

1. The TPH + TCOH were restricted in the surface - 1 to 3 inches

2. Barium + Lead.

3. PCB localized?  
A area - 1 mg/kg - highest surface  
C area - 10 mg/kg - highest surface  
code 5 foot x 10 foot having prog lower conc.  
at 10 foot - 1.18 mg/kg

ENVIRONMENTAL ASSESSMENT  
SOUTHERN PACIFIC TRANSPORTATION COMPANY  
744 HIGH STREET  
OAKLAND, CALIFORNIA

CEH

OOH - highest detected 28 mg/l  
PCB was found - 0.78 ug/l etc - 0.6 ug

September 5, 1989

Lead  
PCB  
OOH  
TPH

Prepared for:

SOUTHERN PACIFIC TRANSPORTATION COMPANY  
One Market Plaza  
San Francisco, California 94105



**ecology and environment, inc.**

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International Specialists in the Environment

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**TABLE OF CONTENTS**

<u>Section</u>	<u>Page</u>
1 INTRODUCTION .....	1-1
2 BACKGROUND INFORMATION .....	2-1
2.1 SITE HISTORY AND SETTING .....	2-1
2.2 ANALYSIS OF EXISTING ON-SITE SOIL DATA .....	2-4
3 ENVIRONMENTAL ASSESSMENT ACTIVITIES .....	3-1
3.1 SOIL SAMPLING .....	3-1
3.2 MONITORING WELL INSTALLATION .....	3-5
3.3 GROUNDWATER SAMPLING .....	3-10
3.4 HYDROGEOLOGIC ASSESSMENT .....	3-10
3.4.1 Stratigraphy .....	3-10
3.4.2 Hydrology .....	3-15
4 ANALYTICAL RESULTS .....	4-1
4.1 SOIL SAMPLE RESULTS .....	4-2
4.1.1 Inorganic Results .....	4-2
4.1.2 Organic Results .....	4-10
4.2 GROUNDWATER RESULTS .....	4-14
4.2.1 Inorganic Results .....	4-14
4.2.2 Organic Results .....	4-14
5 CONCLUSIONS AND RECOMMENDATIONS .....	5-1

Appendices

- A Soil Boring Lithologic Logs
- B ASC Laboratory Analytical Results

## LIST OF ILLUSTRATIONS

<u>Figure</u>		<u>Page</u>
2-1	Southern Pacific Transportation Company High Street Property Location Map .....	2-2
2-2	SPTCo. High Street Property, Site Layout .....	2-3
2-3	SPTCo. High Street Property, Approximate Sample Locations, July 1988 Sampling .....	2-5
3-1	SPTCo. High Street Property Soil Boring and Monitoring Well Locations .....	3-2
3-2	Typical Monitoring Well Construction Diagram .....	3-7
3-3	Hydrogeologic Cross Section Parallel to the East Property Boundary, SPTCo. High Street Property .....	3-13
3-4	Hydrogeologic Cross Section Parallel to the West Property Boundary, SPTCo. High Street Property .....	3-14

## LIST OF TABLES

<u>Table</u>		<u>Page</u>
2-1	Soil Sample Results - July 1988 Sampling .....	2-6
3-1	Soil Sample Compositing Scheme .....	3-3
3-2	Monitoring Well Construction Details .....	3-6
3-3	Well Development Water Quality Observations .....	3-9
3-4	Groundwater Level Elevations .....	3-11
4-1	Initial Soil Sample Results - Metals .....	4-3
4-2	Individual Soil Sample Results - Metals .....	4-4
4-3	Initial Soil Sample Results - Organics .....	4-11
4-4	Individual Soil Sample Results - Organics .....	4-12
4-5	SPTCo. High Street - Groundwater Results .....	4-15

## 1. INTRODUCTION

This environmental assessment report presents the results of Ecology and Environment, Inc.'s (E & E's) soil and groundwater investigation at the Southern Pacific Transportation Company's (SPTCo.'s) property located at 744 High Street in Oakland, California. This investigation was conducted in response to a request by the Alameda County Health Care Services Agency (ACHCSA) to characterize potential soil and groundwater contamination at the property. The ACHCSA request was prompted by the detection of total oil and grease (TOG), polychlorinated biphenyls (PCBs), lead, benzene, and a variety of volatile organic chemicals (VOCs) at levels that may require remediation in surface soils at the property. These constituents were detected during surface soil sampling that was performed in July 1988 by the owner of the former Scrap Metal Supply business that leased a portion of the property from SPTCo.

E & E's investigative activities consisted of reviewing the existing soils data for the property, identifying potential off-site sources of contamination, reconstructing the history of the site, drilling and sampling 12 boreholes, installing and sampling three groundwater monitoring wells, and performing a local groundwater level elevation survey.

The site history and the results of the July 1988 sampling activities are summarized in Section 2; the environmental assessment activities performed by E & E are described in Section 3; the analytical results are presented and discussed in Section 4; and the conclusions and recommendations are presented in Section 5.

*12 Bore Holes*

*3 Groundwater monitoring wells*

## 2. BACKGROUND INFORMATION

In this section the site history, land use, regional environmental setting, sources of potential contamination, and results of previous on-site soil sampling are discussed.

### 2.1 SITE HISTORY AND SETTING

The SPTCo. property located at 744 High Street in Oakland, California, is situated in a light to heavy industrial area approximately 0.4 miles from the Alameda Estuary and San Leandro Bay (see Figure 2-1). The current property layout is shown in Figure 2-2. SPTCo. acquired the property in 1872 from the Contract and Finance Company. Use of the property, if any, prior to 1872 is unknown.

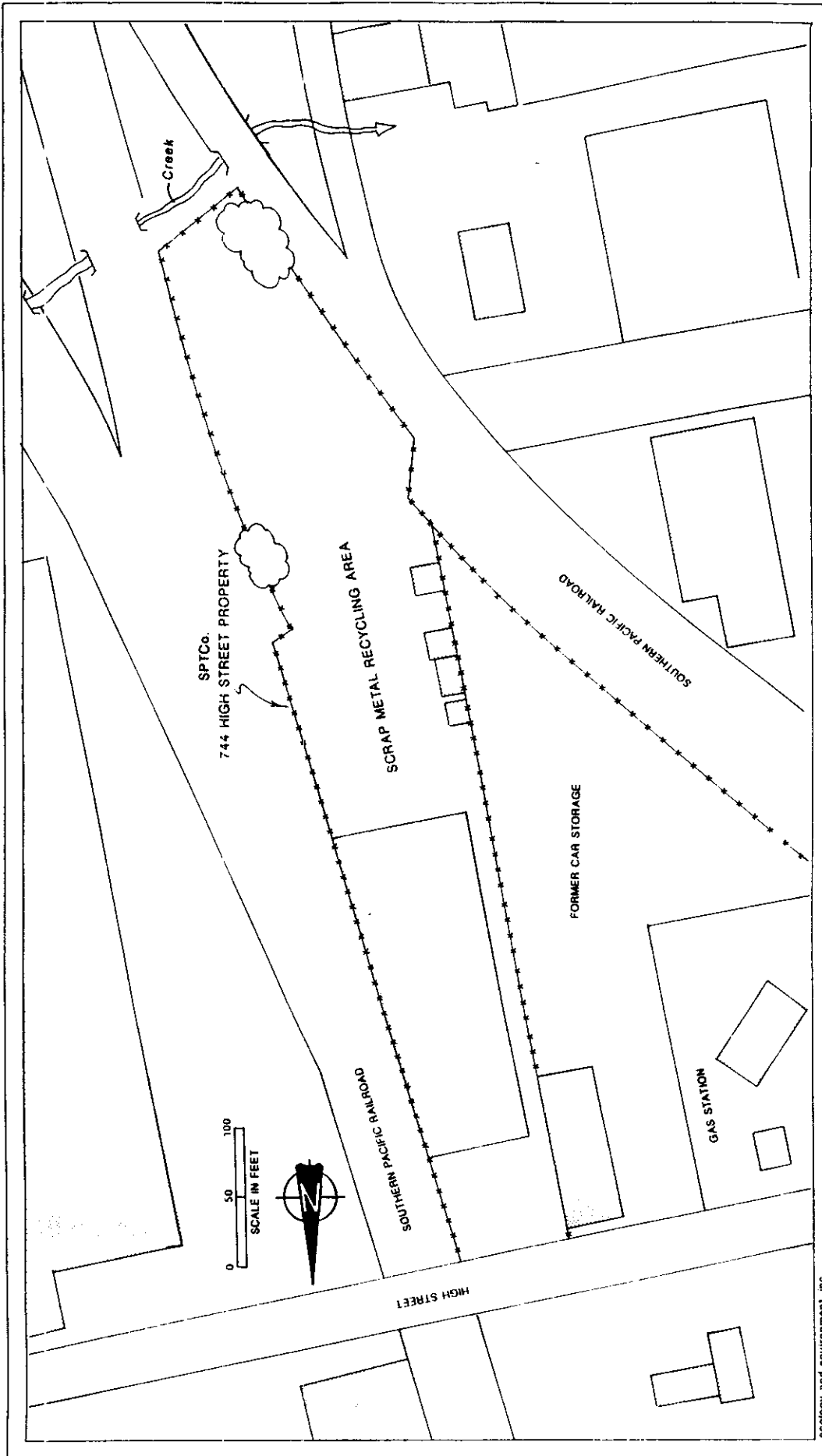
The property is currently occupied by two businesses, Scrap Metal Supply (SMS) and Kayak America Works (KAW). SMS is a metal recycler and occupies approximately 56,000 square feet in the southern part of the property. KAW is an auto body repair shop and occupies approximately 26,000 square feet in the northern part of the property. SMS began leasing the southern portion of the property from SPTCo. in 1956, and SPTCo. began leasing out the northern 26,000 square feet in 1966. Initially, the northern portion was used for a sheet metal plant.

The area in the vicinity of the property is underlain by alluvial deposits that consist predominantly of clays and silts, although layers of generally clayey sands and gravels occur throughout the subsurface. A near-surface clayey gravel layer approximately 5 feet thick was encountered at a depth of about 7 feet below ground surface beneath the property. The next generally coarse-grained layer reportedly occurs at a depth of about 40 feet below ground surface and it is separated from the near-surface coarse-grained layer by silty clay and clayey silt.

Saturated strata was encountered at a depth of approximately 10 feet below ground surface beneath the property. The groundwater flow

*Groundwater flow*  
*10 ft*





ecology and environment, inc.

Figure 2-2 SPTCo. HIGH STREET PROPERTY  
SITE LAYOUT

direction would be expected to be toward the southwest toward the Alameda Estuary and San Leandro Bay. The generally clayey nature of the coarser-grained layers preclude them from being reliable sources of water in the area. No drinking water supply wells have been identified within one mile of the property.

## 2.2 ANALYSIS OF EXISTING ON-SITE SOIL DATA

In July of 1988, at the request of the owner of the Scrap Metal Supply business, Property Commission Control (PCC) conducted soil sampling at the SPTCo. High Street property. A total of seven discrete surface soil samples were collected at the approximate locations shown on Figure 2-3. Samples from four locations were combined to yield one composite analysis (denoted as No. 4 on Figure 2-3). Samples from locations 1, 2, and 3 were analyzed individually. The resulting four samples were analyzed for TOG, PCBs, a variety of metals, and volatile organics. Analytical results are presented in Table 2-1 (only those metals that were detected in a sample are presented in the table).

The sample results reveal that surface soils at the Scrap Metal Supply business contain TOG, PCBs, and a variety of VOCs (primarily toluene, xylene, and other aromatic petroleum compounds). TOG occurred at levels above that which the ACHCSA considers a threat to groundwater quality or public health and safety (1,000 mg/kg) in three of the four analyses. PCBs occurred above the total threshold limit concentration (TTLIC) of 50 mg/kg in one of the four analyses.

The detection of contaminants in surface soils at the property resulted in the ACHCSA requesting SPTCo. to develop and implement a Plan of Correction for the characterization and/or remediation of soil and/or groundwater contamination on the property.

*Groundwater Treatment  
DET. R. M. ...*



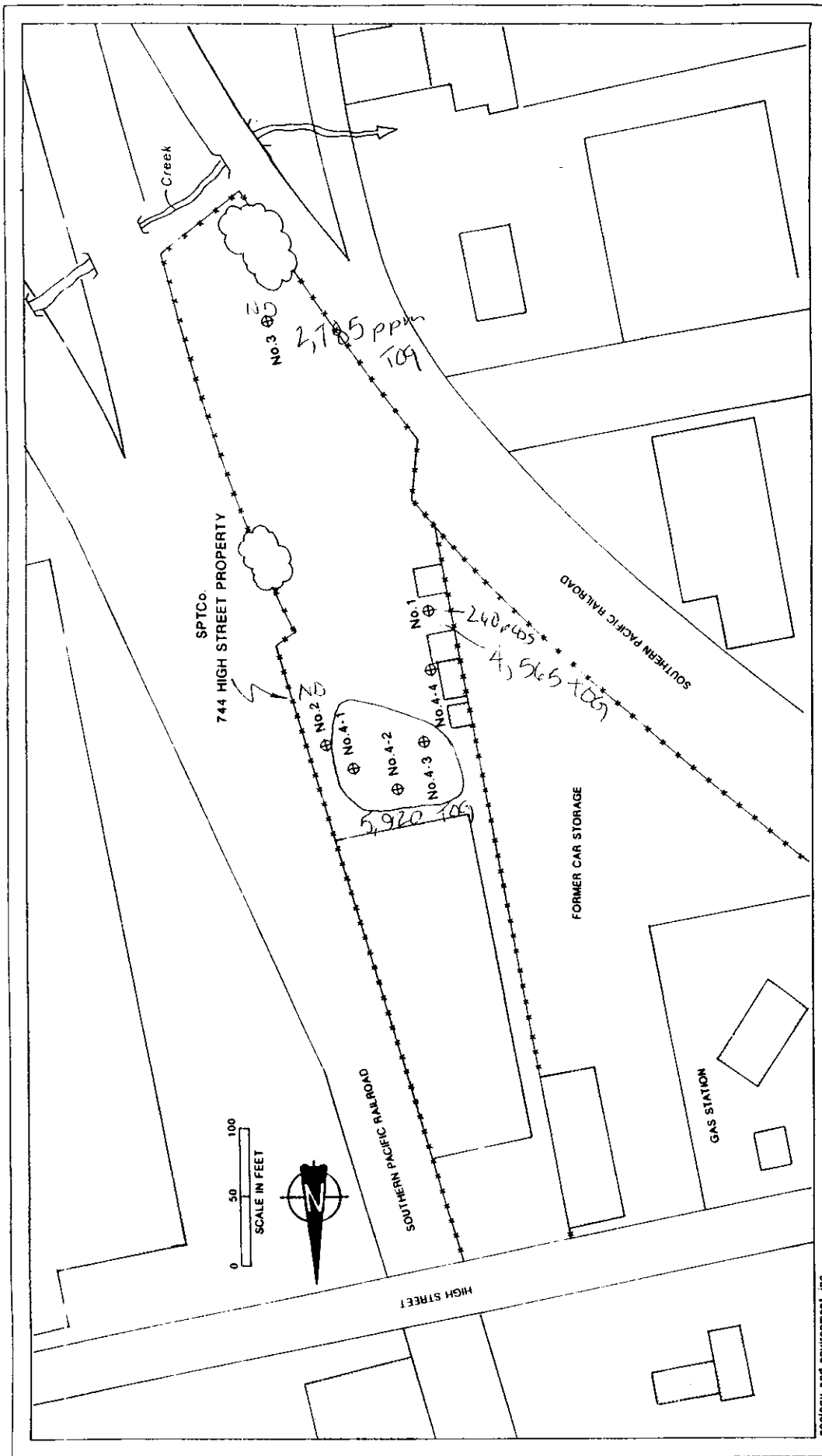


Figure 2-3 SPTCo. HIGH STREET PROPERTY  
 APPROXIMATE SAMPLE LOCATIONS  
 JULY 1988 SAMPLING

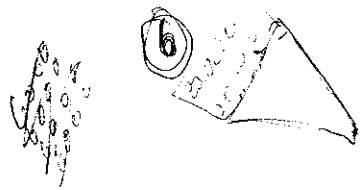
Table 2-1

SOIL SAMPLE RESULTS  
July 1988 Sampling (mg/kg)

Sample Number Type	#1 Discrete	#2 Discrete	#3 Discrete	#4 Composite of Four Locations
Depth	8"	12"	12"	0 to 6"
Total Oil and Grease	4,565	58	2,785	5,920
PCBs	260	ND	ND	836
Barium	0.498	<0.33	<0.33	0.942
Cadmium	0.031	ND	ND	0.143
Copper	0.106	ND	ND	0.525
Iron	0.487	0.253	0.333	0.370
Manganese	0.561	0.082	0.489	6.98
Nickel	<0.17	ND	ND	0.218
Lead	<0.17	ND	ND	0.339
Zinc	3.61	0.553	0.610	27.1
Trichlorofluoromethane	ND	ND	1.0	ND
Carbon Disulfide	7.6	1.2	3.8	ND
Methylene Chloride	8.5	2.1	3.2	ND
Chloroform	1.5	0.9	0.7	ND
Toluene	1.3	ND	ND	ND
Xylene	1.3	ND	ND	ND
1-chloro-2- methylbenzene	<6.25	<6.25	<6.25	ND
1,2,3-Trimethyl- benzene	<6.25	ND	ND	ND
1,2,3-Trimethyl- benzene	<6.25	ND	ND	ND
1,2,4-Trimethyl- benzene	<6.25	ND	ND	ND
1-methyl-3- (methylethyl) benzene	ND	ND	ND	ND

ND = Not Detected.

m/spea/t2-1



### 3. ENVIRONMENTAL ASSESSMENT ACTIVITIES

E & E's environmental assessment activities were designed to evaluate the subsurface extent of potential soil contamination at the site and to determine whether groundwater is contaminated. The environmental assessment activities were implemented in response to a letter request by the ACHCSA, dated March 2, 1989, for a Plan of Correction for the property. The Plan of Correction was submitted to the ACHCSA on April 26, 1989 and was subsequently verbally approved. The scope of work proposed in the Plan of Correction included drilling and sampling 12 soil boreholes, installing and sampling three groundwater monitoring wells, and performing a local groundwater level elevation survey.

*Fig 3.1*

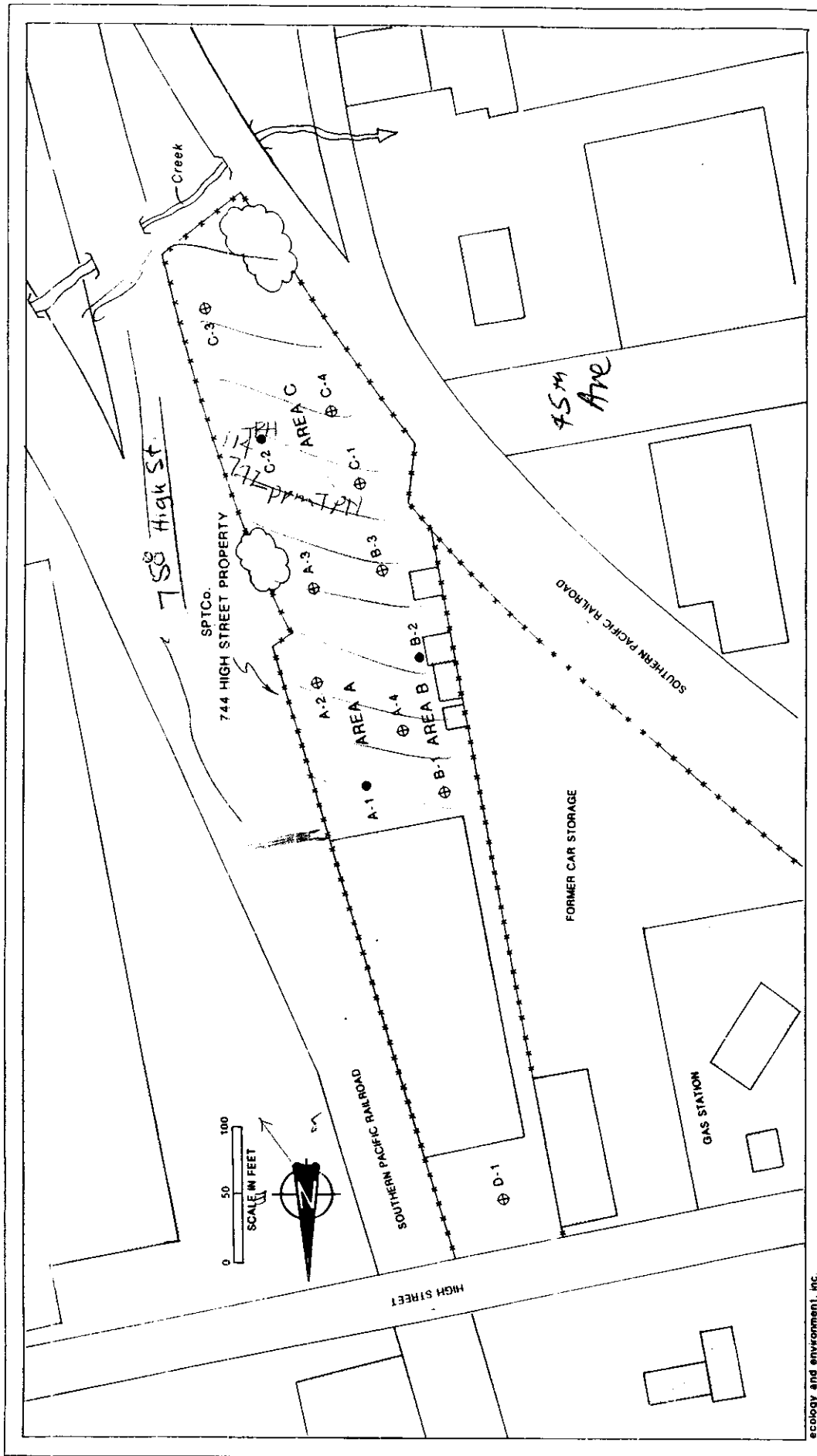
The field investigations proposed in the Plan of Correction were carried out between May 22, 1989, and May 26, 1989. Additional sampling, to augment the results of the initial characterization activities, was conducted on July 28, 1989. This sampling involved collecting and analyzing samples from trenches excavated adjacent to several previously drilled soil borings. A total of eight soil samples (including one duplicate) were collected on July 28, 1989.

#### 3.1 SOIL SAMPLING

A total of 12 soil borings were drilled and sampled on the property at locations shown on Figure 3-1. Soil borings were drilled with a Mobile Drill B-53 rig using 6 7/8-inch-outside-diameter hollow-stem auger flights. Samples were collected from each borehole at 5-foot intervals with a 2-inch diameter, 1.5-foot-long, split-spoon sampler. The sampler was equipped with three, 6-inch-long brass sleeves. Soil boring details and samples collected are presented in Table 3-1. Lithologic logs of soil borings are presented in Appendix A.

