Department of Environmental Health

FEB 18 1997

CORRECTIVE / INTERIM REMEDIAL ACTION PLAN FOR STID 553 -FORMER GRIMIT AUTO AND REPAIR 1970 SEMINARY AVENUE OAKLAND, CALIFORNIA

February 15, 1997

Prepared by

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Geology / Engineering Geology / Environmental Studies

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February 15, 1997

E-10-1B-192B HC CAP/RAP/ClosPlan: Seminary2/97CAP/RAP

Mr. Doyle Grimit 14366 Lark Street San Leandro, California 94578

RE: CORRECTIVE / INTERIM REMEDIAL ACTION PLAN STID 553 - FORMER GRIMIT AUTO AND REPAIR 1970 SEMINARY AVENUE OAKLAND, CALIFORNIA

Dear Mr. Grimit:

As authorized, we have prepared the attached Corrective / Interim Remedial Action Plan, to address the existing site conditions and remedial action objectives, and to establish a costeffective remedial action plan. The goal of the plan is to achieve continued source removal and reduction of health and environmental risks associated with the project site, with the ultimate goal of achieving site closure.

We appreciate the opportunity to provide services to you on this project and trust this report meets your needs at this time. If you have any questions, or require additional information, please do not hesitate to call.

Very truly yours,

HOEXTER CONSULTING, INC.

David F. Hoexter, RG/CEG/REA Principal Geologist

CORRECTIVE / INTERIM REMEDIAL ACTION PLAN

For

STID 553 - Former Grimit Auto and Repair 1970 Seminary Avenue Oakland, California

То

Mr. Doyle Grimit 14366 Lark Street San Leandro, California 94578

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JOB NO. E-10-18-1923

15/87 CAP/ Interim RAP DESCRIPTION

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COMMENTS

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Hoexter

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EXECUTIVE SUMMARY

Ground water contamination extends off-site to a residential area. ASTM RBCA analysis has shown that risk based screening levels (RBSLs) are exceeded by as much as four orders of magnitude. Corrective action is proposed which emphasizes remediation of the site's "source area". Results of an SME statement suggest that SME stands the effective for romediation of coupling the site, but would not successfully remediate upsts and and section of extraction wells and associated equipment operation and maintenance costs. The proposed remediation plan would consist of:

Installation of additional down gradient monitoring wells.

^{*} Installation of oxygen releasing compounds (ORCs) in soil borings throughout the "source area". An initial small scale calibration test is proposed. The ORCs would be installed as a slurry directly into the calibration test soil borings, and would be pressure injected into the "full scale" remediation program.

Installation of a low volume in-situ bioventing vacuum system in existing and proposed monitoring wells.

TABLE OF CONTENTS

Page No.

LETT	ER OF TRANSMITTAL		
TITLE	EPAGE		
EXEC	JUTIVE SUMMARY		
TABL	LE OF CONTENTS		
1.0	INTRODUCTION		1
2.0	BACKGROUND	* * * * * * * * * * * * *	····‡
2.1	Introduction		
2.2	Investigation History	••••••	2
2.3	Subsurface Conditions and Contaminant Source	***********	
2.4	Contaminant Description	• • • • • • • • • • • • •	2
3.0	PREVIOUS REMEDIAL ACTION EVALUATION	• • • • • • • • • • • • •	5
4.0	RISK EVALUATION	*********	U Q
5.0	REMEDIAL OBJECTIVES	••••••	0
6.0	REMEDIAL SYSTEM DESIGN CONSIDER ATIONS	• • • • • • • • • • • • • •	·9 10
6.1	Site Characteristics		10 10
6.2	Soil Vapor Extraction (SVE) Pilot Study (January, 1997)		10
6.3	Evaluation of SVE Effectiveness	•••••••	10
7.0	PROPOSED PLAN	**********	10
7.1	Introduction and Rationale	********	12
7.2	General Remediation Plan	********	12
7.2.1	Small Scale ORC Calibration Test	**********	12
7.2.2	Additional Monitoring Wells		13
7.2.3	Full Scale ORC Remediation	*********	13
7.2.4	In-Situ Bioventing	******	11
8.0	ADDITIONAL INVESTIGATION	••••••	15
9.0	SCHEDULE/DURATION OF TREATMENT	********	16
10.0	ANALYTICAL TESTING AND REPORTING	*********	17
11.0	FINAL REMEDIAL ACTION PLAN	********	12
12.0	LIMITATIONS	•••••••	10
		* * * * * * * * * * * *	***2

REFERENCES

FIGURES

1-	Location Map	

Site Plan and Proposed Remediation 2-

APPENDICES

- A-
- B-
- Analytical Data Summary Tables RBCA Analysis Summary Tables Terra Vac February 5, 1997 Report Ĉ-

CORRECTIVE / INTERIM REMEDIAL ACTION PLAN STID #553 FORMER GRIMIT AUTO AND REPAIR 1970 SEMINARY AVENUE OAKLAND, CALIFORNIA

1.0 INTRODUCTION

This report presents the results of an evaluation of remediation alternatives and provides a preliminary corrective/remedial action plan for the former Grimit Auto and Repair site, located at 1970 Seminary Avenue, Oakland, California. The project location is shown on the Location Map, Figure 1, and the site is shown on the Site Plan, Figure 2. The report was required by the Alameda County Health Care Services Agency, Local Oversight Program, specifically a letter from Mr. Dale Klettke, Hazardous Materials Specialist, to the property owner, Doyle Grimit, dated September 24, 1996.

This investigation supplements a preliminary evaluation of remedial alternatives, prepared by our firm in July, 1996. The preliminary evaluation was conducted in response to a request by Alameda County Health Care Services Agency, dated May 15, 1996. Mr. Klettke's letter requested "a report which evaluates whether remedial action, interim remedial action, or further tier evaluation is warranted for your site".

In the current evaluation, a field test of the effectiveness of soil vapor/ground water extraction was conducted, and a recommendation for interim remedial action was prepared.

A scope of investigation was presented in our proposal dated November 5, 1996. The proposed cost for this evaluation was pre-approved for reimbursement by the State of California Underground Storage Tank Cleanup Fund Program in a letter dated November 21, 1996, and received by our firm on December 14, 1996.

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 2

2.0 BACKGROUND

2.1 Introduction

The project site is located at 1970 Seminary Avenue, at the southern corner of the Seminary Avenue - Harmon Avenue intersection, in Oakland, Alameda County, California. The immediate site vicinity is primarily residential. The site is currently utilized as an automotive repair facility. The property is owned by Mr. Doyle Grimit, and is leased to the repair facility.

The site is approximately 50 by 100 feet in plan dimension. Three former gasoline and one former waste oil tank were removed in 1989. Fuel has not been dispensed since that time. One inactive hydraulic lift remains at the the site within the service building.

A detailed background description is included in our April 22, 1996 report.

2.2 Investigation History

Three exploratory borings and one monitoring well (MW-1) were installed by Kaldveer Associates in August, 1990 (report dated September 28, 1990). The well was sampled once by Kaldveer. Limited soil excavation was subsequently conducted at the location of the former waste oil tank. Hoexter Consulting subsequently sampled the well three times. In January and February, 1994, Hoexter Consulting conducted further subsurface investigation, including installation of two additional wells. Additional monitoring was followed by a supplemental investigation conducted in March, 1996, which included four soil borings and three additional monitoring wells. The resulting April 22, 1996 report included a preliminary ASTM RBCA Tier One evaluation of the data.

The referenced May 15, 1996 Alameda County letter followed and commented upon the April, 1996 subsurface investigation report. More recently, a preliminary evaluation of remedial action alternatives was conducted, and a report issued July 28, 1996. The evaluation report recommended supplemental ground water contaminant plume definition and further soil source delineation, followed by preparation of a remedial action feasibility study, development of a corrective action plan, and initiation of soil / ground water remediation. Finally, two additional quarterly ground water sampling events occurred, reported on October 21, 1996 and January 28, 1997.

2.3 Subsurface Conditions and Contaminant Source

The subsurface investigations indicated complex soil and ground water conditions consisting of interbedded discontinuous relatively thin lenses of silty and clayey sediments, with relatively limited deposits of "clean" sand or gravel. Based on the investigation, there are two separate ground water contamination zones, a "perched" or shallow zone ranging from 7 to 13 feet, and a deeper zone of from 20 to 30 feet. The two zones are probably interconnected. Based on well development and purging data, the strata yield relatively low volumes of water, and there is poor conductivity between strata. There are also two depth zones of soil contamination: "shallower" soils, to approximately 15 feet depth, and "deeper" soils, from 15 feet to the depth explored (35 feet). The "shallower" soils are generally more highly contaminated than the "deeper" soils.

Based on measured ground water levels, which are as shallow as 5-1/2 feet below the ground surface, it is likely that contaminants were discharged from the former gasoline and waste oil USTs and associated piping directly into saturated sediments. Although the USTs have been removed, and some overexcavation conducted in the former waste oil tank

cavity, contaminated soil remains in the "source area" of the former USTs and beneath the service building.

Previous Hoexter Consulting reports (particularly Hoexter Consulting April 22, 1996) discuss subsurface conditions in more detail, and include cross sections portraying our interpretation of the site stratigraphy.

2.4 Contaminant Description

Based on our investigations, contamination consists of oil and grease, gasoline (TPH-G), purgeable aromatic compounds (BTEX), MTBE, and halogenated volatile compounds (HVOC), particularly PCE, TCE, and DCE. These compounds have been detected in soil samples from various locations and in water samples from all six monitoring wells. The greatest soil and ground water contaminant levels have been observed in the general vicinity of the service building, particularly in the vicinity of well MW-1. The data are summarized in Appendix A of this report.

Ground water contaminant levels during the past two years (December, 1994 through January, 1997) have been as follows:

Maximum Water Column Values, 8/90 - 9/94 (see discussion below)

Compound

Values (**ppb***) (ground water)

Gasoline Oil & Grease

23,000,000 880,000

"Average" Water Column Values. 12/94 - present (see discussion below)

Compound

Range of Values (ppb*) (ground water)

Gasoline Oil & Grease

44,000 - 55,000 11,000 - 110,000

Note that more elevated levels (Appendix A, Table 3A) were initially observed (August, 1990 through September, 1994). This may have been related to the sampling method, which consisted of decanting the sample bailer from the top. This method probably incorporated "floating product", and is representative of the upper two feet of the saturated sediments. Subsequent samples (from December, 1994) were obtained from the approximate middle of the water column, and slowly drained from the bailer bottom using a constricted flow tube. These samples are more representative of an "average" value for the water column.

Typical and maximum levels of individual contaminants in the water have been as follows:

Compound	Typical/Max. Values (ppb*) (ground	water)
Benzene	4000/56.000	•
Toluene	4100/61.000	
Ethylbenzene	1600/28.000	
Xylenes	6800/137.000	
MTBE	490	1
PCE (Tetrachloroethene/perchlo	oroethene) 130	
TCE (Trichloroethene)	340	
VCL (Vinyl chloride)	60	
DCE (cis 1,2 Dichloroethene)	320	
DCA (1,2 Dichloroethane)	15	:

* Results in ug/l, equivalent to parts per billion, or ppb

Soil contaminant levels are generally relatively low, with the exception of oil & grease. TPH-G levels are on the order of 900 mg/kg (equivalent to parts per million, ppm), and generally less than 100 ppm. BTEX compounds are non-detect or depressed, with exceptions. Oil and grease levels are variable, with a maximum of 15,000 ppm detected in the former waste oil tank excavation. Maximum soil contaminant values are as follows:

<u>Compound</u>	Maximum Value (ppm*) (soil)
Gasoline	910
Oil\ & Grease	15,000

and the second s

Compound	Maximum Value (ppm*) (soil)
Benzene Toluene Ethylbenzene Xylenes	2.4 3.5 4.2 8.3
MIBE PCE (Tetrachloroethene/perchloroe TCE (Trichloroethene) VCL (Vinyl chloride)	thene) 1.8 0.82
DCE (CIS 1,2 Dichloroethene) DCA (1,2 Dichloroethane) DCB (1,2-Dichlorobenzene)	 1.7

* Results in mg/kg, equivalent to parts per million, or ppm

BTEX and individual HVOC levels exceed California MCLs. MTBE currently exceeds the proposed San Francisco Bay Region Water Quality Control Board standard of 35 ug/l in four of the six wells. The ASTM RBCA Tier 1 analysis (Section 4 and Appendix B) indicates that screening levels are exceeded for leaching potential of subsurface soils to ground water, soil volatilization to the air, soil and ground water vapor intrusion to

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 5

buildings, and ground water ingestion (see discussion in Hoexter Consulting report dated July 28, 1996). The July 28, 1996 report concluded, based on the Tier 1 results and the contaminant levels observed at the site, that Tier 2 evaluation would not produce substantially different conclusions.

