

FOURTH QUARTER 2001
GROUNDWATER SAMPLING REPORT
WITH
AMENDED WORKPLAN TO
AUGMENT THE GROUNDWATER WITH OXYGEN
(APPENDIX G)

AT

FORMER DESERT SITE DP 793
4035 PARK BLVD.
OAKLAND, CA.

FOR

DESERT PETROLEUM

January 7, 2002

BY

-WEGE-
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TABLE OF CONTENTS

1.0 SITE LOCATION AND DESCRIPTION	3
2.0 LOCAL GEOLOGY	3
2.1 Geomorphology	3
2.2 Stratigraphy	3
2.1.1 Station Property	3
2.1.2 Backyard Sewer Lateral Route	4
2.1.3 Brighton Avenue	4
3.0 COLLECTION AND ANALYSIS OF GROUNDWATER SAMPLES	4
3.1 Depth to Water Measurements	4
4.0 RESULTS OF QUARTERLY GROUNDWATER MONITORING	4
4.1 Groundwater Gradient and Flow Direction	4
4.2 Results of Certified Analysis of Groundwater Samples	5
5.0 WEEKLY PURGING OF RECEPTOR TRENCH	5
6.0 PUMPING ON-SITE WELL RS-5	5
7.0 BIODEGRADATION OF HYDROCARBONS	6
7.1 Dissolved Oxygen	6
7.2 Sulfate	6
7.3 Nitrate	7
7.4 Ferrous Iron	7
8.0 SUMMARY	7
9.0 LIMITATIONS	8

List of Tables

1. Groundwater Elevation and Certified Analytical Results
2. Groundwater Elevations and Electron Acceptor Results
3. Wastewater Discharge
4. Groundwater Removal
5. Pounds of Hydrocarbons

List of Figures

1. Area Base Map "Geotracker"
2. Portion of USGS Oakland East 7.5 Minute Quadrangle
3. Sample Location Figure
4. Groundwater Gradient, December 18, 2001
5. Groundwater Plume, TPHg & Benzene, December 18, 2001
- 5-1. Groundwater Plume, TPHg, December 18, 2001
6. Groundwater Plume, TPHg & Benzene, August 26, 1999
7. Dissolved Oxygen, December 18, 2001
- 7-1. Dissolved Oxygen, August 26, 1999
8. Sulfate in Groundwater, December 18, 2001
- 8-1. Sulfate in Groundwater, August 26, 1999
9. Nitrate in Groundwater, December 18, 2001

- 9-1 Nitrate in Groundwater, August 26, 1999
- 10. Ferrous Iron in Groundwater, December 18, 2001
- 10-1 Ferrous Iron in Groundwater, August 26, 1999
- 11 Lithology Cross Section Site along sewer in backyards
- 12 Lithology Cross Section, east gutter Brighton Avenue

List of Appendices

- A. METHODS AND PROCEDURES, QA/QC WITH FIELD NOTES
- B. GROUNDWATER ELEVATION CHART
- C. LABORATORY REPORT
- D. TPHg AND BENZENE CHARTS - WELLS
- E. HACK SPECTROPHOTOMETER FIELD TEST METHODS
- F. WASTEWATER DISCHARGE PERMIT RENEWAL
- G. AMENDED WORKPLAN - AUGMENT BIODEGRADATION



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January 7, 2002

Dear Mr. Thompson:

The following report documents the fourth quarter 2001 sampling and includes the modified workplan (see Appendix G) to augment the groundwater with oxygen at DP793, 4035 Park Blvd., Oakland, California.

1.0 SITE LOCATION AND DESCRIPTION

Former Desert Petroleum #793 is a non-active service station, located on the northwest corner of the intersection of Park Boulevard and Hampel Street at 4035 Park Blvd., Oakland, California (Figure 1). The site is located in projected section 32; T1S; R3W; MDB&M at an approximate elevation of 210 feet above mean sea level (Figure 2).

2.0 LOCAL GEOLOGY

2.1 Geomorphology

The site is located on the western slope of the Berkeley Hills. The Berkeley Hills are a northwest-southeast trending range within the Coastal Range Province of California. Erosion of the Coastal Ranges has filled the valleys within and bordering the Coastal Range with sequences of gravels, silts, sands, and clays.

2.2 Stratigraphy

2.1.1 Station Property

The native soil from surface to 13 feet below ground surface (BGS) consists of dark brown silty clay. The dark brown clay is underlain by light brown stiff clay that includes subrounded to rounded metavolcanic gravel. This clay extends to approximately 23 feet BGS at the northwest corner of the site. A fine to medium sand, clayey sand, and silty sand underlies the gravel and clay.

2.1.2 Backyard Sewer Lateral Route

Assessments performed along the sewer lateral as it leaves the site and routes through the residential area towards Brighton Avenue show the subsurface to consist of fill from a couple of inches thick to two feet thick. Beneath the fill is a sequence of clay formations that vary from light brown to dark gray to approximately the 6 foot depth. Silty clay then extends to approximately the 14-foot depth. Beneath the silty clay is sand with occasional gravel. This sand is 11 feet thick at RS5 and is underlain by silty clay, see Figure 11.

2.1.3 Brighton Avenue

Construction of the receptor trench along the eastern curb area of Brighton Avenue revealed two separate sequences of lithology. North of the storm drain catch basin the sequence consists of, clay to the four foot depth, silty clay to the seven foot depth, fine silty sand to the 9 foot depth, medium sand to the 10 foot depth, silty caly to the 11 ½ foot depth, gravel to the 12 foot depth underlain by clay to the 16 foot depth. South of the storm catch basin is a sequence of silty clays and clays to depth, see Figure 12.

3.0 COLLECTION AND ANALYSIS OF GROUNDWATER SAMPLES

Groundwater samples were collected on December 18, 2001. Samples were analyzed for Total Petroleum Hydrocabons as gasoline, Benzene, Toluene, Ethylbenzene, Xylenes and Methyl tert-Butal Alcohol using EPA method 8260B. During this time all wells sampled were field tested for Dissolved Oxygen, Sulfate, Nitrate and Ferrous Iron, see Tables 1 and 2. Figure 3 shows the positions of the groundwater monitoring wells, the receptor trench and previous sample locations.

3.1 Depth to Water Measurements

On December 18, 2001 depth to water was measured at each well using a product/water interface probe. Measurements are referenced to the surveyed elevation at the top of casing at each well. Table 1 shows the elevation of groundwater with respect to mean sea level for all wells through December 18, 2001.

4.0 RESULTS OF QUARTERLY GROUNDWATER MONITORING

4.1 Groundwater Gradient and Flow Direction

Figure 4 shows the groundwater elevation gradients and flow direction that were derived from the depth to water measurements of the monitor wells on December 18, 2001, prior to purging the wells for sampling, see Table 1 and Appendix A. On February 15, 2001 submersible pump was placed into onsite well RS-5 to try and capture contaminated groundwater beneath the site and adjoining properties. The pump rate was set at approximately 2 gpm. As shown on groundwater elevation charts generated for each well, pumping from RS5 lowered the water levels in RS-8 and RS-10, see Appendix B. Table 1 shows the groundwater elevations for the wells during the assesment of this site.

The current flow direction is northwest to west. The hydraulic gradient averages 0.09 feet/linear foot downgradient of RS-8 to the receptor trench well T1, see Figure 4. The present flow direction and hydraulic gradient are consistent with previous determinations by WEGE.

4.2 Results of Certified Analysis of Groundwater Samples

The results of the certified analyses of groundwater samples collected on December 18, 2001 are shown in Table 1.

TPH-G concentrations in water samples from the eight monitor wells, the receptor trench well and three recovery wells ranged from a maximum of 48 mg/l at trench well T1, to below laboratory lower detection limits of 50 ug/L in wells MW1, RS2, RS10, R1 and R3 respectively. Benzene concentrations ranged from a maximum of 3.7 mg/L in T1 to below the laboratory lower detection limits (0.5 ug/L) at wells MW1, RS2, RS10, R1 and R3, see Appendix C - Laboratory Report.

Analysis results for Oxygenant Methyl-t-Butyl Ether (MTBE) was below the laboratory lower detection limit in wells MW1, RS2, RS5, RS6, RS8, RS9, RS10, R1, R2, and R3. RS7 and the trench well T1 contained 2.3 and 24 ug/L MTBE respectively. During the September 16, 1998 all Fuel Oxygenants; MTBE, Di-isopropyl Ether (DIPE), tertiary Butyl Alcohol (TBA), Ethyl-t-Butyl Ether (ETBE) and t-Amyl Methyl Ether (TAME) were confirmed with EPA Method 8260. These analytes were below laboratory lower detection limits. Figure 5 (December 18, 2001) shows the lateral distribution of the hydrocarbon plume with benzene distinction in groundwater as determined from groundwater samples collected from the monitor wells and from non-certified results from the Soil Probe Surveys. Figure 6 (August 26, 1999) shows the lateral distribution of the hydrocarbon plume with benzene distinction in groundwater prior to pumping from RS-5 and T1. Appendix D contains charts developed for wells MW1, RS2, RS5, RS6, RS7, RS8, RS9, RS10 and trench well T1 showing TPHg & Benzene concentration reductions with time, with the exception of T1 which showed an increase in both TPHg and Benzene for the December 18, 2001 sampling.

5.0 WEEKLY PURGING OF RECEPTOR TRENCH

Commencing on May 4, 2000, weekly pumping of the receptor trench has been performed for approximately 4 hours per week, see Table 3. During purging the depth to water within the trench is lowered an average of one foot. Immediately after purging ceases, the water level in the trench recovers to its original depth. Weekly purging of the receptor trench was suspended on July 19, 2001 at the request of Desert Petroleum. 62,511 gallons of contaminated groundwater had been removed from the trench, processed through two, in series, activated carbon water scrubs and discharged to the sanitary sewer, see Table 3.

6.0 PUMPING ON-SITE WELL RS-5

On February 15, 2001 a submersible pump with a pump bypass was placed into RS-5. The pump rate was adjusted to 1.5 gpm and allowed to continuously pump from RS-5 for one week. 3223 gallons were pumped from RS-5 through the two in series water carbon units and discharged to the sewer. On February 22, 2001 the pump was inspected and showed a slimy growth covering the pump and discharge line that was below the water level. The pump was cleaned and placed back

into RS-5 and continued to discharge from RS-5 through the water carbon units to sewer until July 19, 2001. On July 19, 2001 Desert Petroleum requested suspension of further pumping at the site. The pump was removed and the site secured. From February 15 through July 19, 2001, 78,919 gallons of gasoline contaminated groundwater was recovered from RS-5 and treated through carbon before being discharged to the sewer, see Table 3.

The pumping from RS-5 lowered the groundwater at this well by at least 15 feet, when compared to the previous water measurements. This created a cone of influence out to offsite wells RS-8 and RS-10, see Chart - Appendix B. Recirculating the pumped groundwater, before it leaves the well (RS-5) has increased the dissolved oxygen in RS-5 from 0.7 mg/L (August 26, 1999) to 3.1 mg/L (March 8, 2001). This should aid in the biodegradation of the hydrocarbon plume, see Table 2.

7.0 BIODEGRADATION OF HYDROCARBONS

During the December 18, 2001 sampling of wells field measurements were obtained to determine the availability of electron receptors to aid in the natural attenuation of the hydrocarbon plume. Along with pH, temperature and electrical conductivity, dissolved oxygen, nitrate, sulfate and ferrous iron were also measured. Water samples were obtained after the wells were purged and allowed to recovery and analyzed in the field using a Hach DR/2000 Spectrophotometer, see Appendix. The following methods were used:

- Dissolved Oxygen, high range (0 to 13 mg/L O₂) - Method 8166 for water and wastewater.
- Nitrate, high range (0 to 30 mg/L NO₃) - Method 8039 for water, wastewater and seawater.
- Sulfate, (0 to 70 mg/L SO₄) - Method 8051 for water and wastewater.
- Ferrous Iron, (0 to 3.00 mg/L Fe₂) - Method 8146 for water, wastewater and seawater.

Table 2 represents the results of electron acceptor field analysis obtained December 18, 2001 compared to results obtained August 26, 1999.

7.1 Dissolved Oxygen

Readings for dissolved oxygen obtained prior to pumping the receptor trench and RS5 indicated two areas of oxygen depletion (<1 mg/L), the entire north half of the site (4035 Park Avenue) and the area excavated for the receptor trench along the eastern curb of Brighton Avenue, see Figure 7-1. Readings obtained during the December 18, 2001 monitoring round show that dissolved oxygen has increased substantially and even exceeds 5 mg/L in the over-excavated area on site. The lowest Dissolved Oxygen level encountered is associated with well RS5 at 1.4 mg/L, compared to 0.7 mg/L at RS5 in August 1999. All other current dissolved oxygen measurements are at 2.5 mg/L or greater, see Figure 7.

7.2 Sulfate

Figure 8 represents sulfate measurements obtained during the December 18, 2001 quarterly monitoring. Comparing sulfate measurements obtained in August 1999, Figure 8-1, to the

December 2001 measurements, Figure 8, the sulfate has been depleted at the receptor trench and beneath Brighton Avenue, but is being replenished at well location RS8.

7.3 Nitrate

Figure 9 represents nitrate measurements obtained during the December 18, 2001 quarterly monitoring. Comparing nitrate measurements obtained in August 1999, Figure 9-1, to the December 2001 measurements, Figure 9, the nitrate is being replenished all along the petroleum plume area.

7.4 Ferrous Iron

Figure 10 represents ferrous iron measurements obtained during the December 18, 2001 quarterly monitoring. The measurements obtained in August 1999, Figure 10-1, and the December 2001 measurements, Figure 10, indicate that ferrous iron is oxidized, as the site becomes more aerobic.

8.0 SUMMARY

Since the installation and weekly purging of the receptor trench (T1) the TPHg concentrations in down gradient wells RS-7 and RS-9 have decreased, see Table 1 with charts RS-7. The weekly purging of the receptor trench was limited to a maximum daily discharge of 5 gpm, thus removing approximately 1200 to 2000 gallons per week. Although this does lower the water level in the trench, after pumping has ceased the water level rebounds to its original depth allowing for the gradient migration of TPHg contaminated groundwater to continue.

Pumping from RS-5 has shown to create a cone of influence off-site downgradient out to RS-8 and RS-10. Pumping has increased the dissolved oxygen in RS-5 and hydrocarbon concentrations have declined in R1, R3, RS-5, RS-8 and RS-10.

Previous sampling, September 2, 1999, showed that aerobic bacteria (hydrocarbon degraders) exist in the groundwater associated with the hydrocarbon plume. A workplan to augment the groundwater with oxygen (air sparging) and nutrients (phosphate and ammonium sulfate) dated August 29, 2000 was presented with the August 29, 2000, Third Quarter 2000 report. This workplan along with current conditions was discussed during a meeting at Alameda County Health that involved Mr. Thompson, Desert Petroleum, Mr. Seery, Alameda County Health and Mr. Converse, Western Geo-Engineers. The meeting concluded that nutrient augmentation was not necessary at this time, but enhanced dissolved oxygen was needed. Due to neighborhood concerns, i.e. residential homes and apartments, air sparging and/or using a mechanical delivery device would create too much noise and a more passive oxygen delivery system was warranted, i.e. hydrogen peroxide or Oxygen Release Compound (ORC). The amended workplan is presented in Appendix G of this report and suggests that ORC would be the most beneficial means of enhancing dissolved oxygen in the groundwater plume.

9.0 LIMITATIONS

This report is based upon the following:

- A. The observations of field personnel.
- B. The results of laboratory analyses performed by a state certified laboratory.
- C. Referenced documents.
- D. Our understanding of the regulations of the State of California, Alameda County and the City of Oakland.
- E. Changes in groundwater conditions can occur due to variations in rainfall, temperature, local and regional water use, and local construction practices.
- F. In addition, variations in the soil and groundwater conditions could exist beyond the points explored in this investigation.

State Certified Laboratory analytical results are included in this report. This laboratory follows EPA and State of California approved procedures; however, WEGE is not responsible for errors in these laboratory results. Western Geo-Engineers is a corporation under California Registered Geologist #3037 and/or Contractors License #513857. The services performed by Western Geo-Engineers have been conducted in a manner consistent with the level of care and skill ordinarily exercised by members of our profession currently practicing under similar conditions in the State of California and the Oakland area. Our work and/or supervision of remediation and/or abatement operations, active or preliminary, at this site is in no way meant to imply that we are owners or operators of this site. Known or suspected contamination of soil and/or groundwater must be reported to the appropriate agencies in a timely manner. No other warranty, expressed or implied, is made.

Sincerely,



George Converse
Geologist



Jack E. Napper
Ca. Reg. Geologist #3037

cc: Mr. Scott O. Seery, Alameda County Health (510) 567-6783
Mr. Leroy Griffin, Oakland Fire Dept.

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
RS-1	12/14/89	228.15	24.25	203.9	19000	2600	2700	200	1200	
RS-1	12/90				15000	3500	330	170	760	
RS-1	2/91				6900	910	200	39	540	
RS-1	6/91				1600	56	180	12	26	
RS-1	9/91				4100	730	7.6	5.1	24	
RS-1	12/91				8300	950	160	71	190	
RS-1	11/09/92	228.15	17.05	211.1	1700	730	9.6	16	14	
RS-1	04/07/94	228.15	13	215.15	860	84	12	16	110	
RS-1	06/19/94	228.15	13.37	214.78	1400	150	12	52	87	
RS-1	09/17/94	228.15	16.33	211.82	310	30	1.8	2.8	3.9	
RS-1	03/12/95	228.15	4.66	223.49	ND	ND	ND	ND	ND	
		DESTROYED BY OVER-EXCAVATION OF UST-DISPENSER AREAS (8/14/95								
		REPLACED WITH MW-1 9/5/95.								
MW-1	10/04/95	229.5	12.38	217.12	ND	ND	ND	ND	ND	
MW-1	12/21/95	229.5	13.40	216.1	< 50	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
MW-1	03/27/96	229.5	5.53	223.97	< 50	< 0.5	< 0.5	< 0.5	< 2	< 50
MW-1	06/11/96	229.5	9.02	220.48	< 50	< 0.5	< 0.5	< 0.5	< 2	< 50
MW-1	09/04/96	229.5	11.84	217.66	< 50	< 0.5	< 0.5	< 0.5	< 2	< 5
MW-1	12/11/96	229.5	12.98	216.52	< 50	< 0.5	0.9	< 0.5	< 1	< 0.5
MW-1	2/21/97	229.5	9.50	220	< 50	< 0.5	0.9	< 0.5	< 1	< 0.5
MW-1	5/28/97	229.5	11.18	218.32	< 50	3	3	< 0.5	< 1	< 0.5
MW-1	9/2/97	229.5	13.00	216.5	< 50	5	< 0.5	< 0.5	< 1	< 0.5
MW-1	11/24/97	229.5	14.12	215.38	< 50	5	< 0.5	< 0.5	< 1	< 0.5
MW-1	2/25/98	229.5	6.41	223.09	< 50	< 0.5	< 0.5	< 0.5	< 1	< 0.5
MW-1	7/8/98	229.5	7.28	222.22	< 50	< 0.5	< 0.5	< 0.5	< 1	< 1
MW-1	9/16/98	229.5	10.96	218.54	< 50	< 0.5	< 0.5	< 0.5	< 1	< 1
MW-1	11/24/98	229.5	12.24	217.26	52	2.3	5.2	< 0.5	5.4	11
MW-1	2/23/99	229.5	7.14	222.36	< 50	< 0.5	5	< 0.5	< 1	< 0.5
MW-1	5/5/99	229.5	7.00	222.5	< 50	2	<0.5	< 0.5	< 1	8

6

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)										
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)	
RS-2***	8/26/99	227.39	11.42	215.97	200	15	23	1.7	23	9	
RS-2	11/10/99	227.39	15.94	211.45	< 50	<0.5	<0.5	<0.5	< 1	<0.5	
RS-2	2/9/00	227.39	8.91	218.48	< 50	<0.5	<0.5	<0.5	< 1	<0.5	
RS-2	6/30/00	227.39	9.79	217.60	52	2	<0.5	<0.5	< 1	<0.5	
RS-2	8/8/00	227.39	10.71	216.68	60	<0.5	<0.5	<0.5	< 1	<0.5	
RS-2	11/16/00	227.39	10.39	217.00	< 50	<0.5	<0.5	<0.5	< 1	<0.5	
RS-2	3/8/01	227.39	6.62	220.77	< 50	<0.5	<0.5	<0.5	<0.5	<0.5	
RS-2	5/31/01	227.39	10.09	217.30	< 50	<0.5	<0.5	<0.5	<0.5	<0.5	
RS-2	12/18/01	227.39	6.99	220.40	< 50	<0.5	<0.5	<0.5	<0.5	<0.5	
RS-5	12/14/89	227.61	25.97	201.64	57000	3100	4300	670	3400		
RS-5	2/91	227.61	FLOATING PRODUCT								
RS-5	6/91	227.61	FLOATING PRODUCT								
RS-5	9/91	227.61	FLOATING PRODUCT								
RS-5	12/91	227.61	FLOATING PRODUCT								
RS-5	11/09/92	227.61	20.73	206.88	50000	650	4800	1100	15000		
RS-5	04/07/94	227.61	18.16	209.45	27000	5000	8700	550	2800		
RS-5	06/19/94	227.61	18.11	209.5	20000	2100	5300	470	2500		
RS-5	09/17/94	227.61	19.63	207.98	9300	230	340	110	700		
RS-5	03/12/95	227.61	14.54	213.07	93000	6400	2000	19000	10000		
RS-5	10/04/95	227.61	17.53	210.08	16000	420	2100	320	1800		
RS-5	12/21/95	227.61	17.47	210.14	48000	3500	9200	840	4800	56	
RS-5	03/27/96	227.61	13.51	214.1	68000	4900	18000	1700	11000	< 3000	
RS-5	06/11/96	227.61	14.25	213.36	66000	6300	20000	2100	12000	< 3000	
RS-5	09/04/96	227.61	16.50	211.11	31000	2100	11000	1100	6800	400	
RS-5	12/11/96	227.61	15.88	211.73	85000	7000	21000	1800	8900	570	
RS-5	2/21/97	227.61	13.76	213.85 sh	100000	5000	22000	1700	7300	<0.5	
RS-5	5/28/97	227.61	15.77	211.84	52000	4500	19000	2100	10000	<0.5	
RS-5	9/2/97	227.61	17.47	210.14	38000	2100	9400	1300	5800	<0.5	

11

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DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
RS-5	11/24/97	227.61	18.67	208.94	45000	4000	16000	1900	9700	<0.5 *
RS-5	2/25/98	227.61	10.53	217.08	160000	2700	31000	5300	28000	<0.5 *
RS-5	7/8/98	227.61	13.75	213.86	45000	2800	12000	2000	8500	<10 *
RS-5	9/16/98	227.61	15.80	211.81	49000	1400	7500	1700	8600	<5 *
RS-5	11/24/98	227.61	16.64	210.97	89000	5300	15000	2800	13000	<10 *
RS-5	2/23/99	227.61	12.36	215.25	19000	1900	11000	2500	4800	<25 *
RS-5	5/5/99	227.61	12.78	214.83	78000	2000	10000	3000	15000	540 *
RS-5***	8/26/99	227.61	16.06	211.55	35000	870	4000	1900	8300	<1 *
RS-5	11/10/99	227.61	17.54	210.07	40000	1000	5600	1800	8100	<0.5 *
RS-5	2/9/00	227.61	16.31	211.3	46000	1400	6900	2700	11000	<0.5 *
RS-5	6/30/00	227.61	15.15	212.46	37000	810	5200	2200	9100	<2.5 *
RS-5	8/8/00	227.61	16.10	211.51	14000	330	500	1400	6500	<0.5 *
RS-5	11/16/00	227.61	17.38	210.23	23000	430	2300	1100	4800	<0.5 *
RS-5	3/8/01	227.61	27.72	199.89	11000	360	260	140	1500	2.6 ****
RS-5	5/31/01	227.61	22.96	204.65	7500	26	11	38	470	<5 ****
RS-5	12/18/01	227.61	15.61	212	12000	610	1200	100	1500	<5 ****
RS-6	12/14/89	227.22	22.52	204.7	11000	1400	1700	160	860	
RS-6	2/91	227.22	FLOATING PRODUCT							
RS-6	6/91	227.22			95000	4200	4200	650	3700	
RS-6	9/91	227.22	FLOATING PRODUCT							
RS-6	12/91	227.22			64000	3700	2300	730	4100	
RS-6	11/09/92	227.22	19.43	207.79	19000	1600	710	500	1600	
RS-6	04/07/94	227.22	14.42	212.8	16000	1200	1300	290	1100	
RS-6	06/19/94	227.22	14.45	212.77	23000	1300	2200	590	2200	
RS-6	09/17/94	227.22	19.52	207.7	24000	630	790	250	1100	
RS-6	03/12/95	227.22	8.90	218.32	3200	450	13	82	230	
RS-6	10/04/95	227.22	17.78	209.44	3700	170	250	38	290	
RS-6	12/21/95	227.22	14.98	212.24	3100	120	30	16	150	58

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)										
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)	
RS-6	03/27/96	227.22	10.00	217.22	6900	180	440	79	360	< 300	
RS-6	06/11/96	227.22	12.00	215.22	7400	220	150	30	100	<1000	
RS-6	09/04/96	227.22	15.00	212.22	1400	68	2.6	7.7	9.2	14	
RS-6	12/11/96	227.22	12.36	214.86	1800	39	16	10	18	< 0.5	
RS-6	2/21/97	227.22	10.00	217.22	2100	71	85	25	40	< 0.5 *	
RS-6	5/28/97	227.22	13.56	213.66	1700	34	12	11	16	< 0.5 *	
RS-6	9/2/97	227.22	16.35	210.87	940	34	71	9	55	< 0.5 *	
RS-6	11/24/97	227.22	15.72	211.5	490	9	6	1	7	< 0.5 *	
RS-6	2/25/98	227.22	6.26	220.96	1400	22	47	5	52	< 0.5 *	
RS-6**	7/8/98	227.22	11.41	215.81	1500	83	9	84	2	<10 *	
RS-6	7/30/98	227.22			<50	<0.5	<0.5	<0.5	<1		
RS-6	9/16/98	227.22	13.42	213.8	990	23	<0.5	<0.5	<1	<1 *	
RS-6	11/24/98	227.22	15.91	211.31	3400	5.3	<0.5	<0.5	14	<0.5	
RS-6	2/23/99	227.22	7.00	220.22	1000	3.4	3.2	1.6	7.3	<0.5	
RS-6	5/5/99	227.22	10.29	216.93	1100	50	10	80	15	2	
RS-6***	8/26/99	227.22	13.72	213.5	690	44	2.5	30	31	<5	
RS-6	11/10/99	227.22	13.90	213.32	1800	2	2	0.9	16	< 0.5	
RS-6	2/9/00	227.22	12.77	214.45	410	3	3	4	7	< 0.5	
RS-6	6/30/00	227.22	12.69	214.53	660	7	2	5	6	< 0.5	
RS-6	8/8/00	227.22	14.72	212.5	660	2	3	2	6	< 0.5	
RS-6	11/16/00	227.22	15.28	211.94	560	1	2	1	5	< 0.5	
RS-6	3/8/01	227.22	10.10	217.12	2200	<0.5	<0.5	<0.5	<0.5	<0.5 ****	
RS-6	5/31/01	227.22	12.96	214.26	630	<0.5	<0.5	<0.5	<0.5	<5 ****	
RS-6	12/18/01	227.22	10.88	216.34	56	0.53	<0.5	<0.5	0.56	<0.5 ****	
RS-7	12/14/89	195.99									
RS-7	7/90	195.99			560000	24000	210000	50000	740000		
RS-7	2/91	195.99	FLOATING PRODUCT								
RS-7	6/91	195.99	FLOATING PRODUCT								

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
RS-7	9/91	195.99	FLOATING PRODUCT							
RS-7	12/91	195.99			270000	11000	22000	2000	13000	
RS-7	11/09/92	195.99	4.62	191.37	81000	12000	16000	1900	13000	
RS-7	04/07/94	195.99	4.03	191.96	74000	16000	16000	1400	8500	
RS-7	06/19/94	195.99	4.07	191.92	83000	22000	19000	1500	9500	
RS-7	09/17/94	195.99	4.05	191.94	270000	13000	15000	2100	1100	
RS-7	03/12/95	195.99	3.72	192.27	35000	5100	560	6300	3600	
RS-7	10/04/95	195.99	4.03	191.96	96000	14000	14000	1300	7000	
RS-7	12/21/95	195.99	3.95	192.04	70000	9300	12000	860	5600	210
RS-7	03/27/96	195.99	3.80	192.19	64000	8900	14000	1100	8300	< 3000
RS-7	06/11/96	195.99	3.79	192.2	65000	12000	17000	1600	9700	<5000
RS-7	09/04/96	195.99	3.99	192	20000	4900	2100	670	4400	100
RS-7	12/11/96	195.99	3.78	192.21	17000	4400	7500	570	4600	180
RS-7	2/21/97	195.99	3.82	192.17	93000	31000	47000	3800	23000	<0.5 *
RS-7	5/28/97	195.99	3.82	192.17	52000	12000	8200	2000	11000	<0.5 *
RS-7	9/2/97	195.99	3.96	192.03	28000	6100	2800	950	3800	<50
RS-7	11/24/97	195.99	3.76	192.23	18000	4300	5900	600	2900	<0.5 *
RS-7	2/25/98	195.99	3.70	192.29	13000	4300	7100	1100	5800	<0.5 *
RS-7**	7/8/98	195.99	3.76	192.23	45000	10000	3400	2000	8000	<10 *
RS-7	7/30/98	195.99			72000	12000	2100	2000	9100	
RS-7	9/16/98	195.99	3.83	192.16	5000	6500	160	<2.5	500	<5 *
RS-7	11/24/98	195.99	3.77	192.22	19000	2100	1100	500	2100	<0.5
RS-7	2/23/99	195.99	3.70	192.29	83000	6500	9900	1200	7000	<10
RS-7	5/5/99	195.99	3.88	192.11	47000	7400	4800	1300	7400	540
RS-7***	8/26/99	195.99	4.16	191.83	15000	3400	91	950	970	<5
RS-7	11/10/99	195.99	4.12	191.87	10000	2900	170	630	1200	<0.5
RS-7	2/9/00	195.99	3.98	192.01	9400	1400	120	480	600	<0.5
RS-7	6/30/00	195.99	4.04	191.95	8200	3300	190	430	540	<0.5
RS-7	8/8/00	195.99	4.06	191.93	11000	2300	150	430	520	<0.5

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
RS-7	11/16/00	195.99	4.04	191.95	5400	1500	40	240	200	<0.5
RS-7	3/8/01	195.99	3.94	192.05	12000	3300	260	480	850	17 ****
RS-7	5/31/01	195.99	4.01	191.98	10000	1900	120	320	620	<100 ****
RS-7	12/18/01	195.99	4.81	191.18	2700	450	21	86	120	2.3 ****
RS-8	12/14/89									
RS-8	09/04/96									
RS-8	12/11/96									
RS-8	2/21/97									
RS-8	5/28/97									
RS-8	9/2/97									
RS-8	11/24/97									
RS-8	2/25/98									
RS-8	7/8/98									
RS-8	9/16/98									
RS-8	11/24/98									
RS-8	2/23/99									
RS-8	5/5/99									
RS-8***	8/26/99	214.67	7.25	207.42	160000	24000	35000	4200	24000	<5
RS-8	11/10/99	214.67	8.69	205.98	150000	21000	29000	3000	14000	<0.5
RS-8	2/9/00	214.67	7.23	207.44	14000	1900	3200	270	2300	<0.5
RS-8	6/30/00	214.67	3.99	210.68	6400	570	870	150	770	<0.5
RS-8	8/8/00	214.67	7.52	207.15	100000	24000	40000	2300	9900	<0.5 *
RS-8	11/16/00	214.67	6.14	208.53	110000	14000	21000	2100	9600	<20 *
RS-8	3/8/01	214.67	9.40	205.27	10000	740	840	220	990	<2 ****
RS-8	5/31/01	214.67	6.83	207.84	730	11	29	4.2	31	<5 ****
RS-8	12/18/01	214.67	7.14	207.53	4500	230	370	77	750	<0.5 ****
RS-9	12/14/89									

15

TABLE 1
 GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
 DESERT PETROLEUM, INC. SITE #793
 4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
RS-9***	09/04/96									
RS-9***	12/11/96									
RS-9***	2/21/97									
RS-9***	5/28/97									
RS-9***	9/2/97									
RS-9***	11/24/97									
RS-9***	2/25/98									
RS-9***	7/8/98									
RS-9***	9/16/98									
RS-9***	11/24/98									
RS-9***	2/23/99									
RS-9***	5/5/99									
RS-9***	8/26/99	195.63	7.46	188.17	17000	3500	1200	360	1600	180 *
RS-9	11/10/99	195.63	7.91	187.72	2800	520	62	46	130	<0.5
RS-9	2/9/00	195.63	6.09	189.54	3400	650	74	64	130	<0.5
RS-9	6/30/00	195.63	6.77	188.86	3000	600	79	74	120	<0.5
RS-9	8/8/00	195.63	7.32	188.31	4900	500	430	160	530	<0.5
RS-9	11/16/00	195.63	6.33	189.3	3000	350	220	90	220	<0.5
RS-9	3/8/01	195.63	4.93	190.7	<50	3.4	<0.5	<0.5	<0.5	<0.5 ****
RS-9	5/31/01	195.63	4.01	191.62	510	96	6	6.2	9.1	5.5 ****
RS-9	12/18/01	195.63	4.81	190.82	210	11	1.8	3.9	7.6	<0.5 ****
RS-10	12/14/89									
RS-10***	09/04/96									
RS-10***	12/11/96									
RS-10***	2/21/97									
RS-10***	5/28/97									
RS-10***	9/2/97									
RS-10***	11/24/97									