For analysis, individual soil samples were composited according to a geographic compositing scheme (see Table 3-1). The soil borings were

*compositing laterally*



⊕ SOIL BORING AND MONITORING WELL  
 ● SOIL BORING AND MONITORING WELL  
 12 Boreholes  
 3 DEVELOPED TO WELLS - soil samples/yr

Figure 3-1 SPTCo. HIGH STREET PROPERTY  
 SOIL BORING AND MONITORING WELL LOCATIONS

Table 3-1

## SOIL SAMPLE COMPOSITING SCHEME

Area	Composite Number	Discrete Samples	Sample Depth (feet)	
A	1	A-1a <sup>1</sup>	3.0 - 3.5	
		A-2a	1.0 - 1.5	
		A-3a	1.5 - 2.0	
		A-4a	1.5 - 2.0	
	2	A-1b	5.5 - 6.0	
		A-2b	5.5 - 6.0	
		A-3b	5.5 - 6.0	
		A-4b	5.5 - 6.0	
	3	A-1c	10.5 - 11.0	
		A-2c	10.5 - 11.0	
		A-3c	10.5 - 11.0	
		A-4c	10.5 - 11.0	
B	4	B-1a	1.0 - 1.5	
		B-2a	1.5 - 2.0	
		B-3a	2.5 - 3.0	
	5	B-1b	6.0 - 6.5	
		B-2b	5.5 - 6.0	
		B-3b	5.5 - 6.0	
	6	B-1c	11.0 - 11.5	
		B-2c	10.5 - 11.0	
		B-3c	10.5 - 11.0	
	C	7	C-1a	3.5 - 4.0
			C-2a	2.0 - 2.5
			C-3a	2.5 - 3.0
C-4a			3.0 - 3.5	
8		C-1b	5.5 - 6.0	
		C-2b	6.0 - 6.5	
		C-3b	6.0 - 6.5	
		C-4b	5.5 - 6.0	
9		C-1c	10.5 - 11.0	
		C-2c	11.0 - 11.5	
		C-3c	10.5 - 11.0	
		C-4c	10.5 - 11.0	
D <sup>4</sup>		D1a	1.0 - 1.5	
		D1b, D1c	5.5 - 6.0	
		D-1d	5.5 - 6.0	

1. "a" designation indicates samples from 0.5 - 1.0 feet.
2. "b" designation indicates samples from 5.5 - 6.0 feet.
3. "c" designation indicates samples from 10.5 - 11.0 feet.
4. Samples from this boring were analyzed individually.

divided into three groups based on geographic proximity. Area A included four soil borings (A-1, A-2, A-3, and A-4); area B included three soil borings (B-1, B-2 and B-3); area C included four soil borings (C-1, C-2, C-3, and C-4); and area D included one soil boring (D-1). Soil samples from similar depths from each boring in a specific area were composited in the laboratory for analysis. For example, in area C soil samples from a depth of 5 feet from borings C-1, C-2, C-3, and C-4 were composited to yield one analysis. The soil samples from boring D-1 were analyzed individually. This compositing scheme yielded a total of 13 analyses, including the analysis of one duplicate sample from a depth of 5 feet from boring D-1.

Three soil borings (A-1, B-2, and C-2) were completed as groundwater monitoring wells. Boreholes at these locations were drilled to a depth of 25 feet and samples were collected at 5-foot intervals to that depth. Boreholes that were not completed as monitoring wells were drilled and sampled to a depth of 10 feet, which was estimated to be the approximate depth of the water table. Soil samples below a depth of 10 feet from borings A-1, B-2, and C-2 were stored in the laboratory as discrete samples.

Following receipt of laboratory analytical results on July 14, 1989, for the soil and groundwater sampling that occurred in late May 1989, it was determined that several soil samples making up a composite should be analyzed individually for select parameters, since the composite results indicated that an individual result for a particular parameter could possibly exceed either the total threshold limit concentrations (TTL) (California Code Regulations (CCR), Title 22, Chapter 30), 10 times the soluble threshold limit concentration (STLC CCR, Title 22, Chapter 30), or the state action level for total petroleum hydrocarbons (TPH). Unfortunately, all portions of individual samples making up a composite were used during compositing. It was therefore necessary to collect samples from as close to selected borings and depths as was possible.

On July 28, 1989, a total of eight soil samples were collected (including one duplicate) and analyzed as individual samples. The

samples were collected from the wall of a trench that was excavated with a backhole as close to the original soil boring as was possible. The samples collected were the shallowest samples from the soil borings in areas B and C.

### 3.2 MONITORING WELL INSTALLATION

Soil borings A-1, B-2, and C-3 (see Figure 3-1 for locations) were drilled to a depth of 25 feet and were completed as groundwater monitoring wells. Wells were completed with 2-inch inside diameter Schedule 40 PVC. The screen consisted of 0.020 inch factory manufactured slots with screen lengths of 15 feet. Monitoring well details are presented in Table 3-2 and a typical construction diagram is presented in Figure 3-2. The wells were completed to a depth of approximately 23 feet with the screen extending to about 8 feet. The intent was to screen across the water table to evaluate whether or not floating product was present. Depths to water appeared to range from 8 to 11 feet below ground surface at the time of drilling. Water levels in wells B-2 and C-2 after installation, however, were measured to be shallower than 8 feet, suggesting that the water-bearing zone screened may be partially confined. There was no evidence of floating product flowing into any of the wells during well development when the water level was lowered below the top of the screen.

In each monitoring well, a coarse-grained sand filter pack was installed so as to extend at least 1.5 feet above the top of the screen, and a bentonite pellet seal at least 1.5 feet thick was placed above the filter pack. The remainder of the annular space was filled with a cement grout. Each monitoring well was completed below grade and was enclosed in a christy box completed flush to the ground surface.

Following installation, each monitoring well was developed using a Teflon bailer. Development continued until either the water was clear and the water quality parameters temperature, specific conductance, and pH had stabilized, or the well repeatedly bailed dry. Parameters

Table 3-2

MONITORING WELL CONSTRUCTION DETAILS

Well	Borehole Depth <sup>1</sup>	Bottom of Casing <sup>1</sup>	Screen Interval <sup>1</sup>	Top of Sand <sup>1</sup> Pack <sup>1</sup>	Top of Bentonite Seal <sup>1</sup>
A-1	25.0	22.5	7.5 - 22.5	6.3	5.0
B-2	25.0	23.0	8.0 - 23.0	6.5	5.0
C-2	23.0	23.0	8.0 - 23.0	6.0	4.5

1. Feet below ground surface.



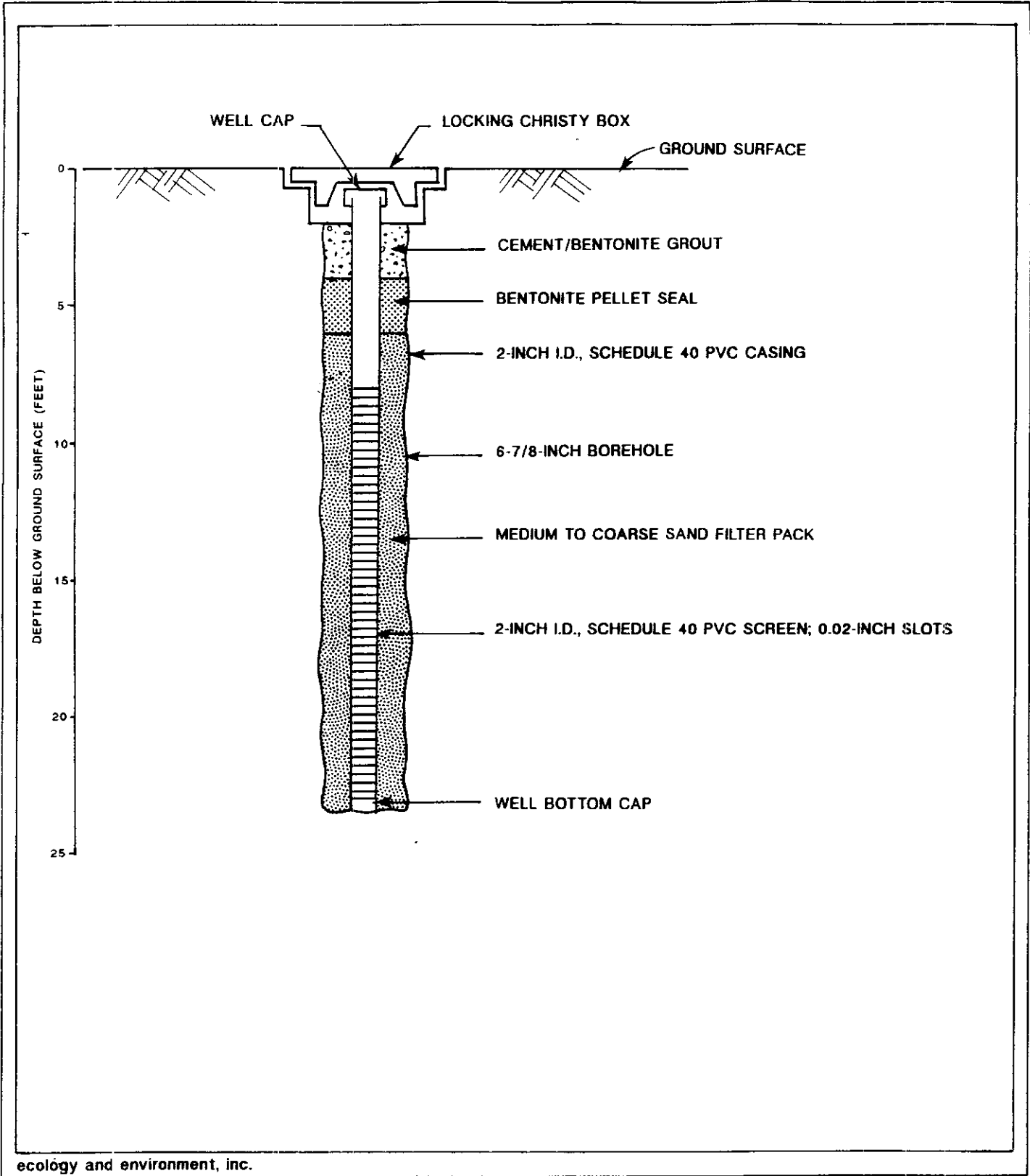


Figure 3-2 TYPICAL MONITORING WELL CONSTRUCTION DIAGRAM

measured during well development are presented in Table 3-3.

During development of monitoring well A-1, 62.5 gallons of groundwater were removed. The well bailed dry twice during development. The well initially bailed dry after 27.5 gallons of water were removed. It was allowed to recover for two hours, after which it bailed dry again after an additional 22.5 gallons of water had been removed. The temperature of the groundwater stabilized at approximately 64<sup>o</sup>F, the specific conductance stabilized at approximately 1,000 umhos/cm, and the pH was 6.

A total of 24.5 gallons of groundwater were removed from monitoring well B-2 during development. The well bailed dry twice during development, initially after 7.5 gallons of water had been removed and again after a cumulative total of 24.5 gallons had been removed. The well was allowed to recover for approximately 5 hours after it initially was bailed dry. After it bailed dry the second time, development was stopped even though the groundwater was still turbid. The temperature of the groundwater in B-2 averaged 62<sup>o</sup>F, the specific conductance was between 800 and 1,100 umhos/cm, and the pH was 6. The specific conductance appeared to increase with increasing turbidity.

A total of 9.5 gallons of water were removed from monitoring well C-2 during development. The well initially bailed dry after 7.5 gallons of water had been removed. It was allowed to recover for 3 hours; however, over this time period it was still not fully recovered and bailed dry again after only 2 more gallons of groundwater had been removed. At this time development was stopped. The temperature of the groundwater was about 63<sup>o</sup>F, the specific conductance was between about 1,300 and 1,500 umhos/cm, and the pH was 6. Continued development was not warranted due to the extremely low yield and the slow recovery rate.

To determine the groundwater flow direction and gradient at the SPTCo. High Street property, the groundwater level in each well was measured on May 26, 1989, and July 28, 1989. The elevation of each well, referenced to mean sea level, was determined to an accuracy of 0.01 feet by a

Table 3-3

## WELL DEVELOPMENT WATER QUALITY OBSERVATIONS

Well	Date	Time	Gallons Bailed	pH	EC	Temp.	Observations
A-1	5/25/89	0830	0	6	1416	67.4	
		0911	10	6	1333	65.6	water is turbid, brown
		0924	15	6	1279	68.3	water is turbid, brown
		0937	20	6	1286	65.9	was is turbid
		0949	25	6	1200	66.1	water is turbid
		0959	27.5	-	-	-	water is turbid
		1015					start bailing again
		1020	30.5	6	1099	70.5	stop bailing
		1240					resume bailing
		1245	33	6	1067	70	water is turbid
		1301	-	6	1062	77.2	
		1304	-	6	975	67.3	
		1305	41				
		1311	46	6	980	69	
		1328	50	6	1067	71.5	water is less turbid
		1541					resume bailing
		1543		6	1018	65	
		1549		6	1002	63.8	
		1554		-	935	63	
		1558		-	952	63.3	moderately turbid
1605	62.5	-	1059	64.7	water is becoming clearer		
B-2	5/25/89	1105	0	6	719	68.9	
		1107	5	6	762	68.4	
		1110	-	6	1066	71	
		1118	7.5	6	1066	71	
		1124	-	6	1222	71	well bailed dry
		1449	-	6	750	65.9	
		1455	-	6	901	64.2	
		1458	-	6	865	62	
		1459	-	6	1074	62	
		1501	-	6	1163	62	
		1508	24.5	6	1171	61.2	water is very turbid, well bailed dry
		C-2	5/25/89	1230	5	6	1564
1232	7.5						well bailed dry
1515				6	1460	63.5	water is clear
1526				6	1340	62.5	
1529	9.5			-	-	-	well bailed dry

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surveyor licensed in California. Groundwater level elevations are summarized in Table 3-4. Measurements were made within about 1 hour of each other so that variations due to tidal fluctuations could be kept to a minimum. The results of the water level measurements are discussed in Section 3.4.2.

### 3.3 GROUNDWATER SAMPLING

Groundwater samples from each well were obtained on May 26, 1989. Samples were analyzed for TOC, PCBs, purgeable organics, priority pollutant metals, and TPH by E & E's Analytical Services Center (ASC). Sampling and shipment procedures followed EPA-approved protocol. In addition to samples from the three monitoring wells, a duplicate sample was collected from monitoring well A-1 and a blank was prepared. The duplicate and blank were analyzed for the same parameters as the environmental samples. Analytical results are discussed in Section 4.

Prior to evacuating the appropriate volumes of water from each well, the static water level was measured in all of the wells. Water levels were measured between 0836 and 0845 on May 26, 1989. Each monitoring well bailed dry during evacuation. Monitoring well A-1 bailed dry after 18 gallons had been evacuated. Monitoring well B-2 bailed dry after 22.5 gallons had been evacuated and monitoring well C-2 bailed dry after approximately 5 gallons had been evacuated. Each well was allowed to recover for approximately 2 hours prior to obtaining a sample.

### 3.4 HYDROGEOLOGIC ASSESSMENT

#### 3.4.1 Stratigraphy

The stratigraphy to a depth of 25 feet beneath the SPTCo. High Street facility is generally uniform with the exception of the extreme southern portion of the property. The stratigraphic sequence encountered, from the surface down, typically consists of 1 to 3 feet of fill, 6 to 10 feet of fine-grained sediment ranging from clay to clayey silt, 3 to 10 feet of coarse-grained sediment consisting of clayey gravel or clayey

Table 3-4

## GROUNDWATER LEVEL ELEVATIONS

Well	Ground Surface Elevation <sup>1</sup>	May 26, 1989		July 28, 1989	
		Depth to Water (feet)	Groundwater Elevation <sup>1</sup>	Depth to Water (feet)	Groundwater Elevation <sup>1</sup>
A-1	15.57	7.10	8.47	8.30	7.27
B-2	14.37	6.37	8.00	8.01	6.36
C-2	15.30	11.24	4.06	14.72	0.58

1. Elevations are referenced to mean sea level.

coarse sandy gravel, and a lower fine-grained layer consisting primarily of clayey silt. Hydrogeologic cross sections are presented in Figures 3-3 and 3-4.

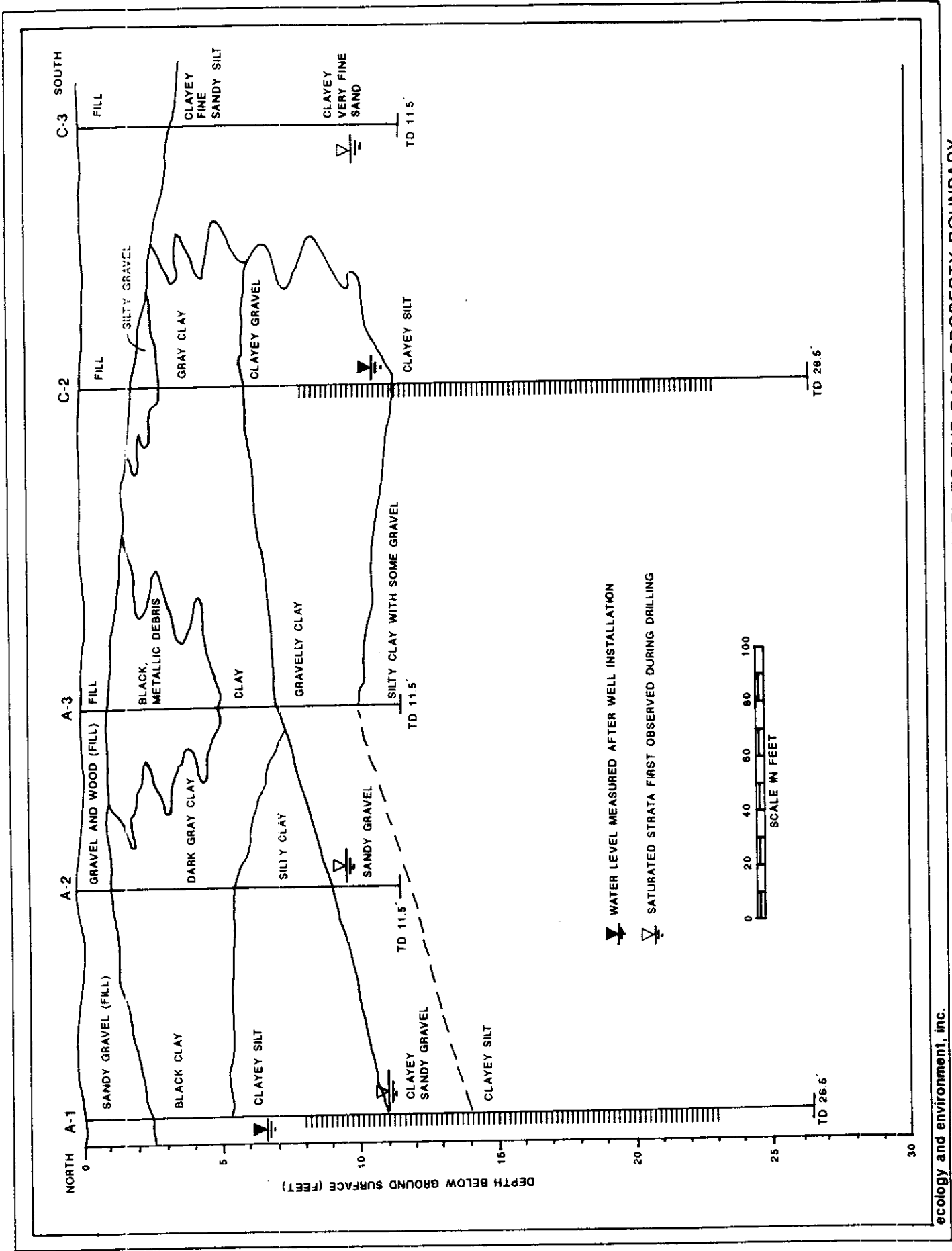
The surface fill generally is thickest in the southern portion of the site. The fill consists of a rubbly mixture of soil, wood debris, metal fragments, and broken pieces of plastic. The fill is underlain by black to dark gray to greenish gray clay that typically is stiff and locally was observed to be mottled, crumbly, and slightly silty.

In the northern portion of the site, the black to gray clay is underlain by tan to orange brown clayey silt to silty clay. To the south this brownish clayey silt was not encountered during drilling, and gray clay typically is underlain directly by clayey gravel or gravelly clay. Both the black to gray clay and the brownish clayey silt contains small rust-colored rootlets.

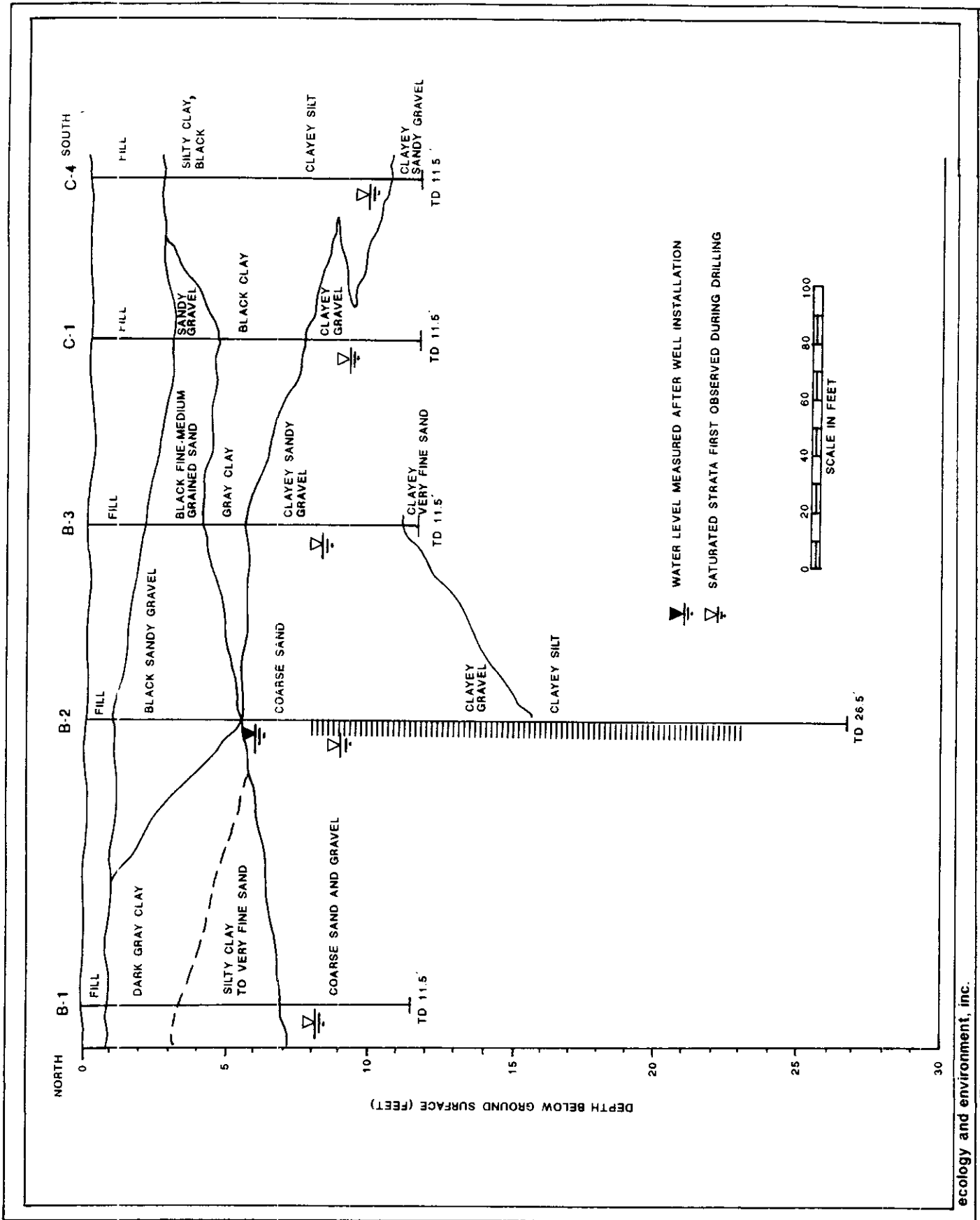
Throughout the site, with the exception of the extreme southern part, the near-surface predominantly clay deposits are underlain by poorly sorted, clayey coarse-grained sediments. These deposits range in thickness from 10 to 15 feet and typically consist of tan to orange-brown clayey gravel, clayey coarse sandy gravel, or clayey coarse sand. These coarse deposits are underlain by gray to gray and brown mottled stiff clayey silt with occasional thin layers of silty very fine sand. Small rust-colored rootlets occur throughout the clayey silt.

