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## GROUNDWATER MONITORING REPORT SR 522 and NE 180th Street RIVERSIDE PROPERTY BOTHELL, WASHINGTON

September 7, 1993

Prepared for:

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#### **EXECUTIVE SUMMARY**

Groundwater Technology, Inc. conducted groundwater monitoring and sampling at the property located near the intersection of State Route 522 and NE 180th Street in Bothell, Washington, known as the Riverside Property. The work was performed to assess the current groundwater conditions underlying the site with respect to substances regulated under the Washington Department of Ecology (WDOE) Model Toxics Control Act (MTCA). Groundwater Technology personnel gauged the depth to water in six groundwater monitoring wells and collected water samples from five. The water samples were analyzed for benzene, toluene, ethylbenzene, xylenes, total petroleum hydrocarbons (TPH)-as-gasoline, TPH, volatile organic compounds and lead.

### Observations and findings:

- The apparent groundwater flow direction is toward the south-southeast at approximately
   2.73 feet per 100 feet.
- Groundwater Samples analyzed for volatile organics, including halogenated hydrocarbons, and gasoline constituents were non-detectable at the respective method detection limits. In addition, total petroleum hydrocarbons (TPH) were non-detectable in the water samples tested.
- Lead concentrations above groundwater MTCA compliance cleanup levels (CCL) were reported in samples collected from monitoring wells MW-1A and MW-3A.



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### GROUNDWATER MONITORING REPORT SR 522 and NE 180TH STREET RIVERSIDE PROPERTY BOTHELL, WASHINGTON

#### 1.0 INTRODUCTION

Presented in this report are the worksteps and results associated with groundwater monitoring and sampling conducted by Groundwater Technology, Inc. at the Texaco Refining and Marketing Inc. Riverside Property located at State Route 522 and NE 180th Street, Bothell, Washington (See Figure 1, Site Location Map). The work was performed to assess the current groundwater conditions underlying the site with respect to substances regulated under the Washington Department of Ecology (WDOE) Model Toxics Control Act (MTCA).

Groundwater Technology previously installed six groundwater monitoring wells and collected groundwater samples in July, 1992. At that time, an open excavation partially filled with water was present at the site. Since then, the excavation was pumped out and petroleum contaminated sediment and soil were excavated from the bottom of the excavation. The excavation was subsequently backfilled above groundwater level with imported clean material. The previous investigation identified two water bearing units at the site; a lower apparently confined water bearing zone, present below a depth of approximately 12 feet, and a shallow perched zone present above a depth of about eight feet. Three of the six groundwater monitoring wells installed in July 1992 (MW-1, MW-2, and MW-3) are completed in the lower zone. The other three wells (MW-1A, MW-2A, and MW-3A) were installed to a depth of eight feet in the perched zone. Tests from the previous groundwater sampling detected concentrations of gasoline constituents, total petroleum hydrocarbons (TPH) as analyzed by EPA method 418.1, trichloroethene, and lead above MTCA Method A compliance cleanup levels (CCL).

### 2.0 SCOPE OF WORK

The following outline summarizes the specific worksteps involved:

- Gauged the depth to water in each of the monitoring wells;
- Purged and sampled 5 of the 6 previously installed groundwater monitoring wells;

Washington Department of Ecology (WAC 173-340)

- Analyzed each water sample from monitoring wells MW-1, MW-2, MW-3, MW-1A and MW-3A for benzene, toluene, ethylbenzene, xylenes (BTEX), total petroleum hydrocarbons-asgasoline (TPH-G), TPH, volatile organic compounds (VOCs) and lead;
- Analyzed, summarized and presented the information obtained in report form.

#### 3.0 WELL MONITORING AND SAMPLING

On February 25, 1993 water levels were measured in groundwater monitoring wells MW-1, MW-1A, MW-2, MW-2A, MW-3 and MW-3A to evaluate groundwater flow direction and gradient. Table 1 shows the relative well-head elevation, measured depth-to-water (DTW), and relative potentiometric groundwater elevations in the monitoring wells. Based on the data, the gradient was approximately 2.73 feet per 100 feet in the lower zone. A groundwater potentiometric surface map was drawn from the data for the lower zone (Figure 3). Water was present in only two of three shallow wells, therefore no gradient was calculated nor map drawn.

Table 1 Groundwater Monitoring Data											
Well Number	MW-1	MW-1A	MW-2	MW-2A	MW-3	MW-3A					
Well-Head Elevation (Feet)	101.82	101.74	101.57	101.47	100.22	100.34					
Date: July 15, 1992											
DTW	8.57	7.63*	9.16	7.70*	10.02	5.63					
Elevation (Feet)	93.25	94.11	94.41	93.77	90.22	94.71					
Date: August 24, 1992											
DTW	9.08	7.89*	9.64	7.69*	10.29	NM					
Relative Elev. (Ft)	92.74	93.85	91.93	93.78	89.93						
Date: August 31, 1992											
DTW	9.18	7.88*	9.75	7.70*	10.32	6.59					
Relative Elev. (Ft)	92.64	93.86	91.85	93.77	89.90	93.75					
Date: February 25, 1993											
DTW	7.87	6.99	9.45	7.74*	9.56	4.45					
Relative Elev. (Ft)	93.95	94.75	92.12	93.73	90.66	95.89					

DTW = Depth to Water

NM = Not Measured (inaccessible)

Note: Elevations are relative based on an arbitrary common datum of 100 feet.



