915022/915/37

PHASE I REMEDIATION FINAL REPORT

Pyramid Co. - Ernst Steel Site Buffalo, New York

prepared for: Whiteman Osterman & Hanna



PHASE I REMEDIATION FINAL REPORT

Pyramid Co. - Ernst Steel Site Buffalo, New York

prepared for:

Whiteman Osterman & Hanna

TABLE OF CONTENTS

		Page
1.0	INTRODUCTION	1
2.0	TEST PIT INVESTIGATION	2
	2.1 SUMMARY	2
	2.2 DECONTAMINATION	5
	2.3 RESULTS	5
3.0	WASTE CHARACTERIZATION SAMPLING	8
	3.1 METHODOLOGY	9
	3.2 DECONTAMINATION	10
	3.3 SAMPLE HANDLING	11
	3.4 ANALYTICAL RESULTS	11
	3.4.1 Metals	11
	3.4.2 Volatiles	13
	3.4.3 Base Neutral/Acid Extractables (BNAs)	13
	3.4.4 PCBs	13
	3.4.5 Pesticides	14
	3.4.6 Quality Control	14
4.0	MONITORING WELL INSTALLATIONS	15
	4.1 METHODOLOGY	16
	4.2 DECONTAMINATION	18
	4.3 WELL DEVELOPMENT	18
	4.4 PHYSICAL SOILS TESTING	20
5.0	SITE GEOLOGY AND HYDROGEOLOGY	21
	5.1 SITE GEOLOGY	21
	5.2 SITE HYDROGEOLOGY	22
6.0	100-FOOT GRID SAMPLING PROGRAM	24
	6.1 METHODOLOGY	25
	6.2 DECONTAMINATION	26

TABLE OF CONTENTS

		<u>Page</u>
	6.3 SAMPLE HANDLING	26
	6.4 ANALYTICAL RESULTS	27
	6.4.1 Metals	27
	6.4.2 Volatiles	28
	6.4.3 BNAs	28
	6.4.4 Pesticides and PCBs	29
	6.4.5 Quality Control	29
7.0	SITE SPECIFIC PARAMETER SELECTION	31
3.0	WORK PLAN SAMPLING PROGRAMS	32
	8.1 MONITORING WELL SAMPLING	32
	8.2 SURFACE SOIL SAMPLING	33
	8.3 SURFACE WATER AND SEDIMENT SAMPLES	34
	8.4 DUPLICATE SAMPLING	36
	8.5 ANALYTICAL RESULTS	36
	8.5.1 TOX and Toluene	36
	8.5.2 Lead and Chromium	37
	8.5.3 Arochlor 1254	39
9.0	HEALTH AND SAFETY	41
	9.1 TEST PIT INVESTIGATION	41
	AND WASTE CHARACTERIZATION SAMPLING	41
	9.2 MONITORING WELL INSTALLATION AND WELL DEVELOPMENT	42
	9.3 WORK PLAN SAMPLING PROGRAMS	43
10.0	UTILITY INVESTIGATION	44.
11.0	CONCLUSIONS	47

TABLE OF CONTENTS

			Page
12.0	FILL	HANDLING PROPOSAL	49
	12.1	AREA PREPARATION	50
	12.2	VOLUME OF FILL	51
	12.3	AVAILABLE DISPOSAL AREA	52
	12.4	EXCAVATION AND HAULING PROCEDURES	53
	12.5	SURFACE WATER CONTROL	54
	12.6	DISPOSAL AREA CONSTRUCTION	55
	12.7	SUMMARY	57

LIST OF FIGURES

			Page
FIGURE	1	TEST PIT LOCATIONS	3
FIGURE	2	FILL TYPE STRATIGRAPHY	5
FIGURE	3	AREAS USED TO QUANTIFY FILL VOLUMES	6
FIGURE	4	FILL CHARACTERIZATION SAMPLING LOCATIONS	10
FIGURE	5	MONITORING WELL LOCATIONS	15
FIGURE	6	100-FOOT GRID SAMPLING LOCATIONS	24
FIGURE	7	TOTAL LEAD CONCENTRATIONS IN ENTIRE FILL STRATA	24
FIGURE	8	EP TOXICITY LEAD CONCENTRATIONS IN ENTIRE FILL STRATA	25
FIGURE	9	TOTAL LEAD CONCENTRATIONS IN NATIVE MATERIAL	25
FIGURE	10	AREA OF FILL PRESENCE	25
FIGURE	11	SURFACE SOIL SAMPLING LOCATIONS	34
FIGURE	12	UNDERGROUND UTILITY INVESTIGATION	44
FIGURE	13	PROPOSED FILL PLACEMENT PLAN	49

LIST OF TABLES

			Following Page
TABLE	1	HNU SCANNING RESULTS TEST PIT SPOILS	3
TABLE	2	TEST PIT INVESTIGATION STRATIGRAPHIC SUMMARY	5
TABLE	3	FILL QUANTIFICATION SUMMARY	6
TABLE	4	SAMPLE COLLECTION SUMMARY	10
TABLE	5	COMPOUNDS DETECTED WASTE CHARACTERIZATION STUDY	11
TABLE	6	HNU SCANNING RESULTS SPLIT SPOON SAMPLES	16
TABLE	7	WELL INSTALLATION DETAIL SUMMARY	17
TABLE	8	GROUNDWATER ELEVATIONS	23
TABLE	9	COMPOUNDS DETECTED 100 FOOT GRID SAMPLING STUDY	27
TABLE	10	PHASE II SITE SPECIFIC PARAMETER RESULTS	36

1.0 INTRODUCTION

A Phase II Site Investigation (SI) was conducted at the Pyramid Company-Ernst Steel Site (Site) between June and September 1988. The program, as described in the report entitled "Work Plan - Phase II Site Investigation" (Work Plan) dated May 1988, was developed by Pyramid and Conestoga-Rovers & Associates (CRA) and was approved by the New York State Department of Environmental Conservation (NYSDEC). This investigation included waste characterization and delineation of the fill areas (excluding the waste piles which were handled separately pursuant to the Interim Remedial Measures Plan), installation of groundwater monitoring wells and sampling of groundwater, soil, surface water and sediment.

The details of these activities are presented in the following sections of this report. The main purpose of this report is to summarize and evaluate the data collected, provide an assessment of the site conditions and a preliminary evaluation of remedial action alternatives.

2.0 TEST PIT INVESTIGATION

The initial activity in the SI was the Test

Pit Investigation. This investigation was conducted with the following purposes in mind:

- i) To delineate the areal and vertical extent of fill material across the Site.
- ii) To determine the type (or types) of fill present at the Site.
- iii) To allow for access to the fill types for waste characterization sampling.
- iv) To gather the data necessary to estimate the total volume of fill present at the Site.

2.1 SUMMARY

Prior to commencing the excavation of test pits, CRA established a grid across the Site to assure that the test pits would be distributed as evenly as conditions would allow.

CRA then contracted SLC, Inc. of Lockport, NY to provide a backhoe and operator to excavate the test pits. The investigation began on June 9, 1988. On June 10, 1988, work was temporarily suspended pending approval of the Order on Consent from the NYSDEC. Following approval, work resumed and was completed on July 6, 1988. NYSDEC site representatives were present for the completion of the Test Pit Investigation. A total of 32 test pit locations were excavated (see Figure 1).

Excavated materials were placed on the ground surface next to the excavation. Upon completion of the test pit, the excavated materials were replaced in the reverse order from which they had been removed.

Test pits were excavated to the depth at which native soil was observed. Each test pit was logged according to stratigraphy encountered and depth to native soil. A representative sample of the materials encountered in each excavation was collected and retained for geologic record.

At the completion of excavation of each test pit, the spoils pile was scanned with an HNU photoionization unit to determine the presence of organic compounds. Table 1 presents the results of the HNU scanning.

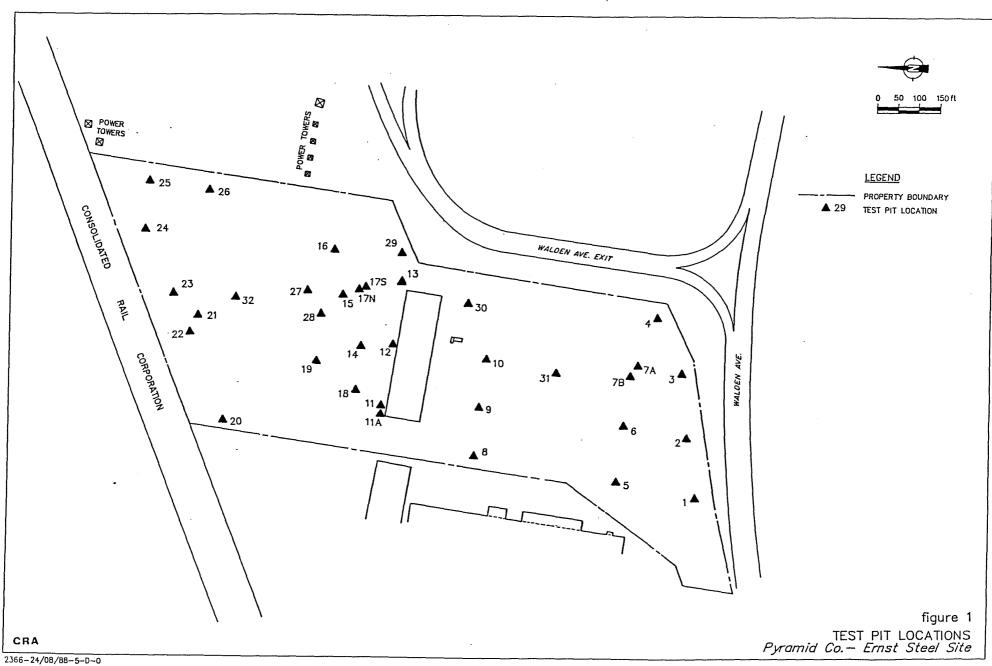


Table 1 HNU Scanning Results Test Pit Spoils Pyramid Co.-Ernst Steel Site

Air Monitoring Results

Test Pit Number	$\frac{\texttt{Background}}{(\texttt{ppm})}$	Results of Scan* (ppm)
1	0.4	0.4
2	0.4	0.4
1 2 3 4	0.4	0.6
4	0.4	0.5
5	0.4	0.5
6	0.4	0.6-0.7
7a	0.4	0.4
7b	0.4	0.8-0.9
8	0.4	0.8
9	0.4	0.7-0.8
10	0.4	0.6
11	0.4	0.9
11A	1.0	0.8
12	0.4	0.8-2.0
12A	1.0	0.8
13	0.4	0.9
14	0.4	0.8
15	0.4	0.6-0.8
16	0.4	. 0.6-0.7
17	0.4	0.4-0.6
18	0.4	0.4-0.6
18A	1.0	0.8
19	0.4	0.4-0.5
20	0.4	0.4-0.6
21	0.5	0.5-0.6
22	0.5	0.5-0.6
23	0.5	0.5
24	0.5	0.4-0.5
25	0.5	0.5
26	0.5	0.6
27	1.0	0.8-1.0
28	1.0	1.0-1.2
29	1.0	1.0
30	1.0	0.2
31	1.0	0.6
32	1.0	0.6-1.0

^{*} including background levels

Based on the results of the HNU scans during the Test Pit Investigation, organic compound contamination at the Site is not anticipated. Virtually all of the readings were within tenths of a ppm of background indicating no significant organic vapors were present. The highest reading recorded was a scan of the spoil pile at Test Pit #12. HNU scan indicated a reading of 0.4-1.6 ppm above background although no odors were detected at this location. Subsequent soil sampling indicated no measurable organics were present. Acetone was observed at 46 ppb but this is within the possible concentration range attributable to laboratory or sampling cross-contamination as exhibited in the blank (12 ppb). Well OW-1 is directly downgradient from Test Pit #12. The groundwater from this well was sampled for the derived Site Specific Parameters. No parameters were measured above Class GA Groundwater Standards and therefore no additional sampling at Test Pit #12 was deemed to be required.

An odor was detected at Test Pit #17. The field notes describe the odor as a sweet, non-distinct smell. However, the HNU did not detect any significant organic vapors.

2.2 DECONTAMINATION

Prior to beginning and at the end of the Test Pit Investigation, the backhoe bucket was cleaned using a water wash.

Wash water from the final cleaning was contained in a 55-gallon drum and staged on-site for later disposal.

2.3 RESULTS

Based on the information gathered during the Test Pit Investigation, the following observations have been made:

1. The Site has been extensively filled. In general, fill is absent or very shallow along the eastern boundary of the Site and deepest in the southern third of the Site.
Table 2 and Figure 2 detail the stratigraphic information collected.

There was one test pit location (TP-11) where fill materials were identified to beyond 6.0 feet in depth. The excavation was not continued beyond 6.0 feet due to the relative uncertainty regarding possible underground

Table 2 Test Pit Investigation Stratigraphic Summary Pyramid/Ernst Steel Site

Test Pit	Fill Depth (ft BGS)	Fill Type	Native Contact (ft BGS)
1	0.0 - 1.0 $1.0 - 3.5$	D B	3.5
2	0.0 - 0.8 $0.8 - 2.0$	D B	2.0
3	0.0 - 1.3	А	1.3
4	N/A	N/A	0
5	0.0 - 0.7 0.7 - 2.5	D A	2.5
6	0.0 - 0.4 $0.4 - 1.1$ $1.1 - 2.1$	D B C	2.1
7A	N/A	N/A	0
7B	0.0 - 0.9 $0.9 - 2.6$	D C	2.6
8	0.0 - 1.2 $1.2 - 1.6$	E A	1.6
9	0.0 - 0.7 0.7 - 1.2	E B	1.2
10	0.0 - 1.0	E	1.0
11	0.0 - 6.0+	D	6.0+
11A	0.0 - 0.9 $0.9 - 1.5$	D A	1.5

A = Black Cindery Fill B = Rust Slag Fill C = Sand Fill

D = Soil Fill

E = Gravel Fill

^{+ =} Did not reach native, greater than 6.0 ft. BGS

Table 2 Test Pit Investigation Stratigraphic Summary Pyramid/Ernst Steel Site

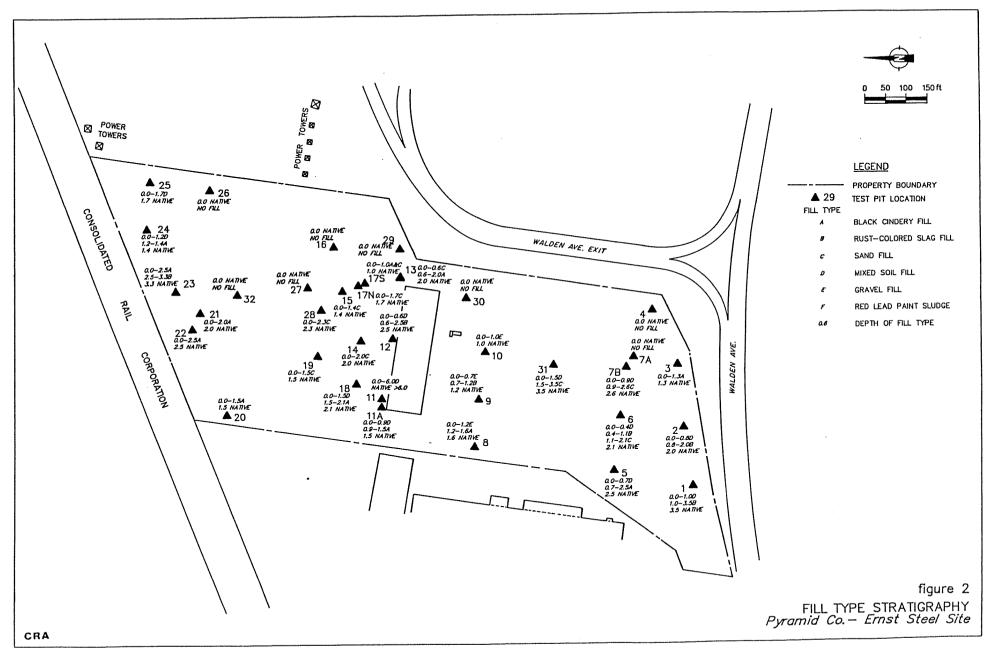
Test Pit	Fill Depth (ft BGS)	Fill Type	Native <u>Contact</u> (ft BGS)
12	0.0 - 0.6 $0.6 - 2.5$	D B	2.5
13	0.0 - 0.6 0.6 - 2.0	C A	2.0
14	0.0 - 2.0	С	2.0
15	0.0 - 1.4	С	1.4
16	N/A		0
17	0.0 - 1.7	С	1.7
18	0.0 - 1.5 $1.5 - 2.1$	D A	2.1
19	0.0 - 1.5	С	1.5
20	0.0 - 1.5	A	1.5
21	0.0 - 2.0	Α	2.0
22	0.0 - 2.5	A	2.5
23	0.0 - 2.5 $2.5 - 3.3$	A B	3.3
24	0.0 - 1.2 $1.2 - 1.4$	D A	1.4
25	0.0 - 1.7	D	1.7
26	N/A		0

A = Black Cindery Fill
B = Rust Slag Fill
C = Sand Fill
D = Soil Fill
E = Gravel Fill

Table 2 Test Pit Investigation Stratigraphic Summary Pyramid/Ernst Steel Site

Test Pit	Fill Depth (ft BGS)	Fill Type	Native Contact (ft BGS)
27	N/A		0
28	0.0 - 2.3	С	2.3
29	N/A		0
30	N/A		0
31	0.0 - 1.5 $1.5 - 3.5$	D C	3.5
32	0.0 - 1.2 $1.2 - 2.6$	C B	2.6

A = Black Cindery Fill
B = Rust Slag Fill
C = Sand Fill
D = Soil Fill
E = Gravel Fill



utilities in the area. Following confirmation that no utilities were present in the area, a confirmation test pit (TP-11A) was excavated a short distance to the west of TP-11 and the fill was present to a depth of 1.5 feet. This indicated that the pocket of fill identified at TP-11 was in fact isolated.

2. Six types of fill were identified visually. They are:

Type A - black cindery fill

Type B - rust-colored slag fill

Type C - sand fill

Type D - mixed soil fill

Type E - gravel fill

Type F - red lead paint sludge.

Based upon the stratigraphic information obtained and presented on Table 2, the volume of fill present at the Site has been estimated. Table 3 summarizes the calculated volume estimates. Figure 3 shows the areas assigned to each test pit to calculate the estimated volume. As can be seen in Table 3, fill types A, B, C and D are present in relatively equal proportions (6,000 to 11,000 cubic yards each). Only 3,000 cubic yards of gravel fill was observed. The total estimated volume of fill present on the Site is on the order of 37,000 cubic yards.

Table 3
Fill Quantification Summary
Pyramid/Ernst Steel Site

Test Pit No.	Test Pit Area	Fill Type A Black Cindery	Fill Type B Rust Slag	Type C Sand Fill	Type D Soil Fill	Type E Gravel Fill
	(ft. ²)	(ft. ³)	(ft• ³)	(ft. ³)	(ft• ³)	(ft. ³)
	(114)	*****		,,,,,,	*****	(11.7)
1	10,313		25,781		10,313	
2	15,375		18,450		12,300	
3	7,350	9,955				
4	No FIII					
5	20,625	37,125			14,438	
6	41,250		28,875	41,250	16,500	
7 a	No Fill					
7ь	11,100			18,870	9,990	
8	35,275	14,110				42,330
9	34,500		17,250			24,150
10	16,500					16,500
`11(1)	8 75	525			788	
12	14,725		27,978		8,835	
13	9,000	12,600	,	5,400	•	
14	9,713	,		19,425		
15	13,313			18,638		
16	No FIII			•		
17N	8,750			14,875		
178	12,900			12,900		
18	35,200	21,120		•	52,800	
19	18,750	•		28,125	·	
20	41,600	62,400		·		
21	9,225	18,450				
22	13,688	34,219				
23	8,100	20,250	6,480			
24	10,925	2,185			13,110	
25	16,150				27,455	
26	No FIII					
27	No FIII					
28	15,300			35,190		
29	No FIII					
30	No FIII					
31	29,250			58,500	43,875	
32*	27,900		39,060	33,480		
Area D	No FIII					
Area E	No FIII					
TOTALS		232,539	163,874	286,653	210,404	82,980

^{*} Only southern portion of Test Pit Area 32 shown on Figure 3 was used for fill estimation. Northern portion was native material.

^{**} Areas 16, 27 and 29 were previously reported as containing fill. Clearing of vegetation allowed redefinition of areas containing fill.

Table 3 Fill Quantification Summary Pyramid/Ernst Steel Site (continued)

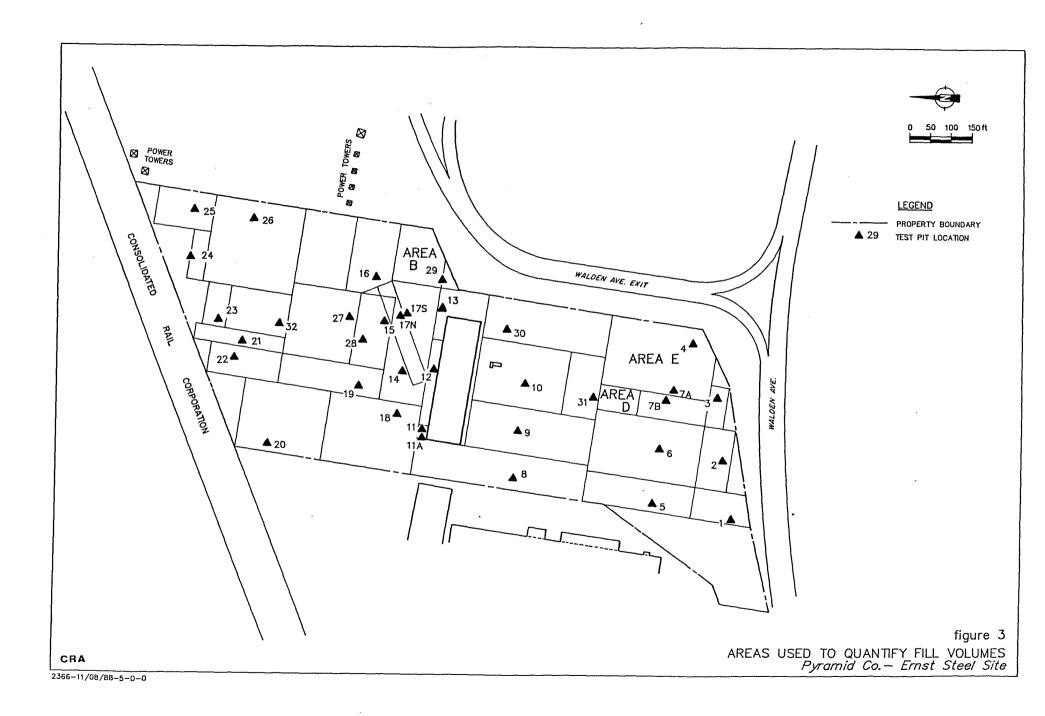
Type A - Black Cindery Fill	232,539 cubic feet	-	9,000 cubic yards
Type B - Rust Slag Fill	163,874 cubic feet	-	6,000 cubic yards
Type C - Sand Fill	286,653 cubic feet	-	11,000 cubic yards
Type D - Soil Fill	210,404 cubic feet	-	8,000 cubic yards
Type E - Gravel Fill	82,980 cubic feet	-	3,000 cubic yards
Total Fill Volume			37,000 cubic yards

Note:

(1)"Type F" fill was identified and sampled on July 6, 1988, however the volume of this fill has not been quantified.

One drum was observed west of Test Pit 28, contents are unknown.

"Type E" fill was not sampled since gravel and crushed slag are materials commonly used in construction activities.



This volume had previously been estimated at 43,000 cubic yards. After clearing the Site of trees and vegetation, it was possible to better define the area containing fill material. As a result of this new data, the volume of fill was recalculated to be 37,000 cubic yards.

The presence of the red lead paint sludge was brought to the attention of CRA personnel by the NYSDEC representative late in the test pit investigation program. Removal of this waste was a concern of the Interim Remedial Measures activities performed previously at the Site. The presence of the paint sludge has been noted and it has been sampled, however, the volume of this waste type has not been quantified.

3.0 WASTE CHARACTERIZATION SAMPLING

After the completion of the Test Pit
Investigation, a waste characterization sampling program was
conducted (July 6, 1988) to determine the site components.
The analytical information from these samples would be used
to develop a list of Site Specific Indicator Parameters
(SSIP) for the subsequent sampling programs. Samples of each
of the identified fill types A, B, C, D and F were collected
for analysis of the Target Compound List (TCL) parameters and
cyanide.

With the concurrence of the NYSDEC site representative, fill type E (gravel fill) was not sampled. Since this fill material is commonly used to construct roads and parking areas it was agreed that it would not contribute to the characterization of this Site.

Waste samples were collected by CRA personnel. NYSDEC site representatives were present throughout the collection of the waste samples. The NYSDEC site representative received splits of all samples collected.

3.1 METHODOLOGY

After reviewing the stratigraphic information collected, the backhoe was used to excavate test pits in areas containing each of the five waste types listed previously for the purpose of sampling.

Test pits were excavated and backfilled in the same manner as during the test pit investigation program.

Samples for waste characterization were collected from test pit excavations in the following manner:

- Using a pre-cleaned garden trowel or stainless steel tablespoon, the wall of the test pit was scraped to remove material which may have come into contact with the backhoe bucket.
- The freshly exposed fill material was then scooped into a pre-cleaned stainless steel bowl using the trowel or spoon until a sufficient volume was collected.
- The sample was mixed thoroughly in the bowl using the sampling tool in order to homogenize the material.
- The homogenized fill was placed into the sample bottles using the spoon or trowel.

Split samples were collected by alternating placement of the fill material between the two sets of sample containers. All glassware was provided by the analytical laboratory.

The sample locations for the preliminary waste characterization program are shown on Figure 4.

Sampling information is summarized in Table 4.

3.2 DECONTAMINATION

Sampling tools were cleaned before each use using the following procedure;

- water wash to remove visible soil
- nitric acid rinse
- water rinse.
- methanol rinse
- hexane rinse
- methanol rinse
- distilled water rinse.
- air dry.

All clean equipment was wrapped in clean aluminum foil for transport between sampling locations. All cleaning fluids were contained and placed into a 55-gallon drum which is staged on-site.

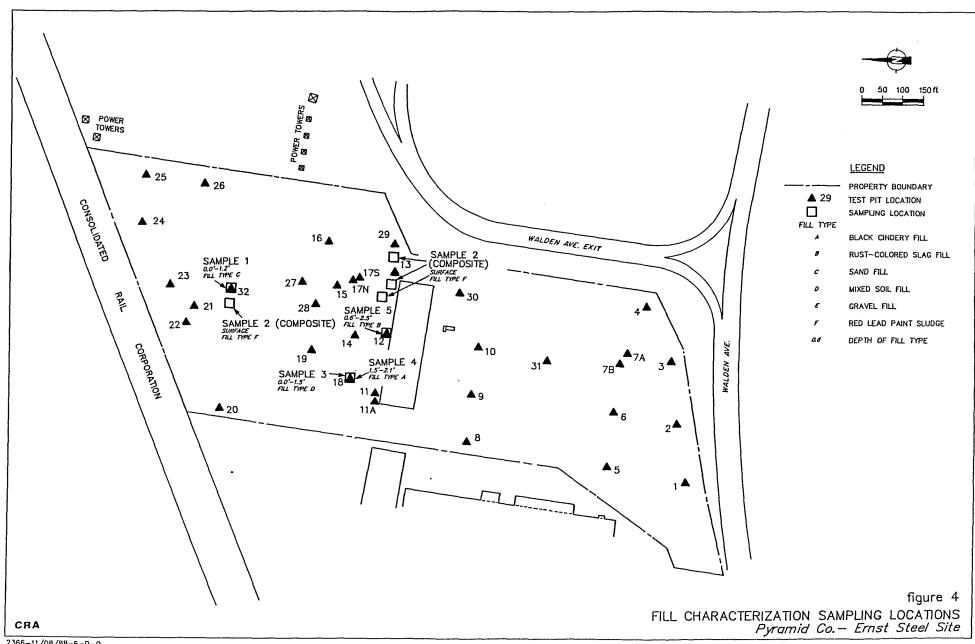


Table 4 Sample Collection Summary Pyramid/Ernst Steel Site

Sample 1

Date: 7/6/88 Time: 1420

Location: Collected in Test Pit 32

Depth: 0 - 1.2 feet Fill Type: C - Sand fill

Sample 2

Date: 7/6/88 Time: 1440

Location: Collected as a composite of surface material observed

west of Test Pit 32, west of Test Pit 13, and between

Test Pit 13 and Test Pit 29.

Depth: Surface

Fill Type: F - Red Lead Paint Sludge

Sample 3

Date: 7/6/88 Time: 1515

Location: Collected in Test Pit 18

Depth: 0 - 1.5 feet Fill Type: D - Soil fill

Sample 4

Date: 7/6/88 Time: 1515

Location: Collected in Test Pit 18

Depth: 1.5 - 2.1 feet

Fill Type: A - Black cindery fill

Sample 5

Date: 7/6/88 Time: 1600

Location: Collected in Test Pit 12

Depth: 0.6 - 2.5 feet Fill Type: B - Rust slag fill

3.3 SAMPLE HANDLING

Immediately after the sample bottles were filled, the jars were labeled and placed into a cooler with ice for storage until shipment to the analytical laboratory. Samples were shipped via Federal Express to EnviroTest Laboratories, Inc. of Newburgh, NY on July 7, 1988. Appropriate Chain of Custody procedures were followed throughout the sampling and shipping process.

3.4 ANALYTICAL RESULTS

The analytical results from the waste characterization study are presented in Appendix A and summarized on Table 5. The following observations have been made.

3.4.1 Metals

The results of the metal analyses presented on Table 5 indicate the following metals above background concentrations:

Table 5
Compounds Detected - Waste Characterization Study
Pyramid/Ernst Steel Site

Metals (ppm)	<u>#1</u>	<u>#2</u>	Sample #3	#4	<u>#5</u>	Background Surface Soil Concentrations (ppm)
Al	3545	7340	10500	3410	5590	4500-100,000
Ва	38.71	546	110	55.43	59.77	10-3000
Be		1.63	0.68			<1-7
Cd		4.29			***	0.41-0.57
Ca	5672	43900	19400	3130	37700	
Cr	9.57	2450	33.98	10.43	15.29	3-1500
Co		29.08	7.50			0.3-50
Cu	184	123	47.39	249	18.74	1-300
Fe	17870	97800	31300	22200	33600	5000-50,000
Pb	208	334	443	376		42-544
Mg	1625	11500	6610		7380	
Mn	769	2810	682	116	3760	7-3000
Hg	0.14	0.12	0.09	0.09		0.01-4.6
Ni	39.78	67.24	25.91	10.87	10.80	<5-200
Κ	559	995				
Ag		1.63				
Na		3188				
Thallium						0.01-2.3
Vanadium	14.52	26.73	32.3	14.67	28 • 28	
Zn	506	6090	139	29.78	26.21	<5-300
Cy an i de		2.04	1.14			
Volatiles (ppb)						
Methylene Chloride	17*	22*	14*	21*	24*	
Acetone		110*	110*		46*	
BNA (ppb)						
Butylbenzylphthal ate	***	580				
bis(2-ethylhexyl)phthalate		3420				
naphthal ene				1260		
2-methyl naphthalene				3330		
dibenzofuran				770		
phen an threne				850		
Pesticides/PCB (ppm)						
Arochior 1254		17				

^{* =} also present in method blank (methylene chloride - 8.9 ppb, acetone - 12 ppb) indicating possible/probable blank contamination.

^{-- =} Not detected above quantifiable limits.

Sample No.	<u>Metal</u>	Concentration (ppm)	Background Concentration (ppm)
l (Sand Fill)	Zinc	506	<5-300
2 (Red Lead Paint Sludge)	Cadmium Chromium Iron Zinc	4.29 2450 97800 6090	0.41-0.57 3-1500 5000-50000 <5-300
5 (Rust Slag Fill)	Manganese	3760	7-3000

Background concentrations are for U.S. soils and were obtained from "Trace Elements in Soils and Plants" by Kabata-Pendias and Pendias (1984) except for lead which was obtained from the Interim Remedial Measures Plan.

The only metal above background concentration in the majority of the samples is zinc with concentrations above background in samples 1 and 2. Of the submitted samples, the red lead paint sludge sample has the largest number of metals above background concentrations. Samples 3 and 4 did not contain any metal concentrations that would be in excess of what could be considered typical background concentrations.

3.4.2 Volatiles

The only volatiles detected, methylene chloride and acetone, were also found in the blank at concentrations of 8.9 ppb and 12 ppb, respectively. These results are indicative of laboratory conditions and not site conditions.

3.4.3 Base Neutral/Acid Extractables (BNAs)

For the BNAs, two phthalate compounds were detected in the red lead paint sludge sample (butylbenzylphthalate at 580 ppb and bis(2-ethylhexyl) phthalate at 3420 ppb) while three PAH compounds (naphthalene at 1260 ppb, 2-methylnaphthalene at 3330 ppb and phenanthrene at 850 ppb) and dibenzofuran (770 ppb) were found in the black cindery fill sample. No soil or groundwater standards could be found for the above compounds. However, the partition coefficient, $K_{\rm OC}$, for the above compounds range from 1,070 to 200,000 indicating that these compounds are highly attenuated in the soils.

3.4.4 PCBs

Arochlor 1254 was found in the red lead paint sludge sample at a concentration of 17 ppm.

3.4.5 Pesticides

No pesticides were detected above quantification limits.

3.4.6 Quality Control

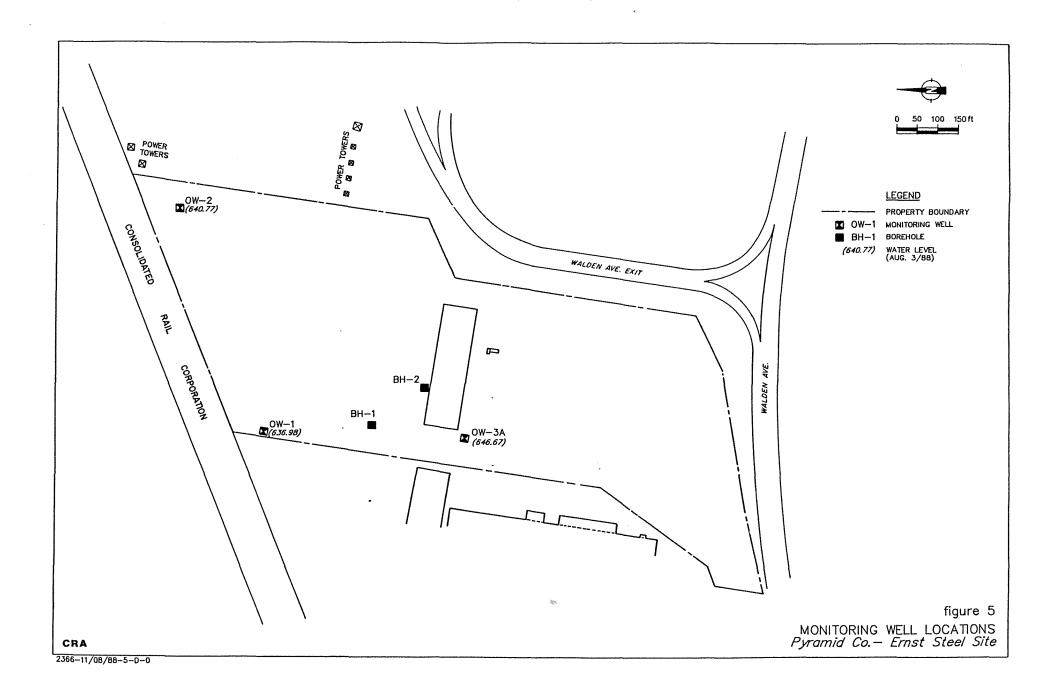
Following receipt of the final analytical data report from EnviroTest, a quality control assessment was conducted by CRA. Appendix A includes this assessment which was sent to EnviroTest for their consideration.