The sedimentary sequence penetrated in boring C-3, located in the extreme southern portion of the property, was distinct from that observed in other areas on the property. At this borehole, approximately 3 feet of dark brown, sandy gravelly fill was underlain by about 4 feet of orange gray-brown, clayey fine sandy silt with some gravel and very fine orange-tan clayey sand that extended to the bottom of the borehole.

In boreholes C-1 and C-2, thin layers (1 to 2 feet thick) of orange-tan



ecology and environment, inc. **Figure 3-3 HYDROGEOLOGIC CROSS-SECTION PARALLEL TO THE EAST PROPERTY BOUNDARY SPTCo. HIGH STREET PROPERTY**



ecology and environment, inc.

Figure 3-4 HYDROGEOLOGIC CROSS-SECTION PARALLEL TO THE WEST PROPERTY BOUNDARY  
SPTCo. HIGH STREET BOUNDARY



sandy to clayey silty gravel were encountered beneath the fill and above the black to gray clay. These thin gravelly deposits do not appear to be widespread and are probably fill.

In boreholes A-3, B-2, and B-3, the surface fill was underlain by black metallic slag. In A-3 the slag consisted of fine to coarse-grained metallic debris. In B-2 it consisted of black sandy gravel with some wood that graded down into black metallic slag. In B-3 it consisted of black fine- to medium-grained slag. These areas represent debris resulting from past scrap metal recycling activities.

At a site located approximately 600 feet to the north of the SPTCo. (C.C. Corp.) property, two soil borings have been drilled to a depth of 240 feet. The coarse-grained layer consisting of clayey gravel to clayey coarse sandy gravel that occurs at a depth of approximately 10 feet beneath the SPTCo. property was also encountered in these nearby borings. Clayey materials predominated in the nearby 240 foot borings; however, coarser grained clayey to clean sands and gravels occurred in the approximate depth ranges of 8 to 18 feet, 40 to 80 feet, 120 to 140 feet, and 200 to 230 feet below ground surface.

At the SPTCo. High Street property, the three borings drilled to a depth of 25 feet penetrated into as much as 15 feet of clayey silt and silty clay beneath the clayey sand and gravel layer. None of the three borings penetrated through this lower fine-grained zone. The deeper borings drilled nearby indicate that another coarser-grained zone occurs at a depth of approximately 40 feet and that the clayey sand and gravel layer encountered at the SPTCo. property is separated from deeper permeable zones by as much as 25 feet of low permeability clayey silts and silty clays.

#### 3.4.2 Hydrology

Saturated strata was encountered at a depth of approximately 10 feet below ground surface throughout most of the site during drilling. Saturated strata was not penetrated in boring D-1 at the north end of

the property so the depth to water in that area is greater than 11.5 feet (the depth of the bottom of the drive sampler when the 10 foot drive sample was collected). In boring C-2, the drill cuttings and drive sampler, to a depth of 26.5 feet did not appear wet, but only slightly moist, so it was difficult to determine when first water was encountered. This lack of water is reflected in the extremely low yield of monitoring well C-2, which bailed dry during development after only 7.5 gallons of water were removed.

The groundwater levels in monitoring wells A-1 and B-2 after installation and development were measured to be above the top of the screens, suggesting that they tap groundwater that is under pressure and that may be partially confined. In monitoring well C-2, the groundwater level was measured to be below the top of the screen and within the permeable zone (clayey gravel) tapped by the well (see Figures 3-3 and 3-4) indicating that the clayey gravel layer is not fully saturated and the groundwater in the well is unconfined.

The groundwater level elevations in the three monitoring wells suggest that there may be a southeasterly flow gradient beneath the property, although the water levels in A-1 and B-2 may not be comparable to the water level in C-2, because monitoring wells A-1 and B-2 appear to be confined and monitoring well C-2 appears to be unconfined.

The horizontal gradient between wells A-1 and B-2 is not less than 0.005 feet per foot. If the water levels in A-1, B-2, and C-2 are comparable, then the horizontal gradient between the northern and southern portions of the property is approximately 0.026 feet per foot in a southeasterly direction.

Between May 26, 1989, and July 28, 1989 the water levels in each well were observed to drop (see Table 3-4). The drops ranged from 1.20 feet in monitoring well A-1 to 3.48 feet in monitoring well C-2. These differences are too great to be attributable to variations caused by tidal fluctuations and probably reflect seasonal changes.

The extremely low yields in the three groundwater monitoring wells at the SPTCo. property indicate that the clayey sand and gravel layer beneath the property has a low hydraulic conductivity. The low hydraulic conductivity and the thin saturated thickness of this layer preclude it from being developed as a usable source of water for any uses in the area.

The shallow water-bearing zone in the vicinity of the SPTCo. property is separated from deeper water-bearing zones by as much as 25 feet of low permeability clayey silts and silty clays. Groundwater monitoring well C-2 indicates that this low permeability zone is virtually non-water bearing. This well was screened primarily in clayey silt. During development and sampling this well rapidly bailed dry and was very slow to recover. The extremely low permeability of this fine-grained zone would tend to prevent the downward migration of any contaminants from the shallow groundwater to deeper water-bearing zones.

Deeper water-bearing zones in the vicinity of the SPTCo. property tend to contain substantial clay components and, as a result, the hydraulic conductivities and yields are low to moderate. There are no high yield, usable aquifers beneath the site to a depth of at least 240 feet.

No wells used for drinking water supply have been identified within a radius of 1 mile of the SPTCo. property. An industrial well has been located approximately 1,500 feet to the southeast of the property. The current status of this well is unknown. It was drilled in 1962 to a depth of 776 feet and reportedly was screened over virtually its entire length. The location of this well is lateral to the groundwater flow direction beneath the SPTCo. property and as a result it would not be a potential receptor of contamination migrating downgradient beneath the SPTCo. property.

#### 4. ANALYTICAL RESULTS

The environmental assessment of the SPTCo. High Street property included both soil and groundwater sampling and analysis. Analytical reports and the quality assurance/quality control (QA/QC) evaluation are presented in Appendix B. Soil sampling and analysis was performed in two stages. Initially, individual samples from similar depths and areas were composited in the laboratory for analysis. The compositing scheme was described in detail in Section 3-1. The compositing resulted in 13 soil analyses (including one duplicate analysis) and included three composite analyses from four borings in area A, three composite analyses from three borings in area B, three composite analysis from four borings in area C, and four individual analyses from boring D. Soil samples were analyzed for TPH (EPA Method 418.1, modified for soils), TOC (EPA Method 413.1, modified for metals), priority pollutant metals (EPA Method 6010), PCBs (EPA Method 8080), and purgeable organics (EPA Method 8240).

The second stage of soil sampling and analysis involved collecting individual samples from depths near borings in areas where the composite results indicated that soluble concentrations of particular metals may potentially be present. This sampling included collecting near-surface samples near the borings in areas B and C. Analyses were dependent on the composite sample results. Individual samples from area B were analyzed for zinc, lead, nickel, copper, and barium; individual samples from area C were analyzed for cadmium, lead, nickel, barium, total petroleum hydrocarbons, and PCBs. Based on the individual sample results, California waste extraction tests (WET), (CCR, Title 22, Chapter 30) were performed on samples B-2A and C-3A. B-2A is being analyzed for zinc and lead and C-3A is being analyzed for lead.

Groundwater samples were collected from three monitoring wells that were installed on the property during this investigation. Groundwater samples were analyzed for TPH (EPA Method 418.1), TOG (EPA Method 413.1), priority pollutant metals (EPA Method 200.7), purgeable organics (EPA Method 624), and PCBs (EPA Method 625).

## 4.1 SOIL SAMPLE RESULTS

### 4.1.1 Inorganic Results

Inorganic analyses consisted of analysis for priority pollutant metals. Results of the initial sampling are presented in Table 4-1; results of the second sampling are presented in Table 4-2.

To determine the environmental significance of the metal results, concentrations have been compared to TTLC and STLC concentrations and also to average concentrations for the western United States presented in "Element Concentrations in Soils and Other Surficial Materials of the Conterminous United States," 1984, U.S. Geological Survey Professional Paper 1270. Concentrations exceeding the TTLC would classify the soil as a hazardous waste for disposal purposes. To determine whether a sample exceeds the STLC criteria for a particular metal, a WET test must be performed. When comparing total concentrations of a particular metal to the STLC criteria, the 10-fold dilution involved in the WET test must be taken into consideration. For example, during the WET test, a 50-gram soil sample is extracted in a total volume of 500 ml of solution, so the original concentration is diluted 10 times, and then the extract is analyzed. The resulting extracted concentration, expressed in mg/l, is then compared to the STLC criteria. If the total concentration of a particular metal is 50 mg/kg, the 10-fold dilution would result in a maximum possible concentration in the extract solution of 5 mg/l (assuming 100% leachability). If the STLC criteria for that particular metal is 5 mg/l, then a total concentration equal to or greater than 50 mg/kg would be required for the STLC to be exceeded during the WET test. Therefore, as a screening tool to determine which total concentrations could potentially yield WET test results that equal or exceed the STLC, a total concentration in mg/kg that is at least 10 times the STLC criteria in mg/l must be present.

#### Arsenic

The concentrations of arsenic appear to be relatively uniform throughout



Table 4-2

## INDIVIDUAL SOIL SAMPLE RESULTS - METALS

SOIL Pb LEVELS = 6.7 PPM.

Sample	Depth (feet)	Barium	Cadmium	Copper	Lead	Nickel	Zinc
B-1A	2	453	NA <sup>1</sup>	193	356	121	1,990
B-2A	2	207	NA	113	425	30.2	3,400
B-3A	2.5	166		46	82.5	16.3	178
C-1A	3	728	1.1	NA	491	76.7	NA
C-2A	2.5	331	3.7	NA	303	85.9	NA
C-21A <sup>2</sup>	2.5	336	5.6	NA	1110	86.7	NA
C-3A	2	112	0.7	NA	71.4	62.2	NA
C-4A	3	951	1.8	NA	187	180	NA
TTLc		10,000	100	2,500	1,000	2,000	5,000
STLc		100	1.0	25	5	20	250
West U.S. Average		670	-	27	20	19	65

1. NA = Not analyzed.
2. C-21A is a duplicate of C-2A.

B2A Pb - 6 mg/L

SURFACE TO 3' SHOW Pb TTLc  
LEVELS AT 10X STLc LIMITCOMPOSITE SC AT 5' SHOWS Pb  
TTLc AT 126

the SPTCo. High Street property. With the exception of the composite analysis of near-surface samples from area B and from the near-surface sample from boring D, the concentration of arsenic ranged from 4.70 mg/kg to 1.86 mg/kg. The highest concentration detected was 9.54 mg/kg and this occurred in the near-surface sample from boring D. Boring D was situated at the north end of the property in a location that was reportedly never used for metal recycling activities, and as such, was selected to represent soil unaffected by metal recycling activities on the property.

None of the concentrations of arsenic are high enough to yield soluble concentrations in excess of the STLC (5 mg/l) and the concentrations are either less than or very similar to the average concentration (7 mg/kg) found throughout the western United States.

In conclusion, none of the concentrations of arsenic detected indicates that former activities at the property resulted in the introduction of arsenic at levels above naturally occurring concentrations and that the concentrations present are not of environmental concern.

#### Barium

Concentrations of barium were observed to range from 63.5 mg/kg to 1,500 mg/kg throughout the property, although only three results exceeded the average concentration in the western United States of 670 mg/kg. Two composite results and one individual result were high enough that barium could potentially exceed the STLC. The two composites were from the near-surface samples from areas B and C. Subsequent sampling and analysis of individual samples for barium at each of the borings in areas B and C did not reveal concentrations that could exceed the STLC. The concentration detected in the near-surface sample from boring D (1,500 mg/kg) potentially could exceed the STLC.

#### Cadmium

Concentrations of cadmium were uniform throughout the property and only



one result (the composite of the samples from a depth of approximately 5 feet in area C) indicated that the concentrations could potentially exceed the STLC. Since this result is a composite of four individual samples, however, and since individual concentrations from the property are lower than leachable concentrations, it is unlikely that individual samples making up the composite would contain concentrations of chromium that could potentially exceed the STLC.

N.B

#### Chromium

No concentrations of chromium were detected that could potentially exceed the STLC in soil samples throughout the property. Concentrations observed were close to the average concentration in the western United States of 56 mg/kg and concentrations observed in boring D, which was drilled in a location apparently not used for metal recycling activities, were similar to those results from locations where metal recycling activities occurred. In conclusion, chromium soils at the property are most likely naturally occurring and do not occur at levels that are of environmental concern.

#### Hexavalent Chromium

Hexavalent chromium does not occur at levels that are of environmental concern at the SPTCo. High Street property. Concentrations that could potentially exceed the STLC do not occur and the results from boring D are generally similar to the results from the area where metal recycling activities occurred.

#### Cobalt

Concentrations of cobalt that could potential exceed the STLC or concentrations that might be of environmental concern were not observed at the SPTCo. High Street property.

## Copper

Concentrations of copper in two composite samples indicated that individual samples making up the composite could potentially contain concentrations that exceed the STLC. Subsequent sampling of the near surface individual samples from the three borings making up the composite, however, indicated that individual samples did not contain copper at concentrations that could exceed the STLC.

Results from the SPTCo. High Street property range from slightly below to slightly above the average concentration for the western United States of 27 mg/kg. The concentrations of copper observed at the SPTCo. High Street property do not appear to be of environmental concern.

## Lead

During the initial sampling, lead was detected at concentrations that potentially exceed the STLC, in one individual analysis and in all of the composite analyses except those of samples from approximately 10 feet in areas A and B. In the individual analyses from boring D, the concentration in the near-surface sample was markedly higher than in the deeper samples. Similar trends were observed in the composite results from areas A and B. In area C, however, the highest composite lead concentration was observed in the analysis of samples from a depth of approximately 5 feet. Subsequent sampling and analysis of individual near-surface samples from borings in area C, however, indicated that the near-surface individual results, with the exception of boring C-3, were higher than the composite result of samples from 5 feet.

Individual samples from near the surface were collected from borings in areas B and C. The individual results show a large variation in concentrations, with a low concentration of 71.4 mg/kg to a high concentration of 1,100 mg/kg. The individual analyses indicate that the lead results are very difficult to reproduce. Individual sample C-2A and its field duplicate C-21A revealed concentrations of 303 mg/kg and 1,110 mg/kg, respectively, and a laboratory duplicate of C-21A resulted

in a concentration of 600 mg/kg. Results above 50 mg/kg represent concentrations that could potentially exceed the STLC. WET tests for lead were performed on two individual samples, B-2A and C-3A. B-2A contained 425 mg/kg lead and C-3A contained 71.4 mg/kg lead. The WET test results reveal an extractable concentration of 6.06 mg/l in B-2A. This concentration exceeds the STLC of 5.0 mg/l for lead. Several individual samples had total lead concentrations similar to B-2A (B-1A, C-1A, and C-3A) and extractable concentrations of lead in these samples might also be expected to exceed the STLC.

Although several near-surface samples contained lead in concentrations that could potentially exceed the STLC, soil samples from depths of 5 and 10 feet do not contain lead at concentrations that would indicate that the STLC could potentially be exceeded. The WET test result for individual sample C-3A indicated an extractable concentration of 2.16 mg/l, below the STLC of 5.0 mg/l. This sample contained a total lead concentration of 71.4 mg/kg. This concentration is similar to that which would be expected to be found in the individual samples making up the composite analyses from depths of 5 and 10 feet throughout the property.

*N.D. COMPOSITE #8 FROM DEPTHS OF 5.5-6.5 FT SHOWED PB AT 126 PPM.*

In conclusion, total concentrations of lead in several near-surface individual samples were at levels that indicated that the STLC may be exceeded. Results from depths of 5 and 10 feet, however, reveal lead at concentrations below which the STLC would be expected to be exceeded. These results indicate that lead is not migrating significantly into the subsurface and concentrations detected at the surface are not a threat to groundwater in the area.

*EXCEPT COMPOSITE #8 126 PPM.*

The generally decreasing concentrations observed with depth and the low levels detected in shallow groundwater (see Section 4.2.1) suggest that lead in soils at the property probably is not an environmental concern in terms of impacting groundwater quality.

*- GROWS AFTER SEASONALLY FLOODED BETWEEN 6-14"*

### Nickel

Soluble concentrations of nickel that could potentially exceed the STLC were not detected in any individual analyses from the SPTCo. High Street property. Nickel in several composite analyses indicated that concentrations that could potentially exceed the STLC may exist in an individual sample making up the composite, but the generally even distribution throughout the property suggests that markedly higher individual concentrations are not likely to occur.

### Vanadium

Vanadium appears to be fairly evenly distributed throughout the property. The highest concentration detected (56.4 mg/kg) is less than the average concentration in the western United States (88 mg/kg). None of the concentrations observed on the SPTCo. High Street property indicate that concentrations that could potentially exceed the STLC occur. In conclusion, vanadium in soils does not appear to be an environmental concern at the SPTCo. High Street property.

### Zinc

Zinc at a concentration that potentially could exceed the STLC was detected only in the composite analysis of near-surface samples from the borings in area A. This analysis was a composite of three individual samples. Resampling near the three borings in area B revealed one sample, B-2A, that could potentially contain concentrations that exceed the STLC. A WET test for zinc was performed on sample B-2A and revealed an extractable concentration of 268 mg/l. This concentration is slightly above the STLC of 250 mg/l for zinc. The total concentration of zinc in this sample is 3,400 mg/kg. This concentration was practically twice as high as any other total concentration detected. Concentrations detected from depths of 5 and 10 feet in composite analyses were low enough that individual samples making up a composite would not be expected to contain zinc at a level above 1,500 mg/kg, and thus extractable concentrations in samples from depths of 5 and 10 feet

would not be expected to exceed the STLC.

In conclusion, the occurrence of zinc at a concentration that resulted in an extractable concentration exceeding the STLC was extremely localized and restricted to the near surface. Concentrations detected in samples from 5 and 10 feet indicate that zinc is not migrating into the subsurface significantly and is not impacting the quality of groundwater.

#### 4.1.2 Organic Results

Organic analyses included analysis for total petroleum hydrocarbons (TPH), total oil and grease (TOG), polychlorinated biphenyls (PCBs), and purgeable organics. Results of the initial sampling are presented in Table 4-3, results of the second sampling are presented in Table 4-4.

##### Total Oil and Grease

The highest concentrations of TOG occurred in the near-surface composite samples from area C and boring D. TOG at 2,800 mg/kg was detected in the near-surface composite from area C and 1,300 mg/kg was detected in the near-surface individual analysis from boring D. These two results are an order of magnitude higher than any other TOG result and compare closely with the concentration of TPHs detected.

##### Total Petroleum Hydrocarbons

The distribution of TPHs was generally similar to that of TOG. The highest concentrations occurred in the near-surface composite from area C (1,600 mg/kg) and the near-surface individual sample from boring D. TPH at 150 mg/kg in the composite of samples from 5 feet from area C was the only sample where detection at depth was greater than the near-surface samples, indicating that TPH is largely restricted to the top 2 to 3 feet throughout the property and that migration into the subsurface is minimal.

*Only surface*

Table 4-3

INITIAL SOIL SAMPLE RESULTS - ORGANICS  
(mg/kg dry weight, ppm)

Composite No.	1	2	3	4	5	6	7	8	9	10	5	5 (dup)	10	STLC	STLC	TTLC
Area	A	A	A	B	B	B	C	C	C	D	D	D	D	D	D	D
Depth (Feet)	0	5	10	0	5	10	0	5	10	0	5	5	10	STLC	STLC	TTLC
<b>GENERAL:</b>																
Oil & Grease	25	ND	ND	15	44	41	2,800	220	26	1,300	ND	ND	ND	ND	ND	ND
TPH	56	ND	ND	ND	ND	ND	1,600	150	ND	90	ND	ND	ND	ND	ND	ND
<b>PURGEABLE ORGANICS:</b>																
Meth. Chloride	<6*	<6*	ND	<6*	<6*	<6*	6	7	ND	<6*	6	<6*	<6*	<6*	<6*	<6*
Acetone	84	55	38	30	58	56	73	83	23	120	74	21	15	15	15	15
Trans, 1,2 DCE	ND	ND	ND	ND	ND	ND	ND	<6*	ND	ND	ND	ND	ND	ND	ND	ND
Toluene	ND	ND	ND	ND	ND	ND	ND	<6*	ND	ND	ND	ND	ND	ND	ND	ND
Ethyl Benzene	ND	ND	ND	ND	ND	ND	ND	<6*	ND	ND	ND	ND	ND	ND	ND	ND
Xylenes	ND	ND	ND	ND	ND	ND	ND	18	ND	ND	ND	ND	ND	ND	ND	ND
<b>PCBs:</b>																
PCB-1242	0.15	<0.06*	<0.06*	ND	ND	ND	2.3	0.77	0.30	ND	ND	ND	ND	ND	ND	ND
PCB-1254	0.19	<0.06*	0.087	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
PCB-1260	ND	ND	ND	ND	ND	ND	14	2.6	0.88	0.09	ND	ND	ND	ND	ND	ND
TOTAL	0.34	tr	0.092	-	-	-	16.3	3.37	1.18	0.09	-	-	-	-	-	50
AROCLORS																
<i>STLC = 50</i>																

\* Compound present but at a level below the measurable detection limit.

Table 4-4  
INDIVIDUAL SOIL SAMPLE RESULTS - ORGANICS (mg/kg)

Samples	Depth (feet)	TPH <sup>1</sup>	PCBs <sup>2</sup>
C-1A	3	ND <sup>4</sup>	0.02
C-2A	2.5	114	0.18
C-21A <sup>3</sup>	2.5	772	0.46
C-3A	2	86	ND
C-4A	3	10	2.8
TTLC			50
STLC			5.0

1. TPH = Total Petroleum Hydrocarbons.
2. PCBs = Polychlorinated Biphenyls.
3. C-21A is a duplicate of C-2A.
4. ND = Not Detected.

RESULTS OF DISCRETE  
SAMPLING IN AREA C  
WHERE COMPOSITE SAMPLING  
FOUND 11.00 N/AH TPD.

// COMPOSITE SAMPLES  
AT SURFACE FOUND:  
2400 TOG, 1100 TPH

AT 5' FOUND  
220 TOG  
150 TPH

Based on the TPH exceeding the state action level of 1,000 mg/kg in the near surface composite from area C, individual samples from similar depths were subsequently collected from near the borings in area C (see Table 4-4 for results). TPHs detected were greater than 100 mg/kg in only one of the four samples, C-2A. This sample and its duplicate, C-21A, had 114 mg/kg and 772 mg/kg, respectively, indicating how extremely variable TPH is in soils on the property. All of the individual results were substantially lower than the composite result. The individual samples were collected from trenches excavated as closely as possible to the original borehole, generally less than 1 foot away. This further indicates the extremely localized and variable nature of TPH on the property.

