated underlying fine-grained soils extends beneath the facility, has a minimum 30-foot thickness, and may locally exceed 55 feet in thickness.

On December 12, 2011, prior to start-up of the first SVE pilot test, groundwater levels were measured in all of the monitoring wells and SVE points at the facility. The depths to groundwater ranged from 25.82 to 40.69 feet below the tops of the well casings, and the groundwater elevations ranged from 253.10 to 270.14 feet above the NAVD 88 datum. The groundwater elevations at several of the wells and SVE-1 are shown on Figure 5. The groundwater table beneath the property occurs as a single, hydraulically continuous saturated zone; however, the depth and elevation of the water table vary significantly beneath the property. The water table position appears to be primarily controlled by the elevation of the uppermost laterally-extensive aquitard. Beneath the northern and eastern portions of the facility and east of the facility, the uppermost aquitard is Unit B, the aquitard elevation is about 260 to 270 feet above the NAVD 88 datum, and the water table intercepts either Unit A sands or underlying Unit B silty sand (this shallower area of the water table is monitored at

MW-4, MW-5, SVE-1, and SVE-2). Beneath the south-central and southern parts of the facility, the uppermost aquitard unit is Unit D, the aquitard elevation is typically below 260 feet, and a deeper area of the water table intercepts Unit C gravel at MW-2 and either Unit A sands or underlying Unit D silts at MW-3 (the deeper water table area). Well MW-1, in the north-central part of the facility (approximately 10 feet southwest of SVE-1), monitors the deeper water table within the aquitard units (Unit B and Unit D).

Within the shallow water table area, groundwater is expected to primarily flow laterally above the aquitard within the relatively permeable Unit A sands towards the deeper water table area. Within the deeper water table area, groundwater is expected to primarily flow laterally above the aquitard within the relatively permeable Unit C gravels. The elevations of both the aquitard and the water table beneath the deeper water table area in the central part of the facility (near MW-2) is about 9 to 15 feet lower than at the northern, southern, and eastern parts of the facility. Based on the known topography of the uppermost aquitard surface, the water table elevations, and the area of petroleum hydrocarbon-impacted groundwater (see Figure 4), there appears to be a groundwater flow component beneath the facility to the south-southwest.

SVE PILOT TESTS

From December 12, 2011 through January 5, 2012, SLR conducted four SVE pilot tests to: 1) assess the potential effectiveness of SVE, with and without groundwater de-watering, at remediating the remaining volatile petroleum hydrocarbon-impacted soil, 2) evaluate if groundwater extraction can effectively de-water the shallower portion of the saturated zone (above the Unit B silty sand), expose the capillary fringe above the deeper water table (above the Unit D silts), and allow SVE to remove additional petroleum hydrocarbon vapors from the shallower saturated zone and the capillary fringe, and 3) obtain the information necessary for potential future design and implementation of an SVE system, with or without groundwater recovery/treatment.

Prior to conducting the first three pilot tests, SLR obtained verbal authorization from the Olympic Region Clean Air Agency (ORCAA) to conduct the three shorter pilot tests without treatment of the extracted soil vapors. However, for the extended pilot test (Pilot Test #4), ORCAA required treatment of the extracted vapors prior to emission to the atmosphere.

Each pilot test consisted of connecting a 3-horsepower Rotron EN523 regenerative blower to the selected extraction point, and applying the maximum vacuum pressure created by the blower. After applying the vacuum, SLR personnel measured the flow rate of the extracted air within a 2-inch-diameter influent pipe to the blower by using an anemometer, and monitored the approximate petroleum hydrocarbon concentrations in the extracted and emitted soil vapors by using a photoionization detector (PID). During each test, SLR also measured the vacuum

pressures at the SVE points and at all of the monitoring wells with magnehelic gauges to evaluate the radius of applied vacuum influence in the subsurface. Prior to beginning and during each test, SLR measured the depths to groundwater in all of the SVE points and monitoring wells by using an electronic water level indicator. The field data collected during each test are described below and copies of the field notes that include the data are attached.

At initiation and completion of Pilot Tests #1 and #2, a sample of the extracted soil vapors was collected for laboratory analysis. During Pilot Test #3, an extracted soil vapor sample was only collected at test initiation, and during Pilot Test #4, the samples were collected after one week of operation and at completion of the test. During the initiation of the first three tests, the samples were collected at the emission stack. Immediately after completing each test, the samples were collected at the influent pipe to the blower. We had planned to collect all of the samples at the influent pipe to the blower; however, the samples were collected by using a gas sampling bulb with new tubing, and due to the high vacuum pressures, the sampling method could not overcome the applied vacuum while the blower was operating. Therefore, we collected the test initiation samples from the emission stack (under positive pressure after the blower). To allow for sample collection under ambient pressure during Pilot Test #4, the blower was temporarily shut off on December 29, 2011, and a sample was collected at the influent pipe instead of the stack. All of the extracted soil vapor samples were submitted to Friedman & Bruya, Inc. (F&B) in Seattle, Washington, for analysis of benzene, toluene, ethylbenzene, and total xylenes (BTEX) by EPA Method 8021B and GRO by Ecology Method NWTPH-Gx. The samples analytical results are described below and presented in Table 1. Copies of the laboratory reports are attached.

Pilot Test #1

Pilot Test #1 was conducted on December 12, 2011, and a vacuum pressure of approximately 97 inches of water column was initially applied to soil vapor extraction point SVE-1. Groundwater de-watering was not conducted during the test. The airflow rate was only 18 cubic feet per minute (cfm) and the PID readings decreased from 89.1 to 12.1 parts per million (ppm) in 70 minutes. The extracted soil vapor sample collected at system initiation (Test1-Sample1) contained a GRO concentration of 30 milligrams per cubic meter [mg/m³; converted to 6.9 ppm by volume (ppm-v)]. To expose more of the SVE point screen during the test, the applied pressure was reduced to 60 inches of water column after the second hour of the test by opening a dilution valve on the influent pipe to the blower. After reducing the pressure, the airflow rate (prior to the dilution valve) decreased to approximately 14 to 16 cfm, and the first PID reading was only 7.1 ppm. The PID readings were measured at the emission stack, except for two occasions during the last 100 minutes of the test when we temporarily shut off the blower and measured the PID readings at the influent pipe to the blower (prior to the dilution valve). The PID readings at the influent pipe to the blower (prior to the dilution valve).

extracted soil vapor sample (Test1-Sample2) at the completion of the test (collected at the influent pipe) only contained a GRO concentration of 47 mg/m³ (10.8 ppm-v).

After almost four hours of operation, the test was terminated. During the test, vacuum pressures were not measured at any of the monitoring wells or at soil vapor extraction point SVE-2.

Pilot Test #2

On December 13, 2011, the pilot test at SVE-1 was repeated; however, to assess the effects of de-watering the shallower saturated zone above the Unit B silty sand, a Grundfos Redi-Flo 2 submersible pump was installed in SVE-1. The extracted water was pumped into a 4,000-gallon tank for temporary storage, and a totalizing flow meter was used to monitor the groundwater pumping rate.

The initial groundwater pumping rate was approximately 2.5 gallons per minute (gpm); however, after removing the water from the well and associated gravel pack, the pumping rate decreased to approximately 0.13 to 0.19 gpm for the rest of the test. During the 9-hour-long test, a total of 77.1 gallons of groundwater were pumped. The groundwater drawdown in SVE-1 was approximately 8.80 feet, and by the end of the test, groundwater drawdowns of 1.62, 0.64, 0.50, and 0.67 feet were observed in SVE-2, MW-1, MW-4, and MW-5, respectively. SVE-2, MW-4, and MW-5, which are at least partially screened within the shallow saturated zone above Unit B, are located approximately 31 to 72 feet from SVE-1 (see Figure 4). MW-1 is screened within the deeper saturated zone above and within the Unit D silts, and it is located approximately 9 feet from SVE-1. Groundwater drawdown was not observed at wells MW-2 or MW-3. MW-2 and MW-3 are screened within the deeper saturated zone above and/or within Unit D, and are located approximately 63 and 150 feet, respectively, from SVE-1. The extensive radius of pumping influence after extracting only 77 gallons of water indicates that recharge to the shallow groundwater above Unit B, within the test area, was limited. The groundwater drawdown in MW-1 demonstrates the hydraulic continuity between the shallower saturated zone above Unit B and the deeper water table above Unit D.

During Pilot Test #2, a vacuum pressure of approximately 97 inches of water column was applied to SVE-1. The airflow rate was 16 to 18 cfm and the PID readings at the emission stack decreased from approximately 80 to 4 ppm during the test. The extracted soil vapor sample (Test2-Sample1) at system initiation (at the stack) did not contain detectable BTEX or GRO concentrations; however, the sample (Test2_Sample2) collected at system termination (at the influent pipe) contained a GRO concentration of 1,900 mg/m³ (437 ppm-v).

During the test, vacuum pressures (0.30 to 1.0 inches of water column) were detected at SVE-2, but not in any of the monitoring wells. SVE-2 is located approximately 31 feet from

SVE-1 (see Figure 4). Vacuum pressures were consistently present at SVE-2 after conducting the test for approximately two hours.

After approximately five hours of the test, SLR collect a sample of the extracted groundwater for laboratory analysis. The sample was submitted to F&B for analysis of BTEX by EPA Method 8021B and GRO by Ecology Method NWTPH-Gx. The sample contained benzene and GRO concentrations of 430 and 23,000 μ g/L, respectively. The water sample analytical results are presented in Table 2 and a copy of the laboratory report is attached.

Pilot Test #3

On December 14, 2011, an additional SVE pilot test with groundwater extraction was conducted at monitoring well MW-1, which is screened through the deeper water table above Unit D. This test was conducted to: 1) assess the potential effectiveness of combined SVE and groundwater extraction at removing gasoline from the capillary fringe of the deeper water table (above Unit D), and 2) further assess the hydraulic connection between the shallow saturated zone above Unit B and the deeper water table above Unit D. This test followed the same procedures as Pilot Test #2.

The initial groundwater pumping rate was approximately 2.5 gpm; however, after removing the water from the well and associated gravel pack, the pumping rate decreased to a rate that could not be measured by the flow meter. After approximately 7 hours, we moved the test to MW-2 to find out if the very low pumping rate at MW-1 was due to localized geologic conditions. At MW-2, the pumping rate quickly decreased from 2.5 to 0.5 gpm; however, after 10 minutes, we took the sample port off of the influent pipe to the blower and the open hole appeared to slightly increase the pumping rate. To assess the possible scenario that a vacuum lock had formed on MW-1 and was inhibiting groundwater flow into the well, we moved the test back to MW-1 and took the sample port off of the influent pipe. After removal of the sample port, the pumping rate did not increase over a period of approximately 20 minutes.

After the pumping rate did not increase at MW-1, we moved the test back to MW-2 to further assess the yields of the deeper saturated zone above Unit D. The pumping rate at MW-2 was approximately 0.2 gpm for a period of one hour. During the approximate 9-hour-long test at MW-1 and MW-2, a total of only 19.8 gallons of groundwater were recovered, and it appears that the yield of the deeper saturated zone above Unit D is up to 0.2 gpm. The lower pumping rate at MW-1 represents the geologic conditions near the well, and not a potential vacuum lock on the well during the test.

The groundwater drawdown in MW-1 was approximately 8.90 feet, and by the end of the test, groundwater drawdown was not observed in the SVE points or any of the other monitoring wells, except MW-2 due to the temporary pumping at that well. There was also no observed

drawdown in the SVE points or any of the other monitoring wells during the pumping at MW-2.

During Pilot Test #3, a vacuum pressure of approximately 95 to 97 inches of water column was applied to MW-1 or MW-2. The airflow rates were 16 to 18 cfm and the PID readings at the emission stack were consistently less than 3 ppm during the test. The extracted soil vapor sample (Test3_Sample1) at system initiation contained a GRO concentration of 85 mg/m³ (19.6 ppm-v). During the test on MW-1 or MW-2, there were no detected vacuum pressures in any of the SVE points or the other monitoring wells.

Pilot Test #4

Based on a comparison of the results of Pilot Test #1 and Pilot Test #2, the de-watering of the shallow saturated zone above Unit B allowed access to more of the hydrocarbon-impacted soil and improved the radius of vacuum influence. To determine if a longer period of de-watering would further enhance the SVE operations, Pilot Test #4 was conducted over a two week period at SVE-1. To minimize the maintenance associated with the groundwater pumping, a bottom-inlet pneumatic pump (a QED AP2B AutoPump) replaced the Grundfos pump. In accordance with ORCAA requirements, SLR installed two, 55-gallon carbon-filled canisters in series to the effluent line from the blower to treat the extracted soil vapors.