<sup>\*</sup> Likely not a representative groundwater elevation, water is trapped in the well casing.

### D. Mark Wells Texaco Environmental Services

Following groundwater monitoring on February 25, 1993, wells MW-1, MW-1A, MW-2, MW-3 and MW-3A were purged of approximately three well volumes and water samples were collected in accordance with the Standard Operating Procedures in Appendix A. Samples collected were designated MW-1, MW-1A, MW-2, MW-3 and MW-3A and sent to an analytical laboratory with proper Chain-of-Custody documentation. A degree of turbidity was noted in each sample; primarily in the shallow samples, MW-1A and MW-3A. Well MW-2A was not sampled, because the well was dry. Purged water was stored on site pending laboratory analysis.

#### 4.0 LABORATORY ANALYSIS

Groundwater samples collected were analyzed at GTEL Environmental Laboratories in Concord, California. Each water sample was analyzed for BTEX and TPH-G by EPA Methods 8020 and modified 8015 (WTPH-G), TPH by EPA Method 418.1 (WTPH), VOCs by EPA Method 8240, and lead by EPA Method 7421. Water analyses results are summarized in Table 2. Complete laboratory results are contained in Appendix C.

	Summ	Sample		Results - bruary 25,		vater .		
ANALYTE	EPA METHOD	MDL	MW-1	MW-1A	MW-2	MW-3	MW-3A	CCL
Benzene	8020	0.3	ND	ND	ND	ND	ND	5
Toluene	8020	0.3	ND	ND	ND	ND	ND	40
Ethylbenzene	8020	0.3	ND	ND	ND	ND	ND	30
Xylene	8020	0.5	ND	ND	ND	ND	ND	20
TPH-G	Mod. 8015	10	ND	ND	ND	ND	ND	1000
TPH	WTPH 418.1	1000	ND	ND	ND	ND	ND	1000
VOCs	8240	varies	ND	ND	ND	ND	ND	varies
Lead	7421	5	ND	56	ND	ND	53	5

MDL = Method Detection Limit

ND = Not Detected at MDL

CCL = WAC-340, Model Toxics Act, Method A Compliance Cleanup Levels - Groundwater

VOCs = Volatile Organic Compounds



### D. Mark Wells Texaco Environmental Services

The only detected compound was lead in samples MW-1A and MW-3A at concentrations of 56 and 53  $\mu$ g/L, respectively. Both concentrations are above the CCL of 5  $\mu$ g/L.

### 5.0 CONCLUSIONS AND RECOMMENDATIONS

Groundwater monitoring and sampling was completed at the Riverside Property, Bothell, Washington in February, 1993. The apparent groundwater gradient was 2.73 feet per 100 feet to the south-southeast on February 25, 1993. Total lead concentrations above MTCA CCLs for groundwater were reported in samples collected from monitoring wells MW-1A and MW-3A.

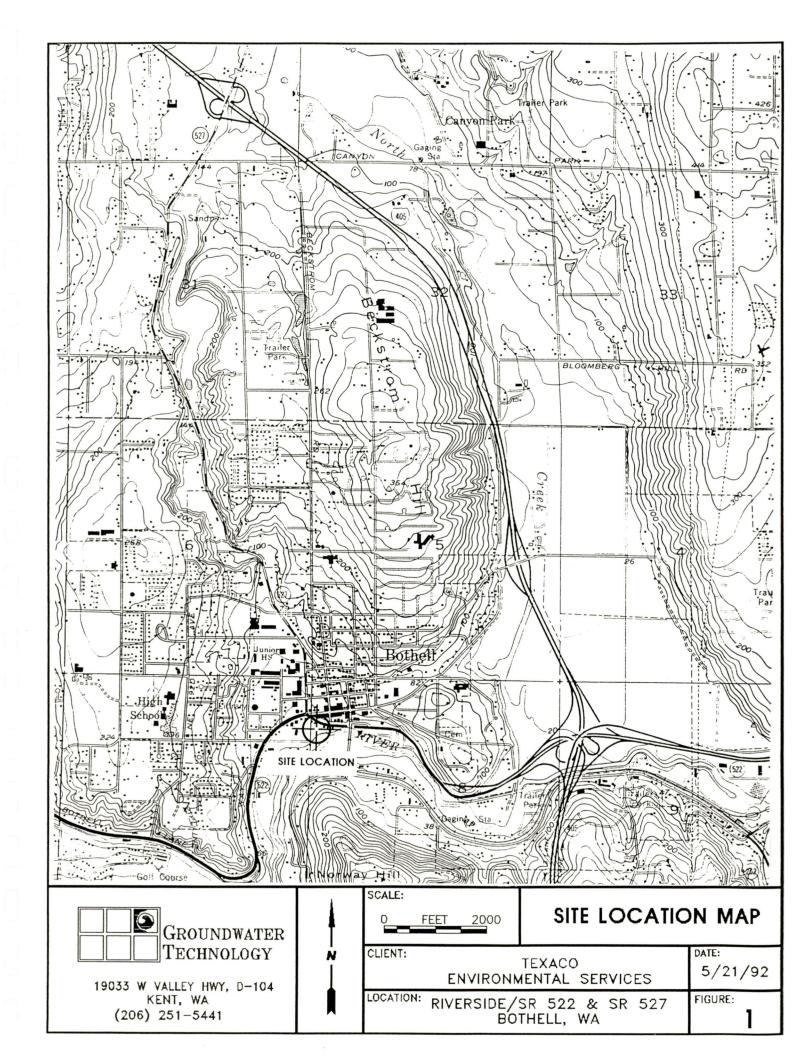
One previous groundwater sampling event was conducted at the site in July, 1992. Groundwater concentrations above MTCA CCLs were reported at that time in monitoring wells MW-1 (TPH), MW-2 (lead), MW-3 (lead) and MW-3A (lead and trichloroethene). Lead was the only compound detected during this sampling event. Due to the turbidity noted in the water samples collected, the reported lead concentrations may not represent actual groundwater conditions. We recommended filtering any future groundwater samples prior to conducting inorganic analyses. This procedure is allowed in accordance with WAC 173-340-720 (8)(a)(i).