EnviroTest provided the missing dates of cyanide distillation indicating all reported values are correct. Due to semi-volatile surrogate analyses outside of the control limits, these samples were reanalyzed and the results are now within their control limits. The poor spike recoveries of lead, mercury, nickel and selenium result in qualifiers for these metals results indicating that the values are estimated. The average response factor for 2,4-dinitrophenol could not be brought into the required range after reanalysis and therefore the resultant values should be qualified as unusable.

4.0 MONITORING WELL INSTALLATIONS

In order to investigate the groundwater quality and flow conditions beneath the Site, three monitoring wells have been installed. OW-l is in the northwest corner of the Site, OW-2 is near the northeast corner of the Site, and OW-3 is located near the center of the Site (see Figure 5). The Monitoring Well Installation Program began on July 13, 1988, and was completed on July 18, 1988. Monitoring well locations were selected to avoid conflicts with future site usage. NYSDEC site representatives were present for the majority of the monitoring well installation program and assisted in the selection of locations.

The three monitoring wells installed to date are in the locations specified in the Work Plan. The necessity for a fourth installation was investigated on September 19, 1988 by the drilling of two additional boreholes at the locations shown on Figure 5. Due to the absence of groundwater and following discussions with the NYSDEC site representative, a fourth monitoring well was determined, with the concurrence of NYSDEC personnel, not to be required.



4.1 METHODOLOGY

Earth Dimensions, Inc. of East Aurora, NY was contracted by CRA to provide drilling services. Using a truck-mounted Mobile Drill B-61 drilling rig, each monitoring well and borehole location was continuously sampled to define the geologic stratigraphy. Samples were obtained in advance of the augering operation using standard penetration test split spoon sampling techniques. Sampling continued until auger refusal (top of bedrock) was encountered.

Each sample was logged according to geologic stratigraphy encountered by CRA personnel and a sample for geologic record was collected. (The Stratigraphic and Instrumentation Logs are included in Appendix B.) In addition, each split spoon sample from the monitoring well installations was scanned with an HNU photoionization detector immediately upon opening to check for the presence of volatile organic compounds.

In general, the sample HNU readings were equal to or only slightly higher than the background level, with the highest above background level (1 ppm) occurring in the sample from the 20 to 22-foot interval at OW-1. Table 6 lists the results of the HNU scanning.

Based on results of the scans of the soil cores obtained by split spoon sampling, organic contamination is not anticipated at the Site.

Table 6
HNU Scanning Results
Split Spoon Samples
Pyramid Co.-Ernst Steel Site

Well No.	Sample No.	Depth (ft BGS)	Background (ppm)	Total Reading (ppm)
OW-1	1 2 3 4 5 6 7 8 9 10 11 12 13 14 15	0-2 2-4 4-6 6-8 8-10 10-12 12-14 14-16 16-18 18-20 20-22 22-24 24-26 26-28 28-29	0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0 0.6-1.0	0.6-1.0 0.6-1.0 1.4 1.4-1.8 1.4-1.8 1.2 1.3 1.4-1.8 1.6 0.6 1.4-2.0 0.6-1.0 1.0 1.0 1.0
OW-2	1 2 3 4 5 6 7 8 9 10 11 12 13 14 15	0-2 2-4 4-6 6-8 8-10 10-12 12-14 14-16 16-18 18-20 20-22 22-24 24-26 26-28 28-29	1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6 1.2-1.6	1.6-1.8 1.4 1.6-2.0 1.2 1.2-1.4 1.2 1.2-1.4 1.0 1.4 1.4 1.4-1.6 1.4-1.6
OW-3	1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17	0-2 2-4 4-6 6-8 8-10 10-12 12-14 14-16 16-18 18-20 20-22 22-24 24-26 26-28 28-30 30-32 32-32.5	0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.8-1.2 0.5-0.8 0.5-0.8 0.5-0.8 0.5-0.8 0.5-0.8 0.5-0.8	1.6 1.4 1.2 1.2-1.5 1.0-1.4 1.4-1.6 1.0 1.0-1.2 1.0-1.2 1.2-1.4 0.8 0.6-0.8 0.8 1.2 1.0 1.0

Following the completion of split spoon sampling, a monitoring well was installed through the augers at three locations. Well construction materials consisted of a 5-foot long, 2-inch diameter #10 slot stainless steel well screen coupled to 2-inch diameter black steel riser pipe. A sandpack composed of #4 and/or #2 QROC quartzite sand was placed around and above the screen to a height determined by observed field conditions. Above the sandpack, a 2-foot bentonite pellet seal was placed. The remainder of the borehole was backfilled to the ground surface with cement/bentonite grout. Each well was fitted with a locking cap and lock. Table 7 summarizes the well installation details.

Prior to the installation of monitoring wells OW-l and OW-2, a 2-foot bentonite pellet plug was placed in each borehole at the bedrock/overburden interface to seal off the potential migration pathway for overburden groundwater into the bedrock regime. At the completion of split spoon sampling at OW-3, the borehole was observed to contain a great deal of sloughed material. In order to install a functional well, borehole OW-3 was grouted to the surface and a new borehole was augered (without sampling) five feet west of the original borehole. Since augering was terminated before the bedrock surface was encountered, an artificial migration pathway was not created. Therefore, a bottom bentonite seal was not necessary at this location.

Table 7
Well Installation Detail Summary
Pyramid/Ernst Steel Site

Well No.	Bentonite Seal Over Top of Bedrock (ft BGS)	Bottom of Screen (ft BGS)	Bottom of Sandpack (ft BGS)	Top of Sandpack (ft BGS)	Bentonite Seal (ft BGS)
OW-1	27.0-29.1	26.5	27.0	18.0	16.0-18.0
OW-2	28.0-29.9	27.0	28.0	16.9	15.0-16.9
OW-3	N/A	28.6	29.0	7.0	5.0- 7.0

4.2 DECONTAMINATION

Before initial use, between monitoring well and borehole locations, and at the completion of the program, all drilling tools were decontaminated using a pressurized steam cleaner.

The well screens and riser pipe were cleaned prior to installation using the steam cleaner. In addition, where deemed necessary by CRA personnel, equipment was decontaminated using nitric acid and solvent rinses as described in Section 3.2. This method of decontamination was performed in order to remove oil or cutting fluids from the riser pipe and well screens.

A mud tub was used to contain liquids generated during equipment decontamination. All wash fluids were then transferred to 55-gallon drums which are staged on-site pending determination of an appropriate disposal method.

4.3 WELL DEVELOPMENT

After the well installation program was completed, the wells were developed following the protocols outlined in the Work Plan.

Monitoring wells OW-1 and OW-2 were found to have insufficient water available for continuous development. Consequently, these wells were developed by hand bailing to dryness on each of three consecutive days. The bailer was also used for surging in an effort to draw into the well any fines which were trapped in the sandpack during well construction. Through development over three consecutive days, OW-1 yielded approximately 10 well volumes (20.5 gallons) while OW-2 yielded approximately six well volumes (6.25 gallons).

Well OW-3 was developed by purging ten well volumes (37 gallons) in one day. An ISCO portable peristaltic pump and a bailer were both used to develop and surge OW-3. Clean teflon tubing connected to a peristaltic pump and a bailer were both inserted to the bottom of the well. By simultaneously pumping the well while raising and lowering the bailer, the well was surged and developed.

All purge water was containerized at the well sites and transferred to 55-gallon drums which are staged at the Site pending determination of an appropriate disposal method.

4.4 PHYSICAL SOILS TESTING

In order to determine the permeability of the on-site soils, selected samples collected during the monitoring well installation program were submitted to a laboratory for physical testing including grain size distribution, Atterberg Limits and permeability. The samples submitted are as follows:

Sample Location	Sample Number	Depth		Description
OW-1	S-5	8-10		Silty-Clay
OW-2	S-6	10-12		Silty-Clay
OW-3	S-10	18-20	(composited with OW3, S-11)	Glacial Till
OW-3	S-11	20-22	(composited with OW3, S-10)	Glacial Till

Appendix C contains the physical testing results. The results for each of the two clay samples were similar. The permeability of the clay appears to be in the low 10^{-8} cm/sec range. The submitted till samples could not be tested for permeability because the samples were not cohesive enough to measure the bulk unit weight. This measurement is required prior to recompaction of the samples for permeability testing. Two additional till samples (OW-3, S-7 and S-8) were sent for analysis and the permeability of the composited till samples was 1.1×10^{-7} cm/sec.

5.0 SITE GEOLOGY AND HYDROGEOLOGY

5.1 SITE GEOLOGY

The site stratigraphic units, in descending order with depth, generally are:

- i) Fill
- ii) Silt
- iii) Clay
- iv) Till

Not all of the above units are present at all locations. As presented in Section 2, the fill ranges in thickness from 0 to 3.5 feet and consists of a variety of materials. Clay generally underlies the fill material except at well OW-l and borehole BH-2 where a thin discontinuous layer of native silt (0.5 to 1.0 feet thick) exists. The clay ranges in thickness from 5.1 to 21.2 feet. Underlying the clay is a silt/clay till ranging in thickness from 6.4 to 26.7 feet.

5.2 SITE HYDROGEOLOGY

The site hydrogeology units, in descending order with depth are:

- i) Fill/silt and
- ii) Clay/Till

The fill/silt and clay/till have been grouped as discrete hydrogeologic units due to similar flow characteristics. This is demonstrated for the clay/till by the physical soils testing data previously presented in Section 4.4 which identifies the clay and till to be similar at least from a hydrogeologic perspective.

The fill/silt layer is a coarser more porous unit that would be expected to be a perched water unit. However, often during the excavation and drilling programs, the fill/silt layer was observed to be dry or moist indicating minimal water presence. No wells are screened in the fill/silt unit since the Work Plan was written with the intent of identifying chemical migration from the Site. The soil sampling programs were designed to evaluate chemical presence within the fill unit itself. The clay/till layer is an aquitard and will be insignificant in terms of groundwater movement as demonstrated by the permeabilities determined in the laboratory (see Appendix C).

Three rounds of groundwater levels have been taken as shown on Table 8. It is noted that the data for OW-2 on July 20, 1988 is not representative of site conditions since this a slow recovering well and had not as yet reached equilibrium on July 20, 1988. The two subsequent rounds of water levels indicate that the limited groundwater flow is to the northwest.

Based upon the geologic stratigraphy descriptions obtained from the borings conducted on the Walden Galleria Site and the Ernst Steel Site, there is considerable information available concerning the subsurface conditions at the Site. Generally, the native soils at the Site are fine grained consisting mostly of clay and silt. Consequently, the native soils are not conducive to extensive groundwater presence or movement. This was confirmed during the excavation of several test pits and holes around the site; many of which remained dry throughout the period for which they were open. As a result, there is little concern or potential for contamination of the groundwater regime beneath the site due to any surface or near-surface activities. The clay and till strata are aquitards and therefore minimal groundwater flow beneath the Site both vertically and horizontally will occur.

TABLE 8

GROUNDWATER ELEVATIONS (ft. ams1)

Monitoring	Top of	5	Sampling Date						
<u>Well</u>	<u>Casing</u>	7/20/88	8/3/88	9/12/88					
OW-1	653.37	639.29	636.98	638.97					
01.7	C = 2 = 20	620.06	C40 88	<i>-</i>					
OW-2	653.72	632.06	640.77	643.82					
OW-3	653.84	647.68	646.67	648.04					

6.0 100-FOOT GRID SAMPLING PROGRAM

In conjunction with the Phase II Site

Investigation and as a result of the Waste Characterization

Sampling Program, further field studies were undertaken to

determine the extent of lead presence in the fill materials

at the Site. This sampling program concentrated on the

extent of lead but also included TCL analysis on selected

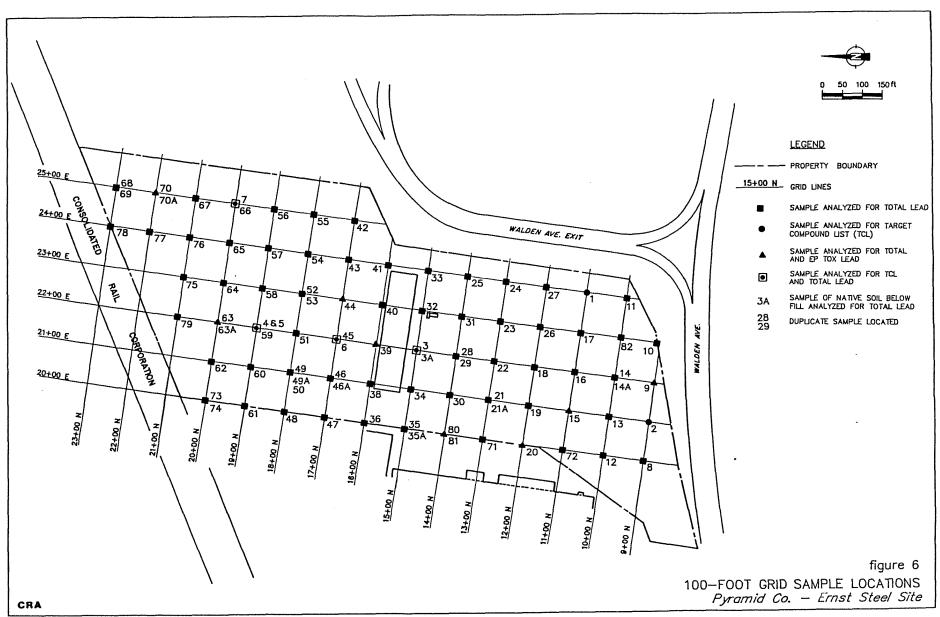
samples. This analytical data was used in conjunction with

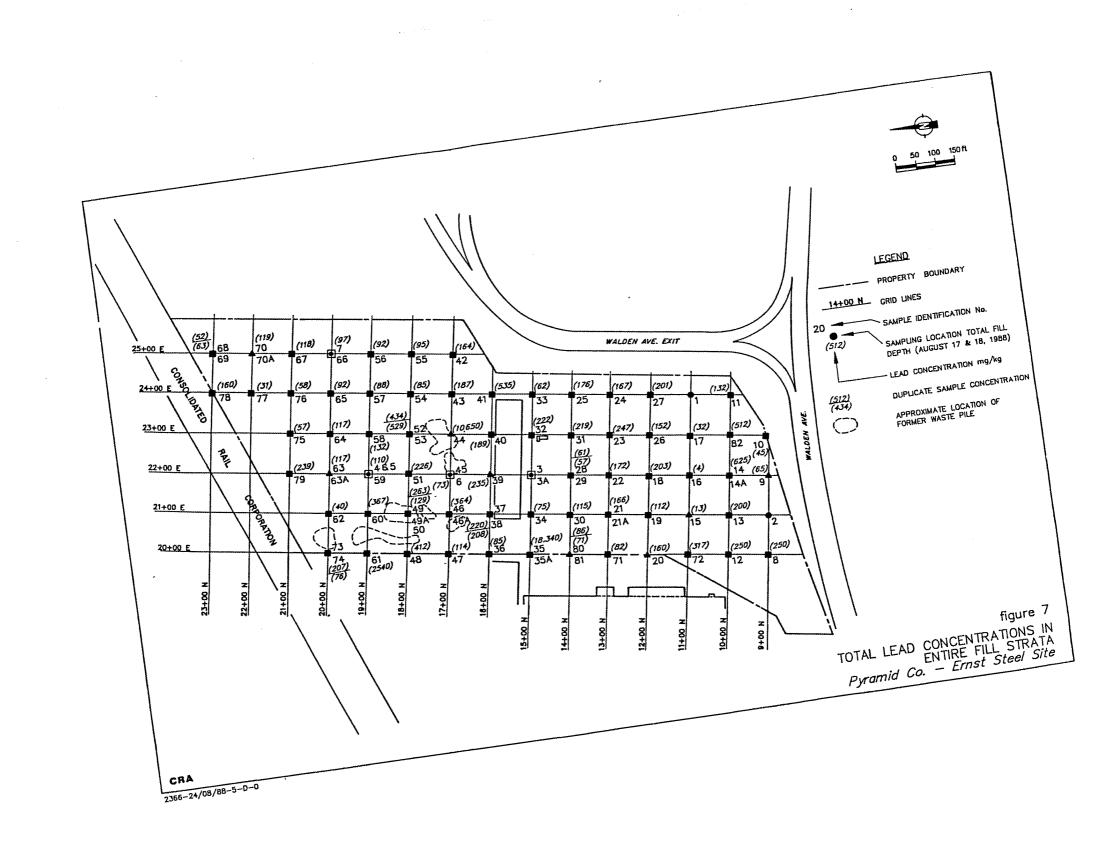
the waste characterization analytical data to develop the

SSIP list for subsequent sampling programs.

On August 17 and 18, 1988, a 100-foot sampling grid was established on the Site. At each grid point, samples of the entire fill horizon extending from just below the surface material to the top of the underlying native soils were collected and analyzed for total lead. The sampling locations (71 locations) are presented in Figure 6. At eight sampling locations, duplicate soil samples were collected for QA/QC purposes for a total of 79 samples. The analytical results are presented in Appendix D and summarized on Figure 7.

At eight of the 71 locations, the sample was analyzed for EP Toxicity lead as well as total lead. The locations of the EP Toxicity sample points and analytical





results are summarized in Figure 8. The analytical results are presented in Appendix D.

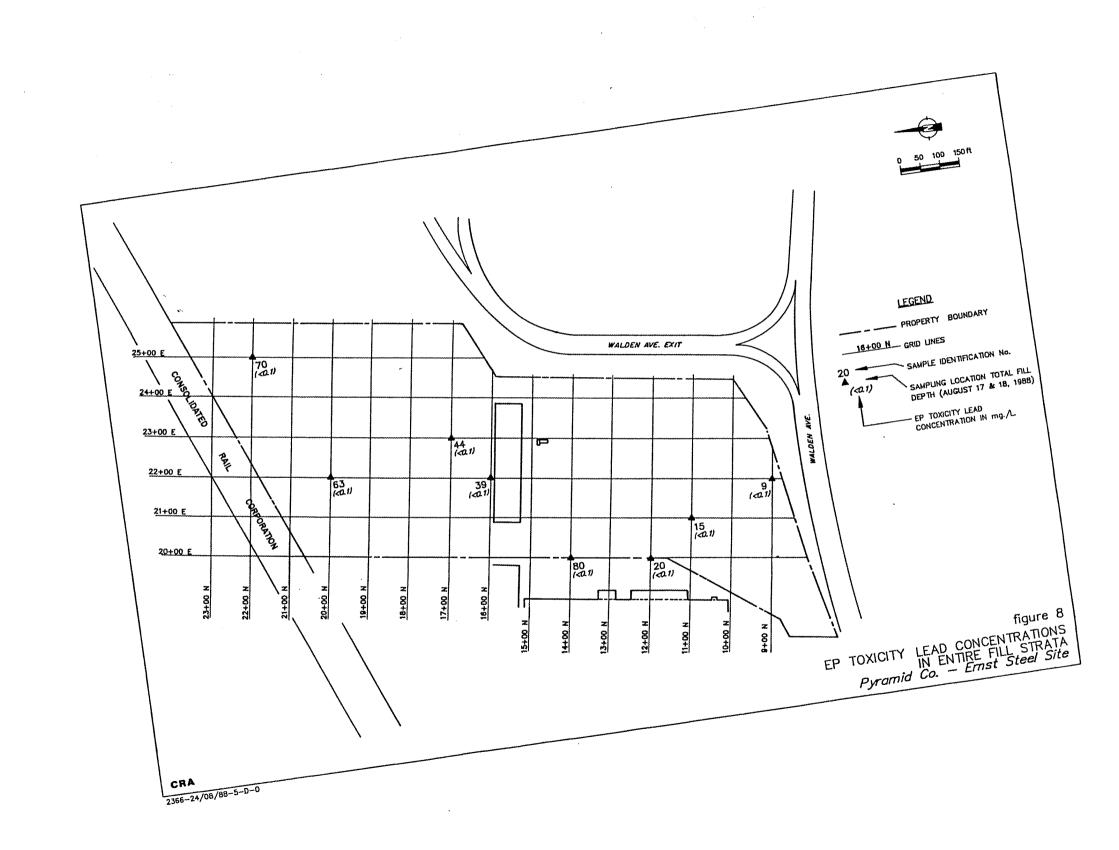
At eight additional locations beneath the area of fill presence, samples of the underlying native material were collected and analyzed for total lead. Figure 9 presents the locations of native soil sample collection as well as the concentrations of total lead in the native materials and overlying fill material at each of these sample points. The analytical results are presented in Appendix D.

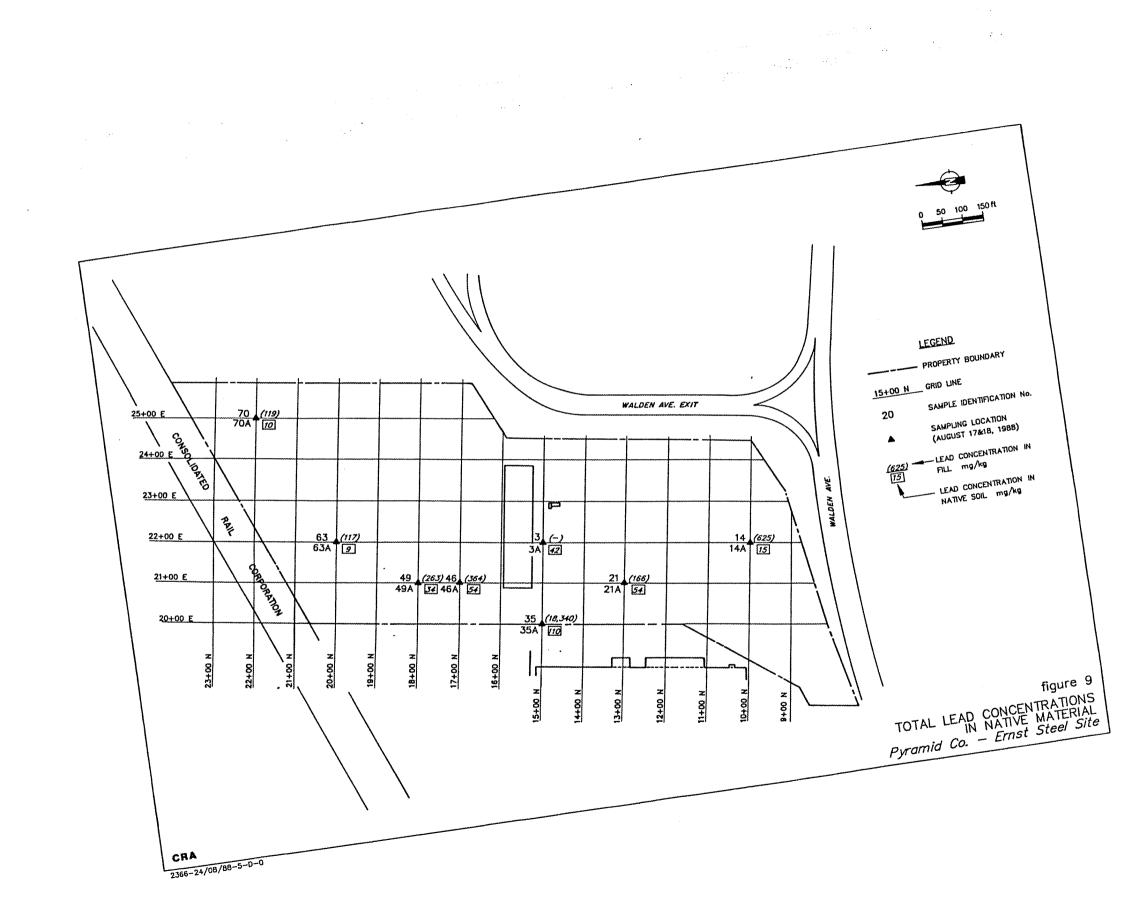
As a result of the site work performed, the areal extent of fill has been delineated. Figure 10 presents the boundaries of this area and the depths of fill.

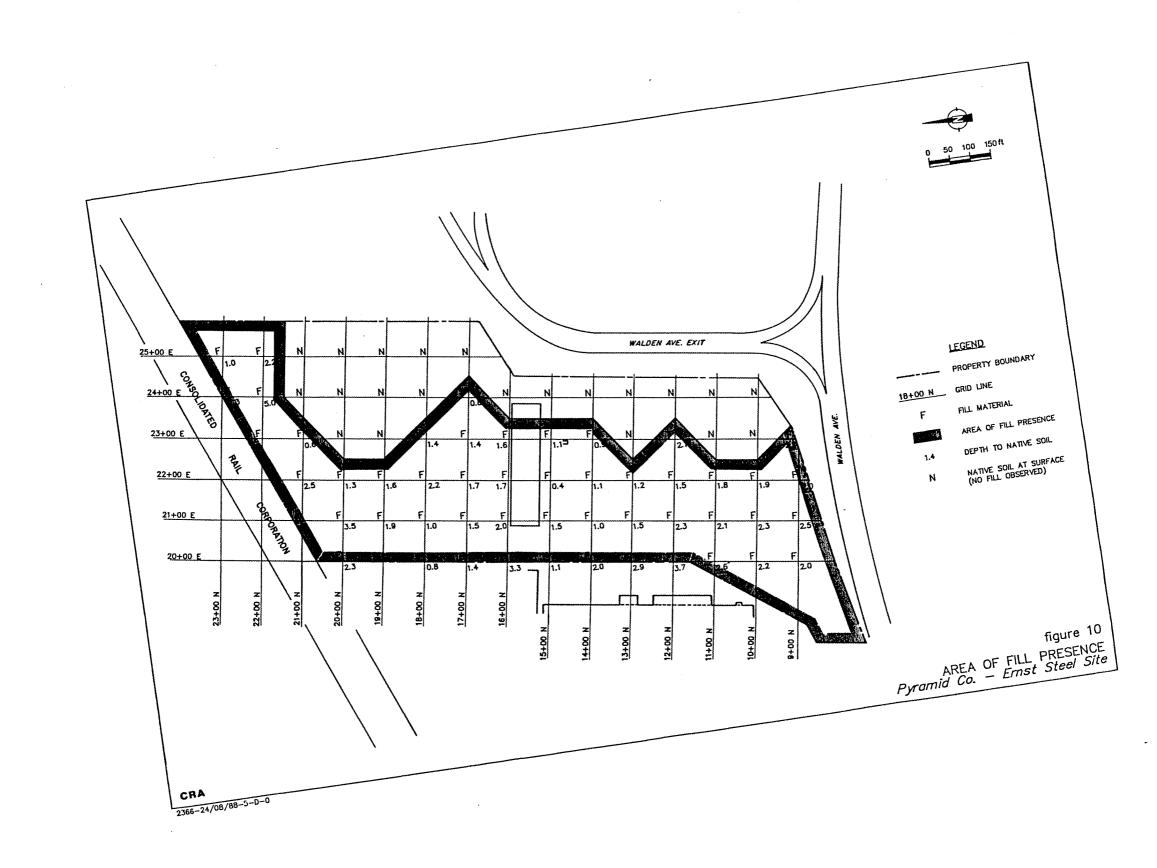
To confirm the waste characterization, additional samples were collected of the entire fill strata at six locations (plus one duplicate sample) and analyzed for the TCL. The sampling locations are presented on Figure 6 and the analytical results are presented in Appendix E.

6.1 METHODOLOGY

At each 100-foot grid sampling location, a backhoe was used to excavate a test pit. The samples of the







fill or native soil at each location were collected in the same manner as the waste characterization sampling described in Section 3.1.

6.2 <u>DECONTAMINATION</u>

Sampling tools were cleaned and wrapped before each use using the protocols described in Section 3.2. All cleaning fluids were contained and placed into a 55-gallon drum which is staged on-site pending determination of an appropriate disposal method.

6.3 SAMPLE HANDLING

Manner as the waste characterization samples (see
Section 3.3). The total lead and EP Toxicity lead samples
were transported to BLT Technical Services, Inc. of Niagara
Falls, New York by CRA personnel. The TCL samples were
shipped via Federal Express to Recra Environmental, Inc. of
Columbia, MD. Appropriate Chain of Custody procedures were
followed throughout the sampling and shipping process.

6.4 ANALYTICAL RESULTS

The analytical results for total lead and EP Toxicity lead are presented in Appendix D and the TCL analytical results from the 100-foot grid sampling study are presented in Appendix E. The TCL data is summarized on Table 9. The following observations have been made.

6.4.1 Metals

The results of the metals analyses indicate the following metals above background concentrations:

<u>Metal</u>	Sample No.	Concentration (ppm)	Background Concentration (ppm)
Cadmium	3 4 7	2.0 0.98 1.2	0.41 - 0.57
Iron	6	52,100	5000 - 50000
Lead	2 6	931 761	42-544
Manganese	2	3,010	7 - 3000
Zinc	6	361	<5 - 300

Background concentrations are for U.S. soils and were obtained from "Trace Elements in Soils and Plants" by Kabata-Pendias and Pendias (1984) except for lead which was obtained from the Interim Remedial Measures Plan.

TABLE 9

COMPOUNDS DETECTED - 100-FOOT GRID SAMPLING STUDY
PYRAMID/ERNST STEEL SITE

Metals (ppm)	<u>#1</u>	#2	#3	#4	<u>#5</u>	#6	<u>#7</u>	Background Concentration
Al	38100	26900	16800	10100	7380	8390	15500	4500 - 100000
As	7.6	8.4	4.9	12	8.9	7.7	9.5	
Ва	106	185	97	77	50	105	70	10 - 3000
Ве	1.5	3.9	2.1	1.1	0.93	0.76	0.95	<1 - 7
Cđ			2.0	0.98			1.2	0.41 - 0.57
Ca	3940	149000	93500	37500	34100	7820	3010	
Cr	21	25	18	16	10	38	16	3 - 1500
Co	4.8	6.2	3.7	7.0	5.9	7.4	3.2	0.3 - 50
Cu	16	36	42	12	35	51	15	1 - 300
Fe	12100	39900	29600	42300	49300	52100	7570	5000 - 50000
Pb	90	931	299	191	105	761	112	42 - 544
Mg	3080	23000	16800	4610	3620	2120	1810	
Mn	51	3010	2590	2330	1340	1610	83	7 - 3000
Ni	12	13	9.5	17	13	19	11	<5 - 200
К	2990	2690	1220	1440	1020	1480	1840	
Na	580	1760	1170	1380	1390	1220	324	
Thallium				0.74			****	0.01 - 2.3
Vanadium	30	16	13	23	15	21	23	
Zn	108	155	88	211	133	361	89	<5 - 300

continued....

TABLE 9

COMPOUNDS DETECTED - 100-FOOT GRID SAMPLING STUDY
PYRAMID/ERNST STEEL SITE

Volatiles (ppb)	#1	#2	#3	#4	#5 ——	#6	#7 	Partition Coefficient K OC
Methylene Chloride*	8	7	9	18	22	23	54	10
Trichlorofluoromethane	5	11		10	15	18	10	182
Chloroform	6	7	7	6	6	6	7	50.2
BNA (ppb)								
Naphthalene						370		1070
Phenanthrene		1100				510		160000
Fluoranthene		1000				370		44000
Pyrene		1700			350	420		44000
Benzo(a)Anthracene		860						220000
Chrysene		930						220000
Benzo(a)Fluoranthene		1100			360	370		630000
Benzo(a)Pyrene		680						630000
Pesticides (ppb)			· 					

^{* =} also present in method blank (5 ppb)

^{-- =} Not detected above quantifiable limits

All the metals which exceed background concentrations do so only by small amounts.

6.4.2 Volatiles

Only three volatiles, methylene chloride (7-54 ppb), trichlorofluoromethane (ND-18 ppb) and chloroform (6-7 ppb), were detected in the seven samples. All of the detected values are low-level.

6.4.3 <u>BNAs</u>

Three of the seven soil samples indicated the presence of several Polynuclear Aromatic Hydrocarbon (PAH) compounds as presented on Table 9. No soil or groundwater standards could be found for the detected compounds except for benzo(a)pyrene. New York State regulations for class GA potable water states that benzo(a)pyrene must be non-detectable. However, the partition coefficient, ${\rm K}_{\rm OC}$, for benzo(a)pyrene is 630,000 indicating that this compound is very highly attenuated in soil. The ${\rm K}_{\rm OC}$ values for the remaining detected PAH compounds range from 1,070 to 630,000 indicating these compounds are also highly attenuated in the soils.

6.4.4 Pesticides and PCBs

No pesticides or PCBs were detected above quantifiable limits.

6.4.5 Quality Control

A quality control assessment of the BLT analytical results for total and EP Toxicity lead was performed by CRA and is presented with the data in Appendix D. All standard laboratory and field QA/QC requirements were adhered to and therefore the presented analytical data is acceptable.

A quality control assessment of the Recra analytical results for TCL parameters was also performed by CRA and is presented with the data in Appendix E. A number of problems were observed which resulted in the following qualifiers to the data being recommended:

a) Poor surrogate recoveries were achieved in the VOA
analyses due to matrix interference and therefore all VOC
results associated with the soil samples should be
qualified as estimated.

- b) Due to poor recoveries on sample spikes for metallic analytes antimony, arsenic, manganese, selenium, silver, thallium and zinc should be qualified as either estimated values or unusable.
- c) On the basis of poor precision established for duplicate analyses of copper and nickel, all associated values for these two analytes should be qualified as estimated.

7.0 SITE SPECIFIC PARAMETER SELECTION

Following the waste characterization and 100-foot grid sampling programs, the TCL analytical data was reviewed for selection of the SSIP. As a result of discussions with the NYSDEC on September 8, 1988, the following set of SSIP has been selected:

<u>Parameter</u>	Method					
	Soil	Water				
Lead (Total)	SW-846-7420	SW-846-7421				
EP TOX Lead (soils only)	SW-846-1310/7420					
Chromium (total)	SW-846-7190	SW-846-7191				
EP TOX Chromium (soils only)	SW-846-1310/7190					
Arochlor 1254	SW-846-8080	SW-846-8080				
Total Organic Halides (TOX)	SW-846-9020	SW-846-9020				
Toluene (water only)		SW-8020				

8.0 WORK PLAN SAMPLING PROGRAMS

Following the determination of the SSIP, the remainder of the sampling programs specified in the Work Plan for the Phase II Site Investigation were conducted on September 12 to 20, 1988.

8.1 MONITORING WELL SAMPLING

The wells were purged and sampled following the protocols outlined in the Work Plan. Prior to purging, water level measurements were taken in each well. The results are presented in Table 8.

Monitoring wells OW-1 and OW-2 were purged to dryness on three consecutive days (September 12 to 14, 1988). On September 15, 1988, both wells were sampled for the SSIP by bottom-loading stainless steel bailer. Split samples collected for the NYSDEC were discarded by CRA personnel at the direction of the NYSDEC site representative. In order to collect samples representative of groundwater flow conditions in a porous media, groundwater samples collected for metals analysis were field-filtered during these sampling activities. However, the NYSDEC requested that non-filtered samples be analyzed. To obtain non-filtered samples, the two wells were again purged to dryness on three consecutive days

(September 17 to 19, 1988) and resampled for unfiltered metals analysis on September 20, 1988. The NYSDEC was informed of the resampling and a NYSDEC site representative was on-site during purging, however the NYSDEC representative was not on-site during sampling and no split samples were collected. The samples were collected by low-rate pumping (peristaltic pump) to collect sediment-free samples.