3.0 PREVIOUS REMEDIAL ACTION EVALUATION

The following options were considered in the July 28, 1996 preliminary remedial action evaluation:

- 1 No Remedial Action
 - 1a Natural Attenuation (no further work).
 - 1b Natural Attenuation (plume definition and quarterly monitoring).
 - 1c ASTM Tier 2 RBCA Evaluation (including plume definition and quarterly monitoring).
- 2 Interim Remediation (additional source delineation and removal).
- 3 Ground Water Extraction.
- 4 Vapor / Ground Water Co-Extraction.
- 5 Vapor / Ground Water Co-Extraction with Air Sparging or ORC.

The July 28, 1996 evaluation concluded that :

- 1a,b The natural attenuation alternatives do not meet current Alameda County remedial criteria.
- 1c Tier 2 evaluation would not be likely to result in acceptable levels of residual contamination in the ground water.
- 2 Interim remediation, while beneficial, would not be sufficient due to constraints imposed by the existing building and adjacent property line.
- 3 Due to complex stratigraphy and hydrogeology, and relatively low permeability materials, ground water extraction would not be cost effective and is not a current remedial solution.
- 4 Vapor / ground water co-extraction could be beneficial for timely remediation.
- 5 Vapor / ground water co-extraction with air sparging or ORC is recommended.

Detailed discussions of these alternatives are included in the July 28, 1996 study. The reader is referred to the original document for the detailed evaluations. The study recommended supplemental ground water contaminant plume definition and further soil source delineation, followed by preparation of a remedial action feasibility study, development of a corrective action plan, and initiation of soil / ground water remediation.

The recommended plume definition, to be primarily located off-site, has been postponed. Further soil source delineation, although beneficial, would result in additional delays, and due to the complex nature of the site, might not provide cost-effective information. A dual soil vapor ground water extraction feasibility study was conducted by a specialty soil vapor extraction (SVE) contractor/consultant. The results have been incorporated into this evaluation. Following approval of the recommendations contained in this report, a detailed corrective action plan will be prepared, and soil/ground water remediation will commence.

The RBCA evaluations conducted for the site indicate that there are elevated health risks for both commercial and residential land uses. Based on the July 28, 1996 study and requirements of the Alameda County Health Department, it was concluded that site remediation be conducted. Based on these findings, Alameda County has required preparation of this document. The following sections discuss remedial objectives, considerations, and the proposed interim remedial action plan.

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 8

4.0 RISK EVALUATION

The exposure pathways included for human health risks for the project site include:

- * Dermal contact/ingestion of soil.
- * Soil leaching potential to ground water.
- * Soil gas volatilization to indoor/outdoor air.
- * Gas volatilization from water to indoor/outdoor air.
- * Ground water ingestion.

Each of these pathways were considered for both commercial and residential land use.

In 1994, the American Society for Testing and Materials (ASTM) issued a **risk based** guidance document for evaluation of the need for **corrective action** ("RBCA") applied primarily to petroleum release sites. The RBCA study (Appendix B) indicated that risk-based screening levels (RBSLs) were exceeded by up to four orders of magnitude. Based on the Alameda County recommended (Alameda County, May 15, 1996 letter) cancer risk of 10⁻⁴ for residential areas, the maximum allowable concentrations of benzene (**ppm**), the most critical compound (corrected by the Regional Water Quality Control Board factor of 0.29), would be as follows:

Exposure Pathway -Residential	<u>Soil</u> *	Ground Water *
* Dermal contact/ingestion of soil.	60.9	
* Soil leaching potential to ground water.	1.36	· · ·
* Soil gas volatilization to indoor air.	0.04	an a
* Soil gas volatilization to outdoor air.	34.8	·
* Gas volatilization from water to indoor air.		0.28
* Gas volatilization from water to outdoor air.		174
* Ground water ingestion.		0.08
Exposure Pathway -Commercial	Soil *	Ground Water *
 Exposure Pathway -Commercial * Dermal contact/ingestion of soil. 	<u>Soil</u> *	Ground Water *
 Exposure Pathway -Commercial * Dermal contact/ingestion of soil. * Soil leaching potential to ground water. 	<u>Soil</u> * >res 4.64	Ground Water *
 Exposure Pathway -Commercial * Dermal contact/ingestion of soil. * Soil leaching potential to ground water. * Soil gas volatilization to indoor air. 	<u>Soil</u> * >res 4.64 0.09	<u>Ground Water</u> *
 Exposure Pathway -Commercial * Dermal contact/ingestion of soil. * Soil leaching potential to ground water. * Soil gas volatilization to indoor air. * Soil gas volatilization to outdoor air. 	<u>Soil</u> * >res 4.64 0.09 37.7	<u>Ground Water</u> *
 Exposure Pathway -Commercial * Dermal contact/ingestion of soil. * Soil leaching potential to ground water. * Soil gas volatilization to indoor air. * Soil gas volatilization to outdoor air. * Gas volatilization from water to indoor air. 	<u>Soil</u> * >res 4.64 0.09 37.7	<u>Ground Water</u> * 0.72
 Exposure Pathway -Commercial * Dermal contact/ingestion of soil. * Soil leaching potential to ground water. * Soil gas volatilization to indoor air. * Soil gas volatilization to outdoor air. * Gas volatilization from water to indoor air. * Gas volatilization from water to outdoor air. 	<u>Soil</u> * >res 4.64 0.09 37.7 	<u>Ground Water</u> * 0.72 243.6

* Results in mg/kg or mg/l, equivalent to parts per million, or ppm

5.0 REMEDIAL OBJECTIVES

The general ground water clean up objective used by the State of California Regional Water Quality Control Board and Alameda County Health Department has been the Federal and California drinking water standards. It is our opinion that these clean up objectives are not justified for the contaminants of concern for the project site. Use of the ASTM-RBCA guideline values for soil and ground water (Section 4.0) represent appropriate clean up objectives.

Most of the contamination appears to originate from the "source area" (former USTs and service building area), particularly the "smear zone' within the highly variable ground water table fluctuation zone. Complete delineation of the ground water contaminant plume emanating from the "source area" has not been conducted, as the plume extends off-site under adjacent residences. It is our opinion that remediation of the off-site contamination is not economically feasible or practical, due to the presence of residences and related improvements on the adjoining properties. In our opinion, reduction of the "source area" contamination within the site is feasible, and the rational alternative to extensive and impractical off-site remediation.

Thus, the principal objective of the proposed remediation will be soil and ground water remediation within the on-site "source area". It is not the objective of the remedial action to achieve non-detectable concentrations of all petroleum or HVOC compounds in the soil or in the ground water; however, it is the objective to further abate the continued leaching of these compounds from the "smear" zone and to reduce the concentrations of these compounds in the ground water to below the ASTM-RBCA Tier 1 evaluation concentration values.

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 10

6.0 REMEDIAL SYSTEM DESIGN CONSIDERATIONS

6.1 Site Characteristics

The site is underlain by a complex series of relatively thinly lensed sediments of limited lateral extent. Silty and clayey deposits predominate, with minor deposits of "clean" sand or gravel. The overall permeability of the sediments is low. For example, the wells require as much as 48 hours to equilibrate when initially uncapped, and have dewatered with removal of as little as three (3) well volumes when purged. Due to the stratigraphic complexity, ground water and air flow (within unsaturated sediments) are most likely irregular.

Ground water levels and gradient flow direction also vary. The two "shifts water water and the source of the sourc

6.2 Soil Vapor Extraction (SVE) Pilot Study (January, 1997)

Terra Vac Corporation, of San Leandro, California, was retained to perform a **deuterepresentation** pilot study of the site. Terra Vac's February 5, 1997 report is included in this report as Appendix C. The Terra Vac study was conducted to assess the feasibility of vapor extraction at the site.

The study consisted of utilizing the existing monitoring well MW-1 as the commensate MW-1 is located near the "source" area, and exhibits the highest levels of contamination of the six on-site wells. A 35 foot long "slurp tube" was extended into the well, to extract ground water simultaneously with vapor.

The test's radius of influence was monitored with two observation points, consisting of driven screened casing located approximately 14 and 25 feet from the extraction well. Approximately 11 cubic feet of air per minute were extracted from the well with an applied vacuum of 12 inches of mercury column. The test ran for approximately three hours. Vacuums of 0.2 and 0.1 inches of water column were observed in observation wells 1 and 2, respectively, at the conclusion of the test.

Total petroleum hydrocarbons were monitored twice during the test. The initial vapor sample, obtained after approximately 20 minutes, indicated a concentration of 39.7 ppm TPH. This decreased to 12.6 ppm TPH at 170 minutes.

The test was conducted at a time of relatively elevated ground water elevation. A total of 130 gallons of water was extracted during the test, corresponding to an average extraction rate of approximately 0.7 gallons per minute.

6.3 Evaluation of SVE Effectiveness

Terra Vac (February 5, 1997) concludes that "The amount of vacuum observed in OB-1 is significant and is indicative of some degree of connectivity between MW-1 and OB-1. There appeared to be some connectivity between MW-1 and OB-2, however the amount of induced vacuum was not as significant." A reduced of the degree of the amount of the pressure during the three hour test can be inferred. However, it is likely that much of the pressure drop in the observation wells was due to a ground water level decline in the wells. Thus,

the degree of airflow between points, as represented by the measured 0.2 and 0.1 inches of water column decline in the observations wells at the three hour conclusion of the test, does not appear to be significant.

In our opinion, SVE could be effective for remediation of gasoline/BTEX and HVOCs at the site, but would require a significant capital investment due to the required density of extraction wells and associated equipment operation and maintenance cost. In addition, SVE, while effective at the site, would not remediate the down-gradient portions of the contaminant plume. Finally, SVE, while effective for the gasoline compounds, would be less effective for the oil and grease and HVOCs incorporated into the waste oil present at the site. Although SVE or of the field induce oxygenation of the soil, and thus would increase natural, passive biodegradation processes of the HVOC and waste oil, the process is anticipated to be show and of minimal overall value.

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 12

7.0 PROPOSED PLAN

7.1 Introduction and Rationale

It is our opinion that SVE and/or dual ground water/soil vapor extraction, although ultimately effective, would not be economically feasible due to the relatively limited area of influence, and thus the need for densely placed vapor extraction points and long-term operating costs. In addition, SVE would be only moderately successful in reducing the concentrations of oil and grease and associated HVOC.

Oxygenation of subsurface soils and ground water have been proven technologies for promoting volatilization and degradation of petroleum hydrocarbon compounds and providing effective site remediation. Based on the subsurface soil and ground water conditions, it is our opinion that application of direct oxygenation of the "shallow" (5 to 20 feet) and the deeper (20 to 35 feet) water bearing zones would result in direct beneficial remediation of the gasoline as well as oil and grease and HVOC compounds present in these zones. This would also provide a positive barrier for potential migration of these contaminants and further promote the bacterial degradation. Oxygen releasing compounds (ORCs), manufactured by Regenesis Bioremediation Products and composed of magnesium peroxide, are recommended for this project application. The principal benefits of this product are the oxygen release combined with the non-hazardous nature of the resulting (oxygen-depleted) compound (magnesium hydroxide).

A passive system of oxygenation, consisting of installation of oxygen releasing compounds (ORCs) to the soil and ground water in conjunction with in-situ bioventing is determined to be the most efficient and cost effective method of source reduction and site remediation.

7.2 General Remediation Plan

The proposed remediation emphasis cleanup of the source area in the vicinity of the former USTs and service area. Complete site cleanup would not be immediately achieved. However, with the mitigation of the "source area", further transport of contaminants from the site would be minimized, and naturally occurring processes would then complete the remediation.

