Limited Site Assessment

Prepared For:

Dan Aalfs 305 East Pioneer Avenue, "Old Homer Tesoro" Homer, Alaska ADEC File #2314.26.031

Prepared By:





Preparation Date:

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ACRONYMS AND ABBREVIATIONS

AAC Alaska Administrative Code

ADEC Alaska Department of Environmental Conservation

BTEX Benzene, Toluene, Ethyl benzene, Xylenes

bgs Below Ground Surface COC Chain of Custody DRO Diesel Range Organics

EPA Environmental Protection Agency

GRO Gasoline Range Organics
PID Photoionization Detector

ppm Parts Per Million

LOQ Limit of Quantification (formerly PQL: Practical Quantitative Limit)

MDL Method Detection Limit

QA/QC Quality Assurance/Quality Control

RRO Residual Range Organics

sf Square Feet SM Silty Sand

SP Poorly Graded Sand SW Well Graded Sand

SGS Societe Generale de Surveillance
USCS Unified Soil Classification System
VOC Volatile Organic Compound

EXECUTIVE SUMMARY

Restoration Science & Engineering, LLC (RSE) is providing this Underground Storage Tank (UST) closure site assessment report to describe site conditions encountered during site characterization activities in accordance with our workplan, approved by the ADEC's Paul Horwath on September 13, 2011. RSE implemented this workplan and, in conjunction with an excavation contractor (Property Improvements), excavated six test pits on the subject property to assess subsurface hydrocarbon impacts. Field activities took place from September 20-21, 2011.

1.0 SITE DESCRIPTION AND HISTORY

The subject property is located at 305 East Pioneer Avenue, Lot 3A1 of Nils O. Svedlund Subdivision No. 8 Plat 90-20, located near the intersection of East Pioneer and Svedlund Street in downtown Homer, Alaska (Attachment A, Figure 1). The site operated as a retail fuel and service station from at least as early as 1973 until 1991. The subject property is irregular in shape with the approximate dimensions of 200 ft by 80-100 ft in width. Homer is located in the Southern Kenai Peninsula on the north side of Kachemak Bay. This property is situated on the south side of East Pioneer Avenue. The property is primarily comprised of a flat gravel fill pad sloping gently to the south before reaching a steep vegetated embankment on the southern edge of the property. The property is bounded by developed commercial properties to the east and west, an embankment descending to an alley to the south, and Pioneer Avenue to the north.

Based on our review of ADEC records, one 3,000-gallon and one 4,000-gallon gasoline USTs, one 8,000-gallon diesel UST, as well as the associated piping and dispensers were removed from the site in 1991 without proper UST closure documentation. In addition to retail fuel sales the site was used for automotive maintenance activities. Maintenance activities occurred in two structures, known as the "Old Garage" and the "New Garage" and are located on the property. Two other structures, known as the "Old Wood Frame Shop" and the "Old Gas Station Office", as well as a gas station canopy have been removed from the site. The Old Wood Frame Shop is also a suspected location of historic automotive maintenance activities.

2.0 AREA HYDROGEOLOGY AND CLIMATE

Homer is located in the Southern Kenai Peninsula on the north side of Kachemak Bay. The Homer area lies on the southern end of the Kenai Peninsula between Cook Inlet and the Kenai Mountains. Significant geologic features in Homer include the escarpment and adjacent upland (Diamond Ridge and Lookout Mountain), the bench area at the foot of the escarpment (Homer Bench area) and the Homer Spit extending into Kachemak Bay. The city of Homer and the subject property lay on an irregular terrace and alluvial fan complex between the bluff and Kachemak Bay referred to as the "Homer Bench".

Lot 3A1 is located on the upper portion of the lower homer bench, and had a moderately steep grade to the south, prior to being filled. The topography of the Homer bench is a result of Pleistocene glaciations and resultant surficial deposits of glacial drift or pro-glacial lake-bottom and deltaic sediments overlain by fan alluvium and colluvium. The depth to groundwater at the site is not known, but is greater than 13 feet below grade based recent test pit observations.

3.0 SITE CHARACTERIZATION ACTIVITIES

RSE supervised the excavation of six test pits throughout the subject property. Test pits were excavated to between 8 ft below ground surface (bgs) and 13 ft bgs. The locations of test pits were chosen using a combination of historic aerial photographs, on-site interviews, and a workplan prepared by International Consulting and Engineering, dated April 2003, that included a diagram indicating the probable locations of USTs and dispenser islands. A scaled Site Diagram indicating the locations of site features and test pits is provided as Figure 2. Field screening results are provided in Table 1. Soil encountered has been described using Unified Soil Classification System (USCS) conventions and symbols.

Test pit TP1 was excavated to a depth of 8 feet bgs in the probable location of the former dispenser islands. A wooden box approximately 8ft x 3ft x 4ft was encountered approximately 3 ft bgs (Photograph 4 in Appendix A). This box may have acted as the foundation for the dispenser island (previously removed). The top of the box was open, and the bottom of this crib was located approximately 7 ft bgs. Material above 7 ft bgs, including within the box, consisted of well graded sand (SW) with silt (SM). The excavator bucket was used to break through the bottom of the crib. A relatively dense layer of silt with poorly graded fine sand (ML-SP) was encountered below the crib at a depth of approximately 7 ft bgs. Impacted soil was encountered heterogeneously throughout the excavation. Two analytical samples were collected from this excavation, one at the depth of 3ft bgs (TP1-3), which exhibited the greatest field screening result, and the other from the ML-SP layer beneath the wooden crib at a depth of approximately 8 ft bgs (TP1-8).

Test pit TP2 was excavated in the reported vicinity of the former tanks. The initial test pit was excavated to a depth of 7.5 ft bgs. No staining or olfactory evidence of hydrocarbons was encountered until 7.5 ft bgs, where impacted soil was encountered. A trench was advanced from TP2 to the south and impacted soil was again encountered at 6.5 ft bgs. A flexible pipe coupled to a copper pipe was encountered within the zone of impacted soil at approximately 7 ft bgs. This line was tentatively identified as the water service to the building. A portion of this line was removed, and a remaining portion was crimped and left in place. The location was marked above grade using an "I" beam and timber. TP2 was completed to a depth of 11 ft bgs. Material above 10 ft bgs appeared to consist of relatively unconsolidated fill materials consisting of a heterogeneous mixture of well graded sand with silt (SW-SM). Occasional large cobbles and debris was encountered. Apparent native material was encountered at 10 ft bgs and consisted of ML-SP. Two analytical samples were collected from this excavation, one at the depth which returned the greatest field screening result at a depth of 6 ft bgs (TP2-6), and the other from native material at a depth of approximately 11 ft bgs (TP2-11).

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Because hydrocarbon impacts were encountered at the former locations of the dispenser island and the USTs, RSE initiated an effort to evaluate the horizontal and vertical extent of hydrocarbon impacts at the subject property. Test pit TP3 was excavated to the north of the former UST location and TP2. Material encountered in TP3 was an unconsolidated and consisted of SW-SM. Some large cobbles were encountered in this area. Perched (non-recharging) water was encountered at approximately 9 ft bgs and impacted soil was encountered at the soil-water interface. One analytical sample (TP3-9) was collected from the just above the soil water interface.