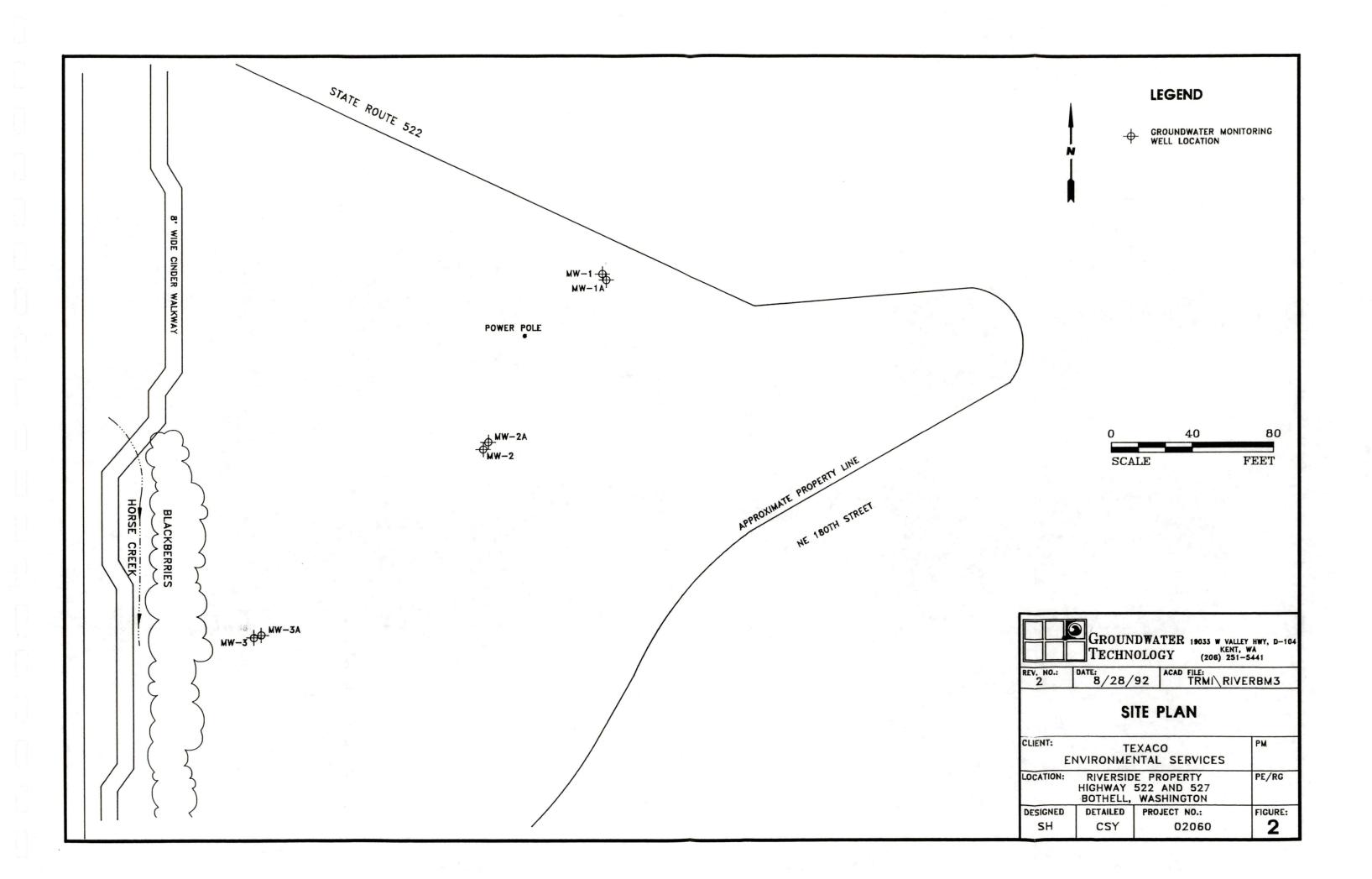


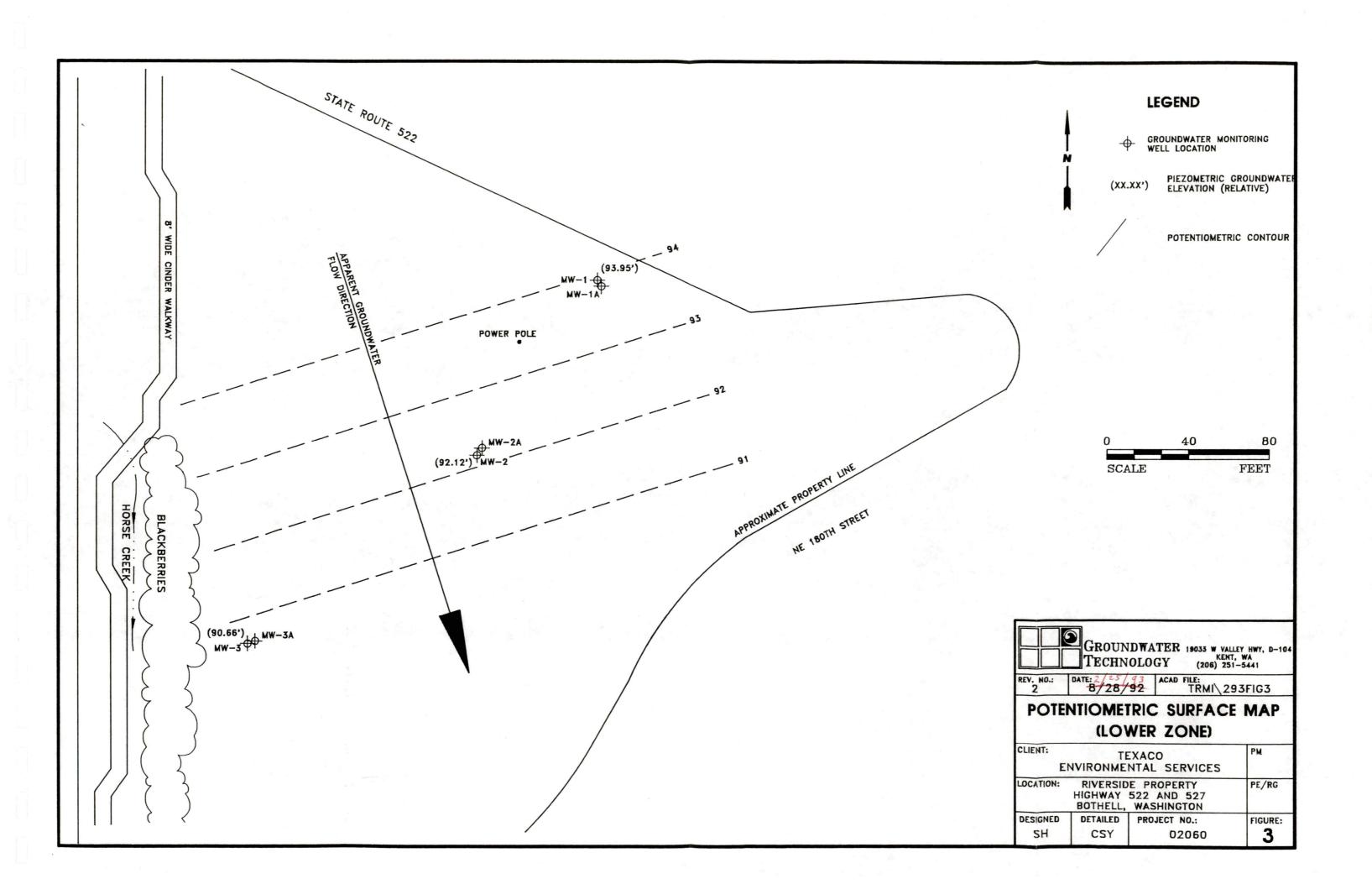


**FIGURES** 











# APPENDIX A STANDARD OPERATING PROCEDURES



#### 3.0 WATER QUALITY SAMPLING

- 3.1 Water samples should not be taken from the stagnant water in the well.
- 3.2 Water samples should be taken in triplicate.
- 3.3 Remove 3 to 5 volumes of water in the well prior to sampling. The water may be removed by bailing, submersible pump, or purge system. Wells with a slow recovery period should be bailed dry and then sampled within 1 hour or when recovered to 80%. Monitor pH, temperature and specific conductivity with each well volume to insure water quality stabilization has occurred. However, this is not necessary at every well or in all circumstances.
- 3.4 Use only Teflon, stainless steel, or glass bailers to obtain the sample. Use Teflon only for sampling water containing chlorinated compounds and also for bacteriological samples. PVC bailers can be used for one-time sampling for other than EPA 624 analysis. Using a bailer for a one-time sampling reduces the possibility for cross-contamination.
- 3.5 When sampling, avoid stirring up any sediments in the well and agitating the water to reduce volitization of any dissolved compounds that may be present.
- 3.6 All sampling equipment must be cleaned following the appropriate procedure to avoid cross contamination from site to site and sample to sample. The sampling equipment should be cleaned before each well sampling, between each sampling, and at the end of each sampling round.
- 3.7 Monitoring wells should be gauged prior to sampling.
- 3.8 If possible, the monitoring wells should be sampled starting with the cleanest well and ending with the most contaminated well.
- 3.9 Wells containing free-phase contaminants should not be sampled.
- 3.10 When filling out the chain of custody form:
  - enter the samples in the order in which they were collected;
  - make a note as to the cleaning fluid used to clean the sampling equipment;
  - attempt to identify which samples are the most contaminated;
  - complete all other requested information.
- 3.11 The laboratory sample identification label should be filled out with a waterproof pen and firmly affixed to each sample container. Typically, identification labels require that the following information be supplied:
  - job name
  - job number
  - sampler's name
  - sample identification
  - date sampled and time
  - analysis requested