19

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)					
					TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
RS-10***	2/25/98									
RS-10***	7/8/98									
RS-10***	9/16/98									
RS-10***	11/24/98									
RS-10***	2/23/99									
RS-10***	5/5/99									
RS-10***	8/26/99	208.46	3.76	204.7	5100	160	340	190	1000	32 *
RS-10	11/10/99	208.46	3.83	204.63	500	7	2	2	4	<0.5
RS-10	2/9/00	208.46	0.31	208.15	100	4	3	1	6	<0.5
RS-10	6/30/00	208.46	2.22	206.24	640	5	2	4	2	<0.5
RS-10	8/8/00	208.46	2.46	206	460	2	2	2	7	<0.5
RS-10	11/16/00	208.46	2.46	206	360	1	1	2	<1	<0.5
RS-10	3/8/01	208.46	2.82	205.64	53	<0.5	<0.5	<0.5	<0.5	<0.5 ****
RS-10	5/31/01	208.46	4.93	203.53	210	<0.5	<0.5	1.5	5	<5 ****
RS-10	12/18/01	208.46	2.10	206.36	<50	<0.5	<0.5	<0.5	<0.5	<0.5 ****
R1	12/14/89									
R1	09/04/96	227.69	15.00	212.69	1800	1100	3	29	< 10	< 30
R1	12/11/96	227.69	10.30	217.39	<50	<0.5	< 0.5	< 0.5	< 1	4
R1	2/21/97	227.69	11.88	215.81	2500	670	9	3	13	<0.5 *
R1	5/28/97	227.69	14.03	213.66	24000	4300	36	2000	370	<0.5 *
R1	9/2/97	227.69	14.98	212.71	4400	320	6	340	72	20
R1	11/24/97	227.69	14.06	213.63	100	39	1	18	10	<0.5
R1	2/25/98	227.69	8.93	218.76	1200	400	8	13	150	<0.5
R1	7/8/98	227.69	11.36	216.33	68	14	< 0.5	< 0.5	< 1	<1 *
R1	9/16/98	227.69	13.30	214.39	16000	3400	92	< 0.5	410	<1 *
R1	11/24/98	227.69	10.72	216.97	340	19	1.6	35	9.7	<0.5
R1	2/23/99	227.69	9.34	218.35	60	16	0.6	5.6	1.2	<0.5
R1	5/5/99	227.69	11.30	216.39	1300	290	3	150	1	15

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
R1	8/26/99	227.69	13.97	213.72	6500	630	<0.5	1300	<1	<1
R1	11/10/99	227.69	13.73	213.96	480	12	4	22	9	<0.5
R1	2/9/00	227.69	13.10	214.59	<50	8	<0.5	1	<1	<0.5
R1	6/30/00	227.69	13.42	214.27	2600	350	35	1900	220	<0.5
R1	8/8/00	227.69	14.25	213.44	10000	910	76	2100	390	<0.5
R1	3/8/01	227.69	13.72	213.97	<50	<0.5	<0.5	<0.5	<0.5	<0.5
R1	3/8/01	227.69	13.72	213.97	<50	<0.5	<0.5	<0.5	<0.5	<0.5
R1	5/31/01	227.69	15.77	211.92	3800	400	16	470	67	<5
R1	12/18/01	227.69	9.90	217.79	<50	<0.5	<0.5	1.5	<0.5	<0.5
R2	12/14/89									
R2	09/04/96	230.68	13.44	217.24	14000	7600	<10	170	190	<100
R2	12/11/96	230.68	12.42	218.26	488	300	1	< 0.5	30	16
R2	2/21/97	230.68	10.50	220.18	5700	2100	5	2	10	3
R2	5/28/97	230.68	13.10	217.58	36000	14000	63	260	220	<0.5
R2	9/2/97	230.68	14.16	216.52	30000	12000	330	1000	790	47
R2	11/24/97	230.68	14.71	215.97	41000	15000	830	1500	4200	<0.5
R2	2/25/98	230.68	7.39	223.29	800	400	<0.5	<0.5	15	<0.5
R2	7/8/98	230.68	11.27	219.41	290	31	< 0.5	1	< 1	2
R2	9/16/98	230.68	13.73	216.95	6600	11000	24	<0.5	35	<1
R2	11/24/98	230.68	11.67	219.01	6100	<0.5	36	<0.5	21	<0.5
R2	2/23/99	230.68	7.55	223.13	1100	310	3	2	26	<0.5
R2	5/5/99	230.68	10.89	219.79	11000	5300	7	36	7	8
R2	8/26/99	227.28	13.14	214.14	6700	940	33	190	240	<1
R2	11/10/99	227.28	14.42	212.86	5100	2600	160	1800	8100	<0.5
R2	2/9/00	227.28	12.45	214.83	4700	1400	110	130	340	<0.5
R2	6/30/00	227.28	12.94	214.34	7100	3200	110	300	480	<0.5
R2	8/8/00	227.28	13.58	213.7	30000	13000	250	1000	2700	<0.5
R2	11/16/00	227.28	14.33	212.95	44000	17000	230	790	3600	<0.5

13

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TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)												
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)			
R2	3/8/01	227.28	11.15	216.13	2300	640	8.6	61	170	<2	****		
R2	5/31/01	227.28	13.38	213.9	2200	580	12	72	100	<25	****		
R2	12/18/01	227.28	12.35	214.93	4900	2000	120	44	280	<5	****		
R3	12/14/89												
R3	09/04/96	230.32	9.90	220.42	<50	<0.5	<0.5	<0.5	<2	<5			
R3	12/11/96	230.32	8.18	222.14	<50	<0.5	<0.5	<0.5	<1	5			
R3	2/21/97	230.32	6.76	223.56	340	35	59	8	54	<0.5	*		
R3	5/28/97	230.32	9.98	220.34	<50	<0.5	<0.5	<0.5	<1	<0.5	*		
R3	9/2/97	230.32	10.86	219.46	<50	4	<0.5	<0.5	<1	<0.5	*		
R3	11/24/97	230.32	11.20	219.12	not enough water to sample. No sample								
R3	2/25/98	230.32	3.42	226.9	<50	<0.5	<0.5	<0.5	<1	<0.5	*		
R3	7/8/98	230.32	8.78	221.54	140	<0.5	<0.5	4	24	<1	*		
R3	9/16/98	230.32	10.38	219.94	<50	<0.5	<0.5	<0.5	<1	<1	*		
R3	11/24/98	230.32	11.12	219.2	not enough water to sample. No sample								
R3	2/23/99	230.32	3.95	226.37	<50	<0.5	<0.5	<0.5	<1	<0.5	*		
R3	5/5/99	230.32	7.58	222.74	80	9	<0.5	<0.5	<1	6			
R3	8/26/99	227.25	10.76	216.49	<50	2	<0.5	<0.5	<1	1	*		
R3	11/10/99	227.25	11.09	216.16	140	3	4	1	11	<0.5			
R3	2/9/00	227.25	8.76	218.49	<50	2	<0.5	<0.5	<1	<0.5			
R3	6/30/00	227.25	9.67	217.58	<50	0.7	<0.5	1	1	<0.5			
R3	8/8/00	227.25	10.44	216.81	72	<0.5	<0.5	<0.5	<1	<0.5			
R3	11/16/00	227.25	10.26	216.99	110	4	1	<0.5	3	<0.5			
R3	3/8/01	227.25	6.54	220.71	<50	<0.5	<0.5	<0.5	<0.5	<0.5	****		
R3	5/31/01	227.25	10.01	217.24	<50	<0.5	<0.5	<0.5	<0.5	<0.5	****		
R3	12/18/01	227.25	6.79	220.46	<50	<0.5	<0.5	<0.5	<0.5	<0.5	****		
T 1	12/14/89												
T 1	09/04/96												

19

TABLE 1
GROUNDWATER ELEVATIONS AND CERTIFIED ANALYTICAL LABAORATAORY RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per billion [ug/L, ppb]) (AMSL = Above mean sea level)									
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	TPH-G (UG/L)	BENZENE (UG/L)	TOLUENE (UG/L)	ETHYL-BENZENE (UG/L)	XYLENES (UG/L)	MTBE (UG/L)
T 1	12/11/96									
T 1	2/21/97									
T 1	5/28/97									
T 1	9/2/97									
T 1	11/24/97									
T 1	2/25/98									
T 1	7/8/98									
T 1	9/16/98									
T 1	11/24/98									
T 1	2/23/99									
T 1	5/5/99									
T 1***	8/26/99	195.11	2.44	192.67	40000	7200	5000	950	8100	53*
T 1	11/10/99	195.11	2.23	192.88	46000	5600	3600	910	6500	<0.5
T 1	2/9/00	195.11	2.22	192.89	35000	2900	5700	720	6600	<0.5
T 1	6/30/00	195.11	2.22	192.89	30000	3400	3200	950	4600	<5
T 1	8/8/00	195.11	2.73	192.38	8900	1600	760	260	870	<5
T 1	11/16/00	195.11	2.72	192.39	4000	1300	92	80	290	<0.5
T 1	3/8/01	195.11	2.12	192.99	25000	4400	3400	770	3200	26****
T 1	5/31/01	195.11	2.30	192.81	8900	940	210	340	1500	<50****
T 1	12/18/01	195.11	2.20	192.91	48000	3700	5500	1200	5300	24****

ND BELOW LABORATORY DETECTION LIMITS

TPH-G TOTAL PETROLEUM HYDROCARBONS AS GASOLINE

* MTBE results confirmed by EPA Method 8260 (GC/MS)

** LAB REPORT HAD RS-6 AND RS-7 MISLABELED, RESAMPLE ON 7/30/98 CONFIRMED.

*** WELL CASING ELEVATION SURVEY 8-27-99, WADE HAMMOND No.6163, BENCH MARK CITY OF OAKLAND #2814

**** SAMPLES ANALYZED USING EPA METHOD 8260B

TABLE 2
GROUNDWATER ELEVATIONS AND ELECTRON ACCEPTOR RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per million [mg/L, ppm] unless otherwise noted) (AMSL = Above mean sea level)															
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	FIELD MEASUREMENTS						CERTIFIED LABORATORY RESULTS DISSOLVED IN WATER					
					DISSOLVED OXYGEN O2 (MG/L)	SULFATE SO4 (MG/L)	NITRATE NO3 (MG/L)	FERROUS IRON FE2 (MG/L)	TEMP-ERATURE (F)	PH	TOTAL PETROLEUM HYDROCARBONS GASOLINE (MG/L)	CARBON DI OXIDE CO2 (MG/L)	METHANE CH4 (MG/L)	AEROBIC HYDROCARBON DEGRADING BACTERIA CFU/ML	ORTHO-PHOSPHATE PO4 (MG/L)	AMMONIA as NITROGEN N (MG/L)
MW-1	8/26/99	229.57	11.41	218.16	4.9	35	0	0.25	75.4	6.55	<0.05	0.13	<0.00001	10	<1	<0.5
	9/2/99	229.57	11.65	217.92					72.9	8.16						
	3/8/01	229.57	12.30	217.27	4.9				67.6	7.33	<0.05					
	12/18/01	229.57	13.74	215.83	4.4	61	7.6	0	67.1	7.63	<0.05					
RS-2	8/26/99	227.39	11.42	215.97	0.7	46	2.7	0.65	80.9	6.97	0.2	nm	nm	nm	nm	nm
	9/2/99	227.39	12.00	215.39					72.9	8.16						
	12/18/01	227.39	6.99	220.4	4.6	>77	11.4	0.07	67.6	7.75	<0.05					
RS-5	8/26/99	227.61	16.06	211.55	0.7	31	1.3	0.92	71.7	7.08	35					
	9/2/99	227.61	16.26	211.35					68.4	7.15		0.16	0.00021	3000	<1	<0.5
	3/8/01	227.61	27.72	199.89	3.1				59.7	7.46	11					
	12/18/01	227.61	15.61	212	1.4	37	8.2	>3.3	66.6	6.83	12					
RS-6	8/26/99	227.22	13.72	213.5	1.2	76	0.3	>3.3	77.8	6.66	0.69					
	9/2/99	227.22	14.14	213.08					69	6.69		0.36	<0.00001	400	<1	<0.5
	12/18/01	227.22	10.88	216.34	4.3	>77	0	0	66.7	6.84	0.056					
RS-7	8/26/99	195.99	4.16	191.83	0.3	>77	0.8	1.27	73.4	6.99	15	nm	nm	nm	nm	nm
	9/2/99	195.99	4.14	191.95					68.4	7.15						
	12/18/01	195.99	4.81	191.18	2.5	1	6	0.87	68.1	6.82	2.7					
RS-8	8/26/99	214.67	7.25	207.42	2.6	0	0	0.54	69.2	6.7	160					
	9/2/99	214.67	7.38	207.29					71.7	5.74		0.058	0.000018	6600	<1	<0.5
	3/8/01	214.67	9.40	205.27	2.2				63.3	6.97	10					
	12/18/01	214.67	7.14	207.53	4.2	49	9.2	0.08	67.3	6.98	0.23					
RS-9	8/26/99	195.63	7.46	188.17	2.1	7	0	0.59	73.5	6.95	17					
	9/2/99	195.63	7.61	188.02					70.9	6.98		0.25	0.0021	10000	<1	<0.5
	3/8/01	195.63	4.93	190.7	8.1				62.7	6.89	<0.05					
	12/18/01	195.63	4.81	190.82	WATER TO CLOUDY, LIGHT GREY				68.3	6.8	0.21					
RS-10	8/26/99	208.46	3.76	204.7	4.2	nm	nm	nm	70.9	8.03	5.1					
	9/2/99	208.46	3.96	204.5					73.3	7.24		0.1	0.000037	8800	<1	<0.5
	3/8/01	208.46	2.82	205.64	3.5				61.5	6.16	0.053					
	12/18/01	208.46	2.10	206.36	4.3	46	4.1	0	66.9	6.54	<0.05					
R1	8/26/99	227.69	13.97	213.72	0.4	9	0	>3.3	70.6	6.38	6.5	nm	nm	nm	nm	nm
	9/2/99	227.69	14.18	213.51					66.4	7.24	<0.05					
	12/18/01	227.69	9.90	217.79	5.2	14	4.2	0								
R2	8/26/99	227.28	13.14	214.14	0.4	>77	0.8	0.3	72.7	6.65	6.7	nm	nm	nm	nm	nm
	9/2/99	227.28	13.23	214.05					66.5	6.69	4.9					
	12/18/01	227.28	12.35	214.93	2.8	>77	1.3	0.07								
R3	8/26/99	230.32	10.76	219.56	2.5	>77	0.7	0.05	75	6.95	<0.05	nm	nm	nm	nm	nm
	9/2/99	230.32	10.87	219.45					67.1	6.91	<0.05					
	12/18/01	230.32	6.79	223.53	5.5	>77	6.2	0								

21

TABLE 2
GROUNDWATER ELEVATIONS AND ELECTRON ACCEPTOR RESULTS FROM WATER SAMPLES
DESERT PETROLEUM, INC. SITE #793
4035 PARK BOULEVARD, OAKLAND, CALIFORNIA

ID#	(All concentrations in parts per million [mg/L, ppm] unless otherwise noted) (AMSL = Above mean sea level)																
	DATE SAMPLED	WELL CASING ELEVATION (FEET AMSL)	DEPTH TO GROUND WATER (FEET)	GROUND WATER ELEVATION (FEET AMSL)	FIELD MEASUREMENTS						CERTIFIED LABORATORY RESULTS DISSOLVED IN WATER						
					DISSOLVED OXYGEN O2 (MG/L)	SULFATE SO4 (MG/L)	NITRATE NO3 (MG/L)	FERROUS IRON FE2 (MG/L)	TEMP-ERATURE (F)	pH	TOTAL PETROLEUM HYDROCARBONS GASOLINE (MG/L)	CARBON DI OXIDE CO2 (MG/L)	METHANE CH4 (MG/L)	AEROBIC HYDROCARBON DEGRADING BACTERIA CFU/ML	ORTHO-PHOSPHATE PO4 (MG/L)	AMMONIA as NITROGEN N (MG/L)	
T 1	8/26/99	195.11	2.44	192.67	0.8	32	0.5	0.03	75.3	7.29	40						
	9/2/99	195.11	2.20	192.91					78.1	7.57		0.11	0.00019	1300	<1	<0.5	
	3/8/01	195.11	2.18	192.93	3.1						25						
	12/18/01	195.11	2.20	192.91	2.8	0	4.3	0.6	66.3	6.52	48						
T 2	8/26/99	195.3	CAR		nm	nm	nm	nm	nm	nm	NA						
	9/2/99	195.3	CAR									nm	nm	nm	nm	nm	
T 3	8/26/99	202.38	CAR		nm	nm	nm	nm	nm	nm	NA						
	9/2/99	202.38	CAR									nm	nm	nm	nm	nm	
T 4	8/26/99	197.48	CAR		nm	nm	nm	nm	nm	nm	NA						
	9/2/99	197.48	CAR									nm	nm	nm	nm	nm	
LF-1	8/26/99	226.59	CAR		nm	nm	nm	nm	nm	nm	NA						
	9/2/99	226.59	CAR									nm	nm	nm	nm	nm	

NA NOT ANALYZED
nm NOT MEASURED
CAR CAR PARKED OVER WELL, NO ACCESS
MG/L milligrams per liter (ppm)
F degrees Fahrenheit
CFU/ML colony forming units per milli:
< below laboratory lower detection limits.
AMSL ABOVE MEAN SEA LEVEL

TABLE 3
 WASTEWATER DISCHARGE PERMIT # 5043550 1
 FORMER DP #793
 4035 PARK BLVD., OAKLAND, CALIFORNIA

WASTEWATER SOURCE ID	DATE	METER READING	NEW METER	GALLONS DISCHARGED BETWEEN VISITS	ACCUMULATIVE GALLONS DISCHARGED	AVERAGE DISCHARGE PER MINUTE IN GALLONS	EPA METHOD 624				7420 LEAD ug/L	
		IN GALLONS #35635668	IN GALLONS #47083426				BENZENE ug/L	TOLUENE ug/L	ETHYL-BENZENE ug/L	XYLENES ug/L		
		314110										
BAKER TANK	1/25/00	314110		0	0	0.00						
BAKER TANK	1/26/00	315050		940	940	0.65	<1	<1	<1	<1	<50	
BAKER TANK	1/28/00	321120	1098330	6070	7010	2.11						
BAKER TANK	2/2/00		1102560	4230	11240	0.59						
BAKER TANK	2/3/00		1107482.2	4922	16162	3.42	<1	<1	<1	<1	<50	
BAKER TANK	2/7/00		1107482.2	0	16162	0.00						
BAKER TANK AND 1/4LY SAMPLES	2/9/00		1109680	2198	18360	0.78	EPA METHOD 624				239.2	
F1 (PSP No. 1)	3/23/00		1109720	40	18400	0.00	<1	<1	<1	<2	<5	
F1 (PSP No. 1)	5/4/00		1110780	1060	19460	0.02						
F1 (PSP No. 1)	5/12/00		1111700	920	20380	0.08						
F1 (PSP No. 1)	5/18/00		1113359	1659	22039	0.19						
F1 (PSP No. 1)	5/25/00		1113840	481	22520	0.05						
F1 (PSP No. 1)	5/31/00		1115111	1271	23791	0.15						
F1 (PSP No. 1)	6/16/00		1115823	712	24503	0.03						
F1 (PSP No. 1)	6/28/00		1116293	470	24973	0.03						
F1 (PSP No. 1)	6/30/00		1116303	10	24983	0.00	EPA METHOD 624				200.7	
F1 (PSP No. 1)	7/5/00		1116313	10	24993	0.00	<1	<1	<1	<2	<2	
F1 (PSP No. 1)	7/13/00		1117816	1503	26496	0.13						
F1 (PSP No. 1)	7/20/00		1118892	1076	27572	0.11						
F1 (PSP No. 1)	7/27/00		1118892	0	27572	0.00						
F1 (PSP No. 1)	8/3/00		1120336	1444	29016	0.14						
F1 (PSP No. 1)	8/10/00		1121041	705	29721	0.07						
F1 (PSP No. 1)	8/17/00		1121041	0	29721	0.00						
F1 (PSP No. 1)	8/24/00		1121860	819	30540	0.08	EPA METHOD 624				200.7	
F1 (PSP No. 1)	8/30/00		1122720	860	31400	0.10	<1	<2	<1	<2	<2	
F1 (PSP No. 1)	9/7/00		1123270	550	31950	0.05						
F1 (PSP No. 1)	9/14/00		1123819	549	32499	0.05						
F1 (PSP No. 1)	9/21/00		1123819	0	32499	0.00						
F1 (PSP No. 1)	10/5/00		1124153	334	32833	0.02						
F1 (PSP No. 1)	10/12/00		1124660	507	33340	0.05						
F1 (PSP No. 1)	10/19/00		1125904.3	1244	34584	0.12						
F1 (PSP No. 1)	10/26/00		1127167	1263	35847	0.13						
F1 (PSP No. 1)	11/9/00		1128367.2	1200	37047	0.06						
F1 (PSP No. 1)	11/16/00		1129779.5	1412	38460	0.14						
F1 (PSP No. 1)	11/22/00		1130940.5	1161	39621	0.13						
F1 (PSP No. 1)	12/1/00		1134147	3207	42827	0.25	EPA METHOD 624				200.7	

23

TABLE 3
 WASTEWATER DISCHARGE PERMIT # 5043550 1
 FORMER DP #793
 4035 PARK BLVD., OAKLAND, CALIFORNIA

WASTEWATER SOURCE ID	DATE	METER READING	NEW METER	GALLONS DISCHARGED	ACCUMULATIVE GALLONS DISCHARGED	AVERAGE DISCHARGE PER MINUTE IN GALLONS	EPA METHOD 624		ETHYL-BENZENE	XYLENES	7420 LEAD
		IN GALLONS #35635668	IN GALLONS #47083426	BETWEEN VISITS			ug/L	ug/L	ug/L	ug/L	ug/L
		314110									
F1 (PSP No. 1)	12/7/00		1134289	142	42969	0.02	<1	<1	<1	<2	<2
F1 (PSP No. 1)	12/14/00		1134431	142	43111	0.01					
F1 (PSP No. 1)	12/21/00		1134573	142	43253	0.01					
F1 (PSP No. 1)	12/28/00		1134714.8	142	43395	0.01					
F1 (PSP No. 1)	1/11/01		1134714.8	0	43395	0.00	no discharge, could not access trench well				
F1 (PSP No. 1)	1/18/01		1135243.8	529	43924	0.05					
F1 (PSP No. 1)	1/25/01		1136144	900	44824	0.09					200.7
F1 (PSP No. 1)	2/8/01		1136659	515	45339	0.03					<2
F1 (PSP No. 1)	2/15/01		1137441.4	782	46121	0.08					
F1 (PSP No. 1)	2/22/01		1141123.6	3682	49804	0.37	start discharge from RS5				
F1 (PSP No. 1)	3/1/01		1150736.5	9613	59417	0.95	EPA METHOD 624				
F1 (PSP No. 1)	3/8/01		1158901.1	8165	67581	0.81	<1	<1	<1	<2	
F1 (PSP No. 1)	3/14/01		1162321.2	3420	71001	0.40					
F1 (PSP No. 1)	3/21/01		1162321.4	0	71001	0.00	no discharge, pump removed for repair				
F1 (PSP No. 1)	4/4/01		1163471.7	1150	72152	0.06					
F1 (PSP No. 1)	4/12/01		1164723.5	1252	73404	0.11	EPA METHOD 8260B				
F1 (PSP No. 1)	4/19/01		1173267	8544	81947	0.85	<0.5	<0.5	<0.5	<0.5	
F1 (PSP No. 1)	5/3/01		1181423.5	8157	90104	0.40					
F1 (PSP No. 1)	5/10/01		1188209.3	6786	96889	0.67					
F1 (PSP No. 1)	5/16/01		1189899.1	1690	98579	0.20					
F1 (PSP No. 1)	5/24/01		1198018.4	8119	106698	0.70					
F1 (PSP No. 1)	5/31/01		1199647.8	1629	108328	0.16					
F1 (PSP No. 1)	6/6/01		1204217.2	4569	112897	0.53					
F1 (PSP No. 1)	6/14/01		1210661.4	6444	119341	0.56					
F1 (PSP No. 1)	6/21/01		1214600	3939	123280	0.39					
F1 (PSP No. 1)	6/28/01		1219387.7	4788	128068	0.47					
F1 (PSP No. 1)	7/5/01		1223625.4	4238	132305	0.42					
F1 (PSP No. 1)	7/12/01		1228500	4875	137180	0.48	EPA METHOD 8260B				
F1 (PSP No. 1)	7/19/01		1232750.7	4251	141431	0.42	<0.5	<0.5	<0.5	<0.5	
REMOVE PUMP AND DISCONTINUE SEWER DISCHARGE ON July 19, 2001. COMMENCE 1/4LY DISCHARGE											
F1 (PSP No. 1)	12/18/01			238	141669	5.00	<0.5	<0.5	<0.5	<0.5	MTBE <0.5
< BELOW LABORATORY LOWER DETECTION LIMITS											
ug/L. micrograms per liter (parts per billion)											

Note: water meter #47083426 did not function during initial test, substitute meter #35635668 used until cleaned and tested. Re-installed January 28, 2000.
 WATER DISCHARGED TO SEWER IS FROM WEEKLY PURGING OF T1, CONTINUOUS DISCHARGE FROM WELL RS5 AND PURGED WATER FROM 1/4LY SAMPLING.

24

TABLE 4
 RECEPTOR TRENCH GROUNDWATER REMOVAL
 FORMER DP #793
 4035 PARK BLVD., OAKLAND, CALIFORNIA

RECEPTOR TRENCH WATER ANALYSIS
 EPA METHOD 8020

PURGING BY	DATE PURGED	METER READING IN GALLONS RS5	METER READING IN GALLONS TRENCH	DEPTH TO TOP OF WATER IN FEET	GALLONS T1	ACCUMULATED GALLONS REMOVED FROM TRENCH GALLONS	Accumulated gallons removed from RS5 Gallons	TOTAL GALLON: REMOVED	RECEPTOR TRENCH WATER ANALYSIS						
									TPHg	BENZENE ug/L	TOLUENE ug/L	ETHYL-BENZENE ug/L	XYLENES ug/L	MTBE ug/L	
WEGE	8/9/99			6.47	200	200									
WEGE	8/10/99			5.02	1730	1930									
WEGE	8/11/99			7.89	960	2890									
WEGE	8/12/99			8.12	800	3690									
WEGE	8/13/99			8.87	600	4290									
WEGE	9/2/99			2.2	3600	7890			40000	7200	5000	950	8100	53	
WEGE	9/16/99			2.27	5131	13021									
WEGE	9/23/99			4.26	3351	16372									
WEGE	9/30/99			4.69	1734	18106									
WEGE	10/7/99			4.78	293	18400									
WEGE	1/25/00				0	18400									
WEGE	1/26/00				0	18400									
WEGE	1/28/00		1098330.0		0	18400									
WEGE	2/23/00		1102560.0		0	18400			35000	2900	5700	720	6600	<0.5	
WEGE	2/29/00		1109680.0	2.22	0	18400									
WEGE	3/23/00		1109720.0		0	18400				1020	6500	1010	5090		
WEGE	5/4/00		1110780.0		1060	19460									
WEGE	5/12/00		1111700.0	2.19	920	20380									
WEGE	5/18/00		1113359.0	2.18	1659	22039									
WEGE	5/25/00		1113840.0		481	22520									
WEGE	5/31/00		1115111.0	2.15	1271	23791									
WEGE	6/18/00		1115823.0		712	24503									
WEGE	6/28/00		1116293.0	2.22	470	24973									
WEGE	6/30/00		1116303.0		10	24983			30000	3400	3200	950	4600	<5	
WEGE	7/5/00		1116313.0		10	24993									
WEGE	7/6/00		1116313.0		0	24993									
WEGE	7/13/00		1117816.0		1503	26496									
WEGE	7/20/00		1118892.0	2.29	1076	27572									
WEGE	7/27/00		1118892.0	2.21	0	27572									
WEGE	8/3/00		1120336.0	2.9	1444	29016									
WEGE	8/10/00		1121041.0	2.75	705	29721			8900	1600	760	260	870	<5	
WEGE	8/17/00		1121041.0	2.73	0	29721									
WEGE	8/24/00		1121860.0	2.75	819	30540									
WEGE	8/30/00		1122720.0	2.75	660	31400									
WEGE	9/7/00		1123270.0	2.78	550	31950									
WEGE	9/14/00		1123810.0	2.79	540	32490									
WEGE	9/21/00		1123810.0		0	32490									
WEGE	10/5/00		1124253.0	2.81	443	32933									
WEGE	10/12/00		1124660.0	2.4	407	33340									
WEGE	10/19/00		1125904.3			1244	34584								
WEGE	10/26/00		1127167.0	2.22	1263	35847									

TABLE 5 POUNDS HYDROCARBONS

DP 793

4035 PARK BLVD, OAKLAND, CA

Pounds TPHg in soil and in groundwater AUGUST 1999.