Test pit TP4 was excavated to the west of the former USTs, near the boundary of the adjoining property to the west. TP4 contained mixed debris (wiring, bricks, concrete and building materials) and fill material consisting of SW-SM. Heterogeneous areas within the excavation were impacted by hydrocarbons. The excavation was completed to a depth of approximately 13 ft bgs, where apparent native soil consisting of brown organic silt (OL) with poorly graded fine sand (SP). Two analytical samples were collected from this excavation, one from the fill material overlying the native soil at 11 ft bgs (TP4-12), and the other from the native soil at a depth of 13 ft bgs (TP4-13). The sample from 13 ft bgs exhibited a relatively high PID result for the site and was chosen to characterize VOCs at the subject property.

Test pit TP5 was excavated to the south of the former USTs. TP5 contained fill material consisting of SW-SM. Mixed debris and heterogeneous regions of hydrocarbon impacted fill material were present throughout the excavation. TP5 was completed to a depth of approximately 13 ft bgs, where apparent native soil consisting of brown OL-SP was encountered. One analytical sample was collected from the completed depth of 13 ft bgs (TP5-13).

Test pit TP6 was excavated to the south of the "New Garage" in the vicinity of the former "Wood Shop" building, which is no longer present on site. TP6 contained mixed debris (including a truck bumper) and fill material consisting of SW-SM. Heterogeneous regions of hydrocarbon impacted fill material were present throughout the excavation. TP6 was completed to a depth of 12 ft bgs, where native material consisting of ML was encountered. One analytical (TP6-12) and one duplicate sample (TP6-XI) were collected from native material at the completed depth in this excavation.

4.0 FIELD AND ANALYTICAL SAMPLING METHODOLOGY

During the site characterization effort the test pits were subjected to field screening and analytical sampling. The primary objective of field screening is to qualitatively test for the presence of contamination on site. The primary objective of analytical sampling is to quantitatively identify contaminants that are present on site, and to ensure accurate enough results that samples can be compared to relevant cleanup criteria, and prudent cleanup or disposal decisions can be determined. The following sections provide details on terminology, methodology, and specific data QA/QC requirements that insure these objectives are met.

4.1 FIELD SCREENING METHODS

A photoionization detector (PID) was used during soil screening. The PID was calibrated prior to use in accordance with the manufacturer's calibration specifications. For this project the PID was a MiniRAE Lite Model PGM 7300, and it was calibrated using 100 part per million isobutylene gas.

The following general headspace screening procedure was used to obtain and analyze field screening samples using the PID:

- 1. A clean ziplock bag was partially filled (one-third to one-half) with the sample to be analyzed;
- 2. Headspace vapors were allowed to develop in the bag for at least 10 minutes but no longer than one hour and bags were agitated for 15 seconds at the beginning and end of the headspace development period to assist volatilization;
- 3. The instrument sampling probe was inserted to a point about one-half the headspace depth;
- 4. After probe insertion the highest meter reading was taken and recorded, typically between two and five seconds after probe insertion.

4.2 ANALYTICAL SAMPLE COLLECTION

Samples were placed in method specific containers, tightly closed, and maintained at temperatures between 2° and 6° Celsius during transport to the analytical laboratory. Analytical samples were transported under chain of custody (COC) protocol and submitted to SGS Environmental Services (SGS) in Anchorage for analyses. All volatile samples were accompanied by a trip blank, to be analyzed by the same method.

5.0 SOIL SAMPLE RESULTS

Soil samples were taken of representative soils remaining in place within the test pits. All samples were analyzed for gasoline range organics (GRO), diesel range organics (DRO), residual range organics (RRO); and benzene, toluene, ethyl benzene, and total xylenes (collectively referred to as BTEX) by Alaska Methods (AK) 101, 102, 103, EPA method SW8021B, respectively. One sample (TP4-13) from a depth exhibiting a relatively high field screening result and that was representative of impacted soil at the site was analyzed for volatile organic compounds (VOCs) by Environmental Protection Agency (EPA) method 8260.

Soil sample results are compared to ADEC 18 AAC 75 Method 2 cleanup levels for migration to groundwater (Table 2). ADEC Method 2 cleanup levels for migration to groundwater are 250 milligrams per kilogram (mg/Kg) for DRO, 11,000 mg/Kg for RRO, 300 mg/Kg for GRO, 25 micrograms per kilogram (μ g/Kg) for benzene, 6,500 μ g/Kg for toluene, 6,900 μ g/Kg for ethylbenzene and 63,000 μ g/Kg for total xylenes.

All soil samples exceeded the ADEC cleanup criteria for at least one constituent, with the exception of the sample TP3-9, which exhibited low concentrations of hydrocarbons, less than the ADEC cleanup criteria. Results from sample TP5-13 were less than the ADEC cleanup criteria for every parameter analyzed, with the exception of benzene, which was present at a concentration of 859 µg/Kg.

All other samples exhibited concentrations of at DRO that were greater than the ADEC cleanup criteria. Many samples exhibited concentrations of other analytes that were greater than the ADEC cleanup criteria. Details regarding the particular hydrocarbons above the ADEC cleanup criteria in each test pit are provided below:

Samples collected from TP1 exhibited concentrations of DRO greater than the ADEC cleanup criteria. The sample collected from 3 ft bgs (TP1-3) exhibited a concentration of DRO of 5,160 mg/Kg, while the sample collected from the underlying native soil at a depth of 8 ft bgs (TP3-8) exhibited a concentration of DRO of 892 mg/Kg. Neither of these samples exhibited concentrations of RRO, GRO, or BTEX constituents that were greater than the ADEC cleanup criteria, though results associated with sample TP1-3 had a method detection limit (MDL) that was greater than the ADEC cleanup criteria of 25 μ g/Kg for benzene and does not eliminate benzene as a contaminant of concern at this depth. The MDL for benzene associated with TP3-8 was less than the ADEC cleanup criteria and benzene was not detected in this sample (MDL = 6.67 μ g/Kg for benzene in this sample).

Samples were collected from depths of 6 ft bgs and 11 ft bgs from TP2 (TP2-6 and TP2-11, respectively). Both samples contained concentrations of DRO, GRO, and all BTEX constituents that were significantly greater than the ADEC cleanup criteria.

The sample collected from the TP3 (TP3-9) did not exhibit concentrations of any analyte that were greater than the applicable ADEC cleanup criteria. This test pit was impacted by hydrocarbons, but at concentrations less than the ADEC cleanup criteria.

Samples were collected from depths of 12 ft bgs and 13 ft bgs from TP-4 (TP4-12 and TP4-13, respectively). Both of these samples exhibited concentrations of DRO, GRO, and BTEX constituents that were greater than the ADEC cleanup criteria. Sample TP4-13 was submitted for VOC analysis by EPA method 8260. VOC results confirmed that BTEX constituents are present at concentrations exceeding ADEC cleanup criteria. One other constituent, 1,2,4-trimethylbenzene, was present at a concentration of 61,700 μ g/Kg, a concentration greater than the ADEC cleanup criteria of 23,000 μ g/Kg. No halogenated hydrocarbons were detected in the VOC analysis of this sample. Several other VOC constituents were detected, but at concentrations that were less than their respective ADEC cleanup criteria (See Table 3).

The sample collected from TP-5 contained benzene at a concentration of 859 μ g/Kg, exceeding the ADEC cleanup criteria of 25 μ g/Kg. All other analytes were detected in this sample, but at concentrations that are less than the applicable ADEC cleanup criteria.

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Two analytical samples were collected from the same sample location from excavation TP-6 (TP6-12 and TP6-XI). TP6-XI was submitted to the lab as a blind duplicate. Both samples exhibited concentrations of DRO and benzene that were greater than their respective ADEC cleanup levels.