*DISCRETES TAKEN AT 25' & COMPOSITE AT SURFACE*

*THOUGH A CORRELATION SHOWN BETWEEN TPH & TOG FINDINGS, ONLY WENT SAMPLES EVALUATED FOR TOG*

#### Purgeable Organics

Organic chemicals other than methylene chloride and acetone were detected only in the composite of 5-foot samples from area C. Methylene chloride and acetone were detected at similar low concentrations in all of the soil samples and they were most likely introduced in the laboratory. No known past activities at the property would have resulted in the widespread low-level distribution of these chemicals.

The organic chemicals detected in the composite of 5-foot samples from the borings in area C are common constituents of fuel (gasoline) or their breakdown products. The low concentrations indicate that subsurface gasoline contamination is not significant.

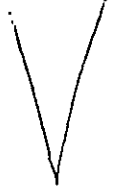
#### Polychlorinated Biphenyls

Three different aroclors were detected in soil samples from the SPTCo. High Street property, PCB-1242, 1254, and 1260. PCB-1242 was detected in area A and C, PCB-1254 was detected only in area A, and PCB-1260 was detected in area C and boring D. PCBs were not detected in samples from area B. For the purposes of the following discussion the different aroclors have been grouped together.



composites 12-5

In areas A and C, total PCBs were detected in composite samples from all depths. In area A, PCBs were not detected at levels above 1 mg/kg. In area C, the highest concentration (16 mg/kg) was detected in the near-surface composite with the 5-foot and 10-foot composites containing progressively lower concentrations.



discrepancies



To further assess the distribution of PCBs in area C, individual near-surface samples were collected from trenches excavated as closely as possible to the original boring. Results of this resampling are presented in Table 4-4. The highest concentration detected was 2.8 mg/kg in sample C-4A. PCBs were not detected in sample C-3A and concentrations less than 1 mg/kg were detected in samples C-1A and C-2A. The lower PCB concentrations detected during the resampling, which were lower than those originally detected, indicate that PCBs are extremely localized on the property. PPC FINDINGS FOR PCB?

#### 4.2 GROUNDWATER RESULTS

##### 4.2.1 Inorganic Results

Groundwater samples were analyzed for Title 22 CCR metals. Results are presented in Table 4-5. In general, similar metals were detected in each sample. Metals detected included barium, copper, lead, and zinc, although copper was not detected in groundwater from monitoring well B-2. A duplicate was collected from monitoring well A-1. Hexavalent chromium, mercury, and nickel were detected in only one of the two samples analyzed from well A-1, so the occurrence of these three metals is questionable.

None of the metals in groundwater at the SPTCo. High Street property exceeds either federal or state drinking water standards.

##### 4.2.2 Organic Results

Organic analyses included analysis for TOG, TPH, purgeable organics, and PCBs. Results are presented in Table 4-5.

Table 4-5

SPTCO. HIGH STREET - GROUNDWATER RESULTS

Well No.	May 26, 1989 Sampling				July 28, 1989 Sampling			EPA Standard
	A-1	A-1 Dup	B-2	C-2	Blank	C-2	C-2 Duplicate	
<b>GENERAL:</b>								
Oil & Grease (mg/l)	2.3	1.4	2.8	2.4	1.6			
TPH (mg/l)	ND <sup>1</sup>	1.4	1.5	ND	ND			
<b>METALS:</b>								
Antimony (mg/l)	ND	ND	ND	ND	ND			
Arsenic (mg/l)	ND	ND	ND	ND	ND			0.05
Barium (mg/l)	0.086	0.072	0.037	0.027	ND		1.0	1.0
Beryllium (mg/l)	ND	ND	ND	ND	ND			
Cadmium (mg/l)	ND	ND	ND	ND	ND			0.01
Chromium (mg/l)	ND	ND	ND	ND	ND			0.05
Cr <sup>6+</sup> (mg/l)	0.063	ND	ND	ND	ND			
Cobalt (mg/l)	ND	ND	ND	ND	ND			
Copper (mg/l)	0.013	0.019	ND	0.014	ND			
Lead (mg/l)	0.030	0.024	0.006	0.019	ND			0.05
Mercury (mg/l)	0.0012	ND	ND	ND	ND			0.002
Molybdenum (mg/l)	ND	ND	ND	ND	ND			
Nickel (mg/l)	0.018	ND	ND	ND	ND			
Selenium (mg/l)	ND	ND	ND	ND	ND			0.01
Silver (mg/l)	ND	ND	ND	ND	ND			0.05
Thallium (mg/l)	ND	ND	ND	ND	ND			
Vanadium (mg/l)	ND	ND	ND	ND	ND			
Zinc (mg/l)	0.085	0.054	0.045	0.110	ND			

mbe/spea/t4-5

540 - 2158

Table 4-5 (Cont.)

Well No.	May 26, 1989 Sampling				July 28, 1989 Sampling			EPA Standard
	A-1	A-1 Dup	B-2	C-2	Blank	C-2	C-2 Duplicate	
PCBS:								
PCB-1260 (ug/l)	ND	ND	ND	1.0	ND	0.61	0.78	ND
<u>PURGEABLE ORGANICS:</u>								
Methylene Chloride (ug/l)	20	18	<5*	<5*	17			40
Acetone (ug/l)	14	<10*	31	55	<10*			

mbe/spea/t4-5

1. ND = Not Detected.

Total Oil and Grease

503 A+E?

Low levels of oil and grease were detected in groundwater from all of the monitoring wells. The maximum level detected was 2.8 mg/l. Oil and grease at 1.6 mg/l was also detected in the travel blank. This occurrence, at levels that were similar to those found in the environmental samples, suggest that the oil and grease may have been introduced either during sampling or in the laboratory.

- Floating product evaluated?  
- How sampled?

Total Petroleum Hydrocarbons

ANALYSIS 8015?

PLS P 5030?

Very low levels of TPHs were detected in groundwater from monitoring well B-2 and in the duplicate from monitoring well A-1. The detection of TPHs at a very low level (1.4 mg/l) in only one of two samples analyzed from monitoring well A-1 makes the environmental occurrence of TPHs in groundwater suspect.

- Floating product evaluated?  
- How sampled?

Purgeable Organics

The only purgeable organic chemicals detected in groundwater were methylene chloride and acetone. Both occurred at low levels and also in the blank. Occurrences of both chemicals are most likely due to introduction in the laboratory.

- Why weren't BTEX looked for  
- Which test method used?

8010 }  
8015 }  
why not 8020 too?  
or 8240

Polychlorinated Biphenyls

During the groundwater sampling of May 26, 1989, Aroclor 1260 at 1.0 ug/l in monitoring well C-2 was the only PCB detected. The laboratory analytical data QA/QC report (see Appendix B), however, concluded that PCB groundwater results were suspect.

Because the recommended maximum contaminant level for PCBs in drinking water is zero, this well was resampled on July 28, 1989. During this sampling PCBs were detected in both the environmental sample (0.61 ug/l) and its duplicate (0.78 ug/l), confirming the detection of May 26, 1989.

Monitoring well C-2 is in area C where the highest levels of PCBs were detected in soil samples. The soil composite of samples from a depth of 10 feet (the approximate water table) in area C contained PCBs at 1.18 mg/kg.

5. CONCLUSIONS AND RECOMMENDATIONS

AGNLE  
Subsurface soil and groundwater sampling at the SPTCo. High Street property revealed that generally low levels of a variety of contaminants are present at the site in apparently localized occurrences.

AGNLE  
Analysis of soil samples for priority pollutant metals did not reveal any metals at levels that exceed the TTLC and, thus, none of the soils classify as hazardous waste for disposal purposes in terms of metal content. WET tests for lead indicate that the STLC may be exceeded in several samples from near the ground surface. Subsurface sample results, however, are lower and indicate that lead is not migrating into the subsurface and is not adversely impacting groundwater quality. A WET test for zinc, performed on one individual sample with a total zinc concentration markedly higher than other samples, revealed that the STLC was slightly exceeded. This markedly high concentration was from the near surface and appeared to be atypical. Subsurface zinc concentrations were substantially lower and indicated that zinc is not migrating into the subsurface and is not adversely impacting groundwater quality.

AGNLE  
Organic chemicals commonly associated with gasoline, in particular toluene, ethylbenzene, total xylenes, and trans 1,2 dichloroethene, were detected in soils at very low levels in the composite of 5-foot samples from area C. The low levels and lack of similar chemicals in groundwater indicate that this occurrence is not a significant environmental concern.

TPHs were observed in the composite of near-surface samples in area C (1,600 mg/kg) and in the near-surface sample from boring D (900 mg/kg). These either are greater than or are very close to the California Water Resources Control Board action level for TPHs of 1,000 mg/kg. During the field activities, several small areas of oil-stained surface soils were observed that most likely had relatively high TPH levels. These

oil-stained surface soils were not necessarily sampled, however, since the focus of E & E's investigation was on evaluating the subsurface migration of contaminants. Data obtained concerning subsurface TPHs and the on-site observations of oil-stained surface soils suggests that the areas with high levels of TPHs are of limited extent both horizontally and vertically.

PCBs were detected in subsurface soil composite samples to a depth of 10 feet in both areas A and C. The concentrations in area A were less than 1 mg/kg and are not considered of environmental concern. Higher PCB concentrations were observed in area C with the near surface composite analysis (made up of four individual samples) containing 16.3 mg/kg. This implies that one of the individual samples making up the composite could potentially have a PCB concentration four times the reported level, or 65.2 mg/kg. A level such as this would exceed the state action level of 25 mg/kg for environmental settings similar to that at the SPTCo. High Street property.

The only constituent detected in groundwater of environmental concern are PCBs at a level approaching 1 ug/l. The RMCL for PCBs in drinking water is zero. This standard may not be applicable to the SPTCo. High Street property because the yields of water are so low as to preclude the shallow groundwater from being developed as a source for drinking water supply.

In conclusion, the occurrence of contaminants of environmental concern in soils at the SPTCo. High Street property appears to be restricted in both horizontal and vertical directions. The constituents of concern are primarily localized and near-surface occurrences of petroleum hydrocarbons and PCBs. Metal concentrations in soils do not appear to be at levels that are of environmental concern. The lack of gasoline-related constituents detected suggests that the petroleum hydrocarbons are primarily in the form of less volatile oils and greases. This conclusion is supported by the close correlation between TPH and TOG concentrations.

PCB  
16.3 mg  
Composite

BTex?

Ab net

?

J.K

The only constituents detected in groundwater that may be of environmental concern are PCBs; however, its environmental significance will be dependent on the potential uses of the shallow groundwater at the property.

### Recommendations

Based on the data obtained during this investigation and on observations made in the field, E & E recommends that the following actions be taken:

- 1) o Excavate and remove any soil visibly stained with oil and also any material that appears to be metallic slag. Excavated soil or slag should be treated as a hazardous waste for disposal purposes.
- 2) o Determine, in conjunction with the appropriate agency, whether remediation is needed in terms of the PCBs detected in groundwater.

- CONTINUE GROUND WATER MONITORING

- EVALUATE AREA FOR BACKGROUND LEVELS OF PCB.

1 - DEFINE DEPTH & ZONES TO BE SCRAPPED



REFERENCE

1. Shacklette, H.T. and Boerngen, J.G., 1984, "Element Concentrations in Soils and Other Surficial Materials of the Conterminous United States," U.S. Geological Survey Professional Paper 1270.

# BORING A-1

DATE DRILLED: 5/23/89

DRILL METHOD: Hollow Stem Auger

SAMPLING METHOD: Split Spoon

DEPTH IN FEET	BLOWS/0.5 FT	SAMPLE NO.	LOCATION	SYMBOLS	DESCRIPTION	NOTES
0					Concrete	
					Black sandy gravel (fill)	
8				CL	Black clay, stiff, mottled, little sand	Sampler would not drive past 1 ft, drilled to 2.5 ft to take sample
10						Sample is wet
14		A-1A				HNu=14ppm
5				ML CL	Tan to orange gray brown clayey silt to silty clay, moderately stiff	HNu=40 ppm
6		A-1B				50% recovery
13					Tan brown clayey silt	Cuttings wet at 7 ft
10						
6						
20		A-1C		GC	Tan to orange gray brown clayey sandy gravel, wet	HNu=8 ppm borehole
26						0 ppm breathing zone
15				ML CL	Gray clayey silt, moist, stiff, mottled, with rootlets	Sampler is wet
2						HNu=3 ppm borehole,
3		A-1D				0 ppm breathing zone
6						
20					Gray and brown mottled clayey silt, stiff	HNu=0 ppm
4						
7		A-1E				
11						
25					Gray and brown mottled clayey silt, stiff	HNu<1 ppm
4						Bottom of hole =
8		A-1F				11.5 feet
14						
30						