Monitoring well OW-3 was purged of five well volumes (20 gallons) prior to sampling for SSIP on September 12, 1988. Purging was conducted by peristaltic pump and sampling was conducted using a bottom-loading stainless steel bailer. Split samples were collected for and retained by the NYSDEC site representative. The NYSDEC split samples were not field-filtered. On September 20, 1988, the well was resampled for unfiltered metals following purging of five well volumes (20 gallons). Both purging and sampling was conducted by peristaltic pump.

Teflon tubing was dedicated to each well.

Bailers were precleaned prior to each sampling event using the protocols described in Section 3.2.

8.2 SURFACE SOIL SAMPLING

Samples of the upper surface of the fill material were collected at the ten locations indicated on

Figure 11. These locations were selected in concurrence with the NYSDEC site representative. Each sample was collected from the upper one-foot of the fill material. This depth was suggested by the NYSDEC field representative as representative of the surface soil. Previous sampling activities have characterized the entire fill strata.

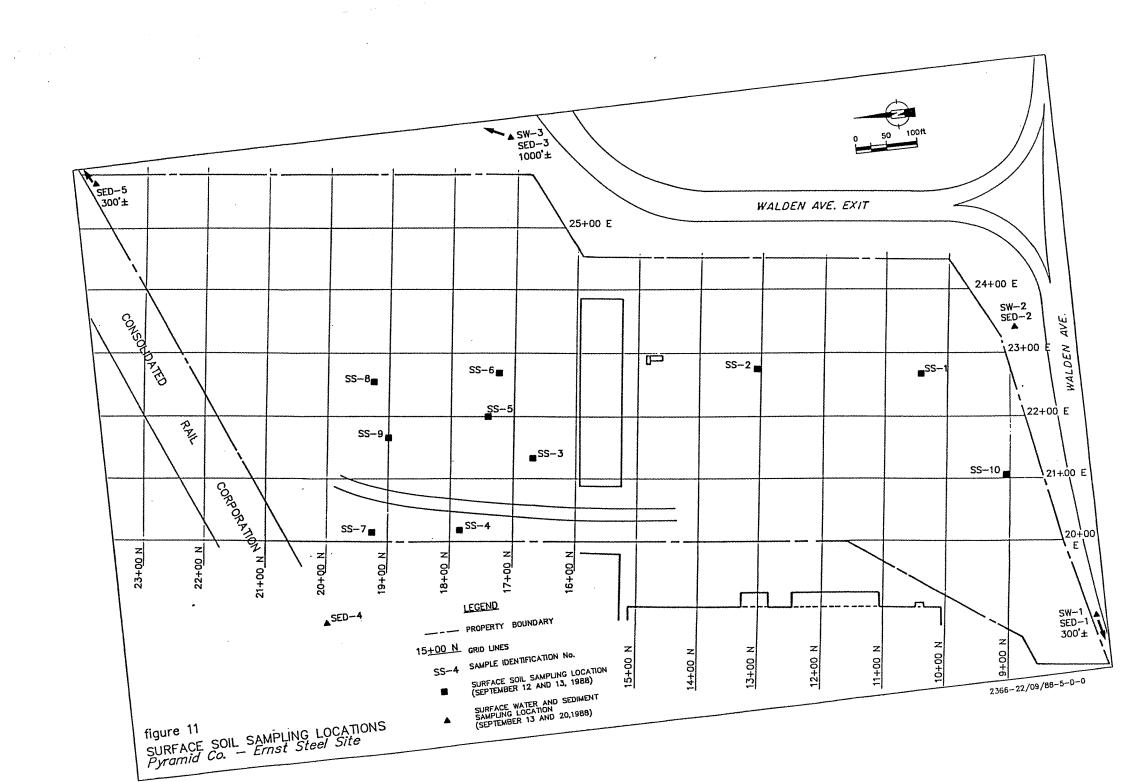
The surface soil samples were collected using the following procedures:

- i) dig a hole with a pre-cleaned shovel to a depth of approximately one foot,
- ii) scrape the side of the hole with a precleaned garden trowel, and
- iii) from the freshly exposed material, scoop and place the sample directly into the sample jar using the trowel.

The sampling tools were cleaned between each sampling location using the protocols described in Section 3.2.

8.3 SURFACE WATER AND SEDIMENT SAMPLES

Surface water and sediment samples were collected from the ditches bordering the Site as specified in the Work Plan. The locations from which samples were collected are presented on Figure 11. The ditch along the



south side of the Site contained surface water at the southwest corner of the Site. A surface water sample and a sediment sample were collected at this location. It is noted that the surface water at this location appeared to flow to the west. The remainder of the south ditch was dry between the southwest corner and the surface drain at the southeast corner of the Site. Surface water flowing from the surface drain was sampled prior to the flow entering the ditch. A sediment sample was collected at this location. then continued along the south ditch to the east and north until discharging into the Scajaquada Creek. Surface water and sediment samples were collected from the ditch prior to discharge into the Creek. The north ditch was dry. Consequently, no surface water samples could be collected in the north ditch, however sediment samples were collected at two locations (Sed-4 and Sed-5).

Surface water samples were collected by dipping the glassware in the water or by catching the cascading water as in the surface drain sample. The sediment samples were collected following the procedure specified in the Work Plan.

8.4 DUPLICATE SAMPLING

For quality control purposes, a duplicate sample was collected for each different sample type: groundwater, surface soil, surface water and sediment. No split samples were required by the NYSDEC site representative.

8.5 ANALYTICAL RESULTS

The laboratory report of the analytical results is presented in Appendix F. A summary of the analytical results is presented in Table 10 for all of the SSIP and the following observations have been made.

8.5.1 TOX and Toluene

Total organic halides (TOX) were detected in both the groundwater and surface water samples. The largest detected TOX concentration in the surface water is 60.4 ppb. In the groundwater, the highest TOX concentration is 28.6 ppb. All of the detected concentrations are low considering the group of organics which comprise TOX. One low-level concentration of toluene was also detected in OWl (0.53 ppb). The New York State Department of Health has recently promulgated Maximum Contaminant Levels (MCL)

TABLE 10

PHASE II SITE SPECIFIC INDICATOR PARAMETER RESULTS

	Lea	a <u>d</u>	EP Toxicity Lead (ppm)	Chr	omium	EP Toxicity Chromium (ppm)	Arochlor 1254 (ppb)	TOX (ppb)	Toluene (ppb)
Groundwater	Filtered (ppm)	Unfiltered (ppm)		Filtered (ppm)	Unfiltered (ppm)				
OW-1	0.003	<.002		0.013	0.007		<1.0	28.6	0.53
OW-2	<.002	<.002		0.006	0.003		<1.0	<10.0	<0.5
OW-102*		<.002			<.003				
OW-3	<.002	<.002		<.003	<.003		<1.0	<10.0	<0.5
OW-103*	<.002			<.003			<1.0	18.6	<0.5
Surface Water									
SW-1	<.002	0.007		0.005	<.003		<1.0	15.4	<0.5
SW-2	<.002	0.008		<.003	0.003		<1.0	60.4	<0.5
SW-102*	0.002			<.003			<1.0	35.4	<0.5
SW-3	<.002	0.002		<.003	<.003		<1.0	49.0	<0.5
Soil							(ppm)	(ppm)	
SS-1		219	0.049	1	6	<.02	<.2	44	
SS-2		507	0.048	. 1	9.3	<.02	<.2	34	
SS-3		181	0.054	2	22.5	<.02	<.2	15	
SS-4		,600	49.9	18		<.02	1.2	34	
SS-5	8,	,980	15.6	18	39	0.12	<.2	75	
SS-6	5,	,570	35.6	47		0.38	0.6	60	
SS-7	14,	,200	132	. 16		0.038	1.6	26	
SS-107*	14,	,200	229	13		0.024	2.2	34	
SS-8		563	0.23		30.1	<.02	<.2	35	
SS-9		312			9.2	<.02	<.2	25	
SS-10	4,	,370	50.4	2	27.8	<.02	<.2	17	
Sediment							(ppm)	(mqq)	
Sed-1		286	0.12		25.1	<.02	<.2	<10	
Sed-2		118	0.057		19.8	0.024	<.2	<10	
Sed-3		38.1	0.035		8.8	<.02	<.2	<10	
Sed-103*		31.5	0.02		9.1	<.02	<.2	<10	
Sed-4		255	0.043		21.2	<.02	<.2	111	
Sed-5		35	<.02	1	6.7	<.02	<.2	13	

^{*} Duplicates identified in this manner, i.e., OW-102 is duplicate of OW-2.

in Part V of the Sanitary Code. The new MCL for toluene in drinking water is 5 ug/L (ppb). It is noted that all of the toluene concentrations measured at the Site were below the MCL.

8.5.2 Lead and Chromium

Lead was detected in the sediment samples ranging from 32 to 286 ppm and in the surface soil samples ranging from 181 to 29,600 ppm. The lead results for six of the ten surface soil sample locations are above background concentrations while five of these locations also had EP Toxicity results above the 5 ppm concentration which defines a hazardous waste pursuant to the EP Toxicity test. None of the lead results for the sediment samples were above background. In fact, the two downgradient samples (Sed-3 and Sed-5) had lead concentrations of only 38 and 35 ppm respectively which are approximately one order of magnitude lower than the concentrations of the upgradient samples. This indicates that the lead presence at the Site has not migrated via the ditches to the Scajaquada Creek.

Lead was not detected above the quantifiable level (0.002 ppm) in any of the unfiltered groundwater or filtered surface water samples. Lead was detected at or above the quantifiable level (0.002 ppm) in all three unfiltered surface water samples SW-1 (0.007 ppm), SW-2 (0.008 ppm) and SW-3 (0.002 ppm) and in one filtered

groundwater sample OW-1 (0.003 ppm). Although one filtered groundwater sample shows a higher concentration than the associated unfiltered sample, the results are essentially identical within the reproducibility capabilities of the analytical techniques and are therefore, consistent. The New York State Class GA standard for lead is 0.025 ppm.

The higher concentrations of chromium found in the surface soil correspond to the locations of the highest surface lead concentrations. Chromium, like lead, was detected at lower concentrations in the downgradient sediment samples than in the upgradient samples. The three surface soil samples (SS-4, SS-5 and SS-6) where EP Toxicity chromium was detected did not exceed the EP Toxicity criteria for chromium (5 ppm). As well, the samples are located in areas where the lead concentration exceeds 1000 ppm. Therefore, remediation of lead contaminated soils will sufficiently address any chromium contamination as well.

Chromium was detected at or above the quantifiable level (0.003 ppm) in two unfiltered groundwater samples (OW-1 at 0.007 ppm and OW-2 at 0.003 ppm), two filtered groundwater samples (OW-1 at 0.013 ppm and OW-2 at 0.006 ppm), one unfiltered surface water sample (SW-2 at 0.003 ppm) and one filtered surface water sample (SW-1 at 0.005 ppm). Although three filtered samples show higher concentrations than the associated unfiltered samples, the

results are essentially within the reproducibility capabilities of the analytical techniques and are therefore, consistent. The New York State Class GA standard for hexavalent chromium is 0.05 ppm. It is noted that analyses were for total chromium, not just the hexavalent isotope.

The isolated single occurrence of lead concentrations above 1,000 ppm in the southern portion of the Site may be an artifact of sampling. During sampling, a small area (approximately 1 square foot) with an orange/red material was noted. Some of this material was placed in the sample. During collection of this sample, the remains of a paint can were found. No other observances of the orange/red material were made in the vicinity of the area of sample collection. During site excavation, the backhoe removed a small volume of soils (including all the orange/red material) which was transported to a secure landfill site for proper disposal.

8.5.3 Arochlor 1254

Arochlor 1254 was detected at four surface soil sampling locations but was not detected in any of the sediment samples above the quantified limit of 0.2 ppm. The locations of Arochlor 1254 presence (0.6-2.2 ppm)

correspond to areas with lead concentrations above 1000 ppm.

Therefore, any Arochlor 1254 contaminated soils will be addressed by the removal of lead contaminated material.

Arochlor 1254 was not detected above the quantified limit (1.0 ppb) in any of the groundwater or surface water samples. The New York State Class GA standard for PCB is 0.1 ppb.

9.0 HEALTH AND SAFETY

The health and safety requirements were as described in the Work Plan and outlined below.

9.1 TEST PIT INVESTIGATION AND WASTE CHARACTERIZATION SAMPLING

Throughout the test pit investigation, air monitoring was performed at 15 minute intervals during active excavation activity. An HNU photoionization instrument was used to scan for organic vapors and a Sibata Dust Indicator was used to measure dust generated by the excavation process. No excursions were noted with either instrument.

CRA personnel were equipped with tyvek coveralls, hard hats, safety glasses, latex gloves and rubber boots. Respirators were on hand but were not worn. A full complement of safety equipment was available for use by the backhoe operator. Since the operator sat in an enclosed cab and did not leave the machine during the test pit investigation, wearing of the protective gear was not necessary.

At the completion of work, the backhoe bucket was cleaned using a water wash. All cleaning water was

contained in a 55-gallon drum and staged temporarily at the Site. All Tyvek, gloves, etc. were collected in plastic garbage bags and also staged temporarily at the Site. During the subsequent site excavation, the staged drums and bags were transported, along with the contaminated soils to a secure landfill site.

9.2 MONITORING WELL INSTALLATION AND WELL DEVELOPMENT

During the drilling and well development programs, the drillers and samplers wore Tyvek coveralls, boots, gloves, hardhats, and safety glasses. Respirators were on hand but were not worn. At each drilling location, a survey site was constructed by placing a ground sheet of polyethylene over the ground. The polyethylene was then covered with sheets of plywood. Finally the survey site was enclosed with plastic construction fencing.

All soil cuttings that were brought to the surface during augering and monitoring well installation were placed into 55-gallon drums. All used Tyvek, gloves, etc. were collected in plastic garbage bags for future disposal. The waste drums have been staged on-site pending analytical work to determine an appropriate method of disposal. The drums and bags were staged until site excavation, at which

time they were transported, along with the contaminated soils, to a secure landfill site.

9.3 WORK PLAN SAMPLING PROGRAMS

During all groundwater, surface water, surface soil and sediment sampling, CRA personnel were equipped with Tyvek coveralls, hard hats, safety glasses, latex gloves and rubber boots. Respirators were on hand but were not worn.

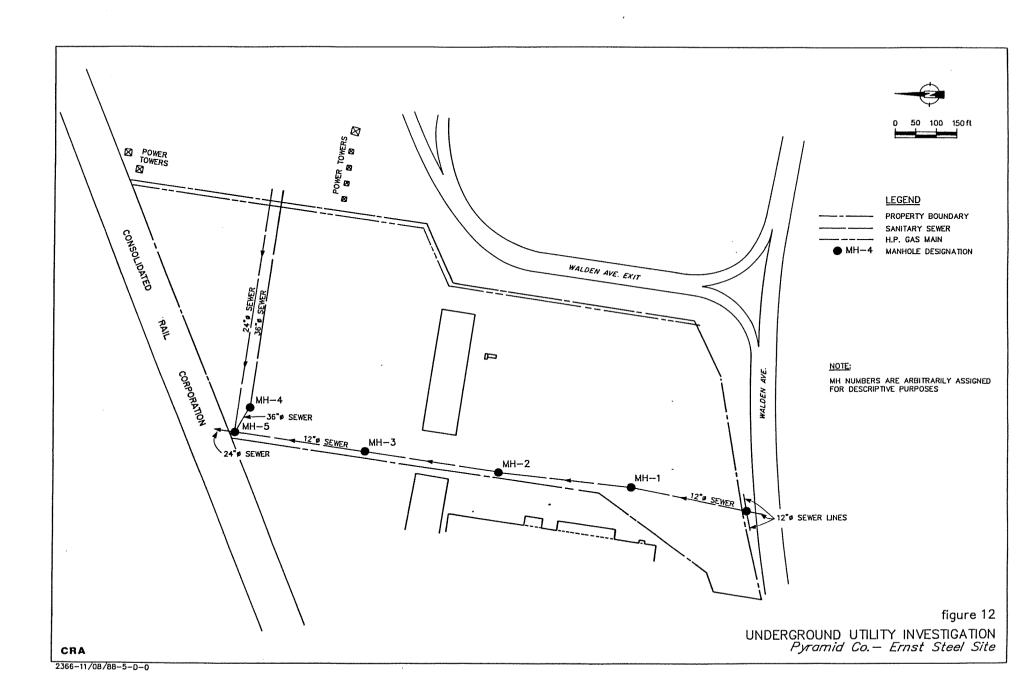
All purge water was staged temporarily at the Site in 55-gallon drums. All Tyvek, gloves, etc. were collected in plastic garbage bags and also staged temporarily at the Site. During the subsequent site excavation, the staged bags were transported, along with the contaminated soils, to a secure landfill site. The drummed water was transported to the same facility for treatment.

10.0 UTILITY INVESTIGATION

To date, the only known underground utilities on-site are a sanitary sewer network and a 24-inch diameter gas main (see Figure 12).

The sewer consists of a north-south segment which is located parallel to the western property line. This 12-inch diameter line flows to the north and empties into a manhole located near the northwest corner of the property (MH-5). In addition there are two lines which appear to run east-west through the Site. One is a 24-inch diameter line which flows from the east into MH-5. The other is a 36-inch diameter line which flows from the east into MH-4 located southeast of MH-5. MH-4 and MH-5 are connected by a 36-inch diameter line.

A visual inspection of MH-1 and MH-3 identified a single flow stream to the north with no other apparent stubs. MH2 has a welded steel plate on top and therefore, could not be inspected. MH-4 has a stub entering from the east and a stub entering from the northwest (to MH-5). Although flow in the sanitary sewer system could be heard in the manhole, there was no detectable flow observed in this manhole. MH-5 has a stub entering from the south. This is the 12-inch diameter north-south line. Effluent from this line cascades to the bottom of MH-5. There is also a



36-inch diameter line entering from the southeast (to MH-4), a 24-inch diameter line entering from the east and a 24-inch diameter line heading nearly due north. Visual observation of flow conditions in MH-5 indicate that the direction of flow leaving MH-5 is through the 24-inch diameter line to the north. This line is the deepest service (lowest invert elevation) in the system.

The 24-inch diameter line (from the east to MH-5) is believed to tie into a manhole located east of the Ernst property. The actual origin/termination of the 36-inch line (from the east to MH-4) is unknown.

The sanitary sewer has been observed to be quite deep. During visual inspection of MH-5 the invert of the north-south line was measured to be approximately 17 feet below the rim of the manhole. The bottom of MH-5 was measured to be approximately 23 feet below the rim.

The natural gas line is a large diameter high pressure gas main which generally runs north-south, roughly parallel to the eastern fence line. The gas main is approximately 5 feet west (inside) of the fence and is believed to be approximately 4 feet below the ground surface.

Chemical migration, if any, via the bedding materials of the underground utilities is expected to be minimal due to:

- i) the current minimal observed impact on the groundwater due to existing site conditions,
- ii) remediation of the Site pursuant to the Interim Remedial Measures Plan reduces the source of chemistry, and
- iii) on-site vaulting (see Section 12) will reduce the impact of soils remaining on-site.

11.0 CONCLUSIONS

The analytical results presented and discussed in this report indicate that off-site migration via surface water, sediments and groundwater has not occurred. The impact on the groundwater from the on-site materials has been minimal. This is as expected due to the protection of the groundwater table by the considerable clay/till unit underlying the site. This unit has a low permeability and there were no observations made of any significant waterbearing lenses in the clay/till. Consequently, there is little potential for contamination of the groundwater regime beneath the Site due to Site conditions. No Class GA standards have been exceeded. This statement is qualified by recognizing that the quantifiable limit for Arochlor 1254 was 1 ppb while the Class GA standard is 0.1 ppb.

Future impact on the groundwater, after site remediation, would be less than that currently observed due to removal of material with lead concentrations above 1000 ppm. It is noted that areas with elevated concentrations for other SSIP coincide with areas designated for excavation pursuant to the Interim Remedial Measures Plan. No excavation or removal involved any tampering with the existing clay/till unit beneath the site keeping in place this massive hydrogeologic protection feature.

The only metal found consistently above background levels in the fill is lead with the red lead paint sludge sample accounting for the majority of the other metals found above background concentrations. This red lead paint sludge was observed to be in small quantities mostly on the surface of the fill materials on site.

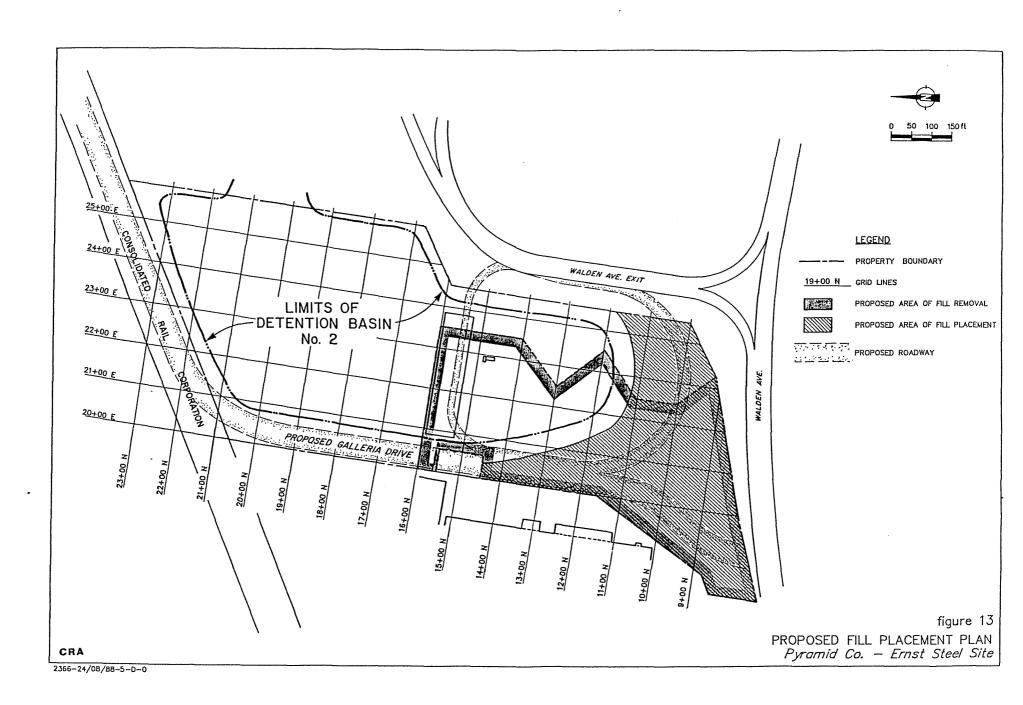
The Interim Remedial Measures Plan has already been established for the Site, i.e. removal and proper disposal of all soils with lead concentrations above 1000 ppm. It is recommended that a small volume of material from the isolated area in the southern portion of the Site also be excavated and appropriately disposed.

All fill materials with lead concentrations below 1000 ppm can remain on site. A fill handling proposal for this material is presented in the subsequent section of this report.

12.0 FILL HANDLING PROPOSAL

Based upon the fill characterization and additional lead analyses that have been performed for the Site, it is apparent that the fill material which does not exceed the established criteria requiring off-site disposal at a permitted waste facility can be secured on the Site in an environmentally sound manner. The proposed plan is to deposit this fill material into an area on the Site where it will not interfere with future plans for Site development. Based upon current plans, the only available area for the deposition of fill is in the vicinity of the proposed off-ramp from the Walden Avenue exit of the New York State Thruway. This off-ramp loops around south of the proposed detention basin No. 3. Figure 13 illustrates the general layout of the area including the roadways, detention basin and proposed disposal area.

The immediate interest in the southern half of the Site is to gain access to the underlying native soils which have been demonstrated to be clean (see Figure 9). At the same time, it is imperative that the plan for final Site remediation be kept in mind to insure that an overall acceptable plan of remediation is attained for the entire Site and not just the southern half. Consequently, although this proposed plan deals only with the fill material from the southern half of the Site, it is anticipated that the scope of the plan could be enlarged to accommodate the handling of



all fill material left on the Site which contains lead at concentrations of less than 1000 ppm.

The proposed plan to dispose of the fill from the southern half of the Site is presented in the following subsections of this report.

12.1 AREA PREPARATION

In order to dispose of the fill from the southern half of the Site, it will be necessary to overexcavate the area shown on Figure 13. Overexcavation is necessary because the proposed final grades for this area are not significantly different than the existing grades.

Therefore, in order to maintain sight lines and surface drainage, it will be necessary to excavate a pit of appropriate depth such that once fill is placed into the pit, the top of the fill would closely match the final proposed grade except for the placement of an overlying cap, the need for which will be discussed in subsequent sections.

The ultimate required depth of the pit/disposal area is dependent upon the following factors:

- Volume of fill to be disposed.
- Available area in which to construct the disposal area.
- Final grades required to accommodate off-ramp construction and surface drainage configuration.
- o Thickness of cap over fill.

In order to gain access to the proposed disposal area, the overlying fill material would be removed and placed onto the area shown for the construction of Detention Pond No. 3. The excavation of native soils from the disposal area would then proceed as expeditiously as possible to minimize the time that the fill materials lie in a disturbed state.

12.2 VOLUME OF FILL

The volume of fill to be disposed in the pit would be as follows:

If only fill from the southern half of the Site is disposed, the volume would be on the order of 13,000 cubic yards. The total fill volume in the southern half of the Site is 16,000 cubic yards. However, approximately 3,000 cubic yards is gravel from the parking area and roadway. This gravel material will be used in other areas of the Mall Project on an as needed basis.

If all fill including the northern half of the Site

(assumes all lead concentrations less than 1,000 ppm) be

placed in the disposal area, the volume would be

37,000 cubic yards (total fill volume estimated to be

present at the Site) less the 3,000 cubic yards of

gravel, less the 850+ cubic yards of fill material

already removed from the Site, less the remaining volume

of fill to be removed from the Site. Thus the maximum

volume of fill to be disposed would be somewhat less

than 33,000 cubic yards.

12.3 AVAILABLE DISPOSAL AREA

The proposed disposal area would extend across the southern site boundary as shown on Figure 13. A 50± foot buffer zone will be maintained between the disposal area and detention pond. Similarly a 10-foot buffer zone will be maintained between the disposal area and southern and western property boundaries. Beneath the off-ramp along the eastern property boundary, additional fill may be placed if needed.

If the entire on-site area shown on Figure 13 is used for waste disposal, approximately 170,000 square feet

of storage area is available. At this size, the average fill depth deposited in the disposal area would be approximately:

- ° 2.1 feet to accommodate 13,000 cubic yards
- ° 5.2 feet to accommodate 33,000 cubic yards.

If the area under proposed Galleria Drive is not included in the disposal area, the disposal area would be reduced to approximately 121,000 square feet which would result in the following impact on fill placement depth:

- ° 2.9 feet to accommodate 13,000 cubic yards
- ° 7.4 feet to accommodate 33,000 cubic yards.

Although there is an overall intent to minimize the depth of excavation, this must be weighed against the need to maintain access and accommodate Site conditions encountered.

12.4 EXCAVATION AND HAULING PROCEDURES

Excavation and hauling of native soils will be accomplished by pan scraper, loader and dump truck or other appropriate means.

Initial stripping of the fill from the disposal area will be accomplished by similar means or by use of a bulldozer. Whatever method is employed, precautions will be taken to minimize dust generation from fill material. Should dust be measured above 150 ug/m3 using air particulate monitoring equipment around the periphery of the excavation operation (i.e. within 200 feet of the excavation or within 50 feet of off-site boundaries) appropriate wetting of the fill will be undertaken to control fugitive dust generation. Measurements will be taken at hourly intervals during fill handling.

Haul routes will be wetted as necessary as is currently done.

12.5 SURFACE WATER CONTROL

During the excavating of native soil, berms will be constructed at the end of each working day and as needed at times of impending threat of rain to keep surface water from draining into the open excavation. This will insure that water accumulating in the excavation is only clean rain water that has fallen directly into the excavation and can therefore be pumped directly into surrounding surface water drainage ditches.

In order to maintain the integrity of the disposal area and consequently the cleanliness of the rainwater falling into the disposal area once fill placement into the disposal area has begun, the following protocols will be followed:

- ° fill will not be placed in the cell during rain events,
- at the conclusion of each day's fill placement or upon threat of impending rain, the exposed fill material will be covered with a thin layer of clay.

To prevent rainfall contact with freshly disturbed fill materials, the fill material initially scraped aside to gain access to the disposal area will be covered with 6 mil polyethylene and secured in place. This material will be the first material placed in the disposal area once the excavation is complete.

12.6 DISPOSAL AREA CONSTRUCTION

The soil beneath the Site has been typically characterized as clay although some silty layers were occasionally encountered and a layer of till was encountered at the monitoring well installed at the southwest corner of

the overhead crane (OW-3). Consequently, groundwater considerations are not expected to cause a concern at the proposed disposal area. Nonetheless, the following precautions will be taken:

- Sidewalls of the disposal area will be overexcavated and a 5-foot compacted clay plug will be installed to cut off the flow of groundwater from any perched water lens observed.
- Any identified clay discontinuity in the base or sidewalls of the disposal area will be overexcavated and a 2-foot layer of clay compacted in place. On-site clay is suitable for this purpose. Two samples submitted to a physical testing laboratory as part of the Phase II Site Investigation obtained remoulded permeabilities of 1.45x10-8 and 3.37x10-8 cm/sec.
- Due to the potential for concern regarding long term contact of groundwater with the fill following closure of the disposal area, the fill from the southern portion of the Site will be placed at the bottom of the disposal area. This material typically has very low lead concentration and would pose much less concern, if any at all, with groundwater contact.

Once all of the fill is in place, the entire disposal area will be capped with 12± inches of compacted clay from the Site. This in turn will be covered with 6± inches of topsoil or road bedding material and asphalt, as appropriate. In order to support any roadways, the fill material would necessarily be compacted as it is placed in accordance with good engineering judgment.

12.7 SUMMARY

Given that the fill materials left at the Site following completion of the waste pile remediation do not constitute hazardous waste, the proposed plan of on-site disposal is environmentally sound and is a responsive solution to the environmental concern.



APPENDIX A

WASTE CHARACTERIZATION SAMPLING RESULTS

INORGANIC ANALYSIS

Conestoga-Rovers & Associates Waterloo, Ontario, Canada

> Project #2366 Lab #66242

Matrix: Soil

August 9. 1988

The control of the second of the commentation of the control of th

Envirotest Laboratories	Lab	#	66242-001
CRA Consulting Engineers	#1		
Date Collected 7/6/88	Log	Date	7/8/88
Job # 2366			

Matr	ix:Soil		Units: mg/kg
1.	Aluminum	3545	13. Magnesium 1625
2.	Antimony	10.75 U	14. Manganese 769
3.	Arsenic	2.15 U	15. Mercury 0.14
4.	Barium	38.71	16. Nickel 39.78
5.	Beryllium	0.54 U	17. Potassium 559
6.	Cadmium	1.08 U	18. Selenium 1.08 U
7.	Calcium	5672	19. Silver 1.08 U
8.	Chromium	9.57	20. Sodium 538 U
9.	Cobalt	3.23 U	21. Thallium 10.75 U
10.	Copper	184	22. Vanadium 14.52
11.	Iron	17870	23. Zinc 506
12.	Lead	208	
	Cyanide	1.08 U	Percent Solids 93
	Phenol		

ICP interelement and background corrections applied? NO If yes, corrections applied before ___ or after ___ generation of raw data.

U = Sample analyzed, result less than the detection limit.

Envirotest Laboratories Lab # 66242-002 CRA Consulting Engineers #2 Date Collected 7/6/88 Log Date 7/8/88 Job # 2366

Matrix:Soil

1.	Aluminum		7340		13.	Magnesium	 11500	
2.	Antimony		10.20	U	14.	Manganese	 2810	
3.	Arsenic		2.04	U	15.	Mercury	 0.12	
4.	Barium		546		16.	Nickel	 67.24	
5.	Beryllium	1	1.63		17.	Potassium	 995	
6.	Cadmium		4.29		18.	Selenium	 1.02	U
7.	Calcium		43900		19.	Silver	 1.63	
8.	Chromium		2450		20.	Sodium	 3188	
9.	Cobalt		29.08		21.	Thallium	 10.20	U
10.	Copper		123		22.	Vanadium	 26.73	
11.	Iron		97800		23.	Zinc	 6090	
12.	Lead		334					
	Cyanide		2.04		Perc	ent Solids	 98	
	Phenol							

Units: mg/kg

ICP interelement and background corrections applied?]
If yes, corrections applied before ___ or after ___ generation of raw data.

U = Sample analyzed, result less than the dettection limit.

Envirotest Laboratories	Lab #	66242-003
CRA Consulting Engineers	#3	
Date Collected 7/6/88	Log Date	7/8/88
Job # 2366		

Matr	ix:Soil			Unit	s: mg/kg		
1.	Aluminum	10500		13.	Magnesium	 6610	
2.	Antimony	11.36	U	14.	Manganese	 682	
3.	Arsenic	2.27	U	15.	Mercury	 0.09	
4.	Barium	110		16.	Nickel	 25.91	
5.	Beryllium	0.68		17.	Potassium	 568	U
6.	Cadmium	1.14	U .	18.	Selenium	 1.14	U
7.	Calcium	19400		19.	Silver	 1.14	U
8.	Chromium	33.98		20.	Sodium	 568	Ū
9.	Cobalt	7.50		21.	Thallium	 11.36	U
10.	Copper	47.39		22.	Vanadium	 32.3	
11.	Iron	31300		23.	Zinc	 139	
12.	Lead	443					
	Cyanide	1.14		Perc	ent Solids	 88	
	Phenol						

ICP interelement and background corrections applied? NO If yes, corrections applied before ___ or after ___ generation of raw data.

U = Sample analyzed, result less than the detection limit.

Envirotest Laboratories	Lab #	66242-004
CRA Consulting Engineers	#4 ·	
Date Collected 7/6/88	Log Date	7/8/88
Job # 2366		

Matr	ix:Soil		Units: mg/kg	
1.	Aluminum	3410	13. Magnesium 544	U
2.	Antimony	10.87 U	14. Manganese 116	
3.	Arsenic	2.17 U	15. Mercury 0.09	
4.	Barium	55.43	16. Nickel 10.87	
5.	Beryllium	0.54 U	17. Potassium 544	U
6.	Cadmium	1.09 U	18. Selenium 1.09	U
7.	Calcium	3130	19. Silver 1.09	U
8.	Chromium	10.43	20. Sodium 544	U
9.	Cobalt	3.26 U	21. Thallium 10.87	U
10.	Copper	249	22. Vanadium 14.67	
11.	Iron	22200	23. Zinc 29.78	
12.	Lead	376		
	Cyanide	1.09 U	Percent Solids 92	
	Phenol		•	

ICP interelement and background corrections applied? NO If yes, corrections applied before ___ or after ___ generation of raw data.

U = Sample analyzed, result less than the detection limit.