6.0 ANALYTICAL METHODOLOGY QA/QC

Specific data quality objectives include the measurement of several parameters to establish target analyte detection limits, data precision, data bias, and method comparability.

MDLs are determined in order to insure that contaminants of concern can be quantified at concentrations less than the applicable ADEC cleanup levels. Target MDLs are set below the applicable cleanup criteria, in accordance with 40 CFR part 36, Appendix B.

Laboratory reports were reviewed for QA/QC sample and performance deviations as outlined in our workplan. Data was flagged appropriately according to EPA 540/R-99/008 National Functional Guidelines. Many of these performance criteria are also included in the ADEC laboratory data review checklist, which was completed for this project. No major QC failures affecting sample usability were recorded for this project. The laboratory did not meet the target MDL for benzene for sample TP1-3, which was not detected in the sample. This "non-detect" result can not be used to establish the absence of benzene as a contaminant of concern at this location and has been "R" flagged as a result of this QC failure. Additional minor QA/QC deviations are detailed in the ADEC Laboratory Data Review Checklist, provided in Appendix C.

Several samples are "J" flagged as estimates because the result was greater than the MDL, but below the LOQ. While the analyte has been positively identified in these samples, the error associated with the measurement is sufficiently that the result should be considered an estimate. LOQs associated with "J" flagged results are less than the ADEC cleanup levels and are sufficient to establish that these contaminants are not present at the site at concentrations greater than the ADEC cleanup criteria.

Sample TP6-XI was submitted to the lab as a blind duplicate sample and was collocated with sample TP6-12. While both samples indicated the presence of the same constituents at concentrations that exceeded the ADEC cleanup criteria, the relative percent difference (RPD) between the sample and the field duplicate exceeded the ADEC guidance of 50% for DRO, RRO, GRO, and xylenes, which exhibited RPDs of 92%, 99%, 76%, and 63%, respectively. The RPD for benzene in the two samples was 42%, which met the ADEC RPD criteria. RPDs greater than 50% indicate a high degree of heterogeneity of impacted soil at the subject property, an observation that is consistent with the heterogeneity of the mixed fill materials observed on site.

7.0 INVESTIGATIVE DERIVED WASTE

Test pits were completed and back filled one at a time. All excavated material was placed back into the test pit from which it came at the approximate depth from which it originated. Small quantities of perched groundwater were encountered in some excavations, especially near the building foundation, where drain tile was also observed. Consistently recharging groundwater was not observed in any test pit and test pits were dry at the time they were backfilled.

8.0 DISCUSSION AND CONCLUSIONS

Limitations

This report was prepared for the exclusive use of our client, Dan Aalfs. Conclusions and statements of site conditions are based upon information provided from observations during RSE's field investigation and conversations with onsite personnel. Based upon this, the data and interpretation presented in this report provide our best engineering judgment regarding the environmental characteristics of the site and do not guarantee that additional pollution is absent from the subject property.

Hydrocarbon Impacts

Hydrocarbon impacts are present throughout the site. Hydrocarbon impacted soil was distributed heterogeneously throughout test pits, and many also contained debris such as concrete and wiring. Fill material in test pits was inconsistent and relatively unconsolidated.

TP3 was located to the north of the former tank area and was relatively clean with the exception of a small lens of impacted soil just above a layer of perched groundwater. This impacted soil contained concentrations of hydrocarbons below the applicable ADEC cleanup criteria. TP3 likely represents the northern boundary of contaminated soil present on site.

TP5 was located to the south of the former tank area and contained soil that was impacted by hydrocarbons, but only exceeded the ADEC cleanup criteria for benzene. Notably, some of the lowest levels of GRO observed at the site were present in TP5, perhaps indicating that benzene was migrating into this location from a source location near TP2, which contained GRO concentrations greater than the ADEC cleanup criteria and concentrations of benzene one-to-two orders of magnitude greater than that observed in TP5.

TP1 was located in the former dispenser area and exhibited concentrations of DRO and benzene at concentrations greater than the ADEC cleanup criteria.

TP2 and TP4 were located in the probable location of the former USTs and to the west of the former USTs, respectively. TP2 contained relatively clean soil down to a depth of 6 ft bgs to 7.5 ft bgs, at which point hydrocarbon impacted soil was encountered. TP4 contained mixed fill materials, debris and impacted soil throughout the excavation. Concentrations of DRO, GRO, and BTEX constituents greater than the ADEC cleanup criteria were present in these locations. It should be noted that only TP2 and TP4

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contained concentrations of GRO greater than the ADEC cleanup criteria, implicating this region as a unique source area.

TP6 was located to the south of the new garage, in the vicinity of a wood shed that was previously removed from the subject property. TP6 contained mixed fill, debris, including a truck bumper, and impacted soil throughout the excavation. It should be noted that DRO was present at concentrations not only exceeding the ADEC cleanup criteria, but also exceeding DRO concentrations encountered in other test pits on site. This test pit also exceeded the ADEC cleanup criteria for benzene. The presence of large quantities of DRO and benzene in the relative absence of GRO, toluene, ethylbenzene, and xylenes in this area may indicate a unique source.

Garage Building Floor Drains

In addition to test pits excavated on the subject property, the garage drain system was checked using tracer dye to confirm that the garage drain system discharged to the public sanitary sewer. Tracer dye and large quantities of water were introduced into the drains. The sanitary sewer was observed at a sanitary sewer manhole located at the top of the bluff near the properties southwest boundary. Observation of tracer dye and dramatic changes in flow rate associated with the introduction of tracer dye and water confirm that the buildings drain system is connected to the City of Homer municipal sanitary sewer system.

Conceptual Site Model

Significant hydrocarbon impacts to subsurface soil are present at the subject property. Media impacts and potential transport mechanism were evaluated using the ADEC Human Health Conceptual Site Model Graphic Form. This form identified potential site works, visitors, and construction workers as potential receptors via incidental soil ingestion, dermal absorption, and inhalation of fugitive dust and organic vapors in outdoor area. The building present on site is currently unused and secured. A copy of the completed Human Health Conceptual Site Model Graphic Form is provided in Appendix B.

Water Supply Well Search

Because of hydrocarbon impacts to subsurface soil and the potential for these hydrocarbons to migrate to groundwater a drinking water well search was performed using the State of Alaska Division of Mining, Land and Water Alaska Hydrologic Survey Well Log Tracking System (WELTS). This search was performed for Township 6 south, Range 13 west, section 20. Results from the WELTS search indicated that no drinking water wells were present in this section. The absence of drinking water wells in the area is consistent with fact that water in the area surrounding the subject property is supplied by the City of Homer from the Bridge Creek Reservoir. The reservoir is located approximately 2.3 miles to the north and at a considerably higher elevation than the subject property. The Bridge Creek Reservoir is not considered a potential receptor for contamination originating at the subject property.

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Estimate Quantity of Impacted Soil

Based on field observations and laboratory results several areas within subject property are impacted by hydrocarbon contamination at concentrations exceeding ADEC cleanup levels. These areas include the former dispenser island, which is impacted by DRO, but not by benzene; The vicinity of the former USTs located to the west of the New Garage building, which is impacted by DRO, GRO and BTEX constituents; and the area south of the New Garage building which is impacted by DRO and benzene. Test pit TP5 was advanced to the southeast of the New Garage building and was only impacted by benzene at concentrations greater than the ADEC cleanup criteria. The total quantity of soil at the subject property that is impacted by hydrocarbons at a concentration greater than the ADEC cleanup criteria is estimated to be approximately 1,600 cubic yards.