Pilot Test #4 was conducted from December 22, 2011 through January 5, 2012, and the pump and blower operated continuously, except for the period from the afternoon of December 30th through the morning of January 3rd when the equipment was inadvertently shut off. During the approximate 10-day-long period of operation, a total of 2,511 gallons of shallow groundwater were pumped, and the average pumping rate was approximately 0.17 gpm. By the end of the test, the groundwater drawdowns in SVE-2, MW-4, and MW-5, which are at least partially screened within the shallow saturated zone above Unit B, were 2.03, 0.70, and 0.89 feet, respectively. The groundwater drawdown in MW-1, which is partially screened within the deeper saturated zone above Unit D, was 2.59 feet. Groundwater drawdown was not observed at deeper saturated zone wells MW-2 or MW-3. Except possibly at well MW-5, it is not known if the groundwater drawdowns in the monitoring points had stabilized by the end of the test.

During Pilot Test #4, a vacuum pressure of approximately 96 inches of water column was initially applied to SVE-1; however, the applied pressure decreased to 86 inches of water column by the end of the test. The airflow rate was initially 53 cfm and it increased during the test to 75 cfm. The decreased pressure and increased airflow rate demonstrate that de-watering of the shallow saturated zone improved the performance of the SVE system. The PID reading at the influent pipe to the blower was initially 922 ppm, and the readings steadily decreased to 250 ppm by the end of the test. However, the extracted soil vapor samples collected on

December 29th (TEST4-Sample1) and January 5th (Test4_Sample2) at the influent pipe contained lower GRO concentrations (6.9 and 41.4 ppm-v, respectively).

By the end of the test, vacuum pressures were detected at all of the monitoring points, except MW-3. The vacuum pressure at SVE-2 was 4.5 inches of water column and the vacuum pressures at MW-1, MW-2, MW-3, MW-4, and MW-5 ranged from 0.03 to 0.70 inches of water column.

On December 29, 2011, SLR pumped some of the water in the temporary storage tank through two, 55-gallon carbon-filled canisters in series for treatment, and collected a sample of the effluent from the second carbon canister. The sample was submitted to F&B for analysis of BTEX and GRO. The sample did not contain BTEX or GRO concentrations greater than the MRLs. After completing Pilot Test #4, OWSI personnel pumped the all of the water in the storage tank through the carbon-filled canisters for treatment. The treated water was pumped into a tanker truck and hauled to the OWSI wastewater treatment plant in Port Ludlow for disposal.

EVALUATION OF PILOT TEST DATA

The results of the pilot tests indicate the following:

- There is limited recharge to the shallow groundwater above Unit B beneath the eastcentral part of the property (the area of concern), and limited pumping can significantly reduce the shallow water levels
- The shallow groundwater above Unit B is hydraulically connected to the deeper water table above Unit D
- De-watering of the shallow groundwater zone above Unit B exposes more of the hydrocarbon-impacted soil to the applied vacuum, and extracts the groundwater that contains the greatest petroleum hydrocarbon concentrations at the property
- The radius of shallow groundwater pumping influence is greater than 70 feet
- Extracted soil vapor sample analytical results and corresponding PID readings from Pilot Tests #2 and #4 were inconsistent and may reflect difficulties collecting a representative sample during operation of the blower

In addition to the results listed above, two key metrics that were used to evaluate the potential effectiveness of SVE/de-watering are the effective radius of vacuum influence and the hydrocarbon mass recovery rates.

Effective Radius of Vacuum Influence

For SVE system design purposes, the effective radius of applied vacuum influence is the maximum distance from an extraction point where the airflow rate is sufficient to remove the volatile petroleum hydrocarbons from vadose zone soils within a reasonable time [Chevron Research and Technology Company (CRTC), 1991]. To determine this radius of vacuum influence, SLR normalized the final vacuum pressure readings from the monitoring points during Pilot Test #4 by dividing the measured pressures by the final applied pressure at SVE-1, and converting to percentage values. The normalized values and distances from SVE-1 were then plotted on semi-log graph paper, and a best fit line of the data was used to represent the spatially averaged radial distribution of vacuum induced by SVE-1. In accordance with CRTC (1991) protocols, we conservatively assumed that the radial distance that corresponds to 1% of the applied vacuum, as interpolated from the fitted vacuum distribution line, represents the effective radius of vacuum influence and the appropriate spacing of SVE points in a full-scale system. Based on the data from Pilot Test #4, the effective radius of vacuum influence at the site is estimated to be 46 feet. The effective radius of influence should increase with a longer operation period because more of the shallow saturated zone will be de-watered. A copy of the plot of the normalized vacuum data is attached.

Hydrocarbon Mass Recovery Rates

To calculate the total hydrocarbon mass removal rates during the extended pilot test, SLR used the airflow and soil vapor sample analytical data collected during Pilot Test #4, after one week of the test and at the termination of the test. The mass removal rates were calculated by using the following equation:

Removal rate (lbs/day) = airflow rate (cfm) x GRO concentration (ppm-v) x molecular weight of GRO (g/mole) x 1.58^{-7} x 24 hours/day (USEPA, 1989)

On December 29, 2011 (7 days after test activation), the hydrocarbon mass removal rate was 0.14 pounds per day. On January 5, 2012 (2 days after reactivating the blower and pump), the hydrocarbon mass removal rate increased to 1.2 pounds per day. These mass recovery rates may be biased low because there were significant discrepancies between the PID readings (392 and 250 ppm-v) and the analyzed GRO concentrations in extracted soil vapor samples (6.9 and 41.4 ppm-v). As stated previously, the differences between the PID readings and the GRO concentrations may be due to difficulties collecting representative samples. When using the PID readings instead of the analyzed concentrations, the hydrocarbon mass removal rates on December 29th and January 5th were 8.05 and 7.27 pounds per day, respectively. The mass removal rate calculation sheet is attached. The

hydrocarbon mass removal rates were limited by the presence of the shallow groundwater, and the rate should increase with continued removal of the groundwater.

CONCLUSIONS

In December 2011 and January 2012, four SVE pilot tests were conducted at the OWSI facility. The purposes of the tests were to: 1) assess the potential effectiveness of SVE, with and without groundwater de-watering, at remediating the remaining volatile petroleum hydrocarbon-impacted soil, 2) determine if groundwater extraction can effectively de-water the shallower portion of the saturated zone (above the Unit B silty sand), expose the capillary fringe above the deeper water table (above the Unit D silts), and allow SVE to remove additional petroleum hydrocarbon vapors from the shallower saturated zone and the capillary fringe, and 3) obtain the information necessary for potential future design and implementation of an SVE system, with or without groundwater recovery/treatment.

Based on a comparison of the applied vacuum pressures, airflow rates, PID readings, and vacuum pressure readings (in the monitoring points) during Pilot Test #1 and Pilot Test #4, de-watering of the shallow saturated zone improves SVE performance at the site. The dewatering reduces resistance to subsurface airflow and exposes more of the hydrocarbonimpacted soil to the applied vacuum. Furthermore, the extracted water sample results, as well as previous shallow groundwater sampling results (SLR, 2011), show that de-watering removes the groundwater that contains the greatest petroleum hydrocarbon concentrations at the site. Based on the interpreted geology and hydrogeology of the facility, it appears that there is a relatively limited volume of water in the shallow saturated zone above Unit B. Based on the groundwater drawdown in each of the shallow groundwater monitoring points during a limited period of pumping, it appears that recharge to the shallow saturated zone is limited, and we believe that this zone can be effectively de-watered within the area of concern. Since the shallow groundwater above Unit B and the deeper water table above Unit D are hydraulically connected, the pumping operations would also lower the deeper water table, exposing the capillary fringe. By combining de-watering with SVE, the higher permeability hydrocarbonimpacted soil at the site should be exposed to an applied vacuum. Due to preferential airflow through the higher permeability soils, some of the hydrocarbons within finer-grained soils may not be accessible to the vacuum.

Based on the limited area of hydrocarbon-impacted soil [approximately 3,140 square feet; maximum length of approximately 65 feet and maximum width of approximately 58 feet (see Figure 3)] and an estimated effective radius of vacuum influence of at least 46 feet, SLR believes that SVE, combined with shallow groundwater de-watering, would be an effective method to remediate the remaining source of the hydrocarbon-impacted groundwater. The petroleum hydrocarbons in the coarser-grained (higher permeability) soils should be

effectively remediated and any remaining hydrocarbons in finer-grained (lower permeability) soils would leach to the groundwater at very low rates due to limited water movement through those soils. The risks associated with any remaining impacted soil would be low because the contamination would occur at depths below 10 feet bgs beneath the garage building and below 15 feet bgs outside of the building footprint.

SLR appreciates the opportunity to provide our services. If you have any questions, please call me at (425) 402-8800.

Sincerely,

SLR International Corp

Michael D. Staton, L.G. Principal Geologist

- Attachments: Limitations References Tables 1 and 2 Figures 1 through 5 Field Notes Laboratory Reports Plot of Normalized Vacuum Data Hydrocarbon Mass Removal Calculations
- cc: Diana Smeland, Port Ludlow Associates Sue Schroader, Olympic Property Group Joe Rehberger, Cascadia Law Group Chip Goodhue, Aspect Consulting Group Tom Kilbane, Short Cressman & Burgess

LIMITATIONS

The services described in this report were performed consistent with generally accepted professional consulting principles and practices. No other warranty, express or implied, is made. These services were performed consistent with our agreement with our client. This report is solely for the use and information of our client unless otherwise noted. Any reliance on this report by a third party is at such party's sole risk.

Opinions and recommendations contained in this report apply to conditions existing when services were performed and are intended only for the client, purposes, locations, time frames, and project parameters indicated. We are not responsible for the impacts of any changes in environmental standards, practices, or regulations subsequent to performance of services. We do not warrant the accuracy of information supplied by others, nor the use of segregated portions of this report.

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TABLES

Table 1Extracted Soil Vapor Sample Analytical ResultsOlympic Water and Sewer PropertyPort Ludlow, Washington

Pilot Test Number/ Location	Sample Name	Sample Location	Sample Date	Active Test Duration Prior to Sample Collection (Hours)	Analytical Results (mg/m ³)				
					Benzene ^a	Toluene ^a	Ethylbenzene ^a	Total Xylenes ^a	Gasoline-Range Organics ^b
Test #1/SVE-1	Test1-Sample1	Stack	12/12/11	0.1	0.26	0.40	<0.1	< 0.3	30
	Test1-Sample2	Influent to Blower ^c	12/12/11	3.7	0.35	0.88	0.30	0.68	47
Test #2/SVE-1	Test2-Sample1	Stack	12/13/11	0.1	< 0.1	< 0.1	<0.1	< 0.3	<10
	Test2_Sample2	Influent to Blower ^c	12/13/11	8.3	14	32	5.0	10	1,900
Test #3/MW-1	Test3_Sample1	Stack	12/14/11	0.1	0.58	1.6	0.62	1.6	85
Test #4/SVE-1	TEST4-Sample1	Influent to Blower ^c	12/29/11	162.3	0.18	0.46	0.53	1.8	30
	Test4_Sample2	Influent to Blower ^c	01/05/12	45	2.5	5.1	1.6	4.6	180

Notes:

 $mg/m^3 = Milligrams$ per cubic meter.

^a = Benzene, toluene, ethylbenzene, and total xylenes by EPA Method 8021B.

^b = Gasoline-range organics by Ecology Method NWTPH-Gx.

^c = The sample was collected after shutting off the blower.