Envirotest Laboratories	Lab #	66242-005
CRA Consulting Engineers	#5	
Date Collected 7/6/88	Log Date	7/8/88
Job # 2366		

Matrix:Soil	Units:	mg/kg

1.	Aluminum		5590		13.	Magnesium	 7380	
2.	Antimony		11.49	U	14.	Manganese	 3760	
3.	Arsenic		2.30	U	15.	Mercury	 0.09	U
4.	Barium		59.77		16.	Nickel	 10.80	
5.	Beryllium	a	0.57	U	17.	Potassium	 575	U
6.	$\mathtt{Cadmium}$		1.15	U	18.	Selenium	 1.15	U
7.	Calcium		37700		19.	Silver	 1.15	U
8.	Chromium		15.29		20.	Sodium	 575	U
9.	Cobalt		3.45	U	21.	Thallium	 11.49	U
10.	Copper		18.74		22.	Vanadium	 28.28	
11.	Iron		33600		23.	Zinc	 26.21	
12.	Lead		100	U				
	Cyanide		1.15	U	Perce	ent Solids	 87	
	Phenol							

ICP interelement and background corrections applied? NO If yes, corrections applied before ___ or after ___ generation of raw data.

U = Sample analyzed, result less than the detection limit.

ORGANIC ANALYSIS

Conestoga-Rovers & Associates Waterloo, Ontario, Canada

> Project #2366 Lab #66242

Matrix: Soil

August 9. 1988

1A VOLATILE ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Assoc. Lab Number: 66242-001

Project Name: #2366, Pyramid/Ernst Steel Site Date Collected: 7/6/88

Sample Location: #1 Date Received: 7/8/88

Matrix: Soil SW846 METHOD 8240

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kg
74-87-3	Chloromethane	11.U	78-87-5	1,2-Dichloropropane	5.4.4
74-83-9	Bromomethane	11.0	10061-02-6	Trans-1,3-Dichloropropene	5.4.0
75-01-4	Vinyl Chloride	11.U	79-01-6	Trichloroethene	5.4.0
75-00-3	Chloroethane	11.U	124-48-1	Dibromochloromethane	5.4 U
75-09-2	Methylene Chloride	17.B	79-00-5	1,1,2-Trichloroethane	5.4.U
67-64-1	Acetone	38.BJ	71-43-2	Benzene	5.4.0
75-15-0	Carbon Disulfide	5.4.0	10061-01-5	cis-1,3-Dichloropropene	5.4.0
75-35-4	1.1-Dichloroethene	5.4.U	110-75-8	2-Chloroethylvinylether	11.0
75-34-3	1.1-Dichloroethane	5.4.0	75-25-2	Bromoform	5.4.0
156-60-5	Trans-1,2-Dichloroethene	5.4.0	108-10-1	4-Methyl-2-Pentanone	11.U
67-66-3	Chloroform	3.6.J	591-78-6	2-Hexanone	11.0
107-02-2	1.2-Dichloroethane	5.4.0	127-18-4	Tetrachloroethene	5.4.0
78-93-3	2-Butanone	11.0	79-34-5	1.1.2,2-Tetrachloroethane	5.4.0
71-55-6	1.1.1-Trichloroethane	5.4.0	108-88-3	Toluene	5.4.0
56-23-6	Carbon Tetrachloride	5.4.U	108-90-7	Chlorobenzene	5.4.0
108-05-4	Vinyl Acetate	11.U	100-41-4	Ethylbenzene	5.4.0
75-27-4	Bromodichloromethane	5.4.0	100-42-5	Styrene	5.4.0
				Total Xylenes	5.4.0

Data Reporting Qualifiers

Value-If the result is a value greater than or equal to the detection limit, report the value.

- U-Indicates compound was analyzed for but not detected.

 Report the minimum detection limit for the sample with the U (e.g. 10U) based on necessary concentration/dilution action (This is not necessarily the instrument detection limit). The footnote should read: U-Compound was analyzed for but not detected. The number is the minimum attainable detection limit for the sample.
- J-Indicates an estimated value. This flag is used either when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed or when the mass spectral data indicated the presence of a compound that meets the identification criteria but the result is < than the specified detection limit but > than zero (e.g. 10J). If limit of detection is 10 ug/l and a concentration of 3.0 ug/l is calculated, report as 3J.
- C-This flag applies to pesticide parameters where the identification has been confirmed by GC/MS. Single component pesticides >= 10 ng/ul in the final extract should be confirmed by GC/MS.
- B-This flag is used when the analyte is found in the blank as well as a sample. It indicates possible/probable blank contamination and warns data the user to take appropriate action.
- Other-Other specific flags and footnotes may be required to properly define the results. If used, they must be fully described and such description attached to the data summary report.

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Associates

Lab Number: 66242-001

Project Name: #2366. Pyramid/Ernst Steel Site

Date Collected: 7/6/88

Sample Location: #1

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8270

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	uọ/kg
108-95-2	Pheno:	360.U	83-32-9	Acenaphthene	360.0
111-44-4	bis(-2-Chloroethyl)Ether	360.U	51-28-5	2,4-Dinitrophenol	360.U
95-57-8	2-Chlorophenol	360.U	100-02-7	4-Nitrophenol	360.U
541-73-1	1,3-Dichlorobenzene	360.U	132-64-9	Dibenzofuran	360.U
106-45-7	1,4-Dichlorobenzene	360.U	121-14-2	2,4-Dinitrotoluene	360.U
100-51-6	Benzyl Alcohol	360.U	606-20-2	2,5-Dinitrotoluene	360.U
95-50-1	1,2-Dichlorobenzene	360.U	84-66-2	Diethylphthalate	360.U
95-48-7	2-Methylphenol	360.0	7005-72-3	4-Chlorophenyl-phenlyether	360.U
39638-32-9	bis(2-chloroisopropyl)Ether	360.U	86-73-7	Fluorene	360.U
106-44-5	4-Methylphenol	360.U	100-01-6	4-Nitroaniline	360.U
621-64-7	N-Nitroso-Di-n-Propylamine	360.U	534-52-1	4.6-Dinitro-2-Methylphenol	360.U
67-72-1	Hexachloroethane	360.U	86-30-6	N-Nitrosodiohenvlamine (1)	- 360.U
98-95-3	Nitrobenzene	360.U	101-55-3	4-Bromophenyl-phenylether	360.U
78-59-1	Isophorone	360.0	119-74-1	Hexachlorobenzene	360.U
88-75-5	2-Nitrophenol	360.0	87-86-5	Pentachlorophenol	360.0
105-67-9	2.4-Dimethylphenol	360.U	85-01-8	Phenanthrene	360.U
65-85-0	Benzoic Acid	360.U	120-12-7	Anthracene	360.0
111-91-1	bis(-2-Chloroethoxy)Methane	360.8	84-74-2	Di-n-Butylphthalate	360.0
120-83-2	2,4-Dichlorophenol	360.0	205-44-0	Fluoranthene	360.U
120-82-1	1.2.4-Trichlorobenzene	369.0	129-00-0	Pyrene	360.U
91-20-3	Naphthalene	360.0	85-69-7	Butyibenzylphthalate	360.U
106-47-8	4-Chloroaniline	360.0	91-94-1	3.3'-Dichlorobenzidine	360.U
87-68-3	Hexachlorobutadiene	360.U	56-55-3	Benzo(a)Anthracene	360.0
59-50-7	4-Chloro-3-Methylphenol	360.U	117-81-7	bis(2-Ethylhexyl)Phthalate	140.J
91-57-6	2-Methylnaphthalene	369.U	218-01-9	Chrysene	360.U
77-47-4	Hexachlorocyclopentadiene	360.U	117-84-0	Di-n-Octyl Phthalate	360.U
88-06-2	2,4,6-Trichlorophenol	360.U	205-99-2	Benzo(b)Fluoranthene	360.U
95-95-4	2,4,5-Trichlorophenol	360.U	207-08-9	Benzo(k)Fluoranthene	360.U
91-58-7	2-Chloronaphthalene	360.U	50-32-8	Benzo(a)Pyrene	360.U
88-74-4	2-Nitroaniline	360.U	193-39-5	Indeno(1,2,3-cd)Pyrene	360.U
131-11-3	Dimethyl Phthalate	360.U	53-70-3	Dibenzo(a.h)Anthracene	360.8
208-96-8	Acenaphthylene	360.U	191-24-2	Benzo(g.h.i)Perylene	360.0
99-09-2	3-Nitroaniline	360.U		-	

⁽¹⁾⁻Cannot be separated from diphenylamine

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers Date Received: 7/8/88

Project: #2366, Pyramid/Ernst Steel Date Collected: 7/6/88

Laboratory Number: 66242-001 Matrix: Soil

Sample Location: #1 Method:SW846-8080

Pesticides/PCBs

CAS NO.	COMPOUND	ug/kg
319-84-6	Alpha-BHC	18.U
319-87-7	Beta-BHC	18.U
319-86-8	Delta-BHC	18.U
58-89-9	Gamma-BHC(Lindane)	18.U
76-44-8	Heptachlor	18.U
309-00-2	Aldrin	18.U
1024-57-3	Heptachlor Epoxide	18.U
959-98-8	Endosulfan I	18.U
60-57-1	Dieldrin	36.U
72-55-9	4.4'-DDE	36.U
72-20-8	Endrin	36.U
	Endosulfan II	36.U
72-54-8	4.4 -DDD	36.U
1031-07-8	Endosulfan Sulfate	36 . U
50-29-3	4.4'-DDT	36.U
72-43-5		180.U
53494-70-5	Endrin Ketone	36.U
57-74-9	Chlordane	180.U
8001-35-2	Toxaphene	360.U
12674-11-2	Arochlor-1016	180.U
11104-28-2	Arachlor-1221	180.U
11141-16-5	Arochlor-1232	180.U
53469-21-9	Arochlor-1242	180.U
12672- 29-6	Arochlor-1248	180.U
11097- 69-1	Arochlor-1254	360.U
11096-82-5	Arochlor-1260	360.U

1A VOLATILE ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Assoc.

Lab Number: 66242-002

Project Name: #2366, Pyramid/Ernst Steel Site

Date Collected: 7/6/88

Sample Location: #2

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8240

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/k
74-8 7-3	Chloromethane	10.U	78-87-5	1,2-Dichloropropane	5.4.0
74-83-9	Bromomethane	10.0	10051-02-6	Trans-1,3-Dichloropropene	5.4.0
75-01-4	Vinyl Chloride	10.U	79-01-6	Trichloroethene	5.4.0
75-00-3	Chloroethane	10.U	124-48-1	Dibromochloromethane	5.4 U
75-09-2	Methylene Chloride	22.B	79-00-5	1,1,2-Trichloroethane	5.4.0
67-64-1	Acetone	110.B	71-43-2	Benzene	5.4.U
75-15-0	Carbon Disulfide	5.1.0	10061-01-5	cis-1,3-Dichloropropene	5.1.0
75-35-4	1.1-Dichloroethene	5.1.0	110-75-8	2-Chloroethylvinylether	10.U
75-34-3	1.1-Dichloroethane	5.1.0	75-25-2	Bromoform	5.1.0
156-50-5	Trans-1,2-Dichloroethene	5.1.0	108-10-1	4-Methyl-2-Pentanone	10.0
67-66-3	Chloroform	3.5.J	591-78-6	2-Hexanone	10.U
107-02-2	1,2-Dichloroethane	5.1.0	127-18-4	Tetrachloroethene	5.1.0
78-93-3	2-Butanone	10.U	79-34-5	1,1,2,2-Tetrachloroethane	5.1.0
71-55-6	1,1,1-Trichloroethane	5.1.0	108-88-3	Toluene	5.1.0
56-23-6	Carbon Tetrachloride	5.1.0	108-90-7	Chlorobenzene	5.1.0
109-05-4	Vinvl Acetate	10.U	100-41-4	Ethvlbenzene	5.1.0
75-27-4	Bromodichloromethane	5.1.0	100-42-5	Styrene	5.1.0
				Total Xylenes	5.1.0

Data Reporting Qualifiers

Value-If the result is a value greater than or equal to the detection limit, report the value.

- U-Indicates compound was analyzed for but not detected. Report the minimum detection limit for the sample with the U (e.g. 10U) based on necessary concentration/dilution action (This is not necessarily the instrument detection limit). The footnote should read: U-Compound was analyzed for but not detected. The number is the minimum attainable detection limit for the sample.
- J-Indicates an estimated value. This flag is used either when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed or when the mass spectral data indicated the presence of a compound that meets the identification criteria but the result is (than the specified detection limit but) than zero (e.g. 10J). If limit of detection is 10 ug/l and a concentration of 3.0 ug/l is calculated, report as 3J.
- C-This flag applies to pesticide parameters where the identification has been confirmed by 6C/MS. Single component pesticides >= 10 ng/ul in the final extract should be confirmed by 6C/MS.
- B-This flag is used when the analyte is found in the blank as well as a sample. It indicates possible/probable blank contamination and warns data the user to take appropriate action.
- Other-Other specific flags and footnotes may be required to properly define the results. If used, they must be fully described and such description attached to the data summary report.

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Associates

Lab Number: 66242-002

Project Name: #2366, Pyramid/Ernst Steel Site

Date Collected: 7/6/89

Sample Location: #2

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8270

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kg	
108-95-2	Phenol Phenol	3 4 0.U	83-32-9	Acenaphthene	3 4 0.U	
111-44-4	bis(-2-Chloroethyl)Ether	340.U	51-28-5	2,4-Dinitrophenal	340.U	
95-57-8	2-Chlorophenol	3 4 0.U	100-02-7	4-Nitrophenol	340.U	
541-73-1	1.3-Dichlorobenzene	340.U	132-64-9	Dibenzofuran	340.U	
106-46-7	1,4-Dichlorobenzene	340.U	121-14-2	2,4-Dinitrotoluene	340.U	
100-51-6	Benzyl Alcohol	340.U	606-20-2	2,6-Dinitrotoluene	340.U	
95-50-1	1,2-Dichlorobenzene	340.U	84-66-2	Diethylphthalate	340.U	
95-48-7	2-Methylphenoi	340.U	7005-72-3	4-Chlorophenyl-phenlyether	340.U	
39638-32-9	bis(2-chloroisopropyl)Ether	3 4 0.U	86-73-7	Fluorene	340.U	
106-44-5	4-Methylphenol	250.J	100-01-6	4-Nitroaniline	340.U	
621-64-7	N-Nitroso-Di-n-Propylamine	3 4 0.U	534-52-1	4.6-Dinitro-2-Methylphenol	340.U	1
67-72-1	Hexachioroethane	340.U	86-30-6	N-Nitrosodiphenylamine (1)	340.U	
98-95-3	Nitrobenzene	340.0	101-55-3	4-Bromophenyl-phenylether	340.U	
78-59-1	Isophorone	3 4 0.U	118-74-1	Hexachiorobenzene	340.U	
88-75-5	2-Nitrophenol	340.U	87-86-5	Pentachlorophenol	340.U	
105-67-9	2.4-Dimethylphenol	340.0	85-01-8	Phenanthrene	3 4 0.U	
65-85-0	Benzoic Acid	340.0	120-12-7	Anthracene	340.U	
111-91-1	bis(-2-Chioroethoxy)Methane	340.U	84-74-2	Di-n-Butylphthalate	240.J	
120-83-2	2,4-Dichlorophenol	340.0	206-44-0	Fluoranthene	150.J	
120-82-1	1,2,4-Trichlorobenzene	340.U	129-00-0	Pyrene	340.U	
91-20-3	Naphthalene	340.U	85-68-7	Butylbenzylphthalate	580	
106-47-8	4-Chloroaniline	340.U	91-94-1	3,3'-Dichlorobenzidine	340.U	
87-68-3	Hexachlorobutadiene	340.U	56-55-3	Benzo(a)Anthracene	340.U	
59-50-7	4-Chloro-3-Methylphenol	340.U	117-81-7	bis(2-Ethylhexyl)Phthalate	3420	
91-57-6	2-Methylnaphthalene	340.U	218-01-9	Chrysene	340.U	
77-47-4	Hexachlorocyclopentadiene	340.U	117-84-0	Di-n-Octyl Phthalate	340.0	
88-06-2	2,4,6-Trichlorophenol	340.U	205-99-2	Benzo(b)Fluoranthene	340.U	
95-95-4	2,4,5-Trichlorophenol	340.U	207-08-9	Benzo(k)Fluoranthene	340.U	
91-58-7	2-Chloronaphthalene	340.U	50-32-8	Benzo(a)Pyrene	340.U	
88-74-4	2-Nitroaniline	3 40. U	193-39-5	Indeno(1.2.3-cd)Pyrene	340.U	
131-11-3	Dimethyl Phthalate	340.U	53-70-3	Dibenzo(a.h)Anthracene	340.U	
208-96-8	Acenaphthylene	3 40. U	191-24-2	Benzo(g,h,i)Pervlene	340.0	
99-09-2	3-Nitroaniline	340.U		- · ·		

⁽¹⁾⁻Cannot be separated from diphenylamine

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers Date Received: 7/8/88

Project: #2366, Pyramid/Ernst Steel Date Collected: 7/6/88

Laboratory Number: 66242-002 Matrix: Soil

Sample Location: #2 Method:SW846-8080

Pesticides/PCBs

CAS NO.	COMPOUND	ua/kg
319-84-6	Alpha-BHC	520.U
319-87-7	Beta-BHC	520.U
319-86-8	Delta-BHC	520.U
58-89-9	Gamma-BHC(Lindane)	520.U
76-44-8	Heptachlor	520.U
309-00-2	Aldrin	520.U
1024-57-3	Heptachlor Epoxide	520.U
959-98-8	Endosulfan I	520.U
60-57-1	Dielarin	1100.U
72-55-9	4.4'-DDE	1100.U
72-20-8	Endrin	1100.U
33213-65-9	Endosulfan II	1100.U
72-54-8	4.4'-DDD	1100.U
1031-07-8	Endosulfan Sulfate	1100.U
50-29-3	4.4'-DDT	1100.U
72-43-5	Methoxychlor	5400.U
53494-70-5	Endrin Hetone	1100.U
57-74-9	Chlordane	5400.U
8001-35-2	Toxaphene	11,000.U
12674-11-2	Arochlor-1016	5400.U
11104-28-2	Arochlor-1221	5400.U
11141-16-5	Arochlor-1232	5400.U
53469-21-9	Arochlor-1242	5400.U
12672-2 9-6	Arochlor-1248	5400.U
11097 -69-1	Arochlor-1254	17,000
11096-82-5	Arochlor-1260	11,000.U

1A VOLATILE ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Assoc.

Lab Number: 66242-003

Project Name: #2366. Pyramid/Ernst Steel Site

Date Collected: 7/6/88

Sample Location: #3

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8240

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kg
7 4- 87-3	Chloromethane	11.U	78-87-5	1.2-Dichloropropane	5.4.U
74-83-9	Bromomethane	11.0	10061-02-6	Trans-1.3-Dichloropropene	5.4.U
75-01-4	Vinyl Chloride	11.U	79-01-6	Trichloroethene	5.4.U
75-00-3	Chloroethane	11.U	124-48-1	Dibromochloromethane	5.4 U
75-09-2	Methylene Chloride	14.B	79-00-5	1,1,2-Trichloroethane	5.4.U
67-64-1	Acetone	110.B	71-43-2	Benzene	5.4.0
75-15-0	Carbon Disulfide	5.7.0	10051-01-5	cis-1,3-Dichloropropene	5.7.0
75-35-4	1,1-Dichloroethene	5.7.0	110-75-8	2-Chloroethylvinylether	11.U
75-34-3	1.1-Dichloroethane	5.7.0	75-25-2	Bronoform	5.7.0
156-60-5	Trans-1,2-Dichloroethene	5.7.0	108-10-1	4-Methyl-2-Pentanone	11.0
67-66-3	Chloroform	3.7.J	591-78-6	2-Hexanone	11.0
107-02-2	1.2-Dichloroethane	5.7.0	127-18-4	Tetrachloroethene	5.7.U
78-93-3	2-Butanone	11.U	79-34-5	1,1,2,2-Tetrachloroethane	5.7.U
71-55-6	1,1,1-Trichloroethane	5.7.8	108-88-3	Toluene	5.7.0
56-23-5	Carbon Tetrachloride	5.7.0	108-90-7	Chlorobenzene	5.7.U
108-05-4	Vinvl Acetate	11.0	100-41-4	Ethylbenzene	5.7.0
75-27-4	Bromodichloromethane	5.7.U	100-42-5	Styrene	5.7.0
				Total Xvlenes	5.7.0

Data Reporting Qualifiers

Value-If the result is a value greater than or equal to the setection limit, report the value.

- U-Indicates compound was analyzed for but not detected.
 Report the minimum detection limit for the sample with
 the U (e.g. 10U) based on necessary concentration/dilution action (This isenot necessarily the instrument
 detection limit). The footnote should read: U-Compound
 was analyzed for but not detected. The number is the
 minimum attainable detection limit for the sample.
- J-Indicates an estimated value. This flag is used either when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed or when the mass spectral data indicated the presence of a compound that meets the identification criteria but the result is < than the specified detection limit but > than zero (e.g. 10J). If limit of detection is 10 ug/l and a concentration of 3.0 ug/l is calculated, report as 3J.
- C-This flag applies to pesticide parameters where the identification has been confirmed by 6C/MS. Single component pesticides >= 10 ng/ul in the final extract should be confirmed by 6C/MS.
- B-This flag is used when the analyte is found in the blank as well as a sample. It indicates possible/probable blank contamination and warns data the user to take appropriate action.
- Other-Other specific flags and footnotes may be required to properly define the results. If used, they must be fully described and such description attached to the data summary report.

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Associates Lab Number: 66242-003

Project Name: #2366, Pyramid/Ernst Steel Site Date Collected: 7/6/88

Sample Location: #3 Date Received: 7/8/88

Matrix: Soil SW846 METHOD 8270

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kg
108-95-2	Phenol	380.0	83-32-9	Acenaphthene	380.U
111-44-4	bis(-2-Chloroethyl)Ether	380.U	51-28-5	2,4-Dinitrophenal	380.U
95-57-8	2-Chlorophenol	360'N	100-02-7	4-Nitrophenol	380.U
541-73-1	1.3-Dichlorobenzene	380.N	132-64-9	Dibenzofuran	380.U
106-46-7	1.4-Dichlorobenzene	380.U	121-14-2	2,4-Dinitrotoluene	380.U
100-51-6	Benzyl Alcohol	380.U	606-20-2	2,6-Dinitrotoluene	380.U
95-50-1	1,2-Dichlorobenzene	380.U	84-66-2	Diethylohthalate	380.U
95-48-7	2-Methylphenol	380.U	7005-72-3	4-Chlorophenyl-phenlyether	380.U
39638-32-9	bis(2-chloroisopropyl)Ether	380.0	86-73-7	Fluorene	380.U
106-44-5	4-Methylphenol	380.U	100-01-6	4-Nitroaniline	380.0
621-64-7	N-Nitroso-Di-n-Propylamine	380.0	534-52-1	4,6-Dinitro-2-Methylphenol	380.U
67-72-1	Hexachloroethane	380.U	86-30-6	N-Nitrosodiphenylamine (1)	380.U
99- 95-3	Nitrobenzene	380.U	101-55-3	4-Bromophenyl-phenylether	380.U
78-59-1	Isophorone	380.U	118-74-1	Hexachiorobenzene	380.U
88-75-5	2-Nitrophenol	380.U	87-86-5	Pentachloropheno!	380.U
105-67-9	2.4-Dimethylohenol	380.U	85-01-8	Phenanthrene	150.J
65-85- 0	Benzoic Acid	380.0	120-12-7	Anthracene	380.U
111-91-1	bis(-2-Chloroethoxy)Methane	380.U	84-74-2	Di-n-Butylohthalate	380.U
120-83-2	2.4-Dichlorophenol	380.0	206-44-0	Fluorantmene	240.J
120-82-1	1.2.4-Trichlorobenzene	380.U	129-00-0	Pvrene	380.U
91-20-3	Naphthalene	280.N	85-68-7	Butylbenzylphthalate	380.U
106-47-8	4-Chloroaniline	380.U	91-94-1	3,3'-Dichlorobenzidine	380.U
87-68-3	Hexachlorobutadiene	380.U	56-55-3	Benzo(a)Anthracene	380.U
59-50-7	4-Chloro-3-Methylphenol	380.U	117-81-7	bis(2-Ethylhexyl)Phthalate	210.J
91-57-6	2-Methylnaphthalene	350.J	218-01-9	Chrysene	220.J
77-47-4	Hexachlorocyclopentadiene	380.U	117-84-0	Di-n-Octyl Phthalate	380.U
88-06-2	2.4,6-Trichlorophenol	380.U	205-99-2	Benzo(b)Fluoranthene	190.J
95-95-4	2.4,5-Trichlorophenol	380.U	207-08-9	Benzo(k)Fluoranthene	380.U
91-58-7	2-Chlosonachthalene	380.U	50-32-8	Benzo(a)Pyrene	150.J
88-74-4	2-Nitroaniline	380.U	193-39-5	Indeno(1,2,3-cd)Pyrene	150.J
131-11-3	Dimethyl Phthalate	380.U	53-70-3	Dibenzo(a,h)Anthracene	380.0
208-96-8	Acenaphthylene	380.U	191-24-2	Benzo(g.h.i)Perylene	380.U
99-09-2	3-Nitroaniline	380.U			

⁽¹⁾⁻Cannot be separated from diphenylamine

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers Date Received: 7/8/88

Project: #2366. Pyramid/Ernst Steel Date Collected: 7/6/88

Laboratory Number: 66242-003 Matrix: Soil

Sample Location: #3 Method:SW846-8080

Pesticides/PCBs

CAS NO.	COMPOUND	ua/kg
319-84-6	Alpha-BHC	39.U
319-87-7	Beta-BHC	39.U
319-86-8	Delta-BHC	39.U
58-89-9	Gamma-BHC(Lindane)	39.U
76-44-8	Heotachlor	39.U
309-00-2	Aldrin	39.U
1024-57-3	Heptachlor Epoxide	39.U
959-98-8	Endosulfan I	39.U
60-57-1	Dieldrin	77.U
72-55-9	4.4'-DDE	77.U
72-20-8	Endrin	77.U
33213-65-9	Endosulfan Il	77.U
72-54-8	4.4 -DDD	77.U
1031-07-8	Endosulfan Sulfate	77.U
50-29-3	4,4'-DDT	77.U
72-43-5	Methoxychlor	390.U
53494-70-5	Endrin Ketone	77.U
57-74-9	Chlordane	390.U
8001-35-2	Toxaphene	770.U
12674-11-2	Arachlor-1016	390.U
11104-28-2	Arochlor-1221	390.U
11141-16-5	Arochlor-1232	390.U
53469-21 -9	Arochlor-1242	390.U
12672-2 9-6	Arochlor-1248	390.U
11097 -69-1	Arochlor-1254	770.U
11096- 82-5	Arochlor-1260	770.U

1A VOLATILE ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Assoc.

Lab Number: 66242-004

Project Name: #2366, Pyramid/Ernst Steel Site

Date Collected: 7/6/88

Sample Location: #4

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8240

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kç
74-87-3	Chloromethane	11.U	78-87-5	1.2-Dichloropropane	5.4.U
74-83-9	Bromomethane	11.0	10061-02-6	Trans-1,3-Dichloropropene	5.4.0
75-01-4	Vinyl Chloride	11.0	79-01-6	Trichloroethene	5.4.0
75-00-3	Chloroethane	11.0	124-48-1	Dibromochloromethane	5.4 U
75-09-2	Methylene Chloride	21.8	79-00-5	1,1,2-Trichloroethane	5.4.0
67-64-1	Acetone	5.0.J	71-43-2	Benzene	5.4.0
75-15-0	Carbon Disulfide	5.4.U	10061-01-5	cis-1,3-Dichloropropene	5.4.0
75-35-4	1,1-Dichloroethene	5.4.0	110-75-8	2-Chloroethylvinylether	11.0
75-34-3	1.1-Dichloroethane	5.4.U	75-25-2	Bromoform	5.4.0
156-60-5	Trans-1,2-Dichloroethene	5.4.U	108-10-1	4-Methyl-2-Pentanone	11.0
67-66-3	Chloroform	4.1.J	591-78-6	2-Hexanone	11.0
107-02-2	1.2-Dichloroethane	5.4.U	127-18-4	Tetrachloroethene	5.4.0
78-93-3	2-Butanone	11.U	79-34-5	1.1.2.2-Tetrachloroethane	5.4.0
71-55-6	1.1.1-Trichloroethane	5.4.0	108-89-3	Toluene	5.4.0
56-23-5	Carbon Tetrachloride	5.4.U	108-90-7	Chlorobenzene	5.4.U
108-05-4	Vinyl Acetate	11.U	100-41-4	Ethvibenzene	5.4.0
75-27-4	Bromodichloromethane	5.4.U	100-42-5	Styrene	5.4.0
				Total Xylenes	5.4.0

Data Reporting Qualifiers

Value-If the result is a value greater than or equal to the detection limit, report the value.

- U-Indicates compound was analyzed for but not detected. Report the minimum detection limit for the sample with the U (e.g. 10U) based on necessary concentration/dilution action (This is not necessarily the instrument detection limit). The footnote should read: U-Compound was analyzed for but not detected. The number is the minimum attainable detection limit for the sample.
- J-Indicates an estimated Value. This flag is used either when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed or when the mass spectral data indicated the presence of a compound that meets the identification criteria but the result is < than the specified detection limit but > than zero (e.g. 10J). If limit of detection is 10 ug/l and a concentration of 3.0 ug/l is calculated, report as 3J.
- C-This flag applies to pesticide parameters where the identification has been confirmed by GC/MS. Single component pesticides >= 10 ng/ul in the final extract should be confirmed by GC/MS.
- B-This flag is used when the analyte is found in the blank as well as a sample. It indicates possible/probable blank contamination and warns data the user to take appropriate action.
- Other-Other specific flags and footnotes may be required to properly define the results. If used, they must be fully described and such description attached to the data summary report.

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Associates Lab Number: 66242-004

Project Name: #2366. Pyramid/Ernst Steel Site Date Collected: 7/6/88

Sample Location: #4 Date Received: 7/8/88

Matrix: Soil SW846 METHOD 8270

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kg
108-95-2	Phenol	360.U	83-32-9	Acenaphthene	360.U
111-44-4	bis(-2-Chloroethyl)Ether	360.U	51-28-5	2,4-Dinitrophenol	360.U
95-57-8	2-Chlorophenol	360.U	100-02-7	4-Nitrophenol	360.U
541-73-1	1.3-Dichlorobenzene	360.U	132-64-9	Dibenzofuran	770
106-46-7	1.4-Dichlorobenzene	360.U	121-14-2	2,4-Dinitrotoluene	360.U
100-51-6	Benzyl Alcohol	360.0	606-20-2	2,6-Dinitrotoluene	360.U
95-50-1	1,2-Dichlorobenzene	360.U	84-66-2	Diethylphthalate	360.U
95-48-7	2-Methylphenol	360.U	7005-72-3	4-Chlorophenyl-phenlyether	360.U
39638-32-9	bis(2-chloroisopropyl)Ether	360.U	86-73-7	Fluorene	360.U
106-44-5	4-Methylphenol	360.U	100-01-6	4-Nitroaniline	360.U
621-64-7	N-Nitroso-Di-n-Propylamine	360.U	534-52-1	4,6-Dinitro-2-Methylphenol	360.U
67-72-1	Hexachloroethane	360.U	86-30-6	N-Nitrosodiphenylamine (1)	360.0
98-95-3	Nitrobenzene	360.U	101-55-3	4-Bromophenyl-phenylether	360.U
78-59-1	Isophorone	360.U	118-74-1	Hexachiorobenzene	360.U
89-75-5	2-Nitrophenol	360.U	87-86-5	Pentachlorophenol	360.U
105-67-9	2.4-Dimethylphenol	360.U	85-01-8	Phenanthrene	850
65-85-0	Benzoic Acid	360.U	120-12-7	Anthracene	360.0
111-91-1	bis(-2-Chloroethoxy)Methane	360.U	84-74-2	Di-n-Butylphthalate	360.U
120-83-2	2,4-Dichlorophenoi	360.U	206-44-0	Fluoranthene	360.U
120-82-1	1,2,4-Trichlorobenzene	360.U	129-00-0	Pyrene	160.J
91-20-3	Naphthalene	1260	85-68-7	Buty!penzylphthalate	360.U
106-47-8	4-Chloroaniline	360.U	91-94-1	3.3 -Dichlorobenzidine	360.U
87-68- 3	Hexachlorobutadiene	360.U	56-55-3	Benzo(a)Anthracene	120.J
59-50-7	4-Chloro-3-Methylphenol	360.U	117-81-7	bis(2-Ethylhexyl)Phthalate	270.J
91-57-6	2-Methylnaphthalene	3330	218-01-9	Chrysene	190.J
77-47-4	Hexachlorocyclopentadiene	360.0	117-84-0	Di-n-Octyl Phthalate	360.U
88-06-2	2,4,6-Trichlorophenol	360.U	205-99-2	Benzo(b)Fluoranthene	360.U
95-95-4	2,4,5-Trichlorophenol	360.U	207-08-9	Benzo(k)Fluoranthene	360.U
91-58-7	2-Chlosonaphthalene	360.U	50-32-8	Benzo(a)Pyrene	360.U
88-74-4	2-Nitroaniline	360.U	193-39-5	Indeno(1,2,3-cd)Pyrene	360.0
131-11-3	Dimethyl Phthalate	360.U	53-70-3	Dibenzo(a.h)Anthracene	360.U
208-96-8	Acenaphthylene	360.U	191-24-2	Benzo(g,h,i)Perylene	360.U
99-09-2	3-Nitroaniline	360.U		- · · · · · ·	

⁽¹⁾⁻Cannot be separated from diphenylamine

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers Date Received: 7/8/88

Project: #2366, Pyramid/Ernst Steel Date Collected: 7/6/88

Laboratory Number: 66242-004 Matrix: Soil

Sample Location: #4 Method:SW846-8080

Pesticides/PCBs

CAS NO.	COMPOUND	uạ/kạ
319-84-6 319-87-7 319-86-8 58-89-9 76-44-8 309-00-2 1024-57-3 959-98-8 60-57-1 72-55-9 72-20-8 33213-65-9 72-54-8 1031-07-8 50-29-3 72-43-5 53494-70-5 57-74-9 8001-35-2 12674-11-2 11104-28-2 11141-16-5 53469-21-9 12672-29-6	Alpha-BHC Beta-BHC Delta-BHC(Lindane) Heptachlor Aldrin Heptachlor Epoxide Endosulfan I Dieldrin 4.4'-DDE Endrin Endosulfan II 4.4'-DDD Endosulfan Sulfate 4.4'-DDT Methoxychlor Endrin Ketone Chlordane Toxaphene Arochlor-121 Arochlor-1242 Arochlor-1248	1.8.U 1.8.U 1.8.B.U 1.
11097-69-1 11096-82-5	Arochlor-1254 Arochlor-1260	36.U 36.U

1A VOLATILE ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Assoc.