The proposed remediation plan would consist of:

- * Installation of additional down gradient monitoring wells."
- Installation of oxygen releasing compounds (ORCs) in soil borings throughout the "source area". An initial small scale calibration test is proposed. The ORCs would be installed as a slurry directly into the calibration test soil borings, and would be placed into pre-drilled borings or pressure injected into the soils in the "full scale" remediation program.
- * Installation of a low volume in-situ bioventing vacuum system in existing and proposed monitoring wells.

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Necessar

As discussed in Section 6.3, SVE would be relatively ineffective in remediating waste oil and the accompanying HVOC compounds. The proposed ORC bioremediation system would, in our opinion, be more effective in remediating the waste oil and HVOC, by directly providing oxygen for the microbial cultures to degrade these more complex hydrocarbons.

The following ORC procedures have been developed with the assistance of Mr. Jack Peabody, manager of Regenesis Bioremediation Products Pleasant Hill, California office. Procedures and volumes of material are recommended by Mr. Peabody based on Regenesis' experience and proprietary software.

7.2.1 Small Scale ORC Calibration Test

A small scale calibration test would be initially conducted. This test would consist of installing four soil borings approximately three (3) feet each from the existing monitoring well MW-1 (Figure 2). The test would provide information which would be used for final design of the larger-scale remediation. In addition, it would provide "immediate" remediation of the most heavily impacted area of the site.

Two-inch diameter borings would be installed by direct push methods, thus eliminating the need to treat or dispose of contaminated soil. A 65 per cent ORC slurry would be tremied to the borings, which would be capped with five (5) feet of neat bentonite cement at the ground surface. Each boring would receive approximately 43 pounds of the ORC compound.

Effectiveness of the test would be monitored with monthly dissolved oxygen (DO) measurements of MW-1, and quarterly sampling of contaminants. Details of the monitoring are provided in Section 10.0 of this report. Note that a decline in TPH and BTEX would be anticipated prior to an increase of DO in the well. This is due to utilization of all available oxygen by the in-situ bacteria during the initial stages of remediation, particularly at highly contaminated areas.

7.2.2 Additional Monitoring Wells

Three additional monitoring wells are proposed. The proposed well locations are shown on Figure 2. The wells are required for plume definition and to monitor remediation progress. One 35 foot "deep" well and one 20 foot "shallow" well would be located on the down-gradient side of the source area along the property line; one 20 foot "shallow" well would be located on the site up-gradient area adjacent to Seminary Avenue. Well reference elevations would be surveyed, and groundwater data from the wells incorporated into evaluations of ground water gradient flow direction.

7.2.3 Full Scale ORC Remediation

The results of the calibration test will be used to complete a final design of the ORC remediation system. The current plan is to install a grid of six to nine additional borings approximately spaced at 8 to 10 foot centers within the "source area" (Figure 2). Additional, more widely spread borings, would be placed within the balance of the site, where contamination levels are less elevated. The final plan may slightly increase the number of borings and decrease the spacing. The ORC slurry would be placed in the same manner as described in Section 7.2.1, or pressure grouted, whichever is less costly.

As discussed above, the purpose of the proposed full scale remediation would be to reduce concentrations of contaminants in the "source area". Peripheral and off-site areas would not be directly addressed; natural biodegradation processes would be allowed to work in these areas.

7.2.4 In-Situ Bioventing

The in-situ bioventing would consist of installing a low-volume vacuum system (one to three inches of mercury) applied to the two existing and the two proposed additional "source area" wells. The system would provide positive control of off-gassing to reduce off-site migration of vapors and help promote air flow through the soil column and "smear" zone", resulting in more efficient site remediation. An activated carbon filtration system would be employed to treat produced vapors.

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 15

8.0 ADDITIONAL INVESTIGATION

Off site definition of the plume, as previously proposed, is in our opinion, still warranted. This investigation would consist of obtaining representative grab ground water samples from public rights of way along adjacent and nearby streets (Harmon Avenue, Seminary Avenue, and Holway Street.

9.0 SCHEDULE/DURATION OF TREATMENT

The proposed "hot spot" pilot test will be initiated following approval of costs by the State UST Fund. Approval by the Fund will require approximately two (2) to four (4) weeks. Permitting, ordering of materials, and installation are anticipated to require approximately three weeks. The pilot test will require an estimated six (6) to nine (9) months of monitoring. A progress reporting will be included with each regularly scheduled "Quarterly" ground water monitoring.

A brief work plan for the installation of the proposed monitoring wells will be prepared following acceptance of this corrective/interim remedial action plan. The wells will be installed in conjunction with the "hot spot" pilot test, pending State UST Fund approval of costs.

10.0 ANALYTICAL TESTING AND REPORTING

Analytical testing of the six current and three recommended wells (including the "hot spot" pilot test well, MW-1) will consist of a continuation of the currently scheduled "Quarterly" monitoring, which includes TPH-G/BTEX/MTBE, oil/grease, halogenated volatile organics (HVOC) (alternate sampling events); and dissolved oxygen (DO), nitrate, sulfate, also $\ell \wedge A$ is and ferrous iron.

In addition, well MW-1 will be monitored monthly for dissolved oxygen (DO) and TPH-G/BTEX/MTBE. The MW-1 sample will be obtained as a grab water sample from the top of the water column, as opposed to purging, which could disrupt the ORC oxygenation process.

A sampling plan for the period following installation of the full-scale ORC and bioventing remediation will be included in the final remedial action plan.

1970 Seminary, Oakland, CA; E-10-1B-192B; February 15, 1997; Page 18

11.0 FINAL REMEDIAL ACTION PLAN

A final remedial action plan (RAP) will be prepared following completion of the pilot test. The RAP will include a detailed plan for installation of the ORC and bioventing systems.

12.0 LIMITATIONS

This report has been prepared according to generally accepted geologic and environmental practices. No other warranty, either expressed or implied as to the methods, results, conclusions or professional advice provided is made. It should be recognized that certain limitations are inherent in the evaluation of subsurface conditions, and that certain conditions may not be detected during an investigation of this type. If you wish to reduce the level of uncertainty associated with this study, we should be contacted for additional consultation.

The analysis, conclusions and recommendations contained in this report are based on site conditions as they existed at the time of our investigation; review of previous reports relevant to the site conditions; and laboratory results from an outside analytical laboratory. Changes in the information or data gained from any of these sources could result in changes in our conclusions or recommendations. If such changes do occur, we should be advised so that we can review our report in light of those changes.

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APPENDIX A

Analytical Data Summary Tables

TABLE 1

GROUND WATER ELEVATION DATA

(All Measurements in Feet)

1

Well Number and Date of Measurement	Reference Elevation (2)	Depth to Water	Relative Ground Water Elevation (2)	
MW-1 ("deep")				
8/6/90	37.0	21.5	15.5	
1/28/92		21.0	16.0	
4/27/92		20.95	16.05	
8/10/92		22.20	14.8	
2/11/94		15.93 (3)	21.07 (3)	
2/28/94		13.85 (4)	23.15 (4)	
9/9/94		20.19	16.81	
12/28/94		14.91	22.09	
4/13/95		14.18	22.82	
11/1/95		20.90	16.10	
3/8/96		11.82	25 18	
3/25-26/96 10/7/96 1/15/97	36.97	13.54 21.41 13.34	23.18 23.43 15.59 23.63	
MW-2 ("deep")				
2/11/94	36.40	14.16 (3)	22.24 (3)	
2/28/94		16.01 (4)	20.39 (4)	
9/9/94		18.96	17.44	
12/28/94		21.42	14.98	
4/13/95		19.69	16.71	
11/1/95		21.91	14.49	
3/8/96		14.56 (6)	21 84 (6)	
3/25-26/96	36.39	10.84	25.55	
10/7/96		18.41	17.98	
1/15/97		10.07	26.32	
MW-3 ("shallow")				
2/11/94	36.94	6.97 (3)	29.97 (3)	
2/28/94		7.74 (4)	29.20 (4)	
9/9/94		9.68	27.26	
12/28/94		8.15	28.79	
4/13/95		8.05	28.89	
11/1/95		7.82	29.12	
3/8/96		5.69	31.25	
3/25-26/96	36.94	6.91	30.03	
10/7/96		9.51	27.43	
1/15/97		6.23	30.71	

Table 1 continued

Well Number and Date of Measurement	Reference Elevation (2)	Depth to Water	Relative Ground Water Elevation (2)
MW-4 ("deep")			
3/25-26/96 10/7/96 1/15/97	36.46	14.14 22.31 13.78	22.32 14.15 22.68
MW-5 ("deep")			
3/25-26/96 10/7/96 1/15/97	36.77	15.63 22.86 17.33	21.14 13.91 19.44
MW-6 ("shallow")			
3/25-26/96 10/7/96 1/15/97	36.42	8.52 12.82 7.72	27.90 23.60 28.70

Notes

- (1) N/A = not applicable.
- (2) Elevations from a survey conducted by Andreas Deak, California Licensed Land Surveyor, March 21, 1996, City of Oakland datum.
- (3) Well under pressure when locking cap removed; water level may not have been stabilized.

(4) Depth to water was measured over a 120 minute period; indicated depths appear to be stabilized readings.

(5) Surveyed elevations of wells MW 1 and MW-2 varied to 0.02 foot on March 21, 1996 survey as compared to February 11, 1994 survey; previously calculated measurements of elevation have **not** been modified to reflect the new survey data.

(6) Well not stabilized (water level rising).

TABLE 2

SOIL

SUMMARY OF ANALYTICAL TEST RESULTS -PETROLEUM HYDROCARBONS

(Results reported in parts per million, mg/kg) (1) (2)

Sample	TPH- Gasoline	e Benzene	Toluene	Ethyl- Benzene	Xylenes	Oil and Grease	HVOC
Initial US	ST Remov	al Confirm	ation Testi	ing			
Gasoline	USTs						
South tank South tank Center tank North tank	22 ND 20 ND 21	ND ND ND 0.068 2.4	ND ND 0.031 ND 2.9	ND ND ND 0.320	ND ND 0.200 ND 1.7	NA NA NA NA	NA NA NA NA
Waste Oil	UST						
1 2	NA NA	0.093 0.160	0.510 0.400	0.480 0.810	1.7 2.4	5500/7 7200/4	60 (6) ND 60 (6) ND
Previous	Kaldveer	Investigati	ion				
EB-1							
16.0 21.0 26.0	4 0.5 50	NA NA NA	NA NA NA	NA NA NA	NA NA NA	NA NA NA	NA NA NA
EB-2							
10.0 16.0	NA NA	NA NA	NA NA	NA NA	NA NA	4,200 ND	NA NA
EB-3							
10.0 16.0	NA NA	NA NA	NA NA	NA NA	NA NA	2,800 150	NA NA

Waste Oil Tank Overexcavation Confirmation Testing	
--	--

1 (south side)	190	ND	ND	0.58	1.3	15,000/270	0 NA
2 (west side)	ND	ND	ND	ND	ND	9,800	NA
3 (east side)	4.4	ND	ND	0.0083	0.021	890 11,000/440	0 NA
4 (north side)	12	0.0042	ND	0.0091	0.021	7,500 410/250	NA
5 (west floor)	270	ND	3.5	1.3	ND	230 5,500/670	NA
6 (east floor)	260	ND	ND	1.2	2.5	3,700	NA
Stockpile	11	0.0031	ND	0.044	0.094 1,000	2,200 1,500/710	
Initial Hoex	ter Inves	tigation					
MW-2							
10.5-11.0 16.0-16.5 20.5 21.0	910 ND	ND ND	0.76 0.022	4.2 ND	6.1 ND	38 ND	NA NA
25.5-26.0 (3)	ND	ND	ND	ND	ND	ND	NA
MW-3							
10.5-11.0 20.5-21.0	ND 1.2	ND 0.17	0.020 0.047	ND ND	ND 0.085	ND NA	NA NA
April, 1996	Hoexter	Investigati	on				
EB-4							
7.5-8.0 14.5-15.0	300 63	ND ND	ND ND	3.3 ND	8.3 0.82	820 3600	ND Det (5)
EB-5							
3.5-4.0 7.5-8.0 12.5-13.0 18.0-18.5	ND 130 120	ND ND ND	ND ND ND	ND 0.55 0.84	ND 1.3 1.4	NA NA NA	NA NA NA
19.5-20.0 (3)	4.5	0.025	0.015	0.028	0.078	240	Det (5)
EB-7							
9.0-9.5 14.0-14.5 20.0-20.5	ND ND	ND ND	ND ND	ND ND	ND ND	ND NA	NA NA
23.0-23.5 (3)	130	ND	0.38	1.9	2.9	620	ND

MW-4

16.0-16.5	13	0.038	0.015	ND	0.023	NA	NA
31.0-31.5 (3) 36.0-36.5	68 5.4	0.21 ND	0.092 0.008	0.15 0.015	0.39 0.011	190 NA	NA NA
MW-5							
11.0-11.5 21.0-21.5 21.0-21.5	9.7 ND	ND ND	0.019 ND	ND ND	0.038 ND	NA NA	NA NA
35.5-36.0 (3)	NA	NA	NA	NA	NA	ND	NA
MW-6							
11.0-11.5 16.0-16.5 (3)	10	0.037	0.033	0.18	0.46	ND	NA

Notes

(1) NE) = non	-detect
--------	---------	---------

- (2) NA = not applicable(3) Composite
- (4) Chromatogram patterns/comments G - gas WG - weathered gas NGM - non-gas mix, > C9 NDM - non-diesel mix, generally C7 - C12/13 (5) Detected: see Table 2B(6) TOG/Motor Oil

Hoexter Consulting, Inc. 734 Torreya Court, Palo Alto, California 94303 (415) 494-2505

.