Table 2Extracted Water Sample Analytical ResultsOlympic Water and Sewer PropertyPort Ludlow, Washington

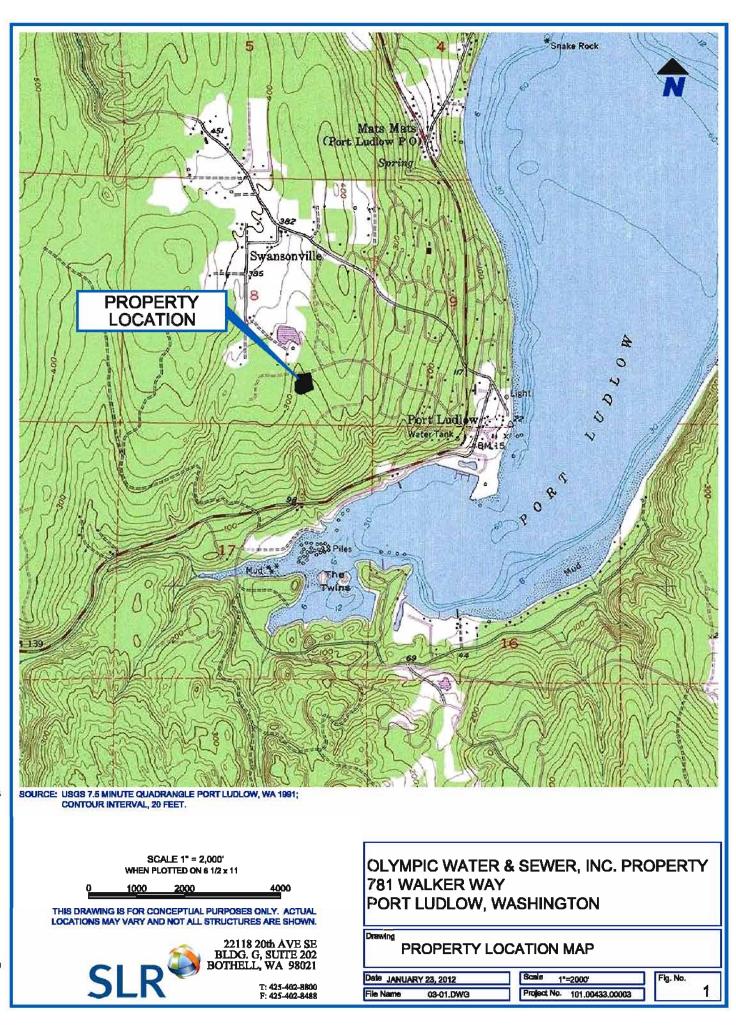
			Analytical Results (µg/L)					
Sample Name	Sample Location	Sample Date	Benzene ^a	Toluene ^a	Ethylbenzene ^a	Total Xylenes ^a	Gasoline-Range Organics ^b	
Effluent_Pre-Carbon	Influent to Storage Tank	12/13/11	430	1,900	<20	2,000	23,000	
System-Effluent_122911	Effluent from Second Carbon Canister	12/29/11	<1	<1	<1	<3	<100	
Notes:								

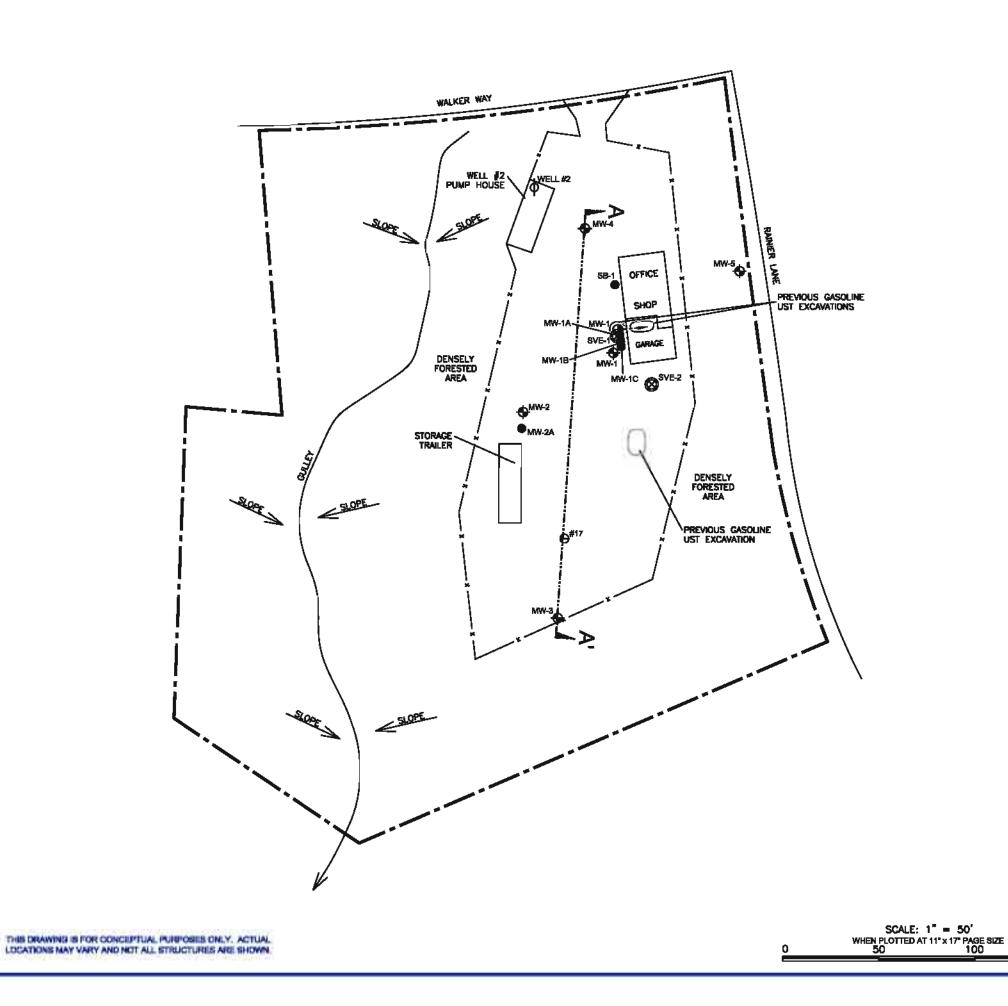
 $\mu g/L = Micrograms$ per liter.

^a = Benzene, toluene, ethylbenzene, and total xylenes by EPA Method 8021B.

^b = Gasoline-range organics by Ecology Method NWTPH-Gx.

FIGURES

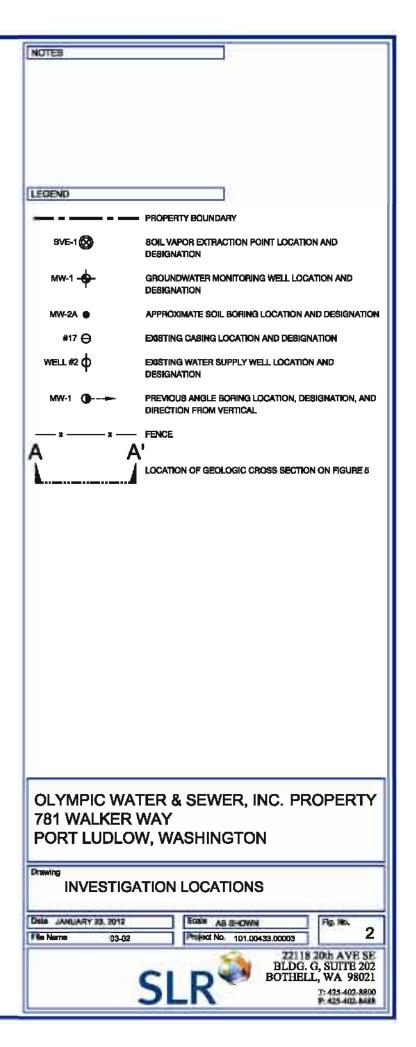


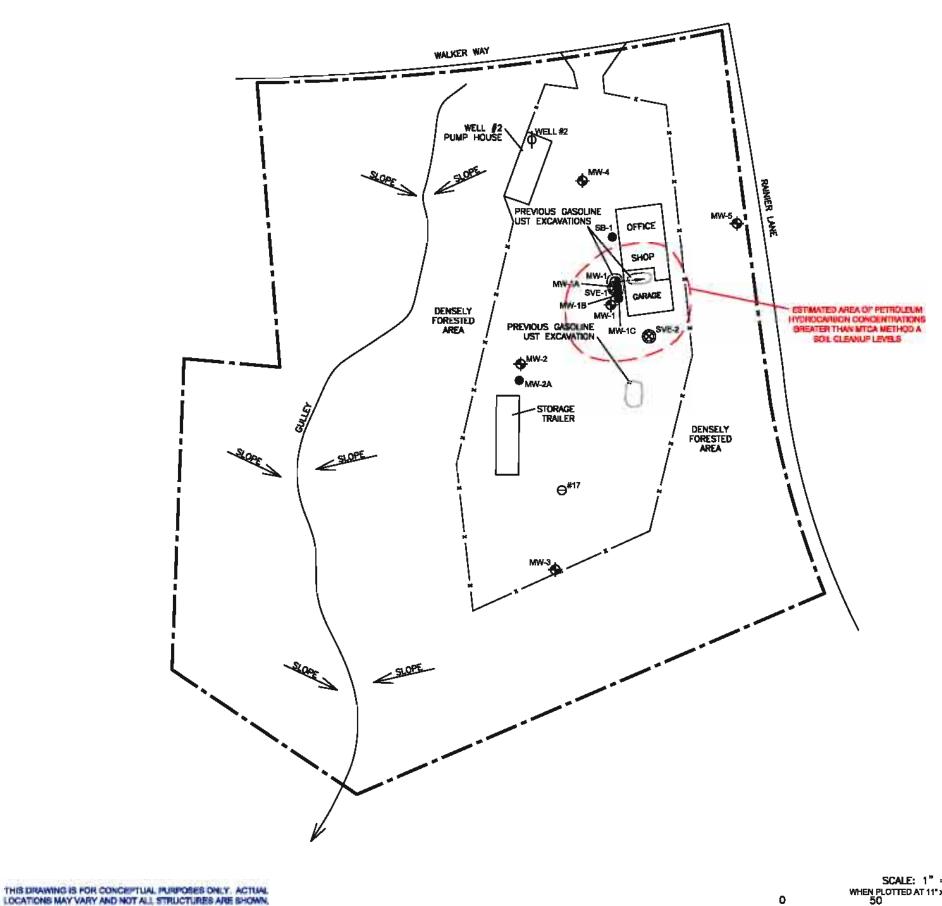


wei: Fridey, April 20, 2012 1:32-13 PM by egoodele Dreefing path: NoPorteedFigurer/Bother/Oympic Weak and SamerUV

R

150'