Lab Number: 66242-005

Project Name: #2365. Pyramid/Ernst Steel Site

Date Collected: 7/6/88

Sample Location: #5

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8240

CAS NO.	COMPOUND	ug/kg	CAS NO.	COMPOUND	ug/kg
74-87-3	Chloromethane	11.U	78-87-5	1,2-Dichloropropane	5.4.U
74-83-9	Bromomethane	11.0	10061-02-6	Trans-1,3-Dichloropropene	5.4.U
75-01-4	Vinyl Chloride	11.0	79-01-6	Trichloroethene	5.4.0
75-00-3	Chloroethane	11.U	124-49-1	Dibromochloromethane	5.4 U
75-09-2	Methylene Chloride	24.8	79-00-5	1,1,2-Trichloroethane	5.4.0
67-64-1	Acetone	46	71-43-2	Benzene	1.5.J
75-15-0	Carbon Disulfide	5.7.0	10061-01-5	cis-1,3-Dichloropropene	5.7.U
75-35-4	1,1-Dichloroethene	5.7.0	110-75-8	2-Chloroethylvinylether	11.0
75-34-3	1,1-Dichloroethane	5.7.0	75-25-2	Bromeform	5.7.0
156-60-5	Trans-1,2-Dichloroethene	5.7.8	108-10-1	4-Methyl-2-Pentanone	11.U
67-66-3	Chloroform	2.7.J	591-78-6	2-Hexanone	11.0
107-02-2	1.2-Dichloroethane	5.7.U	127-18-4	Tetrachloroethene	5.7.0
78-93-3	2-Butanone	11.0	79-34-5	1,1,2,2-Tetrachloroethane	5.7.0
71-55-6	1.1.1-Trichloroethane	5.7.0	108-88-3	Toluene	5.7.0
56-23-6	Carbon Tetrachloride	5.7.0	108-90-7	Chlorobenzene	5.7.0
108-05-4	Vinyl Acetate	11.0	100-41-4	Ethvibenzene	5.7.0
75-27-4	Bromodichloromethane	5.7.0	100-42-5	Styrene	5.7.0
				Total Xylenes	5.7.0

Data Reporting Qualifiers

Value-If the result is a value greater than or equal to the detection limit, report the value.

- U-Indicates compound was analyzed for but not detected. Report the minimum detection limit for the sample with the U (e.g. 10U) based on necessary concentration/dilution action (This is not necessarily the instrument detection limit). The footnote should read: U-Compound was analyzed for but not detected. The number is the minimum attainable detection limit for the sample.
- J-Indicates an estimated value. This flag is used either when estimating a concentration for tentatively identified compounds where a 1:1 response is assumed or when the mass spectral data indicated the presence of a compound that meets the identification criteria but the result is < than the specified detection limit but > than zero (e.g. 10J). If limit of detection is 10 ug/l and a concentration of 3.0 ug/l is calculated, report as 3J.
- C-This flag apolies to pesticide parameters where the identification has been confirmed by GC/MS. Single component pesticides >= 10 ng/ul in the final extract should be confirmed by GC/MS.
- B-This flag is used when the analyte is found in the blank as well as a sample. It indicates possible/probable blank contamination and warns data the user to take appropriate action.
- Other-Other specific flags and footnotes may be required to properly define the results. If used, they must be fully described and such description attached to the data summary report.

ORSANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers & Associates

Lab Number: 66242-005

Project Name: #2366. Pyramid/Ernst Steel Site

Date Collected: 7/6/88

Sample Location: #5

Date Received: 7/8/88

Matrix: Soil

SW846 METHOD 8270

CAS NO.	COMPOUND	ug/kg	CAS NG.	COMPOUND	ug/kg
108-95-2	Phenol	380.U	83-32-9	Acenaphthene	380.U
111-44-4	bis(-2-Chloroethyl)Ether	380.U	51-28-5	2.4-Dinitrophenol	380.U
95-57-8	2-Chlorophenol	380.U	100-02-7	4-Nitrophenol	380.U
541-73-1	1,3-Dichlorobenzene	380.U	132-64-9	Dibenzofuran	380.U
106-46-7	1,4-Dichlorobenzene	390.U	121-14-2	2,4-Dinitrotoluene	380.U
100-51-6	Benzyl Alcohol	380.U	606-20-2	2,6-Dinitrotoluene	380.U
95-50-1	1,2-Dichlorobenzene	380.U	84-66-2	Diethylphthalate	380.U
95-48-7	2-Methylohenol	J80.U	7005-72-3	4-Chlorophenyl-phenlyether	380.U
39638-32-9	bis(2-chloroisopropyl)Ether	380.U	86-73-7	Fluorene	380.U
106-44-5	4-Methylphenol	380.U	100-01-6	4-Nitroaniline	380.U
621-64-7	N-Nitroso-Di-n-Propylamine	380.U	534-52-1	4.6-Dinitro-2-Methylphenol	380.U
67-72-1	Hexachloroethane	380.0	86-30-6	N-Nitrosodiphenylamine (1)	380.U
98-95-3	Nitrobenzene	3 8 0.U	101-55-3	4-Bromochenyl-phenylether	380.U
78-59-1	Isophorone	380.0	118-74-1	Hexachiorobenzene	380.U
88-75-5	2-Nitrophenol	380.U	87-86-5	Pentachlorophenol	380.U
105-67-9	2.4-Dimethylphenol	380.8	85-01-8	Pnenanthrene	380.U
£5-85-0	Benzoic Acid	380.0	120-12-7	Anthracene	380.U
111-91-1	bis:-2-Chloroethoxy)Methane	3 8 0.0	84-74-2	Di-n-Butylphtnalate	380.0
120-83-2	2,4-Dichlorophenol	380.0	206-44-0	Fluoranthene	380.0
120-83-1	1,2,4-Trichlorobenzene	380.U	129-00-0	Pyrene	380.U
91-20-3	Naphthalene	380.0	85-68-7	Butylbenzylphthalate	380.U
106-47-8	4-Chloroaniline	380.U	91-94-1	3.3'-Dichlorobenzidine	380.U
87-68-3	Hexachlorobutadiene	380.U	56-55-3	Benzo(a)Anthracene	380.U
59-5 0-7	4-Chloro-3-Methylphenol	3 0 0.U	117-81-7	bis(2-Ethylhexyl)Phthalate	240.J
91-57-6	2-MethyInaphthalene	380.U	218-01-9	Chrysene	380.U
77-47-4	Hexachlorocyclopentadiene	380.U	117-84-0	Di-n-Octyl Phthalate	380.U
88-06-2	2,4.6-Trichlorophenol	380.U	205-99-2	Benzo(b)Fluoranthene	380.U
95-95-4	2,4,5-Trichlorophenol	380.U	207-08-9	Benzo(k)Fluoranthene	380.U
91-58-7	2-Chloronaphthalene	380.U	50-32-8	Benzo(a)Pyrene	380.U
88-74-4	2-Nitroaniline	380.8	193-39-5	Indeno(1.2.3-cd)Pyrene	380.U
131-11-3	Dimethyl Phthalate	380.U	53-70-3	Dibenzo(a,h)Anthracene	380.U
208-96-8	Acenaphthylene	380.U	191-24-2	Benzo(g,h,i)Pervlene	380.U
99-09-2	3-Nitroaniline	380.U			

⁽¹⁾⁻Cannot be separated from diphenylamine

ORGANICS ANALYSIS DATA SHEET

Client Name: Conestoga-Rovers Date Received: 7/8/88

Project: #2366, Pyramid/Ernst Steel Date Collected: 7/6/88

Laboratory Number: 66242-005 Matrix: Soil

Sample Location: #5 Method:SW846-8080

Pesticides/PCBs

CAS NO.	COMPOUND	ug/kg
319-84-6	Alpha-BHC	58.U
319-87-7	Beta-BHC	58.U
319-86-8	Delta-BHC	58.U
58-89-9	Gamma-BHC(Lindane)	58.U
76-44-8	Heptachlor	58.U
309-00-2	Aldrin	58.U
1024-57-3	Heptachlor Epoxide	58.U
959-98-8	Endosulfan I	58.0
60-57-1	Dieldrin	120.U
72-55-9	4.4'-DDE	120.U
72-20-8	Endrin	120.U
33213- 65-9	Endosulfan II	120.U
71-54-8	4.4'-DDD	120.U
1031-07-8	Endosulfan Sulfate	120.U
50-29-3	4.4'-DDT	120.U
72-43-5	Methoxychlor	580.U
53494-70-5	Endrin ketone	120.U
57-74-9	Chlordane	580.U
8001-35-2	Toxaphene	1200.U
12674-11-2	Arochlor-1016	580.U
11104-28-2	Arochlor-1221	580.U
11141-16-5	Arochlor-1232	580.U
53469-21 -9	Arochlor-1242	580.U
12672- 29-6	Arochlor-1248	580.U
11097 -69-1	Arochlor-1254	1200.U
11096- 82-5	Arochlor-1260	1200.U

MEMO



To:

B. Clegg/K. Schmidtke

Reference No. 2366

From:

T. Misercola

Date:

9/21/88

Re:

Reissue of Memo Dated 8/24/88

Discussion of Results for Pyramid/Ernst Steel

The following memo details as assessment of analytical results reported by Envirotest for soil samples collected from the Pyramid Site. The samples submitted for analysis consisted of the following:

Matrix:

Soil/Fill

Investigative Samples:

5

All samples were submitted for HSL volatile organics, semi-volatile organics, metals and cyanide. CLP-RAS Methods for Organic and Inorganic Analyses were used.

The QA/QC criteria by which these data have been assessed are outlined in the Statement of Work (SOW) for Organic and Inorganic Analyses for Contract Laboratory Programs.

Based on review of this data set and related quality control data, the following are noted:

1. Sample Holding Time

Based on criteria outlined in Functional Guidelines for Evaluating Organic and Inorganic Analysis prepared by the USEPA, the following holding time requirements have been established for the Contract Lab Program analyses:

VOA (solids)

10 days from verified time of sample

receipt

Semi-Volatiles (solids)

10 days from verified time of sample

receipt

Pesticides/PCBs (solids)	10 days from verified time of sample receipt till extraction
Metals (solids)	6 months prior to analysis
Mercury (solids)	28 days prior to analysis
Cyanide (solids)	14 days prior to distillation

By comparing the sampling dates of all samples on the Chain of Custody and the actual dates of sample receipt for the dates of extraction and/or analysis, it is noted that all samples were handled properly in regards to holding time requirements. However, it should be noted that in the laboratory data package there was no mention as to the date of the distillation for the cyanide analyses for all five soil samples.

Action: Until notification from laboratory of actual dates of distillation are known, all results for cyanide shall be flagged as follows:

Sample ID = xxxx-oooo-yyy

x = project numbero = time of collectiony = sample number

Sample ID No.	Result (mg/kg)
2366-1420-001	1.08 UJ
2366-1440-002	2.04 J
2366-1515-003	1.14 J
2366-1515-004	1.09 UJ
2366-1600-005	1.15 UJ

UJ = The material was analyzed for, but was not detected.

The associated value is an estimate and may be inaccurate or inprecise.

J = The associated value is an estimated value

2. Spike Recoveries and Surrogate Recoveries

Laboratory performance on individual samples is established by spiking activities. All samples submitted for volatile, semivolatile and pesticides/PCB analysis were spiked with surrogate compounds prior to sample preparation.

A. <u>Discussion of Results - Organics</u>

All samples submitted for VOA and pesticide/PCB analyses yielded surrogate recoveries within the control limits established by CLP-RAS protocols. Therefore, no qualifications of the VOA data is required on this basis.

However, problems did occur in the semi-volatile surrogate recoveries. All surrogate in the base neutral fraction gave acceptable recoveries but four of the five samples for the acid fraction gave at least two surrogates out of the required control limit.

Action: The samples should have been, by contract requirements, re-extracted and reanalyzed. By not doing so, the laboratory failed to perform satisfactorily. However, the following qualifiers should be noted for samples 2366-02 to 2366-05 for use of this data:

- 2366-1440-002 (1) positive results for acid extractables are flagged as estimated (J)
 - (2) negative results (less than detection limit) are useable because surrogate recovery was less than 10 percent.
- 2366-1515-003 (1) positive results for acid extractables are flagged as estimated (J).
 - (2) negative results (less than detection limit) are unusable because surrogate recovery was less than 10 10 percent.
- 2366-1515-004 (1) positive results for acid extractables are flagged as estimates (J).

- (2) negative results (results less than detection limit) are flagged with the sample quantitation number classified as estimated (UJ).
- 2366-1600-005 (1) positive results for acid extractables are flagged as estimated (J).
 - (2) negative results (less than detection limit) are unusable because surrogate recovery was less than 10 percent.

B. <u>Discussion of Results - Inorganics - Matrix Spike Analysis</u>

The matrix spike sample analysis provides information about the effect of sample matrix on the digestion and measurement methodology.

Acceptable spike recovery limits for metal analyses is established as 75-125 percent as indicated in "Functional Guidelines for Evaluating Inorganic Analyses". The following spike recoveries for the metals listed failed to achieve that criteria in sample 2366-1420-001.

Metal	Percent Recovery	•
Lead	0	
Mercury	60	
Nickel	71	
Selenium	68	

In addition, poor spike recovery was found in copper and zinc analysis. However, Functional Guidelines for Evaluating Inorganic Analysis states that spike recovery limits are not acceptable when sample concentration for the analytes of interest exceed the spike concentration by at least a factor of 4 or more. Action: The following qualifications are made for those metals out of compliance for spike recovery:

Sample I.D. No.	<u>Metal</u>	Result
2366-1420-001	Lead Mercury Nickel Selenium	208 J 0.14 J 39.78 J 1.08 UJ
2366-1440-002	Lead Mercury Nickel Selenium	334 J 0.12 J 67.24 J 1.02 UJ
2366-1515-003	Lead Mercury Nickel Selenium	443 J 109 J 25.91 J 1.14 UJ
2366-1515-004	Lead Mercury Nickel Selenium	376 J 109 J 10.87 J 1.09 UJ
2366-1600-005	Lead Mercury Nickel Selenium	100 UJ 109 UJ 10.80 J 1.15 UJ

J = Associated value is an estimated quantity
 UJ = The material was analyzed for, but not detected.
 The associated value is an estimate and may be inaccurate or inprecise.

3. Laboratory/Reagent Blank Analysis

The assessment of results on blank analyses is for the purpose of determining the existence and magnitude of contamination problems. Laboratory reagent blanks must contain less than the Contract Required Detection Limit (CRDL) for all HSL compounds and less than five times the CRDL of methylene chloride, acetone, toluene and 2-butanone.

All blanks submitted for HSL metals and VOAs met those requirements. However, although no CRDL existed for 1-propanol, high levels of that compound were detected in the following laboratory reagent blanks:

<u>Date</u>	Concentration (ug/l)
6/30/88	5 9
7/08/88	7 5
7/14/88	70

No positive results for 1-propanol are reported unless the concentration in the sample exceeds five times the amount in the blank. Correction action at this point would be to flag all data for 1-propanol in all five samples as non-detected and adjust the sample quantitation limit to the value reported in the sample and flag the limit as estimated (UJ):

ID Number	Results (ug/kg)	Detection Limit			
2366-1420-001	ND	25 UJ			
2366-1440-002	ND	26 UJ			
2366-1515-003	ND	9.1 UJ			
2366-1515-004	ND	63 UJ			
2366-1600-005	ND	37 UJ			

UJ = The material was analyzed for, but not detected. The associated value is an estimate and may be inaccurate or inprecise.

4. Continuing Calibration

"Laboratory Functional Guidelines for Evaluating Organic Analyses" specifies that all systems performance calibration compounds (SPCC's) for semi-volatile analyses have average response factors greater than 0.05. Acid extractable compound 2,4-dinitrophenol showed less than required average response factor values.

Action: All sample results for 2,4-dinitrophenol should be flagged as useable.

5. Matrix Spike/Matrix Spike Duplicate Analysis

In general, no action is taken on MS/MSD data to qualify an entire case as these data alone do not give a proper indication of the precision and accuracy of the analysis of a particular sample.

The precision of an analytical method is demonstrated by the reproducibility of the data. Tables 1A and 1B are the Relative Percent Differences (RPD) between duplicate matrix spike analyses.

All MS/MSD recoveries for metals and VOAs were within the control limits established by CLP-RAS protocols as presented in Tables 1A and 1B. Five of the 11 MS/MSD recoveries for base/neutral and acid compounds were out of control. This may imply a high bias in results for those compounds out of the recovery limits, but all concentrations for the five spike compounds were reported as less than detection limit in each sample analyzed.

6. Summary

In summary, Envirotest Labs have been notified verbally and shortly, in writing, as to the deficiencies in the data presented to CRA. Until such time as we receive notification as to what they plan to do about the deficiencies, the data presented is not acceptable for use for the following reasons:

- a) No date of distillation for cyanide analysis.
- b) At least two surrogate recoveries for the acid fraction in samples 2, 3, 4 and 5 were outside control limit. Lab should re-extract and reanalyze.
- c) Spike recoveries for metals were generally out of control, a post digestion spike is required in this case but no data in the report indicates that it was performed.
- d) Due to high levels of 1-propanol in laboratory blanks, all results given for that compound should be listed as non-detected.

e) Average response factors in the continuing calibration for 2,4-dinitrophenol were not within their specified limits, making all data for that compound unusable.

References:

- 1. "Laboratory Data Validation Functional Guidelines for Evaluating Organic Analyses", Technical Directive Document, Prepared by the USEPA Data Validation Work Group, NUS Corporation Superfund Division.
- 2. "Laboratory Data Validation Functional Guidelines for Evaluating Inorganic Analyses", prepared by the USEPA Office of Emergency and Remedial Response.

TM:jd Attachments

Table 1A
Relative Percent Difference
Duplicate Matrix Spike Analysis
Inorganic Parameters

I.D. Number	Parameter	<u>RPD(1)</u>	Spike Recovery (percent)	Dup./Spike Recovery (percent)
2366-1420-001	Antimony Arsenic	35.2	100	70*
	Barium	27.1 2.4	113 87	86 85
	Beryllium	2.4	86	84
	Cadmium	4.8	86	82
	Chromium	15.8	89	76
	Cobalt	15.4	96	83
	Cyanide		105	
	Mercury	18.2	60*	50*
	Selenium	3.0	68*	66*
	Silver	2.2	90	88
	Thallium	. 0	105	105
	Vanadium	3.6	86	83

⁽¹⁾Relative percent difference between spike duplicate analysis.

* Outside of spike control limits.

Note: Control limits for RPD assumed to be $35\pm$ percent. Control limits for spike recovery are 75-125 percent.

Table 1B Relative Percent Difference Duplicate Matrix Spike Data Organic Parameters

			RPD(2)	On the	Day (Online
I.D. Number	Parameter	<u>RPD</u> (1)	Control Limits	Spike Recovery (percent)	Dup./Spike Recovery (percent)
GC/MS Volatile	e Compounds				
2366-1600-005	1,1-Dichloroethene Trichloroethene Chlorobenzene Toluene Benzene	7.6 3.8 1.0 3.2 4.9	22 24 21 21 21	96 106 103 91 100	89 102 102 94 105
BNA Compounds					
2366-1600-005	1,2,4-Trichlorobenzene Acenapthene 2,4-Dinitrotoluene Pyrene N-Nitroso Di-N-Propylamine 1,4-Dichlorobenzene Pentachlorophenol Phenol 2-Chlorophenol 4-Chloro-3 Methylphenol 4-Nitrophenol	22 21* 23 21 1.7 20 129* 56* 73* 142* 7.4	23 19 47 36 38 27 47 35 50	70 47 92* 52 58 82 9.6* 31 25	56 38 73 42 59 67 45* 55 54
Pesticides	·				
2366-1420-001	Lindane Heptachlor Aldrin Dieldrin Endrin 4,4-DDT	11 7.8 9.6 6.9 2.9	51 31 43 38 45 50	50 74 69 56 67 65	56 80 76 60 69 65

⁽¹⁾ Relative Percent Difference

⁽²⁾Control limits for RPD analysis are detailed in SW-846 (Sept. 1986)
* Values outside QC limits

MEMO



To:

Bruce Clegg

Reference No. 2366

From:

Tony Misercola

Date:

9/21/88

Re:

Envirotest Laboratories - Pyramid/Ernst Steel Site

This memo details Envirotest Laboratories response to the specific concerns on their analytical quality control outlined in my memo dated August 28, 1988.

1. Dates of Cyanide Distillation

On the initial lab report which was done by CLP-RAS protocols, Envirotest failed to report the dates of distillation for total cyanide analyses on the five soil samples submitted. Their response, in letter form, show the dates distillation occurred. All dates of distillation were within the holding time limits established in the SOW for CLP-RAS analyses. Therefore, the qualifier "J" can be removed from my original assessment of this analytical data on cyanide.

Results
1.08 U
2.04
1.14
1.09 U
1.15 U

U = The material was analyzed for, but not detected. The associated value is an estimated sample quantitation limit.

2. Surrogate Recoveries for Semi-Volatile Analyses

As noted in my memo dated 8/24/88, four of the five soil samples submitted for analyses had at least two surrogates outside control limits established in the SOW for semi-volatiles. Envirotests response was to reanalyze three of the samples for surrogate compliance. Sample No. 5 was not reanalyzed per CLP protocol set forth in the EPA Organics Statement of Work in Exhibit D, Section Iv 7.4.5. (10/86), which states that if surrogate recoveries were out of compliance in the sample and spike, reanalysis is not necessary.

All surrogates for reanalysis on semi-volatiles were within their control limits on reanalysis. Although re-extraction occurred outside prescribed holding time criteria set forth for original analyses, there are currently no holding time guidelines established for re-extraction.

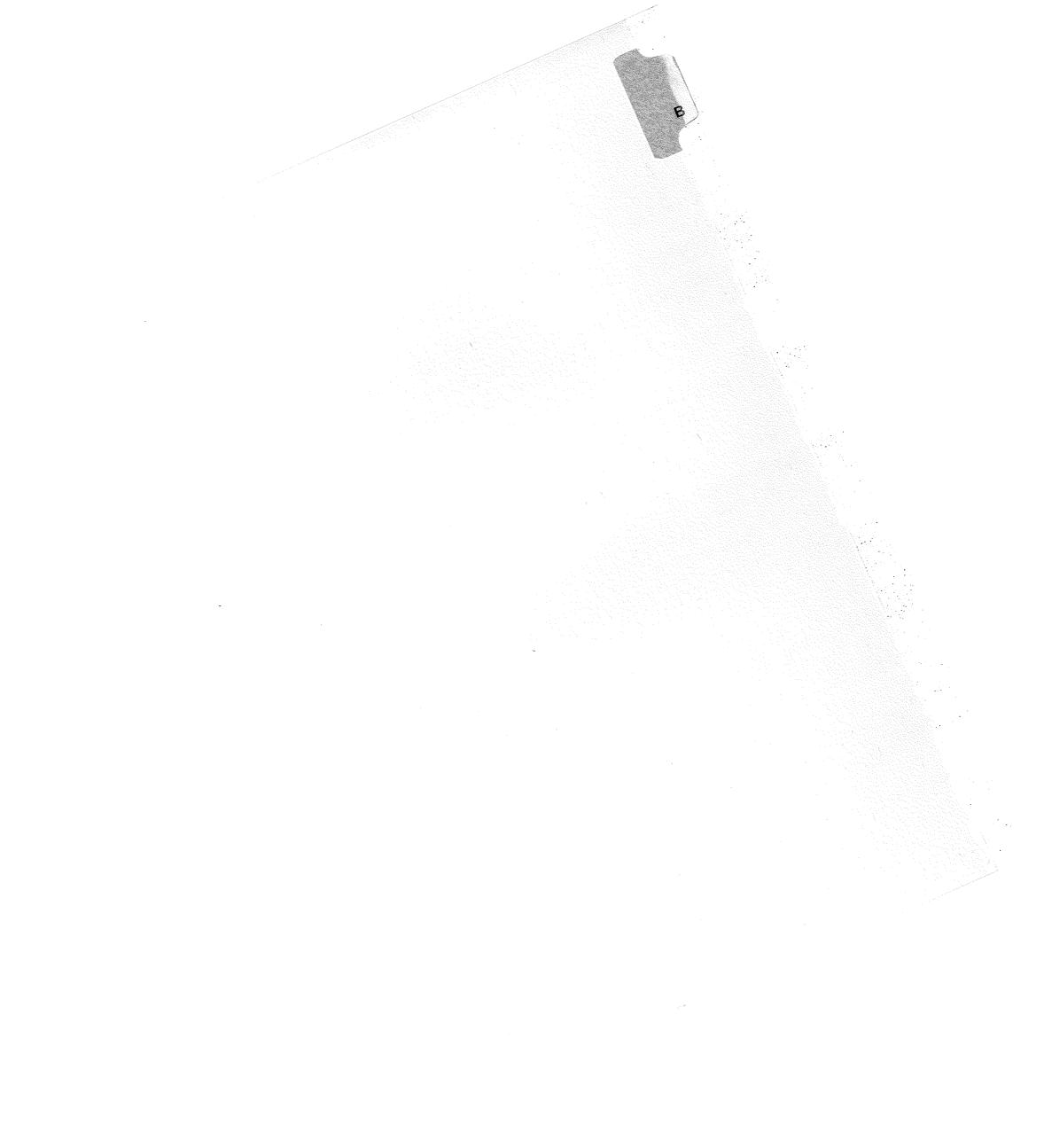
3. Spike Recovery for Metal Analyses

On the initial set of results from Envirotest, spike recovery for lead on Sample 001 was 0 percent. Their response was that they could not explain this but stated that if they could locate the sample it would be reanalyzed. As for the poor recoveries associated with mercury, nickel and selenium, their response was that by flagging their results per CLP protocols, no further action was required. Qualifiers made for those metals out of compliance in my memo of 8/24/88 should not be removed.

4. Continuing Calibration

In the original assessment of Envirotests analytical data, it was noted that system performance calibration compound 2.4-dinitrophenol had lower than required average RF value for that compound in range, but were unsuccessful. Envirotest did reanalyze the samples but were still unable to bring the average RF value for 2,4-dinitrophenol in range. All results for 2,4-dinitrophenol should still be qualified as unusable.

TM:jd



APPENDIX B

STRATIGRAPHIC AND INSTRUMENTATION LOGS

STRATIGRAPHIC AND INSTRUMENTATION LOG

(OVERBURDEN)

PROJECT NAME: PYRAMID

PROJECT NO .:

2366

HOLE DESIGNATION: OW-1 (Page 1 of 2) DATE COMPLETED: JULY 13, 1988

(L-1)

CLIENT:

PYRAMID DRILLING METHOD: HSA 7.5" O.D.

LOCATION:

NORTHWEST CORNER OF PROPERTY

CRA SUPERVISOR: D. OSCAR

i	STRATIGRAPHIC DESCRIPTION & REMARKS	ELEVATION		SAMPLE			
ft BGS		ft AMSL	INSTALLATION	N S U T M A	Ä,		
	REFERENCE ELEVATION (Top of Riser) GROUND ELEVATION	653.37 651.3		M A T E E R	A L U E		
	Fill — brown and gray silt, some fine sand, fine to medium gravel, rock fragments, trace vegetation, dry	000.0		1SS	31		
- 2.0	Fill — gray and rust slag, dry Fill — rust and brown fine to medium sand size material, some slag, trace fine gravel, dry	649.3 648.9 648.4	7.5°s BOREHOLE	255	9		
4.0	Black silt, some clay, trace fine sand, vegetation, moist, native Same, except mottled, gray Brown mottled clay, some silt, trace	647.3		388	23		
- 6.0	brown and gray brown mottled clay, some silt, moist, native Brown and gray brown mottled clay, some silt, moist, native Same, except brown			455	21		
- 8.0	Salits, skeept Breim		SEMENT/ BENTONTE GROUT	555	21		
10.0			2" BLACK STEEL PIPE	655	15		
12.0	Same, except red brown			755	8		
- 14.0				855	6		
- 16.0	some silt, trace fine sand and fine gravel, moist, native	635.3	=PENTONITE AL	955	7		
18.0	Same, except moist Gray brown clayey silt, sand, little fine to medium angular to subangular gravel, moist,	632.8		1088	11		
- 20.0	1 10 7011	631.1		1155	56		
22.0			SAND PACK	1255	118		
- 24.0	Brown fine sand, some silt, trace fine gravel, wet, native, till Brown silt, some fine sand, trace fine angular	627.4 627.3 627.0 626.3		1355	79		
- 26.0	gravel, dry to moist, native, till	625.3	BENTONITE PELLET SEAL	1455	79		
пол	TES: MEASURING POINT ELEVATIONS MAY CHANG	GE; REFER		TABLE			
	GRAIN SIZE ANALYSIS WATER FOUND V STATIC WATER LEVEL V						



PROJECT NAME: PYRAMID

HOLE DESIGNATION: OW-1 (Page 2 of 2) DATE COMPLETED: JULY 13, 1988

PROJECT NO .: 2366

(L-1)

CLIENT:

PYRAMID

DRILLING METHOD: HSA 7.5" O.D.

LOCATION:

NORTHWEST CORNER OF PROPERTY

CRA SUPERVISOR: D. OSCAR

	TOWN TOWN THE STREET SOUNCE OF THE EXTENSION OF THE EXTEN		511/1 551 E1(1)551(:			
DEPTH		ELEVATION			MPLE	
ft BGS		ft AMSL	INSTALLATION	202000	ST AT E	שכר>≺כַ
- 28.0	Brown and gray silt, some clay, trace fine sand gravel, moist, native		₩ WELL SCREEN - SAND PACK - 7.5"ø BOREHOLE	1455	X	44
20.0	Gray brown sllty clay, sand, trace fine gravel, moist, native	622.8 622.4 622.3 622.2	BENTONITE PELLET SEAL	15SS	M	70
- 30.0		622.2	SCREEN DETAILS: Screened Interval: 21.5 to 26.5 ft BGS			
- 32.0	END OF HOLE & 201 ET BCS		Length -5ft Diameter -2in Slot # 10 Material-Stainless Steel			
- 34.0						
- 36.0						
38.0			·			
- 40.0						
42.0						
44.0						
- 46.0	* , · · ·		. 7 /			
- 48.0	•					
50.0						
- 52.0			·			
NO	I TES: MEASURING POINT ELEVATIONS MAY CHAN	GE; REFER	TO CURRENT ELEVATION T	TABLE	.L	
	GRAIN SIZE ANALYSIS WATER I	FOUND 🔽	STATIC WATER LEVEL	Y		

PROJECT NAME: PYRAMID

2366

CLIENT:

PYRAMID

LOCATION:

PROJECT NO .:

NORTHEAST CORNER OF PROPERTY

HOLE DESIGNATION: OW-2 (Page 1 of 2) DATE COMPLETED: JULY 15, 1988

DATE COMPLETED:

DRILLING METHOD: HSA 7.5" O.D.

(L-2)

CRA SUPERVISOR: D. TARNOWSKI

DEPTH	STRATIGRAPHIC DESCRIPTION & REMARKS	ELEVATION		SAMPLE		
ft. BGS		ft. AMSL	INSTALLATION	N Z N		
	REFERENCE ELEVATION (Top of Riser) GROUND ELEVATION	653.72 651.4		M A A A L L L L L L L L L L L L L L L L		
	Fill — Brown silt, vegetation, gravel, dry	651.2				
- 2.0	Fill — Brown silty clay, gravel, orangish tan slag, wood fragments, dry	240.0		1SS 15		
	Red brown, orange, and gray silty mottled clay, silt lenses, dry, native	649.2	CEMENT/ BENTONITE GROUT	2SS 14		
- 4.0	Same, except dry to moist	646.2		3SS X 21		
- 6.0	Red brown silty clay, dry to moist, native	0 70.2	7.5° ø BOREHOLE			
0.0	same, except trace gray silt pockets.			455 24		
- 8.0			2° BLACK STEEL PIPE	5SS 32		
- 10.0	Same, except trace silt seams, softer, moist			6SS 16		
- 12.0	Same, except no silt seams			755 8		
- 14.0				855 5		
- 16.0			- PENTONITE AL	955 6		
- 18.0				1055 4		
- 20.0	Brown silty clay, trace fine gravel, wet, native	631.4	SAND PACK	1155 3		
- 22.0	Brown to red brown silty clay, trace fine gravel, trace sand, wet, native Red brown sandy clay, silt, fine gravel, wet,	629.4		1255 4		
- 24.0	native, Till Same, except brown	627.9				
- 26.0		625.9 625.4	WELL SCREEN	1355 3		
	wet, native, till Brown sandy clay, silt, fine gravel, wet, native, till	625.0		14SS 3		
NOTES: MEASURING POINT ELEVATIONS MAY CHANGE; REFER TO CURRENT ELEVATION TABLE						
	GRAIN SIZE ANALYSIS 💭 WATER I	FOUND 🔽	Z STATIC WATER LEVEL	Y		

PROJECT NAME: PYRAMID

HOLE DESIGNATION: OW-2 (Page 2 of 2) DATE COMPLETED: JULY 15, 1988

(L-2)

CLIENT:

2366

DRILLING METHOD: HSA 7.5" O.D.

PROJECT NO .:

PYRAMID

	LOCATI	ON: NORTHEAST CORNER OF PROPERTY		CRA SUPERVISOR: [D. TARNO	OWSK	a
	EPTH	STRATIGRAPHIC DESCRIPTION & REMARKS	ELEVATION	MONITOR		MPLE	
	. BGS		ft. AMSL	INSTALLATION	M C M	S T A T	, Ņ,
					M E R	•E	A L U E
Γ		Brown silty fine to medium sand, fine gravel, /wet, native, till	625.0	WELL SCREEN	1100	\bigvee	
		Brown sandy clay, silt, fine gravel, wet, native,		SAND PACK	14SS	Ň	3
	28.0	till		BENTONITE		\Box	
			621.6	PELLET SEAL PELLET SEAL BOREHOLE	15SS	X	4.
r	30.0	`augered to 29.9 ft. BGS, no sample / END OF HOLE ❷ 29.9 FT. BGS	621.6 621.5				
		END OF HOLE & 29.9 FT. BGS		SCREEN DETAILS: Screened Interval:			
+	32.0		·	22.0 to 27.0 ft BGS Length -5ft			
				Diameter —2in Slot # 10			
F	34.0			Material-Stainless Steel			
-	36.0						
	•	on y and		. 7			
_	38.0	•					
L	40.0						
	+0.0						
	40.0						
r	42.0						
r	44.0						
十	46.0						
F	48.0						
l		•					
-	50.0						
-	52.0	,					
_							
	тои	ES: MEASURING POINT ELEVATIONS MAY CHAN	GE; REFER	TO CURRENT ELEVATION T	ABLE		
		GRAIN SIZE ANALYSIS WATER	FOUND 🔽	STATIC WATER LEVEL	Y		

PROJECT NAME: PYRAMID

PROJECT NO .:

2366

HOLE DESIGNATION: OW-3 (Page 1 of 2) DATE COMPLETED: JULY 18, 1988

(L-3)

PYRAMID

DRILLING METHOD: HSA 7.5" O.D.