Restoration Science & Engineering, LLC

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References

Waller, R.M., Feulner, A.J., Morris, D.A. "Water Resources and Surficial Geology of the Homer Area, South-Central Alaska" USGS Hydrologic Investigations Atlas HA-187. 1968.

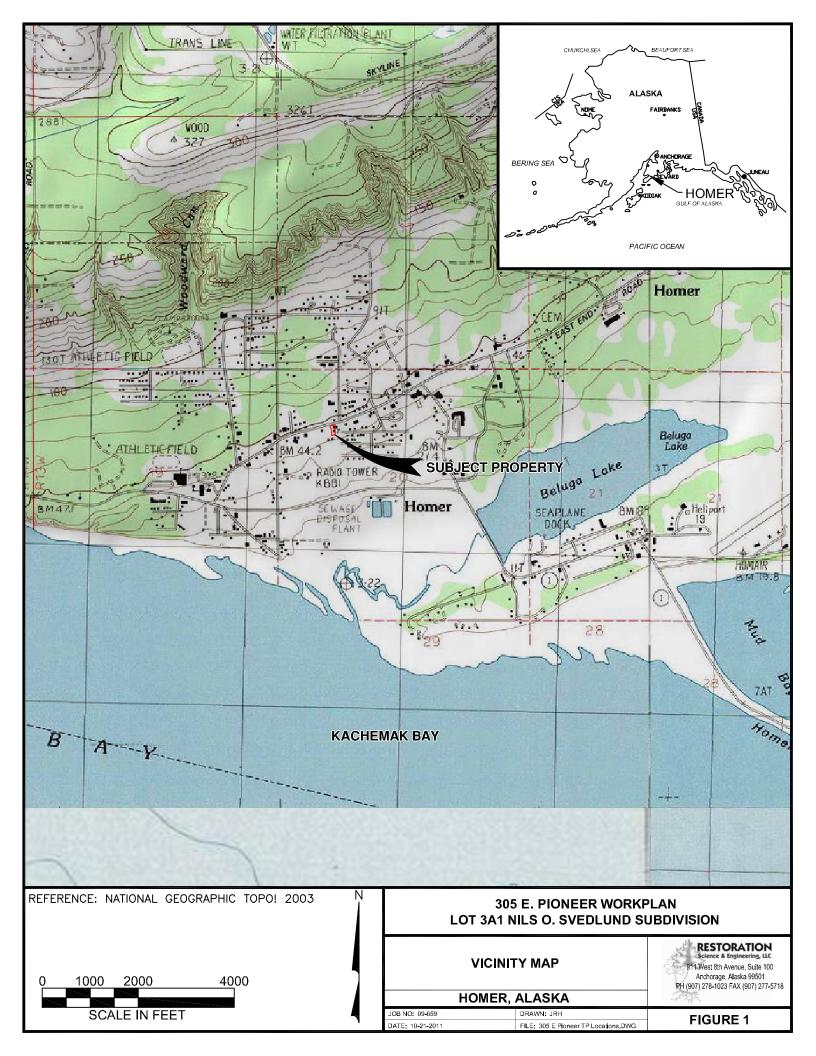
May 18, 1970 Aerial Photograph, Aeromap.

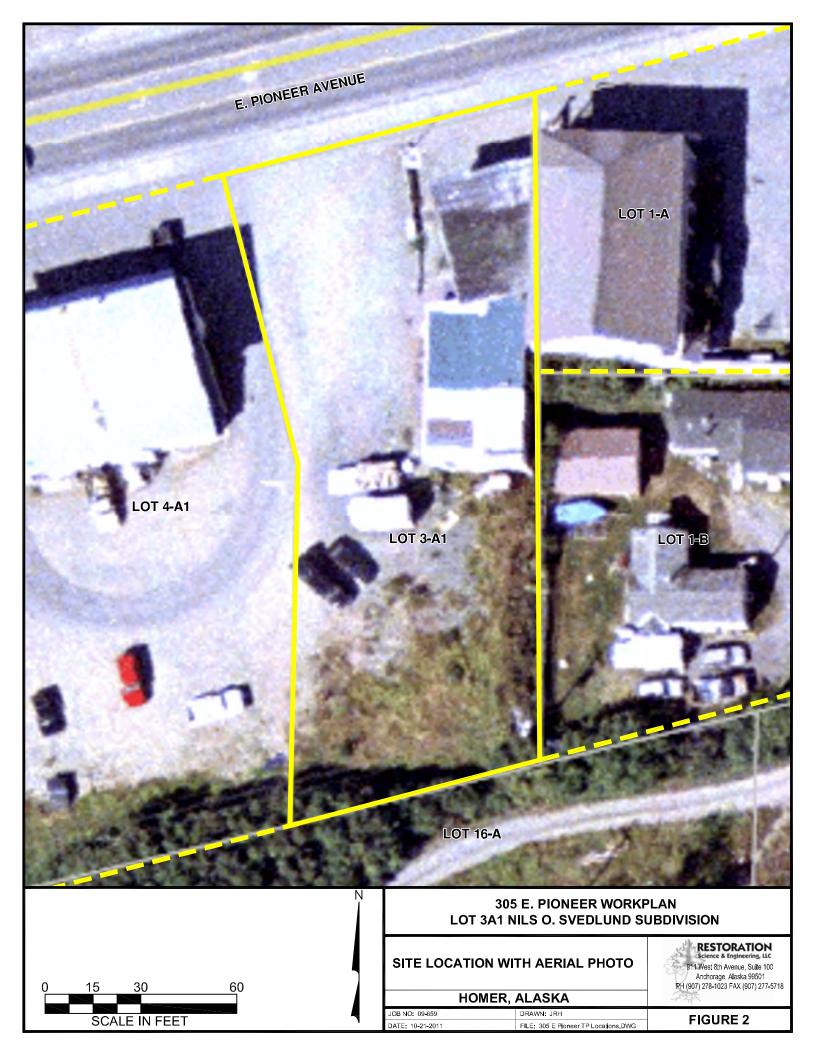
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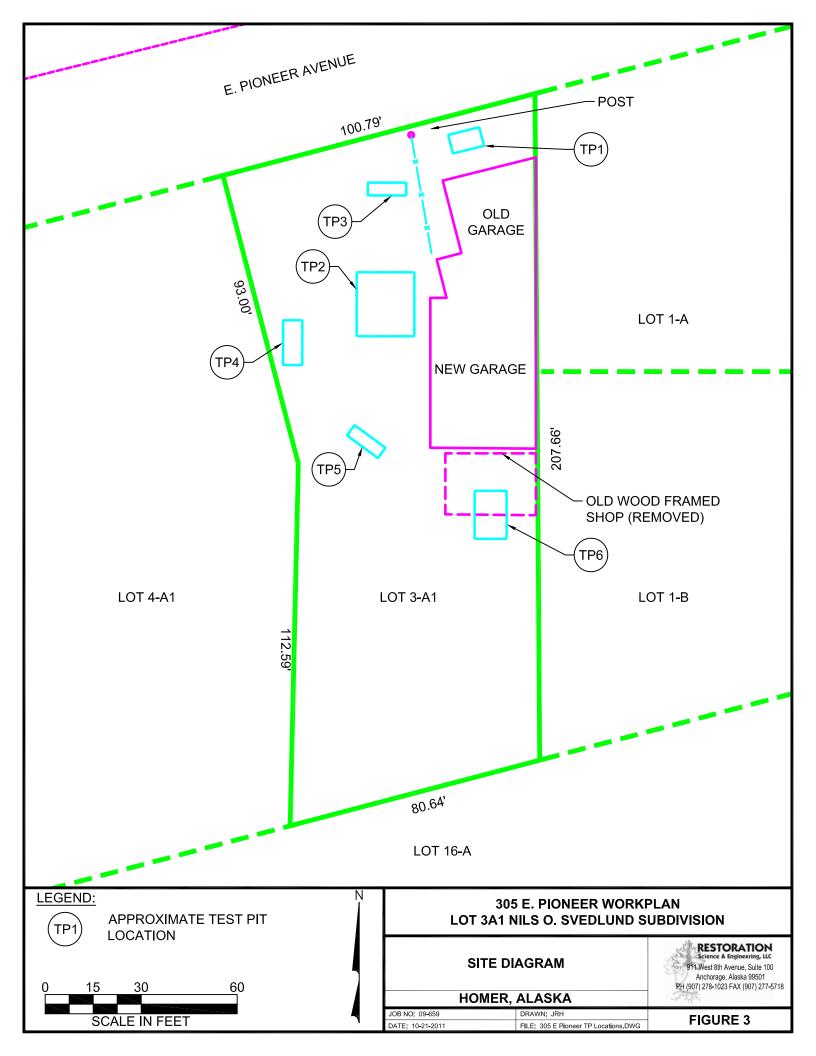
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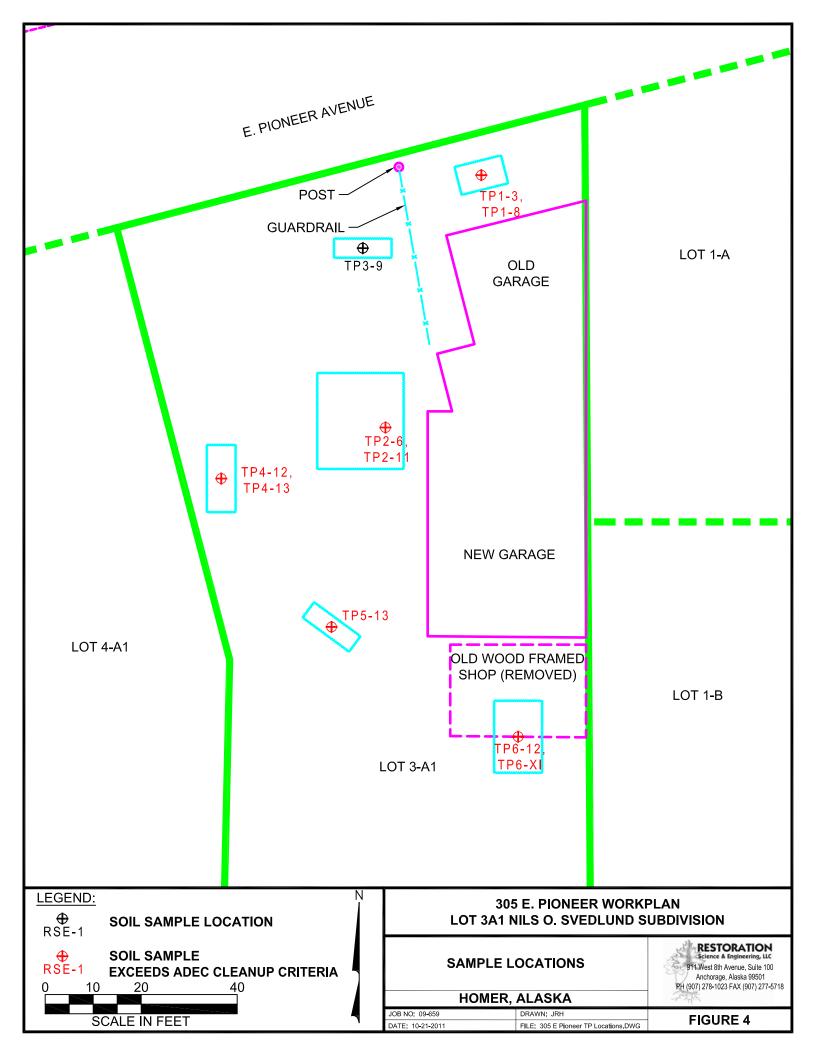
Appendix A: Figures