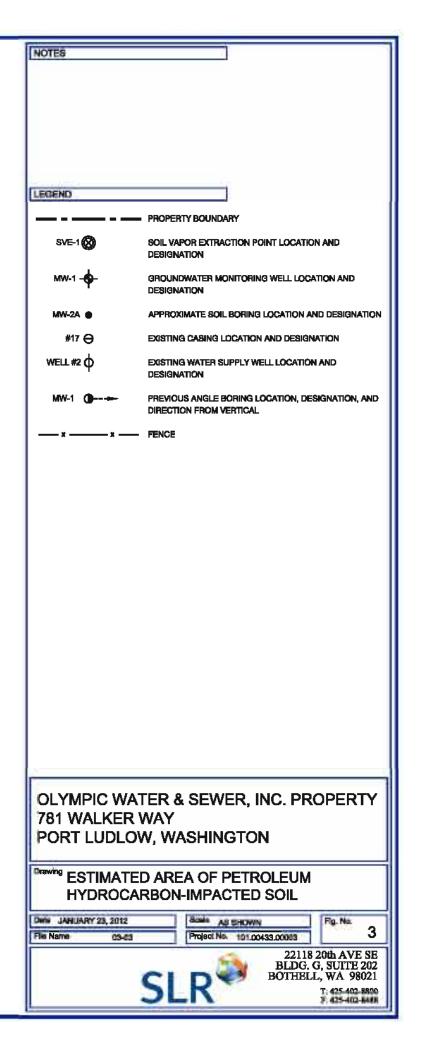


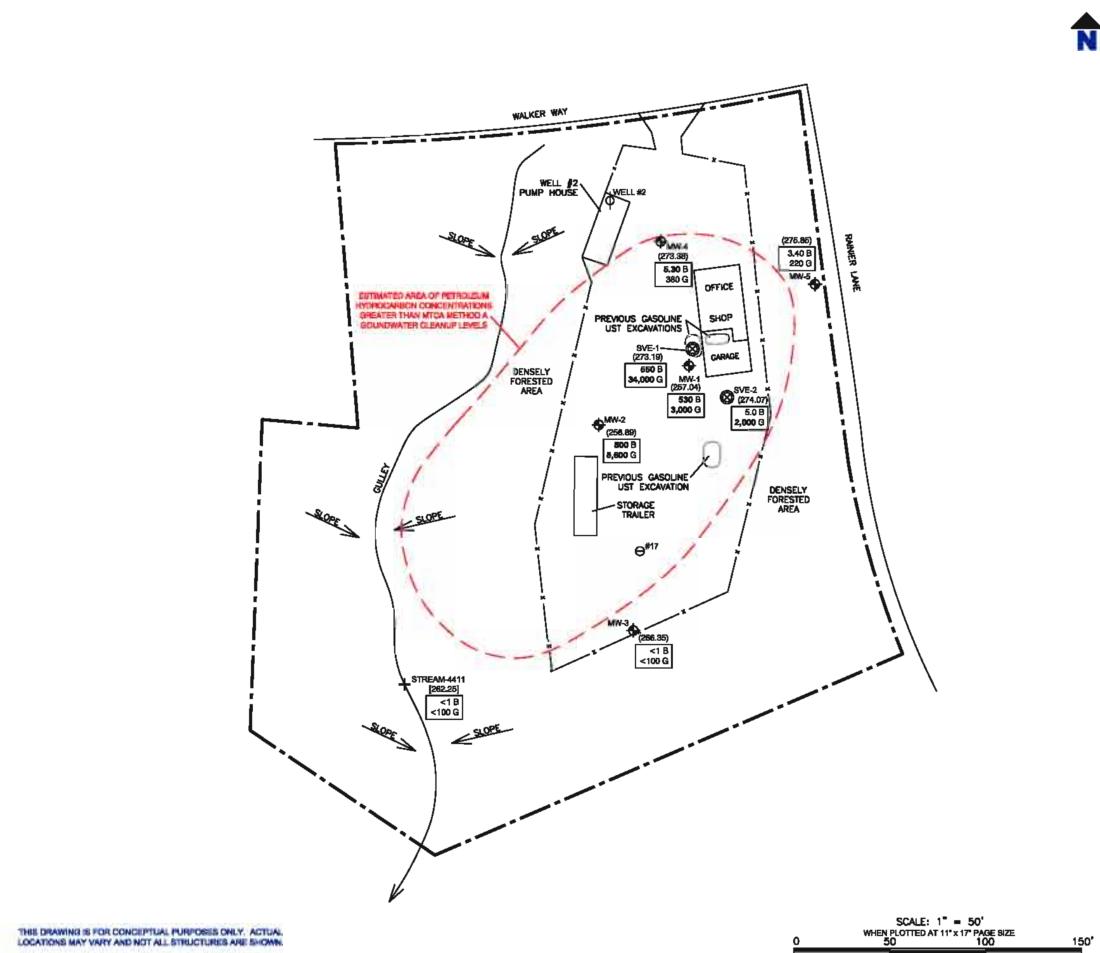
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N

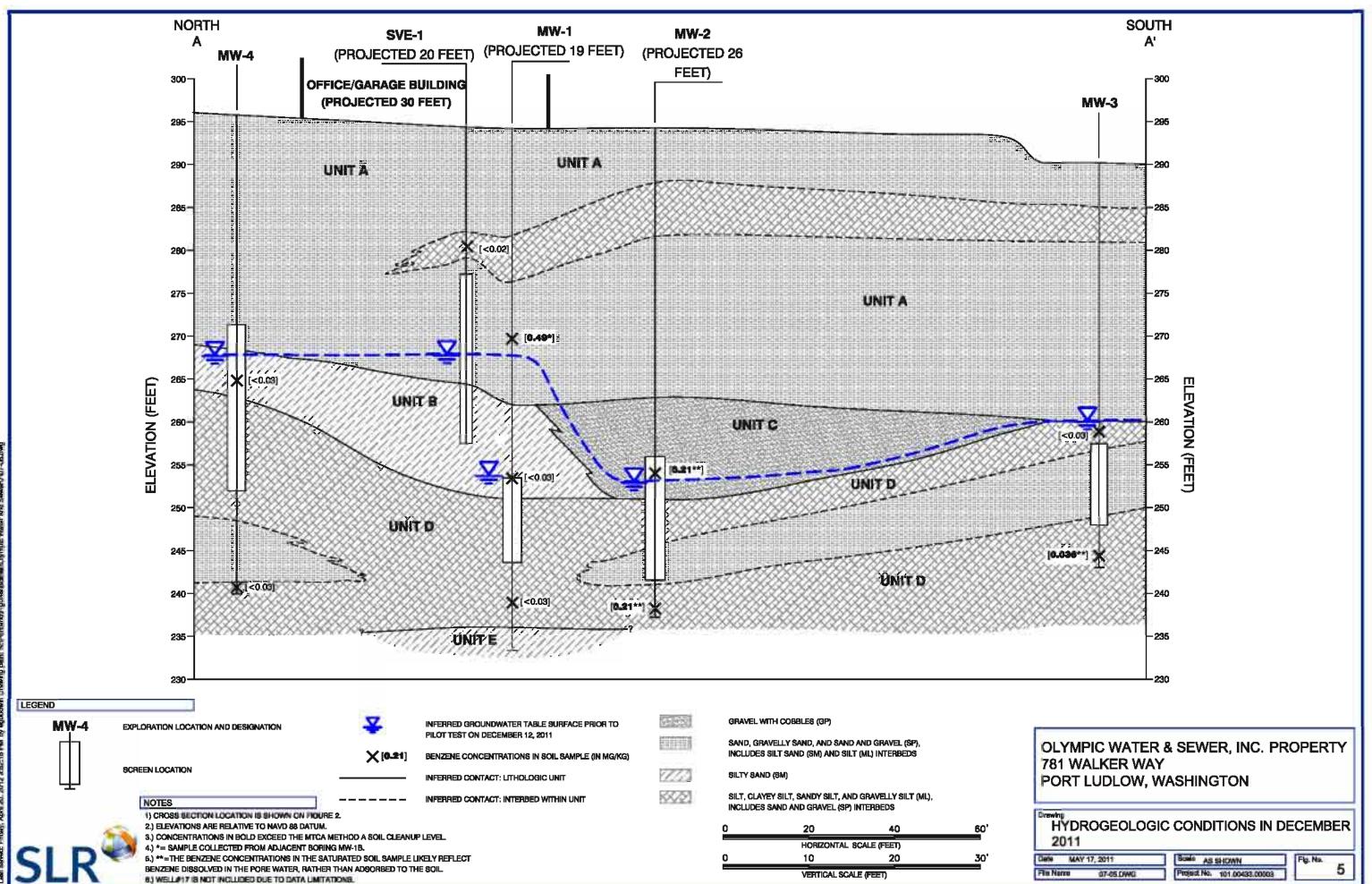
SCALE: 1" = 50' WHEN PLOTTED AT 11" x 17' PAGE SIZE 50 100

150'





NOTES					
1					
LEGEND					
Contraction of the local division of the loc					
	PROPERTY BOUNDARY				
MW-1 - Q -	GROUNDWATER MONITORING WELL LOCATION AND DESIGNATION				
STREAM-4411 +	APPROXIMATE STREAM SAMPLE LOCATION AND DESIGNATION				
SVE-1	SOIL VAPOR EXTRACTION POINT LOCATION AND DESIGNATION				
#17 O	EXISTING CASING LOCATION AND DESIGNATION				
WELL#2 O	EXISTING WATER SUPPLY WELL LOCATION AND DESIGNATION				
x x	FENCE				
(273.19)	SHALLOW GROUNDWATER ELEVATION (IN FEET ABOVE NAVD 86 DATUM) ON APRIL 8, 2011				
[262.25]	APPROXIMATE SURFACE WATER ELEVATION (IN FEET ABOVE NAVD 68 DATUM) ON APRIL 4, 2011				
5.30 B	B-BENZENE CONCENTRATION IN APRIL 2011				
380 G	G=GRO CONCENTRATION IN APRIL 2011 GROUNDWATER SAMPLE (IN µg/L)				
	NOTE: CONCENTRATIONS IN BOLD EXCEED THE MTCA METHOD A GROUNDWATER CLEANUP LEVELS.				
	TER & SEWER, INC. PROPERTY				
781 WALKER WAY PORT LUDLOW, WASHINGTON					
ESTIMATED AREA OF GASOLINE-IMPACTED					
SHALLOW GROUNDWATER					
Deta JANUARY 23, 2012	Scale As shown Ptr. No.				
File Name 03-04.DW					
22118 20th AVE SE BLDG. G, SUITE 202					
SLR BOTHELL, WA 98021 T: 425-402-8800 F: 425-402-8488					
	JLN F: 425-402-8408				



FIELD NOTES

Soil Vapor I	Extraction P	ilot Test - O	Soil Vapor Extraction Pilot Test - Olympic Water and Sewer, Inc. Facility, 781 Walker W	ar and Sewe	r, Inc. Facil	ity, 781 Wa	Iker Way, I	ay, Port Ludiow, WA	1, WA									
SLR interna	tional Corp		Date: 13/	113/11		Weather:	ش)رد در دج	•	Field Staff:			SVE Point:	5-3VE-2		Dewatering?:		(Yes/No	
22118 20th	22118 20th Ave. SE, G-202	202	Start Time:	0001		~4306	~43°E, dry		Chris Lee	<u>,</u> ,		SVE Equipment:		Rotran E	ENESASI	3	regen. b	6/oiver
Bothell, WA 98021	V 98021		End Time:	1830))			Cuclonii		<u>ب</u> و	icriccice +	Park		
			Air Samples					Water	Water Samples			10 A	Aimble, Nober	(u pos		
		Name		Ē	Time	1	Name	me		i=	Time							
「 「 に S に S に S	Г` d		8 -	0001		EFTÚ	UENT-	EFFLUENT - Ree - CARGON	APEN	1430		Dewaterin	Dewatering Equipment: Grand Fes		Į	icentre 2 M	Z	
152 2	ļ	- SUMAR		18:30								-		Baldo	Master Neter	VED Meter Blog	Tokakza	
Time	Applied Vacuum	Air Flow	Water Flow Rate	PID Reading		ş		Water Levels	sl					Vacuu	Vacuum Measurements (in. H ₂ O)	ments		
(24-hr)	(in. 世記)	Rate (fpm)		(mqq)	SVE-1	SVE-2	MW-1	MW-2	MW-3	MW-4	MW-5	SVE-1	SVE-2	MW-1	MW-2	MW-3	MW-4	MW-5
0640	õ,Q	NIA	-		8,86	J X Z					<u> </u>	SYSTEM	P V	2522	C C	11		1
0/1/0	0.00	NA	3.50	NIA	35.30 26.38		ちょう	39.78	E1.12	27.35	38.38	SyS'TE	5	TURNED	\sim			Ŷ
0945	いたん	St.t.	0.00	78.94	35.30		39.94	36-15	39.13		38:98	076	0.00	0.00	0.00	0.00	000	0.00
1005	5709	STOPPED TEST	12	C RE	Paires	ļ												ſ
1090	0.46	839	120	9. S a	変	36.35	39.94	39:78	99.13	32-25	36:38	94.0	0.30	0.00	0.00	0.00	0.00	0.00
1050	0,4%	801	033	ų,0*	33.50	3356 RM -												
1120	のたし	814	0.19	3.4-	35.07 37.13		40.18	S9:70	29.10	A 50	29.35	0.46	0.00	0.00	0.00	0.00	0,00	0.00
- 1 20	0.5%	83-1	0, [9]	29.1%	25.18	14.00	40.31		6060	21.53	29.38	040	1.00	0,00	000	Q Q Q	0°0	80
1330	のたわ	oret.	0.19	4.4.4	35.30	58.30		39.77	39.08	9754	41.400	0.76	0.90	0.00	0.00	0.00	Ċ.C	0.00
1400	oto	1+1	0.(3	6.4.0	8.8	26,26	tein	NM -			Î	0.46	0.9U 0.9U	0,00	0.00	0.00	Q Q Q	000
ju30	0.46	104	0,13	9.04		15:56	40.37	39-78	30.06	37.59	39.23	0,40			0.00	0.00	0,00	0,00
1500	0.4%	801	0.3	11.6 -	36.38	37.5	40.3S	- WN	1 1			97.0				0.00		000
1530	97.0	土土北	0,0		35,00	37.58	40.3%	124.105	そのゆで	37.63	- tht be	でたわ	0.60	0.00	0.00	0,00	0.00	0.00
1600	970	765	Q. 13		35.30	わた	そかった	39.79	39. 16	5-65 (2 20	39.51	97.0	0.50	0000	0.00	0.00	00'0	0,00
1630	いたん	761	0, 13	SILA	35.00	53×82	15.01	39.70	39.19	37.80	39.58	いたし	0,30	0.00	0.00	0,00	0.00	000
1700	97.0	ナナナ	0.13	34	83.30	19791	40,53	39.79	ちとした	37.85	12461	0.40	0,40	0.10	0,00	0,00	0.00	0.00
1730	0.70	764	0.13	3.90	35.30	74.94	40.55	31.40	39,38	58°±€	39.61	97.0	0.30		0.00	0.00	9.8 0	0.00
1800	97.0		0. (5	404	35.20	29497	110.5C	of the	29.31	58:20	99,63	97.0	Ro	0.10	0.00	0,00	0,00	0°0
1830	046	768	0, Ŭ	ų, i a	3530	3% C	50. TK	DF. 65	29,09		20162	つた	0.50	0.10	0,00	CrOC	800	0°Q
·																		
Notes: 7	chull zer	reading	Tokultzer reading @ beginning of Jest:	The frink		32.5	gallens		nd Red	Could Real gallons			019	P10 Reeding	Ş	el e Mel	time ef	
1	irned G	1 collow	Turned Grundfes pump en C	୧୦୦୫୦	12	Top of the	Primp C 35,23	35,22)			Colle	Collection -	Г 1	Ċ,	Same 3)
1000	(00)	C				1	j_Cr24	Some	أمصراب	Same back into 4 505-3	1-2012-1	2	からく	208	Sas worm.			`
	20-		iour 1300 tomps supplied pomping, which is	N Non	nping,	8773 8773	n	Conned.	". cleen	Remised Lowed & Chistolled Dung	anshal.	led our	ç					
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÷7;	-2.6 -2.6	CW57	staff 1	turned off power to volum off, restart	d d o	wer to de restarted	devaterna		5	ccciclentally. red vorum,		. Vacum succed		verter le	back in b to 886 f	into Sue-1 886 prim 4	the form	
	5	how when	¥ .	CLUMM IC	1	r	51.5											
\sim	Soil vapor		and			collected	1		ጜ				(
in Sout	1 vorpor	· Samples	anl	PID reactinus		collected	at na	manitule d	atter f	turning >	sve ble	blever off,	¥					