Square Feet	Thickness	Cubic feet	Upper mg/kg	Lower mg/kg	Average con mg/kg	kg Soil	mg TPHg	pounds TPHg
		soil density=	1.9					
TPHg in soil 5 - 10 foot depth 8/99								
2165	5	10825	50	1	25.5	582409.9	14851452.39	32.7
1945	5	9725	100	50	75	523227.4	39242052.56	86.5
700	5	3500	290	100	195	188308.1	36720069.75	81.0
325	5	1625	1000	100	550	87428.74	48085805.63	106.0
100	5	500	2000	1000	1500	26901.15	40351725	89.0
440	5	2200	4000	1000	2500	118365.1	295912650	652.4
TPHg in soil 10-15 foot depth AUGUST 1999								
5006	5	25030	50	1	25.5	1346672	34340125.01	75.7
1599	5	7995	100	50	75	430149.4	32261204.14	71.1
2815	5	14075	1000	100	550	757267.4	416497054.9	918.2
240	5	1200	1200	1000	1100	64562.76	71019036	156.6
TPHg in soil 15-20 foot depth AUGUST 1999								
1926	5	9630	50	1	25.5	518116.1	13211961.8	29.1
1044	5	5220	100	50	75	280848	21063600.45	46.4
1250	5	6250	1000	100	550	336264.4	184945406.3	407.7
TPHg in soil 20-25 foot depth AUGUST 1999								
2900	5	14500	10	1	5.5	780133.4	4290733.425	9.5
TPHg in soil 25-30 foot depth AUGUST 1999								
410	5	2050	50	1	25.5	110294.7	2812515.233	6.2
175	5	875	100	50	75	47077.01	3530775.938	7.8
145	5	725	1000	100	550	39006.67	21453667.13	47.3
80	5	400	1600	1000	1300	21520.92	27977196	61.7
Total pounds TPHg in Soil AUGUST 1999								2884.9

TABLE 5 POUNDS HYDROCARBONS

DP 793
4035 PARK BLVD, OAKLAND, CA

MASS GROUNDWATER CONTAMINATION AUGUST 1999									
Square Feet	Thickness	Cubic feet	PORES	Upper mg/l	Lower mg/l	Average con mg/l	LITERS water	mg TPHg	pounds TPHg
			0.3						
10520	16	168320		1	0.05	0.525	1429895	750695	1.66
8650	16	138400		10	1	5.5	1175722	6466470	14.26
2000	16	32000		40	10	25	271843	6796080	14.98
2550	16	40800		100	10	55	346600	19063004	42.03
750	16	12000		160	100	130	101941	13252356	29.22
Total Calculated Mass TPHg in Groundwater AUGUST 1999									102.14

MASS GROUNDWATER CONTAMINATION DECEMBER 2001									
Square Feet	Thickness	Cubic feet	PORES	Upper mg/l	Lower mg/l	Average con mg/l	LITERS water	mg TPHg	pounds TPHg
			0.3						
4825	16	77200		1	0.05	0.525	655822	344306	0.76
4984	16	79744		10	1	5.5	677433	3725883	8.21
120	16	1920		12	10	11	16311	179417	0.40
1240	16	19840		48	10	29	168543	4887741	10.78
Total Calculated Mass TPHg in Groundwater DECEMBER 2001									20.14

MASS BENZENE GROUNDWATER CONTAMINATION AUGUST 1999									
assumption: free product contains 287 mg/L of benzene									
Square Feet	Thickness	Cubic feet	PORES	Upper mg/l	Lower mg/l	Average con mg/l	LITERS water	mg Benzene	pounds Benzene
			0.3						
6775	16	108400		0.1	0.0005	0.05025	920869	46274	0.10
7800	16	124800		1	0.1	0.55	1060188	583104	1.29
2000	16	32000		7.2	1	4.1	271843	1114557	2.46
2000	16	32000		10	1	5.5	271843	1495138	3.30
1200	16	19200		24	10	17	163106	2772801	6.11
Total Calculated Mass Benzene in Groundwater 1999									13.25

MASS BENZENE GROUNDWATER CONTAMINATION DECEMBER 2001									
assumption: free product contains 287 mg/L of benzene									
Square Feet	Thickness	Cubic feet	PORES	Upper mg/l	Lower mg/l	Average con mg/l	LITERS water	mg Benzene	pounds Benzene
			0.3						
3940	16	63040		0.001	0.0005	0.00075	535531	402	0.00
5265	16	84240		0.5	0.001	0.2505	715627	179265	0.40
2170	16	34720		1	0.5	0.75	294950	221212	0.49
450	16	7200		2	1	1.5	61165	91747	0.20
970	16	15520		3.7	1	2.35	131844	309833	0.68
Total Calculated Mass Benzene in Groundwater 2001									1.77

ESTIMATED POUNDS GROUNDWATER TPHg REDUCTION FROM 1999 TO 2001 82.0
 ESTIMATED POUNDS GROUNDWATER BENZENE REDUCTION FROM 1999 TO 2001 11.5

ESTIMATED % REDUCTION TPHg 80.3
 ESTIMATED % REDUCTION BENZENE 86.7

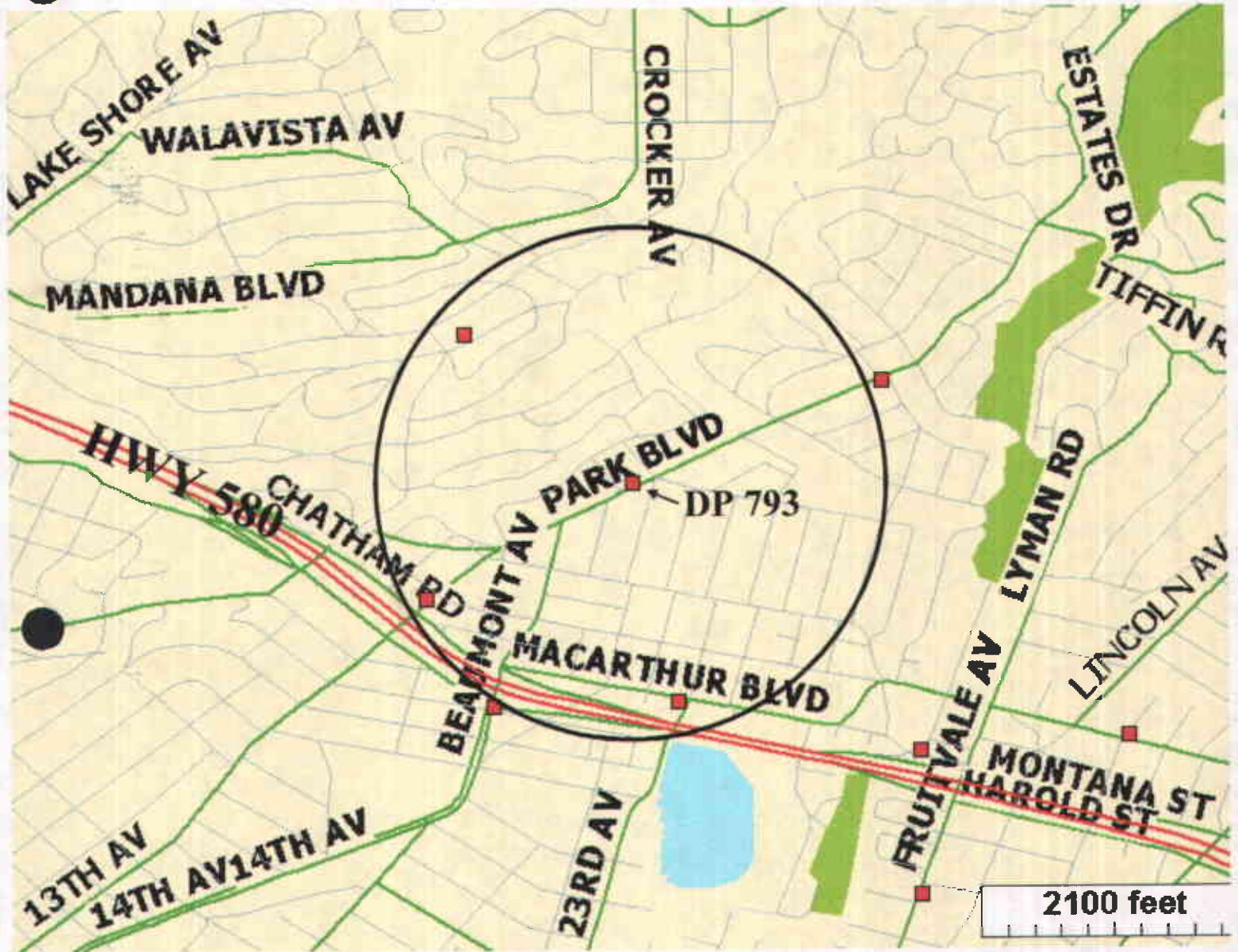


FIGURE 1
 GEOTRACKER
 AREA WELL & LUST MAP
 DP 793
 4035 PARK BLVD.
 OAKLAND, CA

- LUST SITES
- WELLS

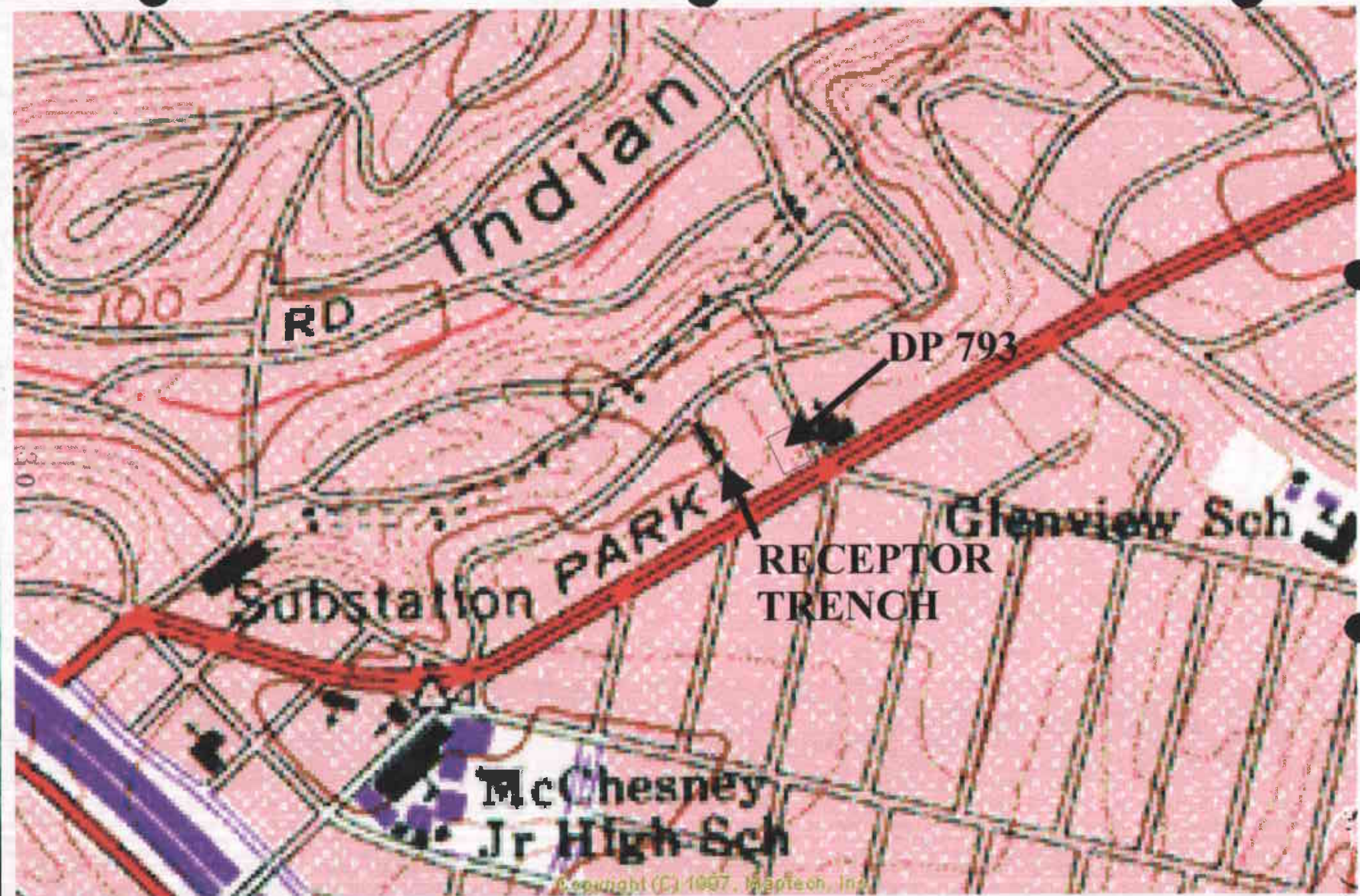
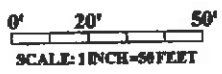
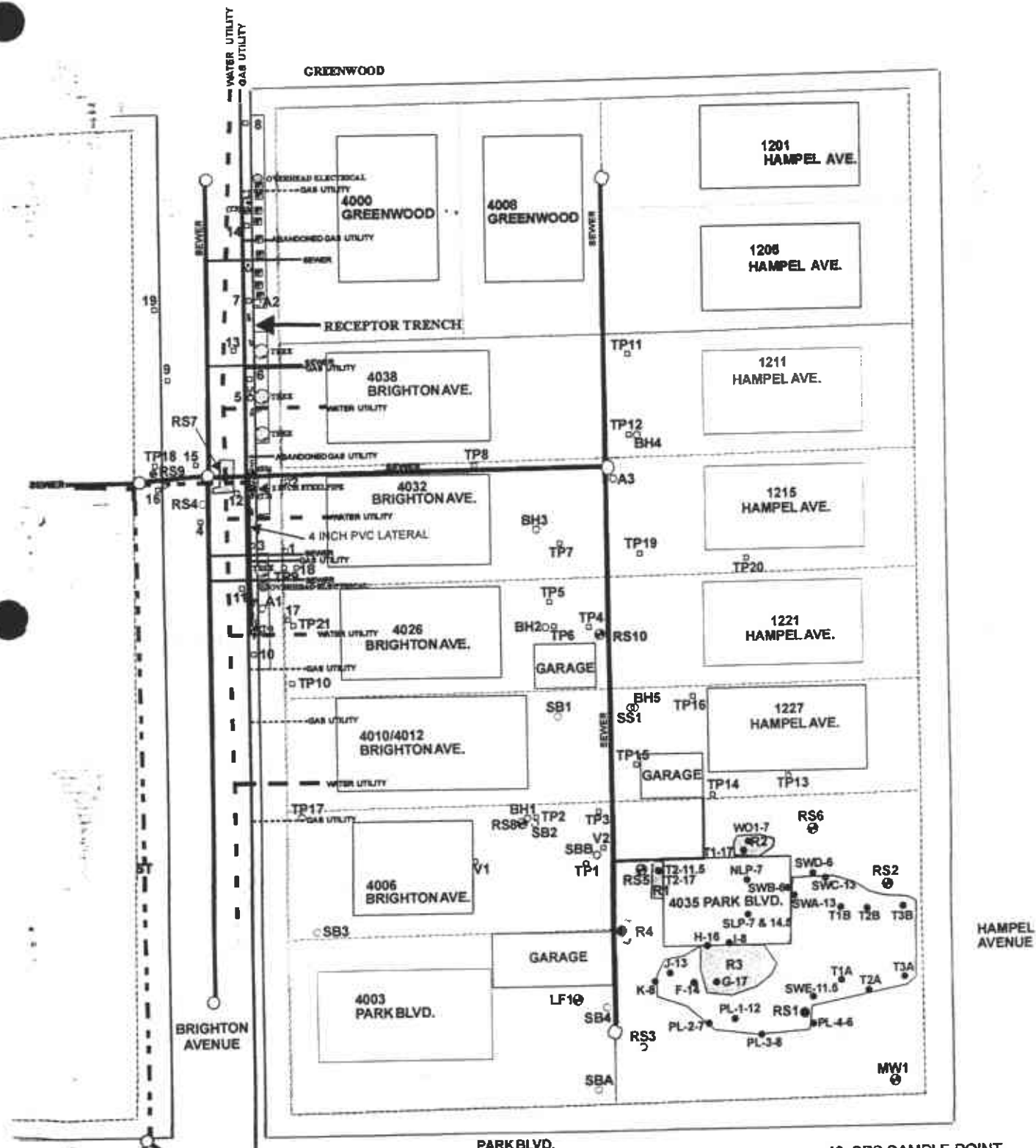


FIGURE 2
PORTION OF OAKLAND EAST 7.5 MINUTE USGS TOPOGRAPHIC MAP





**FIGURE 3-SAMPLE LOCATIONS
SEWER AND FREE PRODUCT
INVESTIGATION FOR
DP793, 4035 PARK BLVD.
OAKLAND, CALIFORNIA**

- TP10 SPS SAMPLE POINT
- SOIL SAMPLE POINT
- SOIL BORING
- ⊙ RECEPTOR TRENCH SAMPLE POINT
- ⊙ RS2 GROUNDWATER MONITORING WELL
- ⊙ RS1● DESTROYED MONITORING WELL

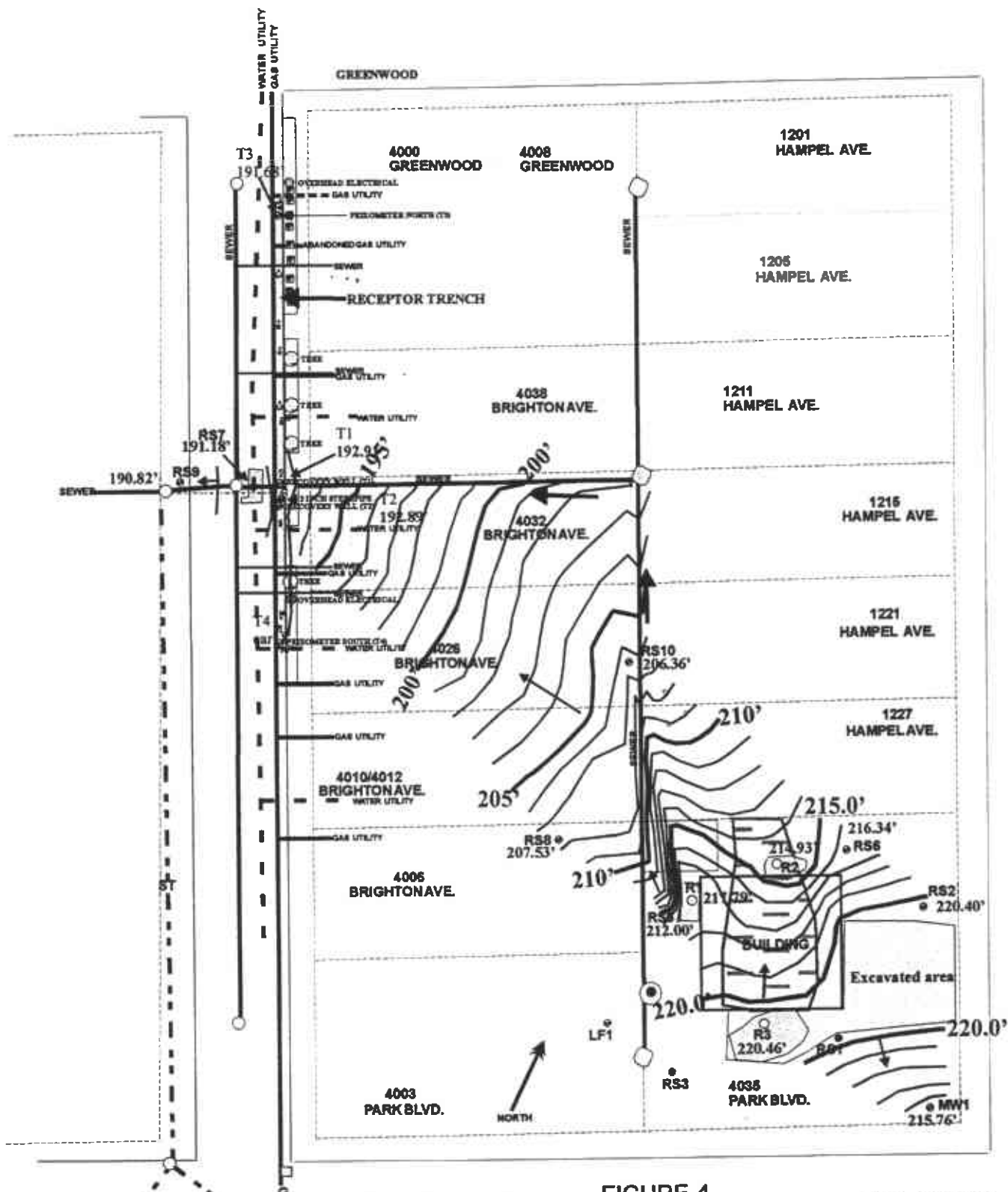
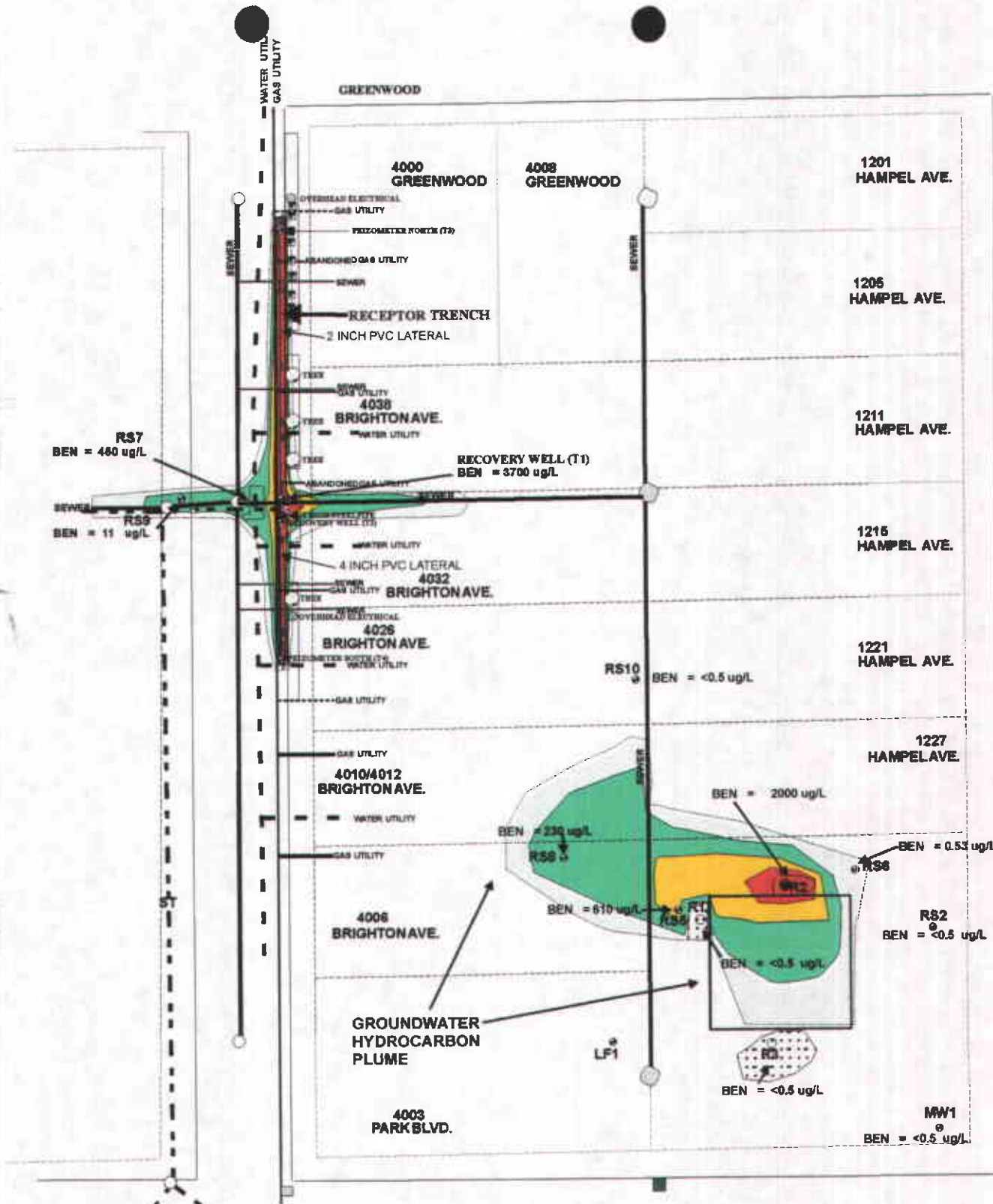


FIGURE 4
 DP 793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA
 GROUNDWATER ELEVATION
 12/18/01.

CONTOURS ARE
 FEET ABOVE SEA
 LEVEL



0' 20' 50'
SCALE: 1 INCH=50 FEET

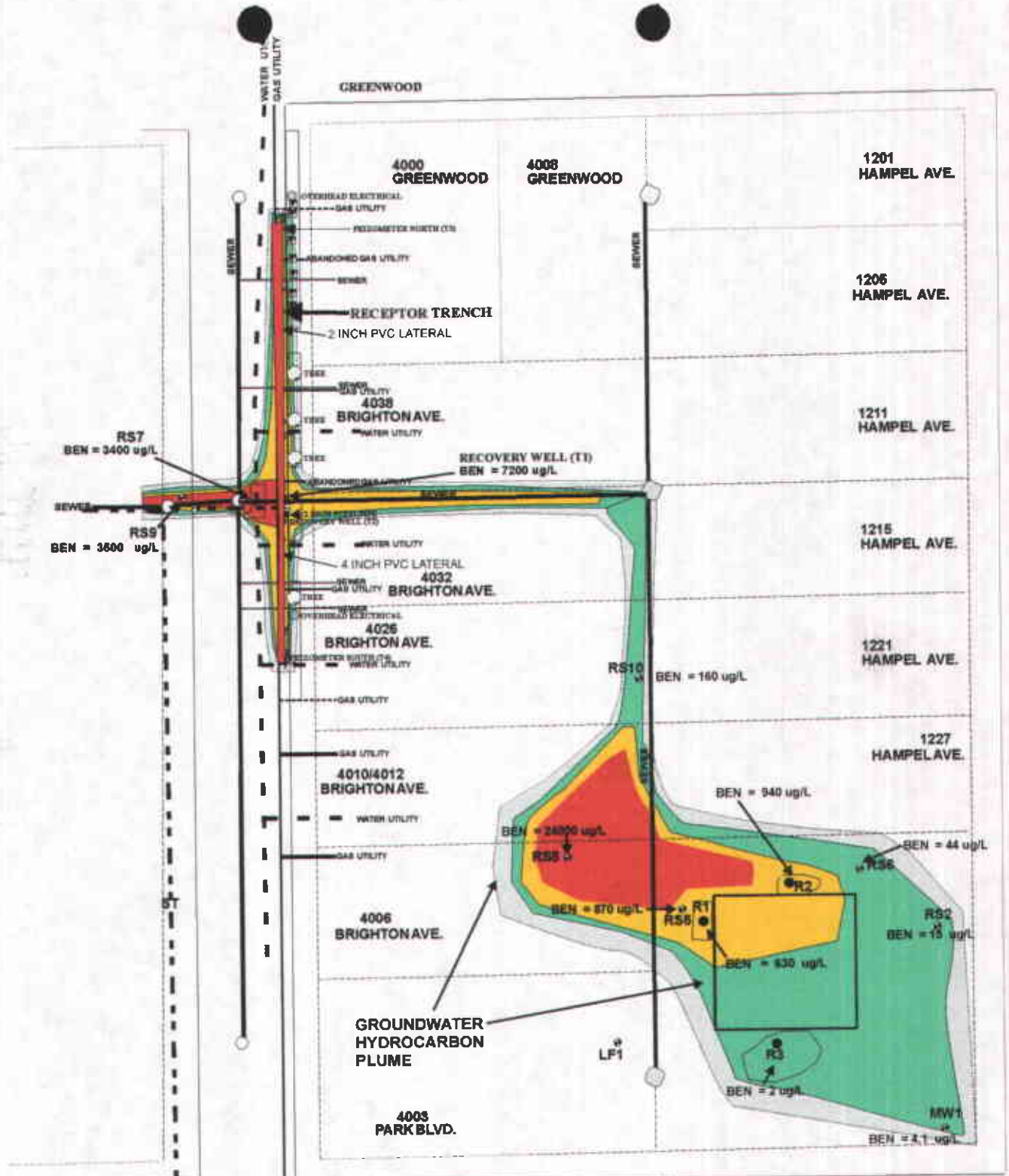


**FIGURE 5
GROUNDWATER
PLUME**

DP 793, 4035 PARK BLVD.
OAKLAND, CALIFORNIA

- RS3 SOIL BORING
- ⊥ TRENCH SAMPLE POINT
- RS2 GROUNDWATER MONITORING WELL
- Benzene > 1000 ug/L
- Benzene > 500 ug/L
- Benzene > 1 ug/L
- TPHg Groundwater Plume

12/01



0' 20' 50'
SCALE: 1 INCH = 20 FEET



FIGURE 6
GROUNDWATER
PLUME
08/26/99

DP 793, 4035 PARK BLVD.
OAKLAND, CALIFORNIA

8/99

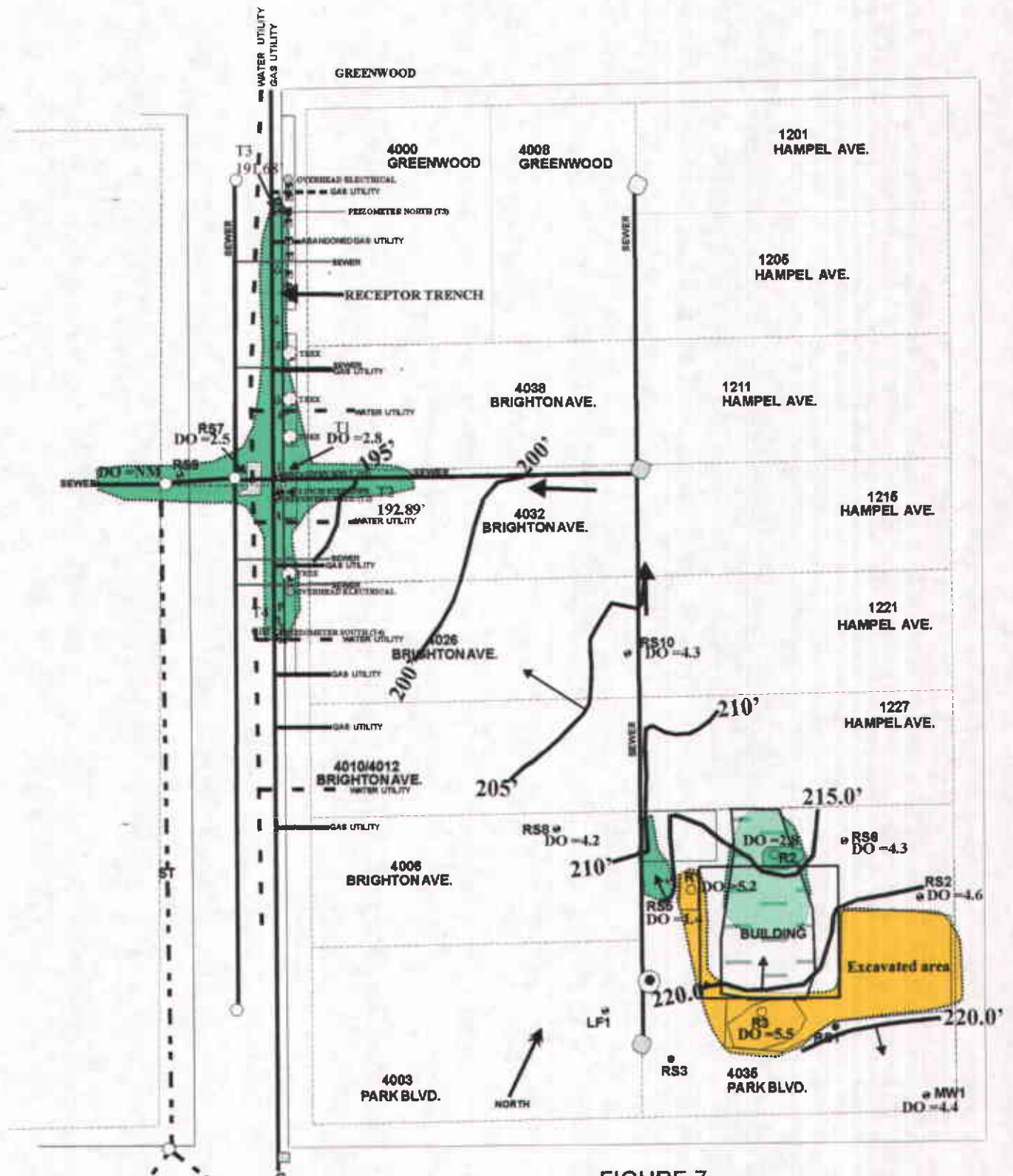




FIGURE 7
 DP 793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA
 GROUNDWATER ELEVATION
 WITH DISSOLVED OXYGEN
 12/18/01.

CONTOURS ARE FEET ABOVE SEA LEVEL

 DO < 3 mg/L

 DO > 5 mg/L

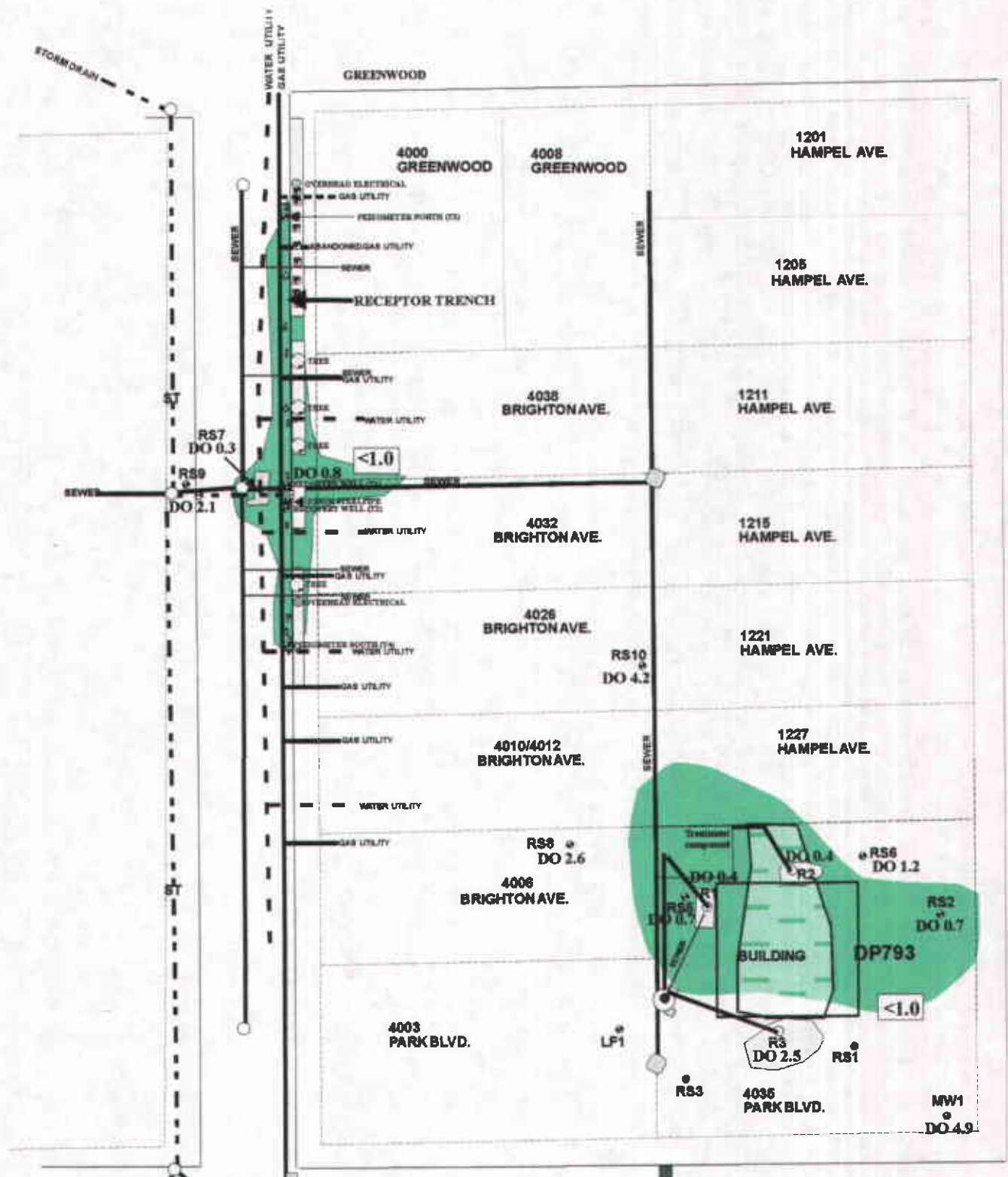


FIGURE 7-1
 DP 793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA
 TOTAL DISSOLVED OXYGEN
 IN GROUNDWATER 8/26/99.