Appendix B: Tables



LIMITED SITE ASSESSMENT, 305 EAST PIONEER HEADSPACE CONCENTRATIONS IN SOIL NOVEMBER 2011 REPORT DATE

		HEADSP	ACE CONCEN	TRATIONS IN SOIL
Screening Sample ID	Date	PID Headspace Concentration (10.2 eV)	Depth Below Ground Surface	NOTES
		(ppmv)	(feet)	
TP1-3	9/20/2011	606	3	(apparent fill) SW-SM
TP1-5	9/20/2011	0.0	5	(apparent fill) SW-SM
TP1-7	9/20/2011	1.2	7	(apparent fill) SW-SM
TP1-8	9/20/2011	44.1	8	(apparent fill) SW-SM
TP1-8	9/20/2011	66	8	(apparent fill) SW-SM
TP1-8	9/20/2011	5.8	8	(native materials) ML-SP
TP2-7.5	9/20/2011	2345	7.5	(apparent fill) SW-SM
TP2-6	9/20/2011	502	6	(apparent fill) SW-SM
TP2-11	9/20/2011	365	11	(native materials) ML-SP
TP3-9	9/20/2011	3.9	9	(apparent fill) SW-SM
TP4-11	9/20/2011	210	11	(apparent fill) SW-SM
TP4-13	9/20/2011	899	13	(native materials) ML-SP
TP5-11	9/20/2011	240	11	(apparent fill) SW-SM
TP5-13	9/20/2011	215	13	(native materials) ML-SP
TP6-6	9/21/2011	396	6	(apparent fill) SW-SM
TP6-10	9/21/2011	1005	10	(apparent fill) SW-SM
TP6-12	9/21/2011	713	12	(native materials) ML-SP
				·

- 1) All field screening performed using a MiniRae Lite, Model PGM 7300 photoionization detector calibrated to 100 ppmv isobutylene.
- 2) Screening samples were warmed to approximately 60 °F before reading.
- 3) PID = photoionization detector, ppmv = parts per million by volume
- 4) Bold samples are associated with laboratory samples of the same name.
- 5) USCS symbols: SW = Well Graded Sand, SP = Poorly Graded Sand, SM = Silty Sand, ML = Silt

TABLE 2 LIMITED SITE ASSESSMENT, 305 EAST PIONEER HYDROCARBON CONCENTRATIONS IN SOIL NOVEMBER 2011 REPORT DATE

SAMPLE ID	DATE	PID RESULTS	DIESEL RANGE ORGANICS	RESIDUAL RANGE ORGANICS	GASOLINE RANGE ORGANICS	BENZENE	TOLUENE	ETHYL- BENZENE	TOTAL XYLENES
		(ppmv)	(mg/Kg)	(mg/Kg)	(mg/Kg)	(μg/Kg)	(μg/Kg)	(μg/Kg)	(μg/Kg)
TP1-3	9/19/2011	606	5160	1340	110	ND (34.1) R	67.7 J	704	991
TP1-8	9/19/2011	5.8	892	1460	1.71 J	ND (6.67)	ND (13.0)	ND (13.0)	ND (83.4)
TP2-6	9/19/2011	502	3680	130	789	8470	1350	38700	131905
TP2-11	9/19/2011	365	6570	612	2050	28000	64000	103000	511000
TP3-9	9/19/2011	3.9	76.5	156	ND (0.634)	5.07 J	ND (6.59)	ND (6.59)	ND (42.3)
TP4-12	9/19/2011	210	1270	81.0	1370	13400	141000	60800	276100
TP5-13	9/19/2011	215	193	764	7.90 J	859	36.2 J	77.0 J	1818
TP6-12	9/19/2011	215	39700	6800	40.0	428	ND (80.4)	91.5 J	4000
TP6-XI	9/19/2011		14700	2290	89.2	653	334	98.1 J	7760
TP4-13 ²	9/19/2011	713	1850	415	720				
EC Method 2		p levels for	250	11000	300	25	6500	6900	63000