TABLE 2B

SOIL

SUMMARY OF ANALYTICAL TEST RESULTS -HALOGENATED VOLATILE ORGANIC COMPOUNDS

(Results reported in parts per million, mg/kg) (1) (2)

Sample	CA	1,2 DCB	1,2 DCA	cis 1,2 DCE	trns 1,2 DCE	1,2 DCP	PCE	TCE	VCL
EB-4									
7.5-8.0 14.5-15.0	ND ND	ND 1.7	ND ND	ND ND	ND ND	ND ND	ND 1.8	ND 0.82	ND ND
EB-5									
18.0-18.5 19.5-20.0 (3)	ND	ND	ND	ND	ND	ND	0.52	ND	ND
EB-7									
20.0-20.5 23.0-23.5 (3)	ND	ND	ND	ND	ND	ND	ND	ND	ND

Notes on following page

Table 2B Notes

- ND = non-detect
 NA = not applicable
 Composite
 Abbreviations as follows:

Chloroethane
1,2 Dichlorobenzene
1,2 Dichloroethane
cis 1,2 Dichloroethene
trans 1,2 Dichloroethene
1,2 Dichloropropane
Tetrachloroethene (perchloroethene)
Trichloroethene
Vinyl chloride

.

TABLE 3A

GROUND WATER

SUMMARY OF ANALYTICAL TEST RESULTS -PETROLEUM HYDROCARBONS (8)

(Results reported in parts per *billion*, ug/l) (1)

Well and Date	TPH Gasoline (8)	Benzene	Toluene	Ethyl- benzene	Xylenes	Oil Gre HVO(& ease C (7)
MW-1	("deep")						
8/6/90 (2) 1/28/92 4/27/92 (3) 4/27/92 (4) 8/10/92 2/11/94 1 9/9/94 23 12/28/94 4/13/95 11/1/95 3/25/96 10/8/96 1/16/97	54,000 2,000,000 500,000 175,000 170,000 ,800,000 ,000,000 55,000 45,000 45,000 45,000 55,000 44,000 45,000 55,000 48,000	3,500 7,400 3,400 4,200 4,200 56,000 3,700 2,800 2,600 3,000 3,300 2,600	3,200 17,000 6,400 4,400 4,200 5,100 61,000 5,300 3,400 3,400 4,100 4,500 3,200	$\begin{array}{c} 1,900\\ 28,000\\ 10,000\\ 3,200\\ 3,300\\ 5,200\\ 9,100\\ 1,400\\ 1,200\\ 1,400\\ 1,600\\ 1,700\\ 1,300 \end{array}$	9,400 120,000 45,000 14,600 15,900 23,900 137,000 5,800 5,100 5,900 6,800 7,100 5,300	7,600 75,000 440,00 N/ 120,000 16,000 880,000 880,000 83,000 52,000 46,000 11,000	0 (5) 00 (6) A (6) (6) (6) (5) (5) (5) (7) (5) (7) (5) (7)
MW-2	("deep")						. ,
2/11/94 9/9/94 12/28/94 4/13/95 11/1/95 3/25/96 10/8/96 1/16/97 MW-3 ("s	130 1,000 330 1300 100 4500 710 330 hallow")	22 89 100 280 9.9 470 1.9 41	1.1 ND 3.8 6.9 ND 57 0.54 2.4	5.2 ND 5.4 33 ND 220 1.0 1.3	7.3 6.9 4.7 23 ND 280 1.0 9.9	ND 5100 ND ND ND ND ND ND ND	(6) (6) 5) (5) (5) (7) (5) (7) (5)
2/11/94 9/9/94 12/28/94 4/13/95 11/1/95 3/25/96 10/8/96 1/16/97	ND 710 2,300 1,700 1,100 2,300 160 1,800	ND 10 7.8 2.9 4.4 4.0 ND 2.8	ND ND ND ND 0.96 0.5 0.68	ND ND 130 61 27 120 1.2 48	ND 3.5 73 24 22 65 0.77 66	ND ND ND ND ND ND ND ND ND ND ND ND ND N	(6) (6) (5) (5) (5) (7) (5) (7) (5)

Table 3A continued

Well and Date	Vell and TPH Date Gasoline Benzen (8)		Ethyl- Toluene benzene Xylenes			Oil & Grease HVOC (7)		
MW-4	("deep")							
3 /26/ 96 10/8/96 1/16/97	9,900 7,800 4,800	4,000 3,900 1,900	40 33 21	71 31 2.5	100 40 27	ND ND 5,200	(5) (7) (5) (7) (5)	
MW-5	("deep")							
3/26/96 10/8/96 1/16/97	1,200 6,700 3,000	43 260 150	8.2 92 68	83 410 190	95 370 180	ND ND ND	(5) (7) (5) (7) (5)	
MW-6 ("shallow")							
3/26/96 10/8/96 1/15/97	9,900 1,300 6,500	1,000 120 570	150 2.3 65	470 1.4 170	720 4.0 630	ND ND ND	(5) (7) (5) (7) (5)	
EB-4								
3/8/96	15,000	780	840	1,300	590	7,500	(5) (7)	
MCL	NA	1	150	700	1750	N	A	

Notes

ND - non-detect; N/A - not applicable
 Kaldveer Associates report, September, 1990
 Sequoia Analytical Laboratory
 Applied Remediation Laboratory
 Gravimetric Method

(6) Infrared Method

(7) HVOC detected: see Table 2C(8) MTBE see Table 2B

TABLE 3B

GROUND WATER

SUMMARY OF ANALYTICAL TEST RESULTS -MTBE

(Results reported in parts per *billion*, ug/l)

Well and Date	MTBE
MW-1 ("deep") 10/8/96 1/16/96	490 310
MW-2 ("deep") 10/8/96 1/16/96	41 12
MW-3 ("shallow") 10/8/96 1/16/96	ND 7.1
MW-4 ("deep") 10/8/96 1/16/96	140 84
MW-5 ("deep") 10/8/96 1/16/96	190 90
MW-6 ("shallow") 10/8/96 1/16/96	57 220

TABLE 3C

GROUND WATER

SUMMARY OF ANALYTICAL TEST RESULTS -HALOGENATED VOLATILE ORGANIC COMPOUNDS

(Results reported in parts per <u>billion</u>, ug/l) (1) (2)

Well and Date	CA	1,2	DCB	1,2	DCA	cis	1,2 DCE	trns 1,2 DCE	1,2 DCP	PCE	TCE	VCL
MW-1 ("deep")												
3/25/96	ND<5		7.2		5.3		82	ND<5	ND<5	ND<5	7.8	25
10/8/96	ND<20		ND<20		ND<	20	45	ND<20	ND<20	ND<20	ND<20	26
1/16/97	NA		NA		NA		NA	NA	NA	NA	NA	NA
MW-2 ("deep")												
3/25/96	ND<0.5		ND<0.5		8.7		11	ND<0.5	1.0	ND<0.5	3.2	0.92
10/8/96	ND<0.5		ND<0.5		15		9.6	ND<0.5	1.1	ND<0.5	6.6	ND<0.5
1/16/97	NA		NA		NA		NA	NA	NA	NA	NA	NA
MW-3 ("shallow	v")											
3/25/96	ND<0.5		ND<0.5		0.56		1.2	ND<0.5	ND<0.5	ND<0.5	ND<0.5	ND<0.5
10/8/96	ND<0.5		ND<0.5		1.1		0.87	ND<0.5	ND<0.5	ND<0.5	ND<0.5	ND<0.5
1/16/97	NA		NA		NA		NA	NA	NA	NA	NA	NA
MW-4 ("deep")												
3/26/96	ND<8		22		ND<	8	300	9.2	ND<8	38	150	44
10/8/96	ND<15		22		4.9		320	ND<15	ND<15	52	130	60
1/16/97	NA		NA		NA		NA	NA	NA	NA	NA	NA
MW-5 ("deep")												
3/26/96	1.4		ND<0.5	5	2.1		6.2	ND<0.5	ND<0.5	ND<0.5	ND<0.5	10
10/8/96	ND<2.5		ND<2.5		4.9		4.4	ND<2.5	ND<2.5	ND<2.5	ND<2.5	9.4
1/16/97	NA		NA		NA		NA	NA	NA	NA	NA	NA
, .,												

Continued following page

Table 3C continued

MW-6 ("shalle 3/26/96 10/8/96 1/16/97	ow") ND<0.5 ND<0.5 NA	ND<0.5 ND<0.5 NA	3.9 2.3 NA	15 9.9 NA	ND<0.5 ND<0.5 NA	1.9 ND<0.5 NA	0.77 ND<0.5 NA	2 0.57 NA	ND<0.5 ND<0.5 NA
EB-4 3/8/96 (grab)	ND	ND	ND	42	ND	ND	130	340	ND
MCL	NA	600	0.5	6	10	5	7	5	0.5

Notes

ND = non-detect
 NA = not applicable
 Composite
 Abbreviations as follows:

CA	Chloroethane	1,2 DCP	1,2 Dichloropropane
1,2 DCB	1,2 Dichlorobenzene	PCE	Tetrachloroethene (perchloroethene)
1,2 DCA	1,2 Dichloroethane	TCE	trichloroethene
cis 1,2 DCE	cis 1,2 Dichloroethene	VCL	vinyl chloride
trans 1,2 DCE	trans 1,2 Dichloroethene	VCL	vinyreinorae

TABLE 3D

GROUND WATER

SUMMARY OF ANALYTICAL TEST RESULTS -ADDITIONAL PARAMETERS

(Results reported in parts per *million*, mg/l) (1)

Well and Date	Dissolved Oxygen	Ferrous Iron	Nitrate	Sulfate
MW-1 ("deep") 10/8/96	1.5	ND	ND	ND
1/16/97	1.4	3.6	ND	ND
MW-2 ("deep")				
10/8/96	3.7	ND	3	25
1/16/97	5.4	0.28	3	25 25
MW-3 ("shallow")				
10/8/96	3.8	ND	ND	5
1/16/97	5.2	ND	ND	5
MW-4 ("deep")				
10/8/96	3.0	ND	ND	ND
1/16/97	4.7	0.75	ND	5
MW-5 ("deep")				
10/8/96	2.8	ND	ND	8
1/16/97	3.4	0.38	ND	9
MW-6 ("shallow")				
10/8/96	2.7	ND	ND	6
1/16/97	2.7	0.28	ND	8

Notes

(1) ND - non-detect; N/A - not applicable

APPENDIX B

RBCA Analysis Summary Tables

RBCA TIER 1/TIER 2 EVALUATION

Output Table 1

Site Name: 1970 Seminary Site Location: Oakland, C Job Identification: E-10-18-1928 Date Completed: 10/3/96 Completed By: David Hoexter Software: GSI RBCA Spreadsheet Version: v10

NOTE: values which differ from Tier 1 default values are shown in bold italics and underlined.