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Soil Vapor Extraction Pilot Test - Olympic Water and Sewer, Inc. Facility, 781 Walker Way, Port Ludlow, WA	traction P	ilot Test - O	lympic Wate	r and Sewe	r, Inc. Facil	ity, 781 Wa	lker Way, F	ort Ludiow	, WA				I					
SLR International Corp	onal Corp		Date: /J/	11/hl		Weather:			Field Staff:			SVE Point:	-MW	~	Dewatering?:		YesyNo	
22118 20th Ave. SE, G-202	we. SE, G-2	202	Start Time: End Time:	(550) 1020		Overcit,	é si	- 40°F	Chris Lee	Ş		SVE Equipi	4		ENSD3MSL, regentative	125, regé	meathe	blower
	77007	Air Samples	nples	044				Water :	Water Samples			Cucleric	e Mus		ちんしらたいす		(10659) YANA	<u> </u>
	Na	Name		Time	ne		Name			Ч	Time	19ª Ac	3" Acmule Noter		extraction huse	sonh m	3	
TEST 3.	SA	SAMPLE 2		1050				$\left \right\rangle$			$\left \right $	Dewaterin	g Equipmer	it: Crund	Dewatering Equipment: Cound Bs Redictio 2- w/ Baldor	1,610 2	~/ Bal	lor
												l's of	Mast Meth	Po Po	VIE HL	エーション	L. W. Leven N.	meter
And a second	-											(C-)	12101 NO12		<u>101 C.</u>	۳h		2
Time	Applied Vacyum	Air Flow	Water Flow Rate	PID Reading				Water Levels (ft. BTOC)	S					Vacuu	vacuum Measurements ⁻ (in. H ₂ O)	ements		
(24-hr)	(in. Hg)	Rate (fpm)	(mdg)	(mdd)	SVE-1	SVE-2	MW-1	MW-2	MW-3	MW-4	8-WM	SVE-1	SVE-2	MW-1	MW-2	MW-3	MW-4	MW-5
1040	0.00	0,00	n b	0,00	2,50	35.84 39.98		3640	39.18	3754	30.05	SX57B	SYSTEM TURINUU		11 11 11			A
1050	97,0	804	0.00	1.6	36.50	25.54	36.St	35-155	37.18	27.54	20-02	0.0	0.00	いたし	0.00	000	(2) (2)	0.00
0.01	いたし	870	0.00	1.9	6hig	25.80	48、87	ROVE	29.18	3753-	88	0,00	000	0.4%	0.0	0.00	0.0	0,00
120	96.0	810	0.00	1.8	ふりう	25.78	06-Sh		29.13		39.03	0.00	0.00	0.0%	0,00	0,00	000	0.00
0000	97.0	808	0.00	ľŮ	2643	25.1%			29.19	White C	30/00	0.00	0.00	のそし	0,00	0,00	$\alpha \omega$	000
1250 9	97,0	TOF	0,d0	1,4			48,87	39.92-29.18		5h.tl	39,00	000	000	0:510	0.00	0.0	0,0	0.00
1390 6	0.7.0	ZUZ	8 0	(((36.39	35.76		39.92	39.18	Str EC	39.00	0.00	0,00	0:46	0,00	0.00	200	0.00
1350 6	97,0	16t	0.0		No.37	25-35	48.87	39,91	29.18		3698	0,00	0.00	のよう	Q'O	0.00	00	0.00
065	0.46	809	0.00			JE: FO-	48-38				38.90	0.00	0.00	0.49	0.0	0.00	0.0	0.00
1450 0	0.26	-78g	0,0	1.9	_					ったせん	76.36	0,00	0-00	98°0		0,00	0.00	0.00
1590 6	97,0	Q6t	0.00	2.1				39.88		とかとも	£6.3C	0.0	0.00	0.36	(C) (C)	0. <i>ČŮ</i>	0.00	0-00
000	0.66	PCA	0.00	0°C	26.34	35.68		39,88	Filo	七ち、七乙	76.36	0.00	0.00	98.0	0.00	0.00	000	0.00
RENED	S S	MEN	1-1-0	R MO	Mover	2	かいず											A
1630	95.0	Th8t	0.00	1.8	26.33	3569	U/8.80	06-5-1	39.06	ST. 43	26.97	0.00	0.0	95.0	000	80	0.0	000
TURNE	Sy Sy	NETTER	日 日 0	N N) VED	B.4eic		2-~1W	ل ل									
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	-																	
																		-
Notes: Tole	Totalizer redu		(begann		of test		96.6 gallins											
-1 -1 -00	ct Ct	-	بغ،98 ⁴)											
-				-						-		-	-			,		
	-) 	Jompletzi	Kunp completer downtard well,					5		Vounn F	pulled.	putted all water in pump tube back	overentia	the purch	.tvbe?		inte NW-1.	1-1. S
ایتر 		it con) juny ~	23 PV	had vounn		Plow sharped	· sharpon ·	いたい		5			1				201
Se. I voper	- •	soundles	and p	PID readings	lings	wore all		collectel	a T S	noten	system extensit stack.	stack	•					
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Soil Vapor Extraction Pilot Test - Olympic Water and Sewer, Inc. Facility, 781 Walker Way, SLR international Corp Date: $\frac{\partial \mathcal{J}}{\partial t}$	n Pilot Test - rp	Olympic Wate Date: $ \mathcal{P} $	er and Sewei -//t//11	r, Inc. Facili	<mark>ity, 781 Wa</mark> l Weather:	lker Way, P		, WA Field Staff:	,		SVE Point	SVE Point: MW~ 2		Dewatering?:	an a	(Yes/No	
22118 20th Ave. SE, G-202 Bothell, WA 98021	G-202	Start Time: End Time:	17400		Orareart,	des,	~ 40°F	Chn's	Lee		SVE Equi.	• •	h	EN5D3MSL	MSL re	N533MSL regon. blower	Jac
	Air S	Air Samples					Water	Water Samples			ار (د (د (•			-	,	
	Name		Time	ne		Name	ne		F	Time	ج ح	nubler f	Texple	extracture	hm 1	husc	
				\setminus							Dewater	ing Equipm.	ant: Gww	Dewatering Equipment: Countles Red Po D Submersble	e Pe P	Submerabl	0
												Mesher Meter	BLOS	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	n nur - nur bionte	to the man	L
Applied	d Air Flow	Water Flow Rate	PID Reading			>	Water Levels (ft. BTOC)	S					Vacu	1 5-	ements		
(24-hr) (in Hg)				SVE-1	SVE-2	MW-1	MW-2	MW-3	MW-4	MW-5	SVE-1	SVE-2	MW-1		MW-3	MW-4	MW-5
1610 0.00	0.00 11 0.00	ດ ທີ່ມ	.6	26.53	26.53 75.76 47.07 26.53 75.75 47.00		39.33	39.16 39.16	247.42C	28.97	00'0	00.0	0000	0.00 0.7=0	000	20°0	000 000
5	NOL-SX	NH S	2	DENOW	106												
「ゆせつ 1000 「ほう 000	194	2 e P Y	+ 1J	10.02 D5 60 U. C.	20 00 A 00 20		1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1	20,00	94.47	41.00 00.14		0000	200,0			000 000 000	3.0
	1	60		De.33 25-68 46.00	H 68				£5.38 Qh-EC	10.30		0,00	, 0 S	920	0000	6 00 00	0.00
Operated s	system @	C-MW	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	sample	Port	by bi	emoved	2	alleviate		Vacoum 1	lack.					
															,		
Notes: Totalizer reading & beginning at test: 97.7 gallons. Totalizer reading C evel of test: 120.7 gallons	realing (ی ہے۔ ای ان	to Ja	ist te	ist test : 97.7 gallens test: 120.7 gallans	7.7 gall	llens										
1740: Rump. developed grand. Tauit & stopped pumping. PID shot up to drawn beek into Mix-2. by vocum.	p. devel	apried gi	mur- i	9- 5- 4 G	ndets	red pr	mpine.	DID	s hot i	б Б	+00%	900+ ppm as		water	fari		
Soil vopor samples and PID readings	amples	and PIC) revolin	ع م ا	2	l coller	sted c	tr Sys	stem	were all collected at system exhaust	L Stack.	Ϋ́κ.					