- 1) Gasoline Range Organics by Method AK 101, Diesel range organics analyses by Method AK 102, Residual Range Organics by Method AK103, BTEX (benzene, toluene, ethylbenzene and total xylenes) analyses by Method EPA 8021B.
- 2) Sample TP4-13 was analyzed for Volatile Organic Compounds by EPA 8260, VOC results are provided in Table 3.
- 3) The J flag indicates that the result is an estimated value.
- 4) ND indicates that the analyte was not detected. The limit of detection is given in parantheses.
- 5) Bold Text indicates that a sample result is greater than the ADEC cleanup levels.
- 6) The R flag indicates that the this data point has been rejected. The limit of detection associated with the non-detection of benzene in this sample is greater than the ADEC cleanup criteria: benzene may be present at concentrations greater than the ADEC cleanup criteria at this location.
- 7) ppmv= parts per million by volume, mg/Kg = milligrams per kilogram, µg/Kg = micrograms per kilogram

TABLE 3 LIMITED SITE ASSESSMENT, 305 EAST PIONEER VOC CONCENTRATIONS IN SOIL NOVEMBER 2011 REPORT DATE

HYDROCARBON CONCENTRATIONS IN SOIL							
SAMPLE ID	VOLATILE ORGANIC COMPOUNDS BY METHOD 8260 ANALYTES	RESULT	LIMIT OF QUANTITATION	LIMIT OF DETECTION	ADEC CLEANUP CRITERIA		
		(μg/Kg)		(μg/Kg)	(μg/Kg)		
TP4-13	1,1,1,2-Tetrachloroethane	ND	272	84.9	NA		
	1,1,1-Trichloroethane	ND	272	84.9	18		
	1,1,2,2-Tetrachloroethane	ND	544	163	17		
	1,1,2-Trichloroethane	ND	272	84.9	18		
	1,1-Dichloroethane	ND	272	84.9	NA		
	1,1-Dichloroethene	ND	272	84.9	NA		
	1,1-Dichloropropene	ND	272	84.9	NA		
	1,2,3-Trichlorobenzene	ND	544	163	NA		
	1,2,3-Trichloropropane	ND	272	84.9	0.53		
	1,2,4-Trichlorobenzene	ND	272	84.9	850		
	1,2,4-Trimethylbenzene	61700	5440	1630	23000		
	1,2-Dibromo-3-chloropropane	ND	1090	338	NA		
	1,2-Dibromoethane	ND	272	84.9	NA		
	1,2-Dichlorobenzene	ND	272	84.9	5100		
	1,2-Dichloroethane	ND	272	84.9	2500		
	1,2-Dichloropropane	ND	272	84.9	18		
	1,3,5-Trimethylbenzene	14000	272	84.9	23000		
	1,3-Dichlorobenzene	ND	272	84.9	33		
	1,3-Dichloropropane	ND	272	84.9	NA		
	1,4-Dichlorobenzene	ND	272	84.9	640		
	2,2-Dichloropropane	ND	272	84.9	NA		
	2-Butanone (MEK)	ND	2720	849	59000		
	2-Chlorotoluene	ND	272	84.9	NA		
	2-Hexanone	ND	2720	849	NA		
	4-Chlorotoluene	ND	272	84.9	NA		
	4-Isopropyltoluene	694	272	849	NA		
	4-Methyl-2-pentanone (MIBK)	ND	2720	849	8100		
	Benzene	14300	136	42.5	25		
	Bromobenzene	ND	272	84.9	NA		
	Bromochloromethane	ND	272	84.9	NA		
	Bromodichloromethane	ND	272	84.9	44		
	Bromoform	ND	272	84.9	340		
	Bromomethane	ND	2180	675	NA		

- 1) The J flag indicates that the result is an estimated value.
- 2) ND indicates that the analyte was not detected. The limit of detection is given in parantheses.
- 3) Bold Text indicates that a sample result is greater than the ADEC cleanup levels.
- **4)** μg/Kg = micrograms per kilogram

TABLE 3 LIMITED SITE ASSESSMENT, 305 EAST PIONEER VOC CONCENTRATIONS IN SOIL NOVEMBER 2011 REPORT DATE

	HYDROCARBON CONCENTRATIONS IN SOIL							
SAMPLE ID	VOLATILE ORGANIC COMPOUNDS BY METHOD 8260 ANALYTES	RESULT	LIMIT OF QUANTITATION	LIMIT OF DETECTION	ADEC CLEANUP CRITERIA			
		(μg/Kg)		(μg/Kg)	(μg/Kg)			
TP4-13	Carbon disulfide	ND	1090	338	12000			
	Carbon tetrachloride	ND	272	84.9	23			
	Chlorobenzene	ND	272	84.9	630			
	Chloroethane	ND	2180	675	32			
	Chloroform	ND	272	84.9	460			
	Chloromethane	ND	272	84.9	NA			
	cis-1,2-Dichloroethene	ND	272	84.9	16			
	cis-1,3-Dichloropropene	ND	272	84.9	33			
	Dibromochloromethane	ND	272	84.9	NA			
	Dibromomethane	ND	272	84.9	NA			
	Dichlorodifluoromethane	ND	544	163	140000			
	Ethylbenzene	39200	2720	849	6900			
	Hexachlorobutadiene	ND	544	163	120			
	Isopropylbenzene (Cumene)	4360	272	84.9	5100			
	Methylene chloride	ND	1090	338	210			
	Methyl-t-butyl ether	ND	1090	338	1300			
	Naphthalene	9810	544	163	20000			
	n-Butylbenzene	3240	272	84.9	15000			
	n-Propylbenzene	11200	272	84.9	15000			
	o-Xylene	58000	2720	849	63000			
	P & M -Xylene	134000	5440	1630	63000			
	sec-Butylbenzene	1120	272	84.9	12000			
	Styrene	ND	272	84.9	960			
	tert-Butylbenzene	ND	272	84.9	12000			
	Tetrachloroethene	ND	136	42.5	24			
	Toluene	122000	2720	849	6500			
	trans-1,2-Dichloroethene	ND	272	84.9	370			
	trans-1,3-Dichloropropene	ND	272	84.9	33			
	Trichloroethene	ND	136	42.5	NA			
	Trichlorofluoromethane	ND	544	163	86000			
	Vinyl chloride	ND	272	84.9	8.5			
	Xylenes (total)	192000	10900	3380	63000			

- 1) The J flag indicates that the result is an estimated value.
- 2) ND indicates that the analyte was not detected. The limit of detection is given in parantheses.
- 3) Bold Text indicates that a sample result is greater than the ADEC cleanup levels.
- 4) µg/Kg = micrograms per kilogram

Appendix C: Selected Site Photographs





Photo 1. Subject Property (Looking South)



Photo 3. Breaking Ground at Test Pit One (Looking South)



Photo 2. Subject Property (Looking Northeast)



Photo 4. Wooden Crib in Test Pit One



Photo 5. Test Pit Two (Looking Southwest).