LOCATION:

CLIENT:

SOUTH OF CRANES, WEST SIDE OF PROPERTY

CRA SUPERVISOR: D. TARNOWSKI

DEPTI	- 1	STRATIGRAPHIC DESCRIPTION & REMARKS	ELEVATION		MONI			MPLE	
ft Bo	3				NSTALL	ATION .	N U	S	ά.
		REFERENCE ELEVATION (Top of Riser) GROUND ELEVATION	653.84 651.4				M B E R	A T E	A L
		Fill — brown silt, gravel, vegetation, dry	651.0					NA	
		Fill - black silt, cinders, slag, gravel, dry	650.5			-CEMENT/ BENTONITE GROUT	155	X	13
- 2.0)	Orange, brown, and gray silty Clay, dry, native	649.4			GROU I		$\langle \cdot \rangle$	
		Olive green, gray, and brown mottled silty clay, trace silt seams, dry to moist, native			-	-7.5 *s BORËHOLE	255	X	19
- 4.0)	Olive green, gray and red brown silty clay, smal fine to medium sand pocket, silt seams, dry to moist, native	647.4				388	M	14
F 6.0	,		645.4		-	BENTONITE SEAL		(
		Red brown sandy clay, silt, fine angular to subangular gravel, moist to wet, native, till	644.4		8333	SEAL	4SS	X	17
- 8.0	ו	Red brown silty fine sand, trace clay, sub- rounded to subangular gravel, wet, native, till Same, except moist	643.0				5SS		14
10.	о.	Red brown silty clay, trace sand, subrounded gravel, moist, native, till	641.9			-2"# BLACK STEEL PIPE		()	
1.0		Red brown silty fine sand, trace clay, subrounded gravel, wet, native, Till Red brown silty clay, orange, olive green, and	640.4			STEEL PIPE	6SS	X	20
12.	ں.	gray silt inclusions, subrounded gravel, dry to moist, native, till	639.0 638.8				7SS	\bigvee	33
14.	.0	Red brown silty fine sand, wet, native Red brown silty clay, some sand, subrounded to subangular Gravel, dry to moist, native, till Same, except trace subangular gravel					855		104
- 16.	.0	Red brown sandy clay, silt, trace subangular gravel, moist, native, till	635.4			-SAND PACK			
1.5		/Red brown silty fine sand, subrounded gravel, \ wet, native / Red brown silty clay, trace sand, subrounded	634.4 634.3 633.2			-3/10 (400	988	X	50
- 18.	.0	gravel, moist, native, till Brown silty fine to medium sand, fine	633.0 632.8 632.6				1055	\bigvee	56
- 20	.0	subrounded gravel, wet, native Red brown silty clay, trace sand, trace fine gravel, moist, native, till	631.4						
_ 22	n	Red brown silty fine sand, trace fine gravel,	629.9 629.4				1155	\triangle	38
		native, till	020.7				1255		54
- 24	.0	Red brown sandy clay, silt, subrounded to subangular gravel, wet, native, till Brown silty fine sand, trace fine gravel, wet,							_
- 26	. 0	Red brown silty clay, trace fine sand and					1355	\triangle	42
		gravel, wet, native, till Red brown silty fine sand, fine gravel, wet,	624.4			-WELL SCREEN	1455	X	60
		native						V \	
N	ОТ	ES: MEASURING POINT ELEVATIONS MAY CHAN		_					
	GRAIN SIZE ANALYSIS WATER FOUND STATIC WATER LEVEL								



(L-3)

PROJECT NAME: PYRAMID

HOLE DESIGNATION: OW-3A

(Page 2 of 2)

PROJECT NO .:

2366

DATE COMPLETED:

JULY 18, 1988

CLIENT:

PYRAMID

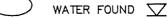
DRILLING METHOD: HSA 7.5" O.D.

LOCATION: SOUTH OF CRANES, WEST SIDE OF PROPERTY

CRA SUPERVISOR: D. TARNOWSKI

DEPTH STRATIGRAPHIC DESCRIPTION & REMARKS ELEVATION MONITOR SAMPLE INSTALLATION ft BG Y. • E Red brown silty clay, trace fine sand and gravel, wet, native, till SAND PACK 624.4 **14SS** 60 624.3 Red brown silty fine sand, fine gravel, wet, WELL SCREEN 28.0 \native Red brown silty clay, trace fine sand and gravel, wet, native, till 7.5 6 BORÉHOLE **15SS** 28 30.0 SCREEN DETAILS: Screened Interval: **16SS** 14 620.3 23.6 to 28.6 BGS Gray silty fine to medium sand, subangular to subrounded gravel, wet, native Same, with rock fragments Length →5ft 32.0 Diameter -2in **17SS** 50 618.9 Slot # 10 618.7 augered to 32.7 ft. BGS, no sample Material - Stainless Steel END OF HOLE @ 32.7 ft. BGS 34.0 NOTE: At completion the initial borehole was grouted to ground surface. In an adjacent borehole (5.0 ft. west) a 2.0" ID observation well was installed to 29.0' BGS (OW-3A). 36.0 38.0 40.0 42.0 44.0 46.0 48.0 50.0 52.0 NOTES: MEASURING POINT ELEVATIONS MAY CHANGE; REFER TO CURRENT ELEVATION TABLE

GRAIN SIZE ANALYSIS



STATIC WATER LEVEL



APPENDIX C

PHYSICAL TESTING RESULTS

9:19AM ;

GEOTECHNICAL TEST RESULTS PYRAMID HALL MEN YORK SELVING AND

REFERENCE NO.: 88-0155

PERFORMED FOR:

61ynn Geotechnical Engineering Septechnical and Civil Engineering Services 6437 Locust Street Extn. Lockport, New York 14094

PERFORMED BY:

J&L Testing Company, Inc. 938 South Central Avenue Canonsburg, PA 15317

AUGUST 10. 1988

	1
ENGINEER MA	rk 61 yan
DATE ASSISHED	7-26-88
DATE BASE	7-30-86

04		340-01	
			<u>Costocheical</u>
	Pyras	id Mal]
	Mar 1	lerk .	-

Noference No.: 88-0155

SUMMARY	OF LABOR	ATORY	TECT I	BCB1 YC
20M MAK 1	UP LABUE		8836 1	

*derne			PERM MARKER ATTERPER LIMITS UNICON-COMMESS. U 1017 SERVING SER		PRIAXIAI													
AMPLE PA	DE PYOS - Face	CLASSIFICATION	om/eac	WATER CONTENT (%)	LIMIT LIMIT	PLASTIC IBMIT	\$7#125 (tal)	STIMUM (m)	(hel) DEL MOI	08.AV(37	348	Mark.	# 140	OHER	uw.	ČĢ.	CELL PRESIMEI (pi))	aack Primbre igad
04-1 S-5	0 -10	Broom Sility Clay		21.8					106.5							·		
0V-2 S-6	10-12	Brown Silty Clay		25.5					90.6									
04-3 S-10	18-29	Clay [f]]		11.2					11/7									
04-3 S-11	29-27	Clay ISTI		12.1	<u> </u>				w									
CH-1 S-5	4-10	Jar-Free Cylindrical Soil So.	1.45x 10-8	20.5	Z 3	20			106.5		•	•						
016-2 S-6	10-12	Jar-From Cylindrical Soil St.	3.37± 16 ⁻⁸	25.7					98.6									
04-3 5-10 04-3		Combined Sample		11.8	20	15					•	•	_	_			<u> </u>	
816-3 5-11																		
		1/P-Not Possible									-							
··········		· .									_							
				<u></u>				<u> </u>										
4 500 1	fact Curves	y tes	TCOMP .	ad CHECK	E D				-	• Fi	est 1	H ##	068	!5\$				

Summary of Triaxial Permeability Test Results

Client:GLYNN GEOTECHNICAL Project Location:PYRAMID MALL

Sample Number: OW-1

Date: 08-10-1988

Job Number:88C340-01

Description: SAMPLE 5

FROM JAR

DIRECT FR. SPLIT SPOON

Cell Number: 16

Fluid: DEAIRED WATER

B-Paraseter: 1.0

Physical Property Data.....

Initial Height(in) :	2.85	Final Height(in) :	2.85
Initial Diameter(in) :	1.35	Final Diameter(in) :	1.35
Initial Wet Weight (gm):	137.40	Final wet Weight(gm):	138.20
Wet Density (pcf) :	128.32	Wet Density (pcf) :	128.94
Moisture Content % :	20.50	Moisture Content * :	21.23
Dry Density (pcf) :	106.49	Dry Density (pcf) :	106.36
Initial Saturation * :	96.88	Final Saturation % :	100.01
Initial Void Ratio :	0.5652	Final Void Ratio :	0.5670

Test Parameters......

Cell Pressur	e (psi):	55.00	0.00	9.00	0.00
Head Water	(pai):	50.00	0.00	0.00	0.00
Tail Water	(psi):	42.00	0.00	0.00	0.00

Permeability Input Data

Flow,Q(cc):	0.50	0.00	0.00	0.00
Length, L(in):	2.85	0.00	0.00	0.00
Area, A(sqin):	1.48	0.00	0.00	0.00
Head, h (psi);	8.00	0.00	0.00	0.00
Time, t(min):	800.00	0.00	0.00	0.00
Temp, T(DegC):	20.0	0.0	0.0	0.0

Computed Permeability.....(cm/sec) at 20 Degrees C

Test 1 k= 1.4515442-08

Test 2 k= 0 Test 3 k= 0

Test 4 k- 0

Summary of Triaxial Permeability Test Results

Client:GLYNN GEOTECHNICAL Project Location:PYRAMID MALL Sample Number:OW-2 Date: 08-10-1988 Job Number: 88C340-01

Description: SAMPLE S FROM JAR

DIRECT PR SPLIT SPOON

Cell Number: 17

Fluid: DEAIRED WATER

B-Parameter: 1.0

Physical Property Data.....

Initial Height(in) :	3.07	Final Height(in) :	3.05
Initial Diameter(in) :	1.44	Final Diameter(in) :	1.48
	183.40	Final Wet Weight(gm):	160.90
Wet Density (pof) :		Wet Density (pof) :	126.79
Moisture Content % :		Moisture Content % :	23.77
Dry Density (pcf) :	98.55	Dry Density (pcf) :	102.44
	98.19	Final Saturation * :	100.01
Initial Void Ratio :	0.7046	Final Void Ratio ;	0.6398

Test Parameters.......

Cell	Pressure	(pai):	55.00	0.00	0.00	0.00
Head	Water	(psi):	50.00	0.00	0.00	0.00
Tail	Water	(bai):	42.00	0.00	0.00	0.00

Permeability Input Data.....

Flow.Q(cc):	1.20	0.00	0.00	0.00
Length.L(in):	3.05	0.00	0.00	0.00
Area, A(sqin):	1.58	0,00	0.00	0.00
Head, h (psi):	8.00	0.00	0.00	0.00
Time, t(min):	800.00	0.00	0.00	0.00
Temp, T(DegC) :=	20.0	0.0	0.0	0.0

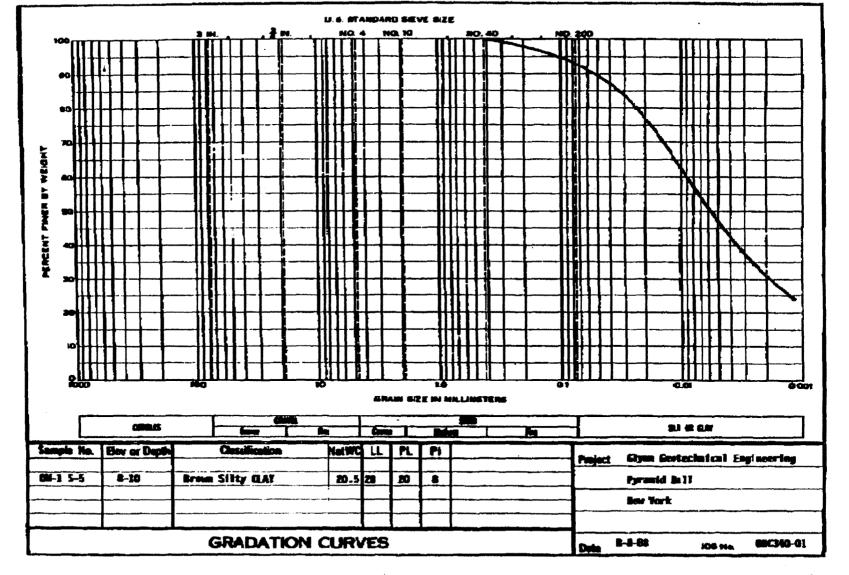
Computed Permembility.....(cm/sec) at 30 Degrees C

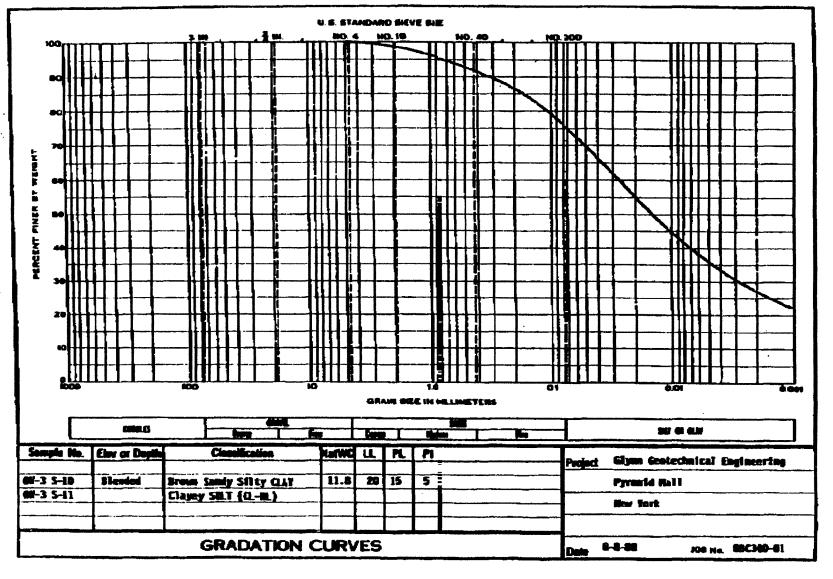
Test 1 k= 3.359672-08

Test 2 k- 0

Test 3 k- o

Test 4 k= 0







LABORATORY TEST RESULTS
CHA-PYRANID MALL
CUMPOSITE SAMPLES
CH-3 S-8
CH-7 S-7
REFERENCE NO. 88-0155

PENFORMED FOR:

Filynn Geotechnical Engineering Geotechnical and Civil Engineering Services 6437 Locust Street Extn. Lockport, New York 14084

PERFORMED BY:

JBL Testing Company, Inc. 050 Couth Control Avanua Famoushman TA 1881T

SEPTEMBER 16, 1988

momer fort Gym	PAR No. 20E340-02	BATERIC . 8-23-88
NTE ASHGHS0 8-23-00	res white Elyun Gootschafeel Engry.	9-02-80
MIE DANRESH	MI - Persold Mill	BC 87 SB
	helevenes No. 86-9155	Promise 1

ф		,	PENM	MARKE		a waas	MCON O	m470E34	WHIT	SPECIME	9	7. B	and the	 .			18 14×14	. &
ord AMRI Ab	SEPIN Face	CLAISM ECATED II	1 -	COMP-0	FRAME FEG NAME	PLANTIC BIMIN	(Pol)	STEALN (%)	etr was (gail)	OWALLA	BARRE	ŧ	971 R	993	44	Ç.	CELL Pars punc (p. 4	BACE Periones (gui)
08-3 08-7	S-8 S-7	Composite Sample of Car	1.03× 10-3	כמ	28	14			127.6		•	٠						
															_	Ļ		
								:			_							
										.								
											L			·				
																		<u> </u>
														<u> </u>				
														П				
		-										П						<u> </u>
																-		
				1					 	1	-					1		

5 10 (6-88)

Test Results

Client: GLYNN GEOTECHNICAL Project Gnostion: PYRAMID MALL Sample Number: COMPOSITE SAMPLE

Date: 09-18-1988 Job Wumber: 886340-01

Description: ON-3 S-8
OW-7 S-7
12-16 FEET

Cell Number: 25

Fluid:DRAIRED WATER

B Parameter:1.0

Physical Property Data.......

Initial Height(in)	ŧ	1.85	Pinal Height(in) :	1.83
Initial Diameter(in)			Final Dismater (in) .	4.40
Initial Wet Weight (gm)	;	420.20	Final Wet Weight (gm);	423.20
Wet Denzity (pof)	ı	140.40	Wat Banaity (puf) ;	144.73
Moisture Content *	:	10.00	Moisture Content % :	10.80
	•	127.64	Dry Density (pof) :	129.72
Initial Saturation %	:	88.39	Final Saturation % :	100.10
Initial Void Ratio	:	0.3102	Final Void Ratio :	0.2891

Test Parameters........

Cell P	ressure	(pai):	85.00	0.00	0.00	0.00
Head W	ater	(pmi):	50.00	0.00	0.00	0.00
Tail W	ater	(bail:	42.00	0.00	0.00	0.00

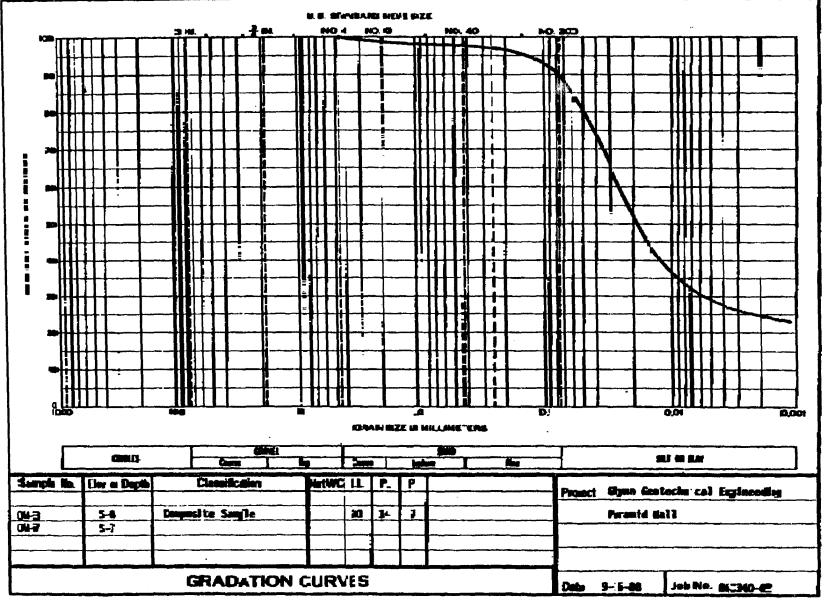
Permeability Input Data......

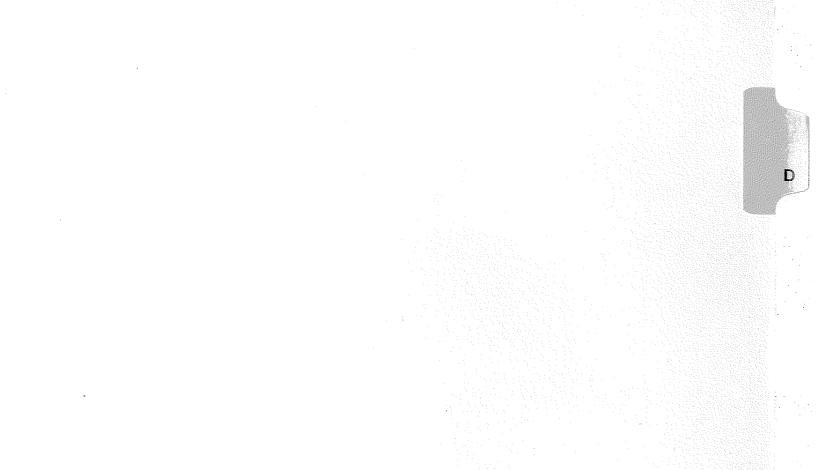
Flow.Q(cc):	17.60	0.00	0.00	0.00
Length, L(in):	1.82	0.00	0.00	0.00
Area, A(sqin):	6.16	0.00	0.00	0.00
Head, h(psi):	8.00	0.00	0.00	0.00
Time.t(min):	500.00	0.00	0.00	0.00
Tamp, T(DagC):	24.6	0.0	0.0	0.0

Computed Permeability.....(cm/sec) at 26 Degrees C

Test 1 k= 1.089194E-07

Test 2 k= 0 Test 3 k= 0 Test 4 k= 0





APPENDIX D

100-FOOT GRID SAMPLING RESULTS
SAMPLES ANALYZED FOR TOTAL AND EP TOX LEAD



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ANALYTICAL RESULTS FOR

CONESTOGA ROVERS & ASSOCIATES

ENVIRONMENTAL LABORATORY ACCREDITATION PROGRAM (ELAP) CERTIFICATION #10797

FIELD INFORMATION

PROJECT NO. 2366

Name of Collector: Steve Supernaut

Site of Collection: Pyramid-Ernst Steel Site Date of Collection: August 17, 1988 (3 - 29)
August 18, 1988 (30 - 82)

ASSIGNED BLT# I.D.	SAMPLE I.D.#	SAMPLE TYPE	Time of Collection
2326-Ø1	3	Soil	113Ø EST
2326-02	8	n	1515 EST
2326-Ø3	9		153Ø EST
2326-04	10	n	1540 EST
2326-Ø5	11	#	1550 EST
2326-Ø6	12	Ħ	1600 EST
2326 – Ø7	13	11	1620 EST
2326 – Ø8	14	Ħ	163Ø EST
2326 – Ø9	14A	n	1640 EST
2326-10	15	Ħ	1655 EST
2326-11	16	11	17Ø5 EST
2326-12	17	Ħ	1715 EST
2326-13	18	Ħ	1730 EST
2326-14	19	n	1740 EST
2326-15	2Ø	n	1750 EST
2326-16	21	Ħ	18Ø5 EST
2326-17	21A	n	1815 EST
2326-18	22	n	1820 EST
2326-19	23	W	1835 EST
2326-20	24	Ħ	1845 EST
2326-21	25	**	1850 EST
2326-22	26	п	1900 EST
2326-23	27	Ħ	1910 EST
2326-24	28	m	193Ø EST
2326-25	29	Ħ	1935 EST
2326-26	3Ø	m	Ø745 EST
2326-27	31	Ħ	Ø8ØØ EST
2326-28	32	n	Ø815 EST



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ANALYTICAL RESULTS FOR

CONESTOGA ROVERS ASSOCIATES

ENVIRONMENTAL LABORATORY ACCREDITATION PROGRAM (ELAP) CERTIFICATION #10797

ASSIGNED			Site, Time and Date
BLT# I.D.	SAMPLE I.D.#	SAMPLE TYPE	of Collection
2326–29	33	n	70.3E 12.00
2326 – 29 2326 – 3Ø	33 34	n	Ø835 EST Ø84Ø EST
2326-31	3 4 35	n	Ø855 EST
2326-32	35A	n	0900 EST
2326-32		H .	
2326 – 33	36 37	**	Ø91Ø EST Ø91Ø EST
2326 – 35	37 38		
2326 – 35		n	Ø915 EST
	39	77	Ø925 EST
2326-37	40		Ø935 EST
2326–38	41	 11	Ø95Ø EST
2326–39	42	 N	Ø955 EST
2326-40	43	 N	1000 EST
2326-41	44	n	1010 EST
2326-42	45		1020 EST
2326-43	46	n	1050 EST
2326-44	47	 n	1100 EST
2326-45	46A		1055 EST
2326-46	48	. " N	1110 EST
2326-47	49	" "	1125 EST
2326-48	49A	7	1136 EST
2326–49	50	# ## ## ## ## ## ## ## ## ## ## ## ## #	1130 EST
2326-50	51	n	1140 EST
2326-51	52		115Ø EST
2326-52	53	n 	1155 EST
2326-53	54	11	1200 EST
2326-54	55	11	1210 EST
2326–55	56	Ħ	1215 EST
2326–56	57	Ħ	1220 EST
2326–57	58	#	1225 EST
2326-58	59	Ħ	1430 EST
2326-59	6Ø	Ħ	1435 EST
2326 – 6Ø	61	Ħ	1450 EST
2326-61	62	n	1510 EST
2326-62	63	Ħ	1521 EST
2326-63	63A	n	1525 EST
2326-64	64	π	1535 EST
2326–65	65	Ħ	1640 EST



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ANALYTICAL RESULTS FOR

CONESTOGA ROVERS ASSOCIATES

ENVIRONMENTAL LABORATORY ACCREDITATION PROGRAM (ELAP) CERTIFICATION #10797

ASSIGNED BLT# I.D.	SAMPLE	I.D.#	SAMPLE TYPE	Site,Time and Date of Collection
2326-66		66	n	1645 EST
2326-67		67	n	1648 EST
2326-68		68	n	1700 EST
2326-69		69	H	17Ø5 EST
2326 – 7Ø		7Ø	Ħ	1710 EST
2326-71		71	n	173Ø EST
2326-72		72	n	1740 EST
2326 - 73		73	n	1750 EST
2326-74		74	n	1755 EST
2326-75		7 5	Ħ	181Ø EST
2326-76		76	Ħ	1820 EST
2326-77		77	, 11	1840 EST
2326-78		7ØA	n	1825 EST
2326 -7 9		78	tī	1855 EST
2326 - 8Ø		8Ø	Ħ	1930 EST
2326-81		81	n	1935 EST
2326-82		82	Ħ	1955 EST
2326-83		79	π	19Ø5 EST

Laboratory Information

Sample ID	Preservation Status Upon Acceptance	Date/1	ime Received
2326-(Ø1 -83)	Properly preserved and collected		August 18, 1988 Not Available

RELEASED BY: Husein Sitabkhan

4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ELAP #10797

ANALYSIS FOR: Conestoga Rovers Associates

Project #2366 Project: Pyramid-Ernst Steel Site

BLT # 2326

SAMPLE ID	Pb, mg/Kg	DUPLICATE
. Method Blank	<10	
EPA 1085 (1)	5.33	
3	42	
8	250	
9	65	
1Ø	45	
11	132	
12	25Ø	
13	210	191
13 Spk (2)	336	346
Method Blank	<10	
EPA 1085	5.27	
14	625	
14A	15	
15	13	
16	4	
17	32	
18	203	
19	112	
2Ø	16Ø	
21	166	
21A	54	
22	17Ø	174
22 SPK (2)	3Ø2	314

- * Dry Weight Basis
- (1) Results in mg/L: TV = 5.00 mg/L
- (2) Spiked with 100 ppm(3) Spiked with 50 ppm
- (4) Results in mg/L : EP Tox

Spikes results are "approximations" due to differences in NOTE weights between original and spiked samples. See attached explanations



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ELAP #10797

ANALYSIS FOR: Conestoga Rovers Associates

Project: Pyramid-Ernst Steel Site

Project #2366

SAMPLE ID	Pb, mg/Kg	DUPLICATE
Method Blank	<10	
EPA 1085	5.27	
23	247	
24	167	
25	176	
26	152	
27	2Ø1	
28	61	
29	54	59
29 SPK (3)	111	89
Method Blank	<1Ø	
EPA 1085	5 . 49	
3Ø	115	
31	219	
32	222	
33	62	
34	75	
35	18,340	
35A	11Ø	
36	85	
37	22Ø	
38	215	2Ø1
38 SPK (3)	266	249
Method Blank	<1Ø	
EPA 1Ø85	5 .3 8	
39	235	
40	189	
41	535	

- * Dry Weight Basis
- (1) Results in mg/L: TV = 5.00 mg/L
 (2) Spiked with 100 ppm
 (3) Spiked with 50 ppm
 (4) Results in mg/L: EP Tox



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ELAP #10797

ANALYSIS FOR: Conestoga Rovers Associates

Project: Pyramid-Ernst Steel Site

Project #2366

SAMPLE ID	Pb, mg/Kg	DUPLICATE
42	164	
43	187	
44	10,650	
45	73	
46	364	
46A	54	
47	1Ø7	120
47 SPK (3)	155	16Ø
Method Blank	<10	
EPA 1085	5.21	
48	412	
49	263	
49A	34	
5Ø	129	
51	226	
52	434	
53	529	
54	85	
55	95	
56	92	92
56 SPK (3)	154	161
Method Blank	<10	
EPA 1085	5.38	
57	88	
58	132	
59 →	110	
60	3 67	
61	2540	

- * Dry Weight Basis
- Results in mg/L: TV = 5.00 mg/L
 Spiked with 100 ppm
 Spiked with 50 ppm
 Results in mg/L: EP Tox

4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ELAP #10797

ANALYSIS FOR: Conestoga Rovers Associates

Project: Pyramid-Ernst Steel Site Project #2366

SAMPLE ID	Pb, mg/Kg	DUPLICATE
62	40.0	
63	117	
63A	9	
64	122	112
64 SPK (3)	19Ø	185
65	92	
Method Blank	<1Ø	
EPA 1085	5 . 44	
66	97	
67	118	
68	52	
69	63	
7Ø	119	
7ØA	10	
71	77	86
71 SPK (3)	134	134
72	317	
73	2Ø7	
74	76	
75	57	
76	58	
77	31	
78	16Ø	
79	239	
8Ø	86	
81	71	72
81 SPK (3)	128	126
Repeat 29	56	5Ø
Repeat 29 SPK	105	94

- * Dry Weight Basis
- (1) Results in mg/L: TV = 5.00 mg/L

- (2) Spiked with 100 ppm(3) Spiked with 50 ppm(4) Results in mg/L : EP Tox

4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ELAP #10797

ANALYSIS FOR: Conestoga Rovers Associates

Project #2366 Project: Pyramid-Ernst Steel Site

BLT # 2326

SAMPLE ID	Pb, mg/Kg	% RPD	% Rec
Method Blank	<10		
EPA 1085	5.27		1Ø5%
82	512		
9 EP Tox (4)	<0.1		
15 EP Tox (4)	<0.1		
20 EP Tox (4)	<0.1		
39 EP Tox (4)	<0.1		
44 EP Tox (4)	<0.1		•
63 EP Tox (4)	<0.1		
70 EP Tox (4)	<0.1		
80 EP Tox (4)	<0.1		
Method Blank	<1Ø		
EPA 1Ø85	5.33		1Ø7%

Sample Preparation Method: EPA SW-846 (3050) Analysis Method: EPA SW-846 (7420)

Samples 3 through 29 Digested 8/18/88 Samples 30 through 82 Digested 8/19/88

All samples analyzed 8/20/88

- * Dry Weight Basis
- (1) Results in mg/L : TV = 5.00 mg/L
- (2) Spiked with 100 ppm(3) Spiked with 50 ppm
- (4) Results in mg/L : EP Tox

4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Date: August 22, 1988

ELAP #10797

ANALYSIS FOR: Conestoga Rovers Associates

Project: Pyramid-Ernst Steel Site

Project #2366

SAMPLE ID	Pb, mg/Kg	DUPLICATE	% RPD
13	210	191	9.48
22	17Ø	174	2.32
29	54	59	. 8.85
38	215	201	6.73
47	107	12Ø	11.5
56	92	92	Ø
64	122	112	8.55
71	7 7	86	11.0
81	71	72	1.40
29 Repeat	56	5Ø	11.3

4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Project #2366

WEIGHT CORRECTED % RECOVERIES

Because of the variation in sample sizes, a more sophisticated methemetical approach to % recovery calculations must be used.

Calculate Total mg Pb in spiked sample:

$$mg(\tau)_{SpX} = Onc_{\bullet}(\frac{mg}{10000c}) \times Sample vol^{\bullet}m(1001cc)$$

$$= \left(\frac{conc}{10}\right)mg = \frac{ppm}{10}mg$$

Calculate sample contribution to spike:

Calculate sample contribution to spike:

$$m_{q}(T)_{smpl} = \frac{m_{q}}{1000.9} \times wt(9)_{wet} \times \frac{9/0.50 \text{ lids}}{100}$$

cont.

3) Recovery =
$$mq(T)_{spk} - mg(T)_{smpl}$$

 $cont \times 100$
Amt. Spike-mg

Amount Spike (mg) =
$$\frac{CONC(PP^m)}{1000.cc} \times Vol.spike(cc)$$



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Project #2366

QA/QC

Spike Results

Sample ID	Pb, mg/Kg	Duplicate	% RPD
13 13 SPK (1) Wt. Sample (2)	210 335 (3.16) 1.1266	191 346 (3.99) 1.0286	9.48
% Recovery	117	134	13.5
22 22 SPK (1) Wt. Sample (2)	170 302 (3.18) 1.2859	174 314 (3.30) 1.3103	2.32
% Recovery	1ø6	68	43.7
29 29 SPK (1) Wt. Sample (2)	54 111 (1.04) 1.0599	59 89 (0.99) 1.2495	8.85
% Recovery	1Ø6	68	43.7
38 38 SPK (3) Wt. Sample (2)	215 266 (2.82) 1.2288	201 249 (2.71) 1.2654	6.73
% Recovery	108	102	5.71
47 47 SPK (3) Wt. Sample (2)	107 155 (1.54) 1.0419	12Ø 16Ø (1.77) 1.16Ø8	11.5
% Recovery	96	88	8 . 7Ø

- * Values in parentheses are conc (mg/L) of spiked solution; Sample volume = 100.0 ml
- (1) Spiked with 10.0 ml of 10 ppm Pb to 100 ml final volume (0.100 mg)
- (2) Weight sample used for spike
- (3) Spiked with 5.0 ml of 5.0 ppm Pb to 100 ml final volume (0.05 mg)



4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

QA/QC

Spike Results

Sample ID	Pb, mg/Kg	Duplicate	% RPD
56 56 SPK (3) Wt. Sample (2)	92 154 (1.32) 1.1660	92 161 (1.21) 1.0190	Ø
% Recovery	1Ø6	104	1.91
64 64 SPK (3) Wt. Sample (2)	122 190 (1.65) 1.2361	112 185 (1.65) 1.2734	8.55
% Recovery	118	13Ø	9.68
71 71 SPK (3) Wt. Sample (2)	77 134 (1.27) 1.1370	86 134 (1.38 1.2221	11.0
% Recovery	108	102	5.71
81 81 SPK (3) Wt. Sample (2)	71 128 (1.15) 1.1233	72 126 (1.21) 1.1908	1.40
% Recovery	102	104	1.94
Repeat 29 29 SPK (3)	56 105 (1.00)	50 94 (1.14)	11.3
Wt. Sample (2)	1.0804	1.3656	16 0
% Recovery	92	1Ø8	16.Ø

^{*} Values in parentheses are conc (mg/L) of spiked solution; Sample volume = 100.0 ml

- (1) Spiked with 10.0 ml of 10 ppm Pb to 100 ml final volume (0.100 mg)
- (2) Weight sample used for spike
- (3) Spiked_with 5.0 ml of 5.0 ppm Pb to 100 ml final volume (0.05 mg)

4626 Royal Avenue, Niagara Falls, New York 14303 • Phone (716) 285-2587

Project #2366

SOLIDS DATA FOR QA/QC

SAMPLE ID	% SOLIDS
13	83.97
22	80.02
29	88.55
38	86.20
47	95.25
56	73.79
64	70.41
71	83.04
81	80.43

WAT. FILE COPY

MEMO

To:

Bruce Clegg

Reference No. 2366

From:

Tony Misercola

Date:

9/14/88

Re:

Analytical Data Validation - Ernst Steel/Pyramid Co.

The following details as assessment and validation of analytical results reported by BLT Technical Services, Inc. for soil/fill samples collected at the Ernst Steel/Pyramid Site on August 17-18, 1988. The samples submitted for analysis consisted of the following:

	Investigative	Field
Matrix	Samples	Duplicates
	-	
Soil/Fill	77	6

All samples collected were submitted for total lead by Method 7420 (Test Method for Evaluating Solid Waste, USEPA SW-846, 3rd Edition, September 1986) and 10 percent of those samples were submitted for EP TOX lead by Method 1310/7420 (USEPA SW-846, 3rd Edition).

The QA/QC criteria by which these data have been evaluated are outlined in the aforementioned methods an the documented entitled "Laboratory Data Validation Functional Guidelines for Evaluating Inorganics Analyses". Based on this data and other related quality control data, the following are noted:

1. Sample Holding Time

Based on the criteria outlined in USEPA SW-846, the holding time for lead analysis on solids has been established as:

<u>Metals</u>

6 months prior to analysis

(Solids)

No preservative required other than storing at 4°C until time of analysis.

It was noted that all soil samples were properly collected and stored in coolers until time of analysis. Actual holding times are established by comparison of the sampling dates specified on the Chain of Custody to the reported dates of analysis. All samples were submitted on a rush basis and analyzed prior to the expiration of their prescribed holding time.

2. Blank Analysis

The assessment of blank analyses results is to determine the existence and magnitude of contamination problems. All 10 method blanks analyzed with each set of samples yielded nondetectable concentrations of lead.