DO 4.9 DISSOLVED OXYGEN
 IN PARTS PER MILLION

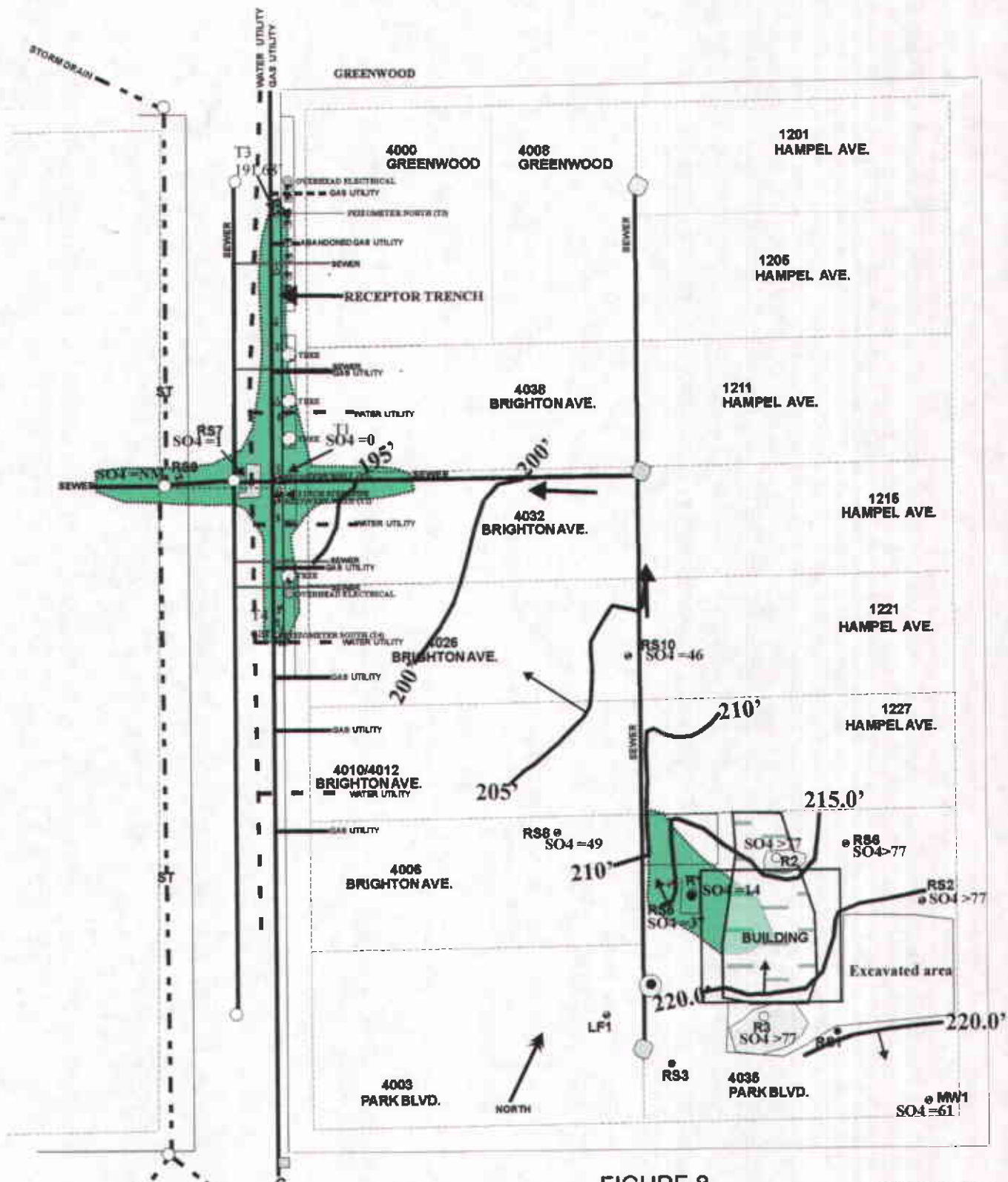



FIGURE 8
 DP 793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA
 GROUNDWATER ELEVATION
 WITH SULFATE
 12/18/01.

CONTOURS ARE
 FEET ABOVE SEA
 LEVEL

 SO4 < 40 mg/L

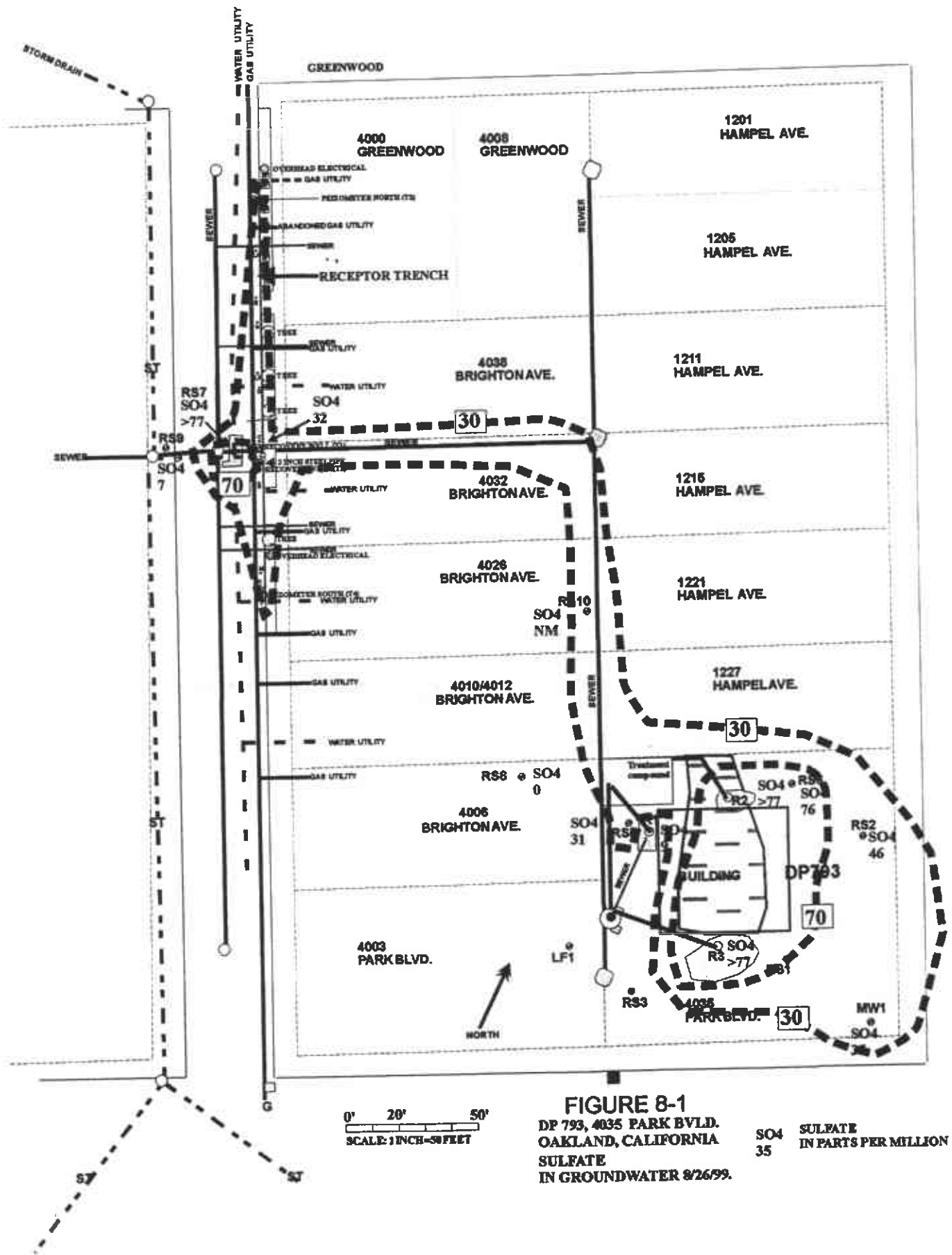


FIGURE 8-1
 DP 793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA
 SULFATE
 IN GROUNDWATER 8/26/99.

SO4 Sulfate
 35 IN PARTS PER MILLION

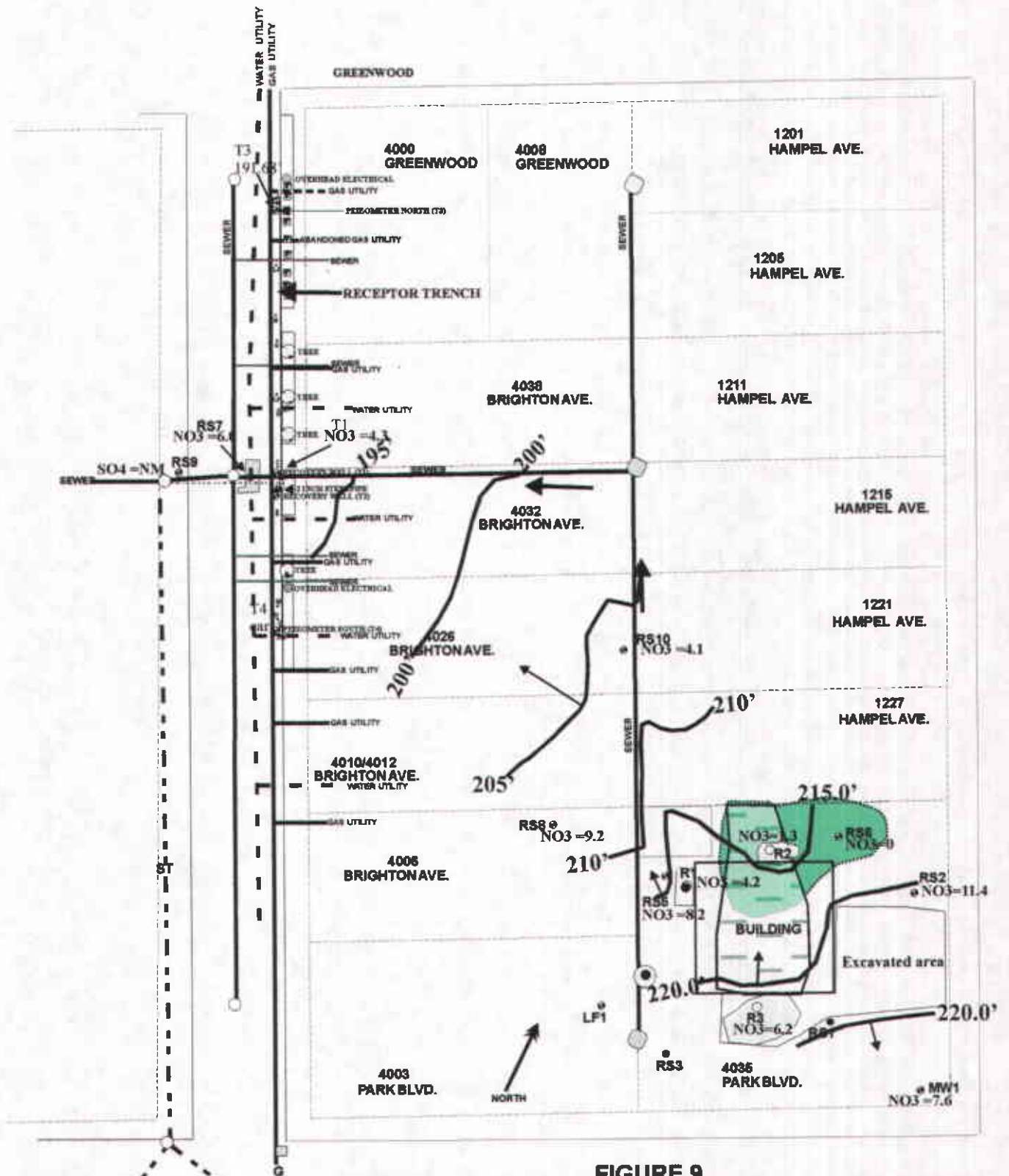



FIGURE 9
 DP 793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA
 GROUNDWATER ELEVATION
 WITH NITRATE
 12/18/01.

CONTOURS ARE
 FEET ABOVE SEA
 LEVEL

 $NO_3 < 2 \text{ mg/L}$

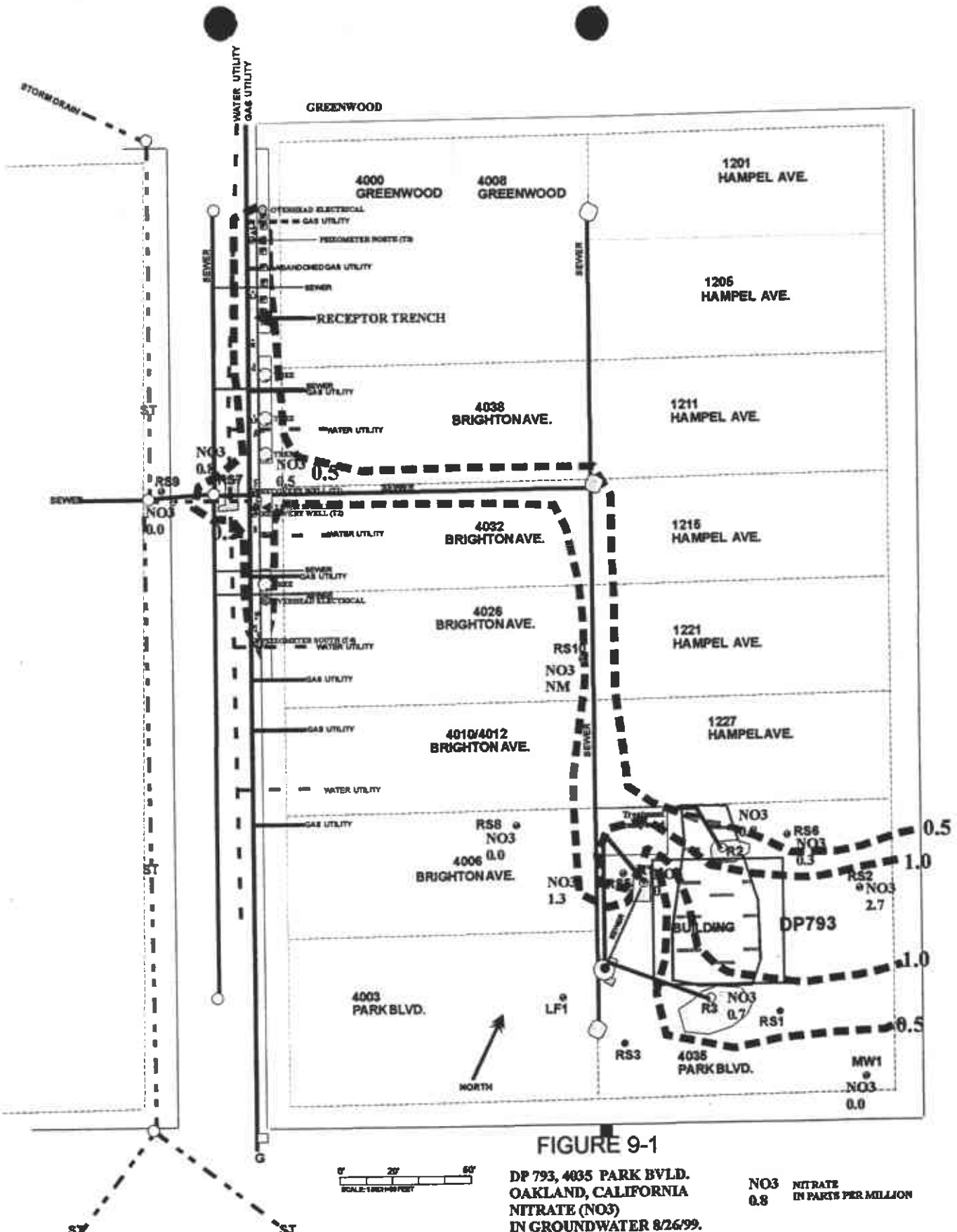


FIGURE 9-1

**DP 793, 4035 PARK BLVD.
OAKLAND, CALIFORNIA
NITRATE (NO3)
IN GROUNDWATER 8/26/99.**

**NO3 NITRATE
0.8 IN PARTS PER MILLION**

230 feet above mean sea level

- BK BLACK
- LB LIGHT BROWN
- DB DARK BROWN
- DG DARK GRAY

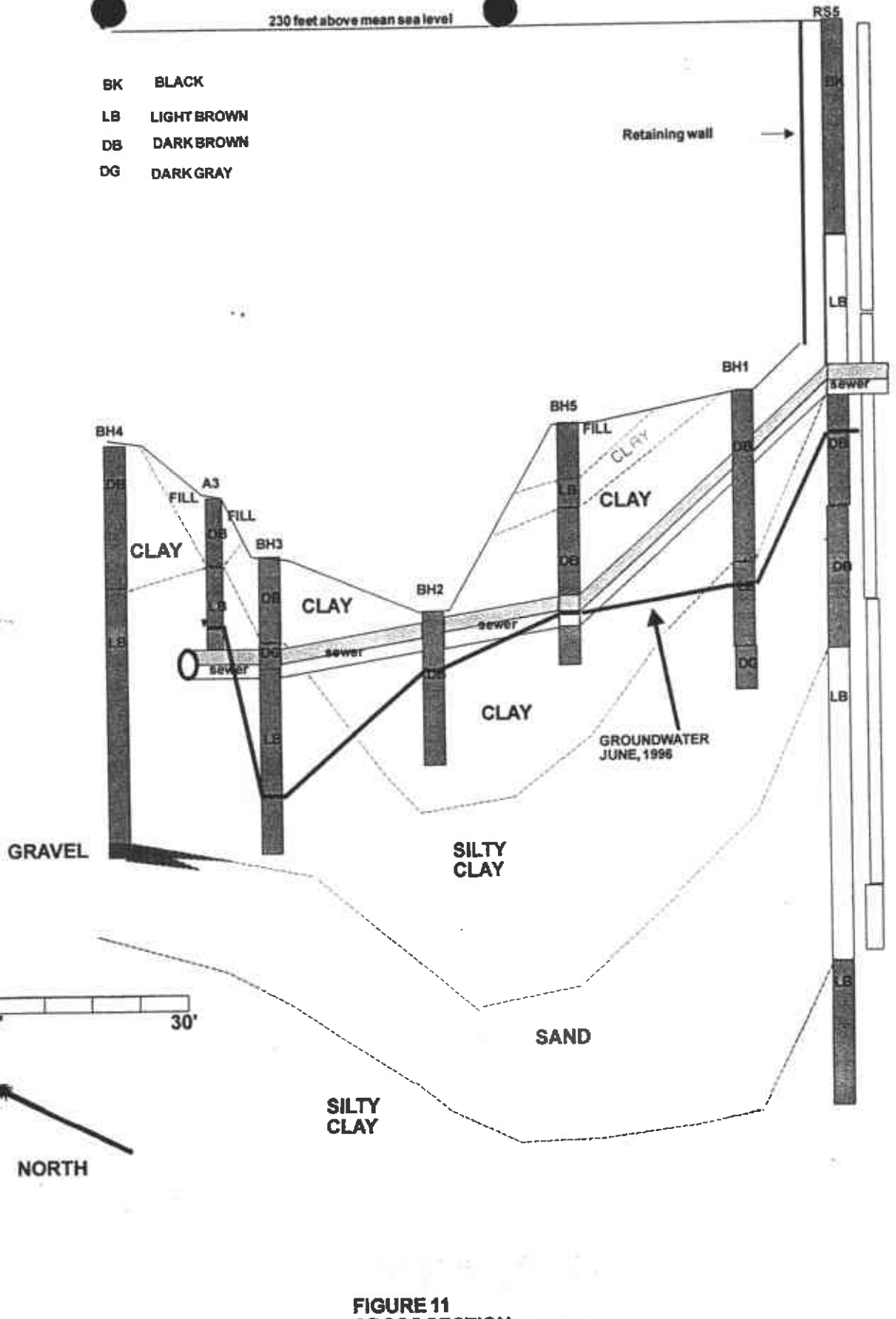
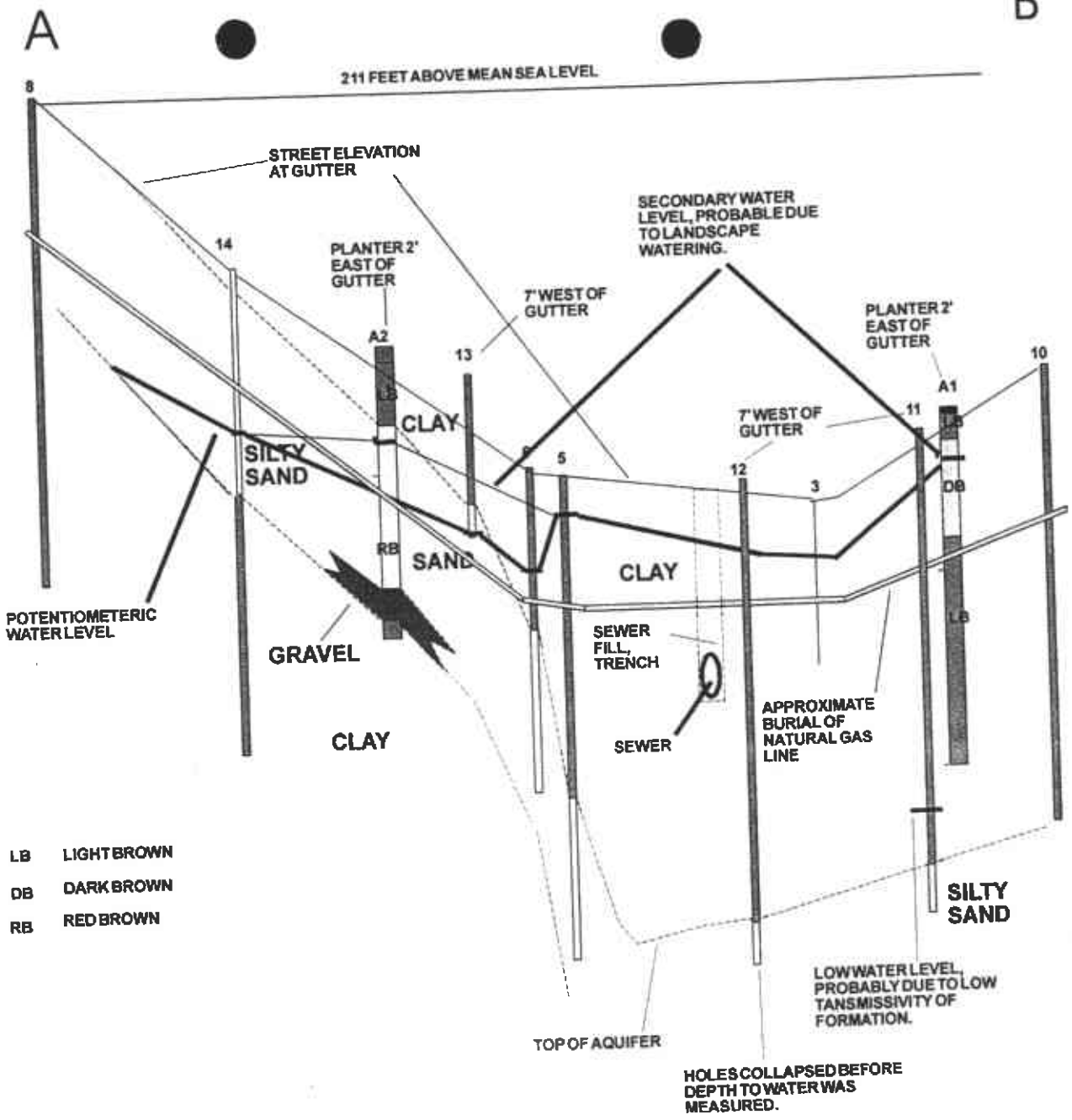


FIGURE 11
CROSS SECTION
FREE PRODUCT INVESTIGATION
FOR
DP793, 4035 PARK BLVD.
OAKLAND, CALIFORNIA



- LB LIGHT BROWN
- DB DARK BROWN
- RB RED BROWN

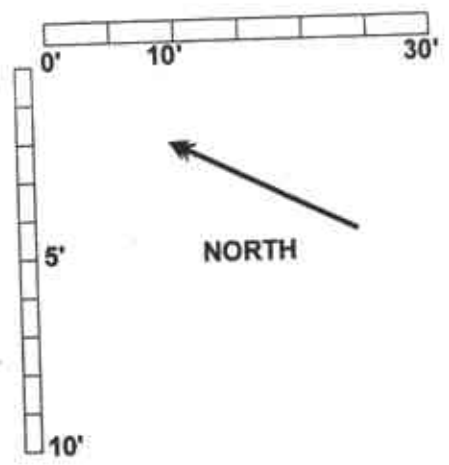


FIGURE 12
 CROSS SECTION
 FREE PRODUCT INVESTIGATION
 FOR
 DP793, 4035 PARK BLVD.
 OAKLAND, CALIFORNIA

APPENDIX A

METHODS AND PROCEDURES, QA/QC WITH FIELD NOTES

APPENDIX A.

METHODS AND PROCEDURES, QA/QC

This Appendix documents the specific methods, procedures, and materials used to collect and analyze ground water samples.

Gauging and Measuring Monitor Wells.

Prior to sampling a well, WEGE personnel obtain two measurements: the depth to ground water and the product thickness using a battery powered depth to water-product interface probe and or by using a specially designed bailer. The probe is lowered into the well casing until the instrument signals that the top of water has been reached. The distance from the top of water to the top of casing is read from the tape calibrated in 0.01 foot intervals for accuracy to 0.01 foot, that is attached to the probe. The measured distance is subtracted from the established elevation at the top of casing to determine the elevation of ground water with respect to mean sea level.

The probe is washed with TSP and rinsed in distilled water before each measurement. WEGE has designed and built bailers that will collect a sample of the contents of a well to show the exact thickness of any floating product.

Purging Standing Water from Monitor Wells

If no product is present, WEGE personnel purge the well. This is accomplished by removing ground water from the well until the water quality parameters (temperature, pH, and conductivity) stabilize, or until the well is emptied of water. Periodic measurements of ground water temperature, pH, and conductivity were taken with a Hydac Monitor or other meter and recorded along with the volume of ground water removed from the well. Purging is done by one or more methods singularly or in combination. Bailers, pneumatic or electric sample pumps, or vacuum pump tanks or trucks may be used. The usual amount of water removed is three well volumes. The water collected during purging is either safely stored onsite for later disposition, transported to an approved onsite or offsite sewer discharge system, or an approved onsite or offsite treatment system.

Collection of Water Sample for Analysis

The well is allowed to recover after purging and a ground water sample is collected. A fresh bailer is used to collect enough water for the requirements of the laboratory for the analyses needed or required. The water samples are decanted from the bailer into the appropriate number and size containers. These containers are furnished pre-cleaned to exact EPA protocols, with and without

preservatives added, by the analytical laboratory or a chemical supply company. The bottles are filled, with no headspace, and then capped with plastic caps with teflon liners.

The vials or bottles containing the ground water samples are labeled with site name, station, date, time, sampler, and analyses to be performed, and documented on a chain of custody form. They were placed in ziplock bags and stored in a chest cooled to 4°C with ice. The preserved samples are chain of custody delivered to the chosen laboratory.

Analytical Results

TPH is the abbreviations used for Total Petroleum Hydrocarbons used by the laboratories for water and soil analyses. The letter following TPH indicates a particular distinction or grouping for the results. The letters "g", "d", "k", or "o" indicates gasoline, diesel, kerosene, or oil, respectively, ie. TPH-d for diesel range TPH.

BTEX or MTBE are acronyms or abbreviations used for Benzene, Toluene, Ethylbenzene and all of the Xylenes (BTEX) and Methyl Tertiary Butyl Ether (MTBE), respectively.

MBTEX is the designation for the combination of the above five compounds.

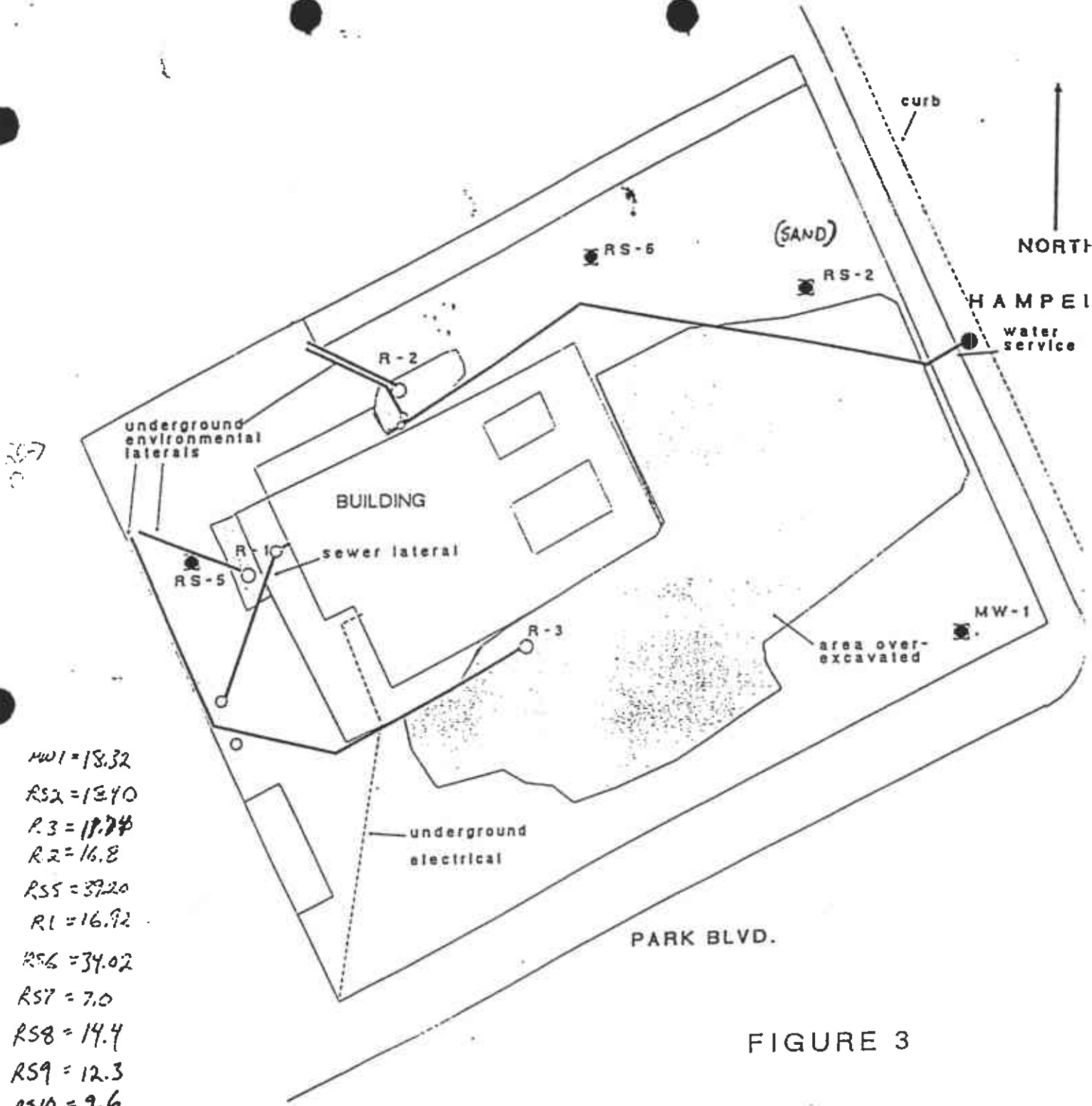
The less than symbol, <, used with a "parts per value" indicates the lower detection limit for a given analytical result and the level, if present, of that particular analyte is below or less than that lower detection limit.

Other abbreviations commonly used are ppm, ppb, mg/Kg, ug/Kg, ml/l and ul/l are parts per million, parts per billion, milligrams per kilogram, micrograms per kilogram, milliliters per liter, microliters per liter, respectively.

Chain of Custody Documentation

All water samples that are collected by WEGE and transported to a certified analytical laboratory are accompanied by chain-of-custody (COC) documentation. This documentation is used to record the movement and custody of a sample from collection in the field to final analysis and storage. Samples to be analyzed at the certified laboratory were logged on the COC sheet provided by the laboratory. The same information provided on the sample labels (site name, sample location, date, time, and analysis to be performed) is also noted on the COC form. Each person relinquishing custody of the sample set signs the COC form indicating the date and time of the transfer to the recipient. A copy of the COC follows the samples or their extracts throughout the laboratory to aid the analyst in identifying the samples and to assure analysis within holding times.

Copies of the COC documentation are included with the laboratory results in Appendix B of this report.