	DEFA	ULT PARA	METERS				
Exposure			Residential		Commercial/Industrial		
Parameter	Definition (Units)	Adult	(1-6yrs)	(1-16 yrs)	Chronic	Constrctn	
ATC	Averaging time for carcinogens (yr)	70					
ATri	Averaging time for non-carcinogens (yr)	30	6	16	25	1	
BW	Body Weight (kg)	70	15	35	70		
ED	Exposure Duration (yr)	30	6	16	25	1	
EF	Exposure Frequency (days/yr)	350			250	180	
EF.Derm	Exposure Frequency for dermal exposure	350			250		
lRgw	Ingestion Rate of Water (I/day)	2			1		
IRs	Ingestion Rate of Soil (mg/day)	100	200		50	100	
IRadj	Adjusted soil ing, rate (mg-yr/kg-d)	1.15+02			9.4E+01		
IRa.in	Inhalation rate indoor (m^3/day)	15			20		
iRa.out	Inhalation rate outdoor (m^3/day)	20			20	10	
SA	Skin surface area (dermal) (cm*2)	5.8E+03		2.0E+03	5.8E+03	5.8E+03	
SAadj	Adjusted dermal area (cm^2•yr/kg)	2.1E+03			1.7E+03		
M	Soil to Skin adherence factor	1					
AAFs	Age adjustment on soil ingestion	TRUE			TRUE		
AAFd	Age adjustment on skin surface area	TRUE			TRUE		
tox	Use EPA tox data for air (or PEL based)	TRUE					
gwMCL?	Use MCL as exposure limit in groundwater?	FALSE					

Matrix of Exposed Persons to		Residential	Commercial/Industrial		
Complete	Exposure Pathways		Chronic	Constrctn	
Groundwa	iter Pathways:				
GW.i	Groundwater Ingestion	TRUE	FALSE		
GW.v	Volatilization to Outdoor Air	TRUE	FALSE		
GW.b	Vapor Intrusion to Buildings	TRUE	FALSE		
Soit Pathy	vays				
S.v	Volatiles from Subsurface Soils	TRUE	FALSE		
SS.v	Volatiles and Particulate Inhalation	TRUE	FALSE	TRUE	
SS.d	Direct Ingestion and Dermal Contact	TRUE	FALSE	TRUE	
S.I	Leaching to Groundwater from all Soils	TRUE	FALSE		
SЪ	Intrusion to Buildings - Subsurface Soils	TRUE	FALSE		

Matrix of I	Receptor Distance	Resi	dential	Commercial/Industrial		
and Locat	tion on- or off-site	Distance	On-Site	Distance	On-Site	
GW	Groundwater receptor (cm)		TRUE		TRUE	
s	Inhalation receptor (cm)		TRUE		TRUE	
Matrix of						
Target Ris	sks	Individual	Cumulative			
TRab	Target Risk (class A&B carcinogens)	<u>1.0E-04</u>				
TRc	Target Risk (class C carcinogens)	1.0E-04				
THQ	Target Hazard Quotient	1.0E+00				
Opt	Calculation Option (1, 2, or 3)	1				

		Commercia	al/Industrial
Definition (Units)	Residential	Chronic	Construction
Exposure duration (yr)	30	25	1
Contaminated soil area (cm^2)	1.9E+06		1.0E+06
Length of affected soil parallel to wind (cm)	1.5E+03		1.0E+03
Length of affected soil parallel to groundwater (c	1.5E+03		
Ambient air velocity in mixing zone (cm/s)	2.3E+02		
Air mixing zone height (cm)	2.0E+02		
Definition of surficial soils (cm)	1.0E+02		
Particulate areal emission rate (g/cm^2/s)	2.2E-10		
Definition (Units)	Value		
Groundwater mixing zone depth (cm)	6.1E+02	-	
Groundwater infiltration rate (cm/yr)	1.56+01		
Groundwater Darcy velocity (cm/yr)	1.2E+03		
Groundwater Transport velocity (cm/yr)	6.6E+03		
Saturated Hydraulic Conductivity(cm/s)			
Groundwater Gradient (cm/cm)			
Width of groundwater source zone (cm)			
Depth of groundwater source zone (cm)			
Biodegradation Capacity (mg/L)			
Is Broattenuation Considered	FALSE		
Effective Porosity in Water-Bearing Unit	3.8E-01		
Fraction organic carbon in water-bearing unit	1 0E-03		
Definition (Units)	Value		
Capillary zone thickness (cm)	1.5E+01		
Vadose zone thickness (cm)	1.6E+02		
Soil density (g/cm^3)	1.7		
Fraction of organic carbon in vadose zone	0.01		
Soil porosity in vadose zone	0.38		
Depth to groundwater (cm)	<u>1.7E+02</u>		
Depth to top of affected soil (cm)	2.1E+02		
Thickness of affected subsurface soils (cm)	<u>8.8E+02</u>		
Soil/groundwater pH	65		
-	capillary	vadose	foundation
Volumetric water content	0.342	0.12	0.12
Volumetric air content	0.038	0.26	0.26
Definition (Units)	Residential	Commercial	
Definition (Units) Building volume/area ratio (cm)	Residential 2.0€+02	Commercial 3.0E+02	
Definition (Units) Building volume/area ratio (cm) Building air exchange rate (s^.1)	Residential 2.0E+02 1 4E-04	Commercial 3.0E+02 2.3E-04	
Definition (Units) Building volume/area ratio (cm) Building air exchange rate (s^.1) Foundation crack thickness (cm)	Residential 2.0E+02 1 4E-04 1.5E+01	Commercial 3.0E+02 2.3E-04	
	Exposure duration (yr) Contaminated soil area (cm^2) Length of affected soil parallel to wind (cm) Length of affected soil parallel to groundwater (c Ambient air velocity in mixing zone (cm/s) Air mixing zone height (cm) Definition of surficial soils (cm) Particulate areal emission rate (g/cm^2/s) Definition (Units) Groundwater mixing zone depth (cm) Groundwater infiltration rate (cm/yr) Groundwater Transport velocity (cm/yr) Groundwater Gradient (cm/cm) Width of groundwater source zone (cm) Biodegradation Capacity (mg/L) Is Bioattenuation Considered Effective Porosity in Water-Bearing Unit Fraction organic carbon in water-bearing unit Definition (Units) Capillary zone thickness (cm) Soil density (g/cm^3) Fraction of organic carbon in vadose zone Soil porosity in vadose zone Depth to po affected soil (cm) Thickness of affected subsurface soils (cm) Soil/groundwater pH Volumetric air content Volumetric air content	Exposure duration (yr)30Contaminated soil area (cm^2)1.9E+06Length of affected soil parallel to wind (cm)1.5E+02Length of affected soil parallel to groundwater (c1.5E+02Ambient air velocity in mixing zone (cm/s)2.3E+02Ari mixing zone height (cm)2.0E+02Definition of suficial soits (cm)1.0E+02Particulate areal emission rate (g/cm^2/s)2.2E+10Definition (Units)ValueGroundwater mixing zone depth (cm)6.1E+02Groundwater infiltration rate (cm/yr)1.2E+03Groundwater Transport velocity (cm/yr)6.6E+03Saturated Hydraulic Conductivity(cm/s)Groundwater source zone (cm)Definition (Units)ValueWidth of groundwater source zone (cm)Biodegradation Capacity (mg/L)Is Bioattenuation ConsideredFALSEEffective Porosity in Water-Bearing Unit3.8E-01Fraction organic carbon in water-bearing unit1.0E+02Soil density (g/cm^3)1.7Fraction of organic carbon in vadose zone0.01Soil porosity in vadose zone0.38Depth to top of affected soil (cm)1.1E+02Soil density (g/cm^3)1.7Fraction of organic carbon in vadose zone0.38Depth to top of affected subsurface soils (cm)8.8E+02Soil groundwater pH6.5Soil/groundwater pH6.5Soil/groundwater pH0.342Volumetric water content0.338	Exposure duration (yr)3025Contaminated soil area (cm^2)1.9E+06Length of affected soil parallel to wind (cm)1.5E+03Length of affected soil parallel to groundwater (c1.5E+02Ambient air velocity in mixing zone (cm/s)2.3E+02Air mixing zone height (cm)2.0E+02Definition of surticial soils (cm)1.0E+02Particulate areal emission rate (g/cm^2/s)2.2E+10Definition (Units)ValueGroundwater mixing zone depth (cm)6.1E+02Groundwater mixing zone depth (cm/yr)1.5E+02Groundwater Infiltration rate (cm/yr)1.5E+03Groundwater Transport velocity (cm/yr)6.6E+03Saturated Hydraulic Conductivity(cm/s)Groundwater Gradient (cm/cm)Width of groundwater source zone (cm)Detfinition (Units)ValueCapillary zone thickness (cm)1.5E+02Soil density (g/cm^3)1.7Fraction of organic carbon in water-bearing unit1.0E+02Soil density (g/cm^3)1.7Fraction of organic carbon in vadose zone0.01Soil porosity in vadose zone0.38Depth to top of affected soil (cm)2.1E+02Soil density (g/cm*3)1.7Fraction of organic carbon in vadose zone0.01Soil porosity in vadose zone0.38Depth to top of affected subsurface soils (cm)2.1E+02Depth to top of affected subsurface soils (cm)2.1E+02Soil groundwater

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dcz

Vertical dispersion coefficient (cm)

RBCA SITE ASSESSMENT

Representative

Tier 1 Worksheet 6.1

Site Name: 1970 Seminary

Site Location: Oakland, C

Completed By: David Hoexter Date Completed: 10/3/1995 Target Risk (Class A & B) 1.0E-4 Targel Risk (Class C) 1.0E-4

MCL exposure limit? D PEL exposure limit? Targel Hazard Quotient 1.0E+0

1 OF 1 Calculation Option: 1

Serial: g-265-vhx-686

SURFACE SOIL RBSL VALUES (< 3 FT BGS)

RBSL Results For Complete Exposure Pathways ("x" if Complete)

CONSTITUENTS OF CONCERN		Concentration	۱			Ingestion, Inhalation Construction RBSL					
			X Soil Leaching to Groundwater		X and De	ermal Contact	X Worker	Applicable RBS	Exceeded ?	Required CRF	
CAS No.	Name	(mg/kg)	Residential (on-site)	Commercial: (on-site)	Regulatory(MCL): (on-site)	Residential: (on-site)	Commercial: (on-site)	Commercial. (on-site)	(mg/kg)	" # " If yes	Only if "yes" left
83-32-9	Acenaphthene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	>Res		<1
120-12-7	Anthracene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	>Res		<1
71-43-2	Benzene	0.0E+0	4.7E+0	NA	NA	2.1E+2	NA	>Res	4.7E+0		<1
75-00-3	Chloroethane	0.0E+0	1.4E+2	NA	NA	>Res	NA	>Res	1.4E+2		<1
95-50-1	Dichlorobenzene (1,2) (-o)	0.0E+0	2.3E+3	NA	NA	2.2E+3	NA	>Res	2.2E+3		<1
106-46-7	Dichlorobenzene, (1,4) (-p)	0.0E+0	2.5E+2	NA	NA	2.5E+2	NA	>Res	2.5E+2		<1
75-34-3	Dichloroethane, 1,1-	0.0E+0	9.0E+1	NA	NA	2.6E+3	NA	3.7E+3	9.0E+1		<1
107-06-2	Dichloroethane, 1,2-	0.0E+0	2.0E+0	NA	NA	6.6E+1	NA	2.6E+3	2.0E+0	0	<1
156-59-2	Dichloroethene, cis-1,2-	0.0E+0	6.2E+0	NA	NA	2.6E+2	NA	,3.4E+2	6.2E+0		<1
156-60-5	Dichloroethene,1,2-trans-	0.0E+0	9.5E+0	NA	NA	>Res	NA	>Res	9.5E+0		<1
100-41-4	Ethylbenzene	0.0E+0	1.3E+2	NA	NA	>Res	ΝA	>Res	1.3E+2		<1
206-44-0	Fluoranthene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	>Res		<1
91-20-3	Naphthalene	0.0E+0	6.2E+1	NA	NA	>Res	NA	>Res	6.2E+1		<1
85-01-8	Phenanthrene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	>Res		<1
129-00-0	Pyrene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	>Res		<1
127-18-4	Tetrachloroethene	0.0E+0	7.1E+3	NA	NA	1.2E+2	NA	6.4E+3	1.2E+2		<1
108-88-3	Toluene	0.0E+0	3.5E+2	NA	NA	>Res	NA	>Res	3.5E+2		<1
71-55-6	Trichloroethane, 1,1,1-	0.0E+0	3.3E+2	NA	NA	2.4E+3	NA	>Res	3.3E+2		<1
79-00-5	Trichloroethane, 1,1,2-	0.0E+0	4.1E-1	NA	NA	1.1E+2	NA	>Res	4.1E-1		<1
79-01-6	Trichloroethene	0.0E+0	2.3E+0	NA	NA	>Res	NA	>Res	2.3E+0		<1
75-01-4	Vinyl chloride	0.0E+0	9.3E-2	NA	NA	3.3E+0	NA	1.6E+2	9.3E-2		<1
1330-20-7	Xylene (mixed isomers)	0.0E+0	>Res	NA	NA	>Res	NA	>Res	>Res		<1