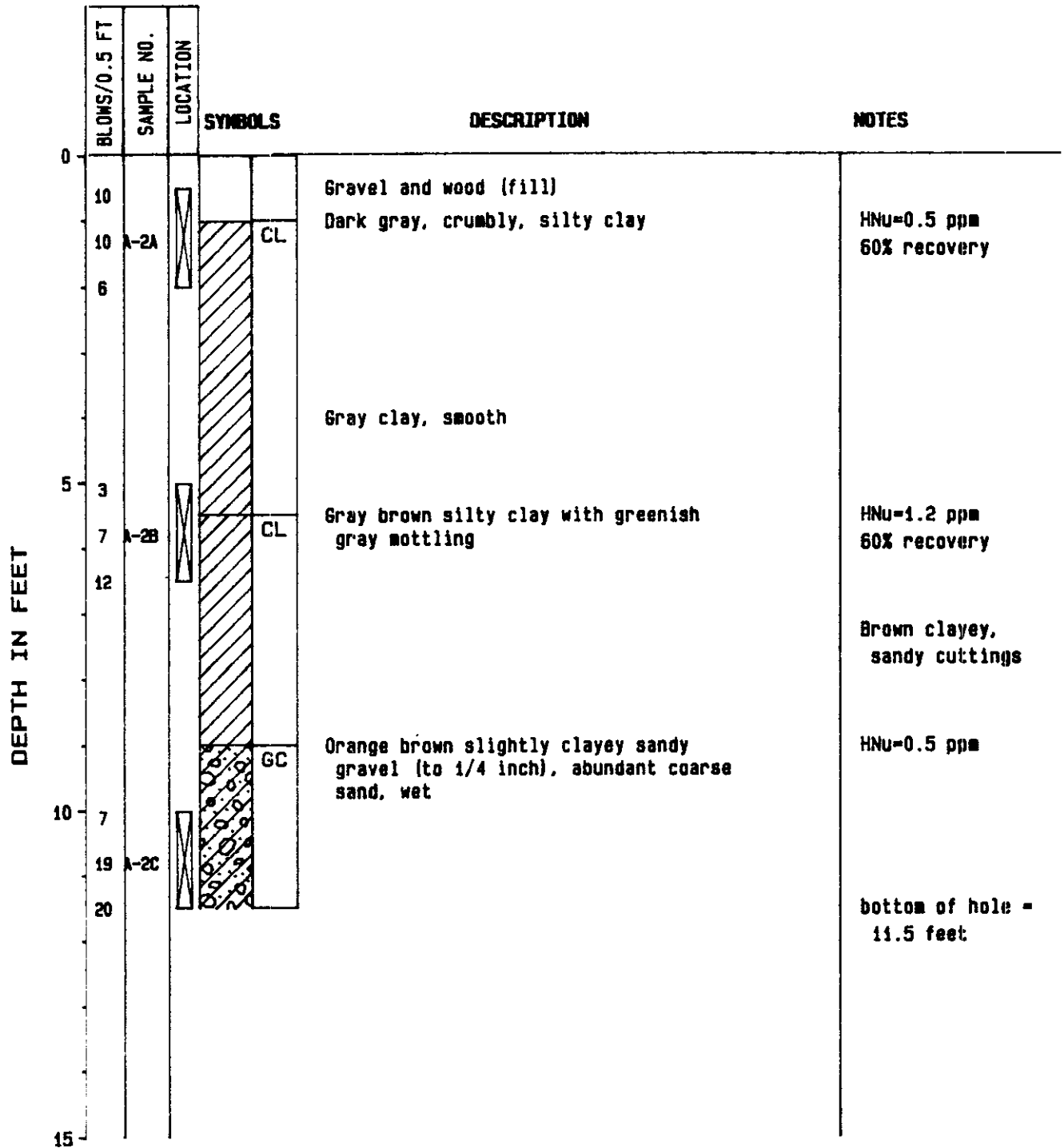
11A

# BORING A-2

DATE DRILLED: 5/22/89

DRILL METHOD: Hollow Stem Auger

SAMPLING METHOD: Split Spoon

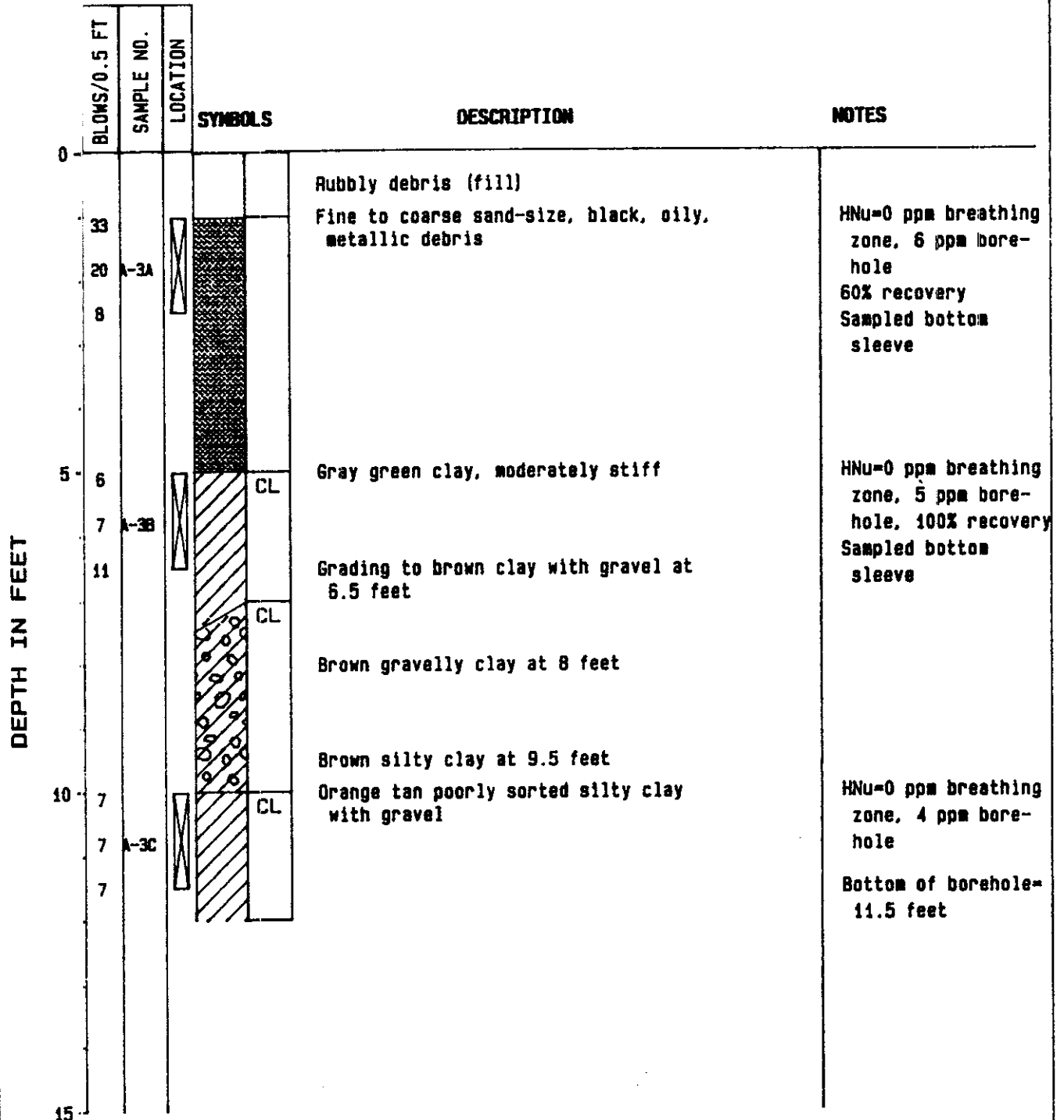


# BORING A-3

DATE DRILLED: 5/23/89

DRILL METHOD: Hollow Stem Auger

SAMPLING METHOD: Split Spoon

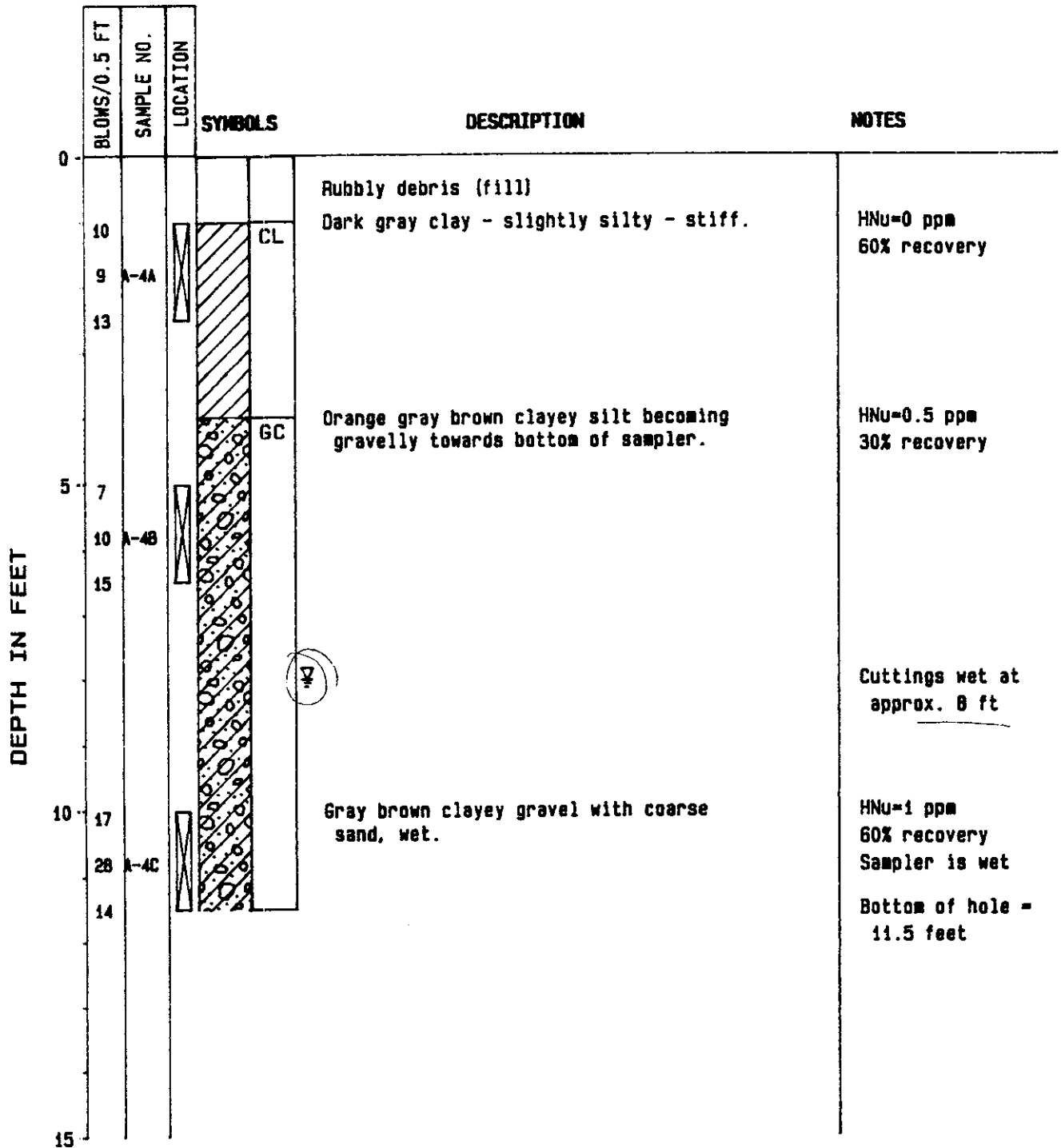


# BORING A-4

DATE DRILLED: 5/22/89

DRILL METHOD: Hollow Stem Auger

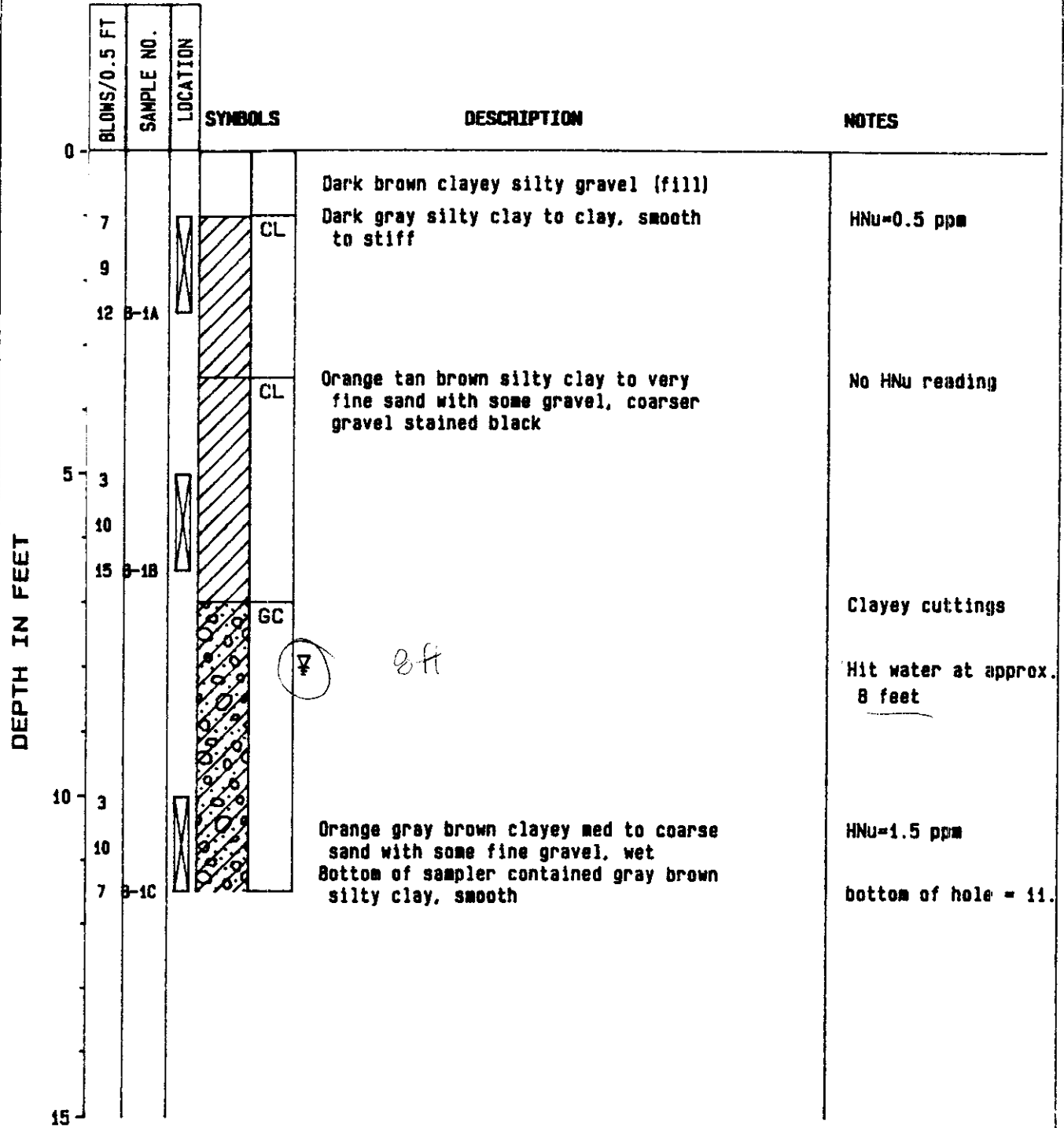
SAMPLING METHOD: Split Spoon



# BORING B-1

DATE DRILLED: 5/22/89

DRILL METHOD: Hollow Stem Auger  
 SAMPLING METHOD: Split Spoon



# BORING B-2

DATE DRILLED: 5/23/89

DRILL METHOD: Hollow Stem Auger  
 SAMPLING METHOD: Split Spoon

DEPTH IN FEET	BLOMS/0.5 FT	SAMPLE NO.	LOCATION	SYMBOLS	DESCRIPTION	NOTES
0				GW	Dark gray to black sandy gravel (fill)	Refusal after 1 ft
13		B-2A		GW	Black sandy gravel with wood, oil stained	Sample collected at 2 feet, metallic slag in sampler HNU=1 ppm cuttings
5	12	B-2B		GC	Orange tan clayey gravelly coarse sand, very poorly sorted	No HNU reading
10	11	B-2C		GC	Orange tan gravelly clayey coarse sand, moist, fine gravel to 1/8 inch	HNU=0.5 ppm Sampler is wet 100% recovery
15	9	B-2D		GC	Tan clayey gravel (1/4 inch), wet	
15	3	B-2E		CL ML	Gray and brown mottled clay, stiff, smooth	HNU=1 ppm Sample collected of gravel 60% recovery
20	5	B-2F			Gray and gray brown mottled silt, clayey silt and very fine sandy silt	HNU=2 ppm 60% recovery
25	4	B-2F			Gray clayey silt, stiff	HNU=0.5 ppm 100% recovery Bottom of hole = 26.5 feet

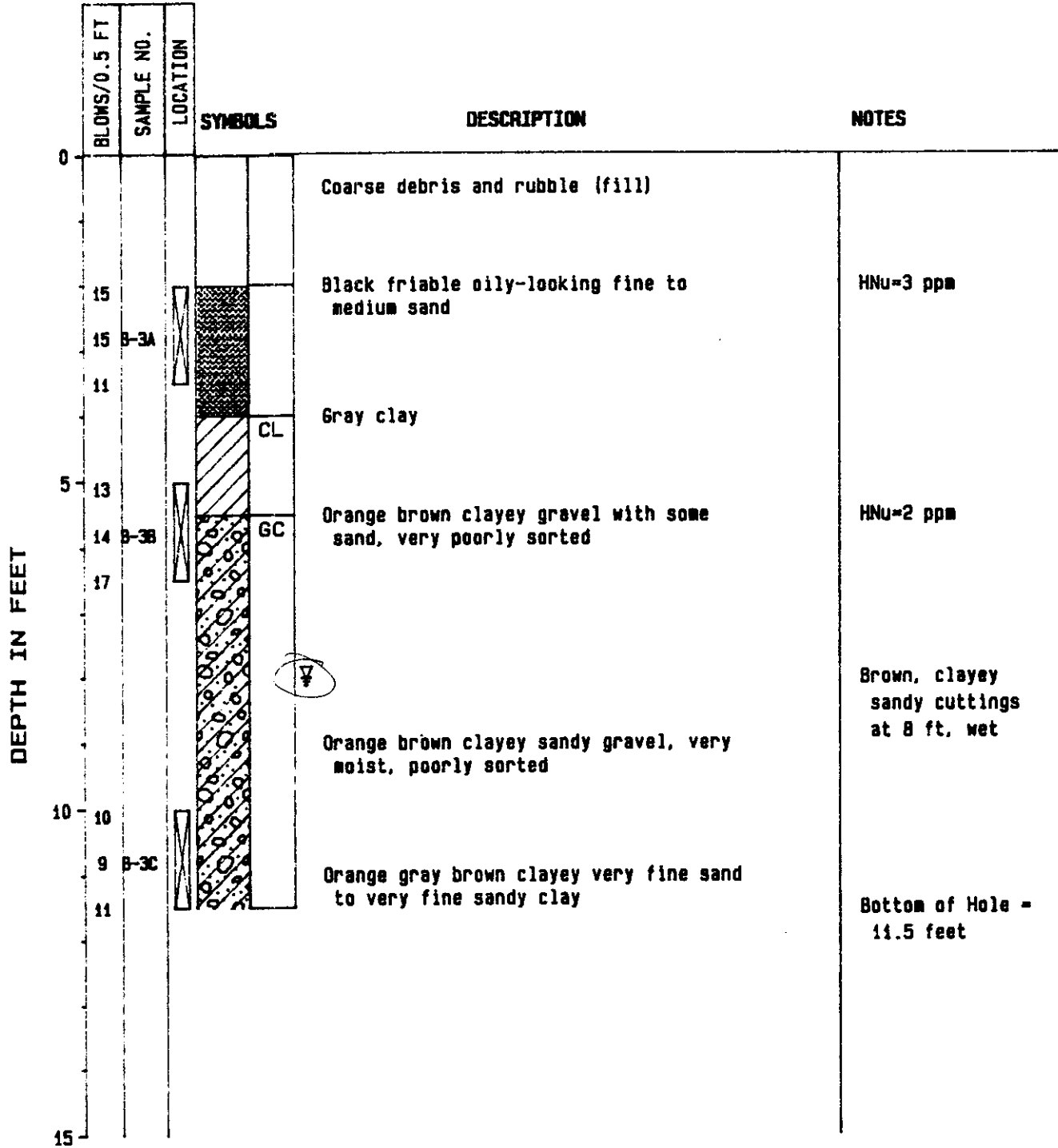
8 ft

# BORING B-3

DATE DRILLED: 5/22/89

DRILL METHOD: Hollow Stem Auger

SAMPLING METHOD: Split Spoon



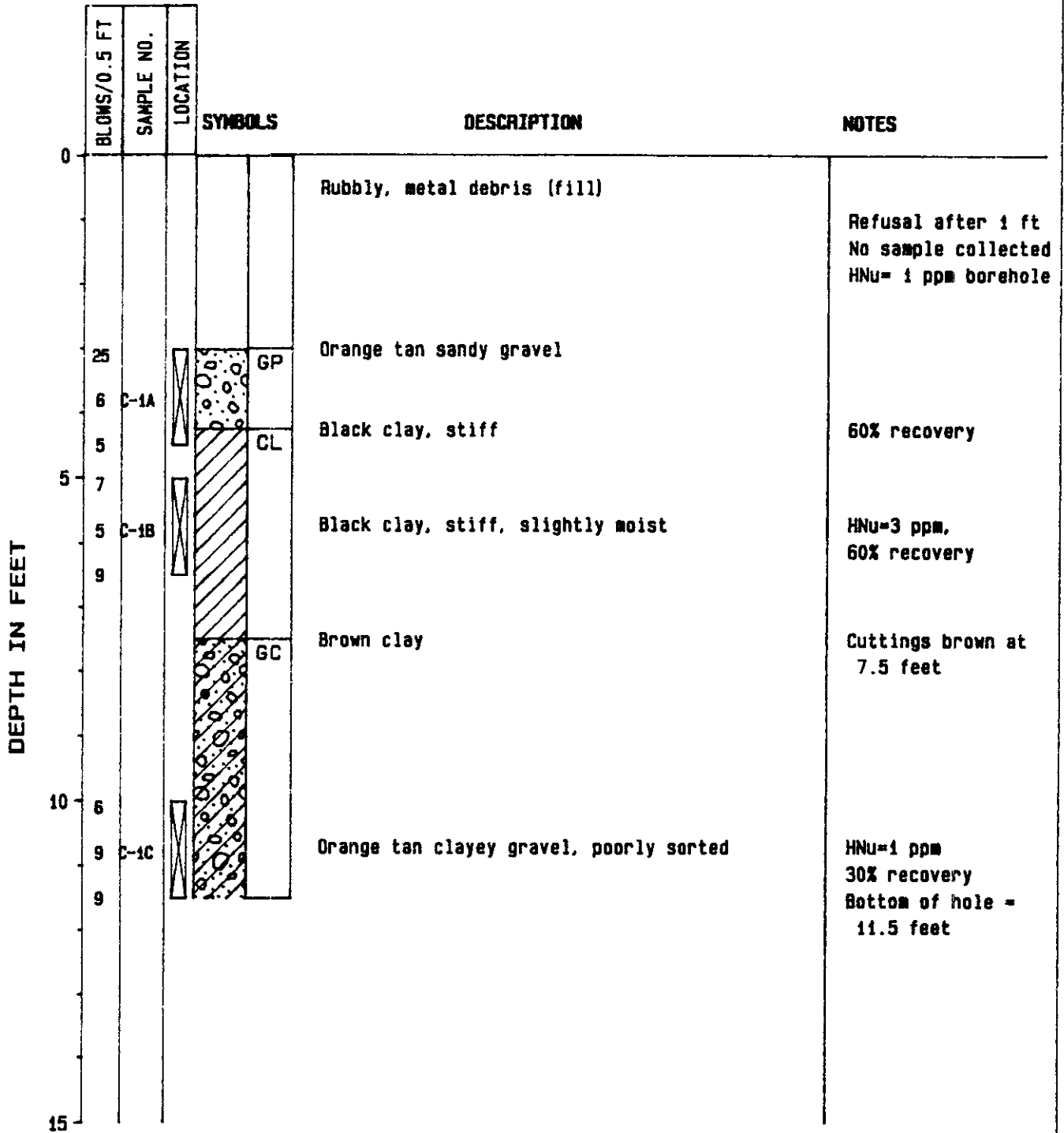
8A



# BORING C-1

DATE DRILLED: 5/24/89

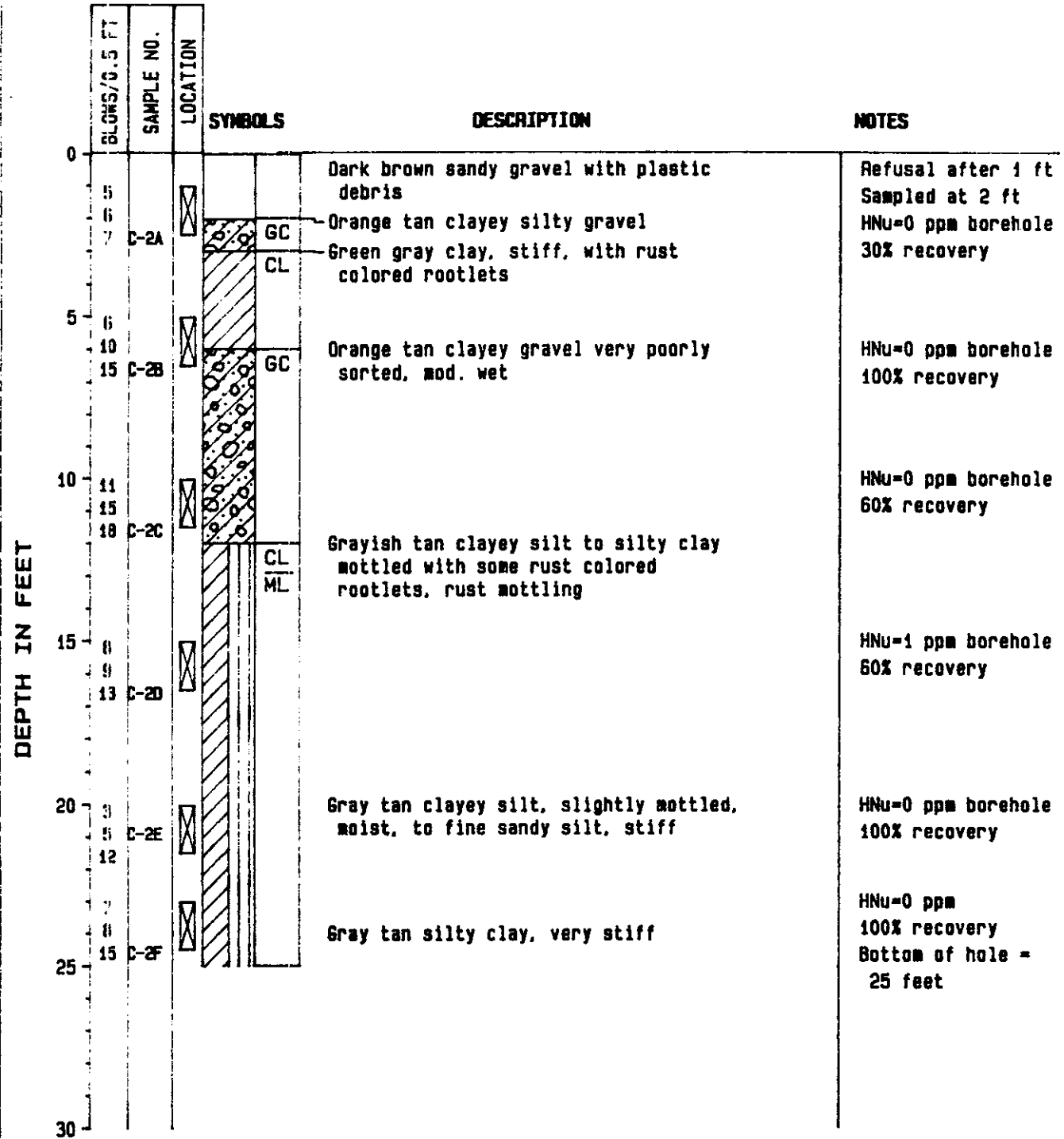
DRILL METHOD: Hollow Stem Auger  
 SAMPLING METHOD: Split Spoon



# BORING C-2

DATE DRILLED: 5/24/89

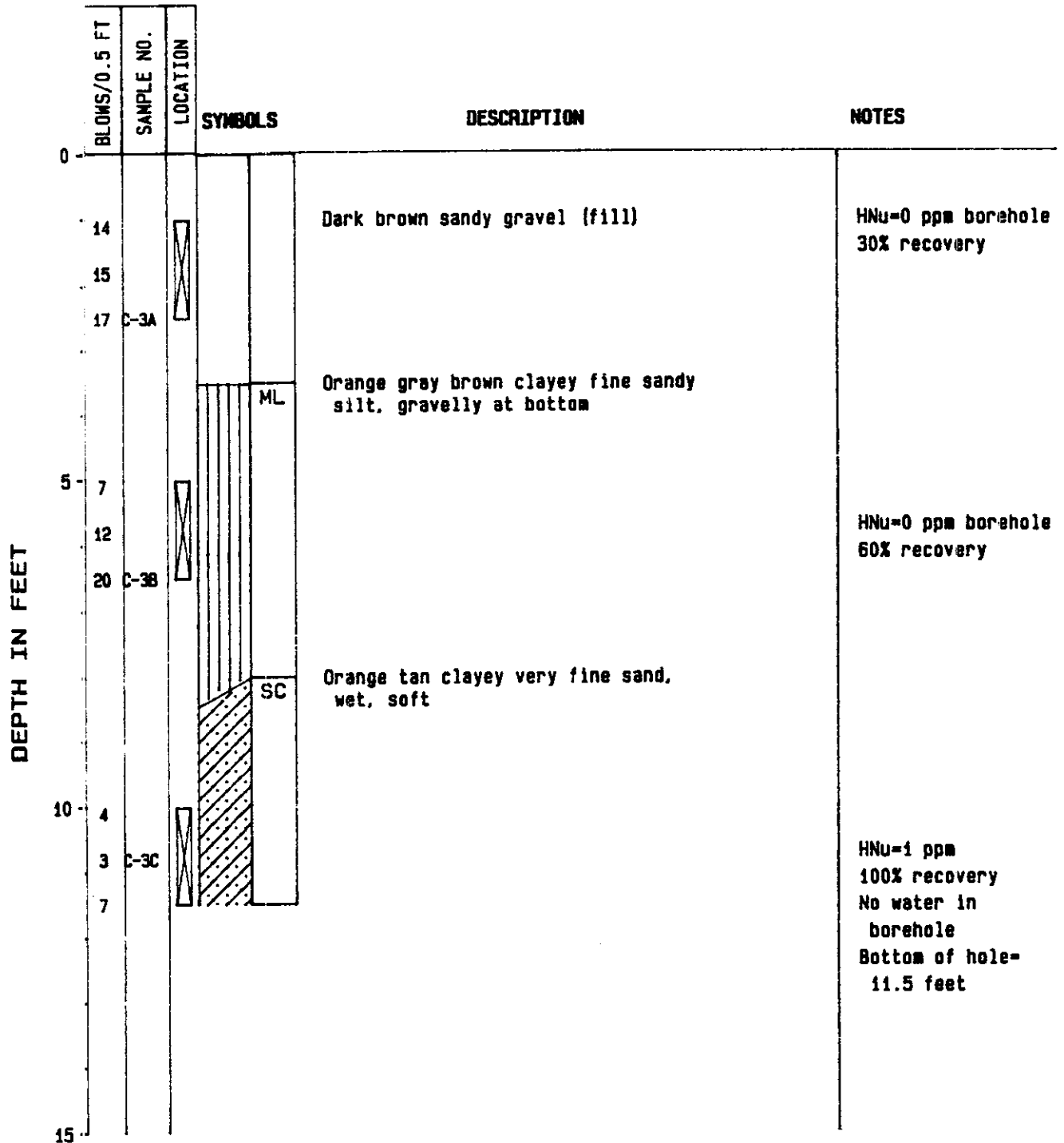
DRILL METHOD: Hollow Stem Auger  
 SAMPLING METHOD: Split Spoon