- 3.12 Acidification is required for samples that will be analyzed by the EPA 624 method. (see Acidification Procedure in this section)
- 3.13 Acidification is recommended for EPA method 601 and 602 samples to preserve them and increase their holding life. (see Acidification Procedure in this section)
- 3.14 Field blanks should be taken as part of each sampling round. A field blank consists of a sample of distilled water which has been collected by putting the distilled water into a sampling bailer after the bailer has been cleaned following the procedure used to clean that bailer during the sampling round. The field blank is stored with the samples. It is not analyzed unless requested by the Project Manager. The field blank should not be identified as such to the laboratory.
- 3.15 Handling of decontaminated equipment:
  - Always use "pristine" gloves (latex, solvex, etc.).
  - Place decontaminated bailers on clean surface (plastic).
  - Do not wipe down bailer with paper towels or cloth. Follow decontamination procedure.
- 3.16 Sample accuracy can be adversely affected by the entrainment of sediment in wells which have not been properly developed. Contaminants adhering to the sediments can be released when samples are acidified for preservation. Therefore, if sediments are present, field filtering of the samples is recommended.
- 3.17 Chemical changes can take place because the sample was oxidized during sampling. It is critical to avoid oxidation of samples when sampling for volatile organic compounds (VOC). Therefore, take care to insure minimal agitation occurs during sampling.
- 3.18 All samples should be properly and promptly preserved.

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- 3.19 All samples should be analyzed quickly; arrangements should be made with the testing laboratory to insure prompt analysis is performed within the allowable times for the specific analyses to be done.
- 3.20 Bailer strings that have contacted water or contaminants should be replaced between each well to avoid contamination from a bailer string which has absorbed contamination. A good practice is to replace the string between wells. <u>Caution</u>: some bailer strings are treated with a fungicide which may be detected in priority pollutant analysis.
- 3.21 Notify laboratory that samples are being shipped in advance of sampling to insure proper delivery and turnaround.
- 3.22 On the chain of custody, note what type of decontamination or preservation fluids, chemicals were used.



- 4.0 ACIDIFICATION PROCEDURE (EPA Methods 601,602, and 624)
  - 4.1 At the start of each sampling round, the amount of acid required to lower a sampling container of water to be sampled to a pH of less than 2 should be determined.
  - 4.2 After removing 3 to 5 well volumes from the first well to be sampled, put 5-10 drops of 50% HCL into a 40 ml sample vial (larger sampling container will require more acid) and fill the vial with water form the well; determine the pH of water in the vial with pH paper; if the pH is too high, repeat the procedure using 15-20 drops of acid in the vial; repeat until the pH of the water in the sample vial is a pH of less than 2 on the pH paper. Note the amount of acid required to lower the pH of the volume of water in the sampling vial. (pH paper should not be placed into sampling container. Pour sample onto pH paper to check for proper pH.)
  - 4.3 Discard the practice acidified sample.
  - 4.4 Once the amount of acid required to reach a pH of <2 is known, the acid can be routinely added to each sample container directly; the water to be analyzed is added to vial or container containing the appropriate amount of acid.
  - 4.5 Note that the amount of acid required is site specific and should be noted on the Chain of Custody form.
  - 4.6 The procedure should be repeated for each site at the start of each sampling round.
  - 4.7 Equipment
    - Bailer or other means to remove 3 to 5 well volumes
    - Sampling bailer
    - Polyethylene squirt bottle of 50% hydrochloric (HCL) acid
    - Narrow range pH paper (1.0 2.5 pH range)
    - Paper towels
    - Waterproof pen
    - Laboratory sample identification labels
    - Cooler with ice
    - Chain of custody forms
    - Sample containers (usually 40 ml glass vials with teflon faced septums)
    - Alconox solution and/or methanol
    - Distilled water
    - Safety equipment (gloves, etc.)
    - Dissolved oxygen meter (sometimes used in limited biorec projects in conjunction with bacteriological testing)





# APPENDIX B LABORATORY ANALYTICAL RESULTS







Client Number: 020603778 Project ID: Bothell, WA Work Order Number: C3-02-0162

### Northwest Region

4080-C Pike Lane Concord, CA 94520 (510) 685-7852 (800) 544-3422 from inside California (800) 423-7143 from outside California (510) 825-0720 (FAX)

March 11, 1993

Mark Nichols Groundwater Technology, Inc. 19033 W. Valley Hwy., D-104 Kent, WA 98032

Enclosed please find the analytical results for samples received by GTEL Environmental Laboratories, Inc. on 02/26/93, under chain of custody record 25920.