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Soil Vandr	Evtrartion I	Pilot Test . (Soil Vanor Extraction Bilot Test - Olymnic Water and Sewer. Inc. Facility. 281 Walker Way. Port Ludlow. WA	er and Sewe	r. Inc. Facil	itv. 781 Wa	Iter Way P	ort Ludlow	L WA							Раде	/ of /	
SLR Interna	SLR International Corp		Date: /3	11/00/01		Weather:	Weather: Overcost,	Ŷ,	Field Staff:			SVE Point:	SUB.	L.	Dewaterin	g?: <	1.8	
22118 20th Ave. SE Botheil, WA '98021	22118 20th Ave. SE, G-202 Bothell, WA '98021	-202	Start Time: End Time:	1345		ry50f	0 (4)	dr.	Ohri	lec		SVE Equip	ment: Rof	She BN	SVE Equipment: Rotom BNSD3MSi regon. O'Science morsh- knockcort Parts	1 Tegin	k blow	- ml
		Air St	Air Samples					Water	Water Samples			1 2" ~	bber	Rexuble	- CXT	rection	husc	
	Ž	Name		Time	ne		Nai	Name ·		Tir	Time	Totalizer R	eading At S	tart: $/\mathcal{I}$	0 O	At End: _/		
												Dewaterin	g Equipmer	Dewatering Equipment:	APA (tome	AP De Terrechalmen Dono	022
														13105		waler	melar	
E	Applied	Air Elow	Water	PID				Water Levels	ls ·					Vacuu	as	ments		
(24-hr)	(inHg)	<u> </u>			SVE-1	SVE-2	1-WM	MW-2	MW-3	MW-4	MW-5	SVE-1	SVE-2	T-WW	MW-2	MW-3	MW-4	MW-5
1330	00	00	1.5	0.0	<u>N</u> N	96.13		40.35	11700	1.	99.33	S 8	0.0	00	00	00	0.0	00
1350	<i>60</i> ,02			6766			30,08	40.3S		かたと	99.J 3	80.0	0.0	0,Û	0.0	00	0.0	0.0
1410	96.0			4537	Ŵ	<i>86.1</i> 3	20.98	40.35	Í	フカートー	39.33	%. N	00	00	0,0	0.0	0.0	0.0
1430	0%	96.0 2.468	0 Ŭ	26 S	MN	36,35 40.03	5003	(10.30	27.08	27:45 29:23	29.23	96.0	0.0	0.0	0,0	0.0	0.0	0.0
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Notes: 73	Notes: Total zer redug		before starking pump: 117.0	Parking &	: Jum	1.7.0												
12	To halven r	reedury	schr s.	the for the second	nondi. Pr			FJ 61	starting blacer, after		burk C	ompletel	7 dewe	itered u	Completely dewatered well : . 120.5	120.5		
6	Tokuluzur r	redy	the hour	1 1044	Site:	124.0)						
1300 (5	contect s	celiment	1300 Barled sectiment Run Sire-I prove to inscribing pump.	7-37	pror 1	to micrit	m Pure	i.				١	-	с. ,				<u> </u>
1330 5	far-teck 1	hand z	allerer	ta N	1 yus	frequenci	3 05-0	miching	Scies	Ceesed	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	ç Ç	ien sk	scheel	うでう			
10	Top of pump	dund	15 38" above bottom	a abo	K C	2 Hon	of well	Υl,										
	-				· · ·			[;				•						
	reading	S Let S S P F S	b) PID readings uses all collected at b) PID readings at Exhaust stock		actives	0	t atter training SI	ि 5 (1 2 2 2 2	SVO 60000	consistenty		0,0 00m				~		
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/ of /	(Yeš/No	regoneration - Party	1050	1798	APQ remedation	Ĺ		MW-4	0'0							/			ţ	n c'annst			
Page		Ste 15	fan 1	At End: /	west -	valur meler	nents	MW-3	00											carter			
	Dewatering?:	Rotran ENESJ3MS2 1	extrae	γ	604	rah	Vacuum Measurements (in. H ₂ O)	MW-2	0.0										1	Ahraugh carlen canister (3).			A.
	<u> </u>	in the	cable	rt: , 深ゴ,	QED	8005	Vacuum	T-WM	00											to gravity flow			planer
	NE - 1	F. Rota	er A	ling At Stai	quipment:			SVE-2	0.											d Garit	, >		
	E Point: S	SVE Equipment: Rotron w/ cyclone morst	2" robber	talizer Read	Dewatering Equipment: QED			SVE-1	90.0									-	IJ	ater h	From VO.A->		tuning
ť	SV	SV &	Ce	Tot	De		(the		27.03											and men.	defacult is eliminate from Vod		ather
,				Time	/0.30		9		-		 	_		 		_			2	air sampic tran	elimina		ntold
	aff:	s lee			2			3 MW-4	37,00 38.03				-					 -		50MF	5.2 +		at manifald
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LABORATORY REPORTS

ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

December 20, 2011

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the results from the testing of material submitted on December 13, 2011 from the Olympic Water & Sewer, Inc, 101.00433.00003, F&BI 112183 project. There are 4 pages included in this report. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures mcp/KJ SLR1220R.DOC

ENVIRONMENTAL CHEMISTS

CASE NARRATIVE

This case narrative encompasses samples received on December 13, 2011 by Friedman & Bruya, Inc. from the SLR International Corp. Olympic Water & Sewer, Inc, 101.00433.00003 project. Samples were logged in under the laboratory ID's listed below.

<u>Laboratory ID</u>	SLR International Corp.
112183 -01	Test1-Sample1
112183 -02	Test1-Sample2
112183 -03	Test2-Sample1

All quality control requirements were acceptable.

ENVIRONMENTAL CHEMISTS

Date of Report: 12/20/11 Date Received: 12/13/11 Project: Olympic Water & Sewer, Inc, 101.00433.00003, F&BI 112183 Date Extracted: 12/14/11 Date Analyzed: 12/14/11

RESULTS FROM THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Results Reported as mg/m³

<u>Sample ID</u> Laboratory ID	<u>Benzene</u>	<u>Toluene</u>	Ethyl <u>Benzene</u>	Total <u>Xylenes</u>	Gasoline <u>Range</u>	Surrogate (<u>% Recovery)</u> (Limit 50-150)
Test1-Sample1 112183-01	0.26	0.40	<0.1	<0.3	30	97
Test1-Sample2 112183-02	0.35	0.88	0.30	0.68	47	93
Test2-Sample1 112183-03	<0.1	<0.1	<0.1	<0.3	<10	98
Method Blank 01-2210 MB	<0.1	<0.1	<0.1	<0.3	<10	96

ENVIRONMENTAL CHEMISTS

Date of Report: 12/20/11 Date Received: 12/13/11 Project: Olympic Water & Sewer, Inc, 101.00433.00003, F&BI 112183

QUALITY ASSURANCE RESULTS FOR THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES, AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Laboratory Code: 112183-03 (Duplicate)

Eaboracory couci	LIBIOO OO (D'uphie	100)		
	Reporting	Sample	Duplicate	Relative Percent Difference
Analyte	Ūnits	Result	Result	(Limit 20)
Benzene	mg/m³	<0.1	<0.1	nm
Toluene	mg/m³	<0.1	< 0.1	nm
Ethylbenzene	mg/m³	< 0.1	< 0.1	nm
Xylenes	mg/m³	< 0.3	< 0.3	nm
Gasoline	mg/m³	<10	<10	nm

Laboratory Code: Laboratory Control Sample

			Percent	
	Reporting	Spike	Recovery	Acceptance
Analyte	Units	Level	LCS	Criteria
Benzene	mg/m³	5.0	89	70-130
Toluene	mg/m ³	5.0	93	70-130
Ethylbenzene	mg/m ³	5.0	91	70-130
Xylenes	mg/m ³	15	90	70-130
Gasoline	mg/m ³	100	124	70-130

ENVIRONMENTAL CHEMISTS

Data Qualifiers & Definitions

a - The analyte was detected at a level less than five times the reporting limit. The RPD results may not provide reliable information on the variability of the analysis.

A1 – More than one compound of similar molecule structure was identified with equal probability.

b - The analyte was spiked at a level that was less than five times that present in the sample. Matrix spike recoveries may not be meaningful.

ca - The calibration results for this range fell outside of acceptance criteria. The value reported is an estimate.

c - The presence of the analyte indicated may be due to carryover from previous sample injections.

d - The sample was diluted. Detection limits may be raised due to dilution.

ds - The sample was diluted. Detection limits are raised due to dilution and surrogate recoveries may not be meaningful.

dv - Insufficient sample was available to achieve normal reporting limits and limits are raised accordingly.

fb - Analyte present in the blank and the sample.

fc – The compound is a common laboratory and field contaminant.

hr - The sample and duplicate were reextracted and reanalyzed. RPD results were still outside of control limits. The variability is attributed to sample inhomogeneity.

ht - Analysis performed outside the method or client-specified holding time requirement.

ip - Recovery fell outside of normal control limits. Compounds in the sample matrix interfered with the quantitation of the analyte.

j – The result is below normal reporting limits. The value reported is an estimate.

J - The internal standard associated with the analyte is out of control limits. The reported concentration is an estimate.

jl - The analyte result in the laboratory control sample is out of control limits. The reported concentration should be considered an estimate.

jr - The rpd result in laboratory control sample associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

js - The surrogate associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

lc - The presence of the compound indicated is likely due to laboratory contamination.

L - The reported concentration was generated from a library search.

nm - The analyte was not detected in one or more of the duplicate analyses. Therefore, calculation of the RPD is not applicable.

 $\rm pc-The\ sample\ was\ received\ in\ a\ container\ not\ approved\ by\ the\ method.\ The\ value\ reported\ should\ be\ considered\ an\ estimate.$

 $\ensuremath{\text{pr}}$ – The sample was received with incorrect preservation. The value reported should be considered an estimate.

ve - Estimated concentration calculated for an analyte response above the valid instrument calibration range. A dilution is required to obtain an accurate quantification of the analyte.

vo - The value reported fell outside the control limits established for this analyte.

x - The sample chromatographic pattern does not resemble the fuel standard used for quantitation.

12183 SAMPLE CHAIN OF CUSTODY HE 12-13-11	Dire Station SAMPLERS (signature) International Corp PROJECT NAMENO. International Corp Olympic Water & Sewer, Inc. 20714 Aver SE G-300-	City, State, ZIP <u>SortHEL</u> , <u>WA</u> 9800-1 Phone # (435) 403-8800 Fax # (435) 403-8485	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$	Bruya, Inc. SIGNATURE PRINT NAME COMPANY DATE TIME Bruue West Relinquished by: CHAIS, LEE SLR PAM 13.05 8282 Relinquished by: Eart Mun TAB Samples receivel at 16.06
112183	Send Report To MIKE Company SLR INTE Address 221(8 207	City, State, ZIP <u>SOTHSL</u> , WA Phone # <u>(435)</u> 403-8800 Fax # <u>6</u>	Sample ID TEST 2 - SAMPLE 2 TEST 2 - SAMPLE 2 TEST 2 - SAMPLE 2	Friedman & Bruya, Inc. 3012 16th Avenue West R Seattle, WA 98119-2029 Ph. (206) 285-8282 Fax (206) 283-5044

ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

December 20, 2011

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the results from the testing of material submitted on December 15, 2011 from the Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112222 project. There are 4 pages included in this report. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures SLR1220R.DOC

ENVIRONMENTAL CHEMISTS

CASE NARRATIVE

This case narrative encompasses samples received on December 15, 2011 by Friedman & Bruya, Inc. from the SLR International Corp. Olympic Water & Sewer, Inc, PO 101.00433.00003 project. Samples were logged in under the laboratory ID's listed below.

Laboratory ID	SLR International Corp.
112222 -01	Test2_Sample2
112222 -02	Test3_Sample1

All quality control requirements were acceptable.

ENVIRONMENTAL CHEMISTS

Date of Report: 12/20/11 Date Received: 12/15/11 Project: Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112222 Date Extracted: 12/16/11 Date Analyzed: 12/16/11

RESULTS FROM THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Results Reported as mg/m³

<u>Sample ID</u> Laboratory ID	<u>Benzene</u>	<u>Toluene</u>	Ethyl <u>Benzene</u>	Total <u>Xylenes</u>	Gasoline <u>Range</u>	Surrogate (<u>% Recovery)</u> (Limit 50-150)
Test2_Sample2 112222-01 1/5	14	32	5.0	10	1,900	96
Test3_Sample1 112222-02	0.58	1.6	0.62	1.6	85	95
Method Blank 01-2227 MB	<0.1	<0.1	<0.1	<0.3	<10	94

ENVIRONMENTAL CHEMISTS

Date of Report: 12/20/11 Date Received: 12/15/11 Project: Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112222

QUALITY ASSURANCE RESULTS FOR THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES, AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Laboratory Code: 112230-01 (Duplicate)

Analyte	Reporting Units	Sample Result	Duplicate Result	Relative Percent Difference (Limit 20)
Benzene	mg/m ³	<0.1	< 0.1	nm —
Toluene	mg/m ³	< 0.1	< 0.1	nm
Ethylbenzene	mg/m ³	< 0.1	< 0.1	nm
Xylenes	mg/m ³	< 0.3	< 0.3	nm
Gasoline	mg/m ³	<10	<10	nm

Laboratory Code: Laboratory Control Sample

			Percent	
	Reporting	Spike	Recovery	Acceptance
Analyte	Units	Level	LCS	Criteria
Benzene	mg/m³	5.0	87	70-130
Toluene	mg/m³	5.0	91	70-130
Ethylbenzene	mg/m ³	5.0	92	70-130
Xylenes	mg/m ³	15	91	70-130
Gasoline	mg/m ³	100	124	70-130

ENVIRONMENTAL CHEMISTS

Data Qualifiers & Definitions

a - The analyte was detected at a level less than five times the reporting limit. The RPD results may not provide reliable information on the variability of the analysis.

A1 – More than one compound of similar molecule structure was identified with equal probability.

b - The analyte was spiked at a level that was less than five times that present in the sample. Matrix spike recoveries may not be meaningful.

ca - The calibration results for this range fell outside of acceptance criteria. The value reported is an estimate.

c - The presence of the analyte indicated may be due to carryover from previous sample injections.

d - The sample was diluted. Detection limits may be raised due to dilution.

ds - The sample was diluted. Detection limits are raised due to dilution and surrogate recoveries may not be meaningful.

dv - Insufficient sample was available to achieve normal reporting limits and limits are raised accordingly.

fb - Analyte present in the blank and the sample.

fc – The compound is a common laboratory and field contaminant.

hr - The sample and duplicate were reextracted and reanalyzed. RPD results were still outside of control limits. The variability is attributed to sample inhomogeneity.

ht - Analysis performed outside the method or client-specified holding time requirement.

ip - Recovery fell outside of normal control limits. Compounds in the sample matrix interfered with the quantitation of the analyte.

j – The result is below normal reporting limits. The value reported is an estimate.