# BORING C-3

DATE DRILLED: 5/24/89

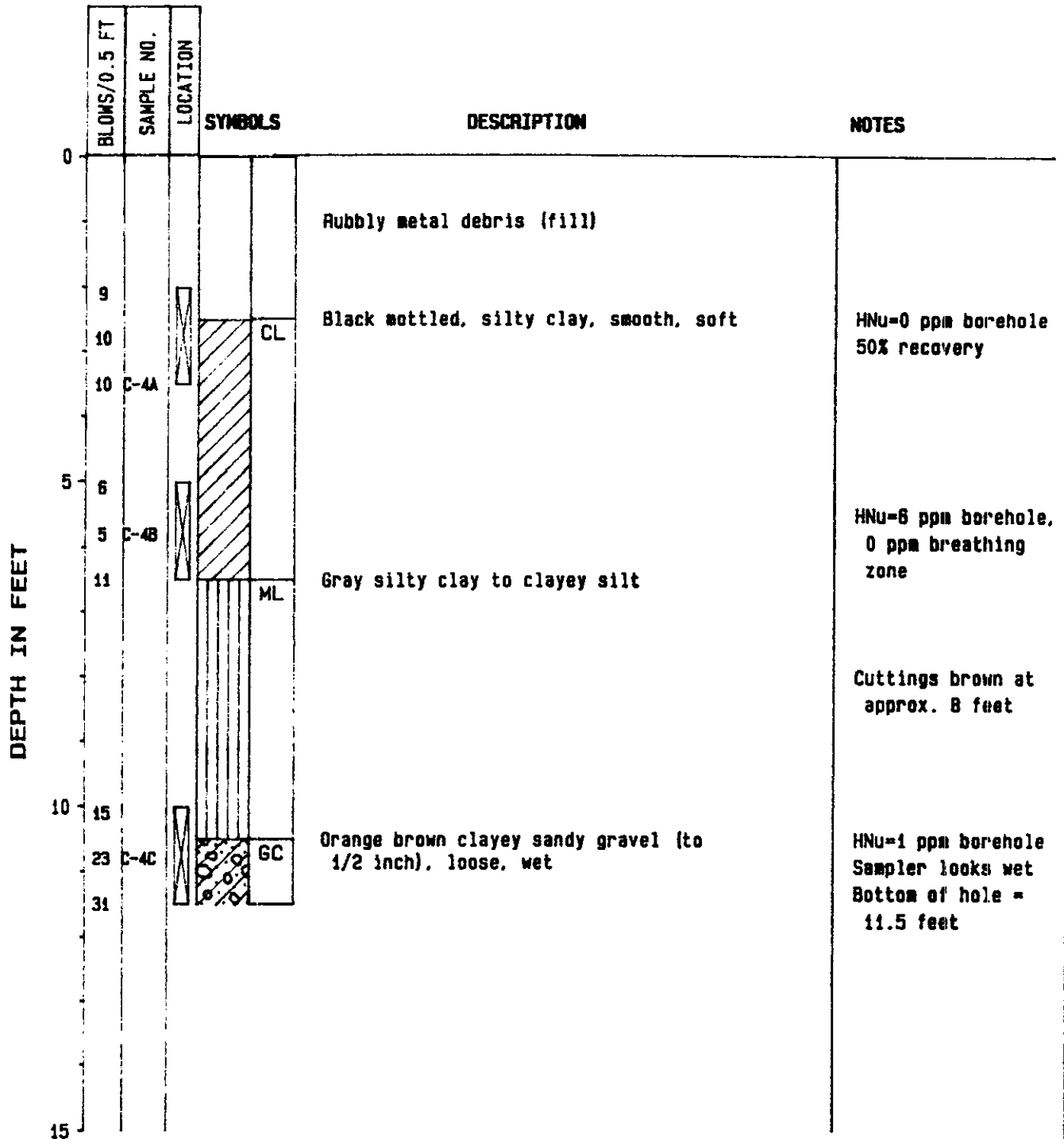
DRILL METHOD: Hollow Stem Auger  
 SAMPLING METHOD: Split Spoon



# BORING C-4

DATE DRILLED: 5/24/89

DRILL METHOD: Hollow Stem Auger  
 SAMPLING METHOD: Split Spoon

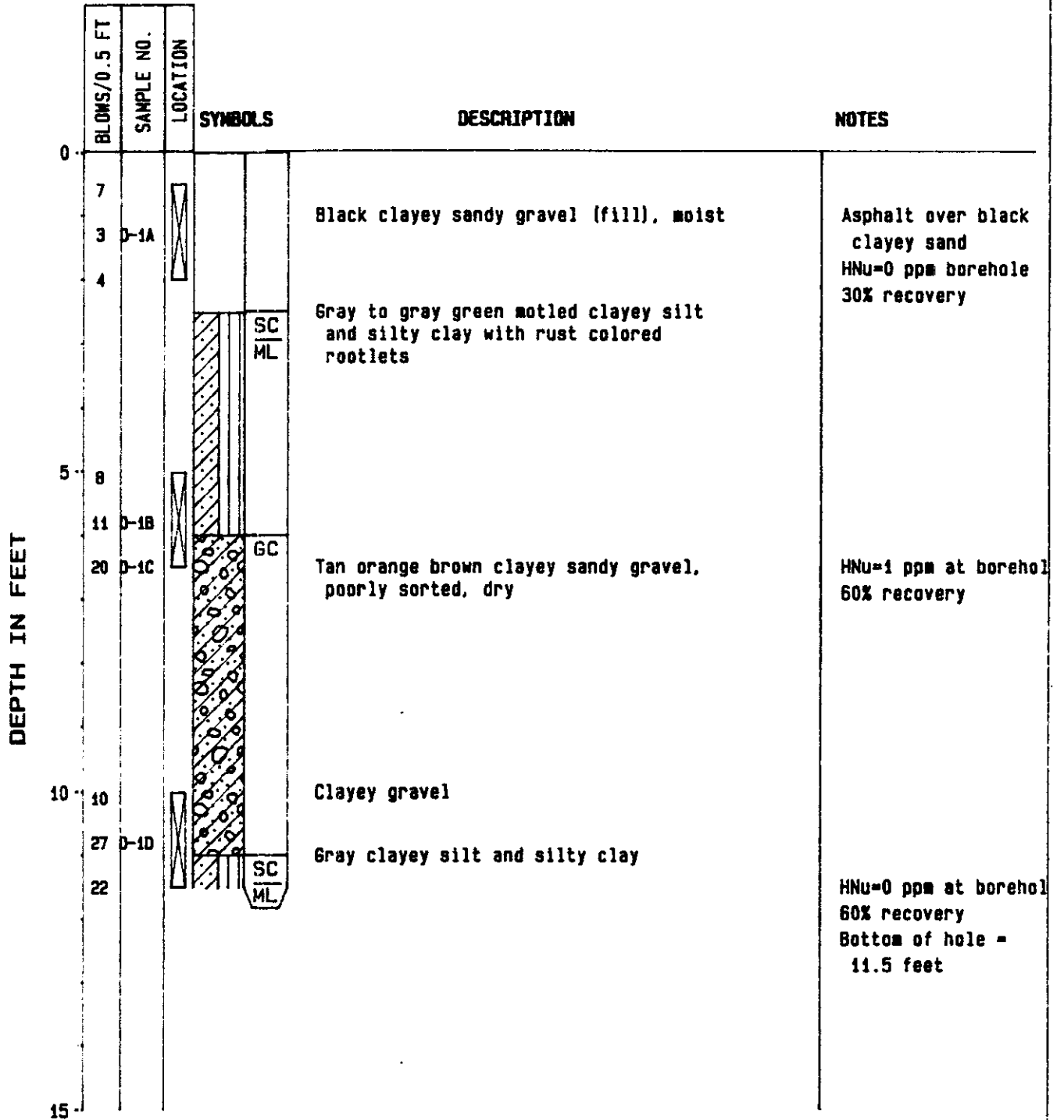


# BORING D-1

DATE DRILLED: 5/24/89

DRILL METHOD: Hollow Stem Auger

SAMPLING METHOD: Split Spoon



## Appendix B

### LABORATORY ANALYTICAL DATA QA/QC

Quality assurance/quality control (QA/QC) protocols for the analysis of the test parameters in groundwater and soil samples were performed to monitor laboratory precision and accuracy. Precision is tested by analyzing samples in replicate which provides a calculation of a relative percent difference (RPD). Accuracy is measured by spiking the sample with the same analyte or a similar compound (e.g., surrogate chemicals) and calculating a percent recovery (%R). These QA/QC results are presented along with the laboratory reports.

For the soil samples, oil and grease and total recoverable petroleum hydrocarbons were within precision and accuracy criteria. Therefore, these results, presented in Table 4-3, are considered valid.

The metals results presented in Table 4-1 had several parameters outside QC limits because of the heterogeneous nature of the soils. EPA methodology require QC limits for RPD to be +20% for concentrations five times above the method detection limit and %R to be 75-125% for acceptable QC limits. Antimony had a low %R of 49%, however, the sample results were all non-detected. This out of control recovery is considered acceptable. Precision QC for arsenic was outside the limits, however, this was due to the low concentration in the sample (less than five times the method detection limit). Arsenic results are, therefore, considered valid. Barium, chromium, copper, lead, and zinc were outside QC limits for precision because the results are not reproducible. Hexavalent chromium, copper, molybdenum, and zinc were outside QC limits for accuracy because the results are not reproducible. Molybdenum results, however, are considered acceptable because the %R was 132% and sample concentrations were all <3.5 ug/kg.

The results presented on Table 4-3 for organic analysis of soils had QC testing within acceptable limits except for method blanks. Methylene chloride blank concentrations were at trace levels (<5 ug/kg) and

acetone blank concentrations were at 11, 10, and 10 ug/kg. Sample results must be at least 10 times greater than the blank concentrations, therefore all methylene chloride and acetone results are considered invalid due to laboratory contamination.

For the groundwater sample results presented on Table 4-5, QC was acceptable for total oil and grease and total recoverable petroleum hydrocarbons. The travel blank contained 1.6 ug/l total oil and grease which may be attributable to field techniques and/or laboratory contamination. All of the total oil and grease groundwater results are approximately the same concentration as the travel blank and, therefore, the total oil and grease in groundwater should be considered of questionable validity.

Precision and accuracy for metals analysis in groundwater (see Table 4-5) were within QC criteria except for copper and zinc (RPD of 33% and 57%, respectively) and lead and selenium (%R of 73% and 60%, respectively). Results for these four metals should be considered suspect except for selenium because it was not detected in any sample.

Organic analysis of groundwater (Table 4-5) had several QC problems. PCB analysis was out of control for accuracy with a %R of 38%, therefore, PCB water results should be considered suspect. The results for methylene chloride and acetone are considered invalid because the method blanks had methylene chloride concentrations of <5 and 7 ug/l and acetone concentrations of 11 ug/l. This was due to laboratory contamination.



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### LABORATORY REPORT

FOR

SOUTHERN PACIFIC RAILROAD-HIGH STREET

S7/850/001

Job No.: 850.001		RE: SP8030				
Sample Date: 5/25/89		P.O. No.:				
Date Received: 5/25/89		Sampled By: E & E, Inc.				
Sample Type: Solid		Delivered By: Federal Express				
E & E Lab. No. 89-	40611	40612	40613	40614	40615	40616
Customer No.	Comp A 1A-4A	Comp A 1B-4B	Comp A 1C-4C	Comp B 1A-3A	Comp B 1B-3B	Comp B 1C-3C
Sample Identity						
Results in: mg/kg dry weight unless noted						
Oil and Grease	25	<6.0	<6.0	15	44	41
Total Recoverable Petroleum Hydrocarbons	56	<6.0	<6.0	<6.0	<6.0	<6.0
Total Solids, %	85	84	87	87	88	86

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: J. Halperin

Date: 7/14/89





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### LABORATORY REPORT

FOR

SOUTHERN PACIFIC RAILROAD-HIGH STREET

S7/850/001.1

Job No.: 850.001		RE: SP8030				
Sample Date: 5/24, 25/89		P.O. No.:				
Date Received: 5/25/89		Sampled By: E & E, Inc.				
Sample Type: Solid		Delivered By: Federal Express				
E & E Lab. No. 89-	40617	40618	40619	40620	40621	40622
Customer No.	Comp C 1A-C4A	Comp C 1B-C4B	Comp C 1C-C4C	D-1A	D-1B	D-1C
Sample Identity						
Results In: mg/kg dry weight unless noted						
Oil and Grease	2,800	220	26	1,300	<6.0	<6.0
Total Recoverable Petroleum Hydrocarbons	1,600	150	<6.0	900	<6.0	<6.0
Total Solids, %	88	84	86	81	86	90

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: *M. Hahn*  
 Date: 7/14/89



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**LABORATORY REPORT**

FOR

SOUTHERN PACIFIC RAILROAD-HIGH STREET

S7/850/001.2

Job No.: 850,001		RE: SP8030				
Sample Date: 5/24/89		P.O. No.:				
Date Received: 5/25/89		Sampled By: E & E, Inc.				
Sample Type: Solid		Delivered By: Federal Express				
E & E Lab. No. 89-	40623					
Customer No.	D-1D					
Sample Identity						
Results In: mg/kg dry weight unless noted						
Oil and Grease	<6.0					
Total Recoverable Petroleum Hydrocarbons	<6.0					
Total Solids, %	86					

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: J. Halm  
Date: 7/14/89

QUALITY CONTROL FOR PRECISION  
 RESULTS OF ANALYSIS OF REPLICATE  
 ANALYSES OF SOLID SAMPLES

S7/850/001.3

Parameter	E & E Laboratory No. 89-	(mg/kg)		Relative Percent Difference (RPD)
		Original Analysis	Replicate Analysis	
Total Solids, %	40619	86	86	0
Oil and Grease	40623	<6.0	<6.0	-
Total Recoverable Petroleum Hydrocarbons	40623	<6.0	<6.0	-

QUALITY CONTROL FOR ACCURACY: PERCENT RECOVERY  
FOR SPIKED SOLID SAMPLES

S7/850/001.4

Parameter	E & E Laboratory No. 89-	Original Value	Amount Added	Amount Determined	Percent Recovery
		(mg/kg)			
Oil and Grease	40623	<6.0	980	1,000	102
Total Recoverable Petroleum Hydrocarbons	40623	<6.0	980	1,100	112



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### LABORATORY REPORT

FOR

SOUTHERN PACIFIC RAILROAD-HIGH STREET

S7/850/001.5

Job No.:	850,001			RE:	SP8030		
Sample Date:	5/25/89			P.O. No.:			
Date Received:	5/25/89			Sampled By:	E & E, Inc.		
Sample Type:	Solid			Delivered By:	Federal Express		
E & E Lab. No. 89-	40611	40612	40613	40614	40615	40616	
Customer No.	Comp A 1A-4A	Comp A 1B-4B	Comp A 1C-4C	Comp B 1A-3A	Comp B 1B-3B	Comp B 1C-3C	
Sample Identity							
Results in: mg/kg dry weight unless noted							
Antimony	<7.00	<7.10	<6.90	<6.90	<6.80	<7.00	
Arsenic	3.52	3.44	2.99	8.79	3.11	3.24	
Barium	186	108	187	347	182	116	
Beryllium	<0.240	<0.240	<0.230	<0.230	<0.230	<0.230	
Cadmium	1.08	1.17	1.01	1.70	1.15	1.16	
Chromium	43.8	61.7	49.8	152	79.1	62.9	
Hexavalent Chromium	<0.012	<0.012	0.028	0.022	0.084	0.099	
Cobalt	13.8	12.1	12.0	6.00	11.8	12.1	
Copper	30.4	20.7	40.1	103	100	30.2	
Lead	32.9	17.1	12.3	538	18.9	8.40	
Mercury*	<0.12	<0.12	<0.11	<0.11	<0.11	<0.12	
Molybdenum	<3.50	<3.60	<3.40	<3.40	<3.40	<3.50	
Nickel	52.5	130	95.5	35.0	115	92.2	
Selenium	<0.60	<0.60	<0.60	<0.60	<0.60	<0.60	
Silver	<1.20	<1.20	<1.10	<1.10	<1.10	<1.20	
Thallium	<0.60	<0.60	<0.60	<0.60	<0.60	<0.60	
Vanadium	36.0	40.8	34.2	29.6	41.8	32.4	
Zinc	42.2	53.1	62.9	1,500	105	52.2	

\*analyzed one day beyond recommended holding time

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: J. Halim  
 Date: 7/14/89



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**LABORATORY REPORT**

FOR

SOUTHERN PACIFIC RAILROAD-HIGH STREET

S7/850/001.6

Job No.:	850.001						RE:	SP8030					
Sample Date:	5/24, 25/89						P.O. No.:						
Date Received:	5/25/89						Sampled By:	E & E, Inc.					
Sample Type:	Solid						Delivered By:	Federal Express					
E & E Lab. No. 89-	40617	40618	40619	40620	40621	40622							
Customer No.	Comp C 1A-C4A	Comp C 1B-C4B	Comp C 1C-C4C	D-1A	D-1B	D-1C							
Sample Identity													
Results In: mg/kg dry weight unless noted													
Antimony	<6.80	<7.10	<7.00	<7.40	<7.00	<6.70							
Arsenic	3.39	4.70	2.71	9.54	1.86	2.27							
Barium	63.5	367	141	1,500	217	137							
Beryllium	<0.230	<0.240	<0.230	<0.250	<0.230	<0.220							
Cadmium	0.580	3.14	2.00	3.06	1.08	1.20							
Chromium	20.4	71.8	100	50.1	68.7	103							
Hexavalent Chromium	<0.011	0.012	<0.012	0.034	<0.012	<0.011							
Cobalt	3.64	33.0	17.0	8.89	14.9	17.0							
Copper	19.2	56.5	37.1	112	32.8	56.0							
Lead	60.8	126	26.2	198	13.0	12.8							
Mercury*	<0.11	<0.12	<0.12	<0.12	<0.12	<0.11							
Molybdenum	<3.40	<3.60	<3.50	<3.70	<3.50	<3.30							
Nickel	29.6	161	140	53.8	94.5	141							
Selenium	<0.60	<0.60	<0.60	<0.60	<0.60	<0.60							
Silver	<1.10	<1.20	<1.20	<1.20	<1.20	<1.10							
Thallium	<0.60	<0.60	<0.60	<0.60	<0.60	<0.60							
Vanadium	10.2	41.9	56.4	27.5	44.2	44.7							
Zinc	78.9	362	83.4	538	50.1	71.6							

\*analyzed one day beyond recommended holding time

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: J. Halim  
Date: 7/14/89



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### LABORATORY REPORT

FOR

SOUTHERN PACIFIC RAILROAD-HIGH STREET

S7/805/001.7

Job No.:	850,001	RE:	SP8030
Sample Date:	5/24/89	P.O. No.:	
Date Received:	5/25/89	Sampled By:	E & E, Inc.
Sample Type:	Solid	Delivered By:	Federal Express
E & E Lab. No. 89-	40623		
Customer No.	D-1D		
Sample Identity			
Results in: mg/kg dry weight unless noted			
Antimony	<7.00		
Arsenic	3.01		
Barium	153		
Beryllium	<0.230		
Cadmium	1.63		
Chromium	87.6		
Hexavalent Chromium	0.013		
Cobalt	20.7		
Copper	108		
Lead	15.3		
Mercury*	<0.12		
Molybdenum	<3.50		
Nickel	186		
Selenium	<0.60		
Silver	<1.20		
Thallium	<0.60		
Vanadium	54.4		
Zinc	137		

\*Analyzed one day beyond recommended holding time

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: E. Halim  
Date: 7/14/89

QUALITY CONTROL FOR PRECISION  
 RESULTS OF ANALYSIS OF REPLICATE  
 ANALYSES OF SOLID SAMPLES

S7/850/001.8

Parameter	E & E Laboratory No. 89- 40623	(mg/kg)		Relative Percent Difference (RPD)
		Original Analysis	Replicate Analysis	
Antimony		<7.00	<7.00	--
Arsenic		3.01	2.15	33
Barium		153	116	28
Beryllium		<0.230	<0.230	--
Cadmium		1.63	1.52	7
Chromium		87.6	67.2	26
Hexavalent Chromium		0.013	0.012	8
Cobalt		20.7	17.6	16
Copper		108	38.4	95
Lead		15.3	12.2	22
Molybdenum		<3.50	<3.50	--
Nickel		186	160	15
Selenium		<0.60	<0.60	--
Silver		<1.20	<1.20	--
Thallium		<0.60	<0.60	--
Vanadium		54.4	44.6	20
Zinc		137	67.3	68



QUALITY CONTROL FOR ACCURACY: PERCENT RECOVERY  
FOR SPIKED SOLID SAMPLES

S7/850/001.9

Parameter	E & E Laboratory No. 89- 40623	Original Value	Amount Added	Amount Determined	Percent Recovery
		(mg/kg)			
Antimony		<7.00	57.0	28.1	49
Arsenic		3.01	4.56	6.70	81
Barium		153	228	415	115
Beryllium		<0.230	5.70	6.07	106
Cadmium		1.63	5.70	7.38	101
Chromium		87.6	22.8	105	76
Hexavalent Chromium		0.013	0.12	0.024	9
Cobalt		20.7	57.0	81.6	107
Copper		38.4	29.1	85.7	162
Lead		15.3	57.0	77.9	110
Molybdenum		<3.50	57.0	75.5	132
Nickel		157	57.0	226	121
Selenium		<0.60	1.14	1.19	104
Silver		<1.20	5.70	5.94	104
Thallium		<0.60	5.70	6.70	118
Vanadium		54.4	57.0	109	96
Zinc		137	57.0	176	68

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
PURGEABLE ORGANIC COMPOUNDS AND HSL COMPOUNDS BY GC/MS

(all results in ug/kg dry weight)

S7/850/001.10

Compound	E & E Lab. No. 89-	40611	40612	40613	40614	40615
	Sample Identity	Comp A 1A-4A	Comp A 1B-4B	Comp A 1C-4C	Comp B 1A-3A	Comp B 1B-3B
chloromethane		<12	<12	<11	<11	<11
bromomethane		<12	<12	<11	<11	<11
vinyl chloride		<12	<12	<11	<11	<11
chloroethane		<12	<12	<11	<11	<11
methylene chloride		<6*	<6*	<6	<6*	<6*
acetone		84	55	38	30	58
carbon disulfide		<6	<6	<6	<6	<6
1,1-dichloroethene		<6	<6	<6	<6	<6
1,1-dichloroethane		<6	<6	<6	<6	<6
trans-1,2-dichloroethene		<6	<6	<6	<6	<6
chloroform		<6	<6	<6	<6	<6
1,2-dichloroethane		<6	<6	<6	<6	<6
2-butanone		<12	<12	<11	<11	<11
1,1,1-trichloroethane		<6	<6	<6	<6	<6
carbon tetrachloride		<6	<6	<6	<6	<6
vinyl acetate		<12	<12	<11	<11	<11
bromodichloromethane		<6	<6	<6	<6	<6
1,2-dichloropropane		<6	<6	<6	<6	<6
trans-1,3-dichloropropene		<6	<6	<6	<6	<6
trichloroethene		<6	<6	<6	<6	<6
dibromochloromethane		<6	<6	<6	<6	<6
1,1,2-trichloroethane		<6	<6	<6	<6	<6
benzene		<6	<6	<6	<6	<6
cis-1,3-dichloropropene		<6	<6	<6	<6	<6
2-chloroethylvinyl ether		<12	<12	<11	<11	<11
bromoform		<6	<6	<6	<6	<6
4-methyl-2-pentanone		<12	<12	<11	<11	<11
2-hexanone		<12	<12	<11	<11	<11
tetrachloroethene		<6	<6	<6	<6	<6
1,1,2,2-tetrachloroethane		<6	<6	<6	<6	<6
toluene		<6	<6	<6	<6	<6
chlorobenzene		<6	<6	<6	<6	<6
ethyl benzene		<6	<6	<6	<6	<6
styrene		<6	<6	<6	<6	<6
total xylenes		<6	<6	<6	<6	<6

\*Compound present below measurable detection limit.