A formal Quality Assurance/Quality Control (QA/QC) program is maintained by GTEL, which is designed to meet or exceed the EPA requirements. Analytical work for this project met QA/QC criteria, unless otherwise stated in the footnotes.

GTEL is certified by the California State Department of Health Services, Laboratory certificate numbers 194 and 1075, to perform analyses for drinking water, wastewater, and hazardous waste materials according to EPA protocols.

If you have any questions concerning this analysis or if we can be of further assistance, please call our Customer Service Representative.

Sincerely,

GTEL Environmental Laboratories, Inc.

E Seen F. Bullen R.M.

Eileen F. Bullen

Laboratory Director

Client Number: 020603778 Project ID: Bothell, WA Work Order Number: C3-02-0162

### Table 1

## **ANALYTICAL RESULTS**

# Aromatic Volatile Organics and Total Petroleum Hydrocarbons as Gasoline in Water

## EPA Methods 5030, 8020, and WTPH-Ga

GTEL Sample Number		01	02	03	04
Client Identification		MW-1	MW-1A	MW-2	MW-3
Date Sampled		02/25/93	02/25/93	02/25/93	02/25/93
Date Analyzed		03/07/93	03/07/93	03/07/93	03/07/93
Analyte	Detection Limit, ug/L		Concentra	ation, ug/L	
Benzene	0.3	< 0.3	< 0.3	< 0.3	< 0.3
Toluene	0.3	< 0.3	< 0.3	< 0.3	< 0.3
Ethylbenzene	0.3	< 0.3	< 0.3	< 0.3	< 0.3
Xylene, total	0.5	<0.5	< 0.5	< 0.5	<0.5
BTEX, total					
TPH as Gasoline	10	<10	<10	<10	<10
Detection Limit Multiplier		1	1	1	1

a. Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, US EPA November 1986. Modification for TPH as gasoline as per the State of Washington Department of Ecology, Appendix L, April, 1992.



Client Number: 020603778 Project ID: Bothell, WA Work Order Number: C3-02-0162

## Table 1 (Continued)

## **ANALYTICAL RESULTS**

# Aromatic Volatile Organics and Total Petroleum Hydrocarbons as Gasoline in Water

## EPA Methods 5030, 8020, and WTPH-Ga

GTEL Sample Number		05			
Client Identification		MW-3A			
Date Sampled		02/25/93			
Date Analyzed		03/07/93			
Analyte	Detection Limit, ug/L		Concentration	on, ug/L	
Benzene	0.3	< 0.3			
Toluene	0.3	< 0.3			
Ethylbenzene	0.3	< 0.3			
Xylene, total	0.5	<0.5			
BTEX, total					
TPH as Gasoline	10	<10			
Detection Limit Multiplier		1			

Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, US EPA November 1986. Modification for TPH as gasoline as per the State of Washington Department of Ecology, Appendix L, April, 1992.



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Field Sample ID	GTEL Lab # (Lab use only)	# Containers	WATER		SLUDGE PRODUCT	ОТНЕЯ	Pı	Meth rese OS <sup>2</sup> H	rve	d	(SPECIFY) DATE	mpling B	BTEX/602 □ 8020	BTEX/Gas Hydroca	Hydrocarbons GC/F	Hydrocarbon Profile (SIMDIS) □	Oil and Grease 413	TPH/IR 418.1 □ SM 503 □	EDB by 504 □ DBCP by 504	EPA 503.1 □ EPA 502.2 □	EPA 601 □ EPA 8010 □	EPA 602 □ EPA 8020 □	EPA 608 □ 8080 □ PCB only □	EPA 624/PPL □ 82	EPA 625/PPL □ 82	EPA 610 □ 8310 □	EP TOX Metals □ F	TCLP Metals □ VO	EPA Metals - Priorit	CAM Metals TTLC ☐ STLC ☐	Lead 239.2 🗆 200.7	Organic Lead	Corrosivity   Flas		
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**Northwest Region** 

4080-C Pike Lane Concord, CA 94520 (510) 685-7852 (800) 544-3422 from inside California (800) 423-7143 from outside California (510) 825-0720 (FAX) Client Number: 020603778 Consultant Project Number: FDMW156 Project ID: Bothell, WA Work Order Number: C3-04-0014

April 13, 1993

Mark Nichols Groundwater Technology, Inc. 19033 West Valley Hwy., D-104 Kent, WA 98032

Enclosed please find the analytical results for samples received by GTEL Environmental Laboratories, Inc. on 04/01/93, under chain of custody record 25819.

A formal Quality Assurance/Quality Control (QA/QC) program is maintained by GTEL, which is designed to meet or exceed the EPA requirements. Analytical work for this project met QA/QC criteria, unless otherwise stated in the footnotes.

GTEL is certified by the California State Department of Health Services, Laboratory certificate numbers 194 and 1075, to perform analyses for drinking water, wastewater, and hazardous waste materials according to EPA protocols.

If you have any questions concerning this analysis or if we can be of further assistance, please call our Customer Service Representative.