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Software: GSI RBCA Spreadsheet Version: v 1.0

	RBCA SITE	ASSESS	MENT					-1	fi er 1 Wor ksh	eet 6.2	
Site Name: 1970 Seminary		Completed	By: David Hoe	der							
Site Location: Oakland, C		Date Compl	eted: 10/3/199	6							1 OF 1
		Target Ris	sk (Class A & B)	1.0E-4	MCL exp	osure limit?		Calci	lation Option	c 1	
SUBSURFACE SOIL RBSL	VALUES	Targe	t Risk (Class C)) 1.0E-4	PEL expo	osure limit?					
(>3 F F BGS)		Target	Hazard Quotient	t 1.0E+0							
	Representative		Real	Results For Comp	ete Exposure l	Pathways ("x" if	Complete)			Pasi	
	Concentration	v		-	Soil Ve	platilization to	Soil V	olatilization to	Applicable	Exceeded	
		A So Remdontial	Commercial:	Groundwater Regulates (MCL)	X in	idoor Air	A O	utdoor Air	RBSL	?	Required CRF
CAS No. Name	(mg/kg)	(on-site)	(on-site)	(on-site)	(on-site)	(on-site)	(on-site)	(on-site)	(mg/kg)	"■" If yes	Only if "yes" left
83-32-9 Acenaphthene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	NA	>Res		<1
120-12-7 Anthracene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	NA	>Res		<1
71-43-2 Benzene	2.4E+0	4.7E+0	NA	NA	1.5E-1	NA	1.2E+2	NA	1.5E-1	-	1.6E+01
75-00-3 Chloroethane	0.0E+0	1.4E+2	NA	NA	2.5E+2	NA	>Res	NA	1.4E+2		<1
95-50-1 Dichlorobenzene (1,2) (-o)	1.7E+0	2.3E+3	NA	NA	4.5E+2	NA	>Res	NA	4.5E+2		<1
106-46-7 Dichlorobenzene, (1,4) (-p)	0.0E+0	2.5E+2	NA	NA	1.4E+2	NA	>Res	NA	1.4E+2		<1
75-34-3 Dichloroethane, 1,1-	0.0E+0	9.0E+1	NA	NA	1.2E+1	NA	>Res	NA	1.2E+1		<1
107-06-2 Dichloroethane, 1,2-	0.0E+0	2.0E+0	NA	NA	9.7E-1	NA	1.7E+2	NA	9.7E-1		<1
156-59-2 Dichloroethene, cis-1,2-	0.0E+0	6.2E+0	NA	NA	8.6E-1	NA	>Res	NA	8.6E-1		<1
156-60-5 Dichloroethene, 1, 2-trans-	0.0E+0	9.5E+0	NA	NA	1.7E+0	NA	>Res	NA	1.7E+0		<1
100-41-4 Ethylbenzene	4.2E+0	1.3E+2	NA	NA	3.8E+1	NA	>Res	NA	3.8E+1		<1
206-44-0 Fluoranthene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	NA	>Res		<1
91-20-3 Naphthalene	0.0E+0	6.2E+1	NA	NA	4.4E+1	NA	>Res	NA	4.4E+1		<1
85-01-8 Phenanthrene	0.0E+0	>Res	NA	NA	2.0E+2	NA	>Res	NA	2.0E+2		<1
129-00-0 Pyrene	0.0E+0	>Res	NA	NA	>Res	NA	>Res	NA	>Res		<1
127-18-4 Tetrachloroethene	1.8E+0	7.1E+3	NA	NA	5.2E+3	NA	>Res	NA	5.2E+3		<1
108-88-3 Toluene	3.5E+0	3.5E+2	NA	NA	2.3E+1	NA	>Res	NA	2.3E+1		<1
71-55-6 Trichloroethane, 1,1,1-	0.0E+0	3.3E+2	NA	NA	4.6E+1	NA	>Res	NA	4.6E+1		<1
79-00-5 Trichloroethane, 1,1,2-	0.0E+0	4.1E-1	NA	NA	4.4E-1	NA	2.8E+2	NA	4.1E-1		<1
79-01-6 Trichloroethene	8.2E-1	2.3E+D	NA	NA	3.3E+0	NA	>Res	NA	2.3E+0		<1
75-01-4 Vinyl chloride	0.0E+0	9.3E-2	NA	NA	6.7E-2	NA	5.3E+1	NA	6.7E-2		<1
1330-20-7 Xylene (mixed isomers)	8.3E+0	>Res	NA	NA	>Res	NA	>Res	NA	>Res		<1

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Software: GSI RBCA Spreadsheet Serial: g-265-vhx-686 Version: v 1.0 —

RBCA SITE ASSESSMENT

Representative

Site Name: 1970 Seminary Site Location: Oakland, C

GROUNDWATER RBSL VALUES

Completed By: David Hoexter Date Completed: 10/3/1996

> Target Risk (Class A & B) 1.0E-4 Target Risk (Class C) 1.0E-4

MCL exposure limit? D PEL exposure limit? **Calculation Option: 1**

Target Hazard Quotient 1.0E+0

RBSL Results For Complete Exposure Pathways ("x" if Complete)

		Concentration	X Groundwater Ingestion		Groundwater Volatilization X to Indoor Air		Groundwater Volatilization X to Outdoor Air		Applicable RBSL	RBSL Exceeded ?	Required CRF	
CAS No.	Name	(mg/L)	Residential (on-site)	Commercial: (on-site)	Regulatory(MCL): (on-site)	Residential: (on-site)	Commercial: (on-site)	Residential (on-site)	Commercial: (on-site)	(mg/L	"∎" If yes	Only if "yes" left
83-32-9	Acenaphthene	0.0E+0	>Sol	NA	NA	>Sol	NA	>Sol	NA	>Sol		<1 ,
120-12-7	Anthracene	0.0E+0	>Sol	NA	NA	>Sol	NA	>Sol	NA	>Sol		<1
· 71-43-2	Benzene	5.6E+1	2.9E-1	NA	NA	9.8E-1	NA	6.0E+2	NA	2.9E-1		1.9E+02
75-00-3	Chloroethane	1.4E-3	1 5E+1	NA	NA	1.2E+3	NA	>Sol	NA	1.5E+1		<1
95-50-1	Dichlorobenzene (1,2) (-o)	2.2E-2	3.3E+0	NA	NA	5.8E+1	NA	>Sol	NA	3.3E+0		<1
106-46-7	Dichlorobenzene, (1,4) (-p)	0.0E+0	3 5E-1	NA	NA	1.6E+1	NA	>Sol	NA	3.5E-1		<1
75-34-3	Dichloroethane, 1,1-	0.0E+0	3.7E+0	NA	NA	4.5E+1	NA	>Sol	NA	3.7E+0		<1
107-06-2	Dichloroethane, 1,2-	1.5E-2	9.4E-2	NA	NA	3.5E+0	NA	1.7E+3	NA	9.4E-2		<1
156-59-2	Dichloroethene, cis-1,2-	3.2E-1	3.7E-1	NA	NA	1.7E+0	NA	>Sol	NA	3.7E-1		<1
156-60-5	Dichloroethene,1,2-trans-	9.2E-3	7.3E-1	NA	NA	1.3E+1	NA	>Sol	NA	7.3E-1		<1
100-41-4	Ethylbenzene	2.8E+1	3.7E+0	NA	NA	>Sol	NA	>Sol	NA	3.7E+0		8.0E+00
206-44-0	Fluoranthene	0.0E+0	>Sol	NA	NA	>Sol	NA	>Sol	NA	>Sol		<1
91-20-3	Naphthalene	0.0E+0	1.5E-1	NA	NA	6.9E+0	NA	>Sol	NA	1.5E-1		<1
85-01-8	Phenanthrene	0.0E+0	1.5E-1	NA	NA	>Sol	NA	>Sof	NA	1.5E-1		<1
129-00-0	Pyrene	0.0E+0	>Sol	NA	NA	>Sol	NA	>Sol	NA	>Sol		<1
127-18-4	Tetrachloroethene	1.3E-1	1.6E-1	NA	NA	2.2E+1	NA	>Sol	NA	1.6E-1		<1
108-88-3	Toluene	6.1E+1	7.3E+0	NA	NA	6.6E+1	NA	>Sol	NA	7.3E+0		8.0E+00
71-55-6	Trichloroethane, 1,1,1-	0.0E+0	3.3E+0	NA	NA	8.1E+1	NA	>Sol	NA	3.3E+0		<1
79-00-5	Trichloroethane, 1,1,2-	0.0E+0	1.5E-1	NA	NA	9.1E+0	NA	3.7E+3	NA	1.5E-1		<1
79-01-6	Trichloroethene	3.4E-1	2.2E-1	NA	NA	6.6E+0	NA	>Sol	NA	2.2E-1		2.0E+00
75-01-4	Vinyl chloride	6.0E-2	4,5E-3	NA	NA	3.7E-2	NA	2.5E+1	NA	4.5E-3		1.3E+01
1330-20-7	Xylene (mixed isomers)	1.45+2	7 3E+1	NA	NA	>Sol	NA	>Sol	NA	7.3E+1		2.0E+00

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Serial: g-265-vhx-686

Tier 1 Worksheet 6.3

1 OF 1

RBCA TIER 1/TIER 2 EVALUATION

Output Table 1

Residential

Commercial/Industrial

Chronic Construction

Site Name: 1970 Seminary Site Location: Oakland, C Job Identification: E-10-1B-192B Date Completed: 10/3/96 Completed By: David Hoexter Software: GSI RBCA Spreadsheet Version: v 1.0

Parameters Definition (Units)

Surface

NOTE: values which differ from Tier 1 default values are shown in bold italics and underlined.

	DEFA	ULT PARA	METERS				
Exposure			Residential		Commercial/Industrial		
Parameter	Definition (Units)	Adult	(1-6yrs)	(1-16 yrs)	Chronic	Constrctn	
ATc	Averaging time for carcinogens (yr)	70					
ATn	Averaging time for non-carcinogens (yr)	30	6	16	25	1	
BW	Body Weight (kg)	70	15	35	70		
ED	Exposure Duration (yr)	30	6	16	25	1	
EF	Exposure Frequency (days/yr)	350			250	180	
EF Derm	Exposure Frequency for dermal exposure	350			250		
lRgw	Ingestion Rate of Water (I/day)	2			1		
IRs	Ingestion Rate of Soil (mg/day)	100	200		50	100	
Radj	Adjusted soil ing rate (mg-yr/kg-d)	1.1E+02			9.4E+01		
Ralin	inhalation rate indoor (m^3/day)	15			20		
Ra.out	Inhalation rate outdoor (m^3/day)	20			20	10	
SA	Skin surface area (dermal) (cm ²)	5.8E+03		2.0E+03	5.8E+03	5.8E+03	
SAadj	Adjusted dermal area (cm^2•yr/kg)	2.1E+03			1.7E+03		
M	Soil to Skin adherence factor	1					
AAFs	Age adjustment on soil ingestion	TRUE			TRUE		
AAFd	Age adjustment on skin surface area	TRUE			TRUE		
tax	Use EPA tox data for air (or PEL based)	TRUE					
gwMCL?	Use MCL as exposure limit in groundwater?	FALSE					

Matrix of Exposed Persons to		posed Persons to Residential		Commercial/Industrial		
Complete	Exposure Pathways		Chronic	Constrctn		
Groundwa	iter Pathways:					
GW.i	Groundwater Ingestion	FALSE	TRUE			
GW.v	Volatilization to Outdoor Air	FALSE	TRUE			
GW b	Vapor Intrusion to Buildings	FALSE	TRUE			
Soil Pathy	vays					
S.v	Volatiles from Subsurface Soils	FALSE	TRUE			
SS.v	Volatiles and Particulate Inhalation	FALSE	TRUE	TRUE		
SS.d	Direct Ingestion and Dermal Contact	FALSE	TRUE	TRUE		
S.I	Leaching to Groundwater from all Soils	FALSE	TRUE			
S.b	Intrusion to Buildings - Subsurface Soils	FALSE	TRUE			