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
PURGEABLE ORGANIC COMPOUNDS AND HSL COMPOUNDS BY GC/MS

(all results in ug/kg dry weight)

S7/850/001.11

Compound	E & E Lab. No. 89-	40616	40617	40618	40619	40620
	Sample Identity	Comp B 1C-3C	Comp C 1A-C4A	Comp C 1B-C4B	Comp C 1C-C4C	D-1A
chloromethane		<12	<11	<12	<12	<12
bronomethane		<12	<11	<12	<12	<12
vinyl chloride		<12	<11	<12	<12	<12
chloroethane		<12	<11	<12	<12	<12
methylene chloride		<6*	6	7	<6	<6*
acetone		56	73	83	23	120
carbon disulfide		<6	<6	<6	<6	<6
1,1-dichloroethene		<6	<6	<6	<6	<6
1,1-dichloroethane		<6	<6	<6	<6	<6
trans-1,2-dichloroethene		<6	<6	<6*	<6	<6
chloroform		<6	<6	<6	<6	<6
1,2-dichloroethane		<6	<6	<6	<6	<6
2-butanone		<12	<11	<12	<12	<12
1,1,1-trichloroethane		<6	<6	<6	<6	<6
carbon tetrachloride		<6	<6	<6	<6	<6
vinyl acetate		<12	<11	<12	<12	<12
bromodichloromethane		<6	<6	<6	<6	<6
1,2-dichloropropane		<6	<6	<6	<6	<6
trans-1,3-dichloropropene		<6	<6	<6	<6	<6
trichloroethene		<6	<6	<6	<6	<6
ditromochloromethane		<6	<6	<6	<6	<6
1,1,2-trichloroethane		<6	<6	<6	<6	<6
benzene		<6	<6	<6	<6	<6
cis-1,3-dichloropropene		<6	<6	<6	<6	<6
2-chloroethylvinyl ether		<12	<11	<12	<12	<12
bromoform		<6	<6	<6	<6	<6
4-methyl-2-pentanone		<12	<11	<12	<12	<12
2-hexanone		<12	<11	<12	<12	<12
tetrachloroethene		<6	<6	<6	<6	<6
1,1,2,2-tetrachloroethane		<6	<6	<6	<6	<6
to uene		<6	<6	<6*	<6	<6
chlorobenzene		<6	<6	<6	<6	<6
ethylbenzene		<6	<6	<6*	<6	<6
styrene		<6	<6	<6	<6	<6
total xylenes		<6	<6	18	<6	<6

\*Compound present below measurable detection limit.

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
PURGEABLE ORGANIC COMPOUNDS AND HSL COMPOUNDS BY GC/MS

(all results in ug/kg dry weight)

S7/850/001.12

Compound	E & E Lab. No. 89-	40621	40622	40623	Method Blank 1	Method Blank 2
	Sample Identity	D-1B	D-1C	D-1D		
chloromethane		<12	<11	<12	<10	<10
bromomethane		<12	<11	<12	<10	<10
vinyl chloride		<12	<11	<12	<10	<10
chloroethane		<12	<11	<12	<10	<10
methylene chloride		6	<6*	<6*	<5*	<5*
acetone		74	21	15	11	10
carbon disulfide		<6	<6	<6	<5	<5
1,1-dichloroethene		<6	<6	<6	<5	<5
1,1-dichloroethane		<6	<6	<6	<5	<5
trans-1,2-dichloroethene		<6	<6	<6	<5	<5
chloroform		<6	<6	<6	<5	<5
1,2-dichloroethane		<6	<6	<6	<5	<5
2-butanone		<12	<11	<12	<10	<10
1,1,1-trichloroethane		<6	<6	<6	<5	<5
carbon tetrachloride		<6	<6	<6	<5	<5
vinyl acetate		<12	<11	<12	<10	<10
bromodichloromethane		<6	<6	<6	<5	<5
1,2-dichloropropane		<6	<6	<6	<5	<5
trans-1,3-dichloropropene		<6	<6	<6	<5	<5
trichloroethene		<6	<6	<6	<5	<5
dibromochloromethane		<6	<6	<6	<5	<5
1,1,2-trichloroethane		<6	<6	<6	<5	<5
benzene		<6	<6	<6	<5	<5
cis-1,3-dichloropropene		<6	<6	<6	<5	<5
2-chloroethylvinyl ether		<12	<11	<12	<10	<10
bromoform		<6	<6	<6	<5	<5
4-methyl-2-pentanone		<12	<11	<12	<10	<10
2-hexanone		<12	<11	<12	<10	<10
tetrachloroethene		<6	<6	<6	<5	<5
1,1,2,2-tetrachloroethane		<6	<6	<6	<5	<5
toluene		<6	<6	<6	<5	<5
chlorobenzene		<6	<6	<6	<5	<5
ethylbenzene		<6	<6	<6	<5	<5
styrene		<6	<6	<6	<5	<5
total xylenes		<6	<6	<6	<5	<5

\*Compound present below measurable detection limit.

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
PURGEABLE ORGANIC COMPOUNDS AND HSL COMPOUNDS BY GC/MS

(all results in ug/kg)

S7/850/001.13

Compound	E & E Lab. No. 89-	Method Blank 3				
	Sample Identity					
chloromethane		<10				
bromomethane		<10				
vinyl chloride		<10				
chloroethane		<10				
methylene chloride		<5*				
acetone		10				
carbon disulfide		<5				
1,1-dichloroethene		<5				
1,1-dichloroethane		<5				
trans-1,2-dichloroethene		<5				
chloroform		<5				
1,2-dichloroethane		<5				
2-butanone		<10				
1,1,1-trichloroethane		<5				
carbon tetrachloride		<5				
vinyl acetate		<10				
bromodichloromethane		<5				
1,2-dichloropropane		<5				
trans-1,3-dichloropropene		<5				
trichloroethene		<5				
dibromochloromethane		<5				
1,1,2-trichloroethane		<5				
benzene		<5				
cis-1,3-dichloropropene		<5				
2-chloroethylvinyl ether		<10				
bromoform		<5				
4-methyl-2-pentanone		<10				
2-hexanone		<10				
tetrachloroethene		<5				
1,1,2,2-tetrachloroethane		<5				
toluene		<5				
chlorobenzene		<5				
ethylbenzene		<5				
styrene		<5				
total xylenes		<5				

\*Compound present below measurable detection limit.

QUALITY CONTROL FOR ACCURACY: PERCENT  
RECOVERY OF SURROGATE SPIKES

S7/850/001.14

Compound	E & E Laboratory No. 89-	Amount Added	Amount Determined	Percent Recovery
		(ngs)		
1,2-dichloroethane-D4	40611	250	288	115
	40612	250	221	88
	40613	250	301	120
	40614	250	243	97
	40615	250	248	99
	40616	250	251	100
	40617	250	240	96
	40618	250	274	110
toluene-D8	40611	250	272	109
	40612	250	209	84
	40613	250	276	110
	40614	250	260	104
	40615	250	265	106
	40616	250	265	106
	40617	250	263	105
	40618	250	283	113
4-bromofluorobenzene	40611	250	258	103
	40612	250	216	86
	40613	250	283	113
	40614	250	254	102
	40615	250	260	104
	40616	250	253	101
	40617	250	243	97
	40618	250	238	95

These recoveries are acceptable to E & E, Inc. guidelines.

QUALITY CONTROL FOR ACCURACY: PERCENT  
RECOVERY OF SURROGATE SPIKES  
(Cont.)

S7/850/001.15

Compound	E & E Laboratory No. 89-	Amount Added	Amount Determined	Percent Recovery
		(ngs)		
1,2-dichloroethane-D4	40619	250	258	103
	40620	250	246	98
	40621	250	256	102
	40622	250	255	102
	40623	250	244	98
	Method Blank1	250	261	104
	Method Blank2	250	235	94
	Method Blank3	250	241	96
toluene-D8	40619	250	230	92
	40620	250	266	106
	40621	250	254	102
	40622	250	257	103
	40623	250	246	98
	Method Blank1	250	254	102
	Method Blank2	250	249	100
	Method Blank3	250	248	99
4-bromofluorobenzene	40619	250	223	89
	40620	250	252	101
	40621	250	251	100
	40622	250	278	111
	40623	250	245	98
	Method Blank1	250	255	102
	Method Blank2	250	248	99
	Method Blank3	250	245	98

These recoveries are acceptable to E & E, Inc. guidelines.

QUALITY CONTROL FOR ACCURACY AND PRECISION:  
 PERCENT RECOVERY AND RELATIVE PERCENT DIFFERENCE (RPD)  
 OF SOIL MATRIX SPIKE (MS) AND MATRIX SPIKE DUPLICATE (MSD)  
 (Sample #40617)

S7/850/001.16

Compound	(ngs)				Percent Recovery		RPD
	Original Result	Amount Added	Amount Determined		MS	MSD	
			MS	MSD			
1,1-Dichloroethene	<6	250	216	244	86	98	13
Trichloroethene	<6	250	199	231	80	92	14
Chlorobenzene	<6	250	270	303	108	121	11
Toluene	<6	250	251	341	100	136	31
Benzene	<6	250	257	304	103	122	17

These recoveries and RPDs are within E & E, Inc. limits.



ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in mg/kg dry weight)

S7/850/001.17

Compound	E & E Lab. No. 89-	40611	40612	40613	40614	40615
	Sample Identity	Comp A 1A-4A	Comp A 1B-4B	Comp A 1C-4C	Comp B 1A-3A	Comp B 1B-3B
PCB-1242		<u>0.15</u>	<0.06*	<0.06*	<0.06	<0.06
PCB-1254		<u>0.19</u>	<0.06*	<u>0.087+</u>	<0.06	<0.06
PCB-1221		<0.06	<0.06	<0.06	<0.06	<0.06
PCB-1232		<0.06	<0.06	<0.06	<0.06	<0.06
PCB-1248		<0.06	<0.06	<0.06	<0.06	<0.06
PCB-1260		<0.06	<0.06	<0.06	<0.06	<0.06
PCB-1016		<0.06	<0.06	<0.06	<0.06	<0.06

\*compound present below measurable detection limit  
+sample identification best fit as PCB-1254

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in mg/kg dry weight)

S7/850/001.18

Compound	E & E Lab. No. 89-	40616	40617	40618	40619	40620
	Sample identity	Comp B 1C-3C	Comp C 1A-C4A	Comp C 1B-C4B	Comp C 1C-C4C	D-1A
PCB-1242		<0.06	<u>2.3</u>	<u>0.77</u>	<u>0.30</u>	<0.06
PCB-1254		<0.06	<0.60	<0.12	<0.06	<0.06
PCB-1221		<0.06	<0.60	<0.12	<0.06	<0.06
PCB-1232		<0.06	<0.60	<0.12	<0.06	<0.06
PCB-1248		<0.06	<0.60	<0.12	<0.06	<0.06
PCB-1260		<0.06	<u>14</u>	<u>2.6</u>	<u>0.88</u>	<u>0.09+</u>
PCB-1016		<0.06	<0.60	<0.12	<0.06	<0.06

+sample identification best fit as PCB-1260

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in mg/kg dry weight)

S7/850/001.19

Compound	E & E Lab. No. 89-	40621	40622	40623	Method Blank	
	Sample Identity	D-1B	D-1C	D-1D		
PCB-1242		<0.06	<0.06	<0.06	<0.05	
PCB-1254		<0.06	<0.06	<0.06	<0.05	
PCB-1221		<0.06	<0.06	<0.06	<0.05	
PCB-1232		<0.06	<0.06	<0.06	<0.05	
PCB-1248		<0.06	<0.06	<0.06	<0.05	
PCB-1260		<0.06	<0.06	<0.06	<0.05	
PCB-1016		<0.06	<0.06	<0.06	<0.05	

QUALITY CONTROL FOR ACCURACY: PERCENT RECOVERY  
FOR SPIKED SOIL SAMPLES

S7/850/001.20

Parameter	E & E Laboratory No. 89- 40623	Original Value	Amount Added	Amount Determined	Percent Recovery
		(mg/kg)			
PCB 1242		<0.06	10	11	110

10542

850.001

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 Interventional Specialists in the Environment

**CHAIN-OF-CUSTODY RECORD**

Project No.: SP-8030		Project Name: Southern Pacific - High Street		Project Manager: Colin May				
Sampler: (Signatures) Robert H. Siskel		Field Team Leader: Bob Entekoball		Ship Via: Federal Express				
STATION NUMBER	DATE	TIME	SAMPLE TYPE	SAMPLE INFORMATION		STATION LOCATION	NUMBER OF CONTAINERS	REMARKS
				COMP	GRA			
A-1A	5/23		X		low	3.5 to 4.0	1	Composite with A-2A, A-3A, A-4A
A-1B	5/23		X			6.0 to 6.5	1	Composite with A-2B, A-3B, A-4B
A-1C	5/23		X			11.0 to 11.5	1	Composite with A-2C, A-3C, A-4C
A-1D	5/23		X		40024	16.0 to 16.5	1	Archive Sample
A-1E	5/23		X		2.5	21.0 to 21.5	1	Archive Sample
A-1F	5/23		X		40026	26.0 to 26.5	1	Archive Sample
A-2A	5/22		X			1.0 to 1.5	1	Composite with A-1A, A-3A, A-4A
A-2B	5/22		X			6.0 to 6.5	1	Composite with A-1B, A-3B, A-4B
A-2C	5/22		X			11.0 to 11.5	1	Composite with A-1C, A-3C, A-4C
A-3A	5/23		X			2.0 to 2.5	1	Composite with A-1A, A-2A, A-4A
A-3B	5/23		X			6.0 to 6.5	1	Composite with A-1B, A-2B, A-4B
A-3C	5/23		X			11.0 to 11.5	1	Composite with A-1C, A-2C, A-4C
Relinquished By: (Signature) B. Entekoball		Date/Time: 5/14/89 11:08	Received By: (Signature) FED. EXP	Date/Time: 5/25-89	Received For Laboratory By: (Signature) Rich March			
Relinquished By: (Signature)		Date/Time:	Received By: (Signature)	Date/Time:	Received For Laboratory By: (Signature)			
Relinquished By: (Signature)		Date/Time:	Received By: (Signature)	Date/Time:	Received For Laboratory By: (Signature)			

BU/Airbill Number: 4518273281 Date: 5/24/89

234055

\*See CONCENTRATION RANGE on back of form.

**Environment, inc.**

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 Environmental Specialists in the Environment

**CHAIN-OF-CUSTODY RECORD**

Project No.: SP-8030		Project Name: Southern Pacific High Street		Project Manager: Colin May		Field Team Leader: Bob Enkeboell		STATION LOCATION	NUMBER OF COM. CONTAINERS	REMARKS
Samplers: (Signatures) <i>Alex H. Siskel</i>		SAMPLE INFORMATION		EXPECTED COMPOUNDS (Concentration)*		DATE				
STATION NUMBER	DATE	TIME	COM. TYPE	CONC.	EXPCD.	DATE	TIME			
A-4A	5/22		X		Low			2.0 to 2.5	1	Composite with A-1A, A-2A, A-3A
A-4B	5/22		X					6.0 to 6.5	1	Composite with A-1B, A-2B, A-3B
A-4C	5/22		X					11.0 to 11.5	1	Composite with A-1C, A-2C, A-3C
B-1A	5/22		X					1.0 to 1.5	1	Composite with B-2A, B-3A
B-1B	5/22		X					6.0 to 6.5	1	Composite with B-2B, B-3B
B-1C	5/22		X					11.0 to 11.5	1	Composite with B-2C, B-3C
B-2A	5/23		X					2.0 to 2.5	1	Composite with B-1A, B-3A
B-2B	5/23		X					6.0 to 6.5	1	Composite with B-1B, B-3B
B-2C	5/23		X					11.0 to 11.5	1	Composite with B-1C, B-3C
B-2D	5/23		X		40627			16.0 to 16.5	1	Archive Sample
B-2E	5/23		X		128			21.0 to 21.5	1	Archive Sample
B-2F	5/23		X		40629			26.0 to 26.5	1	Archive Sample
RECEIVED BY: (Signature) <i>B. Enkeboell</i> Date/Time: 5/24/89 1608      Received By: (Signature) <i>FED. EXP.</i> Date/Time:      Ship Via: <i>Federal Express</i> RECEIVED BY: (Signature)      Date/Time:      Received By: (Signature)      Date/Time:      PL/Airbill Number: <i>4518273281</i> RECEIVED BY: (Signature)      Date/Time: 5/25/89      Received For Laboratory By: (Signature) <i>Pat March</i> Date/Time:      Date: 5/24/89										

\*See CONCENTRATION RANGE on back of form.

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 Regional Specialists in the Environment

**CHAIN-OF-CUSTODY RECORD**

Page 3 of 4

STATION NUMBER	DATE	TIME	SAMPLE TYPE		STATION LOCATION	NUMBER OF CONTAINERS	REMARKS
			GRAB	24			
<p>Project No.: <u>SP-8030</u> Project Name: <u>Southern Pacific High Street</u> Project Manager: <u>Calin May</u>                      Samplers: (Signatures) <u>Robert H. Enkelbelle</u> <u>Tom Cochran</u> Field Team Leader: <u>Bob Enkelbelle</u></p>							
B-3A	5/22		X		2.0 to 3.5	1	Composite with B-1A, B-2A, B-2B, B-2C, B-2D
B-3B	5/22		X		6.0 to 6.5	1	Composite with B-1B, B-2B, B-2C, B-2D
B-3C	5/22		X		11.0 to 11.5	1	Composite with B-1C, B-2C, B-2D
C-1A	5/24		X			1	Composite with C-2A, C-3A, C-4A
C-1B	5/24		X			1	Composite with C-2B, C-3B, C-4B
C-1C	5/24		X			1	Composite with C-2C, C-3C, C-4C
C-2A	5/24		X			1	Composite with C-1A, C-3A, C-4A
C-2B	5/24		X			1	Composite with C-1B, C-3B, C-4B
C-2C	5/24		X			1	Composite with C-1C, C-3C, C-4C
C-2D	5/24		X			1	Archive Sample
C-2E	5/24		X			1	Archive Sample
C-2F	5/24		X			1	Archive Sample
<p>40630 31 40632</p>							
Relinquished By: (Signature) <u>B. Enkelbelle</u>		Date/Time: <u>5/24/89</u>	Received By: (Signature) <u>FED EXP.</u>	Date/Time: <u>5/25/89</u>	Relinquished By: (Signature)	Date/Time:	Received By: (Signature)
Relinquished By: (Signature)		Date/Time:	Received By: (Signature)	Date/Time:	Relinquished By: (Signature)	Date/Time:	Received For Laboratory By: (Signature)
Relinquished By: (Signature) <u>FED. EXP.</u>		Date/Time: <u>5/25/89</u>	Received For Laboratory By: (Signature) <u>Bob Mark</u>	Date/Time:	Relinquished By: (Signature)	Date/Time:	Received For Laboratory By: (Signature)
Ship Via: <u>Federal Express</u>						BL/Airbill Number: <u>4518273281</u>	Date: <u>5/24/89</u>

Distribution: Original Accompanies Shipment; Copy to Coordinator Field Files  
 \*See CONCENTRATION RANGE on back of form.

234055

**Environment, inc.**

100 BOX D, BUFFALO, N.Y., 14225. TEL. 716-632-4481

National Specialists in the Environment

**CHAIN-OF-CUSTODY RECORD**

STATION NUMBER	DATE	TIME	SAMPLE TYPE		STATION LOCATION	NUMBER OF CONTAINERS	REMARKS
			COMP	AIR			
Project Name: <u>Southern Pacific High Street</u> Project Manager: <u>Colin S. Moy</u> Field Team Leader: <u>Bob Enkeball</u>							
C-2A	5/24		X			1	Composite with C-2A, C-1A, C-4A Composite with C-1B, C-2B, C-4B Composite with C-1C, C-2C, C-4C Composite with C-1A, C-2A, C-3A Composite with C-1B, C-2B, C-3B Composite with C-1C, C-2C, C-3C analyt individually analyt individually analyt individually analyt individually
C-3B	5/24		X			1	
C-3C	5/24		X			1	
C-4A	5/24		X			1	
C-4B	5/24		X			1	
C-4C	5/24		X			1	
D-1A	5/24		X			1	
D-1B	5/24		X			1	
D-1C	5/24		X			1	
D-1D	5/24		X			1	
SAMPLE INFORMATION EXPECTED COMPOUNDS (Concentration)* Cold 40620 21 22 40623							
Relinquished By: (Signature)		Date/Time		Received By: (Signature)		Date/Time	
<u>B. Enkeball</u>		<u>5/24/89, 1608</u>		<u>FED. EXP.</u>		<u>5/24/89</u>	
Relinquished By: (Signature)		Date/Time		Received By: (Signature)		Date/Time	
Relinquished By: (Signature)		Date/Time		Received For Laboratory By: (Signature)		Date/Time	
<u>FED. EXP.</u>		<u>5/24/89</u>		<u>Bob Enkeball</u>		<u>5/24/89</u>	
Shipped Via: <u>Federal Express</u>				BL/Airbill Number: <u>4518273281</u>			

\*See CONCENTRATION RANGE on back of form.





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International Specialists in the Environment

### LABORATORY REPORT

FOR

SOUTHERN PACIFIC RAILROAD - HIGH STREET

S8/850/002

Job No.: 850.002		RE: SP8030				
Sample Date: 5/26/89		P.O. No.:				
Date Received: 5/30/89		Sampled By: E & E, Inc.				
Sample Type: Water		Delivered By: Federal Express				
E & E Lab. No. 89-	40759	40760	40761	40762	40763	
Customer No.	A-1	A-2	A-3	B-2	C-2	
Sample Identity						
Results In: mg/L unless noted						
Oil and Grease	2.3	1.4	1.6	2.8	2.4	
Total Recoverable Petroleum Hydrocarbons	<1.0	1.4	<1.0	1.5	<1.0	

Analytical References:

"Methods for the Chemical Analysis of Water and Wastes," EPA-600/4-79-020, March 1983.

Supervising Analyst: M. Halpern

Date: 7/14/89



**ecology and environment, inc.**

International Specialists in the Environment

### LABORATORY REPORT

FOR

SOUTHERN PACIFIC RAILROAD - HIGH STREET

S8/850/002.1

Job No.: 850.002		RE: SP8030				
Sample Date: 5/26/89		P.O. No.:				
Date Received: 5/30/89		Sampled By: E & E, Inc.				
Sample Type: Water		Delivered By: Federal Express				
E & E Lab. No. 89-	40759	40760	40761	40762	40763	
Customer No.	A-1	A-2	A-3	B-2	C-2	
Sample Identity						
Results in: mg/L unless noted						
Barium	0.086	0.072	<0.010	0.037	0.027	
Hexavalent Chromium	0.063	<0.010	<0.010	<0.010	<0.010	
Cobalt	<0.010	<0.010	<0.010	<0.010	<0.010	
Molybdenum	<0.010	<0.010	<0.010	<0.010	<0.010	
Vanadium	<0.010	<0.010	<0.010	<0.010	<0.010	

Analytical References:

"Methods for the Chemical Analysis of Water and Wastes," EPA-600/4-79-020, March 1983.