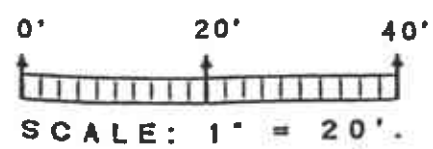


- MW1 = 18.32
- RS2 = 13.40
- R3 = 17.74
- R2 = 16.8
- RS5 = 39.20
- R1 = 16.92
- RS6 = 34.02
- RS7 = 7.0
- RS8 = 14.4
- RS9 = 12.3
- RS10 = 9.6

FIGURE 3

SITE BASE MAP

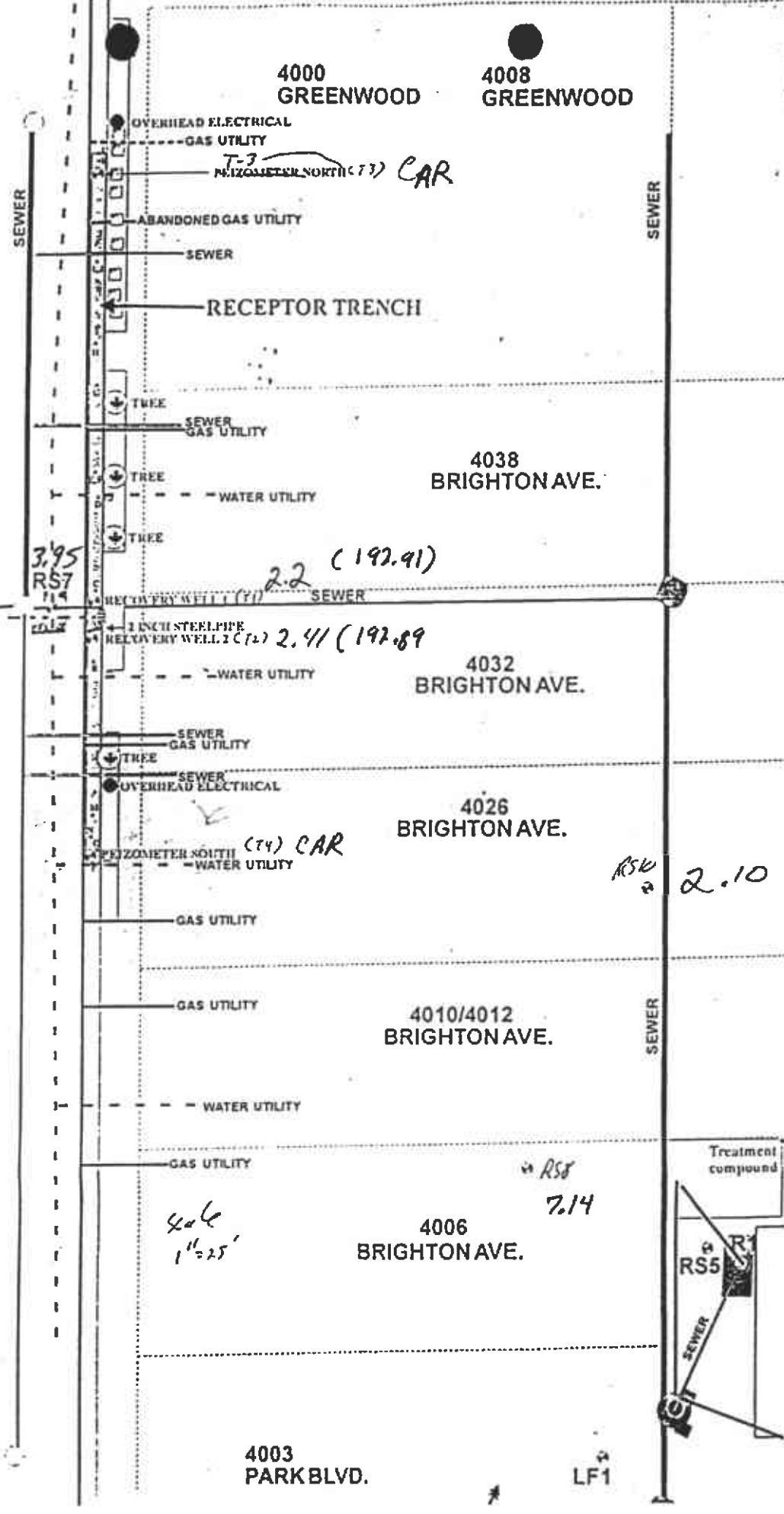
DESERT PETROLEUM STATION #793
 4035 PARK BLVD..
 OAKLAND, CALIFORNIA 94602



30-113 → 24 Hazard

0.829

RS7 = 7.0
 RS8 = 14
 RS9 = 12
 RS10 = 9
 T1 = 14.6
 T2 = 13.9



4.81 109

3.95
RS7
7.14

2.2 (192.91)

2.41 (192.89)

RS4 2.10

RS8 7.14

4.06
1" = 25'

LF1

WELL SAMPLING DATA SHEET

SITE	DP 793	DATE	12-18-01	TIME	11:37
WELL	R55	SAMPLED BY.	BROADWAY		
WELL ELEVATION					
PRODUCT THICKNESS					
DEPTH TO WATER	15.61	DTB	39.2		
FLUID ELEVATION	212.0				
BAILER TYPE	Disposable Bailer				
PUMP	David Pittman				

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
1139	1 Bailer	63.1	7.14	.34
1155	50 gal	65.8	6.91	.36
1157	1	66.4	6.85	.36
1159	1	66.6	6.83	.36

FINAL VOLUME PURGED	52 gal
TIME SAMPLED	1200
SAMPLE ID.	R55
SAMPLE CONTAINERS	3/40cc VOA's
ANALYSIS TO BE RUN	TP11g BTEX / MTRF
LABORATORY	NSE
NOTES:	1st Bailer partic Strong Odor
OR =	1.4
NO ₃ =	8.2
Fe ₂ =	3.30 X
SO ₄ =	37

WELL SAMPLING DATA SHEET

SITE <i>DP 793</i>	DATE <i>12-18-07</i>	TIME <i>955</i>
WELL <i>R57</i>	SAMPLED BY. <i>BROADWAY</i>	
WELL ELEVATION		
PRODUCT THICKNESS		
DEPTH TO WATER	<i>4.81</i>	DTB <i>2.0</i>
FLUID ELEVATION	<i>191.18</i>	
BAILER TYPE	<i>Disposable Bailer</i>	
PUMP	<i>David Pittman</i>	

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
<i>957</i>	<i>1 Bailer</i>	<i>61.3</i>	<i>7.58</i>	<i>.24</i>
<i>959</i>	<i>6 gal</i>	<i>66.9</i>	<i>7.22</i>	<i>.27</i>
<i>1001</i>	<i>1</i>	<i>67.4</i>	<i>7.04</i>	<i>.28</i>
<i>1003</i>	<i>1</i>	<i>67.3</i>	<i>6.85</i>	<i>.29</i>
<i>1005</i>	<i>1</i>	<i>68.1</i>	<i>6.82</i>	<i>.29</i>

FINAL VOLUME PURGED	<i>9 gal</i>
TIME SAMPLED	<i>1006</i>
SAMPLE ID.	<i>R57</i>
SAMPLE CONTAINERS	<i>3/40cc VOA's</i>
ANALYSIS TO BE RUN	<i>TP11g BTEX / MTBE</i>
LABORATORY	<i>NSC</i>
NOTES:	<i>1st Bailer Clear</i> <i>STRONG ODOOR</i>
<i>O₂ = 2.5</i>	
<i>NO₃ = 6.0</i>	
<i>Fe₂ = .87</i>	
<i>SO₄ = 1.</i>	

WELL SAMPLING DATA SHEET

SITE <i>OP 793</i>	DATE <i>12-18-01</i>	TIME <i>928</i>
WELL <i>RS10</i>	SAMPLED BY. <i>BROADWAY</i>	
WELL ELEVATION		
PRODUCT THICKNESS		
DEPTH TO WATER	<i>2.1</i>	DTB <i>9-</i>
FLUID ELEVATION	<i>206.36</i>	
BAILER TYPE	<i>Disposable Bailer</i>	
PUMP	<i>David Pittman</i>	

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
<i>929</i>	<i>1 Bailer</i>	<i>62.4</i>	<i>7.36</i>	<i>.22</i>
<i>931</i>	<i>1 gal</i>	<i>65.8</i>	<i>7.05</i>	<i>.25</i>
<i>933</i>	<i>1</i>	<i>66.3</i>	<i>6.72</i>	<i>.26</i>
<i>935</i>	<i>1</i>	<i>67.0</i>	<i>6.56</i>	<i>.26</i>
<i>937</i>	<i>1</i>	<i>66.9</i>	<i>6.54</i>	<i>.26</i>

FINAL VOLUME PURGED	<i>4 gal</i>
TIME SAMPLED	<i>938</i>
SAMPLE ID.	<i>RS10</i>
SAMPLE CONTAINERS	<i>3/40cc VOA's</i>
ANALYSIS TO BE RUN	<i>TP11g BTEX/MTBE</i>
LABORATORY	<i>USE</i>
NOTES:	<i>1st Bailer Clear No Odors</i>
<i>O₂ = 4.8</i>	
<i>NO₂ = 4.1</i>	
<i>Fe = 0</i>	
<i>SO₄²⁻ = 46</i>	

WELL SAMPLING DATA SHEET

SITE <i>DP 793</i>	DATE <i>12-18-01</i>	TIME <i>1201</i>
WELL <i>R1</i>	SAMPLED BY. <i>BROADWAY</i>	
WELL ELEVATION		
PRODUCT THICKNESS		
DEPTH TO WATER	<i>9.90 DTB 16.92</i>	
FLUID ELEVATION	<i>217.79</i>	
BAILER TYPE	<i>Disposable Bailer</i>	
PUMP	<i>David Pittman</i>	

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
<i>1202</i>	<i>1 Bailer</i>	<i>63.1</i>	<i>7.15</i>	<i>.10</i>
<i>1212</i>	<i>30 gal</i>	<i>64.0</i>	<i>7.17</i>	<i>.11</i>
<i>1214</i>	<i>1</i>	<i>66.2</i>	<i>7.25</i>	<i>.11</i>
<i>1216</i>	<i>1</i>	<i>66.4</i>	<i>7.24</i>	<i>.11</i>

FINAL VOLUME PURGED	<i>32 gal</i>
TIME SAMPLED	<i>1217</i>
SAMPLE ID.	<i>R1</i>
SAMPLE CONTAINERS	<i>3/40cc VOA's</i>
ANALYSIS TO BE RUN	<i>TP11g BTEX / MTBE</i>
LABORATORY	<i>NSC</i>
NOTES:	<i>1st Bailer Clear slight odor</i>
<i>O₂</i>	<i>= 5.2</i>
<i>NO₂</i>	<i>= 4.2</i>
<i>Fe₂</i>	<i>= 0</i>
<i>SO₄</i>	<i>= 14</i>

WELL SAMPLING DATA SHEET

SITE <i>OP 793</i>	DATE <i>12-18-01</i>	TIME <i>1115</i>
WELL <i>R2</i>	SAMPLED BY. <i>BROADWAY</i>	
WELL ELEVATION		
PRODUCT THICKNESS		
DEPTH TO WATER	<i>12.35</i>	DTB <i>16.8</i>
FLUID ELEVATION	<i>214.93</i>	
BAILER TYPE	<i>Disposable Bailer</i>	
PUMP	<i>David Pittman</i>	

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
<i>1119</i>	<i>1 Bailer</i>	<i>63.4</i>	<i>6.91</i>	<i>.43</i>
<i>1126</i>	<i>20 gal</i>	<i>65.0</i>	<i>6.88</i>	<i>.48</i>
<i>1128</i>	<i>1</i>	<i>66.3</i>	<i>6.71</i>	<i>.49</i>
<i>1130</i>	<i>1</i>	<i>66.5</i>	<i>6.69</i>	<i>.49</i>

FINAL VOLUME PURGED	<i>22 gal</i>
TIME SAMPLED	<i>1131</i>
SAMPLE ID.	<i>R2</i>
SAMPLE CONTAINERS	<i>3/40cc VOA's</i>
ANALYSIS TO BE RUN	<i>TP11g BTEX/MTBE</i>
LABORATORY	<i>NSC</i>
NOTES:	<i>1st Bailer Clear Some Max</i>
<i>O₂ = 2.8</i>	
<i>NO₃ = 1.3</i>	
<i>Fe₂ = .07</i>	
<i>SO₄ = 77%</i>	

WELL SAMPLING DATA SHEET

SITE <i>OP 793</i>	DATE <i>12-18-01</i>	TIME <i>1230</i>
WELL <i>R3</i>	SAMPLED BY. <i>BROADWAY</i>	
WELL ELEVATION		
PRODUCT THICKNESS		
DEPTH TO WATER	<i>6.79</i>	DTB <i>11.74</i>
FLUID ELEVATION	<i>220.46</i>	
BAILER TYPE	<i>Disposable Bailer</i>	
PUMP	<i>David Pittman</i>	

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
<i>1250</i>	<i>1 Bailer</i>	<i>63.5</i>	<i>7.15</i>	<i>.35</i>
<i>1258</i>	<i>20 gal</i>	<i>65.9</i>	<i>7.04</i>	<i>.38</i>
<i>1300</i>	<i>1</i>	<i>66.8</i>	<i>6.89</i>	<i>.39</i>
<i>1302</i>	<i>1</i>	<i>67.1</i>	<i>6.91</i>	<i>.39</i>

FINAL VOLUME PURGED	<i>22 gal</i>
TIME SAMPLED	<i>1305</i>
SAMPLE ID.	<i>R3</i>
SAMPLE CONTAINERS	<i>140cc VOA's</i>
ANALYSIS TO BE RUN	<i>TP11g BTEX / MTBE</i>
LABORATORY	<i>NSF</i>
NOTES:	<i>1st Bailer Clear No Odor</i>
<i>O₂ = 5.5</i>	
<i>NO₃ = 6.2</i>	
<i>Fe₂ = 0</i>	
<i>SU₄ = 77#</i>	

WELL SAMPLING DATA SHEET

SITE 00793	DATE 12-18-01	TIME 1006
WELL T1	SAMPLED BY. BROADWAY	
WELL ELEVATION		
PRODUCT THICKNESS		
DEPTH TO WATER	2.2	DTB 14.6
FLUID ELEVATION	192.91	
BAILER TYPE	Disposable Bailer	
PUMP	David Pittman	

WELL PURGING RECORD				
TIME	VOLUME REMOVED	TEMP. F°	pH	COND. X1000
1006	1 Bailer	61.5	7.01	.39
1008	40 gal	65.6	6.83	.38
1010	'	66.3	6.61	.38
1012	'	66.3	6.52	.39
			6.52	.38

FINAL VOLUME PURGED	42 gal
TIME SAMPLED	10/12
SAMPLE ID.	T1
SAMPLE CONTAINERS	40cc VOA's
ANALYSIS TO BE RUN	TP11g BTEX / MTBE
LABORATORY	NSC
NOTES:	1 st Bailer Turbid Strong Odor
O ₂ = 2.8	
NO ₃ = 4.3	
Fe ₂ = .60	
SO ₄ = 0	

Project Manager: George Converse Phone No.: 530-668-5300

Chain-of-Custody Record and Analysis Request

Company/Address: Woodland WEGE 1736 E. Bunker CA 95776 FAX No.: 530-662-0273

Analysis Request

Project Number: _____ P.O. No.: _____ Email Address: wege@metlex.com
 .pdf .xls .doc other

Project Name/Location: DP 793 PARK HUB Sampler Signature: [Signature]

Sample Designation	Sampling		Container (Type/Amount)		Method Preserved				Matrix	Analysis Request										TAT	For Lab Use Only			
	Date	Time	40 ml VOA	SLEEVE	HCl	HNO ₃	ICE	NONE	WATER/SOIL	BTEX (8021B)	BTEX/TPH Gas/MTBE (8021B/M8015)	TPH as Diesel (M8015)	TPH as Motor Oil (M8015)	TPH Gas/BTEX/MTBE (8260B)	5 Oxygenates/TPH Gas/BTEX (8260B)	7 Oxygenates/TPH Gas/BTEX (8260B)	5 Oxygenates (8260B)	7 Oxygenates (8260B)	Lead Scav. (1,2 DCA & 1,2 EDB - 8260B)	EPA 8260B (Full List)	Volatile Halocarbons (EPA 8260B)	Lead (7421/239.2) TOTAL (X) W.E.T. (X)	12 hr/24 hr/48 hr/72 hr/1 wk	12 hr = Results by 9 a.m. of the next bus. day 24 hr = Results by 5 p.m. of the next bus. day 48 hr = Results by 5 p.m. of the 2nd bus. day 72 hr = Results by 5 p.m. of the 3rd bus. day 1 wk = Results by 5 p.m. of the 1st bus. day
MW1	12/18/01	1040	3																					
R1		1217																						
R2		1131																						
R3		1305																						
R52		1101																						
R55		1200																						
R56		1116																						
R57		1006																						
R58		915																						
R59		951																						

Relinquished by: [Signature] Date: 12/18/01 Time: 1240 Received by: _____ Remarks: _____

Relinquished by: _____ Date: _____ Time: _____ Received by: _____

Relinquished by: _____ Date: 12/19/01 Time: 1240 Received by Laboratory: Martin, He / KIFF Analytical Bill to: _____



720 Olive Drive, Suite D
 Davis, CA 95616
 Lab: 530.297.4800
 Fax: 530.297.4803

Lab No. _____

Page 2 of 2

Project Manager: George Comstock Phone No.: _____

Company/Address: WCCG FAX No.: _____

Project Number: _____ P.O. No.: _____ Email Address: _____
 .pdf .xls .doc other

Project Name: ADP793 Sampler Signature: [Signature]

Project Location: Park Blvd Oakland

Chain-of-Custody Record and Analysis Request

Analysis Request

Sample Designation	Sampling		Container		Preservative				Matrix		BTEX (8021B)	BTEX/TPH Gas/MTBE (8021B/MB015)	TPH as Diesel (MB015)	TPH as Motor Oil (MB015)	TPH Gas/BTEX/MTBE (8260B)	5 Oxygenates/TPH Gas/BTEX (8260B)	7 Oxygenates/TPH Gas/BTEX (8260B)	5 Oxygenates (8260B)	7 Oxygenates (8260B)	Lead Scav. (1,2 DCA & 1,2 ED6 - 8260B)	EPA 8260B (Full List)	Volatile Halocarbons (EPA 8260B)	Lead (7421/239.2) TOTAL (X) W.E.T. (X)	TAT	For Lab Use Only	
	Date	Time	40 ml VOA	SLEEVE	HCl	HNO ₃	ICE	NONE	WATER	SOIL																
RS10	12/18/11	9:38	3																							
T1		10:12																								
CARBON Sample discharge		13:50																								

Relinquished by: [Signature] Date: 12/18/11 Time: 12:40 Received by: _____

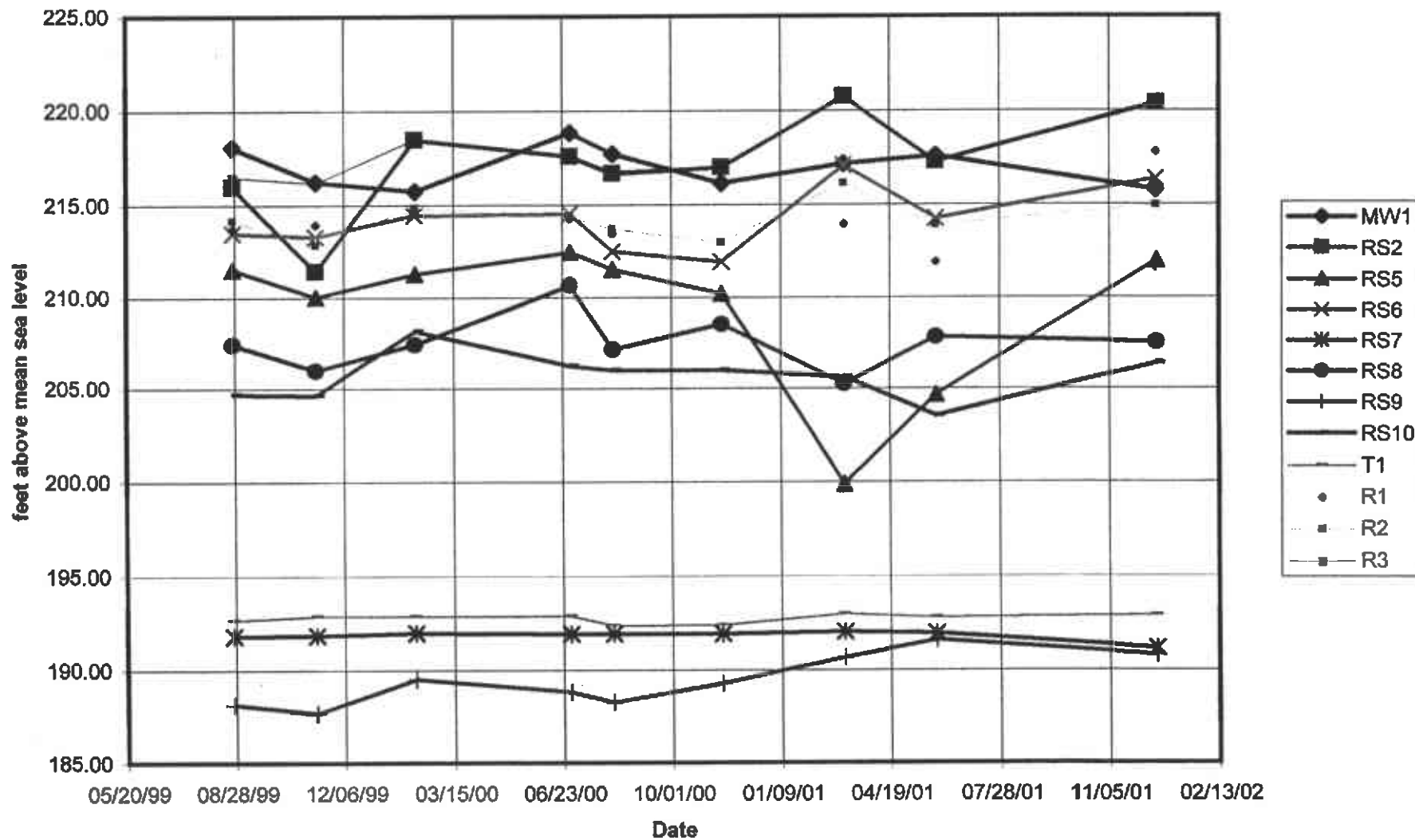
Relinquished by: _____ Date: _____ Time: _____ Received by: _____

Relinquished by: _____ Date: 12/19/11 Time: 12:10 Received by Laboratory: Mark [Signature] / K.I.F. Analytical Bill to: _____

Remarks: _____

APPENDIX B.
GROUNDWATER ELEVATION CHART

Groundwater Elevation





Report Number : 23991

Date : 1/2/2002

George Converse
Western Geo-Engineers
1386 East Beamer St.
Woodland, CA 95776

Subject : 13 Water Samples
Project Name : DP793 PARK BLVD
Project Number :

Dear Mr. Converse,

Chemical analysis of the samples referenced above has been completed. Summaries of the data are contained on the following pages. Sample(s) were received under documented chain-of-custody. US EPA protocols for sample storage and preservation were followed.

Kiff Analytical is certified by the State of California (# 2236). If you have any questions regarding procedures or results, please call me at 530-297-4800.

Sincerely,


Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : MW1

Matrix : Water

Lab Number : 23991-01

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	102		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	103		% Recovery	EPA 8260B	12/26/2001

Sample : R1

Matrix : Water

Lab Number : 23991-02

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	1.5	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	102		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	104		% Recovery	EPA 8260B	12/26/2001

Approved By:  Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : R2

Matrix : Water

Lab Number : 23991-03

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	2000	5.0	ug/L	EPA 8260B	12/29/2001
Toluene	120	5.0	ug/L	EPA 8260B	12/29/2001
Ethylbenzene	44	5.0	ug/L	EPA 8260B	12/29/2001
Total Xylenes	280	5.0	ug/L	EPA 8260B	12/29/2001
Methyl-t-butyl ether (MTBE)	< 5.0	5.0	ug/L	EPA 8260B	12/29/2001
TPH as Gasoline	4900	500	ug/L	EPA 8260B	12/29/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/29/2001
4-Bromofluorobenzene (Surr)	109		% Recovery	EPA 8260B	12/29/2001

Sample : R3

Matrix : Water

Lab Number : 23991-04

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	104		% Recovery	EPA 8260B	12/26/2001

Approved By:  Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : RS2

Matrix : Water

Lab Number : 23991-05

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	103		% Recovery	EPA 8260B	12/26/2001

Sample : RS5

Matrix : Water

Lab Number : 23991-06

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	610	5.0	ug/L	EPA 8260B	12/29/2001
Toluene	1200	5.0	ug/L	EPA 8260B	12/29/2001
Ethylbenzene	100	5.0	ug/L	EPA 8260B	12/29/2001
Total Xylenes	1500	5.0	ug/L	EPA 8260B	12/29/2001
Methyl-t-butyl ether (MTBE)	< 5.0	5.0	ug/L	EPA 8260B	12/29/2001
TPH as Gasoline	12000	500	ug/L	EPA 8260B	12/29/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/29/2001
4-Bromofluorobenzene (Surr)	109		% Recovery	EPA 8260B	12/29/2001

Approved By:  Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : RS6

Matrix : Water

Lab Number : 23991-07

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	0.53	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	0.56	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	56	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	104		% Recovery	EPA 8260B	12/26/2001

Sample : RS7

Matrix : Water

Lab Number : 23991-08

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	450	2.0	ug/L	EPA 8260B	1/1/2002
Toluene	21	2.0	ug/L	EPA 8260B	1/1/2002
Ethylbenzene	86	2.0	ug/L	EPA 8260B	1/1/2002
Total Xylenes	120	2.0	ug/L	EPA 8260B	1/1/2002
Methyl-t-butyl ether (MTBE)	2.3	2.0	ug/L	EPA 8260B	1/1/2002
TPH as Gasoline	2700	200	ug/L	EPA 8260B	1/1/2002
Toluene - d8 (Surr)	100		% Recovery	EPA 8260B	1/1/2002
4-Bromofluorobenzene (Surr)	104		% Recovery	EPA 8260B	1/1/2002

Approved By:  Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : RS8

Matrix : Water

Lab Number : 23991-09

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	230	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	370	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	77	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	750	2.0	ug/L	EPA 8260B	12/28/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	4500	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	102		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	105		% Recovery	EPA 8260B	12/26/2001

Sample : RS9

Matrix : Water

Lab Number : 23991-10

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	11	0.50	ug/L	EPA 8260B	12/28/2001
Toluene	1.8	0.50	ug/L	EPA 8260B	12/28/2001
Ethylbenzene	3.9	0.50	ug/L	EPA 8260B	12/28/2001
Total Xylenes	7.6	0.50	ug/L	EPA 8260B	12/28/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/28/2001
TPH as Gasoline	210	50	ug/L	EPA 8260B	12/28/2001
Toluene - d8 (Surr)	99.6		% Recovery	EPA 8260B	12/28/2001
4-Bromofluorobenzene (Surr)	104		% Recovery	EPA 8260B	12/28/2001

Approved By:  Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : RS10

Matrix : Water

Lab Number : 23991-11

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/26/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	103		% Recovery	EPA 8260B	12/26/2001

Sample : T1

Matrix : Water

Lab Number : 23991-12

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	3700	20	ug/L	EPA 8260B	1/1/2002
Toluene	5500	20	ug/L	EPA 8260B	1/1/2002
Ethylbenzene	1200	20	ug/L	EPA 8260B	1/1/2002
Total Xylenes	5300	20	ug/L	EPA 8260B	1/1/2002
Methyl-t-butyl ether (MTBE)	24	20	ug/L	EPA 8260B	1/1/2002
TPH as Gasoline	48000	2000	ug/L	EPA 8260B	1/1/2002
Toluene - d8 (Surr)	98.4		% Recovery	EPA 8260B	1/1/2002
4-Bromofluorobenzene (Surr)	102		% Recovery	EPA 8260B	1/1/2002

Approved By:  Joel Kiff



Report Number : 23991

Date : 1/2/2002

Project Name : DP793 PARK BLVD

Project Number :

Sample : CARBON DISCHARGE

Matrix : Water

Lab Number : 23991-13

Sample Date :12/18/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/29/2001
Toluene - d8 (Surr)	101		% Recovery	EPA 8260B	12/29/2001
4-Bromofluorobenzene (Surr)	110		% Recovery	EPA 8260B	12/29/2001

Approved By:  Joel Kiff

Report Number : 23991

Date : 1/2/2002

QC Report : Method Blank Data

Project Name : DP793 PARK BLVD

Project Number :

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/29/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/29/2001
Toluene - dB (Surr)	98.3		%	EPA 8260B	12/29/2001
4-Bromofluorobenzene (Surr)	96.9		%	EPA 8260B	12/29/2001
Benzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Toluene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Ethylbenzene	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Total Xylenes	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
Methyl-t-butyl ether (MTBE)	< 0.50	0.50	ug/L	EPA 8260B	12/26/2001
TPH as Gasoline	< 50	50	ug/L	EPA 8260B	12/26/2001
Toluene - dB (Surr)	101		%	EPA 8260B	12/26/2001
4-Bromofluorobenzene (Surr)	104		%	EPA 8260B	12/26/2001

Parameter	Measured Value	Method Reporting Limit	Units	Analysis Method	Date Analyzed
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Approved By: Joel Kiff

Report Number : 23991


Date : 1/2/2002

QC Report : Matrix Spike/ Matrix Spike Duplicate

Project Name : DP793 PARK BLVD

Project Number :

Parameter	Spiked Sample	Sample Value	Spike Level	Spike Dup. Level	Spiked Sample Value	Duplicate Spiked Sample Value	Units	Analysis Method	Date Analyzed	Spiked Sample Percent Recov.	Duplicate Spiked Sample Percent Recov.	Relative Percent Diff.	Spiked Sample Percent Recov. Limit	Relative Percent Diff. Limit
Benzene	23991-13	<0.50	19.2	18.8	18.3	17.7	ug/L	EPA 8260B	12/29/2009	95.0	94.5	0.501	70-130	25
Toluene	23991-13	<0.50	19.2	18.8	18.7	18.2	ug/L	EPA 8260B	12/29/2009	97.1	96.9	0.180	70-130	25
Tert-Butanol	23991-13	25	96.2	93.8	125	118	ug/L	EPA 8260B	12/29/2001	104	98.8	4.90	70-130	25
Methyl-t-Butyl Ether	23991-13	<0.50	19.2	18.8	17.4	16.6	ug/L	EPA 8260B	12/29/2009	90.8	88.2	2.82	70-130	25
Benzene	23952-01	<0.50	40.0	40.0	37.8	37.0	ug/L	EPA 8260B	12/26/2009	94.4	92.6	1.95	70-130	25
Toluene	23952-01	<0.50	40.0	40.0	38.8	38.0	ug/L	EPA 8260B	12/26/2009	97.0	95.0	2.06	70-130	25
Tert-Butanol	23952-01	<5.0	200	200	186	190	ug/L	EPA 8260B	12/26/2009	92.8	95.0	2.30	70-130	25
Methyl-t-Butyl Ether	23952-01	210	40.0	40.0	252	252	ug/L	EPA 8260B	12/26/2001	102	103	1.15	70-130	25

Approved By:  Joel Kiff

KIFF ANALYTICAL, LLC

720 Olive Drive, Suite D Davis, CA 95616 530-297-4800

Report Number : 23991

Date : 1/2/2002

QC Report : Laboratory Control Sample (LCS)

Project Name : DP793 PARK BLVD

Project Number :

Parameter	Spike Level	Units	Analysis Method	Date Analyzed	LCS Percent Recov.	LCS Percent Recov. Limit
Benzene	40.0	ug/L	EPA 8260B	12/29/200	93.0	70-130
Toluene	40.0	ug/L	EPA 8260B	12/29/200	94.9	70-130
Tert-Butanol	200	ug/L	EPA 8260B	12/29/200	101	70-130
Methyl-t-Butyl Ether	40.0	ug/L	EPA 8260B	12/29/200	86.6	70-130
Benzene	40.0	ug/L	EPA 8260B	12/26/200	94.5	70-130
Toluene	40.0	ug/L	EPA 8260B	12/26/200	95.8	70-130
Tert-Butanol	200	ug/L	EPA 8260B	12/26/200	89.4	70-130
Methyl-t-Butyl Ether	40.0	ug/L	EPA 8260B	12/26/200	94.2	70-130

KIFF ANALYTICAL, LLC

720 Olive Drive, Suite D Davis, CA 95616 530-297-4800

Approved By:


Joel Kiff

Project Manager: George Converse
 Company/Address: WEGE 1386 E. Baymer Woodland CA 95776
 Project Number: _____ P.O. No.: _____
 Project Name/Location: DP 793 PARK BLVD

Phone No.: 530-668-5300
 FAX No.: 530-662-0273
 Email Address: wege@mtl.com
 .pdf .xls .doc other
 Sampler Signature: [Signature]

Chain-of-Custody Record and Analysis Request

Sample Designation	Sampling		Container (Type/Amount)		Method Preserved				Matrix	Analysis Request										TAT	For Lab Use Only					
	Date	Time	40 ml VOA	SLEEVE	HCl	HNO ₃	ICE	NONE	WATER/SOIL	BTEX (8021B)	BTEX/TPH Gas/MTBE (8021B/M8015)	TPH as Diesel (M8015)	TPH as Motor Oil (M8015)	TPH Gas/BTEX/MTBE (8260B)	5 Oxygenates/TPH Gas/BTEX (8260B)	7 Oxygenates/TPH Gas/BTEX (8260B)	5 Oxygenates (8260B)	7 Oxygenates (8260B)	Lead Scav. (1,2 DCA & 1,2 EDB - 8260B)	EPA 8260B (Full List)	Volatile Halocarbons (EPA 8260B)	Lead (7421/239.2) TOTAL (X) W.E.T. (X)	12 hr / 24 hr / 48 hr / 72 hr / 1 wk			
MW1	12/18/01	1040	3																						-01	
R1		1217																								-02
R2		1131																								-03
R3		1305																								-04
R52		1101																								-05
R55		1200																								-06
R56		1116																								-07
R57		1006																								-08
R58		915																								-09
R59		951																								-10