Matrix of F	Receptor Distance	Resi	dential	Commercial/Industrial		
and Locati	tion an- ar off-site	Distance	On-Site	Distance	On-Site	
G₩	Groundwater receptor (cm)		TRUE		TRUE	
S	Inhalation receptor (cm)		TRUE		TRUE	
Matrix of						
Target Ris	sks	Individual	Cumulative			
TRab	Target Risk (class A&B carcinogens)	1.0E-04				
TRo	Target Risk (class C carcinogens)	1.0E-04				
THQ	Target Hazard Quotient	1.0E+00				
Opt	Calculation Option (1, 2, or 3)	1				
Tier	RBCA Tier	1				

t	Exposure duration (yr)	30	25	1
A	Contaminated soil area (cm^2)	1.9E+06		1.0E+06
w	Length of affected soit parallel to wind (cm)	1.5E+03		1.0E+03
W.gw	Length of affected soit parallel to groundwater (c	1.5E+03		
Uair	Ambient air velocity in mixing zone (cm/s)	2.3E+02		
delta	Air mixing zone height (cm)	2.0E+02		
Lss	Definition of surficial soils (cm)	1.0E+02		
Pe	Particulate areal emission rate (g/cm*2/s)	2.2E-10		
Groundwater	Definition (Units)	Value		
delta.gw	Groundwater mixing zone depth (cm)	6.1E+02	-	
Ī	Groundwater infiltration rate (cm/yr)	1.5E+01		
Ugw	Groundwater Darcy velocity (cm/yr)	1.2E+03		
Ugw tr	Groundwater Transport velocity (cm/yr)	6 6E+03		
Кš	Saturated Hydraulic Conductivity(cm/s)			
grad	Groundwater Gradient (cm/cm)			
Sw	Width of groundwater source zone (cm)			
Sd	Depth of groundwater source zone (cm)			
BC	Biodegradation Capacity (mg/L)			
BIO?	Is Bioattenuation Considered	FALSE		
phi.eff	Effective Porosity in Water-Bearing Unit	3.8E-01		
foc.sat	Fraction organic carbon in water-bearing unit	1.0E-03		
Soil	Definition (Units)	Value		
hc	Capillary zone thickness (cm)	1.5E+01	-	
hv	Vadose zone thickness (cm)	1.6E+02		
rho	Soil density (g/cm^3)	1.7		
foc	Fraction of organic carbon in vadose zone	0.01		
phi .	Soil porosity in vadose zone	0.38		
Lgw	Depth to groundwater (cm)	<u>1.7E+02</u>		
Ls	Depth to top of affected soil (cm)	2.1E+02		
Lsubs	Thickness of affected subsurface soils (cm)	8.8E+02		
pН	Soil/groundwater pH	6.5		
		capillary	vadose	foundation
phi.w	Volumetric water content	0.342	0.12	0.12
phi.a	Volumetric air content	0.038	0.26	0.26
Building	Definition (Units)	Residential	Commercial	
Lb	Building volume/area ratio (cm)	2.0E+02	3.0E+02	
ER	Building air exchange rate (s^-1)	1.4E-04	2.3E-04	
Lork	Foundation crack thickness (cm)	1.5E+01		
eta	Foundation crack fraction	0.01		
Dispersive Tr	Participation (Inclusion)	Decidenti-1	Operatoria - 1	
rarameters	Cennition (Units)	Residential	Commercial	
Groundwater	n an star al coal al construction constants de la serie de la s			
ax	Longitudinal dispersion coefficient (cm)			
ay	Transverse dispersion coefficient (cm)			
az	Vertical dispersion coefficient (cm)			

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Vapor dcy

dcz

Transverse dispersion coefficient (cm) Vertical dispersion coefficient (cm)

RBCA SITE ASSESSMENT

Representative

. Tier 1 Worksheet 6.1

Calculation Option: 1

Site Name: 1970 Seminary Site Location: Oakland, C

c	Completed By: David Hoexter
σ	Date Completed: 10/3/1996
	Target Risk (Class A & B) 1.0E-4
	Target Risk (Clase C) 1.0E-4
	Target Hazard Quotient 1.0E+0

MCL exposure limit?
 PEL exposure limit?

1 OF 1

SURFACE SOIL RBSL VALUES (< 3 FT BGS)

RBSL Results For Complete Exposure Pathways ("x" if Complete)

CONSTITUENTS OF CONCERN		Concentration			Ingestion, Inhalation			Construction RBSL				
			X So	il Leaching to	Leaching to Groundwater		nd Dermal Contact	Х	Worker	Applicable RBS	L Exceeded ?	Required CRF
CAS No.	Name	(mg/kg)	Residential: (on-site)	Commercial: (on-site)	Regulatory(MCL): (on-site)	Resider (on-sil	tiał: Commercial le) (on-site)	C	ommercial: (on-site)	(mg/kg)	" = " If yes	Only if "yes" left
83-32-9	Acenaphthene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
120-12-7	Anthracene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
71-43-2	Benzene	0.0E+0	NA	1.6E+1	NA	NA	3.6E+2		>Res	1.6E+1		<1
75-00-3	Chloroethane	0.0E+0	NA	3.8E+2	NA	NA	>Res		>Res	3.8E+2		<1
95-50-1	Dichlorobenzene (1,2) (-o)	0.0E+D	NA	>Res	NA	NA	>Res		>Res	>Res		<1
106-46-7	Dichlorobenzene, (1,4) (-p)	0.0E+0	NA	8.4E+2	NA	NA	4.3E+2		>Res	4.3E+2		<1
75-34-3	Dichloroethane, 1,1-	0.0E+0	NA	2.5E+2	NA	NA	3.8E+3		3.7E+3	2.5E+2		<1
107-06-2	Dichloroethane, 1,2-	0.0E+0	NA	6.8E+0	NA	NA	1.1E+2		2.6E+3	6.8E+0	D	<1
156-59-2	Dichloroethene, cis-1,2-	0.0E+0	NA	1.7E+1	NA	NA	3.7E+2		3.4E+2	1.7E+1		<1
156-60-5	Dichloroethene,1,2-trans-	0.0E+0	NA	2.6E+1	NA	NA	>Res		>Res	2.6E+1		<1
100-41-4	Ethylbenzene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
206-44-0	Fluoranthene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
91-20-3	Naphthalene	0.0E+0	NA	1.7E+2	NA	NA	>Res		>Res	1.7E+2		<1
85-01-8	Phenanthrene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
129-00-0	Pyrene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
127-18-4	Tetrachioroethene	0.0E+0	NA	2.4E+4	NA	NA	2.1E+2		6.4E+3	2.1E+2		<1
108-88-3	Toluene	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1
71-55-6	Trichloroethane, 1,1,1-	0.0E+0	NA	9.1E+2	NA	NA	3.5E+3		>Res	9.1E+2		<1
79-00-5	Trichloroethane, 1,1,2-	0.0E+0	NA	1.2E+0	NA	NA	1.8E+2		>Res	1.2E+0		<1
79-01-6	Trichloroethene	0.0E+0	NA	6.4E+0	NA	NA	>Res		>Res	6.4E+0		<1
75-01-4	Vinyl chloride	0.0E+0	NA	3.1E-1	NA	NA	5.7E+0		1.6E+2	3.1E-1		<1
1330-20-7	Xylene (mixed isomers)	0.0E+0	NA	>Res	NA	NA	>Res		>Res	>Res		<1

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	RBCA SITE	E ASSESSI	MENT					τ	lier 1 Worksh	eet 6.2	
Site Name: 1970 Seminary		Completed E	By: David Hoe	xter							
Site Location: Oakland, C		Date Comple	eted: 10/3/199	6		1 14 0					1 OF 1
SURGURFACE SOIL DRSI	VALUES	Target Risk (Class A & B) 1.0E-4			MCL exposure limit?		Calculation Option			: 1	
	VALUES	Target	Risk (Class C	1.UE-4	LI PEL expos	surê limit?					
(= 311 863)		i arget F	RBSI	[1.05+0 Results For Como	ole Exposure P	athwave ("y" if	Complete)				
	Representative	KEOL KARRING FBL COUD			ere Exposure r	allina jo (x ii	completey	lete)			
CONSTITUENTS OF CONCERN	Concentration	Y Soil Leaching to Group		Groundwater	Soil Volatilization to X Indeer Air		Soil Volatilization to X Outdoor Air		Applicable	Exceeded	Required CRE
		Residential:	Commercial:	Regulatory(MCL):	Residential;	Commercial	Residential:	Commercial	ROOL	;	Requied City
CAS No. Name	(mg/kg)	(or-sile)	(on-site)	(on-site)	(on-site)	(on-site)	(on-site)	(on-site)	(mg/kg)	"🖤" If yes	Only if "yes" left
83-32-9 Acenaphthene	0.0E+0	NA	>Res	NA	NA	>Res	NA	>Res	>Res		<1
120-12-7 Anthracene	0.0E+0	NA	>Res	NA	NA	>Res	NA	>Res	>Res		<1
71-43-2 Benzene	2.4E+0	NA	1.6E+1	NA	NA	3.1E-1	NA	1.3E+2	3.1E-1		8.0E+00
75-00-3 Chloroethane	0.0E+0	NA	3.8E+2	NA	NA	5.3E+2	NA	>Res	3.8E+2		<1
95-50-1 Dichlorobenzene (1,2) (-o)	1.7E+0	NA	>Res	NA	NA	1.2E+3	NA	>Res	1.2E+3		<1
106-46-7 Dichlorobenzene, (1,4) (-p)	0.0E+0	NA	8.4E+2	NA	NA	4.2E+2	NA	>Res	4.2E+2		<1
75-34-3 Dichloroethane, 1,1-	0.0E+0	NA	2.5E+2	NA	NA	2.6E+1	NA	>Res	2.6E+1		<1
107-06-2 Dichloroethane, 1,2-	0.0E+0	NA	6.8E+0	NA	NA	2.8E+0	NA	2.3E+2	2.8E+0		<1
156-59-2 Dichloroethene, cis-1,2-	0.0E+0	NA	1.7E+1	NA	NA	1.8E+0	NA	>Res	1.8E+0		<1
156-60-5 Dichloroethene,1.2-trans-	0.0E+D	NA	2.6E+1	NA	NA	3.9E+0	NA	>Res	3.9E+0		<1
100-41-4 Ethylbenzene	4.2E+0	NA	>Res	NA	NA	9.8E+1	NA	>Res	9.8E+1		<1
206-44-0 Fluoranthene	0.0E+0	NA	>Res	NA	NA	>Res	NA	>Res	>Res		<1
91-20-3 Naphthalene	0.0E+0	NA	1.7E+2	NA	NA	1.1E+2	NA	>Res	1.1E+2		<1
85-01-8 Phenanthrene	0.0E+0	NA	>Res	NA	NA	>Res	NA	>Res	>Res		<1
129-00-0 Pyrene	0.0E+0	NA	>Res	NA	NA	>Res	NA	>Res	>Res		<1
127-18-4 Tetrachioroethene	1.8E+0	NA	2.4E+4	NA	NA	1.6E+4	NA	>Res	1.6E+4		<1
108-88-3 Toluene	3.5E+0	NA	>Res	NA	NA	5.8E+1	NA	>Res	5.8E+1		<1
71-55-6 Trichloroethane, 1,1,1-	0.0E+0	NA	9.1E+2	NA	NA	1.2E+2	NA	>Res	1.2E+2		<1
79-00-5 Trichloroethane, 1,1,2-	0.0E+0	NA	1.2E+0	NA	NA	1.4E+0	NA	3.9E+2	1.2E+0		<1
79-01-6 Trichloroethene	8.2E-1	NA	6.4E+0	NA	NA	8.6E+0	NA	>Res	6.4E+0	α	<1
75-01-4 Vinyl chloride	0.0E+0	NA	3.1E-1	NA	NA	1.7E-1	NA	7.4E+1	1.7E-1		<1
1220 20 7 Vulana (mivad isomore)	8 36+0		>	R1A		- Paa	NIA	• D- •	- D		- 1

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	RBCA SITE ASSESSMENT		Tier 1 Worksheet 6.3			
Site Name: 1970 Seminary	Completed By: David Hoexter					
Site Location: Oakland, C	Date Completed: 10/3/1996			1 OF 1		
	Target Risk (Class A & B) 1.0E-4	MCL exposure limit?	Calculation Option: 1			
GROUNDWATER RBSL VALUES	Target Risk (Class C) 1.0E-4	PEL exposure limit?				