3. <u>Laboratory Control Samples</u>

To establish the ability to generate acceptable laboratory accuracy and precision, the laboratory must perform a quality control check sample. In all, 10 EPA check samples were analyzed and demonstrated relatively high accuracy from adequate results of their percent recoveries. Precision was achieved through the low standard deviation between percent recoveries. See Table 1

4. Duplicate Sample Analyses

Duplicate analyses of individual samples are indicators of laboratory precision based on each sample matrix. As outlined in "Functional Guidelines for Evaluating Inorganics Analyses", the RPD Control (relative percent difference) limits is ± 35 percent. Table 2 lists the results of the laboratory selected duplicates and their corresponding RPD values. It is noted that the analyses of these duplicates indicated that acceptable precision was attained for lead.

5. Field OA/OC

Field duplicate samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision, therefore the results may have more variability than laboratory selected duplicates which measure

only lab performance. It is also expected that soil duplicate results will have a greater variance than aqueous matrices due to difficulties associated with collected identical field samples. Small changes in the matrix may have a substantial effect on the reproducibility of the analytical data.

Table 3 lists the results of field duplicate analyses for lead.

The results of field duplicates yielded satisfactory reproducibility with the exception of duplicates collected at Location 49 and 73. The sample collected at Location 49 yielded twice as much lead as its duplicate, while the sample at 73 yielded almost three times as much lead as its duplicate. The discrepancies in these data may be attributed to the inhomogeneity of the sample matrix.

6. Matrix Spike/Matrix Spike Duplicate (MS/MSD) Analyses

In general, no action is taken on MS/MSD data to qualify an entire case as these data alone do not give a proper indication of the precision and accuracy of the analyses of a particular sample.

RPD (relative percent difference) control limits of ± 20 percent (outlined in USEPA SW-86) for inorganic parameters was determined from waste water analyses. RPD values for eight out of 10 MS/MSD analyses on the soil/fill samples attained acceptable precision, while two RPD values were relatively high (43.7 percent). Again, the discrepancies in the data may be attributed to inhomogeneity in the sample matrix.

The control limits established for spike percent recovery as outlined in USEPA SW-846, have generally been developed for wastewater analyses. The samples being analyzed were a rough mixture of soil and fill and considered a very difficult matrix to deal with. Of the 20 spike analyses, three were outside the control limits of 75-125 percent. Considering the nature of the matrix, the percent recoveries indicate satisfactory accuracy among the analyses. Table 4 presents the precision and accuracy data obtained from the MS/MSD data.

7. Summary

In summary, standard laboratory and field QA/QC were adhered to, making the data acceptable for use with the following qualifications:

- a. There were no holding time violations.
- b. USEPA SW-846 does not give proper control limits for spike recoveries for solids, nor proper control limits for RPD values for solid MS/MSD samples in conjunction with metals analyses. Even so, most of the quality control done by the Lab was acceptable by wastewater standards established in USEPA SW-846.

References

"Laboratory Data Validation Functional Guidelines for Evaluating Inorganics Analyses", prepared by USEPA Office of Emergency and Remedial Response.

TM:jd Attachment

TABLE 1

Summary of QC Check Sample Data

Identification No.

EPA 1085

Parameter

Lead

Matrix

Water

Standard Deviation

1.7 Percent

Percent Recovery

106.6

105.4

105.4

109.8

107.6

104.2

107.6

108.8

105.4

106.6

TABLE 2

Comparison of Laboratory Duplicates for Lead Analysis

	Original		RPD Control		
I.D. No.	Value	Duplicate	Limit	RPD	<u> Matrix</u>
,	(mg/kg)	(mg/kg)	(%)	(%)	
2366-13	210	191	3 5	9.5	soil/fill
2366-22	170	174	3 5	2.3	
2366-29	54	59	3 5	8.8	
2366-38	215	201	3 5	6.7	
2366-47	107	120	3 5	11.5	
2366-56	92	92	3 5	0	
2366-64	122	112	3 5	8.5	
2366-71	77	86	3 5	11.0	
2366-81	71	72	35	1.4	

TABLE 3

Lead Results for Field Duplicates

	Original		Duplicate	
I.D. No.	Concentration	I.D. No.	Concentration	<u>RPD</u>
	(mg/Kg)		(mg/kg)	(%)
2366-28	61	2366-29	57	6.8
2366-37	220	2366-38	208	5.6
2366-49	263	2366-50	129	68.4
2366-52	434	2366-53	529	19.7
2366-68	52	2366-69	63	15.9
2366-73	207	2366-74	76	92.6

TABLE 4

Relative Percent Difference Between
Duplicate Matrix Spike Analysis

I.D. No.	Parameter	RPD	Spike ¹ Recovery	Spike/Dup. ¹ Recovery
			(Percent)	(Percent)
2366-13	Lead	13.5	117	134
2366-22	Lead	43.7*	106	68
2366-29	Lead	43.7*	106	68*
2366-38	Lead	5.7	108	102
2366-47	Lead	8.7	96	88
2366-56	Lead	1.9	106	104
2366-64	Lead .	9.7	118	130*
2366-71	Lead	5.7	108	102
2366-81	Lead	1.9	102	104
2366-29	Lead	16	92	108
Repeat				

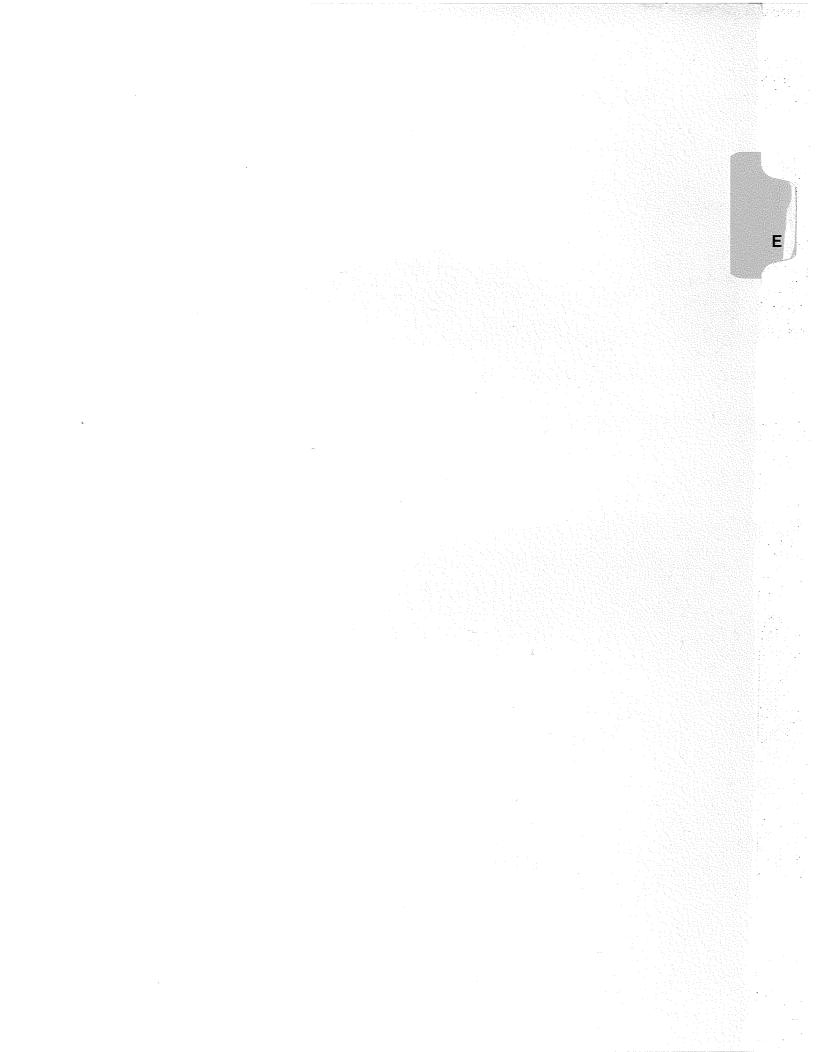
¹Control limits for percent recovery for lead (75-125 percent) USEPA SW-846, 3rd Edition, September 1986.

RPD = Relative percent difference between spike and duplicate analyses.

* Outside control limits

Note:

Control limits for RPD assumed to be ± 20 percent. Matrix for all samples - soil/fill



APPENDIX E

100-FOOT GRID SAMPLING RESULTS
SAMPLES ANALYZED FOR TARGET COMPOUND LIST

CLIENT:CRASAMPLE#1SAMPLE ID:SAMPLE#1		E:5GM G/KG):_UG/KG_
VOLATILE COMPOUNDS	I CONCENTRATION	I DETECTION LIMIT
Chloromethane	L AUD	- _ 10
Bromomethane	1 ND	
Vinyl Chloride	1 ND	1 10
Chloroethane	I ND	
Methylene Chloride	i	
Trichlorofluoromethane	15	
1,1-Dichloroethene	I ND	i
1,1-Dichloroethane	i ND	
Trans-1,2-Dichloroethene	i ND	_ i
Chloroform	16	- '
Chloroform	i ND	5
1,1,1-Trichloroethane	i ND	5
Carbon Tetrachloride	I ND	- '
Bromodichloromethane	I ND	_ '
1,2-Dichloropropane	I ND	- 1 5
Trans-1,3-Dichloropropene	I ND	_'
Trichloroethene	I ND	
Trichloroethene	I ND	- <u>'</u>
1,1,2-Trichloroethane	l ND ·	_
Benzene	I ND	_ i5
cis-1,3-Dichloropropene	l ND	
2-Chloroethylvinylether	I ND	
Bromoform	I ND	
Tetrachloroethene	i ND	
1,1,2,2-Tetrachloroethane	I ND	- '
Toluene		5
Chlorobenzene	I ND	5
		- i
Ethylbenzene		_ i
1,2-Dichlorobenzene		i
1,4-Dichlorobenzene		
	1	
SURROGATE	RECOVERY DATA	
SURROGATE COMPOUND QC	LIMITS	PERCENT RECOVERY
WATER	SOIL	
Toluene-d8	574-121	84

ND: NOT DETECTED

LABORATORY NUMBER:8801272 S DATE COLLECTED/RECEIVED:8/18/88 L DATE OF ANALYSIS:8/18/88 C	MATRIX: (SOIL/WAT SAMPLE WT/VOLUME: UNIT (UG/L OR UG/)ILUTION FACTOR:_ CONCENTRATION I	5GH /KG):_UG/KG_ 1
	CUNCENTRATION	DETECTION CITIES
Chloromethane	ND I	10
Bromomethane 1	ND I	10
Vinyl Chloride	ND I	10
Chloroethane		10
Methylene ChlorideI	7	5
Trichlorofluoromethane	11I	5 / 1
1,1-Dichloroethene	NDI	5I
1,1-Dichloroethane	ND1	5 <u>/</u> I
Trans-1,2-Dichloroethene	NDI	<u> </u>
Chloroform	71	[5I
1,2-Dichloroethane	NDI	5l
1,1,1-Trichloroethanel	ND1	اا
Carbon Tetrachloride		51
Bromodichloromethane	ND1	5I
1,2-Dichloropropane	ND1	5I
Trans-1,3-Dichloropropene	NDI	5I
Trichloroethene	NDI	5I
Dibromochloromethane	ND1	5l
1,1,2-Trichloroethane	ND I	5I
Benzene	ND	l5l
cis-1,3-Dichloropropene	ND1	5I
2-Chloroethylvinylether	ND	101
Bromoform	NDI	5I
Tetrachloroethene	ND!	اا
l 1,1,2,2-Tetrachloroethanel_	NDI	5I
Toluene	ND	ا <u> </u>
Chlorobenzene	ו טאו	5l
EthylbenzeneI		١١
I 1,3-DichlorobenzeneI	NDI	5I
1,2-Dichlorobenzene	ND1	51
1,4-Dichlorobenzene	NDI	51

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIMITS		PERCENT RECOVERY	
	WATER	SOIL		
Toluene-d8	88-110	81-117		
Bromofluorobenzene				

ND: NOT DETECTED

CLIENT:CRASAMPLE#3	MATRIX: (SOIL/WATER)_SUIL SAMPLE WT/VOLUME:5GM UNIT (UG/L OR UG/KG):_UG/KG_ DILUTION FACTOR:1		
VOLATILE COMPOUNDS	CONCENTRATION	I DETECTION LIMIT!	
Chloromethane	I ND	<u> </u>	
Bromomethane			
Vinyl Chloride	I ND	1101	
Chloroethane	IND		
Methylene Chloride	!9	_11	
Trichlorofluoromethane	IND	151	
1,1-Dichloroethene	IND		
1,1-Dichloroethane	1ND	- ' '	
Trans-1,2-Dichloroethene	IND	151	
Chloroform	_1		
1,2-Dichloroethane	iND		
1,1,1-Trichloroethane	_!ND		
Carbon Tetrachloride	_!ND	5	
Bromodichloromethane		_!	
1,2-Dichloropropane			
Trans-1,3-Dichloropropene	!ND		
Trichloroethene	IND		
Dibromochloromethane	IND I ND	5	
1,1,2-Trichloroethane	1 ND	15	
Benzene cis-1,3-Dichloropropene		5	
1 2-Chloroethylvinylether			
Promoform	I NU	- i 5 I	
Bromoform_ Tetrachloroethene	I ND	5	
1 1,1,2,2-Tetrachloroethane	I ND	1 5	
Toluene	I ND	1 5 1	
Chlorobenzene		5 1	
I Ethylbenzene	I ND	· · · · · · · · · · · · · · · · · · ·	
l 1,3-Dichlorobenzene	IND	_I	
l 1,2-Dichlorobenzene	IND	_151	
1,4-Dichlorobenzene		5	
SURROGATE RECOVERSURROGATE COMPOUND QC LIMIT	======	PERCENT RECOVERY	
WATER	 SOIL		
		•	
Toluene-d8	74-121		
ND: NOT DETECTED			

CLIENT:CRASAMPLE#4SAMPLE ID:SAMPLE#4	SAMPLE WT/VOLUME UNIT (UG/L OR UG	MATRIX: (SUIL/WATER)_SUIL SAMPLE WT/VOLUME:5GM UNIT (UG/L OR UG/KG):_UG/KG_ DILUTION FACTOR:1		
VOLATILE COMPOUNDS	CONCENTRATION	I DETECTION LIMIT!		
Chloromethane	IND	10		
Bromomethane	I ND	10 1		
Vinyl Chloride	I ND	10 1		
Vinyl Chloride	I ND	10		
i Methylene Chioride	18	5		
Trichlorofluoromethane	I10	i5i		
1,1-Dichloroethene	IND			
1 1,1-Dichloroethane	IND	l <u> </u>		
Trans-1,2-Dichloroethene	1ND	.		
Chloroform	16	51		
l 1,2-Dichloroethane	IND	51		
1 1,1,1-Trichloroethane	I ND	l5l		
Carbon Tetrachloride	IND	l1		
Bromodichloromethane	1ND	l51		
1,2-Dichloropropane	IND	l1		
Trans-1,3-Dichloropropene	I NU	l5l		
Trichloroethene	I ND	J 5		
Dibromochloromethane	I ND_	l 5l		
1,1,2-Trichloroethane		5 1		
Benzene		5		
l cis-1,3-Dichloropropene	I ND	1 5 1		
1 2-Chloroethylvinylether	I ND	10		
l Bromoform	IND	5I		
Tetrachloroethene	I ND	151		
1 1,1,2,2-Tetrachloroethane		I5I		
I Toluene	1ND	l		
Chlorobenzene	IND	l1		
I Ethylbenzene		l		
I 1.3-Dichlorobenzene	I ND	1 5 1		

SURROGATE RECOVERY DATA

SURROGATE COMPUUND	QC LIMIT	S	PERCENT RECOVERY
	WATER	SOIL	
Toluene-d8	86-115	74-121	

ND.

ND: NOT DETECTED

1,2-Dichlorobenzene_

1,4-Dichlorobenzene_

CLIENT:CRA	
SAMPLE ID:SAMPLE#5	MATRIX: (SOIL/WATER)_SUIL
LABORATORY NUMBER:8801275	SAMPLE WT/VOLUME:5GM
DATE COLLECTED/RECEIVED:8/18/88	UNIT (UG/L OR UG/KG):_UG/KG_
DATE OF ANALYSIS:8/18/88	DILUTION FACTOR:1
	•
	:

VOLATILE COMPOUNDS	CONCENTRATION	I DETECTION LIMIT
Chloromethane	IND	1101
Bromomethane	ו אט_	101
Vinyl Chloride		1101
Chloroethane	IND	101
Methylene Chloride	122	l5l
Trichlorofluoromethane		_l5
l 1,1-Dichloroethene	1NU	_l
l 1,1-Dichloroethane	INÜ	_I5/
Trans-1,2-Dichloroethene	1NÜ	.l5!
Chloroform		15
I 1,2-Dichloroethane		l5!
l 1,1,1-Trichloroethane		15
l Cárbon Tetrachloride	I ND	15
Bromodichloromethane	I ND	15
1,2-Dichloropropane		5
Trans-1,3-Dichloropropene	I ND	15
Trichlorosthene	I ND	15
Dibromochloromethane		15
1 1,1,2-Trichloroethane		5
I Bénzene		5
l cis-1,3-Dichloropropene	I ND	5
1 2-Chloroethylvinylether	I ND	10
l Bromoform		15
Tetrachloroethene		5
1 1,1,2,2-Tetrachloroethane		5
I Toluene	I ND	5
Chlorobenzene		1 5
I Ethylbenzene		5
1,3-Dichlorobenzene		5
1 1,2-Dichlorobenzene		1 5
1 1,4-Dichlorobenzene		- 5
1		

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIMI	TS	PERCENT RECOVERY
	WATER	SOIL	; }
Toluene-d8	88-110	81-117	
Bromofluorobenzene	86-115	74-121	
1,2-Dichloroethane-d4	76-114	70-121	98

ND: NOT DETECTED

CLIENT:CRASAMPLE#6	MATRIX: (SUIL/W	ATERN SHILL
LABORATORY NUMBER:8801276	SAMPLE WT/VOLUM	
DATE COLLECTED/RECEIVED: 8/18/88	UNIT (UG/L OR U	
DATE OF ANALYSIS:8/18/88	DILUTION FACTOR	
VOLATILE COMPOUNDS	CONCENTRATION	DETECTION LIMIT
Chloromethane	INU	_ _
Bromomethane	IND_	
Vinyl Chloride	1ND	II V
Chloroethane	IND	1101
Methylene Chloride	123	I5I
Trichlorofluoromethane	118	_15
1,1-Dichloroethene	1ND	_1'51
1,1-Dichloroethane	1ND	151
Trans-1,2-Dichloroethene	IND	
Chloroform	16	_151
1,2-Dichloroethane	1ND	_151
1,1,1-Trichloroethane	1ND	11
Carbon Tetrachloride	IND	_1
Bromodichloromethane	1ND	
1,2-Dichloropropane	IND	
Trans-1,3-Dichloropropene	1NU	5
Trichloroethene	IND	I
Dibromochloromethane	1NU	151
1,1,2-Trichloroethane	1ND	
Benzene	1ND	
cis-1,3-Dichloropropene	IND	_15
2-Chloroethylvinylether		_110
Bromoform	1ND	
Tetrachloroethene		_!5
1,1,2,2-Tetrachloroethane		_ !5
Toluene		
Chlorobenzene		_ [
Ethylbenzene	IU	
1,3-Dichlorobenzene	1ND	
1,2-Dichlorobenzene	IND	
l 1,4-Dichlorobenzene	!ND	5
SURROGATE RECO	DVERY DATA	
SURROGATE COMPOUND QC LIMI	ITS	PERCENT RECOVERY
UATER	SOIL	
Toluene-d8	81 - 117	93
1,2-Dichloroethane-d476-114	/₩=#&# 7n_191</td><td>97</td></tr><tr><td>1,2-0101101 00 (110110-04/0-114</td><td>, , , , / U=141, , , , , , , , , , , , , , , , , , ,</td><td> //</td></tr></tbody></table>	

ND: NOT DETECTED

CLIENT:CRA	
SAMPLE ID:SAMPLE#7	MATRIX: (SUIL/WATER)_SUIL
LABORATORY NUMBER:8801277	SAMPLE WIZUOLUME:5GH
DATE COLLECTED/RECEIVED:8/18/88	UNIT (UG/L OR UG/KG):_UG/KG_
DATE OF ANALYSIS:8/18/88	DILUTION FACTOR:1

1	VOLATILE COMPOUNDS	I CONCENTRATION	1 DETECTION LIMIT!
 	Chloromethane	IND	1101
l	Bromomethane	1ND	10
ļ	Vinul Chloride	I ND	101
l	Chloroethane	I ND	11 U
1	Methylene Chloride	154	l5
1	Trichlorofluoromethane	1111U	_15
1	1,1-Dichloroethene	IND	_I5
1	1,1-Dichloroethane	INU	_15
i	Trans-1,2-Dichloroethene	1ND	l5
1	Chloroform		_15
1	1,2-Dichlornethane	111ID	_15
1	1,1,1-Trichlorgethane	1NU	_15
1	Carbon Tetrachloride	INU	15
1	Bromodichloromethane	1ND	_I5
1	1,2-Dichloropropane		_15
1	Trans-1,3-Dichloropropene	1NU	_I5
1	Trichloroethene	IND	_15
1	Dibromochloromethane	1ND	_15
ŧ	1,1,2-Trichloroethane	IND	_15
1	Benzene	IND	_15
i	cis-1,3-Dichloropropene	IND	_15
١	2-Chloroethylvinylether	1_ND	_1LU
l	Bromoform		_15
ł	Tetrachloroethene		_15
١	1,1,2,2-Tetrachloroethane	1ND	_15
1	To luene	1ND	_15
1	Chlorobenzene	INU	_I5
ł	Ethylbenzene		_15
١	1,3-Dichlorobenzene		l5
ı	1,2-Dichlorobenzene		15
1	1,4-Dichlorabenzene		_!5

SURROGATE RECOVERY DATA

SURROGATE COMPUUND	QC LIMITS	õ	PERCENT RECOVERY
	WATER	SOIL	
			·
Toluene-d8	88-110	81-117	124
Bromofluorobenzene	86-115	74-121	
1.2-Dichlorgethane-d4	76-114		

ND: NOT DETECTED

PECPO EMPLIPHMENTAL, INC. COLUMBIA, MARYLAND VOCATTLE OPERANTOS ANALYSIS DATA SHEET

DATE COLLECTED/RECEIVED:			E: UBZL	
DATE OF HORICAGO:	37) (3 / (48)	Lilland Burk Field in 1625		
I VOLATILE COMPOUNDS		1 CUNCEMPATION	L DESERTION LIMIT	
Chloromethane	-	MD	111	
Bromomethane		1111	<u> </u>	
Vinyl Chloride		115	111	
Chloroethane		N1:	1 11	
Methylene Chloride			· ' —— —— · · —— —— —— ·	
Trichlorofluoromethane 1,1-Dichloroethene		1 <u>1 11)</u>		
1 1,1-Dichloroethane		1 10	5	
Trans-1,2-Dichlorgethene			· · · · · · · · · · · · · · · · · · ·	
			· · · · · · · · · · · · · · · · · · ·	
Chloroform_ 1,2-Dichloroethane			1 6.	
1 1,1,1-Trichloroethane		1 111:	5	
Carbon Tetrachloride) F.	
Bromodichloromethane		1 5111	1 5	
1 1,2-Dichloropropane			, <u> </u>	
Trans-1,3-Dichloropropen	<u> </u>		1 5	
Trichloroethene	'-'	h:f)	1	
Dibromochloromethane		1 110	1 6	
1 1,1,2-Trichlorosthans			f-	
Benzene			5	
cis-1,3-Dichloropropens_			1 15	
1 2-Chloroethylvinylether_		1 100	1 1.0	
Bromoform_		14[1	1 E	
Tetrachloroethene		htp.	1 19	
1 1,1,2,2-letrachloroethan		1 140	5	
I Toluene		1	1 5	
Chlorobenzene		111.)	1	
Ethylbenzene		110	1	
1 1,3-Dichlorobenzene		(<u> </u>	
1 1,2-Dichlorobenzene		11	[
1,4-Dichlorobenzene				
Su	RRUGATE REC			
SURROGATE COMPOUND	ÇC L11		PERCENT RECOURRY	
	WATER	SOIL	and the same and t	
Toluene-d3	86-115	74-121	1 02	

ND: NOT DETECTED

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND VOLATILE ORGANICS ANALYSIS DATA SHEET

CLIENT:CRA_ SAMPLE ID:SAMPLE#1(MS)_ LABORATORY NUMBER:8801271_ DATE COLLECTED/RECEIVED:8/18/88 DATE OF ANALYSIS:8/18/88	MATRIX: (SUIL/WA SAMPLE WT/VOLUME UNIT (UG/L OR UG DILUTION FACTOR:	:5GM SZKGJ:_UGZKG_
VOLATILE COMPOUNDS	I CONCENTRATION	I DETECTION LIMIT
Chloromethane	1 ND	1 10
Bromomethane	I ND	10
Vinyl Chloride	I ND	10
i Unioroethane	IND	_I1
Methylene Chloride	16	5
Trichlorofluoromethane	l6	l:5
1,1-Dichloroethene	1	_l`5
1,1-Dichloroethane		_15/
Trans-1,2-Dichloroethene	IND	_I5
l Chloroform	I	_I5
1 1,2-Dichloroethane	IND	I5
l 1,1,1-Trichloroethane	1ND	_15
Carbon Tetrachloride	IND	_I5
Bromodichloromethane	1ND	_l5
1,2-Dichloropropane		_I5
1 Trans-1,3-Dichloropropene		_15
Trichloroethene		l 5
Dibromochloromethane		_I
1 1,1,2-Trichloroethane		_I5
Benzene	163	_15
cis-1,3-Dichloropropene	1ND	_I5

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIM	ITS	PERCENT RECOVERY
	WATER	SOIL	:
Toluene-d8			
Bromofluorobenzene 1,2-Dichloroethane-d4			

1υ_

5

5

5.

5_

5_

ND_

ND

ND_

NU_

68

5 U

ND.

ND

ND

ND_

ND: NOT DETECTED

* : PROBABLE CONTAMINATION

2-Chloroethylvinylether_

1,1,2,2-Tetrachloroethane_

Tetrachloroethene_

1,3-Dichlorobenzene_

1,2-Dichlorobenzene_

1,4-Dichlorobenzene_

Bromoform_

Toluene___

Chlorobenzene

Ethylbenzene_

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND VULATILE ORGANICS ANALYSIS DATA SHEET

CLIENT:CRA		
SAMPLE ID: SAMPLE#1(MSD)	_ MATRIX: (SOIL/WA	ATER) SOIL
LABORATORY NUMBER:8801271	SAMPLE WT/VOLUME	
DATE COLLECTED/RECEIVED: 8/18/88	UNIT (UG/L OR UG	
DATE OF ANALYSIS:8/18/88		
	_	;
I VOLATILE COMPOUNDS	I CONCENTRATION	IDETECTION LIMITE
		.
Chloromethane		
I Bromomethane	1ND	_!!
Vinyl Chloride	IND	_1101
Chloroethane	1NÜ	_
Methylene Chloride	IL U	_151
Trichlorofluoromethane	1ND	_151
1,1-Dichloroethene	181	_1
1 1,1-Dichloroethane	UUU	_151
I Trans-1,2-Dichloroethene	IND	_11
Chloroform	18	_!!
1,2-Dichloroethane	IND	_[
1,1,1-Trichloroethane	1ND	_
Carbon Tetrachloride	IND	51
Bromodichloromethane	1ND	_11
1,2-Dichloropropane		5
I Trans-1,3-Dichloropropene		_II
Trichloroethene		J5J
Dibromochloromethane		J 5 I
I 1,1,2-Trichloroethane		151
I Benzene		_ I
cis-1,3-Dichloropropene		
1 2-Chloroethylvinylether	IND	101
1 Bromoform		
Tetrachloroethene	I ND	
1 1,1,2,2-Tetrachloroethane		
I Toluene		
Chlorobenzene		1
I Ethylbenzene		5
1,3-Dichlorobenzene	I ND	1 5 1
I 1,2-Dichlorobenzene	I NU	1 5 1
1 1,4-Dichlorobenzene	I ND	1 5
		1
SURROGATE RE	COVERY DATA	
SURROGATE COMPOUND QC LI	MITS	PERCENT RECOVERY
	· · · · · · -	

WATER

SOIL

ND: NOT DEFECTED

^{*:} PROBABLE CONTAMINATION

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SUIL

Sample Identification: SAMPLE # 1	Ulient: UHA
Laboratory Humber: 88#1271	Dilution Factor: _1
Date Collected/Reserved:_8/18/88	Date of Analysis:_8/19/88

	1 Concentration	 1 Detection Limit
Semivolatile Compounds	l (ug/kg)	t (ug/kg)
N-Nitrosodimethylamine		330
Fhenol		_1330
bis(-2-Chloroethyl)Ether		330
2-Chlorophenol	!ND	339/
1,3-Dichlorobenzene		330/
1,4-Dichlorobenzene	!t1D	_!330
1,2-Dichlorobenzene	ND	350
bis(2-Chloroisopropy))ether		133U
N-Nitroso-Di-n-propylamine	!ND	1330
Hexachloroethane	NO	330
Nitrobenzene	1ND	133.0
Isophorane	1ND	1330
2-Nitrophenol	IN()	1830
2,4-Dimethylphenol	111	1830
bis(-2-Chloroethoxy)Methane	1ON	1330
2,4-Dichlorophenol	1ND	1330
1,2,4-Trichlorobenzene	1NO	1339
Naphthalene	1 <u>NO</u>	3311
Hexachlorobutadiene	1ND	1330
4-Chloro-3-methylphenol	1 10	1 8311
Hexachlorocyclopentadiene	1 10	370
2,4,6-Trichlorophenol	I NO	330
2-Chloronaphthalene	1 10	330
Dimethyl Phthalate	I ND	1 350
Acenaphthylene	I MO	330
Acenaphthene	I ND	330
2,4-Dinitrophenol	1 ND	1 830
4-Nitrophenol	I ND	1 830
2,4-Dinitrotoluene	T ND	330
2,6-Dinitrotoluene	I ND	1 330
Diethylphthalate	ND	330
4-Chlorophenyl-phenylether	I ND	330
Fluorene	I ND	330
4,6-Dinitro-2-methylphenol	I ND	1 8311
N-Nitrosodiphenylamine	1 110	330
4-Bromophenyl-phenylether	1 14D	1 330
Hexachlorobenzene	1 NU.	3.50

Sample Identification:_SAMPLE # 1_____

 Semivolatile Compounds 	Concentration (ug/kg) 	Detection Limit
 Pentachlorophenol	I ND	1 830 1
I Phananthrene	I ND	330
i Anthracene	1 NO	330
Di-n-Butylphthalate	1ND	330
Fluoranthene	IND	350 1
1 Pyrene	1ND	_1 330 1
Benzidine	110	1 830 :
Butylbenzylphthalate	1ND	_13301
1 3,3'-Dichlorobenzidine	IND	_1 <u>830</u> 1
Benzo(a)Anthracene	1ND	_13301
Bis(2-Ethylhexyl)Phthalate	IND	11
Chrysene	IND	_173,01
Di-n-octyl phthalate	111)	_13301
l'Benzo(b)fluoranthene	1ND	13301
Benzo(k)fluoranthene	110	330
Benzo(a)Pyrene	1ND	330
Indeno(1,2,3-cd)Pyrene	1ND	330
Dibenzo(a,h)Anthracene	1ND	_13301
Benzo(g,h,i)Perylene	INC:	330

SUPRUGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Recovery
Nitrobenzene-d5	(23-120)	73
2-Fluorobiphenyl		
Terphenyl-d14		
Phenol-d5		
2-Fluorophenol	(25-121)	48
2,4,6-Tribromophenol.	(19-122)	

ND: NOT DETECTED

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SUIL

Sample Identification: SAMPLE # 2	Client: 000
Laboratory Number: 8801272	Milution Factor: t
	Date of Analysis: 8/19/88

	Concentration	I Detection Limit
Semivolatile Compounds	l (ug/kg) l	l Gugakgi T
N-Nitrosodimethylamine	1 NO	1 1 330
Pheno!	1 115	330
bis(-2-Chloroethyl)Ether	ND	3311
2-Chlorophenol	I ND	330
1,3-Dichlorobenzene	ND	330 /
1,4-Dichlorobenzene	1 140	330/
1,2-Dichlorobenzene	1 140	330
bis(2-Chlorossopropylether	1 110	330
N-Nitroso-Di-n-propylamine	NU	3311
Hexachloroethane	NO NO	330
Nitrobenzene	1 140	330
Isophorone	I ND	330
2-Nitrophenol	NC NC	1 8311
2,4-Dimethylphenol	1 11(1)	1 830
bis(-2-Chloroethoxy)Methane	I MD	371)
2,4-Dichlorophenol	1 14()	330
1,2,4-Trichlorobenzene	I ND	1 330
Naphthalene	I NLI	330
Hexachlorobutadiene	1 40	1 63.0
4-Chloro-3-methylphenol	ND	830
Hexachlorocyclopentadiene	T NO	330
2,4,6-Trichlorophenol	I ND	350
2-Chloronaphthalene	1 110:	330
Dimethyl Phthalate	1 110	330
Acenaphthylene	1 10	331)
Acenaphthene	I ND	330
2,4-Dinitrophenol		1 870
4-Nitrophenol	1 110	1 830
2,4-Dinitrotoluene	1 140	(331)
2,6-Dinitrotoluene	i 11D	330
Diethylphthalate	1 10	1 371)
4-Chlorophenyl-phenylether	1 110	1 3311
Fluorene	I NO	339
4,6-Dinitro-2-methylphenol	I ND	1 8311
N-Nitrosudiphenylamina	I ND	3 3 1)
4-Bromophenyl-phenylether	I NO	330
Hexach Lonobenzene	NO.	330

Sample Identification: SAMPLE # 2_____

	1 Concentration	1 Detection Limit
Semivolatile Compounds	l (ug∠kg)	l (ug/kg) i
	1	
1 Pentachlorophenol	140	830_11
I Phenanthrene	11100	13301
Anthracene	IND	7.31)
I Di-n-Butylphthalate	1_ND	1350
Fluoranthene	11000	13301
I Pyrene	I1700	13301
Benzidine	110	183 0
Butylbenzylphthalate	ttdD	1338
1 3,3'-Dichlorobenzidine	1 140	1 830
Benzo(a)Anthracene	1 860	3311
Bis(2-Ethythexyl)Phthalate	1140	351/
1 Chrysene	1930	133,0
Di-n-octyl phthalate	11	1330
I-Benzo(b)fluorantheme	1100	1 330
Benzo(k)fluoranthene	I NO	330
I Benzo(a)Pyrene	1 680	330
I Indeno(1,2,3-cd)Pyrene	1 140	330
Dibenzo(a,h)Anthracene		3 1 0
Benzo(g,h,i)Perylene	1140	330
	1	

SUPPOGATE RECOVERY DATA

Surrogate Compound	QC Limits	Perpent Recov⇔ry
Nitrobenzene-d ⁶	(23-128)	
2-Fluorobiphenyl	(30-115)	
Terphenyl-dl4 Phenol-d9		
2-Fluorophenol		
2,4,6-Tribromophenol.	(19-122)	