J - The internal standard associated with the analyte is out of control limits. The reported concentration is an estimate.

jl - The analyte result in the laboratory control sample is out of control limits. The reported concentration should be considered an estimate.

jr - The rpd result in laboratory control sample associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

js - The surrogate associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

lc - The presence of the compound indicated is likely due to laboratory contamination.

L - The reported concentration was generated from a library search.

nm - The analyte was not detected in one or more of the duplicate analyses. Therefore, calculation of the RPD is not applicable.

 $\rm pc$ – The sample was received in a container not approved by the method. The value reported should be considered an estimate.

 $\rm pr$ – The sample was received with incorrect preservation. The value reported should be considered an estimate.

ve - Estimated concentration calculated for an analyte response above the valid instrument calibration range. A dilution is required to obtain an accurate quantification of the analyte.

vo - The value reported fell outside the control limits established for this analyte.

x - The sample chromatographic pattern does not resemble the fuel standard used for quantitation.

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ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

January 6, 2012

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the results from the testing of material submitted on December 30, 2011 from the Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112402 project. There are 4 pages included in this report. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures SLR0106R.DOC

ENVIRONMENTAL CHEMISTS

CASE NARRATIVE

This case narrative encompasses samples received on December 30, 2011 by Friedman & Bruya, Inc. from the SLR International Corp. Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112402 project. Samples were logged in under the laboratory ID's listed below.

Laboratory ID	SLR International Corp.
112402-01	TEST4-Sample1

Please note that the relative percent difference (RPD) of the analysis of ethylbenzene and the xylenes in the laboratory control sample and laboratory control duplicate fell outside of the established control limits. The results have been flagged accordingly.

All other quality control requirements were acceptable.

ENVIRONMENTAL CHEMISTS

Date of Report: 01/06/12 Date Received: 12/30/11 Project: Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112402 Date Extracted: 12/30/11 Date Analyzed: 12/30/11

RESULTS FROM THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Results Reported as mg/m³

<u>Sample ID</u> Laboratory ID	<u>Benzene</u>	<u>Toluene</u>	Ethyl <u>Benzene</u>	Total <u>Xylenes</u>	Gasoline <u>Range</u>	Surrogate (<u>% Recovery)</u> (Limit 50-150)
TEST4-Sample1 112402-01	0.18	0.46	0.53	1.8	30	83
Method Blank 01-2313 MB	< 0.1	< 0.1	<0.1	<0.3	<10	90

ENVIRONMENTAL CHEMISTS

Date of Report: 01/06/12 Date Received: 12/30/11 Project: Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112402

QUALITY ASSURANCE RESULTS FOR THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES, AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Laboratory Code: 112402-01 (Duplicate)

Euseracery could	r -	,		Relative Percent
	Reporting	Sample	Duplicate	Difference
Analyte	Units	Result	Result	(Limit 20)
Benzene	mg/m³	0.18 a	0.34 a	60 a
Toluene	mg/m³	0.46 a	0.83	57 a
Ethylbenzene	mg/m ³	0.53	0.97	58 vo
Xylenes	mg/m ³	1.8	3.2	57 vo
Gasoline	mg/m ³	30 a	64	72 a

Laboratory Code: Laboratory Control Sample

			Percent	
	Reporting	Spike	Recover	Acceptance
Analyte	Units	Level	y LCS	Criteria
Benzene	mg/m ³	5.0	80	70-130
Toluene	mg/m³	5.0	80	70-130
Ethylbenzene	mg/m ³	5.0	85	70-130
Xylenes	mg/m ³	15	81	70-130
Gasoline	mg/m ³	100	116	70-130

ENVIRONMENTAL CHEMISTS

Data Qualifiers & Definitions

a - The analyte was detected at a level less than five times the reporting limit. The RPD results may not provide reliable information on the variability of the analysis.

A1 – More than one compound of similar molecule structure was identified with equal probability.

b - The analyte was spiked at a level that was less than five times that present in the sample. Matrix spike recoveries may not be meaningful.

ca - The calibration results for this range fell outside of acceptance criteria. The value reported is an estimate.

c - The presence of the analyte indicated may be due to carryover from previous sample injections.

d - The sample was diluted. Detection limits may be raised due to dilution.

ds - The sample was diluted. Detection limits are raised due to dilution and surrogate recoveries may not be meaningful.

dv - Insufficient sample was available to achieve normal reporting limits and limits are raised accordingly.

fb - Analyte present in the blank and the sample.

fc – The compound is a common laboratory and field contaminant.

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ip - Recovery fell outside of normal control limits. Compounds in the sample matrix interfered with the quantitation of the analyte.

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jl - The analyte result in the laboratory control sample is out of control limits. The reported concentration should be considered an estimate.

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lc - The presence of the compound indicated is likely due to laboratory contamination.

L - The reported concentration was generated from a library search.

nm - The analyte was not detected in one or more of the duplicate analyses. Therefore, calculation of the RPD is not applicable.

pc – The sample was received in a container not approved by the method. The value reported should be considered an estimate.

pr – The sample was received with incorrect preservation. The value reported should be considered an estimate.

ve - Estimated concentration calculated for an analyte response above the valid instrument calibration range. A dilution is required to obtain an accurate quantification of the analyte.

vo - The value reported fell outside the control limits established for this analyte.

x - The sample chromatographic pattern does not resemble the fuel standard used for quantitation.

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ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

January 10, 2012

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the results from the testing of material submitted on January 6, 2012 from the Olympic Water & Sewer 101.00433.00003, F&BI 201053 project. There are 4 pages included in this report. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures mcp/KJ SLR0110R.DOC

ENVIRONMENTAL CHEMISTS

CASE NARRATIVE

This case narrative encompasses samples received on January 6, 2012 by Friedman & Bruya, Inc. from the SLR International Corp. Olympic Water & Sewer 101.00433.00003, F&BI 201053 project. Samples were logged in under the laboratory ID's listed below.

Laboratory ID	<u>S</u>
201053-01	T

<u>SLR International Corp.</u> Test 4_Sample 2

All quality control requirements were acceptable.

ENVIRONMENTAL CHEMISTS

Date of Report: 01/10/12 Date Received: 01/06/12 Project: Olympic Water & Sewer 101.00433.00003, F&BI 201053 Date Extracted: 01/06/12 Date Analyzed: 01/06/12

RESULTS FROM THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Results Reported as mg/m³

<u>Sample ID</u> Laboratory ID	<u>Benzene</u>	<u>Toluene</u>	Ethyl <u>Benzene</u>	Total <u>Xylenes</u>	Gasoline <u>Range</u>	Surrogate (<u>% Recovery)</u> (Limit 50-150)
Test 4_Sample 2 201053-01	2.5	5.1	1.6	4.6	180	87
Method Blank 02-0038 MB	<0.1	<0.1	<0.1	<0.3	<10	88

ENVIRONMENTAL CHEMISTS

Date of Report: 01/10/12 Date Received: 01/06/12 Project: Olympic Water & Sewer 101.00433.00003, F&BI 201053

QUALITY ASSURANCE RESULTS FOR THE ANALYSIS OF VAPOR SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES, AND TPH AS GASOLINE USING MODIFIED EPA METHOD 8021B AND NWTPH-Gx

Laboratory Code: 201013-02 (Duplicate)

Analyte	Reporting Units	Sample Result	Duplicate Result	Relative Percent Difference (Limit 20)
Benzene	mg/m³	< 0.1	< 0.1	nm
Toluene	mg/m ³	< 0.1	< 0.1	nm
Ethylbenzene	mg/m ³	< 0.1	< 0.1	nm
Xylenes	mg/m ³	< 0.3	< 0.3	nm
Gasoline	mg/m ³	<10	<10	nm

Laboratory Code: Laboratory Control Sample

			Percent	
	Reporting	Spike	Recovery	Acceptance
Analyte	Units	Level	LCS	Criteria
Benzene	mg/m³	5.0	95	70-130
Toluene	mg/m ³	5.0	96	70-130
Ethylbenzene	mg/m ³	5.0	98	70-130
Xylenes	mg/m ³	15	97	70-130
Gasoline	mg/m ³	100	123	70-130

ENVIRONMENTAL CHEMISTS

Data Qualifiers & Definitions

a - The analyte was detected at a level less than five times the reporting limit. The RPD results may not provide reliable information on the variability of the analysis.

A1 – More than one compound of similar molecule structure was identified with equal probability.

b - The analyte was spiked at a level that was less than five times that present in the sample. Matrix spike recoveries may not be meaningful.

ca - The calibration results for this range fell outside of acceptance criteria. The value reported is an estimate.

c - The presence of the analyte indicated may be due to carryover from previous sample injections.

d - The sample was diluted. Detection limits may be raised due to dilution.

ds - The sample was diluted. Detection limits are raised due to dilution and surrogate recoveries may not be meaningful.

dv - Insufficient sample was available to achieve normal reporting limits and limits are raised accordingly.

fb - Analyte present in the blank and the sample.

fc – The compound is a common laboratory and field contaminant.

hr - The sample and duplicate were reextracted and reanalyzed. RPD results were still outside of control limits. The variability is attributed to sample inhomogeneity.

ht - Analysis performed outside the method or client-specified holding time requirement.

ip - Recovery fell outside of normal control limits. Compounds in the sample matrix interfered with the quantitation of the analyte.

j – The result is below normal reporting limits. The value reported is an estimate.

J - The internal standard associated with the analyte is out of control limits. The reported concentration is an estimate.

jl - The analyte result in the laboratory control sample is out of control limits. The reported concentration should be considered an estimate.

jr - The rpd result in laboratory control sample associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

js - The surrogate associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

lc - The presence of the compound indicated is likely due to laboratory contamination.

L - The reported concentration was generated from a library search.

nm - The analyte was not detected in one or more of the duplicate analyses. Therefore, calculation of the RPD is not applicable.

pc – The sample was received in a container not approved by the method. The value reported should be considered an estimate.

 $\ensuremath{\text{pr}}$ – The sample was received with incorrect preservation. The value reported should be considered an estimate.

ve - Estimated concentration calculated for an analyte response above the valid instrument calibration range. A dilution is required to obtain an accurate quantification of the analyte.

vo - The value reported fell outside the control limits established for this analyte.

x - The sample chromatographic pattern does not resemble the fuel standard used for quantitation.

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ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

January 3, 2012

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the amended results from the testing of material submitted on December 15, 2011 from the Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112223 project. As requested, the results from the analysis of the sample Effluent_Pre-Carbon for benzene, toluene, ethylbenzene, and the xylenes (BTEX) have been reported.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures mcp/KJ SLR1220R.DOC

ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

December 20, 2011

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the results from the testing of material submitted on December 15, 2011 from the Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112223 project. There are 4 pages included in this report. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures mcp/KJ SLR1220R.DOC

ENVIRONMENTAL CHEMISTS

CASE NARRATIVE

This case narrative encompasses samples received on December 15, 2011 by Friedman & Bruya, Inc. from the SLR International Corp. Olympic Water & Sewer, Inc, PO 101.00433.00003 project. Samples were logged in under the laboratory ID's listed below.

<u>Laboratory ID</u>	SLR International Corp.
112223 -01	Effluent_Pre-Carbon

All quality control requirements were acceptable.