Target Hazard Quotient 1.0E+0

RBSL Results For Complete Exposure Pathways ("x" if Complete)

	Representative Concentration				Groundwater Volatilization		Groundwater Volatilization		Applicable	RBSL	
		x	Groundwater Ingestion		X to Indoor Air		X to Outdoor Air		RBSL	Exceeded ?	Required CRF
CAS No. Name	(mg/L)	Residential (on-site)	Commercial: (on-site)	Regulatory(MCL): (on-site)	Residential: (on-site)	Commercial: (on-site)	Residential (on-site)	Commercial: (on-site)	(mg/L	" \$ " If yes	Only if "yes" left
83-32-9 Acenaphthene	0.0E+0	NA	>Sol	NA	NA	>Sol	NA	>Sol	>Sol		<1
120-12-7 Anthracene	0.0E+0	NA	>Sol	NA	NA	>Sol	NA	>Sol	>Sol		<1
71-43-2 Benzene	5.6E+1	NA	9.9E-1	NA	NA	2.5E+0	NA	8.4E+2	9.9E-1		5.7E+01
75-00-3 Chloroethane	1.4E-3	NA	4.1E+1	NA	NA	3.1E+3	NA	>Sol	4.1E+1		<1
95-50-1 Dichlorobenzene (1,2) (-o)	2.2E-2	NA	9.2E+0	NA	NA	1.5E+2	NA	>Sol	9.2E+0		<1
106-46-7 Dichlorobenzene, (1,4) (-p)	0.0E+0	NA	1.2E+0	NA	NA	4.8E+1	NA	>Sol	1.2E+0		<1
75-34-3 Dichloroethane, 1,1-	0.0E+0	NA	1.0E+1	NA	NA	1.2E+2	NA	>Sol	1.0E+1		<1
107-06-2 Dichloroethane, 1,2-	1.5E-2	NA	3.1E-1	NA	NA	1.0E+1	NA	2.7E+3	3.1E-1		<1
156-59-2 Dichloroethene, cis-1,2-	3.2E-1	NA	1.0E+0	NA	NA	4.4E+0	NA	>Soł	1.0E+0		<1
156-60-5 Dichloroethene, 1, 2-trans-	9.2E-3	NA	2.0E+0	NA	NA	3.4E+1	NA	>Sol	2.0E+0		<1
100-41-4 Ethylbenzene	2.8E+1	NA	1.0E+1	NA	NA	>Sol	NA	>Sol	1 0E+1		3.0E+00
206-44-0 Fluoranthene	0.0E+0	NA	>Sol	NA	NA	>Sol	NA	>Sol	>Sol		<1
91-20-3 Naphthalene	0.0E+0	NA	4.1E-1	NA	NA	1.8E+1	NA	>Sol	4.1E-1		<1
85-01-8 Phenanthrene	0.0E+0	NA	4.1E-1	NA	NA	>Sol	NA	>Sol	4.1E-1		<1
129-00-0 Pyrene	0.0E+0	NA	>Sol	NA	NA	>Sol	NA	>Sol	>Sol		<1
127-18-4 Tetrachloroethene	1.3E-1	NA	5.5E-1	NA	NA	7.0E+1	NA	>Sol	5.5E-1		<1
108-88-3 Toluene	6.1E+1	NA	2.0E+1	NA	NA	1.7E+2	NA	>Sol	2.0E+1		3.0E+00
71-55-6 Trichloroethane, 1,1,1-	0.0E+0	NA	9.2E+0	NA	NA	2.1E+2	NA	>Sol	9.2E+0		<1
79-00-5 Trichloroethane, 1,1,2-	0.0E+0	NA	4.1E-1	NA	NA	2.8E+1	NA	>Sol	4.1E-1		<1
79-01-6 Trichloroethene	3.4E-1	NA	6.1E-1	NA	NA	2.0E+1	NA	>Sol	6.1E-1		<1
75-01-4 Vinyl chloride	6.0E-2	NA	1.5E-2	NA	NA	1.2E-1	NA	4.3E+1	1.5E-2		4.0E+00
1330-20-7 Xylene (mixed isomers)	1.4E+2	NA	>Sol	NA	NA	>Sol	NA	>Sol	>Sol		<1

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Serial: g-265-vhx-686

APPENDIX C

Terra Vac Corporation February 5, 1997 Report

DUAL VAPOR EXTRACTION PILOT STUDY GRIMIT AUTO AND REPAIR 1970 SEMINARY AVENUE OAKLAND, CALIFORNIA

PROJECT 30-0195

DUAL VAPOR EXTRACTION PILOT STUDY GRIMIT AUTO AND REPAIR 1970 SEMINARY AVENUE OAKLAND, CALIFORNIA

Prepared For:

Doyle Grimit 14366 Lark Street San Leandro, California 94578

Prepared By:

Terra Vac Corporation 1651 Alvarado Street San Leandro, California 94577

Robert Tarr

Staff Engineer

for

Mark P. Frye Project Engineer

February 5,1997

DUAL VAPOR EXTRACTION PILOT STUDY GRIMIT AUTO AND REPAIR 1970 SEMINARY AVENUE OAKLAND, CALIFORNIA

1.0 Introduction

At the request of Doyle Grimit, Terra Vac performed a dual vapor extraction pilot study at the Grimit Auto and Repair site. The purpose of the study was to collect data on the performance of dual vapor extraction technology when applied at the site. Terra Vac understands that this report will be used to evaluate remedial options for addressing hydrocarbon impacted soil and groundwater beneath the site.

2.0 Site Description

The project site is located at 1970 Seminary Avenue in Oakland, California. The neighborhood generally consists of residential houses with nearby one, two, or three-story apartment buildings. The property is bordered by Seminary Avenue on the northwest and Harmon Avenue on the northeast. The site comprises an automobile service building with an office, an attached canopy, and a small detached storage building.

The site is paved throughout with the exception of an approximate 900 square foot area where the former underground storage tanks (UST) were located. The UST area was over-excavated and clean soil was used as backfill.

3.0 Pilot Study Summary

The pilot study was conducted to determine; (a) the radius of influence of an applied vacuum to an existing well on-site, and (b) the resultant groundwater flow rate from that well. To complete this, Terra Vac mobilized a system which comprised of:

- 10HP blower;
- Generator;
- Carbon canister;
- Well head adapting equipment;
- Knock out pot; and
- other miscellaneous equipment.

An existing on-site well was used as the extraction well for this pilot study. The extraction well was adapted with fittings for the 10HP blower to induce a vacuum of approximately 12" Hg. The fittings included a slurp tube that extended down the well that was used to extract

groundwater. The groundwater removed from the extraction well is separated from the knock out pot. Monitoring well MW-1 was selected because the screened interval allowed soil vapors to be drawn from the surrounding subsurface area.

The radius of influence was monitored from two 1-inch black iron pipes driven into the subsurface. The driven pipes are hereinafter, referred to as observation points. Vacuum gauges connected to the observation points were used to measure the amount of vacuum produced in the soil at different distances from the extraction well.

Monitoring well MW-1 has a two-inch casing and is screened across the interval extending approximately 15 to 35 feet below grade. Prior to the start of the study, groundwater was encountered at a depth of approximately 14 feet below grade. The observation points, OB-1 and OB-2, were driven approximately six feet into the subsurface. The locations of MW-1, OB-1, and OB-2 are shown on Figure 1.

Terra Vac mobilized test equipment to the site on January 28, 1997. A 34 foot-long slurp tube was set in MW-1 and the dual vapor extraction system was operated for slightly over three hours. Throughout the duration of the study, Terra Vac monitored the vacuum applied to the slurp tube, induced air flow rates out of the extraction well, the amount of vacuum applied to the well casing and formation, and the resultant vacuum at the observation points. The rate at which groundwater was extracted from MW-1 was also noted. Two samples of the extracted soil vapors were collected and analyzed by Terra Vac for total petroleum hydrocarbons and benzene, toluene, ethylbenzene, and xylenes. Tabulated field data is presented in Table 1.

4.0 Pilot Study Results

An evaluation of the monitoring data indicates the following:

- The induced air flow rate from the extraction well was approximately 11 standard cubic feet per minute with an applied vacuum of 12 inches of mercury column.
- A significant amount of bleed air was required to maintain air flow and groundwater removal within the extraction well casing. Extraction flow rates are expected to increase significantly with continuous application of vacuum to the low permeable materials as a result of dewatering. Wells screened exclusively for dual vacuum extraction will also enhance flow rates.
- The vacuum effectively applied to the well casing and formation was approximately 4 inches of mercury column.
- A vacuum of approximately 0.2 inches of water column was observed in OB-1 at the end of three hours of test operation. OB-1 was located at a distance of approximately 14 feet from MW-1. At the same time, a vacuum of approximately 0.1 inches of water column

was observed in OB-2 which was located at a distance of approximately 25 feet from MW-1. The amount of vacuum observed in OB-1 is significant and is indicative of some degree of connectivity between MW-1 and OB-1. There appeared to be some connectively between MW-1 and OB-2, however the amount of induced vacuum was not as significant.

• A total of 130 gallons of groundwater were extracted during three hours of testing corresponding to an overall groundwater extraction rate of approximately 0.7 gallons per minute.

5.0 Conclusion

The radius of influence of operating the dual vapor extraction system extended to at least 14 feet, with a trace influence at approximately 25 feet from MW-1. The initial TPH-g concentrations decreased from 39.7 mg/L to to 12.6 mg/L during this study. Based on these facts, Terra Vac believes Dual Vapor Extraction, the process of extracting vapor and groundwater simultaneously, is a viable alternative to effectively and rapidly remove the subsurface contaminants at the Grimit Auto and Repair site.



Table 1 Grimit Auto and Repair Pilot Test Field Data 28 January 1997

_	Time	Vacuum ("Hg)	MW-1("Hg)	Ob-1 ("H2O))b-1 ("H2O) Ob-2 ("H2O) f		Pitot	Remarks	
	1213	13.0	0.0	0.00	0.03	Open	0	Drawing H2O down	
	1214	()	0.0	0.00	()	Open	()	Stop system	
	1217	12.0	0.0	0.00	0.03	Open	0	Start system	
	1220	12.8	4.0	0.00	0.03	1/4 Open	0.8	Moderate water flow	
	1223	12.0	3.5	0.00	0.02	1/2 Open	0.3	Took vapor sample 1	
	1227	12.0	3.5	0.06	0.02	3/4 Open	0.2	Low water flow	
	1232	12.0	3.5	0.00	0.02	3/4 Open	0.6	Moderate water flow	
	1242	12.0	3.8	0.00	0.02	3/4 Open	0.2	Moderate water flow	
	1257	12.0	4.0	0.00	0.02	3/4 Open	0.2	Moderate water flow	
	1304	12.0	4.0	0.18	0.04	3/4 Open	0.2	Moderate water flow	
	1325	12.0	4.0	0.15	0.04	3/4 Open	0.4	Moderate water flow	
	1348	12.0	4.0	0.00	0.01	3/4 Open	0.3	Low water flow	
	1412	12.0	4.0	0.00	0.05	3/4 Open	3.6	Low water flow	
	1430	12.0	4.0	0.00	0.00	3/4 Open	5.6	Moderate water flow	
	1445	12.0	4.0	0.00	0.00	3/4 Open	5	Moderate water flow	
	1500	12.0	4.0	0.04	0.10	3/4 Open	5	Moderate water flow	
	1503	()	()	()	()	()	()	Took vapor sample 2	
	1504	12.0	4.0	0.10	0.10	3/4 Open	0.3	Moderate water flow	
	1510	12.0	4.0	0.20	0.09	3/4 Open	0.5	Moderate water flow	
	1515	12.0	4.0	0.18	0.08	3/4 Open	0.5	Moderate water flow	
	1520	12.0	4.0	0.18	0.08	3/4 Open	0.5	Moderate water flow	