Sincerely,

GTEL Environmental Laboratories, Inc.

illen F. Bullen

Eileen F. Bullen

**Laboratory Director** 

Client Number: 020603778
Consultant Project Number: FDMW156
Project ID: Bothell, WA
Work Order Number: C3-04-0014

### Table 1

## **ANALYTICAL RESULTS**

# Total Petroleum Hydrocarbons in Water by Infrared Spectrometry

## WTPH 418.11(SM 5520 FC2)

Methods for Chemical Analysis of Water and Wastes, EPA 600/4-79-202, Revised March 1983, U.S. Environmental

Protection Agency.

Standard Methods for the Examination of Water and Wastewater, 17th ed., 1989, American Public Health Association. "Modification in TPH Methods as per the state of Washington Department of Ecology, Appendix L, April, 1992."

GTEL Sample Number		01	02	03	04
Client Identification		MW1	MW1A	MW2	MW3
Date Sampled		03/30/93	03/30/93	03/30/93	03/30/93
Date Prepared		04/09/93	04/09/93	04/09/93	04/09/93
Date Analyzed		04/09/93	04/09/93	04/09/93	04/09/93
Analyte	Detection Limit, mg/L		Concentra	tion, mg/L	
Total Petroleum Hydrocarbons	1	<1	<1	<1	<1
Detection Limit Multiplier		1	1	1	1

GTEL Sample Number		05			
Client Identification		MW3A			
Date Sampled		03/30/93			
Date Prepared		04/09/93			
Date Analyzed		04/09/93			
Analyte	Detection Limit, mg/L		Concentra	ition, mg/L	
Total Petroleum Hydrocarbons	1	<1			
Detection Limit Multiplier		1			



Client Number: 020603778 Consultant Project Number: FDMW156 Project ID: Bothell, WA Work Order Number: C3-04-0014

Table 1

## **ANALYTICAL RESULTS**

## Volatile Organics in Water

GTEL Sample Number		01	02	03	04
Client Identification		MW-1	MW-1A	MW-2	MW-3
Date Sampled		03/30/93	03/30/93	03/30/93	03/30/93
Date Analyzed		04/01/93	04/01/93	04/01/93	04/01/93
Analyte	Detection Limit, ug/L		Concentration	on, ug/L	
Chloromethane	10	<10	<10	<10	<10
Bromomethane	10	<10	<10	<10	<10
Vinyl chloride	10	<10	<10	<10	<10
Chloroethane	10	<10	<10	<10	<10
Methylene chloride	5	<5	<5	<5	<5
Acetone	20	<20	<20	<20	<20
Carbon disulfide	5	<5	<5	<5	<5
1,1-Dichloroethene	5	<5	<5	<5	<5
1,1-Dichloroethane	5	<5	<5	<5	<5
1,2-Dichloroethene, total	5	<5	<5	<5	<5
Chloroform	5	<5	<5	<5	<5
1,2-Dichloroethane	5	<5	<5	<5	<5
2-Butanone	20	<20	<20	<20	<20
1,1,1-Trichloroethane	5	<5	<5	<5	<5
Carbon tetrachloride	5	<5	<5	<5	<5
Vinyl acetate	50	<50	<50	<50	<50
Bromodichloromethane	5	<5	<5	<5	<5
1,2-Dichloropropane	5	<5	<5	<5	<5
cis-1,3-Dichloropropene	5	<5	<5	<5	<5
Trichloroethene	5	<5	<5	<5	<5

Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, US EPA November 1986 (method modified for additional compounds). Sample introduction by EPA Method 5030.



## Table 1 (Continued)

## **ANALYTICAL RESULTS**

## Volatile Organics in Water

GTEL Sample Number		01	02	03	04
Client Identification		MW-1	MW-1A	MW-2	MW-3
Date Sampled		03/30/93	03/30/93	03/30/93	03/30/93
Date Analyzed		04/01/93	04/01/93	04/01/93	04/01/93
Analyte	Detection Limit, ug/L		Concentration	on, ug/L	
Dibromochloromethane	5	<5	<5	<5	<5
1,1,2-Trichloroethane	5	<5	<5	<5	<5
Benzene	5	<5	<5	<5	<5
trans-1,3-Dichloropropene	5	<5	<5	<5	<5
2-Chloroethylvinyl ether	10	<10	<10	<10	<10
Bromoform	5	<5	<5	<5	<5
4-Methyl-2-pentanone	20	<20	<20	<20	<20
2-Hexanone	20	<20	<20	<20	<20
Tetrachloroethene	5	<5	<5	<5	<5
1,1,2,2-Tetrachloroethane	5	<5	<5	<5	<5
Toluene	5	<5	<5	<5	<5
Chlorobenzene	5	<5	<5	<5	<5
Ethylbenzene	5	<5	<5	<5	<5
Styrene	5	< 5	<5	<5	<5
1,2-Dichlorobenzene	5	<5	<5	<5	<5
1,3-Dichlorobenzene	5	<5	<5	<5	<5
1,4-Dichlorobenzene	5	<5	<5	<5	<5
Xylene, total	5	<5	<5	<5	<5
Trichlorofluoromethane	5	<5	<5	<5	<5
Detection Limit Multiplier		. 1	. 1	1	1

Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, US EPA November 1986 (method modified for additional compounds). Sample introduction by EPA Method 5030.