ENVIRONMENTAL CHEMISTS

Date of Report: 12/20/11 Date Received: 12/15/11 Project: Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112223 Date Extracted: 12/16/11 Date Analyzed: 12/16/11

RESULTS FROM THE ANALYSIS OF WATER SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES AND TPH AS GASOLINE USING EPA METHOD 8021B AND NWTPH-Gx

Results Reported as ug/L (ppb)

<u>Sample ID</u> Laboratory ID	<u>Benzene</u>	<u>Toluene</u>	Ethyl <u>Benzene</u>	Total <u>Xylenes</u>	Gasoline <u>Range</u>	Surrogate (<u>% Recovery</u>) (Limit 52-124)
Effluent_Pre-Carbon 112223-01 1/20	430	1,900	<20	2,000	23,000	101
Method Blank 01-2224 MB	<1	<1	<1	<3	<100	94

ENVIRONMENTAL CHEMISTS

Date of Report: 12/20/11 Date Received: 12/15/11 Project: Olympic Water & Sewer, Inc, PO 101.00433.00003, F&BI 112223

QUALITY ASSURANCE RESULTS FOR THE ANALYSIS OF WATER SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES, AND TPH AS GASOLINE USING EPA METHOD 8021B AND NWTPH-Gx

Laboratory Code: 112205-01 (Duplicate)

	Reporting	Śample	Duplicate	Relative Percent Difference
Analyte	Units	Result	Result	(Limit 20)
Benzene	ug/L (ppb)	<1	<1	nm
Toluene	ug/L (ppb)	<1	<1	nm
Ethylbenzene	ug/L (ppb)	<1	<1	nm
Xylenes	ug/L (ppb)	<3	<3	nm
Gasoline	ug/L (ppb)	<100	<100	nm

Laboratory Code: Laboratory Control Sample

			Percent	
	Reporting	Spike	Recovery	Acceptance
Analyte	Units	Level	LCS	Criteria
Benzene	ug/L (ppb)	50	88	72-119
Toluene	ug/L (ppb)	50	89	71-113
Ethylbenzene	ug/L (ppb)	50	89	72-114
Xylenes	ug/L (ppb)	150	84	72-113
Gasoline	ug/L (ppb)	1,000	99	70-119

ENVIRONMENTAL CHEMISTS

Data Qualifiers & Definitions

a - The analyte was detected at a level less than five times the reporting limit. The RPD results may not provide reliable information on the variability of the analysis.

A1 – More than one compound of similar molecule structure was identified with equal probability.

b - The analyte was spiked at a level that was less than five times that present in the sample. Matrix spike recoveries may not be meaningful.

ca - The calibration results for this range fell outside of acceptance criteria. The value reported is an estimate.

c - The presence of the analyte indicated may be due to carryover from previous sample injections.

d - The sample was diluted. Detection limits may be raised due to dilution.

ds - The sample was diluted. Detection limits are raised due to dilution and surrogate recoveries may not be meaningful.

dv - Insufficient sample was available to achieve normal reporting limits and limits are raised accordingly.

fb - Analyte present in the blank and the sample.

fc – The compound is a common laboratory and field contaminant.

hr - The sample and duplicate were reextracted and reanalyzed. RPD results were still outside of control limits. The variability is attributed to sample inhomogeneity.

ht - Analysis performed outside the method or client-specified holding time requirement.

ip - Recovery fell outside of normal control limits. Compounds in the sample matrix interfered with the quantitation of the analyte.

j – The result is below normal reporting limits. The value reported is an estimate.

J - The internal standard associated with the analyte is out of control limits. The reported concentration is an estimate.

jl - The analyte result in the laboratory control sample is out of control limits. The reported concentration should be considered an estimate.

jr - The rpd result in laboratory control sample associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

js - The surrogate associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

lc - The presence of the compound indicated is likely due to laboratory contamination.

L - The reported concentration was generated from a library search.

nm - The analyte was not detected in one or more of the duplicate analyses. Therefore, calculation of the RPD is not applicable.

 $\rm pc$ – The sample was received in a container not approved by the method. The value reported should be considered an estimate.

 $\ensuremath{\text{pr}}$ – The sample was received with incorrect preservation. The value reported should be considered an estimate.

ve - Estimated concentration calculated for an analyte response above the valid instrument calibration range. A dilution is required to obtain an accurate quantification of the analyte.

vo - The value reported fell outside the control limits established for this analyte.

x - The sample chromatographic pattern does not resemble the fuel standard used for quantitation.

112223				SAMPLE CHAIN OF CUSTODY	CHAIN O	F C	Ŭ.SU	lao		2y	Intel 2	<u>S</u>			-		5
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ENVIRONMENTAL CHEMISTS

James E. Bruya, Ph.D. Yelena Aravkina, M.S. Bradley T. Benson, B.S. Kurt Johnson, B.S. 3012 16th Avenue West Seattle, WA 98119-2029 TEL: (206) 285-8282 e-mail: fbi@isomedia.com

January 6, 2012

Mike Staton, Project Manager SLR International Corp. 22118 20th Ave. SE., G-202 Bothell, WA 98021

Dear Mr. Staton:

Included are the results from the testing of material submitted on December 30, 2011 from the Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112401 project. There are 4 pages included in this report. Any samples that may remain are currently scheduled for disposal in 30 days. If you would like us to return your samples or arrange for long term storage at our offices, please contact us as soon as possible.

We appreciate this opportunity to be of service to you and hope you will call if you should have any questions.

Sincerely,

FRIEDMAN & BRUYA, INC.

Kurt Johnson Chemist

Enclosures mcp/KJ SLR0106R.DOC

ENVIRONMENTAL CHEMISTS

CASE NARRATIVE

This case narrative encompasses samples received on December 30, 2011 by Friedman & Bruya, Inc. from the SLR International Corp. Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112401 project. Samples were logged in under the laboratory ID's listed below.

<u>Laboratory ID</u>	SLR International Corp.
112401-01	System-Effluent-122911

All quality control requirements were acceptable.

ENVIRONMENTAL CHEMISTS

Date of Report: 01/06/12 Date Received: 12/30/11 Project: Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112401 Date Extracted: 01/03/12 Date Analyzed: 01/03/12

RESULTS FROM THE ANALYSIS OF WATER SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES AND TPH AS GASOLINE USING EPA METHOD 8021B AND NWTPH-Gx

Results Reported as ug/L (ppb)

<u>Sample ID</u> Laboratory ID	<u>Benzene</u>	Toluene	Ethyl <u>Benzene</u>	Total <u>Xylenes</u>	Gasoline <u>Range</u>	Surrogate (<u>% Recovery</u>) (Limit 52-124)
System-Effluent-122 112401-01	911 <1	<1	<1	<3	<100	89
Method Blank 02-0013 MB	<1	<1	<1	<3	<100	93

ENVIRONMENTAL CHEMISTS

Date of Report: 01/06/12 Date Received: 12/30/11 Project: Olympic Water & Sewer, Inc. 101.00433.00003, F&BI 112401

QUALITY ASSURANCE RESULTS FOR THE ANALYSIS OF WATER SAMPLES FOR BENZENE, TOLUENE, ETHYLBENZENE, XYLENES, AND TPH AS GASOLINE USING EPA METHOD 8021B AND NWTPH-Gx

Laboratory Code: 112337-01 (Duplicate)

Analyte	Reporting Units	Sample Result	Duplicate Result	Relative Percent Difference (Limit 20)
Benzene	ug/L (ppb)	<1	<1	nm
Toluene	ug/L (ppb)	<1	<1	nm
Ethylbenzene	ug/L (ppb)	<1	<1	nm
Xylenes	ug/L (ppb)	<3	<3	nm
Gasoline	ug/L (ppb)	<100	<100	nm

Laboratory Code: Laboratory Control Sample

			Percent	
	Reporting	Spike	Recovery	Acceptance
Analyte	Units	Level	LCS	Criteria
Benzene	ug/L (ppb)	50	94	65-118
Toluene	ug/L (ppb)	50	95	72-122
Ethylbenzene	ug/L (ppb)	50	100	73-126
Xylenes	ug/L (ppb)	150	93	74-118
Gasoline	ug/L (ppb)	1,000	99	69-134

ENVIRONMENTAL CHEMISTS

Data Qualifiers & Definitions

a - The analyte was detected at a level less than five times the reporting limit. The RPD results may not provide reliable information on the variability of the analysis.

A1 – More than one compound of similar molecule structure was identified with equal probability.

b - The analyte was spiked at a level that was less than five times that present in the sample. Matrix spike recoveries may not be meaningful.

ca - The calibration results for this range fell outside of acceptance criteria. The value reported is an estimate.

c - The presence of the analyte indicated may be due to carryover from previous sample injections.

d - The sample was diluted. Detection limits may be raised due to dilution.

ds - The sample was diluted. Detection limits are raised due to dilution and surrogate recoveries may not be meaningful.

dv - Insufficient sample was available to achieve normal reporting limits and limits are raised accordingly.

fb - Analyte present in the blank and the sample.

fc - The compound is a common laboratory and field contaminant.

hr - The sample and duplicate were reextracted and reanalyzed. RPD results were still outside of control limits. The variability is attributed to sample inhomogeneity.

ht - Analysis performed outside the method or client-specified holding time requirement.

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j – The result is below normal reporting limits. The value reported is an estimate.

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jl - The analyte result in the laboratory control sample is out of control limits. The reported concentration should be considered an estimate.

jr - The rpd result in laboratory control sample associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

js - The surrogate associated with the analyte is out of control limits. The reported concentration should be considered an estimate.

lc - The presence of the compound indicated is likely due to laboratory contamination.

L - The reported concentration was generated from a library search.

nm - The analyte was not detected in one or more of the duplicate analyses. Therefore, calculation of the RPD is not applicable.

pc – The sample was received in a container not approved by the method. The value reported should be considered an estimate.

 $\ensuremath{\text{pr}}$ – The sample was received with incorrect preservation. The value reported should be considered an estimate.

ve - Estimated concentration calculated for an analyte response above the valid instrument calibration range. A dilution is required to obtain an accurate quantification of the analyte.

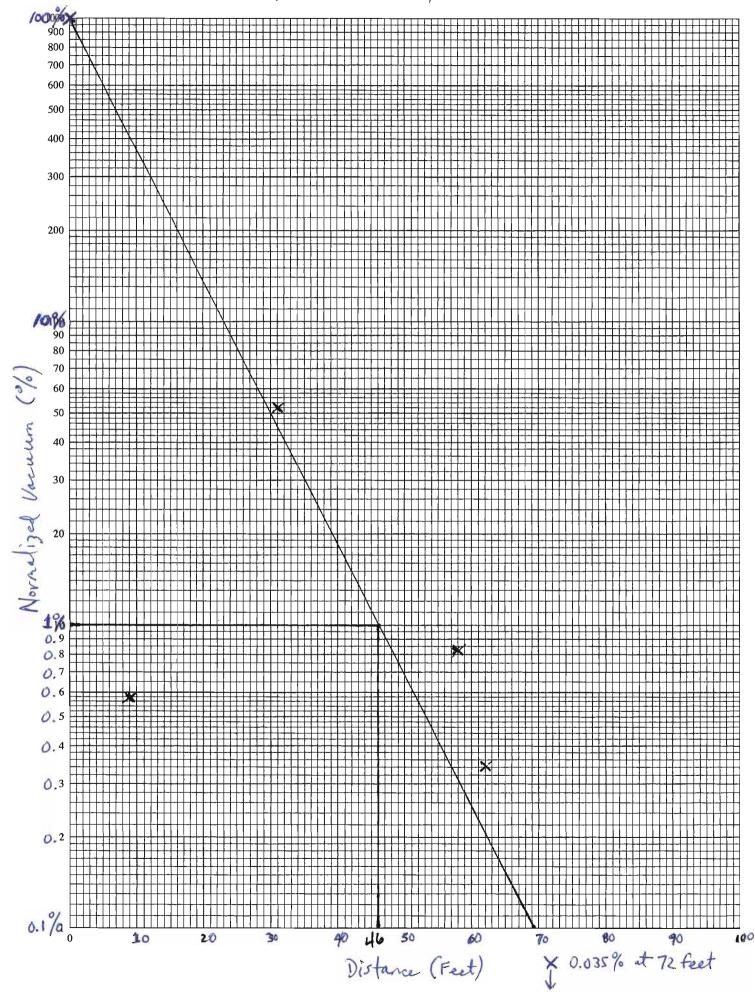
vo - The value reported fell outside the control limits established for this analyte.

x - The sample chromatographic pattern does not resemble the fuel standard used for quantitation.

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Friedman & Bruya, Inc. 3012 16th Avenue West	Relinquished by:	SIGN	SIGNATURE			PRINT NAME	AME		i	COM	COMPANY		DATE DATE	TIME
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· Ph. (206) 285-8282	Relinquished by:	-) 		2	5		Ş			•	
Fax (206) 283-5044	Received by:	by:												*
FORMS/COC/COC/DOC														

PLOT OF NORMALIZED VACUUM DATA

NORMALIZED VACUUM PLOT PILOT TEST #4



HYDROCARBON MASS REMOVAL CALCULATIONS