Relinquished by: [Signature] Date: 12/18/01 Time: 1240 Received by: _____
 Relinquished by: _____ Date: _____ Time: _____ Received by: _____
 Relinquished by: _____ Date: 12/19/01 Time: 1240 Received by Laboratory: Manita Lee / KIFF Analytical

Analysis Request

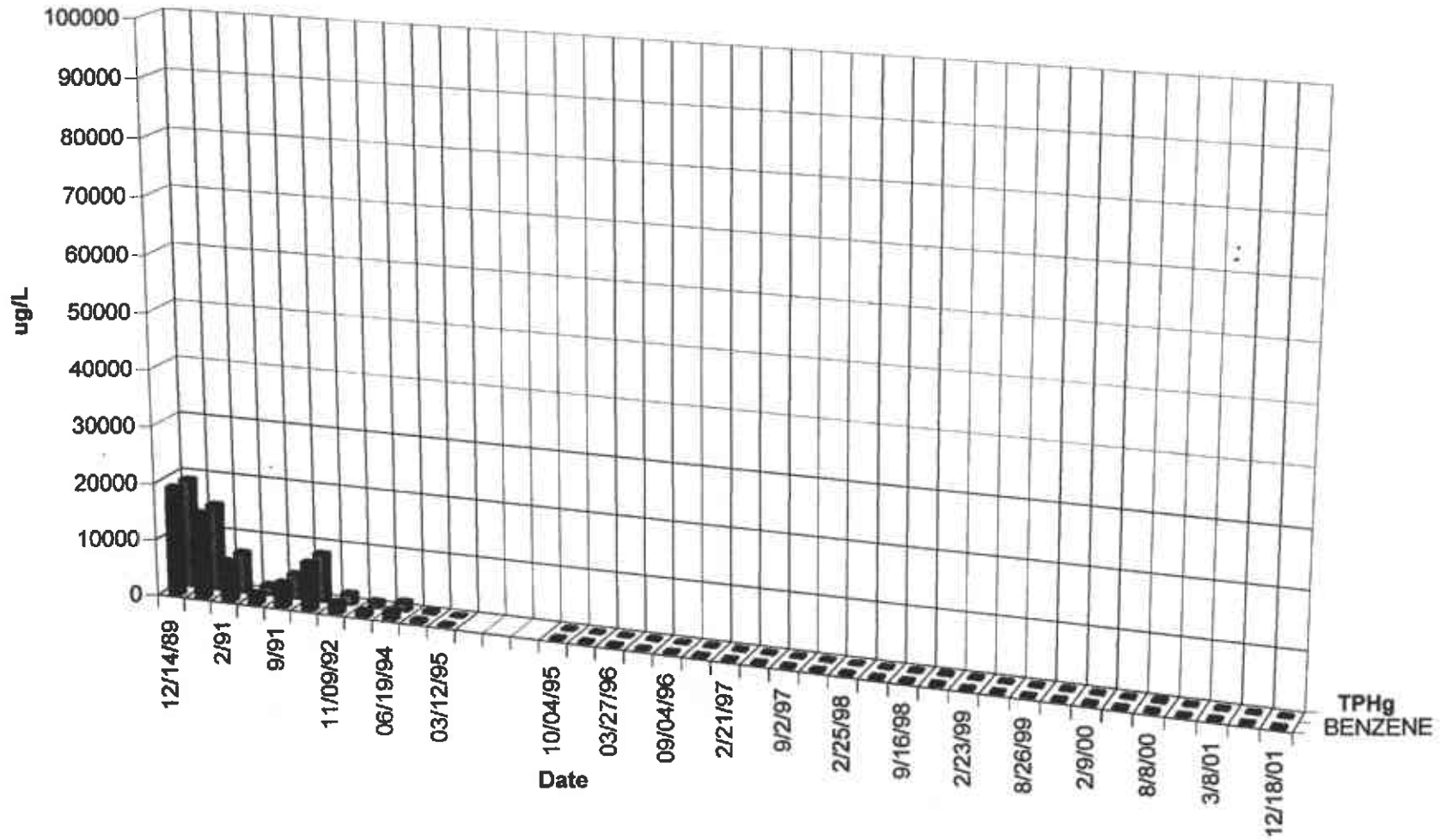
TAT: _____

For Lab Use Only

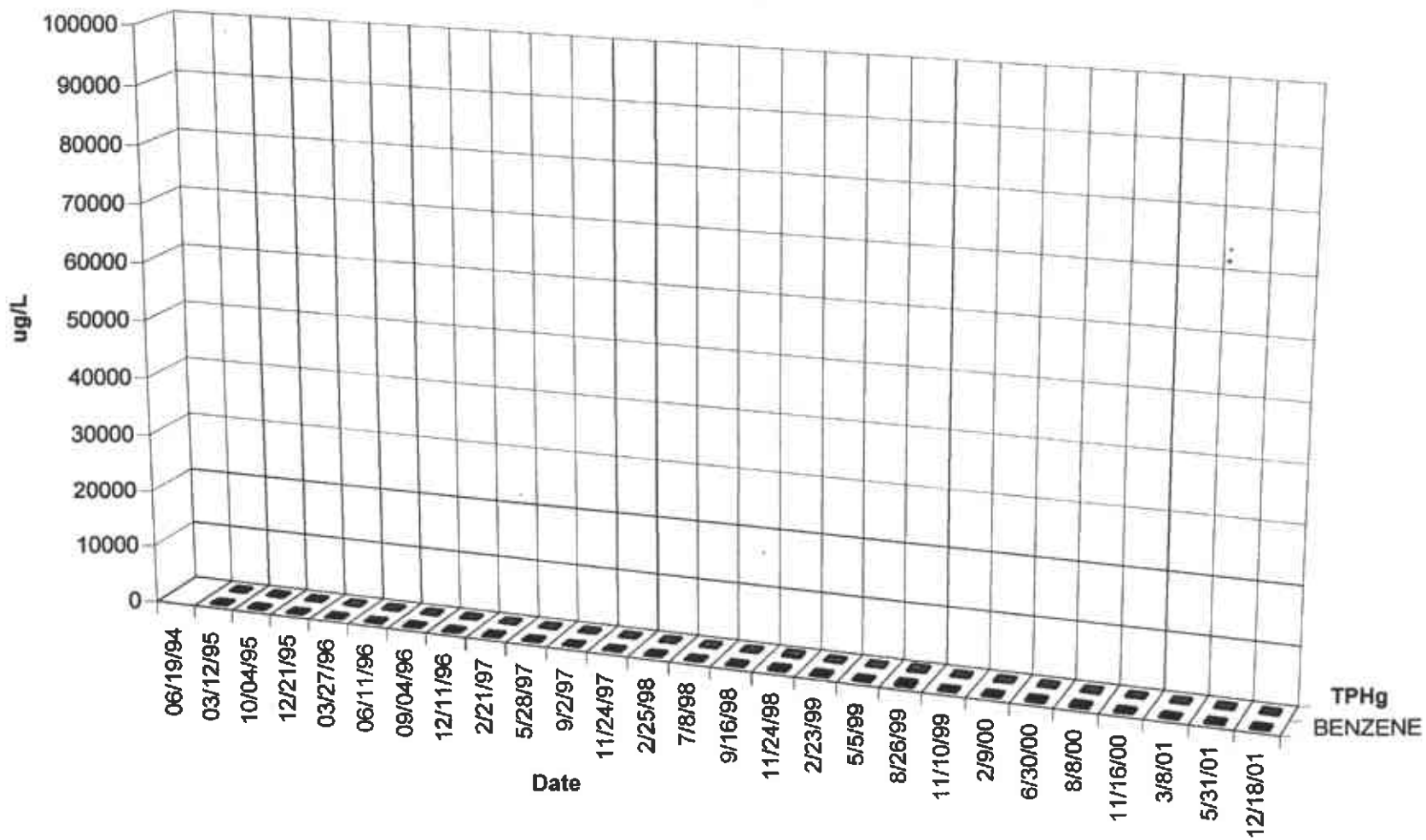
12 hr = Results by 9 a.m. of the next bus. day
 24 hr = Results by 5 p.m. of the next bus. day
 48 hr = Results by 5 p.m. of the 2nd bus. day
 72 hr = Results by 5 p.m. of the 3rd bus. day
 1 wk = Results by 5 p.m. of the 5th bus. day

APPENDIX D.
TPHg AND BENZENE CHARTS

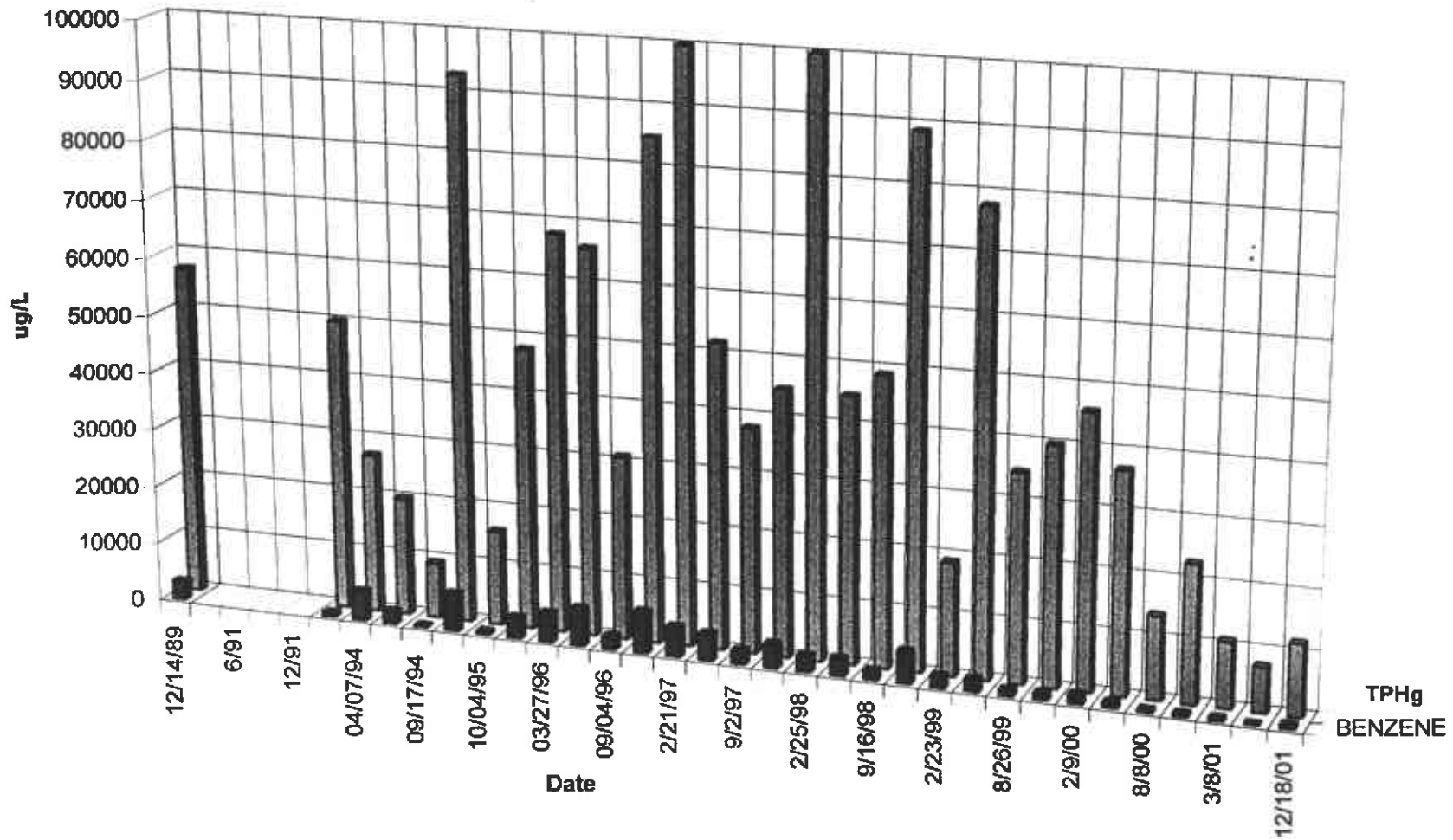
RS-1/MW-1 TPHg



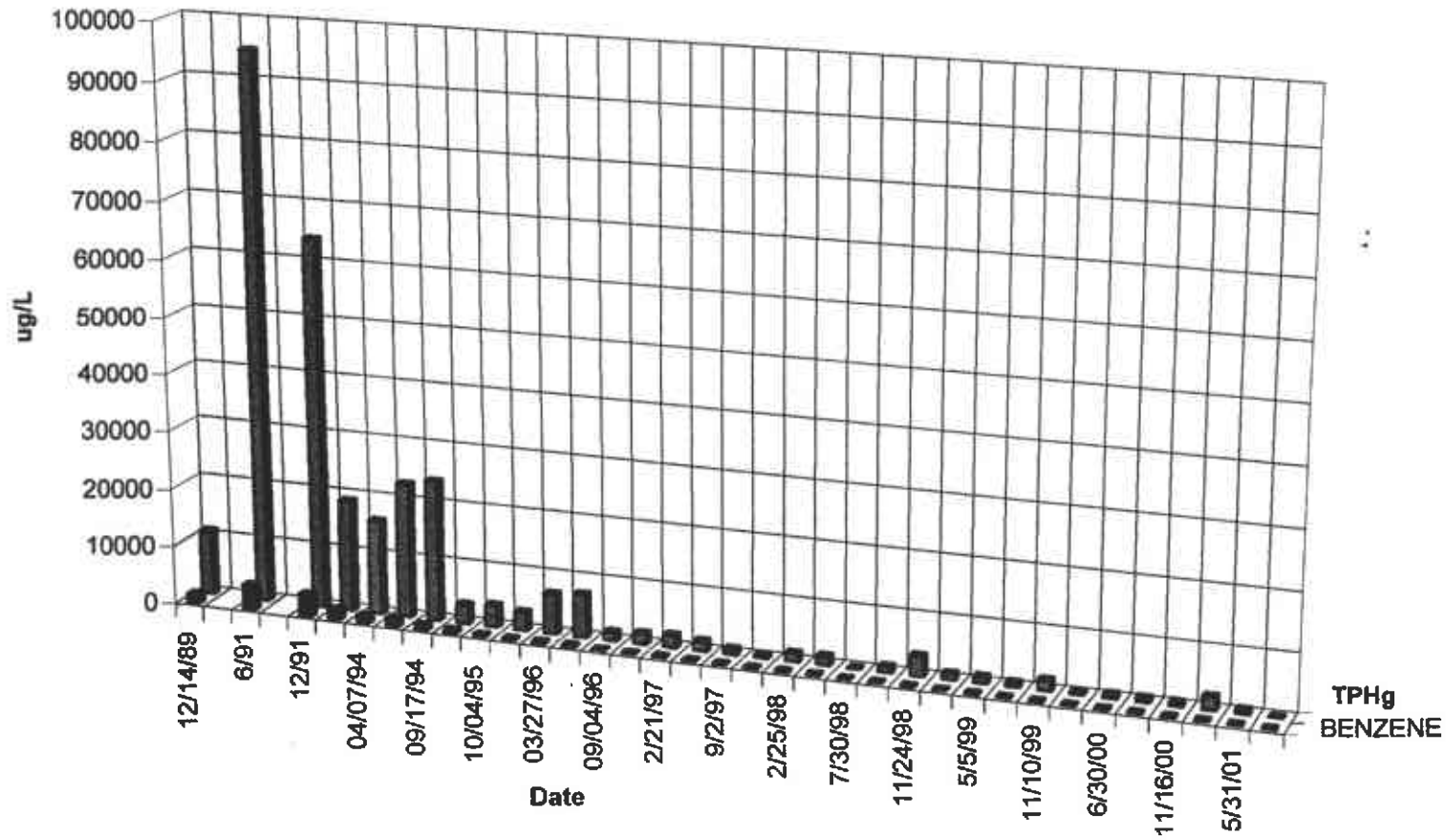
RS-2 TPHg



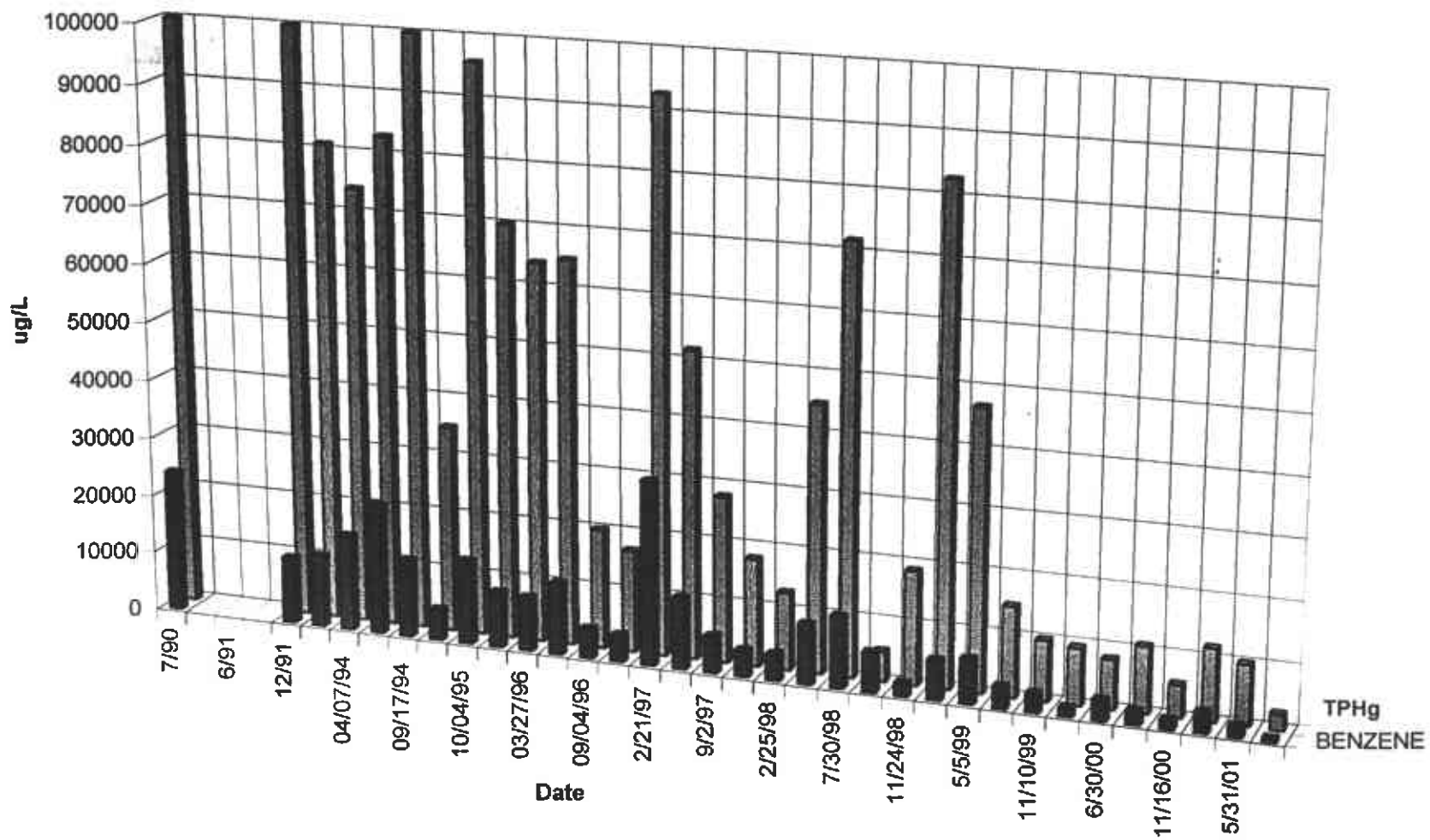
RS-5



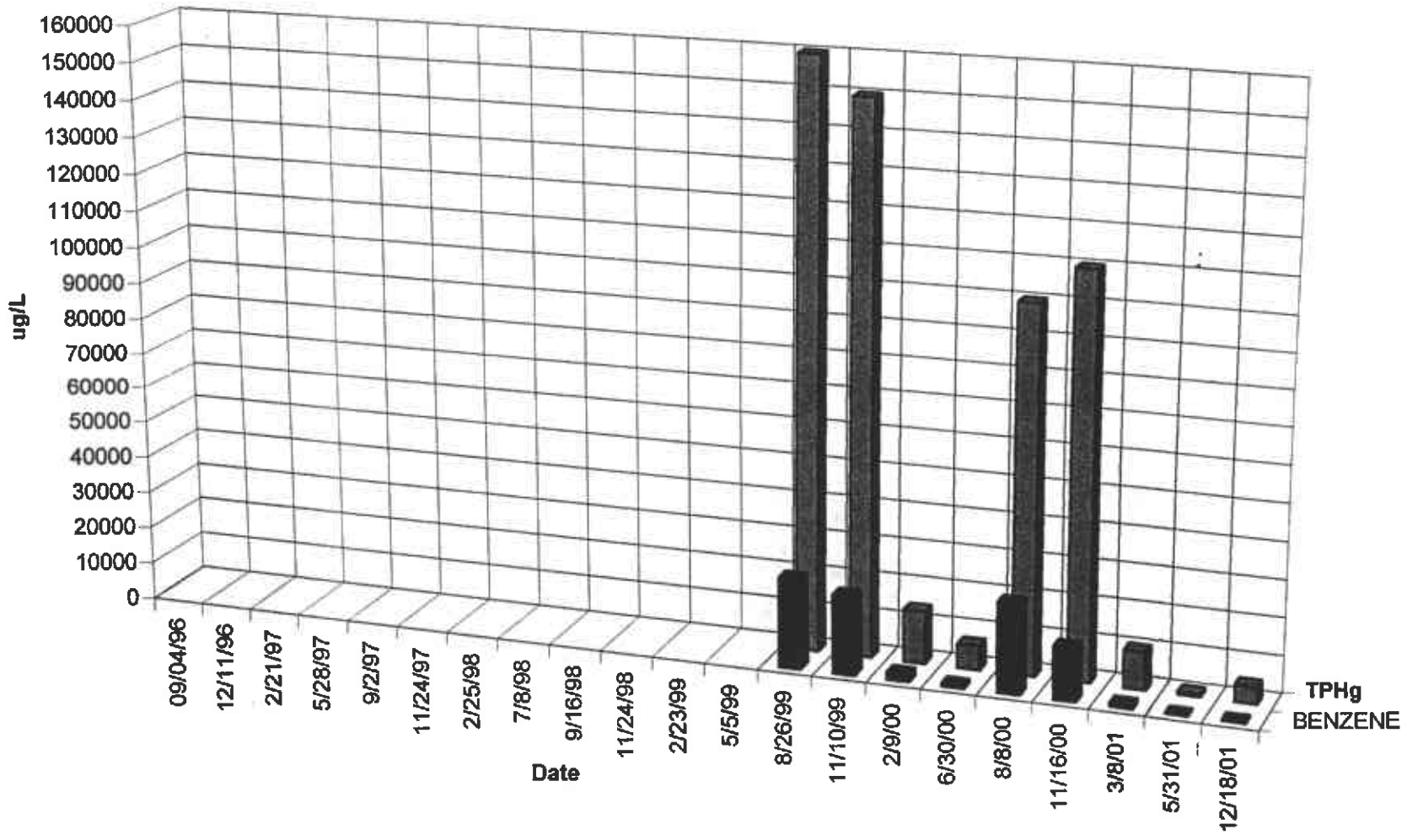
RS-6



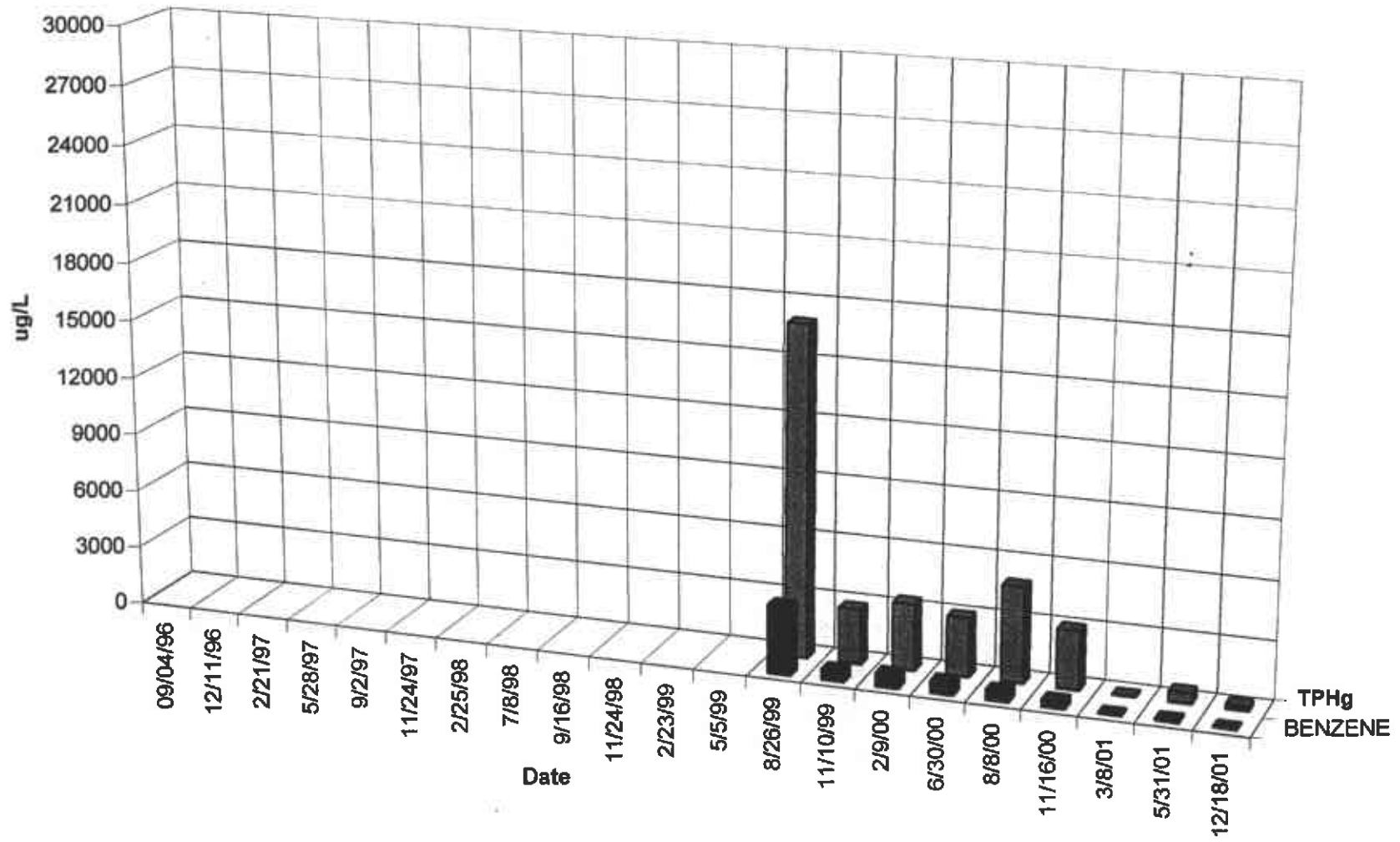
RS-7



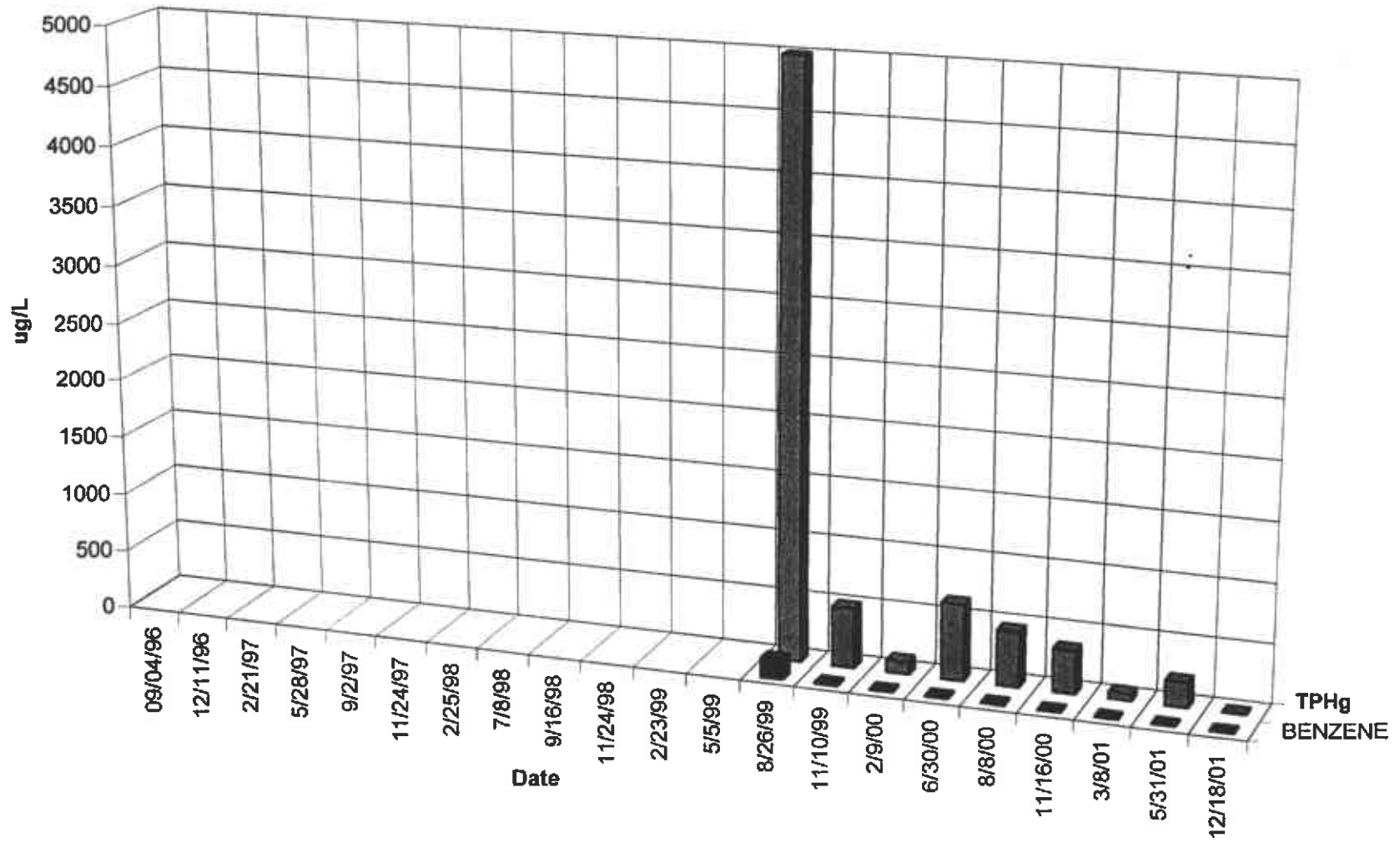
RS-8



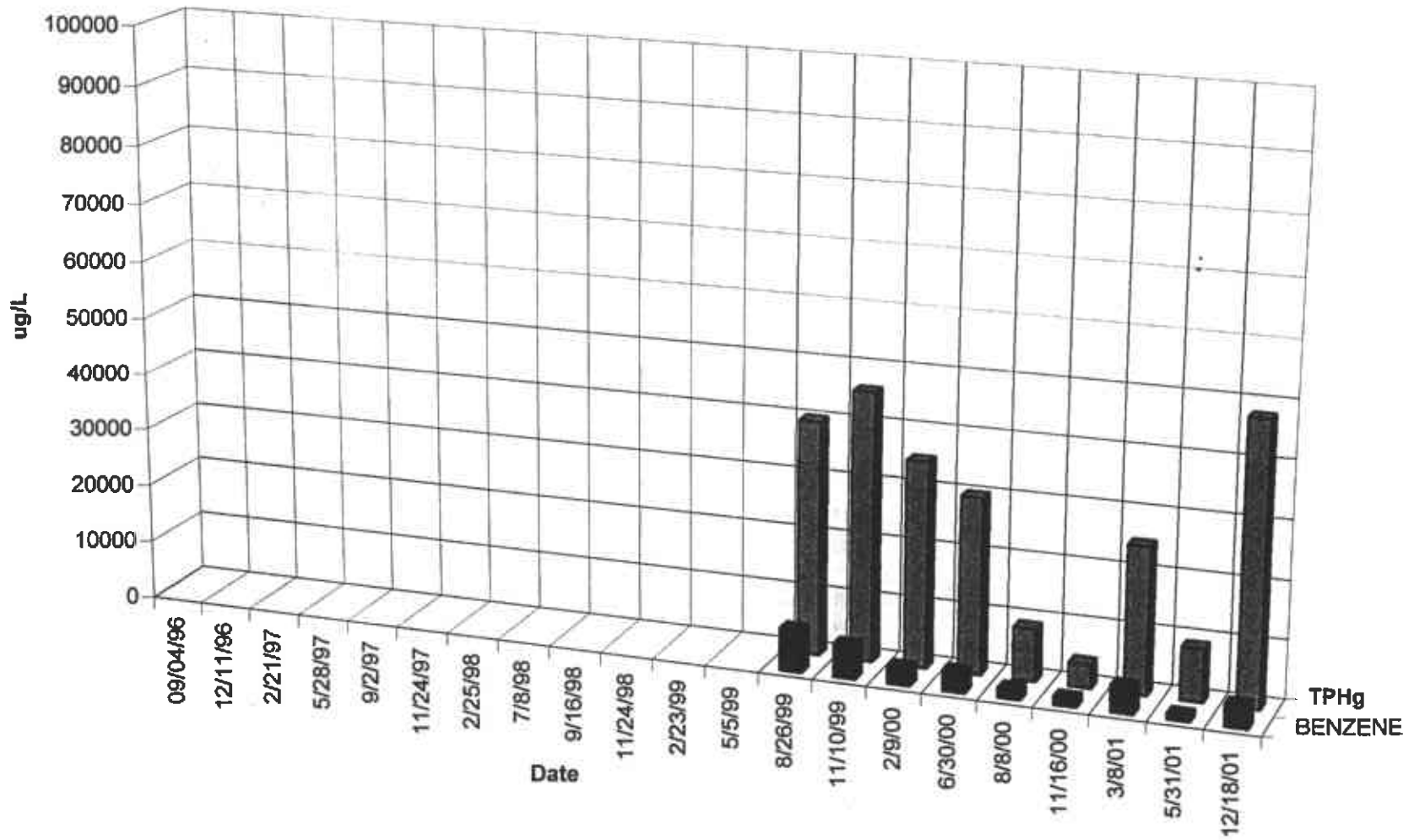
RS-9



RS-10

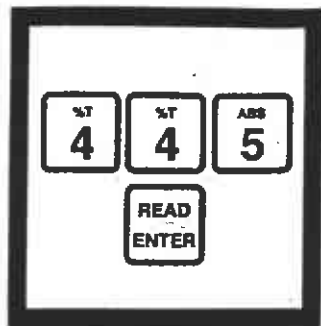


T-1



APPENDIX E
HACH SPECTROPHOTOMETER
FIELD TEST METHODS

HRDO Method



1. Enter the stored program number for dissolved oxygen.

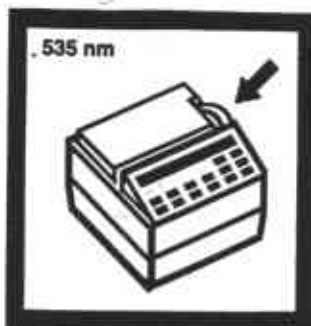
Press: **4 4 5 READ/ENTER**

The display will show:
DIAL nm TO 535

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

Note: Samples must be analyzed on site and cannot be stored; see Sampling and Storage following these steps.