Photo 7. Breaking Ground at Test Pit Six located to the South of the Garage (Looking North)



Photo 6. Typical Test Pit containing debris, including wiring, concrete, building materials and bricks.



Photo 8. Debris and impacted soil encountered in Test Pit Six.

Appendix D: Human Health Conceptual Site Model Graphic Form



HUMAN HEALTH CONCEPTUAL SITE MODEL GRAPHIC FORM

Current & Future Receptors exposure pathway: Enter "C" for current receptors "F" for future receptors, "C/F" for both current and dentify the receptors potentially affected by each future receptors, or "I" for insignificant exposure. Revised, 10/01/2010 Other Subsistence consumers Farmers or subsistence harvesters Instructions: Follow the numbered directions below. Do not consider contaminant concentrations or engineering/land Sife visitors, trespassers, or recreational users × × × × Commercial or industrial workers × × × Residents (adults or children) × \times \times × × use controls when describing pathways. Dermal Absorption of Contaminants in Surface Water Dermal Absorption of Contaminants in Groundwater ☐ Inhalation of Volatile Compounds in Tap Water Inhalation of Volatile Compounds in Tap Water Dermal Absorption of Contaminants from Soil The pathways identified in this column **must Exposure Pathway/Route** Check all pathways that could be complete. agree with Sections 2 and 3 of the Human Ingestion of Wild or Farmed Foods Health CSM Scoping Form Direct Contact with Sediment Ingestion of Surface Water ✓ Inhalation of Fugitive Dust Inhalation of Fugitive Dust Ingestion of Groundwater ✓ Inhalation of Outdoor Air ✓ Incidental Soil Ingestion Inhalation of Indoor Air **Exposure Media** media identified in (2) surface water ✓ groundwater Check all exposure sediment biota ල soil aj $\overline{\ }$ > check biota check groundwater) check soil check soil check surface water check groundwater check groundwate check biota check sedimer For each medium identified in (1), follow the mechanisms. Check additional media under if the media acts as a secondary source. **Transport Mechanisms** top arrow <u>and</u> check possible transport 305 East Pioneer Avenue, Homer, Alaska Resuspension, runoff, or erosion Uptake by plants or animals Uptake by plants or animals 📙 Uptake by plants or animals Uptake by plants or animals [Uptake by plants or animals Flow to surface water body Direct release to subsurface soil ✓ Migration to groundwater[Direct release to surface water Migration to groundwater Direct release to groundwater Migration to subsurface Direct release to surface soil Direct release to sediment ADEC file # 2314.26.031 Runoff or erosion Flow to sediment[Completed By: Nick Braman Sedimentation Volatilization [Volatilization [Volatilization Volatilization Date Completed: 10/20/11 Other (list): Other (list): Other (list): Other (list): Other (list): could be directly affected Check the media that (2-15 ft bgs) Subsurface (0-2 ft bgs) Media Sediment by the release. Surface Ground-Surface Water water Soil > Site:

Appendix E:
ADEC Laboratory Data Review Checklist
And Laboratory Data



Laboratory Data Review Checklist

Completed by:		Nick Braman					
Title:		Environmental	Scientist/Chemis	st	Date:	Oct 19, 2011	
CS Re	eport Name:	UST Closure A	Assessment, 305 I	E. Pioneer, Homer	Report Date:		
Consu	ıltant Firm:	Restoration Sc	ience and Engine	ering			
Labor	atory Name:	SGS		Laboratory Report Number: 1114619			
ADEC	File Number:	2314.26.031		ADEC RecKey Number:			
1. <u>L</u>	aboratory a. Did an A	ADEC CS appro	oved laboratory r	eceive and perform all o	of the submitted	sample analyses?	
	• Yes	○ No	○ NA (Plea	-	Comments:	r	
		-		r "network" laboratory og the analyses ADEC C		d to an alternate	
	○ Yes	○ No • NA (Please explain)		e explain)	Comments:		
	Samples were n	ot transferred.					
2. <u>Cl</u>	nain of Custody	(COC)					
	a. COC infor	mation complet	ed, signed, and d	ated (including released	/received by)?		
r	• Yes	○ No	ONA (Pleas	e explain)	in) Comments:		
		alyses requeste					
г	• Yes	○ No	○ NA (Plea	ase explain)	Comments:		
3. <u>La</u>	aboratory Sampl	e Receipt Docu	mentation				
	a. Sample/co	oler temperatur	e documented and	d within range at receipt	$\pm (4^{\circ} \pm 2^{\circ} \text{ C})$?		
	• Yes	○ No	○NA (Ple	ase explain)	Comments:		

	servation acceptorinated Solve	-	preserved VOC soil (GRO, BTEX,
• Yes	○ No	○ NA (Please explain)	Comments:
c. Sample con	dition docume	nted - broken, leaking (Methanol),	zero headspace (VOC vials)?
○ Yes	○ No	NA (Please explain)	Comments:
All sample condi	tions were ade	quate for the analyses ordered.	
		•	r example, incorrect sample containers nsufficient or missing samples, etc.?
Yes	○ No	ONA (Please explain)	Comments:
provided on the la	boratory repor	t are one day ahead of the actual co	than they actually were, sample dates ellection date.
e. Data quanty	y or usaomity a	ffected? (Please explain)	Comments:
Data quality is no	ot affected by ϵ	erroneous sample dates.	
Case Narrative a. Present and • Yes	understandabl	e? ○ NA (Please explain)	Comments:
b. Discrepance	ies, errors or Q	C failures identified by the lab?	
• Yes	○ No	○NA (Please explain)	Comments:
c. Were all co	rrective action	s documented? ○ NA (Please explain)	Comments:
d. What is the	effect on data	quality/usability according to the c	ase narrative? Comments:
See details on OO	 C failures prese	ented in later sections.	

	○ No	ONA (Please explain)	Comments:
b. All applicab	le holding tim	es met?	
• Yes	○ No	ONA (Please explain)	Comments:
c. All soils rep	orted on a dry	weight basis?	
• Yes	○ No	○ NA (Please explain)	Comments:
d. Are the repo	orted PQLs les	s than the Cleanup Level or the min	nimum required detection level for the
○ Yes	No	ONA (Please explain)	Comments:
greater than the N	IDL but less t	antification (LOQ) and a method do han the LOQ are presented, but J flatfected? (Please explain)	etection limit (MDL). Sample resultagged. Comments:
		le concentration of benzene with a lablish that benzene is not present at	MDL greater than the ADEC cleanut concentrations that may require
-	k		
a. Method Blan		ported per matrix, analysis and 20 sa	amples?
a. Method Blan	thod blank rep	oorted per matrix, analysis and 20 sa	Comments:
a. Method Blan i. One me	thod blank rep	○ NA (Please explain)	•
i. One me	thod blank rep No No	ONA (Please explain)	Comments:
a. Method Blan i. One me	thod blank rep No No	○ NA (Please explain)	•