Supervising Analyst: J. Hansen

Date: 7/14/89

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF WATER ANALYSES  
FOR PRIORITY POLLUTANT METALS

(all results in mg/L)

S8/850/002.2

	E & E Lab. No. 89-	40759	40760	40761	40762	40763
	Sample Identity	A-1	A-2	A-3	B-2	C-2
Antimony		<0.060	<0.060	<0.060	<0.060	<0.060
Arsenic		<0.010	<0.005	<0.005	<0.005	<0.010
Beryllium		<0.002	<0.002	<0.002	<0.002	<0.002
Cadmium		<0.005	<0.005	<0.005	<0.005	<0.005
Chromium		<0.010	<0.010	<0.010	<0.010	<0.010
Copper		0.013	0.019	<0.010	<0.010	0.014
Lead		0.030	0.024	<0.005	0.006	0.019
Mercury		0.0012	<0.0002	<0.0002	<0.0002	<0.0002
Nickel		0.018	<0.015	<0.015	<0.015	<0.015
Selenium		<0.005	<0.005	<0.005	<0.005	<0.005
Silver		<0.010	<0.010	<0.010	<0.010	<0.010
Thallium		<0.005	<0.005	<0.005	<0.005	<0.005
Zinc		0.085	0.054	<0.010	0.045	0.110

QUALITY CONTROL FOR PRECISION  
RESULTS OF ANALYSIS OF REPLICATE  
ANALYSES OF WATER SAMPLES

S8/850/002.3

Parameter	E & E Laboratory No. 89- 40763	(mg/L)		Relative Percent Difference (RPD)
		Original Analysis	Replicate Analysis	
Antimony		<0.060	<0.060	--
Arsenic		<0.010	<0.010	--
Barium		0.027	0.027	0
Beryllium		<0.002	<0.002	--
Cadmium		<0.005	<0.005	--
Chromium		<0.010	<0.010	--
Hexavalent Chromium		<0.010	<0.010	--
Cobalt		<0.010	<0.010	--
Copper		0.014	0.010	33
Lead		0.019	0.016	17
Molybdenum		<0.010	<0.010	--
Nickel		<0.015	<0.015	--
Selenium		<0.005	<0.005	--
Silver		<0.010	<0.010	--
Thallium		<0.005	<0.005	--
Vanadium		<0.010	<0.010	--
Zinc		0.110	0.061	57

QUALITY CONTROL FOR ACCURACY: PERCENT RECOVERY  
FOR SPIKED WATER SAMPLES

S8/850/002.4

Parameter	E & E Laboratory No. 89- 40763	Original Value	Amount Added	Amount Determined	Percent Recovery
		(mg/L)			
Antimony		<0.060	0.50	0.436	87
Arsenic		<0.010	0.04	0.039	98
Barium		0.027	2.0	1.85	91
Beryllium		<0.002	0.05	0.044	88
Cadmium		<0.005	0.05	0.043	86
Chromium		<0.010	0.20	0.171	86
Hexavalent Chromium		<0.010	1.0	0.104	104
Cobalt		<0.010	0.50	0.448	90
Copper		0.014	0.25	0.221	83
Lead		0.019	0.04	0.048	73
Molybdenum		<0.010	0.50	0.454	91
Nickel		<0.015	0.50	0.439	88
Selenium		<0.005	0.01	0.006	60
Silver		<0.010	0.05	0.040	80
Thallium		<0.005	0.05	0.039	78
Vanadium		<0.010	0.50	0.450	90
Zinc		0.110	0.50	0.498	78

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF WATER ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in ug/L)

S8/850/002.5

Compound	E & E Lab. No. 89-	40759	40760	40761	40762	40763
	Sample Identity	A-1	A-2	A-3	B-2	C-2
PCB-1242		<0.50	<0.50	<0.50	<0.50	<0.50
PCB-1254		<0.50	<0.50	<0.50	<0.50	<0.50
PCB-1221		<0.50	<0.50	<0.50	<0.50	<0.50
PCB-1232		<0.50	<0.50	<0.50	<0.50	<0.50
PCB-1248		<0.50	<0.50	<0.50	<0.50	<0.50
PCB-1260		<0.50	<0.50	<0.50	<0.50	<u>1.0</u>
PCB-1016		<0.50	<0.50	<0.50	<0.50	<0.50

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF WATER ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in ug/L)

S8/850/002.6

	E & E Lab. No. 89-	Method Blank				
Compound	Sample Identity					
PCB-1242		<0.50				
PCB-1254		<0.50				
PCB-1221		<0.50				
PCB-1232		<0.50				
PCB-1248		<0.50				
PCB-1260		<0.50				
PCB-1016		<0.50				

QUALITY CONTROL FOR ACCURACY: PERCENT RECOVERY  
FOR SPIKED WATER SAMPLES

S8/850/002.7

Parameter	E & E Laboratory No. 89- 40759	Original Value	Amount Added	Amount Determined	Percent Recovery
		(ug/L)			
PCB 1242		<0.50	50	19	38



RESULTS OF WATER ANALYSIS FOR PRIORITY POLLUTANT  
PURGEABLE ORGANIC COMPOUNDS AND HSL COMPOUNDS BY GC/MS

(all results in ug/L)

58/850/002.8

Compound	E & E Lab. No. 89-	40759	40760	40761	40762	40763
	Sample Identity	A-1	A-2	A-3	B-2	C-2
chloromethane		<10	<10	<10	<10	<10
bromomethane		<10	<10	<10	<10	<10
vinyl chloride		<10	<10	<10	<10	<10
chloroethane		<10	<10	<10	<10	<10
methylene chloride		20	18	17	<5*	<5*
acetone		14	<10*	<10*	31	55
carbon disulfide		<5	<5	<5	<5	<5
1,1-dichloroethene		<5	<5	<5	<5	<5
1,1-dichloroethane		<5	<5	<5	<5	<5
trans-1,2-dichloroethene		<5	<5	<5	<5	<5
chloroform		<5	<5	<5	<5	<5
1,2-dichloroethane		<5	<5	<5	<5	<5
2-butanone		<10	<10	<10	<10	<10
1,1,1-trichloroethane		<5	<5	<5	<5	<5
carbon tetrachloride		<5	<5	<5	<5	<5
vinyl acetate		<10	<10	<10	<10	<10
bromodichloromethane		<5	<5	<5	<5	<5
1,2-dichloropropane		<5	<5	<5	<5	<5
trans-1,3-dichloropropene		<5	<5	<5	<5	<5
trichloroethene		<5	<5	<5	<5	<5
dibromochloromethane		<5	<5	<5	<5	<5
1,1,2-trichloroethane		<5	<5	<5	<5	<5
benzene		<5	<5	<5	<5	<5
cis-1,3-dichloropropene		<5	<5	<5	<5	<5
2-chloroethylvinyl ether		<10	<10	<10	<10	<10
bromoform		<5	<5	<5	<5	<5
4-methyl-2-pentanone		<10	<10	<10	<10	<10
2-hexanone		<10	<10	<10	<10	<10
tetrachloroethene		<5	<5	<5	<5	<5
1,1,2,2-tetrachloroethane		<5	<5	<5	<5	<5
toluene		<5	<5	<5	<5	<5
chlorobenzene		<5	<5	<5	<5	<5
ethylbenzene		<5	<5	<5	<5	<5
styrene		<5	<5	<5	<5	<5
total xylenes		<5	<5	<5	<5	<5

\*Compound present below measurable detection limit.

RESULTS OF WATER ANALYSIS FOR PRIORITY POLLUTANT  
PURGEABLE ORGANIC COMPOUNDS AND HSL COMPOUNDS BY GC/MS

(all results in ug/L)

S8/850/002.9

	E & E Lab. No. 89-	Method Blank 1	Method Blank 2			
Compound	Sample Identity					
chloromethane		<10	<10			
bromomethane		<10	<10			
vinyl chloride		<10	<10			
chloroethane		<10	<10			
methylene chloride		<5*	7			
acetone		11	<10			
carbon disulfide		<5	<5			
1,1-dichloroethene		<5	<5			
1,1-dichloroethane		<5	<5			
trans-1,2-dichloroethene		<5	<5			
chloroform		<5	<5			
1,2-dichloroethane		<5	<5			
2-butanone		<10	<10			
1,1,1-trichloroethane		<5	<5			
carbon tetrachloride		<5	<5			
vinyl acetate		<10	<10			
bromodichloromethane		<5	<5			
1,2-dichloropropane		<5	<5			
trans-1,3-dichloropropene		<5	<5			
trichloroethene		<5	<5			
dibromochloromethane		<5	<5			
1,1,2-trichloroethane		<5	<5			
benzene		<5	<5			
cis-1,3-dichloropropene		<5	<5			
2-chloroethylvinyl ether		<10	<10			
bromoform		<5	<5			
4-methyl-2-pentanone		<10	<10			
2-hexanone		<10	<10			
tetrachloroethene		<5	<5			
1,1,2,2-tetrachloroethane		<5	<5			
toluene		<5	<5			
chlorobenzene		<5	<5			
ethylbenzene		<5	<5			
styrene		<5	<5			
total xylenes		<5	<5			

\*Compound present below measurable detection limit.

QUALITY CONTROL FOR ACCURACY: PERCENT  
RECOVERY OF SURROGATE SPIKES

S8/850/002.10

Compound	E & E Laboratory No. 89-	Amount Added	Amount Determined	Percent Recovery
		(ug/L)		
1,2-dichloroethane-D4	40759	50	45	90
	40760	50	43	86
	40761	50	42	84
	40762	50	50	100
	40763	50	48	96
	Method Blank 1	50	47	94
	Method Blank 2	50	48	96
toluene-D8	40759	50	54	108
	40760	50	50	100
	40761	50	50	100
	40762	50	54	108
	40763	50	48	96
	Method Blank 1	50	51	102
	Method Blank 2	50	51	102
4-bromofluorobenzene	40759	50	44	88
	40760	50	44	88
	40761	50	44	88
	40762	50	53	106
	40763	50	49	98
	Method Blank 1	50	50	100
	Method Blank 2	50	50	100

These recoveries are acceptable to E & E, Inc. guidelines.

QUALITY CONTROL FOR PRECISION  
RESULTS OF ANALYSIS OF REPLICATE  
ANALYSES OF WATER SAMPLES

S8/850/002.11

Compound	E & E Lab. No. 89- 40761	ug/L		Relative Percent Difference (RPD)
		Original Analysis	Replicate Analysis	
chloromethane		<10	<10	--
bromomethane		<10	<10	--
vinyl chloride		<10	<10	--
chloroethane		<10	<10	--
methylene chloride		17	20	16
acetone		<10*	<10*	--
carbon disulfide		<5	<5	--
1,1-dichloroethene		<5	<5	--
1,1-dichloroethane		<5	<5	--
trans-1,2-dichloroethene		<5	<5	--
chloroform		<5	<5	--
1,2-dichloroethane		<5	<5	--
2-butanone		<10	<10	--
1,1,1-trichloroethane		<5	<5	--
carbon tetrachloride		<5	<5	--
vinyl acetate		<10	<10	--
bromodichloromethane		<5	<5	--
1,2-dichloropropane		<5	<5	--
trans-1,3-dichloropropene		<5	<5	--
trichloroethene		<5	<5	--
dibromochloromethane		<5	<5	--
1,1,2-trichloroethane		<5	<5	--
benzene		<5	<5	--
cis-1,3-dichloropropene		<5	<5	--
2-chloroethylvinyl ether		<10	<10	--
bromoform		<5	<5	--
4-methyl-2-pentanone		<10	<10	--
2-hexanone		<10	<10	--
tetrachloroethene		<5	<5	--
1,1,2,2-tetrachloroethane		<5	<5	--
toluene		<5	<5	--
chlorobenzene		<5	<5	--
ethylbenzene		<5	<5	--
styrene		<5	<5	--
total xylenes		<5	<5	--







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**LABORATORY REPORT**

FOR

SOUTHERN PACIFIC TRANSPORT

S10/927/001

Job No.: 927.001		RE: SP 8050				
Sample Date: 7/25, 28/89		P.O. No.:				
Date Received: 7/29/89		Sampled By: E & E, Inc.				
Sample Type: Solid		Delivered By: Federal Express				
E & E Lab. No. 89--	45954	45955	45956	45957	45958	45959
Customer No.	B-1A	B-2A	B-3A	C-1A	C-2A	C-3A
Sample Identity						
Results in: mg/kg dry weight unless noted						
Total Solids, %	82	89	89	82	83	94
Total Recoverable Petroleum Hydrocarbons	NR	NR	NR	<6.1	140	92

NR: Analysis not requested

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: *Henry Adams*  
Date: 8/17/89



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**LABORATORY REPORT**

FOR

SOUTHERN PACIFIC TRANSPORT

S10/927/001.1

Job No.: 927.001		RE: SP 8050			
Sample Date: 7/28/89		P.O. No.:			
Date Received: 7/29/89		Sampled By: E & E, Inc.			
Sample Type: Solid		Delivered By: Federal Express			
E & E Lab. No. 89-	45960	45961			
Customer No.	C-4A	C-21A			
Sample Identity					
Results in: mg/kg dry weight unless noted					
Total Solids, %	78	86			
Total Recoverable Petroleum Hydrocarbons	13	900			

Analytical References:

Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: *[Signature]*

Date: 8/17/89





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## LABORATORY REPORT

FOR

SOUTHERN PACIFIC TRANSPORT

S10/927/001.2

Job No.: 927.001		RE: SP 8050				
Sample Date: 7/25, 28/89		P.O. No.:				
Date Received: 7/29/89		Sampled By: E & E, Inc.				
Sample Type: Solid		Delivered By: Federal Express				
E & E Lab. No. 89-	45954	45955	45956	45957	45958	45959
Customer No.	B-1A	B-2A	B-3A	C-1A	C-2A	C-3A
Sample Identity						
Results in: mg/kg dry weight unless noted						
Barium	552	232	186	888	399	119
Cadmium	NR	NR	NR	1.39	4.48	0.780
Copper	235	127	51.7	NR	NR	NR
Lead	434	478	92.7	599	365	76.0
Nickel	148	33.9	18.3	93.4	103	66.2
Zinc	2,420	3,800	200	NR	NR	NR

NR: Analysis not requested

### Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: *[Signature]*

Date: 8/17/89



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### LABORATORY REPORT

FOR

SOUTHERN PACIFIC TRANSPORT

S10/927/001.3

Job No.: 927,001			RE: SP 8050			
Sample Date: 7/28/89			P.O. No.:			
Date Received: 7/29/89			Sampled By: E & E, Inc.			
Sample Type: Solid			Delivered By: Federal Express			
E & E Lab. No. 89-	45960	45961				
Customer No.	C-4A	C-21A				
Sample Identity						
Results in: mg/kg dry weight unless noted						
Barium	1,220	391				
Cadmium	2.26	6.52				
Lead	240	1,290				
Nickel	231	101				

Analytical References:

"Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," SW-846, Second Edition, U.S. EPA, 1982.

Supervising Analyst: *Gaughan*

Date: 8/17/89

QUALITY CONTROL FOR PRECISION  
 RESULTS OF ANALYSIS OF REPLICATE  
 ANALYSES OF SOLID SAMPLES

S10/927/001.4

Parameter	E & E Laboratory No. 89-	(mg/kg)		Relative Percent Difference (RPD)
		Original Analysis	Replicate Analysis	
Total Solids, %	45960	78	77	1.3
	45958	83	82	1.2
Barium	45961	391	207	62
Cadmium	45961	6.52	3.85	51
Lead	45961	1,290	686	61
Nickel	45961	101	103	2.0

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in mg/kg dry weight)

S10/927/001.5

Compound	E & E Lab. No. 89--	45957	45958	45959	45960	45961
	Sample Identity	C-1A	C-2A	C-3A	C-4A	C-21A
PCB-1242		<0.024	<0.024	<0.021	<0.205	<0.023
PCB-1254		<0.024	<0.024	<0.021	<u>3.6</u>	<0.023
PCB-1221		<0.024	<0.024	<0.021	<0.205	<0.023
PCB-1232		<0.024	<0.024	<0.021	<0.205	<0.023
PCB-1248		<0.024	<0.024	<0.021	<0.205	<0.023
PCB-1260		<u>0.024</u>	<u>0.22</u>	<0.021	<0.205	<u>0.53</u>
PCB-1016		<0.024	<0.024	<0.021	<0.205	<0.023

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF SOIL ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in mg/kg)

S10/927/001.6

Compound	E & E Lab. No. 89--	Method Blank				
	Sample Identity					
PCB-1242		<0.02				
PCB-1254		<0.02				
PCB-1221		<0.02				
PCB-1232		<0.02				
PCB-1248		<0.02				
PCB-1260		<0.02				
PCB-1016		<0.02				

ECOLOGY AND ENVIRONMENT'S, INC.  
ANALYTICAL SERVICES CENTER

RESULTS OF WATER ANALYSIS FOR PRIORITY POLLUTANT  
POLYCHLORINATED BIPHENYLS

(all results in ug/L)

S10/927/001.7

Compound	E & E Lab. No. 89-	45962	45963	Method Blank		
	Sample Identity	C-2	C-21			
PCB-1242		<0.50	<0.50	<0.50		
PCB-1254		<0.50	<0.50	<0.50		
PCB-1221		<0.50	<0.50	<0.50		
PCB-1232		<0.50	<0.50	<0.50		
PCB-1248		<0.50	<0.50	<0.50		
PCB-1260		<u>0.61</u>	<u>0.78</u>	<0.50		
PCB-1016		<0.50	<0.50	<0.50		

QUALITY CONTROL FOR ACCURACY AND PRECISION:  
 PERCENT RECOVERY AND RELATIVE PERCENT DIFFERENCE (RPD)  
 OF SOIL MATRIX SPIKE (MS) AND MATRIX SPIKE DUPLICATE (MSD)  
 (Sample #45960)

S10/927/001.8

Compound	Original Result	Amount Added	Amount Determined		Percent Recovery		RPD
			MS	MSD	MS	MSD	
PCB-1242	<0.205	1.7	1.8	1.5	106	88	18

11004

927



**ecology and environment, inc.**  
195 SUGG ROAD, P.O. BOX D, BUFFALO, N.Y., 14225, TEL. 716-632-4491  
International Specialists in the Environment

A-27

**CHAIN-OF-CUSTODY RECORD**

Project No.: SP8650		Project Name: SPTCO. - High Street		Project Manager: C. MOY		Field Team Leader: C. MOY		STATION LOCATION		NUMBER OF CON-TAINERS	REMARKS
Samplers: (Signatures) Colts - Moy		SAMPLE INFORMATION		EXPECTED COMPOUNDS (Concentration)*		DATE		TIME			
STATION NUMBER	DATE	TIME	SAMPLE TYPE	EXPECTED COMPOUNDS (Concentration)*		DATE	TIME	DATE	TIME	RECEIVED BY: (Signature)	DATE
B-1A	7/25/91	1155	X	Low		7/25/91	1155	7/25/91	1155	X	TPH (Non-metals) PCBS
B-2A	7/28/91	1015	X			7/28/91	1015	7/28/91	1015	X	Note: pushed on TAT - 1015
B-3A	7/28/91	1025	X			7/28/91	1025	7/28/91	1025	X	on ver. - 1015
C-1A	7/28/91	1035	X			7/28/91	1035	7/28/91	1035	X	COOLERS REL
C-2A	7/28/91	1100	X			7/28/91	1100	7/28/91	1100	X	3
C-3A	7/28/91	1053	X			7/28/91	1053	7/28/91	1053	X	PCBS
C-4A	7/28/91	1045	X			7/28/91	1045	7/28/91	1045	X	
C-21A	7/28/91	1105	X			7/28/91	1105	7/28/91	1105	X	
C-2	7/28/91	1106	X	Muta; 1 ppb PCBS		7/28/91	1106	7/28/91	1106	X	
C-21	7/28/91	1130	X			7/28/91	1130	7/28/91	1130	X	

Ship Via: FEDEX - Sat, Delivery  
 BL/Airbill Number: 9870772284  
 Date: 7/28/91

Received By: (Signature) Received By: (Signature)  
 Relinquished By: (Signature) Relinquished By: (Signature)  
 Received For Laboratory By: (Signature) Received For Laboratory By: (Signature)

Distribution: Original Accompanies Shipment; Copy to Coordinator Field Files  
 \*See CONCENTRATION RANGE on back of form.





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**LABORATORY REPORT**

FOR

**SOUTHERN PACIFIC TRANSPORT**

JOB NO:	927.002	P.O. NO:
SAMPLE DATE:	07/28/89	RE: SP8050
DATE RECEIVED:	07/29/89	SAMPLED BY: E & E, Inc.
SAMPLE TYPE:	Solid	DELIVERED BY: Federal Express

E & E LAB		
NO. 89-	46418	46419

CLIENT	B-2A	C-3A
SAMPLE ID-	45955	45959

Results in: mg/L unless noted

Results of Chemical  
Analysis of Extracts  
from California WET  
Test

Lead	6.06	2.16
Zinc	268	NR

NR: Analysis not requested

Analytical Reference: California Administrative Code, Title 22, Chapter 30, "Minimum Standards for Management of Hazardous and Extremely Hazardous Waste", 1984.

Supervising Analyst: Gary DeLuca  
Date: 8/18/89



**TO:** Madulla Logan, Alameda County Hazardous Materials Unit  
Richard Hlett, Regional Water Quality Control Board  
John Bacon, owner of 750 High Street, Oakland

**FROM:** M. Papineau on behalf of John Bacon *MP*

**DATE:** November 8, 1993

**SUBJECT:** 750 High Street, Oakland, California (Economy Lumber)

This note is to confirm our meeting appointment for Tuesday November 9, 1993 at 2 P.M. at the Alameda County Health Department office at 80 Swan Way in Oakland. The purpose of the meeting is to review the status of environmental investigation and remediation at 750 High Street and consideration case closure.

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Engineering  
& Testing®  
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Services

7000 Marina Boulevard  
4th Floor  
Brisbane, CA 94005  
415-742-9900  
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A  GROUP COMPANY



7000 MARINA BOULEVARD, 4TH FLOOR, BRISBANE, CA 94005

FAX 415-742-1033

TO: MS. MADULLA LOGAN DATE: 11/8/93

COMPANY: ALAMEDA COUNTY HAZARDOUS MATERIALS

FAX NUMBER: (510) 568-3706 569 4757

FROM: M. PAPINEAU

CONFIDENTIAL:  YES  NO

TOTAL PAGES SENT (INCLUDING THIS PAGE): 4

REGARDING: Meeting 11/9/93 at Alameda County  
80 Swan Way 2 P.M.

IF THERE ARE ANY PROBLEMS WITH RECEIPT OF THIS TELEFAX,  
PLEASE CALL 415-742-9900

CHRONOLOGY

750 High Street, Oakland, California

*due to presence, RB, lead  
total oil  
benzene*

<u>Date</u>	<u>Index</u>	<u>Event/Activity</u>
5/26/89		Ground water sampling event #1 by Ecology & Environment (E & E) on behalf of Southern Pacific Transportation Company (SPTC) ✓
7/28/89		Ground-water sampling event #2 by E & E for SPTC ✓
12/4/89		Ground-water sampling event #3 by E & E for SPTC ✓
4/30/90	1	First day of soil excavation by IT Corporation ✓
5/8/90	1	Second day of soil excavation ✓
6/12/90	1	Off-haul of 161 cubic yards of petroleum-affected soil from 744 High Street
6/25/90		Ground-water sampling event #4 by E & E for SPTC ✓
7/16/90	2	Report of Post-Excavation Sampling by E & E for SPTC ✓
9/6/90	3	Ground-water sampling event # 5 by E & E for SPTC
10/4/90	3	Report of Ground-Water Sampling Event #5 by E & E
2/5/92	4	Letter to Mr. Bacon from Lester Feldman ✓
4/8/92	5	Ground-water sampling event #8 for well C-5 only by Earth Metrics Incorporated for Mr. John Bacon
6/16/92	6	Level Two Environmental Site Assessment by Certified Engineering & Testing Company, Inc. (CERTIFIED) for John Bacon ✓
5/18/93	7	Revised summary letter and request for review of same by Mr. Britt Johnson (Alameda County Hazardous Materials Unit by CERTIFIED on behalf of John Bacon

CONTINUED ON PAGE TWO

CHRONOLOGY  
Page 2 of 2

<u>Date</u>	<u>Index</u>	<u>Event/Activity</u>
8/19/93	8	Ground-water sampling event #7 by CERTIFIED for Mr. John Bacon
9/13/93	8	Report of Ground-Water Sampling Event #8 by CERTIFIED ✓
END		