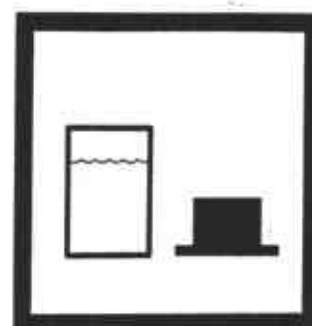
2. Rotate the wavelength dial until the small display shows:

535 nm



3. Press: **READ/ENTER**

The display will show:
mg/l O₂ HRDO



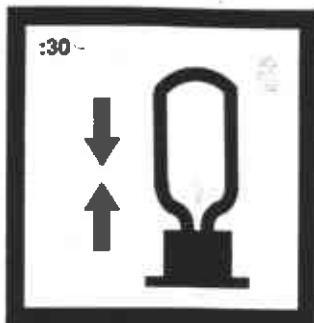
4. Fill a zeroing vial (the blank) with at least 10 mL of sample. Fill a blue ampul cap with sample.

OXYGEN, DISSOLVED, HR, continued



5. Fill a High Range Dissolved Oxygen AccuVac Ampul with sample.

Note: Keep the tip immersed while the ampul fills completely.



6. Without inverting the ampul, immediately place the ampul cap that has been filled with sample securely over the tip of the ampul. Shake the ampul for approximately 30 seconds.

Note: A small amount of the undissolved HRDO Reagent does not affect results.

Note: The cap prevents contamination with atmospheric oxygen.



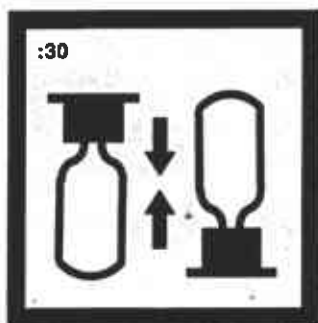
7. Press: **SHIFT TIMER**

A 2-minute reaction period enables oxygen, which was degassed during aspiration, to redissolve and react.



8. Place the AccuVac Vial Adapter into the cell holder.

Note: Place the grip tab at the rear of the cell holder.



9. When the timer beeps, the display will show:
mg/l O₂ HRDO
Shake the ampul for 30 seconds.



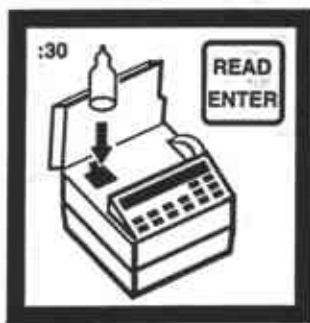
10. Place the blank into the cell holder. Close the light shield.



11. Press: **ZERO**

The display will show:
WAIT

then:
0.0 mg/l O₂ HRDO



12. Place the AccuVac ampul into the cell holder. Close the light shield. Wait approximately 30 seconds for the air bubbles to disperse from the light path.

Press: **READ/ENTER**

The display will show:
WAIT
then the result in mg/L dissolved oxygen will be displayed.

Note: In the constant-on mode, pressing READ/ENTER is not required. WAIT will not appear. When the display stabilizes, read the result.

SAMPLING AND STORAGE

The primary consideration in sampling with the High Range Dissolved Oxygen Ampul is to prevent the sample from becoming contaminated with atmospheric oxygen. This is accomplished by capping the ampul with an ampul cap in the interval between breaking open the ampul and reading the absorbance. If the ampul is securely capped, the ampul should be safe from contamination for several hours. The absorbance will decrease by approximately 3% during the first hour and will not change significantly afterwards.

Sampling and sample handling are important considerations in obtaining meaningful results. The dissolved oxygen content of the water being tested can be expected to change with depth, turbulence, temperature, sludge deposits, light, microbial action, mixing, travel time and other factors. A single dissolved oxygen test rarely reflects the accurate over-all condition of a body of water. Several samples taken at different times, locations and depths are recommended for most reliable results. Samples must be tested immediately upon collection although only a small error results if the absorbance reading is taken several hours later.

ACCURACY CHECK

The results of this procedure may be compared with the results of a titrimetric procedure or dissolved oxygen meter.

PRECISION

In a single laboratory, using a standard solution of 7.22 mg/L O₂ determined by the Winkler method and two representative lots of reagent with the DR/2000, a single operator obtained a standard deviation of ±0.20 mg/L O₂.

INTERFERENCES

The following do not interfere at a level of 10 mg/L which is in excess of naturally occurring levels: Cr³⁺, Mn²⁺, Fe²⁺, Ni²⁺, Cu²⁺ and NO₂⁻.

SUMMARY OF METHOD

The High Range Dissolved Oxygen AccuVac ampul contains reagent that is vacuum sealed in a 12-mL ampul. When the AccuVac ampul is broken open in a sample containing dissolved oxygen, it forms a yellow color which turns purple. The purple color development is proportional to the concentration of dissolved oxygen.

REQUIRED REAGENTS

Description	Quantity Required Per Test	Units	Cat. No.
High Range Dissolved Oxygen AccuVac ampuls, with 2 reusable ampul caps	1 ampul	25/pkg	25150-25

REQUIRED APPARATUS

AccuVac Dissolved Oxygen Sampler	1	each	24051-00
Adapter, AccuVac Vial	1	each	43784-00
Beaker, 50 mL	1	each	500-41
Caps, ampul, blue	varies	6/pkg	1731-06
Vial, zeroing	1	each	21228-00

OPTIONAL APPARATUS

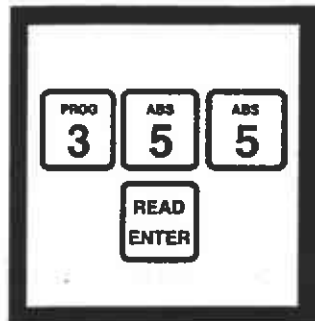
AccuVac Snapper Kit		each	24052-00
BOD bottle and stopper, 300 mL		each	621-00

Dissolved oxygen may also be determined by titrimetric methods. Request Publication 8042 for additional information.

For additional ordering information, see final section.
In the U.S.A. call 800-227-4224 to place an order.

Cadmium Reduction Method (Powder Pillows or AccuVac Ampuls)

USING POWDER PILLOWS



1. Enter the stored program number for high range nitrate nitrogen (NO₃⁻-N)—powder pillows.

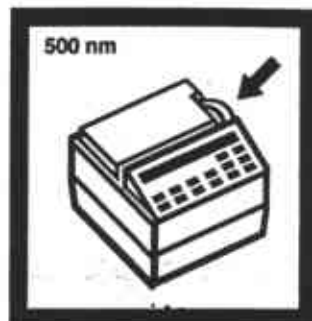
Press: **3 5 5 READ/ENTER**

The display will show:
DIAL nm TO 500

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

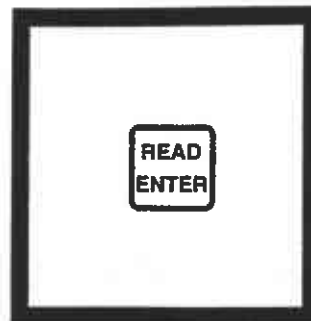
Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

Note: If sample cannot be analyzed immediately, see Sampling and Storage following these steps. Adjust the pH of stored samples before analysis.



2. Rotate the wavelength dial until the small display shows:

500 nm



3. Press: **READ/ENTER**

The display will show:
mg/l N NO₃⁻ H



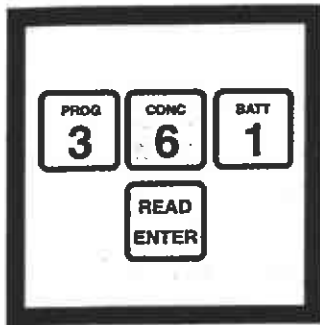
4. Fill a sample cell with 25 mL of sample.

Note: For proof of accuracy, use a 10 mg/L Nitrate Nitrogen Standard Solution (listed under Optional Reagents) in place of the sample.

Note: A reagent blank must be determined on each new lot of NitraVer 5. Perform Steps 4 to 12 using deionized water as the sample. Subtract this value from each result obtained with this lot of reagent.

*For seawater, a manual calibration is required; see Interferences.

USING ACCUVAC AMPULS



1. Enter the stored program number for high range nitrate nitrogen (NO_3^- -N)-AccuVac ampuls.

Press: **3 6 1 READ/ENTER**

The display will show:
DIAL nm TO 500

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

Note: If your instrument does not have program number 361, see Instrument Setup following these steps.



2. Rotate the wavelength dial until the small display shows:

500 nm



3. Press: **READ/ENTER**

The display will show:
mg/l N NO_3^- H AV

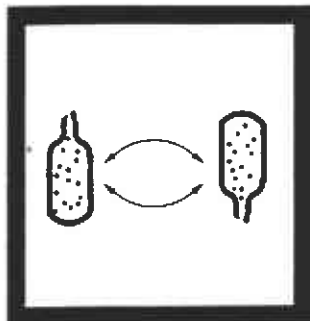


4. Collect at least 40 mL of sample in a 50-mL beaker. Fill a NitraVer 5 Nitrate AccuVac Ampul with sample.

Note: Keep the tip immersed while the ampul fills completely.

Note: For proof of accuracy, use a 10 mg/L Nitrate Nitrogen Standard Solution (listed under Optional Reagents) in place of the sample.

Note: A reagent blank must be determined on each new lot of NitraVer 5. Repeat Steps 4 to 12 using deionized water as the sample. Subtract this value from each result obtained with this lot of reagent.



5. Press: SHIFT TIMER

A one-minute mixing period will begin. Invert the ampul repeatedly until the timer beeps. Wipe off any liquid or fingerprints.

Note: Shaking time and technique influence color development. For most accurate results, make successive tests on a 10 mg/L Nitrate Nitrogen Standard Solution listed under Optional Reagents. Adjust the shaking time to obtain the correct result.

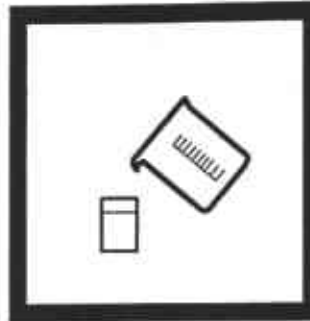


6. When the timer beeps, press: SHIFT TIMER

A 5-minute reaction period will begin.

Note: A deposit of unoxidized metal will remain after the NitraVer 5 Nitrate Reagent Powder dissolves. this deposit will have no effect on test results.

Note: An amber color will develop if nitrate nitrogen is present.



7. Fill a zeroing vial with at least 10 mL of sample (the blank).



8. Place the AccuVac Vial Adapter into the cell holder.

Note: Place the grip tab at the rear of the cell holder.



9. When the timer beeps, the display will show:

mg/l N NO₃⁻ H AV
Place the blank into the cell holder. close the light shield.



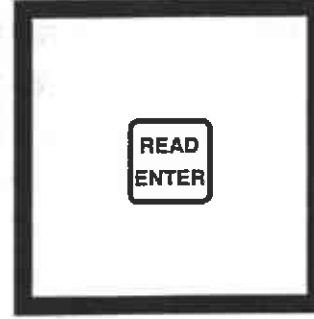
10. Press: ZERO

The display will show:
WAIT

then
0.0 mg/l N NO₃⁻ H AV



11. Place the AccuVac ampul into the cell holder. Close the light shield.



12. Press: READ/ENTER

The display will show:
WAIT
then the nitrate result in mg/L nitrate nitrogen (NO₃⁻-N) will be displayed.

Note: The results can be expressed as mg/L nitrate (NO₃⁻) by multiplying the mg/L nitrate nitrogen (NO₃⁻-N) by 4.4.

Note: In the constant-on mode, pressing READ/ENTER is not required. WAIT will not appear. When the display stabilizes, read the result.

ACCURACY CHECK

Standard Additions Method

a) Snap the neck off a fresh High Range Nitrate Nitrogen Voluette Ampule Standard, 500 mg/L NO_3^- -N.

b) Use the TenSette Pipet to add 0.1, 0.2, and 0.3 mL of standard to three 25-mL samples. Mix each thoroughly. (For AccuVac ampuls, use 50-mL beakers.)

c) Analyze each sample as described above. The nitrogen concentration should increase 2.0 mg/L for each 0.1 mL of standard added.

d) If these increases do not occur, see Standard Additions (Section I) for more information.

Standard Solution Method

Use a 10.0 mg/L Nitrate Nitrogen Standard Solution listed under Optional Reagents to check test accuracy. Or, this can be prepared by diluting 1.00 mL of solution from a High Range Nitrate Nitrogen Voluette Ampule Standard Solution, 500 mg/L NO_3^- -N, to 50.0 mL with deionized water.

PRECISION

In a single laboratory, using standard solutions of 20.0 mg/L nitrate nitrogen (NO_3^- -N) and two representative lots of reagent with the DR/2000, a single operator obtained a standard deviation of ± 0.8 mg/L nitrate nitrogen.

Using standard solutions of 30.0 mg/L (NO_3^- -N) and one representative lot of AccuVac ampuls with the DR/2000, a single operator obtained a standard deviation of ± 2.3 mg/L nitrate nitrogen.

INTERFERENCES

Compensate for nitrite interference as follows:

a) Add Bromine Water, 30 g/L, drop-wise to the sample in Step 4 until a yellow color remains.

b) Add one drop of Phenol Solution, 30 g/L, to destroy the color.

c) Proceed with Step 4. Report results as total nitrate and nitrite.

Strong oxidizing and reducing substances will interfere. Ferric iron causes high results and must be absent. Chloride concentrations above 100 mg/L will cause low results. The test may be used at high chloride levels (i.e., seawater), but a calibration must be performed using standards spiked to the same chloride concentration. See User Stored Programs in the DR/2000 Instrument Manual for more information.

Highly buffered samples or extreme sample pH may exceed the buffering capacity of the reagents and require sample pretreatment; see Interferences, pH (Section I).

SUMMARY OF METHOD

Cadmium metal reduces nitrates present in the sample to nitrite. The nitrite ion reacts in an acidic medium with sulfanilic acid to form an intermediate diazonium salt. This salt couples to gentisic acid to form an amber-colored product. Nitrate can be determined directly using the Nitrate Ion Selective Electrode (Cat. No. 44560-71).

REQUIRED REAGENTS (Using Powder Pillows)

Description	Quantity Required		Cat. No.
	Per Test	Units	
NitraVer 5 Nitrate Reagent Powder Pillows	1 pillow	50/pkg	14034-66

REQUIRED REAGENTS (Using AccuVac Ampuls)

NitraVer 5 Nitrate Reagent AccuVac Ampul	1 ampul	25/pkg	25110-25
--	---------	--------	----------

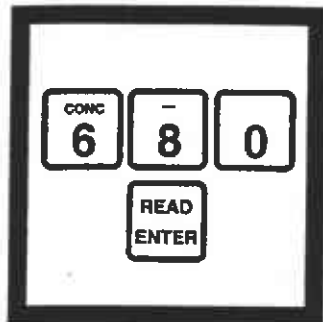
REQUIRED APPARATUS (Using Powder Pillows)

Clippers, for opening powder pillows	1	each	968-00
Stopper, rubber, size 2	1	12/pkg	2118-02

REQUIRED APPARATUS (Using AccuVac Ampuls)

Adapter, AccuVac Vial	1	each	43784-00
Beaker, 50 mL	1	each	500-41
Zeroing Vial	1	each	21228-00

SulfaVer 4 Method*, USEPA approved for reporting** USING POWDER PILLOWS



1. Enter the stored program for sulfate (SO_4^{2-}).

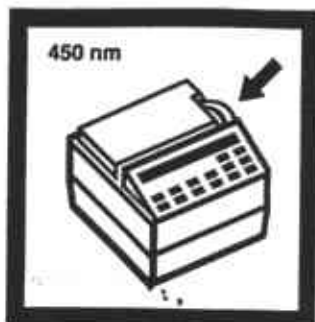
Press: **6 8 0 READ/ENTER**

The display will show:
DIAL nm to 450

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

Note: If samples cannot be analyzed immediately, see Sampling and Storage following these steps.



2. Rotate the wavelength dial until the small display shows:

450 nm

Note: For greater accuracy, prepare an instrument calibration for each new lot of SulfaVer 4 Sulfate Reagent Powder Pillows; see Calibration following these steps.



3. Press: **READ/ENTER**

The display will show:
mg/l SO_4^{2-}



4. Fill a clean sample cell with sample with 25 mL of sample.

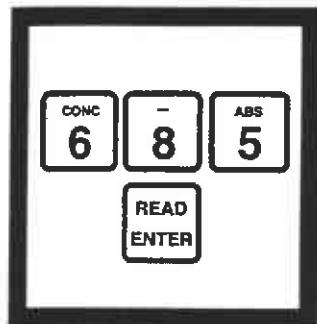
Note: Filter highly turbid or colored samples. Use filtered sample in this step and Step 6. Use labware listed under Optional Apparatus.

Note: For proof of accuracy, use a 50 mg/L sulfate standard solution (see Accuracy Check) in place of the sample.

*Adapted from *Standard Methods for the Examination of Water and Wastewater*

**Procedure is equivalent to USEPA method 375.4 for wastewater.

USING ACCUVAC AMPULS



1. Enter the stored program for sulfate (SO_4^{2-}) – AccuVac Ampuls.

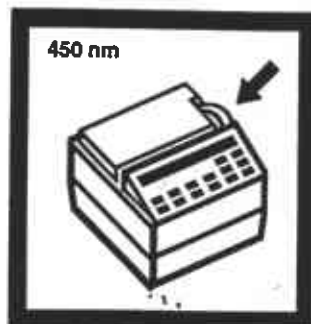
Press: **6 8 5 READ/ENTER**

The display will show:
DIAL nm to 450

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

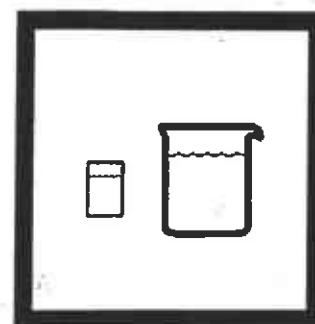
Note: If samples cannot be analyzed immediately, see Sampling and Storage following these steps.



2. Rotate the wavelength dial until the small display shows:
450 nm



3. Press: **READ/ENTER**
The display will show:
mg/l SO_4^{2-} AV



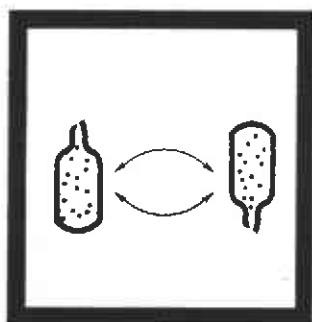
4. Fill a zeroing vial (the blank) with at least 10 mL of sample. Collect at least 40 mL of sample in a 50-mL beaker.

Note: Filter highly turbid or colored samples. Use filtered sample in this step and Step 5. Use labware listed under Optional Apparatus.



5. Fill a SulfaVer 4 Sulfate AccuVac ampul with sample.

Note: Keep tip immersed until the ampul fills completely.



6. Quickly invert the ampul several times to mix. Wipe off any liquid or fingerprints.

Note: A white turbidity will develop if sulfate is present.

Note: Accuracy is not affected by undissolved powder.



7. Press: **SHIFT TIMER**
A 5-minute reaction period will begin



8. Place the AccuVac Vial Adapter into the cell holder.

Note: Place the grip tab at the rear of the cell holder.



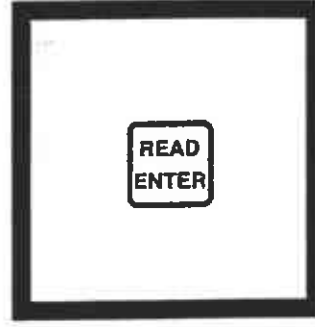
9. When the timer beeps, the display will show:
 $\text{mg/L SO}_4^{2-} \text{ AV}$
 Place the blank into the cell holder. Close the light shield.



10. Press: **ZERO**
 The display will show:
WAIT
 then:
 $0. \text{ mg/L SO}_4^{2-} \text{ AV}$



11. Within five minutes after the timer beeps, place the AccuVac ampul into the cell holder. Close the light shield.



12. Press: **READ/ENTER**
 The display will show:
WAIT
 then the results in mg/L SO_4^{2-} will be displayed.

Note: In the constant-on mode, pressing READ/ENTER is not required. WAIT will not appear. When the display stabilizes, read the result.

CALIBRATION

A new calibration should be performed for each new lot of SulfaVer 4 Sulfate Reagent Powder Pillows as follows:

a) Prepare standards of 0, 10, 20, 30, 40, 50 and 60 mg/L sulfate by diluting 0, 0.1, 0.2, 0.3, 0.4, 0.5 and 0.6 mL of the contents of a Sulfate Voluette Ampule Standard, 2500 mg/L , to 25 mL with deionized water in graduated mixing cylinders. Use a TenSette Pipet to measure the standard. Mix well.

Or, pipet 0, 1.0, 2.0, 3.0, 4.0, 5.0, and 6.0 mL of a 1000- mg/L sulfate Standard Solution into 100-mL volumetric flasks. Dilute to volume. Mix well. Transfer 25 mL to each test cylinder.

b) Store the calibration in the instrument memory using the procedure in the Operation section of the instrument manual. Follow the procedure described, choosing a wavelength of 450 nm, the decimal position as 0000, units as mg/L SO_4^{2-} and a Timer 1 interval of 05:00. Note the program number assigned to the procedure.

c) Add the reagents to the deionized water (0 standard-reagent blank) and to the 10 mg/L standard as described in Steps 4 to 6 of the powder pillow procedure above, using the deionized water blank to perform the zero calibration. Enter the sulfate concentration of the first standard (10 mg/L) and measure the absorbance as directed by the instrument manual. React and measure the remaining standards.

d) Use this stored program number in the powder pillow procedure above. Prepare a new calibration for each new lot of reagent, using the same stored program number.

SAMPLING AND STORAGE

Collect samples in clean plastic or glass bottles. Samples may be stored up to 7 days by cooling to 4 °C (39 °F) or lower. Warm to room temperature before analysis.

ACCURACY CHECK

Standard Additions Method

a) Snap the neck off a Sulfate Voluette Ampule Standard Solution, 2500 mg/L .

b) Use a TenSette Pipet to add 0.1, 0.2 and 0.3 mL of standard to three 25-mL samples. Mix thoroughly. For AccuVac Ampuls, use 50-mL beakers.

c) Analyze each sample as described above. The sulfate concentration should increase 10 mg/L for each 0.1 mL of standard added.

d) If these increases do not occur, see Standard Additions (Section I) for more information.

Standard Solution Method

Check the accuracy of the test by using the Sulfate Standard Solution, 50 mg/L , listed under Optional Reagents. Or, prepare this solution by pipetting 1.0 mL of the contents of a Voluette Ampule Standard for Sulfate into a 50-mL volumetric flask. Dilute to volume with deionized water.

PRECISION

In a single laboratory, using a standard solution of 50 mg/L sulfate and two representative lots of powder pillows with the DR/2000, a single operator obtained a standard deviation of ± 0.9 mg/L sulfate.

In a single laboratory, using a standard solution of 50 mg/L sulfate and two representative lots of AccuVac ampuls with the DR/2000, a single operator obtained a standard deviation of ± 2.2 mg/L sulfate.

INTERFERENCES

The following interfere at levels above those concentrations listed:

Calcium	20,000 mg/L as CaCO ₃
Chloride	40,000 mg/L as CaCO ₃
Magnesium	10,000 mg/L as CaCO ₃
Silica	500 mg/L as CaCO ₃

SUMMARY OF METHOD

Sulfate ions in the sample react with barium in the SulfaVer 4 and form a precipitate of barium sulfate. The amount of turbidity formed is proportional to the sulfate concentration. The SulfaVer 4 also contains a stabilizing agent to hold the precipitate in suspension.

REQUIRED REAGENTS (Using Powder Pillows)

Description	Quantity Required Per Test	Units	Cat. No.
SulfaVer 4 Sulfate Reagent Powder Pillows	1 pillow	50/pkg	12065-66

REQUIRED REAGENTS (Using AccuVac Ampuls)

SulfaVer 4 Sulfate AccuVac Ampuls	1 ampul	25/pkg	25090-25
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REQUIRED APPARATUS (Using Powder Pillows)

Clippers, for opening powder pillows	1	each	968-00
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REQUIRED APPARATUS (Using AccuVac Ampuls)

Adapter, AccuVac Vial	1	each	43784-00
Brush	1	each	690-00
Vial, zeroing	1	each	21228-00

OPTIONAL REAGENTS

Sulfate Standard Solution, 50 mg/L	500 mL	2578-49
Sulfate Standard Solution, 1000 mg/L	500 mL	21757-49
Sulfate Standard Solution, Voluette Ampule, 2500 mg/L, 10 mL	16/pkg	14252-10
Water, deionized	3.78 L	272-17

OPTIONAL APPARATUS

AccuVac Snapper Kit	each	24052-00
Ampule Breaker Kit	each	21968-00
Beaker, 50 mL	each	500-41
Filter Paper, folded, 12.5 cm	100/pkg	1894-57
Flask, volumetric, 50 mL, Class A	each	14574-41
Flask, volumetric, 100 mL, Class A	each	14574-42
Funnel, poly, 65 mm	each	1083-67
Pipet, TenSette, 0.1 to 1.0 mL	each	19700-01
Pipet Tips, for 19700-01 Pipet	50/pkg	21856-96
Pipet, volumetric, Class A, 1.00 mL	each	14515-35
Pipet Filler, safety bulb	each	14651-00

For additional ordering information, see final section.
In the U.S.A. call 800-227-4224 to place an order.

IRON, FERROUS (0 to 3.00 mg/L)

For water, wastewater and seawater

**1,10 Phenanthroline Method* (Powder Pillows or AccuVac Ampuls)
USING POWDER PILLOWS**

1. Enter the stored program number for ferrous iron, (Fe²⁺)-powder pillows.

Press: **2 5 5 READ/ENTER**

The display will show:
DIAL nm TO 510

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

Note: Analyze samples as soon as possible to prevent air oxidation of ferrous iron to ferric iron, which is not determined.



2. Rotate the wavelength dial until the small display shows:

510 nm



3. Press: **READ/ENTER**

The display will show:
mg/l Fe²⁺



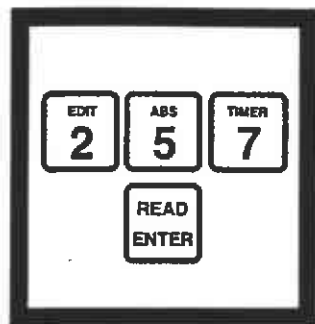
4. Fill a sample cell with 25 mL of sample.

Note: For proof of accuracy, use a 1.0 mg/L ferrous iron standard solution (preparation given in the Accuracy Check) in place of the sample.

*Adapted from *Standard Methods for the Examination of Water and Wastewater*

IRON, FERROUS, continued

USING ACCUVAC AMPULS



1. Enter the stored program number for ferrous iron (Fe^{2+})—AccuVac ampuls.

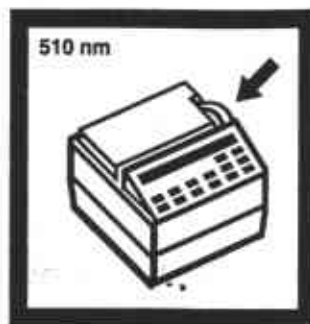
Press: **2 5 7 READ/ENTER**

The display will show:
DIAL nm TO 510

Note: DR/2000s with software versions 3.0 and greater will display "P" and the program number.

Note: Instruments with software versions 3.0 and greater will not display "DIAL nm TO" message if the wavelength is already set correctly. The display will show the message in Step 3. Proceed with Step 4.

Note: Analyze samples as soon as possible to prevent air oxidation of ferrous iron to ferric iron, which is not determined.



2. Rotate the wavelength dial until the small display shows:

510 nm



3. Press: **READ/ENTER**

The display will show:
mg/l Fe^{2+} AV



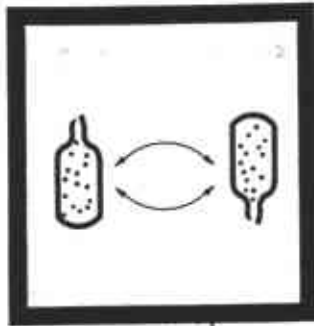
4. Fill a zeroing vial (the blank) with at least 10 mL of sample. Collect at least 40 mL of sample in a 50-mL beaker.

Note: For proof of accuracy, a 1.0 mg/L ferrous iron standard solution (preparation given in the Accuracy Check) can be used in place of the sample.



5. Fill a Ferrous Iron AccuVac Ampul with sample.

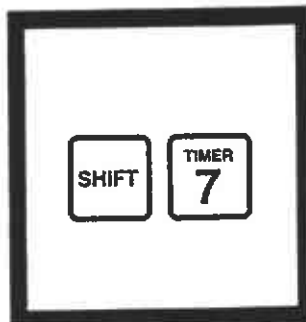
Note: Keep the tip immersed while the ampul fills completely.



6. Quickly invert the ampul several times to mix. Wipe off any liquid or fingerprints.

Note: An orange color will form if ferrous iron is present.

Note: Undissolved powder does not affect accuracy.



7. Press: **SHIFT TIMER**

A 3-minute reaction period will begin.



8. Place the AccuVac Vial Adapter into the cell holder.

Note: Place the grip tab at the rear of the cell holder.



9. When the timer beeps, the display will show:
mg/l Fe²⁺ AV
Place the blank into the cell holder. Close the light shield.



10. Press: **ZERO**

The display will show:
WAIT
then:
0.00 mg/l Fe²⁺ AV



11. Place the AccuVac ampul into the cell holder. Close the light shield.



12. Press: **READ/ENTER**

The display will show:
WAIT
then the result in mg/L Fe²⁺ will be displayed.

Note: In the constant-on mode, pressing READ/ENTER is not required. WAIT will not appear. When the display stabilizes, read the result.

ACCURACY CHECK

Standard Solution Method

Prepare a ferrous iron stock solution (100 mg/L Fe) by dissolving 0.7022 grams of ferrous ammonium sulfate, hexahydrate, in deionized water. Dilute to 1 liter. Prepare immediately before use. Dilute 1.00 mL of this solution to 100 mL with deionized water to make a 1.0 mg/L standard solution. Prepare this immediately before use.

PRECISION

In a single laboratory, using an iron standard solution of 1.000 mg/L Fe²⁺ and two representative lots of reagent with the DR/2000, a single operator obtained a standard deviation of ±0.006 mg/L Fe²⁺.

In a single laboratory using a standard solution of 1.000 mg/L Fe²⁺ and two representative lots of AccuVac ampuls with the DR/2000, a single operator obtained a standard deviation of ±0.009 mg/L Fe²⁺.

SUMMARY OF METHOD

The 1,10 phenanthroline indicator in Ferrous Iron Reagent reacts with ferrous iron in the sample to form an orange color in proportion to the iron concentration. Ferric iron does not react. The ferric iron (Fe³⁺) concentration can be determined by subtracting the ferrous iron concentration from the results of a total iron test.