5. <u>Samples Results</u>

○ Yes	○ No	NA (Please explain)	Comments:			
Sample met me	thod blank crit	eria.				
v. Data q	uality or usabi	lity affected? (Please explain)	Comments:			
b. Laboratory	y Control Sam	ple/Duplicate (LCS/LCSD)				
_	i. Organics - One LCS/LCSD reported per matrix, analysis and 20 samples? (LCS/LCSD required per AK methods, LCS required per SW846)					
• Yes	○ No	○ NA (Please explain)	Comments:			
ii. Metals samples?	•	One LCS and one sample duplicate r	reported per matrix, analysis and 20			
○ Yes	○ No	NA (Please explain)	Comments:			
No metals or in	organic analys	is was performed on these samples.				
project sp	pecified DQOs	ent recoveries (%R) reported and wi , if applicable. (AK Petroleum meth %-120%; all other analyses see the la				
○ Yes	No	ONA (Please explain)	Comments:			
Several analyte	recoveries we	quantified due to a lack of surrogate re biased high in LCS/LCSD sample tates that MIBK recovery was biased	es associated with the 8260 VOC			
limits? A	and project spec	cified DQOs, if applicable. RPD rep	ed and less than method or laboratory ported from LCS/LCSD, MS/DMSD, and all other analyses see the laboratory QC			
• Yes	○ No	ONA (Please explain)	Comments:			
v. If %R	or RPD is outs	ide of acceptable limits, what sample	les are affected? Comments:			

O Y	es	• No	○ NA (Please explain)	Comments:
vii. E) ata qu	ality or usab	oility affected? (Please explain)	Comments:
LCS result usability is		_	, but the analyte was not detected in t	the affected samples, therefore data
c. Surrog	ates -	Organics On	ıly	
i. Are	surro	gate recoveri	es reported for organic analyses - fie	ld, QC and laboratory samples?
⊚ Y	es	○ No	CNA (Please explain)	Comments:
proje	ct spec		`	in method or laboratory limits? And ds 50-150 %R; all other analyses see
0	Yes	No	○ NA (Please explain)	Comments:
by co-elution	on of s optes s	urrogate wit withpleighscott	h target compounds (biased high) and	urrogate recoveries are affected both d due to sample dilutions necassary to ve data flags? If so, are the data flags
clear	ly defi	ned?		
\bigcirc Y	es	No	○ NA (Please explain)	Comments:
associated v	vith fa	iled surrogat	to failed surrogate recoveries because e recoveries were often significantly ility affected? (Use the comment box	to explain.).
				Comments:
l			for establishing that hydrocarbons at ne respective sample locations.	concentrations greater than the ADEC
-	lank -	Volatile ana	lyses only (GRO, BTEX, Volatile Cl	hlorinated Solvents, etc.): Water and
	-	olank reporte r explanation	ed per matrix, analysis and for each con below.)	ooler containing volatile samples?
• Ye	S	○ No	○ NA (Please explain.)	Comments:
			transport the trip blank and VOA same plaining why must be entered below.	<u> </u>
○ Ye	S	○ No	• NA (Please explain.)	Comments:
One cooler v	vas us	ed for all sar	nples.	

• Yes	O No	O NA (Please explain.)	Comments:
iv. If abov	ve PQL, what	samples are affected?	
			Comments:
v. Data qu	ıality or usabil	ity affected? (Please explain.)	
			Comments:
e. Field Duplic			
1. One field	a duplicate sut	omitted per matrix, analysis and 10 p	project samples?
• Yes	○ No	ONA (Please explain)	Comments:
ii. Submit	tted blind to la	b?	
• Yes	○ No	O NA (Please explain.)	Comments:
• Yes	○ No	O NA (Please explain.)	Comments:
• Yes	○ No	O NA (Please explain.)	Comments:
iii. Precisi	ion - All relati	ve percent differences (RPD) less th	
iii. Precisi	ion - All relati		
iii. Precisi	ion - All relation nmended: 30%	ve percent differences (RPD) less th % water, 50% soil) RPD (%) = Absolute Value of: (R ₁ -	an specified DQOs? $\frac{R_2}{x}$ 100
iii. Precisi (Recor	ion - All relati nmended: 30% F	we percent differences (RPD) less the water, 50% soil) RPD (%) = Absolute Value of: (R_{1+}, R_{2+})	an specified DQOs? $\frac{R_2}{x}$ 100
iii. Precisi (Recor	ion - All relation mmended: 30% $R_1 = Sample Co$	we percent differences (RPD) less the water, 50% soil) RPD (%) = Absolute Value of: (R_{1+}, R_{2+})	an specified DQOs? $\frac{R_2}{x}$ 100
iii. Precisi (Recor Where R	ion - All relative mmended: 30% From $R_1 = Sample Color = Sampl$	we percent differences (RPD) less the water, 50% soil) RPD (%) = Absolute Value of: (R_{1-}) ((R_{1+} Reported to the concentration icate Concentration)	an specified DQOs? R ₂) x 100 2/2)
iii. Precisi (Recor	ion - All relation mmended: 30% $R_1 = Sample Co$	we percent differences (RPD) less the water, 50% soil) RPD (%) = Absolute Value of: (R_{1+}, R_{2}) concentration	an specified DQOs? $\frac{R_2}{x}$ 100
iii. Precisi (Recor Where R R	ion - All relation mmended: 30% F R ₁ = Sample Co ₂ = Field Dupl • No	we percent differences (RPD) less the water, 50% soil) RPD (%) = Absolute Value of: (R_{1-}) ((R_{1+} Reported to the concentration icate Concentration)	an specified DQOs? R ₂)_x 100 2)/2) Comments:

f	f. Decontamination or Equipment Blank (if applicable)							
	○ Yes	No	ONA (Please explain)	Comments:				
	i. All resul	ts less than PC	QL?					
	○ Yes	○ No	NA (Please explain)	Comments:				
No o	No decon blank was perfomed for this project.							
	ii. If above PQL, what samples are affected? Comments:							
	iii. Data qu	ality or usabil	ity affected? (Please explain.)	Comments:				
	Data Flags/Qu	`	DE, AFCEE, Lab Specific, etc.)					
	○ Yes		ONA (Please explain)	Comments:				
than dete exco MD	SGS presents both a limit of quantification (LOQ) and a method detection limit (MDL). Samples greater than the MDL but less than the LOQ are presented with "J" flags, indicating the quantity is an estimate detected below the LOQ. In all of these cases the MDL is less than the applicable cleanup standards, except as explained in 5(e), above. Many samples containing large quantities of BTEX constituents had MDLs or LOQs that were greater than the ADEC cleanup levels due to sample dilution Reset Form samples with high quantities of analytes. These samples are not affected by PQLs greater than the TEDE							

cleanup criteria, becuase the results themselves are greater than the ADEC cleanup criteria.

Appendix F: Field Notes

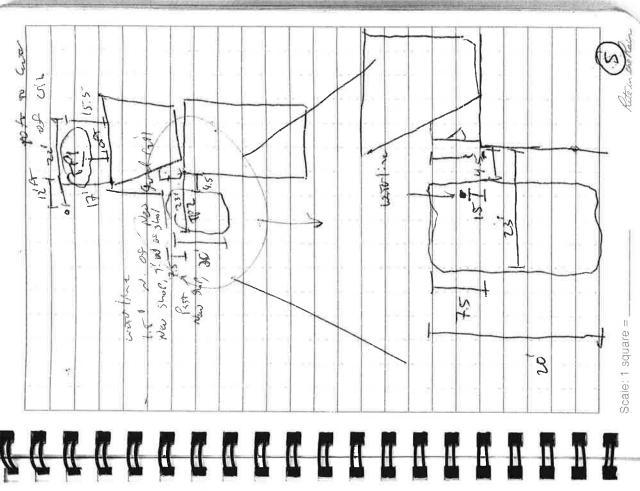


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	DATE				
CONTENTS	REFERENCE				
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fill TRY. PR. 11	51-hdh	(3 - 13' 103 , Ser, U.	195-13 218 4:32	being of ster, filled in excessioners	2/2	4.6 (3:30		Scale: 1 square =
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