# Table 1 (Continued)

## ANALYTICAL RESULTS

# Volatile Organics in Water

GTEL Sample Number		05			
Client Identification		MW-3A			
Date Sampled		03/30/93			
Date Analyzed		04/01/93			
Analyte	Detection Limit, ug/L		Concentrati	on, ug/L	17
Chloromethane	10	<10			
Bromomethane	10	<10	1		
Vinyl chloride	10	<10			
Chloroethane	10	<10			
Methylene chloride	5	<5			
Acetone	20	<20			
Carbon disulfide	5	<5			
1,1-Dichloroethene	5	<5			
1,1-Dichloroethane	5	<5			
1,2-Dichloroethene, total	5	<5			
Chloroform	5	<5			
1,2-Dichloroethane	5	<5			
2-Butanone	20	<20			
1,1,1-Trichloroethane	5	<5			
Carbon tetrachloride	5	<5			
Vinyl acetate	50	<50			
Bromodichloromethane	5	<5			
1,2-Dichloropropane	5	<5			
cis-1,3-Dichloropropene	5	< 5			
Trichloroethene	5	<5			

Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, US EPA November 1986 (method modified for additional compounds). Sample introduction by EPA Method 5030.



# Table 1 (Continued)

## **ANALYTICAL RESULTS**

# Volatile Organics in Water

GTEL Sample Number		05			
Client Identification		MW-3A			
Date Sampled		03/30/93			
Date Analyzed		04/01/93			
Analyte	Detection Limit, ug/L		Concentrati	on, ug/L	
Dibromochloromethane	5	<5			
1,1,2-Trichloroethane	5	<5			
Benzene	5	<5			
trans-1,3-Dichloropropene	5	<5			
2-Chloroethylvinyl ether	10	<10			
Bromoform	5	<5			
4-Methyl-2-pentanone	20	<20			·
2-Hexanone	20	<20			
Tetrachloroethene	5	<5	2 19 3		
1,1,2,2-Tetrachloroethane	5	<5			
Toluene	5	<5			
Chlorobenzene	5	<5			
Ethylbenzene	5	<5			
Styrene	5	<5			
1,2-Dichlorobenzene	5	<5			
1,3-Dichlorobenzene	5	<5			
1,4-Dichlorobenzene	5	<5			
Xylene, total	5	<5			
Trichlorofluoromethane	5	<5			
Detection Limit Multiplier		1			

Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, US EPA November 1986 (method modified for additional compounds). Sample introduction by EPA Method 5030.



Client Number: 020603778
Consultant Project Number: FDMW156
Project ID: Bothell, WA
Work Order Number: C3-04-0014

Table 1

## **ANALYTICAL RESULTS**

# Lead in Water by Graphite Furnace AA

# EPA Methods 7421<sup>1</sup>/3005<sup>2</sup>

GTEL Sample Number		01	02	03	04			
Client Identification		MW-1	MW-1A	MW-2	MW-3			
Date Sampled		03/30/93	03/30/93	03/30/93	03/30/93			
Date Prepared		04/05/93	04/05/93	04/05/93	04/05/93			
Date Analyzed		04/07/93	7/93 04/07/93 04/07/93 0					
Analyte	Detection Limit, ug/L							
Lead, total	5	<5	56	<5	<5			
Detection Limit Multiplier		1	1 1 1					

- 1. Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, November 1986.
- 2. Sample preparation by EPA Method 3005.



Client Number: 020603778 Consultant Project Number: FDMW156 Project ID: Bothell, WA Work Order Number: C3-04-0014

## Table 1 (Continued)

## ANALYTICAL RESULTS

## Lead in Water by Graphite Furnace AA

## EPA Methods 7421<sup>1</sup>/3005<sup>2</sup>

GTEL Sample Number		05			
Client Identification		MW-3A			
Date Sampled		03/30/93			
Date Prepared		04/05/93			
Date Analyzed		04/07/93			
Analyte	Detection Limit, ug/L		Concentra	ation, ug/L	
Lead, total	5	53			
Detection Limit Multiplier		. 1			

- 1. Test Methods for Evaluating Solid Waste, SW-846, Third Edition, Revision 0, November 1986.
- 2. Sample preparation by EPA Method 3005.



GTEL	4080 PIKE LANE, SUITE C CONCORD, CA 94520 (510) 685-7852 (800) 423-7143							CHAIN-OF-CUSTODY RECORD 25819																									
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Company Name:  Cround Water  Company Address: 19033	Name:  Phone #: Z51-5441  200-  FAX #: Z51-8452  Address: 19033 w Valley D.104 Site location: Bothell Wa.									with MTBE	Screen			and and	166. 22	-Audel pa	412						Herb										
MA/K N/2  I attest that the proper field s	The street that the proper field sampling  Which old (NAME) TX-Riverside  Sampler Name (Print):					with MTBE	BTEX/Gas Hydrocarbons PID/FID ☐ with N	as □ Diesel □	D(SIQ)	Oil and Grease 413.1 □ 413.2 □ SM 503 □		504 🗆					EPA 624/PPL LI 8240/IAL A NBS (+15) LL	EFA 023/FFL	EP TOX Metals   Destinides   Herbinides	TCLP Metals   VOA   Semi-VOA   Pest   Herb	EPA Metals - Priority Pollutant □ TAL □ RCRA	STLC	Lead 239.2 □ 200.7 □ 7420 □ 7421🕱 6010 □		int   Reactivity								
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