REQUIRED REAGENTS (Using Powder Pillows)

Description	Quantity Required Per Test	Units	Cat. No.
Ferrous Iron Reagent Powder Pillows	1 pillow	100/pkg	1037-69

REQUIRED REAGENTS (Using AccuVac Ampuls)

Ferrous Iron Reagent AccuVac Ampuls	1 ampul	25/pkg	25140-25
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REQUIRED APPARATUS (Using Powder Pillows)

Clippers, for opening powder pillows	1	each	968-00
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REQUIRED APPARATUS (Using AccuVac Ampuls)

Adapter, AccuVac Vial	1	each	43784-00
Beaker, 50 mL	1	each	500-41
Sample Cell, 10 mL, with cap	1	each	21228-00

OPTIONAL REAGENTS

Ferrous Ammonium Sulfate, hexahydrate, ACS	113 g		11256-14
Water, deionized	3.78 L		272-17

OPTIONAL APPARATUS

AccuVac Snapper Kit		each	24052-00
Clippers, shears, 7-1/4"		each	23694-00
Flask, volumetric, 100 mL, Class B		each	547-42
Flask, volumetric, 1000 mL, Class B		each	547-53
Pipet, volumetric, 1 mL		each	515-35
Pipet Filler, safety bulb		each	14651-00
Pour-Thru Cell Assembly Kit		each	45215-00

For additional ordering information, see final section.
In the U.S.A. call 800-227-4224 to place an order.

APPENDIX F.

WASTEWATER DISCHARGE PERMIT RENEWAL.

 **EAST BAY COMPLIANCE EVENT REMINDER NOTICE**
MUNICIPAL UTILITY DISTRICT

DAVID R. WILLIAMS
DIRECTOR OF WASTEWATER

December 3, 2001

DESERT PETROLEUM, INC.
P.O. Box 1601
Oxnard, CA 93032
Attention: George Converse

Re: Wastewater Discharge Permit No. 50435501

As a compliance reporting condition of your Wastewater Discharge Permit, you are required to submit periodic reports. The purpose of this letter is to remind you that your facility must submit a(n) Self-Monitoring Report by 01/31/02.

The report must contain the proper certification statement and be signed by an authorized person. Details concerning these items may be found in Section B of STANDARD TERMS AND CONDITIONS. This report must be completed and mailed to the Source Control Division by this date. A violation follow up fee may be assessed for late, incomplete or failure to submit this report.

Sincerely,



Marie A. Kulka
Wastewater Control Representative
(510) 287-1632

EBMUD - Mail Slot #702
Source Control Division
P.O. Box 24055
Oakland, CA 94623-1055

cc: George Converse
1386 East Beamer Street
Woodland, CA 95776



WESTERN
GEO-ENGINEERS
CALIF. CONTRACTOR #513857
REGISTERED GEOLOGISTS

1386 EAST BEAMER STREET
WOODLAND CA 95776-6003
532 (916) 668-5300,
FAX (916) 662-0273
Wege@mother.com

December 5, 2001

Mr. Bill Thompson
DESERT PETROLEUM
P.O. Box 1601
Oxnard, Ca 93032

RE: FORMER DESERT #793. 4035 PARK BLVD., OAKLAND, CALIFORNIA.

Dear Bill:

Enclosed please find the renewal application for wastewater discharge permit for the above referenced site. Please review, sign, date and mail back to us. If you have any questions please call.

Sincerely,

George L. Converse
Project Geologist

Enclosure:



WASTEWATER DISCHARGE PERMIT APPLICATION CHECKLIST AND CERTIFICATION

Permit: Wastewater Discharge

Permit No.: 50435501 Renewal

Check the appropriate box for each page of the Wastewater Discharge Permit Application. If you are completing a page and including it in the application, initial the box in the "Page Completed" column. If you are able to certify that no change has occurred to a particular page from the previous application, initial the box in the "No Change" column. Sign and date the checklist/certification. Submit the checklist/certification with your application package.

Page No.	Page Title	Initial if Page Completed	Initial if No Change
1	Applicant Information	<i>SEC</i>	
2	Process Description	<i>SEC</i>	
3	Schematic Flow Diagram	<i>SEC</i>	
4	Building Layout Diagram	<i>SEC</i>	<i>SEC</i>
5	Water Source and Use ("Water Balance")	<i>SEC</i>	
6	Strength Summary	<i>SEC</i>	

I certify under penalty of law that by initialing a page(s) above in the "No Change" column that the most recently submitted application page fully describes conditions at the facility at the present time and for the expected duration of the renewed permit.

I certify under penalty of law that this document and all attachments were prepared under my direction or supervision in accordance with a system designed to assure that the qualified personnel properly gather and evaluate the information submitted. Based on my inquiry of the person or persons who manage the system, or those persons directly responsible for gathering information, the information submitted is, to the best of my knowledge and belief, true, accurate, and complete. I am aware that there are significant penalties for submitting false information, including the possibility of fine and imprisonment for knowing violations.

Gary Corneil / Project Manager
Signature/Title

12-5-01
Date




WASTEWATER DISCHARGE PERMIT

Terms and Conditions

PERMIT NUMBER 50435501

APPLICANT INFORMATION

APPLICANT BUSINESS NAME Desert Petroleum	
PERSON TO BE CONTACTED IN EVENT OF EMERGENCY George Converse <small>Name</small> (530) 668-5300 <small>Day Phone</small> (530) 668-5300 m <small>Night Phone</small> (530) 662-0273 <small>Fax Number</small>	ADDRESS OF PREMISES DISCHARGING WASTEWATER 4035 Park Blvd. <small>Street Address</small> Oakland, CA 94602 <small>City</small> <small>Zip Code</small>
PERSON TO BE CONTACTED ABOUT THIS APPLICATION George Converse <small>Name</small> Project Manager <small>Title</small> (530) 668-5300 (530) 662-0273 <small>Day Phone</small> <small>Fax Number</small>	FACILITY MAILING ADDRESS P.O. Box 1601 <small>Street Address</small> Oxnard, CA 93002 <small>City</small> <small>Zip Code</small> almccowen@aol.com <small>Electronic Mail Address (E-Mail)</small>
CHIEF EXECUTIVE OFFICER/DULY AUTHORIZED REPRESENTATIVE Mr. Bill Thompson President <small>Name (printed)</small> <small>Title</small> 2060 Knoll Drive, Suite 100 Ventura, CA 93003 <small>Street Address</small> <small>City</small> <small>Zip Code</small>	
CERTIFICATION	
<p>I certify under penalty of law that this document and all attachments were prepared under my direction or supervision in accordance with a system designed to assure that the qualified personnel properly gather and evaluate the information submitted. Based on my inquiry of the person or persons who manage the system, or those persons directly responsible for gathering information, the information submitted is, to the best of my knowledge and belief, true, accurate, and complete. I am aware that there are significant penalties for submitting false information, including the possibility of fine and imprisonment for knowing violations.</p>	
 <small>Signature (see certification requirements on reverse)</small>	
12/12/01 <small>Date</small>	



WASTEWATER DISCHARGE PERMIT

Terms and Conditions
PROCESS DESCRIPTION

FACILITY NAME DP793

PURPOSE - The Process Description is intended to provide a description of the primary business activities and the substances which may enter into the wastewater from the business activity. Permit Number
50435501

BUSINESS ACTIVITY Groundwater Remediation	Standard Industrial Classification	Business Classification Code
---	------------------------------------	------------------------------

TYPE OF PRODUCT OR BRAND NAME	QUANTITIES - INDICATE UNITS			
	Past Year		Estimated This Year	
	Mo. Year	Mo. Year	Mo. Year	Mo. Year
Extracted Groundwater	7 / 00	7 / 01	12 / 01	12 / 02
	116,438 gallons		2450 gallons	

Process Description <small>List all wastewater generating operations</small>	Characteristics <small>List all substances that may be discharged to the sewer</small>	Process Number <small>From Schematic</small>
Pump Groundwater	Benzene, Toluene, Ethylbenzene, Xylene, TPH gasoline	1

PRETREATMENT FACILITIES

Pretreatment: Check the type of treatment, if any, given wastewater before it is discharged to the community sewer:

None
 holding tank
 grease trap
 oil and water separator
 grinding
 sedimentation
 pH adjustment
 biological treatment
 screening
 chlorination
 other (describe) activated carbon

Description: Describe the loading rates, design capacity, physical size, etc. of each pretreatment facility checked above. Identify the side sewer to which treated wastewater is discharged.

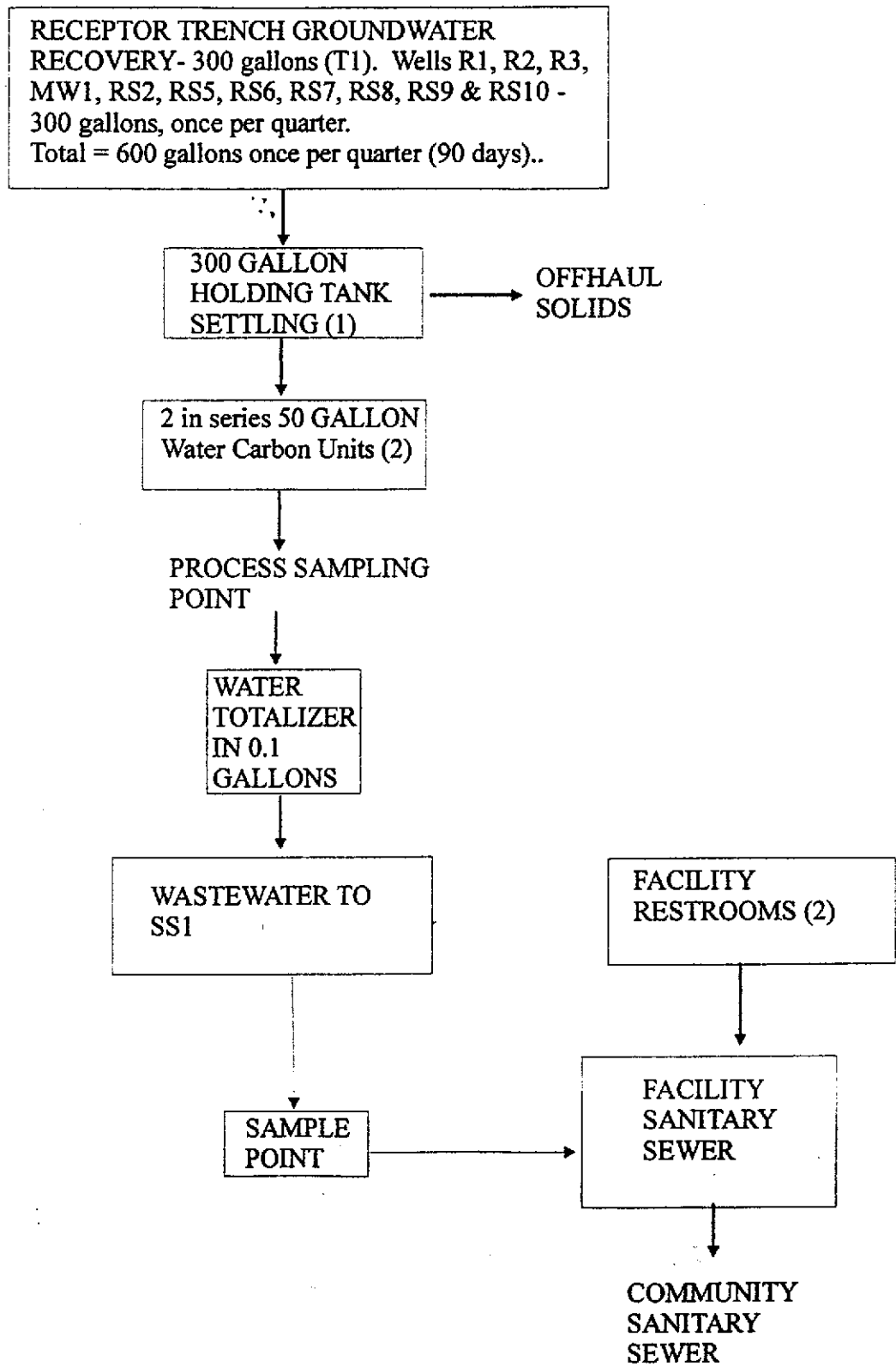
300 gallon capacity poly tank feeds 2 in series 50 gallon (160lbs carbon each) at 5gpm, gravity flow discharge to side sewer SS1 (see figure)

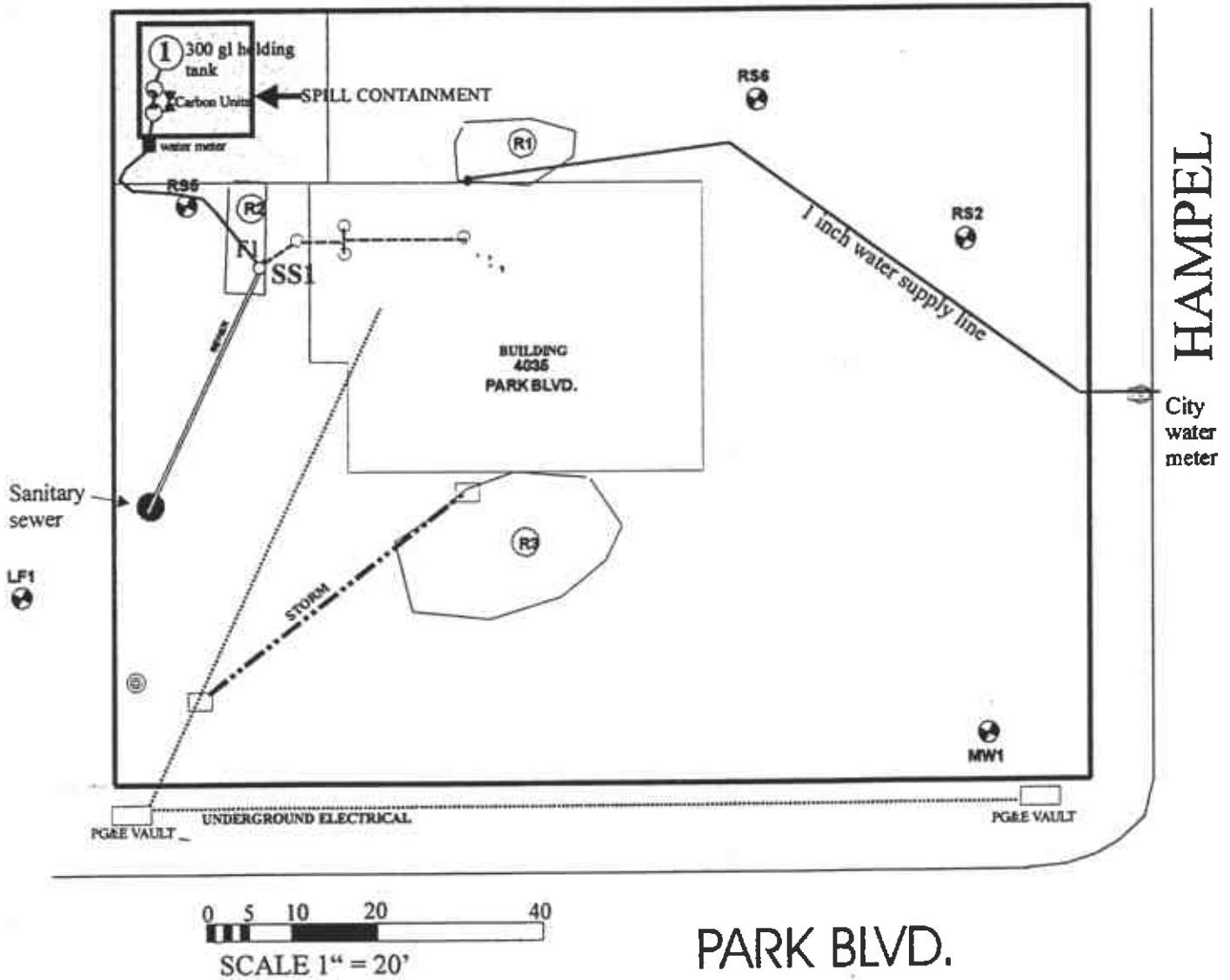
OTHER WASTES: List the type and volume of liquid waste and sludge removed from the premises by means other than the community sewer.

Facility EPA Generator I.D. Number _____

Waste removed by Name, address, State Transporter I.D. No.	Type of Waste Example: Alkaline cleaners, Organic solvents	EPA Waste No.	State Waste No.	Quantity generated lbs. or gal. /month

Figure 1 (Revised December 5, 2001)
Activity: GROUNDWATER RECOVERY AND DISCHARGE SYSTEM
FORMER DESERT PETROLEUM SITE DP 793.






-  MONITOR WELL
- 1 300 gallon poly tank
- 2 2 in series 55 gallon carbon filters.

FIGURE 2
SEWER DISCHARGE
TREATMENT COMPOUND
WASTEWATER DISCHARGE
PERMIT # 5043550 1



WASTEWATER DISCHARGE PERMIT

Terms and Conditions

WATER SOURCE AND USE

FACILITY NAME DP793

PURPOSE: This information will enable EBMUD to evaluate the volumes and source(s) of wastewater discharged to the community sewer.

Permit Number
50435501

Water Use and Disposition Estimate the average quantity of water received and wastewater discharged daily.
NOTE: Show on a separate sheet the **METHOD AND CALCULATIONS** used to determine the quantities shown on the table.

WATER USED FOR:	Supply From			Discharged To		
	EBMUD gal/day	Other (1) gal/day	code	Community Sewer gal/day	Other (2) gal/day	code
SANITARY						
PROCESSES						
BOILER						
COOLING						
WASHING						
IRRIGATION						
OTHER (3)		600	A	600		
TOTAL	0			0		

Notes:

- Enter the quantity and the appropriate code letter indicating the source:
a. well b. creek c. estuary d. bay e. stormwater f. reclaimed water
- Enter the quantity and the appropriate code letter indicating the discharge point:
a. well b. creek c. estuary d. bay e. stormdrain f. rail, truck, barge g. evaporation h. product
- Describe: Pumping wells during groundwater sampling events

Total Number of Employees Total 0

	Office		Production (number of employees per shift)					
	No.	Hours	Day Shift		Swing shift		Night shift	
			No.	Hours	No.	Hours	No.	Hours
Weekday		to		to		to		to
Saturday		to		to		to		to
Sunday		to		to		to		to

Source of Wastewater Discharged

Water Meter Number	Use Code (see reverse)	Percent (%) discharged to: Side Sewer									Total % Disch. to all side sewers
		No. 1	No. 2	No. 3	No. 4	No. 5	No. 6	No. 7	No. 8	No. 9	
47083426	w	100%									100%



WASTEWATER DISCHARGE PERMIT

Terms and Conditions

STRENGTH SUMMARY

FACILITY NAME DF793

PURPOSE: This information will identify for EBMUD the variation in flow rate and the type of constituents and characteristics of the discharge for each side sewer.

Permit Number
50435501

Side Sewer No. SS1 Side Sewer Location west side of building

Wastewater Flow Rate

Peak Hourly (gallons/minute)	Maximum Daily (gallons/day)	Annual Daily Average (gallons/day)	Max. Monthly (CCF *)
5	650	7	1.0

* CCF = hundred cubic feet = 748 gallons

Discharge Frequency

<p>Discharge Period</p> <p><input type="checkbox"/> Continuous <input type="checkbox"/> 24 hrs./day</p> <p><input type="checkbox"/> 365 day/year, or</p> <p>a. Time of day from _____ to _____</p> <p>b. Days of the week _____</p>	<p>Batch Discharge(s)</p> <p>a. Day(s) of the week <u>Tue, Wed, Thb.</u> Time(s) of the day <u>noon-2pm</u></p> <p>c. Volume discharged <u>600g1</u> d. Rate of Discharge <u>5gpm</u></p>
---	---

Stormwater Area - Total area in square feet exposed to stormwater, rainwater, and groundwater and draining to this side sewer
0 or 11,000 sq. ft. There is one storm drain for 11,000 sq. ft. of property
Storm drain may or may not be part of sewer.

Wastewater Strength Estimates - Enter the average annual and maximum wastewater strength for this side sewer for each of the following elements of wastewater strength for the period covered by the Permit. These values will become the basis for sewage disposal charges and are the average and maximum limits on the elements of the discharger's wastewater strength.

Elements of Wastewater Strength	Unit	Average	Maximum
Total Suspended Solids (TSS)	mg/L	N/A	N/A
Filtered Chemical Oxygen Demand (CODF)	mg/L	N/A	N/A

Provide the name and address of the laboratory and the State of California, Department of Health Services, Environmental Laboratory Accreditation Program Certificate Number of the laboratory performing self-monitoring analyses.

Name Kiff Analytical LLC Telephone (530) 297-4800

Street 720 Olive Drive, Suite D City Davis State CA Zip 95616

Certificate Number 2236

APPENDIX G.

WORKPLAN (amended) – AUGMENT BIODEGRADATION

WORKPLAN TO AUGMENT BIODEGRADATION
BY INTRODUCING
OXYGEN
INTO SELECTED WELLS

August 29, 2000
Amended January 7, 2002

Mr. Bill Thompson
Desert Petroleum
P.O. Box 1601
Oxnard, California 93032
(805) 644-6784 FAX (805) 654-0720

Dear Mr. Thompson:

The following is the amended Workplan for oxygen augmentation into wells R3, RS5, RS7 and trench wells T1, T2, T3 and T4 at former Desert Petroleum Station #793.

1.0 SITE LOCATION AND DESCRIPTION

Former Desert Petroleum #793 is a non-active service station, located on the northwest corner of the intersection of Park Boulevard and Hampel Street at 4035 Park Blvd., Oakland, California. The site is located in projected section 32; T1S; R3W; MDB&M at an approximate elevation of 210 feet above mean sea level.

2.0 INTRODUCTION

The following amends the August 29, 2000 workplan to augment the ongoing natural bioremediation at former Desert Petroleum Inc. station DP 793. Gasoline was discovered trickling into a sewer manway on Brighton Avenue on November 30, 1989. The station was closed and all contents removed from the tanks by December 7, 1989. The UST's were removed on June 23, 1994 and over-excavation of on-site contaminated soils occurred August 14, 1995. Various assessments have delineated the gasoline plume and show that the gasoline release followed the sewer main from the station to Brighton Avenue sewer manway. A receptor trench was installed along Brighton Avenue on August 12, 1999. On August 26, 1999 during the 1/4ly sampling round selected wells were also tested for the potential of natural biodegradation, see Western Geo-Engineers report "Further Assessment, Installation of Brighton Avenue Receptor Trench and 3rd Quarter 1999 Groundwater monitoring". The August 29, 2000 workplan was developed to enhance the natural biodegradation that is occurring along the parameters of the groundwater plume associated with this site. Groundwater pumped from the receptor trench and well RS5 was treated with activated carbon and dumped to the sanitary sewer located at the site (4035 Park Blvd.). Comparing the groundwater plume prior to pumping and sewer discharge (August 1999) to the groundwater plume after pumping and sewer discharge (May 2001) showed a dramatic reduction in the groundwater plume. A meeting to discuss "if further remedial actions were warranted at or near the site" was held at Alameda County Health. Review and discussion of the actions that have been taken and the

results obtained indicated that nutrient augmentation was probably not warranted, but of greater benefit would be enhancement of the dissolved oxygen. This workplan has been developed to provide the means for enhancing dissolved oxygen near and within the hydrocarbon plume.

3.0 BIOREMEDIATION

Bacteria native to the soil at hydrocarbon contamination sites normally degrade hydrocarbons. The most effective hydrocarbon degraders (eaters) are the aerobic (oxygen using) bacteria. The amount of available dissolved oxygen is usually the factor controlling the rate that these bacteria degrade the gasoline.

A much slower degradation process starts when the dissolved oxygen is depleted. The plume begins to become anaerobic and the bacterium commences to reduce nitrate, ferric iron, and sulfate to further degrade the hydrocarbons. Eventually, as these compounds and the oxygen are used, the bacteria begin methogenesis, in which the hydrocarbons are converted to methane.

The results of prior bioremediation sampling (August and September 1999) indicate that natural attenuation/bioremediation is active at this site and methogenesis is occurring. In a number of the wells the biodegradation has proceeded to the point that nearly all of the electron acceptors and the nutrients that bacteria require to degrade gasoline have been consumed while a significant amount of hydrocarbons remain.

The most current bioremediation sampling (December 2001) shows similar results to that of the 1999 sampling but indicate that nutrient augmentation is not necessary at this time and oxygen enhancement would be beneficial.

All of the wells show the impact of active biodegradation. It is not therefore possible to determine a background level of the compounds. The closest approximation for background levels at this site is the highest concentrations of electron acceptors and the lowest levels of by-products.

Compound	Function	Concentration		Well
		1999	2001	
Dissolved Oxygen (O ₂)	Electron Acceptor	4.9 mg/l	5.5 mg/L	MW1 / R3
Nitrate	Electron Acceptor	2.7 mg/l	11.4 mg/L	RS2
Sulfate	Electron Acceptor	>77 mg/l	>77 mg/L	R2 & R3 / RS2, RS6, R2 & R3
Ferrous Iron	By-product	0.25 mg/l	0 mg/L	MW1
Methane	By-product	<0.00001 mg/l		MW1
Carbon Dioxide	By-product	0.058 mg/l		RS8
Aerobic Bacteria	By-product	10 CFU/ml		MW1

mg/L milligrams per liter (parts per million)
CFU/ml plate count per milliliter

All of the tested wells have reduced levels of dissolved oxygen.

Since the 1999 sampling Both Nitrate and Sulfate have rebounded, with Nitrate increasing from 0.8 to 6 mg/L in down gradient well RS7 and Sulfate increasing from non-detectable levels to 49 mg/L in well RS8.

The presence of Ferrous Iron and Methane in the wells indicates that biodegradation has progressed to the point that the system is oxygen deficient and the bacteria have started to reduce the iron to provide oxygen for the degradation. The Ferrous Iron outline has been reduced in size, indicating the groundwater near and associated with the hydrocarbon plume is not as oxygen deficient.

The carbon dioxide (CO₂) levels in the groundwater indicate that a portion of the hydrocarbons have been degraded. CO₂ and water are the final byproducts of the biodegradation of hydrocarbons. The carbon in CO₂ results from the oxidation of the hydrocarbon radical CH₂ and as such 1 mg of CO₂ = 0.41 mg of CH₂ (CH₂ (12+1+1 = 14) / (CO₂ (12+16+16=34).

The presence of methane (CH₄) indicates that a number of the wells have progressed into methogenesis.

The levels of electron acceptors present and the presence of the reaction products, carbon dioxide, methane and ferrous iron indicate that the bacteria in the soil and the compounds in the groundwater have the capability to consume a significant amount of hydrocarbons.

Comparison of the hydrocarbon degrader counts (amount of bacteria) to the TPHg concentration, electron acceptors, byproducts, and nutrients, indicate that the addition of oxygen can have a significant effect on the bioactivity, see Table 3 of Third Quarter 2000 report and Table 2 of Fourth Quarter 2001 report.

The current TPHg concentrations show that only two of the wells, compared to five wells in 1999, contain greater than 10 mg/l of TPHg with a high of 48 mg/L at T1. This indicates that it would be beneficial to increase the biodegradation capacity of the aquifer. The most cost beneficial way to do this is by adding dissolved oxygen.

The current groundwater plume covers an area of approximately 10160 square feet. The affected water bearing strata is an estimated 16 feet thick and extends from 10 to 26 feet below the surface at well RS5. The plume volume is 162,560 cubic feet. Assuming a porosity of 0.3, the plume contains 48,768 cubic feet, 364,785 gallons or 1,380,711 liters of water.

4.0 BIO -AUGMENTATION

4.1.1 Air Sparging

Do to the close proximity of residential multistory apartments to the treatment compound, mechanical equipment, such as air pumps, compressors, etc would create a noise problem. The alternative is to chemically add the oxygen to the groundwater.

4.1.2 Hydrogen Peroxide

A 21% solution of hydrogen peroxide can be handled without special permits. The solution would be metered and/or batch dumped into select wells. Special personnel protective equipment would be necessary for the individual monitoring and/or handling the hydrogen peroxide. Due to the highly oxidizing nature of hydrogen peroxide care against splash/spillage of the solution would be necessary along with labeling for emergency response personnel, i.e. fire and police etc. A special compound would need to be constructed to store the hydrogen peroxide containers, metering pumps and distribution lines.

4.1.3 Oxygen Release Compound

Oxygen Release Compound "(ORC) is a proprietary formulation of magnesium peroxide intercalated with food-grade phosphates." The ORC socks are placed into select wells and once exposed to the groundwater allow for a continual release of oxygen for a period of up to a year (dependent upon the groundwater conditions). These socks once placed do not need maintenance and special handling protective equipment is not required.

R2? RS2? R3?

Oxygen augmentation is suggested upgradient the station building at R1, upgradient of RS8 at RS5, along the receptor trench at T1, T2, T3 and T4 and upgradient of RS9 at RS7. Ideally this will cause the water near the well to become saturated with oxygen.

RS8, too?

4.1.4 NUTRIENT ADDITION

As found during the September 2, 1999 sampling round the phosphate and ammonia levels are reduced to levels that negatively effects the biodegradation rate. After approximately 6 months using ORC an evaluation will be made to whether additional nutrient augmentation is necessary. If it is necessary to augment with supplemental nutrients sodium hexametaphosphate and ammonium sulfate maybe added to the wells in order to augment the levels of these compounds.

4.1.4.1 Phosphate

Research of the current literature indicates that the direct addition of orthophosphate may cause the precipitation of insoluble phosphate salts, thus plugging the infiltration wells and the surrounding aquifer.

In sodium hexametaphosphate (SHMP), the phosphate is in the form of polyphosphate which and forms complex ions with the calcium and iron ions and does not precipitate out of solution. Additional sodium SHMP has a neutral pH of 7 and may be used in the treatment of potable water.

4.1.4.2 Ammonium Sulfate

In addition to the SHMP, ammonium sulfate as an ammonia source could be used. In addition to ammonia, this will also supply some sulfate, an important electron acceptor, to the system.

5.0 HYDROCARBON CONTAMINATION

The primary mass of hydrocarbon contamination remaining after the over-excavating and the installation of the receptor trench was found to be 2885 pounds in the soil with significant amounts (102 pounds) to be found in the groundwater, August 1999. The soil contamination is present in three phases; absorbed onto the soil, vapor and free phase. The free phase product has been found either coating the sand grains or as a floating product layer. Presently there is no significant floating product plume associated with this site.

The amount bound to the soil (2885 pounds) was found by contouring the results of the soil samples taken during test borings to find the resulting areas and volumes, see Table 5 and Figures 5 - 9 of Third Quarter 2000 report.

The mass in groundwater (102 pounds, August 1999) was found by contouring the August 24, 1999 ground water results and calculating the volume of contaminated water, as shown in Table 5 and Figures 10 and 11 of Third Quarter 2000 report. Current conditions (December 2001) indicate that this mass has been significantly reduced to 20.14 pounds TPHg and 1.77 pounds Benzene, see Figures 5 and 5-1 and Table 5.

Past experience has shown that significant levels of soil hydrocarbons can be removed through vapor extraction. Examinations of lithology beneath this site and along Brighton Avenue show that the formation is too clayey for this technology to have any degree of success.

Groundwater remediation will greatly benefit from elevating the dissolved oxygen levels in the groundwater within and near the plume. The least intrusive means to accomplish this would be to place socks containing ORC into selected wells (R3, RS5, RS7, T1, T2, T3 and T4) and monitor the levels of dissolved oxygen in the wells associated with the site periodically to evaluate if additional wells should be used for ORC placement.

6.0 INTERESTED PARTIES

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Mr. Leroy Griffin Oakland Fire Dept. OES Haz Mat Mgmt Program 1605 Martin Luther King Jr. Drive Oakland, CA 94612	Steve Marquez SWRCB, Cleanup Fund 2014 T Street Sacramento, CA 95814
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4.0 LIMITATIONS

This report is based upon the following:

- The observations of field personnel.
- The results of laboratory analyses performed by a state certified laboratory.
- Referenced documents.
- Our understanding of the regulations of the State of California and Alameda County, Hazardous Materials Section and/or City of Oakland, California.


Changes in groundwater conditions can occur due to variations in rainfall, temperature, local and regional water usage and local construction practices. In addition, variations in the soil and groundwater conditions could exist beyond the points explored in this investigation.

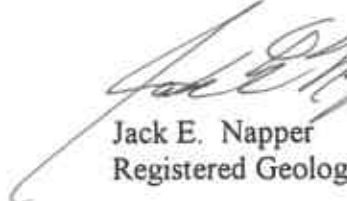

State certified analytical results are included in this report. This laboratory follows EPA and State of California approved procedures; however, WEGE is not responsible for errors in these laboratory results.

The services performed by Western Geo-Engineers, a corporation, under California Registered Geologist #3037 and/or Contractors License #513857, have been conducted in a manner consistent with the level of care and skill ordinarily exercised by members of our profession currently practicing under similar conditions in the State of California and the Oakland area. Our work and/or supervision of remediation and/or abatement operations, active or preliminary, at this site is in no way meant to imply that we are owners or operators of this site. Please note that known contamination of soil and/or groundwater must be reported to the appropriate agencies in a timely manner. No other warranty, expressed or implied, is made.

If you have any questions concerning this report or if we can be of further assistance, please don't hesitate to contact us at (530) 668-5300.

Respectfully,


 George Converse
 Project Geologist



 Jack E. Napper
 Registered Geologist #3037