ND: NOT DETECTED

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification: SAMPLE # 3	Client: DFA
Laboratory Number: _8801273	Dilution Factor:1
Date Collected/Received:_8/18/88	Date of Analysis:_8/18/88

	l Concentration	1 Detection Limit
Semivolatile Compounds	(ug/kg) 	(ug/kg)
N-Nitrosodimethylamine		1 330
Phenol	TI ND	330
bis(-2-Chloroethyl)Ether	NO NO	3.31)
2-Chlorophenol		330
1,3-Dichlorobenzene	110	330/
1,4-Dichlorobenzene	111	330
1,2-Dichlorobenzene	1NO	1 330
bis(2-Chloroisopropyl)ether	110	_1330
N-Nitroso-Di-n-propylamine	IMO	331)
Hexachloroethane	1HD	330
Nitrobenzene	1NO	1330
Isophorone	1ND	<u>1330</u>
2-Nitrophenol	1101	1930
2,4-Dimethylphenol	1ND	1 830
bis(-2-Chloroethoxy)Methane	1NO	1330
2,4-Dichlorophenol	1 ND	1 330
1,2,4-Trichlorobenzene	IND	1 330
Naphthalene	1100	
Hexachlorobutadiene	110	1 ×3 n
4-Chloro-3-methylphenol	I MD	1 830
Hexachlorocyclopentadiene	I NO	330
2,4,6-Trichlorophenal	I NO	330
2-Chloronaphthalene	TI HD	330
Dimethyl Phthalate	1 ND	3311
Acenaphthylene	1 NO	1 330
Acenaphthene	I NO	1 330
2,4-Dinitrophenol	I ND	1 830
4-Nitrophenol	I NO	1 830
2,4-Dinitrotoluene	I NO	1 331
2,6-Dinitrataluene	1 . NO	331
Diethylphthalate		330
4-Chlorophenyl-phenylether	1 ND 1	730
Fluorene	1 10	330
4,6-Dinitro-2-methylphenol	I ND	1 <u>83</u> 0
N-Nitrosodiphenylamine	I ND	330
4-Bromophenyl-phenylether	1 ND	
Hexach Lorobenzene	ND	3311

Sample Identification: SAMPLE # 3_____

	l Concentration	1 Detection Limit
Semivolatile Compounds	l (ug/kg)	(ug/kg) (
Pentachlorophenol	110	1830
Phenanthrene	1ND	I330I
Anthracene	1ND	133.0
Di-n-Butylphthalate	1140	1
Fluoranthene	1ND	1331)
I Pyrene	11D	133U
Benzidine	10M1	1
Butylbenzylphthalate	IND	1330
1 3,3'-Dichlorobenzidine	1ND	1830
Benzo(a)Anthracene	1110	l
Bis(2-Ethylhexyl)Phthalate	1ND	133 <u>1/</u> _
Chrysene	1110	133,0
Di-n-octyl phthalate	1110	1
l, Benzo(b) fluoranthere	IND	1330
Benzo(k)Fluoranthene	1111111	1 330
Benzo(a)Pyrene	1ND	_1330
Indeno(1,2,3-od)Pyrene	IND	1330
Dibenzo(a,h)Anthracene	11ID	_1330
Benzo(g,h,i)Perylane	1NO	730
1		

SURPOGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Recovery
·		
•		61

ND: NOT DETECTED

PECRA ENGINUMENTAL, INC. CULUMBIA, MARYLAND

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification: SAMPLE # 4	Client: CPA
Laboratory Number: 8801274	Dilution Factor: 1
Date Collected/Reneived: 8/18/88	Date of Analysis: 8/18/88

	Concentration	I Detection Limit
Semivolatile Compounds	l (ug/kg)	(ug/kg)
		,)
N-Nitrosodimethylamine		330
Pheno I	1ND	1
bis(-2-Chloroethyl)Ether		13311
2-Chlorophenol	ND	_1330
1,3-Dichlorobenzene	1	1
1,4-Dichlorobenzene	114D	_1350/
1,2-Dichlorobenzene		13311
bis(2-Chloroisopropyl)ether	ND	1330
N-Nitroso-Di-n-propylamine	IND	13311
Hexachlorgethane	IND	1330
Nitrobenzene	INU	1330
Isophorone	1ND	133.0
2-Nitrophenol	1NO	_1830
2,4-Dimethylphenol	1ND	1830
bis(-2-Chloroethoxy)Methane	I ND	1 3311
2,4-Dichlarophenol	II	330
1,2,4-Trichlorobenzene	1 140	330
Naphthalene	I ND	330
Hexachlorobutadiene	I ND	330
4-Chloro-3-methylphenol	1 140	B30
Hexachlorocyclopentadiene	I ND	350
2,4,6-Trichlorophenol	1 ND	330
2-Chloronaphthalene	I ND	330
Dimethyl Phthalate	1 110	33.0
Acenaphthylene	NO NO	331)
Acenaph thene	ND	330
2,4-Dinitrophenol	NO NO	830
4-Nitrophenol	l ND	1 8311
2,4-Dinitrotoluene	N()	3 0
2,6-Dinitrotoluene	1 - 14D	330
Diethylphthalate	1 140	330
4-Chlorophenyl-phenylether	1 140	1 330
Fluorene	I ND	1 3311
4 4 Dimitus 9 sethulahasal	l ND	1 830
N-Nitrosodiphenylamine	I ND	3.70
4-Bromophenyl-phenylether	ND	330
Hexach Lorobenzene		330

Sample Identification: SAMPLE # 4_____

	Concentration	1 Defection Limit
Semivolatile Compounds	l (ug/kg)	(ug/kg)
		1
Pentachlorophenol	1N0	1
Phenanthrene	1ND	1330
Anthracene	1110	
Di-n-Butylphthalate	1110	1
Fluoranthene	IND	1330
Pyrene	1	330
Benzidine	IND	830
Butylbenzylphthalate	1ND	1330
3,3'-Dichlorobenzidine	1ND	1830
Benzo(a)Anthracene	I!JD	(<u></u>
Bis(2-Ethylhexyl)Phthalate	1ND	I33 <i>I</i> /
Chrysene	1ND	1
Di-n-octyl phthalate	IND	1330
-Benzo(b)fluoranthene	1ND	1
Benzo(k)fluoranthene	ND	1330
Benzo(a)Pyrene	110	1350
I Indeno(1,2,3-cd)Pyrene	1 110	330
Dibenzo(a,h)Anthracene	114D	1330
Benzo(g,h,i)Perylene	INU	

SURRUGATE RECOVERY DATA

Surrogate Compound	QC Limits	Pencent Panovery
Nitrobenzene-d5		
2-Fluorobiphenyl	(30-115)	,
Terphenyl-dl4		
Phenol-d5	(24-113)	, 63
2-Fluorophenol	(25-121)	
2,4,6-Tribromophenol.	(19-122)	102

ND: NOT DETECTED

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification: SAMPLE # 5	Client:_U/A
Laboratory Number:_8801275	Dilution Factor:1
Date Collected/Received: 8/18/89	Date of Acalysis: 8/19/88

0	1 Concentration	Detection Limit
Semivolatile Compounds	(ug/kg) 	(ug/kg) (
N-Nitrosodiwethylamine	1 1MD	1 _1330
Pheno I	IID	330
bis(-2-Chloroethyl)Ether	1 140	330
2-Chlorophenol	IND	t330 <u>/</u>
1,3-Dichlorobenzene	1ND	1330 <u>/</u>
1,4-Dichlorobenzene	1NO	133 <i>tr</i> _
1,2-Dichlorobenzene	1ND	1330
<pre>bis(2-Chloroisopropyl)ether</pre>	1ND	_1330
N-Nitroso-Di-n-propylamine	1 10	330
Hexachloroethane	110	1 330
Nitrobenzene	I NO	330
Isophorone	1ND	330
2-Nitrophenol	1 140	830
2,4-Dimethylphenol	l ND	830
bis(-2-Chloroethoxy)Methane	I NO	330
2,4-Dichlorophenol	1 110	3311
1,2,4-Trichlorobenzene	I MD	330
Naphthalene	1 1/2)	330
Hexachlorobutadiene	1 110	330
4-Chloro-3-methylphenol	1 ND	830
Hexachlorocyclopentadiene	I ND	330
2,4,6-Trichlorophenol	1 ND	330
2-Chloronaphthalene	I ND	330
Dimethyl Phthalate	1 110	330
Acenaphthylene	1 NO	330
Acenaphthene	ND ND	1 330
2,4-Dinitrophenol	I NO	1 830
4-Nitrophenol	l ND	830
2,4-Dinitrotoluene	I ND	330
2,6-Dinitrotoluene	I NU	330
Diethylphthalate	I NO	3 5 11
4-Chlorophenyl-phenylether	I ND	330
Fluorene	1 140	330
4,6-Dinitro-2-methylphenol	I ND	1 830
N-Nitrosodiphenylamine	I NO	330
4-Bromophenyl-phenylether	1 110	330
Hexach lorobenzene	ND -	330

Sample Identification: SAMPLE # 5_____

	1 Concentration	1 Detection Limit 1
Semivolatile Compounds	t (ug∠kg) L	(ug/kg) 1
Pentachlorophenol	IND	1930!
Phenanthrene	IHD	
Anthracene	1ND	1
Di-n-Butylphthalate	1ND	tsant
Fluoranthena	1NO	3311
Pyrene	1	1
Benzidine	1110	1
Butylbenzylphthalate	1ND	330
3,3'-Dichlorobenzidine	1NU	1830
Benzo(a)Anthracene	110	1
Bis(2-Ethylhexyl)Phthalate	1NO	1
Chrysene	1ND	133.b
Di-n-octyl phthalate	1NO	
l'Benzo(b)fluoranthene	1360	1330
Benzo(k)fluoranthene	1ND	1
Benzo(a)Pyrene	I	1330
Indeno(1,2,3-cd)Pyrene	1ND	1330
Dibenzo(a,h)Anthracene	11D	1330
Benzo(g,h,ı)Ferylene	INU	330

SUBROGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Pacovery
2-Fluorobiphenyl	(30-115)	
Phenol-d5	(24-113)	50 50

ND: NOT DETECTED

SEMIVOLATILE URGANICS ANALYSIS DATA SHEET FOR SUIL

Sample Identification: SAMPLE # 6	Client: CRA
Laboratory Number:_8801276	Dilution Factor: 1
Date Collected/Received:_8/18/88	Date of Analysis: 8/18/88

	l Concentration	1 Detection Limit
Semivolatile Compounds	l (ug∕kg)	l (ug∠kg)
		·····
N-Nitrosodimethylamine	1ND	370
PhenoI	I NO	_133·0
bis(-2-Chloroethyl)Ether	1ND	330
2-Chlorophenol	I ND	330
1,3-Dichlorobenzene	1 NO	33 y <u>/</u>
1,4-Dichlorobenzene	1ND	330/
1,2-Dichtorobenzene	1NU	1 330
bis(2-Chloroisopropyl)ether	1ND	330
N-Nitroso-Di-n-propylamine	1ND	330
Hexachloroethane	ND ND	_1330
Nitrobenzene	IU	330
Isophonone	I ND	330
2-Nitrophenol	1ND	830
2,4-Dimethylphenol	1 110	830
bis(-2-Chloroethoxy)Methane	I NO	330
2,4-Dichlorophenol	1 140	330
1,2,4-Trichlorobenzene	1 110	330
Naphthalene	370	330
Hexachlorobutadiene	I NO	320
4-Chloro-3-methylphenol	ND ND	830
Hexachlorocyclopentadiene	I NO	3 3 11
2,4,6-Trichlorophenol	I ND	1 33 U
2-Chloronaphthalene	I ND	3311
Dimethyl Phthalate	I NO	330
Acenaphthylene	TO NO	1 330
Acenaphthene	I ND	1 33ú
2,4-Dinitrophenol	I NC	1 830
4-Nitrophenol	1 110	8.40
2,4-Dinitrotoluene	I ND	1 33 u
2,6-Dinitrotoluene	I NU	330
Diethylphthalate	1 110	330
4-Chlorophenyl-phenylether	I ND	330
Fluorene	1 NO	330
4,6-Dinitro-2-methylphenol	I ND	830
N-Nitrosodiphenylamine	I ND	330
4-Bromophenyl-phenylether	I ND	1 330
Hexachlorobenzene	I ND	330

Sample Identification: SAMPLE # 6_____

1	Semivolatile Compounds	Concentration (ug/kg) 	Detection Limit
1			
ı	Pentachlorophenol	1ND	930
1	Phenanthrene	I510	330
ı	Anthracene	1NO	1 330
ŧ	Di-n-ButyIphthalate	1ND	330
1	Fluoranthene	1320	1 330
1	Pyrene	1 420	330
1	Benzidine	IND	1 830
1	Butylbenzylphthalate	IND	330
1	3,3'-Dichlorobenzidine	1ND	1830
ı	Benzo(a)Anthracene	1ND	_1330_/
i	Bis(2-Ethylhexyl)Phthalate	1	<u>. 133p(</u>
1	Chrysene	1N0	133/0
١	Di-n-octyl phthalate	IND	1330
ŀ	Benzo(b)flugranthene	1320	330
1	Benzo(k)fluoranthene	1N0	1330
1	Benzo(a)Pyrene	1ND	330
1	<pre>Indeno(1,2,3-cd)Pyrene</pre>	I ND	1330
ı	Dibenzo(a,h)Anthracene	NDND	1330
1	Benzo(g,h,i)Perylene	IND	3.50

SURHOGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Recovery
Nitrobenzene-d5	(23-120)	83
2-Fluorobiphenyl	(30-115)	
Terphenyl-d14	(18-137)	
Phenol-d5	(24-113)	
2-Fluorophenol	(25-121)	
2.4.6-Tribromophenol.	(19-122)	

ND: NOT DETECTED

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification: SAMPLE # 7	Client:_UPA
Laboratory Number:_8801272	Dilution Factor:1
Date Collected/Received: 8/18/88	Date of Analysis: 8/19/88

	1 Concentration	1 Detection Limit
Semivolatile Compounds	l (ug/kg)	l (ug/kg)
At his		
N-Nitrosodimethylamine		
Phenol		!33 U
bis(-2-Chloroethyl)Ether	NO	30
2-Chlorophenol_	ND	_1330
1,3-Dichlorobenzene	ND	330/
1,4-Dichlarobenzene		331/
1,2-Dichtorobenzene		3.50
bis(2-Chloroisopropyl)ether	ND	330
N-Nitroso-Di-n-propylamine	ND	1330
Hexachloroethane		330
Nitrobenzene	140	_1330
Isophorone	NDND	1330
2-Nitrophenol	INÜ	B30
2,4-Dimethylphenol	1110	I830
bis(-2-Chloroethoxy)Methane	1NÜ	13311
2,4-Dichlorophenol	1ND	1330
1,2,4-Trichlorobenzene	1NO	1
Naphthalene	111	_1330
Hexachiorobutadiene	1NO1	330
4-Chloro-3-methy/pheno/	110	970
Hexachlorocyclopentadiene	1 NÖ	330
2,4,6-Trichlorophenol	1ND	331
2-Chloronaphthalene	1NU	330
Dimethyl Phthalate	I ND	1 330
Acenaphthylene	1 NO	330
Acenaphthene	1 <u>ND</u>	1330
2,4-Dinitrophenol	11MD1	830
4-Nitrophenol	1ND	1 830
2,4-Dinitrotoluene	1NO	1 330
2,6-Dinitrotoluene	IND	1330
Diethylphthalate	1 110	3.30
4-Chlorophenyl-phenylether	1 110	330
Fluorene	I ND	330
4,6-Dinitro-2-methylphenol	1 ND	830
N-Nitrosodiphenylamine	I NU	330
4-Bromophenyl-phenylether	I ND	33 u
Hexachlorobenzene	I ND	330

Sample Identification: SAMPLE \$ 7

	Concentration	Detection Limit
Semivolatile Compounds -	l (ugzkg) I	l (ug∠kg) / I
Don't and Lauran and L	1 146	1 830
Pentachlorophenol	1 140	
Phenanthrens		3311
Anthracene	I ND	330
Di-n-Butylphthalate		330
Fluoranthene	1 14E)	
Pyrene		
Benzidine		930
Butylbenzy)phthalate	<u> </u>	
3,3'-Dichlorobenzidine	ND	[H31]
Benzo(a)Anthracene	ND	-1
Bis(2-Ethylhexyl)Phthalate	!ND	
Chrysene	!ND	_1330
Di-n-octyl phthalate	!HD	13311
Benzo(b)fluoranthene	1ND	330
Benzo(k)fluoranthene	ND	330
l Benzo(a)Pyrene	1ND	1
Indeno(1,2,3-cd)Pyrene	1ND	1330
Dibenzo(a,h)Anthracene	1ND	
Benzo(g,h,ı)Perylene	1ND	1330
l	<u>.</u>	t

SURROGATE RECOVERY DATA

Surrogate Compound	QU Limits	Percent Recovery
Nitrobenzene-d5		80
2-Fluorobiphenyl	(30-115)	80
Terphenyl-d14	(18-137)	
Pheno 1-d5	(24-113)	63
2-Fluorophenol	(25-121)	
2.4.6-Tribromophenol.		

ND: NOT DETECTED

SEMIUOLATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification: SAMPLE # 7 MS	Client:_UPA
Laboratory Number: _8801227 MS	Dilution Factor:1
Date Collected/Received:_8/18/88	Date of Analysis: 8/19/88

	Concentration	I Detection Limit
Semivolatile Compounds	(ug/kg \ ((ug/kg) (
N-Nitrosodimethylamine	 	11 1330
Pheno 1	13100 (M)	13311
bis(-2-Chloroethyl)Ether	1ND	1330
2-Chlorophenol	I2500 (M)	_1330 <u></u>
1,3-Dichlorobenzene	IND	_1330_/
1,4-Dichlorobenzene	I1300 (M)	_133n <u>/</u>
1,2-Dichlorobenzene	1ND	1330
bis(2-Chloroisopropyl)ether	1ND	_1330
N-Nitroso-Di-n-propylamine	I1200 (M)	
Hexachloroethane	110	
Nitrobenzene	1ND	1330
Isophorone	IND	1331J
2-Nitrophenol	1ND	_ (원칙()
2,4-Dimethylphenol	1ND	_1830
bis(-2-Chloroethoxy)Methane	I ND	13311
2,4-Dichlorophenol	1HD	1330
1,2,4-Trichlorobenzene	11100 (M)	1330
Naphthalene	11	_13311
Hexachlorobutadiene	1ND	1330
4-Chloro-3-methylphenol	12600 (M)	1830
Hexachlorocyclopentadiene	1ND	1330
2,4,6-Trichlorophenol	I ND	1330
2-Chloronaphthalene	1ND	1330
Dimethyl Phthalate	IND	1 330
Acenaphthylene	I NÜ	1 330
Acenaphthene	1500 (M)	1 330
2,4-Dinitrophenol	T NO	830
4-Nitrophenol	880 (M)	1 830
2,4-Dinitrotoluene	1 1100 (11)	
2,6-Dinitrataluene	I - ND	1330
Diethylphthalate	. I ND	330
4-Chlorophenyl-phenylether		330
Fluorene	T NO	330
4,6-Dinitro-2-methylphenol		1 830
N-Nitrosodiphenylamine	I ND	1 330
4-Bromophenyl-phenylether		330
Hexachlorobenzene	1 ND	330

Sample Identification: SAMPLE # 7_MS____

 Semivolatile Compounds 	Concentration (ug/kg) 	Defection Limit
 Pentachlorophenol	1 3000 (M)	1 830 I
I Phenanthrene	1 110	330 1
Anthraceme	1NO	3301
Di-n-Buty-lphthalate	110	330
Fluoranthene	1101	330 1
Fyrene	1 2140 (M)	330 1
Benzidine	1101	1 930 1
Butylbenzylpfithalate	IND	13301
1 3,3'-Dichlorobenzidine	11	
Benzo(a)Anthracene	1ND	3301
Bis(2-Ethylhexyl)Phthalate	IND	1133/d1
Chrysene	1ND	11
Di-n-octyl phthalate	1ND	13301
l Benzo(b) fluoranthene	1ND	3301
Benza(k)fluoranthene	1ND	13301
l Benzo(a)Pyrrene	IND	13301
Indeno(1,2,3-cd)Pyrene	1N0	13301
Dibenzo(a,h)Anthracena	I ND	133U1
Benzo(g,h,1)Perylene	NO	3301

SURROGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Pecovery
Nitrobenzene-d5	(23-120)	
2-Fluorobiphenyl	(30-115)	81
Terphenyl-d14		
Phenol-d5		
2-Fluorophenol		
2,4,6-Tribromophenol.	(19-122)	.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,

M : MATRIX SPIKE COMPOUND

ND: NOT DETECTED

Sample Identification: SAMPLE # 7_MSD____

 Semivolatile Compounds 	Concentration (ug/kg) 	Defection Limit
 Pentachlorophenol		830
I Phenanthrene	1ND	_13301
Anthracene	INU	_11
Di-n-Butylphthalate	1ND	13301
Fluoranthene	1ND	13301
I Pyrene	12400 (M)	330
Benzidine	110	1 830 1
Butylbenzylphthalate	1ND	330 1
3,3'-Dichlorobenzidine	IND	_18301
Benzo(a)Anthracene	1ND	330
Bis(2-Ethylhexyl)Phthalate	1ND	331/
Chrysene	1NO	330
[Di-n-octyl phthalate	1ND	330
Benzo(b)fluoranthene	1 10	330 1
Benzo(k)fluoranthene	IND	330
Benzo(a)Pyrene	1ND	330
Indeno(1,2,3-cd)Pyrene	IND	330
Dibenzo(a,h)Anthracene	1140	330
Benzo(g,h,i)Perylene 	IND	330

SURROGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Mecquery
Nitrobenzene-d5	(23-120)	
2-Fluorobiphenyl		
Terphenyl-dl4	(18-132)	
Phenol-d5	(24-113)	
2-Fluorophenol	(25-121)	
2,4,6-Tribromophenol.		

M : MATRIX SPIKE COMPOUND

ND: NOT DETECTED

PECRA EMPTRUMMENTAL, IDC. COLUMBIA, MARYLAND

SEMIVOLATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification: SAMPLE # 7_MSD	Client:_CBA
Laboratory Number: 8801272_MSD	Dilution Factor:l
Date Collected/Received:_8/18/88	Date of Analysis:_0/19/88

	1 Concentration	I Detection Limit
Semivolatile Compounds	l (ug/kg) l	il (ug/kg) il
N. N. A. a. a. d. i. a. k. a. l. a. a. a. a.	I NO	1 3311
N-Nitrosodimethylamine		
Phenolbis(-2-Chloroethyl)Ether	1 32110 (H1) NO	330
		3311 /
2-Chlorophenol		330/
1,4-Dichlorobenzene		
1,2-Dichlorobenzene		
bis(2-Chloroisopropyl)ether	110	1331 <u></u> 133U
N Nitross D. n. sassilaria	1 1900 (M)	730
N-Nitroso-Di-n-propylamine		
Hexachioroethane		
Nitrobenzene		330
Isophorone		
2-Nitrophenol	ND	930
2,4-Dimethylphenol		83.0
bis(-2-Chloroethoxy)Methane	!	1330
2,4-Dichlorophenol	111D	330
1,2,4-Trichlorobenzene	1200 (11)	3.50
Naphthalene	ND	1330
Hexachlorobutadiene	1NU	1330
4-Chloro-3-methylphenol	12700 (M)	1930
Hexachlorocyclopentadiene	1NU	1330
2,4,6-Trichlorophenol	1ND	_137,0
2-Chloronaphthalene	1ND	1330
Dimethyl Phthalate	IHD	1330
Acenaphthylene	tND	I33 U
Acenaph thene	1_11600 (11)	I
2,4-Dinitrophenol	11111	1830
4-Nitrophenol	1800 (11)	
2,4-Dinitrotoluene	t1100 (M)	1330
2,6-Dinitrotoluene	1ND	
Diethylphthalate	1NO	330
4-Chlorophenyl-phenylether	1 140	3311
Fluorene	l NO	3311
4,6-Dinitro-2-methylphenol	I ND	830
N-Mitrosodiphenylamine	ND ND	330
4-Bromophenyl-phenylether	I ND	33 U
Hexachlorobenzene	I ND	330

PECRA EMUIPOMMENTAL, INC. COLUMBIA, MARYLAND

SEMIVULATILE ORGANICS ANALYSIS DATA SHEET FOR SOIL

Sample Identification:_METHOD BLANK	Client:_UBA
Laboratory Number: # 231	Dilution Factor: 1
Date Collected/Received:_8/18/88	Date of Analysis:_8/18/83

	l Concentration	1 Detection Limit
Semivolatile Compounds	(ug/kg) 	(ug/kg) _t
N-Nitrosodimethylamine	1 1ND	1 133!!
Pheno1	11	_1330
bis(-2-Chloroethyl)Ether	1ND	330
2-Chlorophenol	1NO	1 330
1,3-Dichlorobenzene	1ND	1
1,4-Dichlorobenzene	1ND	133u [/]
1,2-Dichlorobenzene	110	1 330
bis(2-Ch)oroisopropyl)ether	1ND	_13311
N-Nitroso-Di-n-propylamine	1ND	1 330
Hexachloroethane	1 ND	330
Nitrobenzene	I ND	330
Isophorone	1NO	3311
2-Nitrophenol	t tho	1 8311
2,4-Dimethylphenol	I ND	1 930
bis(-2-Chloroethoxy)Methane	1 140	1 330
2,4-Dichlorophenol	I ND	1 330
1,2,4-Trichlorobenzene	1 NO	3.30
Naphthalene	I ND	330
Hexachlorobutadiene	IND	320
4-Chloro-3-methylphenol	110	1 830
Hexachlorocyclopentadiene	1ND	330
2,4,6-Trichlorophenol	11601	1 330
2-Chloronaphthalene	11	1 330
Dimethyl Phthalate	I ND	1 330
Acenaphthylene	111111	1 3311
Acenaphthene	I ND	1 330
2,4-Dinitrophenol	1ND	1830
4-Nitrophenol	110	_1830
2,4-Dinitrotoluene	NO	133.0
2,6-Dinitrotoluene	1ND	_13311
Diethylphthalate	1ND	1330
4-Chlorophenyl-phenylether	IND	133#
Fluorene	I NO	1 331
4,6-Dinitro-2-methylphenol	ND	1 830
N-Nitrosodiphenylamine	I NO	330
4-Bromophenyl-phenylether	INDI	1 330
Hexachlorobenzene	I ND	1330

Sample Identification: _NETHOD 8: AND # 271

	Concentration	I Untection Limit
Semivolatile Compounds	(ug/kg)	! (eg/kg)
	1	
Pentachlorophenol	1100	830
1 Phenanthrene	1 HO	330
Anthracene	110	330
Di-n-Butylphthalate	IND	1330
Fluoranthene	1ND	330
Pyrene	1110	1330
Benzidine	I	I83U
Butylbenzylphthalate	IND	_1330
3,3'-Dichlorobenzidine	1NO	1830 <u></u>
Benzo(a)Anthracene	1!ID	1
Bis(2-Ethylhexyl)Phthalate		133 <u></u>
Chrysene	1_ND	13.50
Di-n-octyl phthalate	1114D1	33.0
Benzo(b)fluoranthene	1ND	13?0
Benzo(k)Fluoranthene	1ND	_1330
Benzo(a)Pyrene	IND	
Indeno(1,2,3-od)Pyrene	1140	_13330
Dibenzo(a,h)Anthracene	ND	1330
Benzo(g,h,i)Perylene	IND	1
		1

SURRUGATE RECOVERY DATA

Surrogate Compound	QC Limits	Percent Macovery
Nitrohenzene-d5 2-Fluorobiphenyl		
Terphenyl-d14	(18-137)	93
Phenol-d5		
2,4,6-Tribromophenol		

ND: NOT DETECTED

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 1	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801271	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/1 or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8 0000
beta-BHC	ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND	8.000
Aldrin	ND ·	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND ND	16.00
Endrin	ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16.00
Endrin Aldehyde	ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND -	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16.0

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 2	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801272	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/l or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND	8.000
Aldrin	ND	8 000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND	16.00
Endrin	ND	. 16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16.00
Endrin Aldehyde	ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND .	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16.0

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 3	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801273	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/l or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND	8.000
delta BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND	8.000
Aldrin	ND	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND	16.00
Endrin	ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16.00
Endrin Aldehyde	ND ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16.0

ND - NOT DETECTED

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 4	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801274	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/l or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND	8.000
Aldrin	ND	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8 000
P,P'-DDE	ND	16.00
Dieldrin	ND	16 00
Endrin	ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16 00
Endrin Aldehyde	ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16.0

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 5	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801275	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/l or ug/kg): _ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND	8.000
Aldrin	ND	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND	16.00
Endrin	ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND ND	16.00
Endrin Aldehyde	ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Ch1ordane Ch1ordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16.0

ND - NOT DETECTED

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 6	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801276	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/l or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND ND	80.00
Heptachlor	ND	8.000
Aldrin	ND	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND ND	16.00
Endrin	_ ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16.00
Endrin Aldehyde	ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND ND	800.0
Aroclor 1221	_ ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND	80 0
Aroclor 1254	ND .	160.0
Aroclor 1260	ND	160.0
Endrin Kętone	ND	16.0

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION Sample 7	MATRIX: (soil/water) Soil
LABORATORY NUMBER 8801277	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED 8/19/88	UNIT (ug/l or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND	8.000
Aldrin	ND	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND	16.00
Endrin	ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16.00
Endrin Aldehyde	ND	16 00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND .	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16 0

PESTICIDE ORGANICS ANALYSIS DATA SHEET FOR SOIL

SAMPLE IDENTIFICATION N/A	MATRIX: (soil/water) Soil
LABORATORY NUMBER Lab Blank	SAMPLE WT/VOLUME: 30 g
DATE COLLECTED/RECEIVED N/A	UNIT (ug/l or ug/kg): ug/kg
DATE OF ANALYSIS 8/20/88	

		/
PESTICIDE COMPOUND	CONCENTRATION	DETECTION LIMIT
alpha-BHC	ND	8.0000
beta-BHC	ND	8.000
delta-BHC	ND	8.000
gamma-BHC	ND	80.00
Heptachlor	ND ND	8.000
Aldrin	ND	8.000
Heptachlor epoxide	ND	8.000
Endosulfan I	ND	8.000
P,P'-DDE	ND	16.00
Dieldrin	ND	16.00
Endrin	ND	16.00
P,P'-DDD & Endo. II	ND	16.00
P,P'-DDT	ND	16.00
Endrin Aldehyde	ND	16.00
Endosulfan Sulfate	ND	16.00
Methoxychlor	ND	80.0
Chlordane	ND	80.0
Toxaphene	ND	160.0
Aroclor 1061	ND	800.0
Aroclor 1221	ND	800.0
Aroclor 1232	ND	800.0
Aroclor 1242	ND	800.0
Aroclor 1248	ND .	80.0
Aroclor 1254	ND	160.0
Aroclor 1260	ND	160.0
Endrin Ketone	ND	16.0

Inorganic Data Comment Page

Lab Name RECRA ENVIRONMENTAL, INC.
ICP interelement and background corrections applied? Yes X No
If yes, corrections applied before or after generation of raw data
Footnotes:
NR - Not required by contract at this time
Form I:
Value - If the result is a value greater than or equal to the instrument detection limit but less than the contract required detection limit, report the value in brackets (i.e., [10]). Indicate the analytical method used with P (for ICP), A (for Flame AA) or F (for Furnace AA).
U - Indicates element was analyzed for but not detected. Report with the detection limit value (e.g., 10U).
E - Indicates a value estimated or not reported due to the presence of interference. Explanatory note included on cover page.
S - Indicates value determined by Method of Standard Addition.
N - Indicates spike sample recovery is not within control limits.
* - Indicates duplicate analysis is not within control limits.
+ - Indicates the correlation coefficient for method of standard addition is less than 0.995.
M - Indicates duplicate injection results exceeded control limits.
Indicate method used: P for ICP; A for Flame AA and F for Furnace.
COMMENTS:

EPA Sample No.	1
SAMPLE 1	1

LAB NAME _	RECRA	ENVIRON	ENTAL,]	INC.		CASE NO.		88-	1287	
CONTRACT N	UMBER					LAB RECEI	PT DAT	E{	3/17/	88
LAB SAMPLE	ID.	NO4	023,4024	<u> </u>		QC REPORT	' NO	88-3	1287	<u>¢c</u>
		<u> </u>	lements	Ident:	ified a	nd Measured	<u>l</u>			
Concentrat	ion:	Low				Medium _				
Matrix:	Wate	r	Soil	<u> </u>	s	ludge		Othe	er	
		/-	/2	,						
		ug/I	or mg/l	d gry	weight	(Circle Or	æ)			
1. Alumi 2. Antim		38,100 0.79 U		<u> </u>	13. 14.				N	P .
3. Arsen	ic	7.6		7		Mercury	0.24	U		CV
4. Bariu		106		4	16.		12	*		P
5. Beryl		1.5				Potassium				<u>P</u>
6. <u>Cadmi</u>		0.94 U		3		Selenium	0.79		N	F
7. Calci		3,940 21	<u>1</u>		19.	Silver	1.6 580	U	N	P
9. Cobal		4.8	<u> </u>	<u> </u>	21.	Sodium Thallium	0.79	U	N	A F
10. Coppe		16 *		<u>-</u>		Vanadium			TA	P
11. Iron		12,100		1 —		Zinc			N	A
l2. Lead		90		Ì		ent Solids		64.0		
Cyanide		NR					7.7			
	For as d	reporting lefined or	Cover I	Page. ed. De	Additi efiniti	ndard resul onal flags on of such wer.	or foo	tnote	es ex	plaining
Comments:	CV	- cold va	por							

EPA	Sample	No.	
SA	MPLE 2		-

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO. 88-1287
CONTRACT NUMBER	LAB RECEIPT DATE 8/17/88
LAB SAMPLE ID. NO. 4026	QC REPORT NO. 88-1287 QC
Elements Identified a Concentration: Low Matrix: Water Soil _X S	Medium
ug/L or mg/kg dry weight	(Circle One)
2. Antimony 0.57 U N F 14. 3. Arsenic 8.4 N F 15. 4. Barium 185 A 16. 5. Beryllium 3.9 P 17. 6. Cadmium 0.68 U A 18. 7. Calcium 149,000 P 19. 8. Chromium 25 A 20. 9. Cobalt 6.2 P 21. 10. Copper 36 * A 22. 11. Iron 39,900 A 23. 12. Lead 931 A Perce Cyanide NR	onal flags or footnotes explaining on of such flags must be explicit
Comments: CV - cold vapor	· · · · · · · · · · · · · · · · · · ·

EPA Sample N	w.
SAMPLE 3	

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO.	88-1287
CONTRACT NUMBER	LAB RECEIPT DAT	E <u>8/17/88</u>
LAB SAMPLE ID. NO. 4027	QC REPORT NO	88-1287gc
Dlamanta Idantified on	ad Managers A	
Elements Identified ar		
Concentration: Low	Medium	,
Matrix: Water SoilX SI	ludge	Other
2. Antimony 0.53 U N F 14. N 3. Arsenic 4.9 N F 15. N 4. Barium 97 A 16. N 5. Beryllium 2.1 P 17. N 6. Cadmium 2.0 A 18. S 7. Calcium 93,500 P 19. S 8. Chromium 18. D	(Circle One) Magnesium 16,800 Manganese 2,590 Mercury 0.16 Nickel 9.5 Potassium 1,220 Selenium 0,53 Silver 1.1 Sodium 1,170	N A U CV * P P U N F U N P
9. Cobalt 3.7 P 21.	Thallium 0.53	UNF
10. Copper 42 * A 22. T	Vanadium 13	P
11. Iron 29,600 A 23. 2	Zinc 88	N A
12. Lead 299 A Percer Cyanide NR	nt Solids $(%)$ 9	23.4
Footnote: For reporting results to EPA, star as defined on Cover Page. Addition results are encouraged. Definition and contained on Cover Page, however	onal flags or foo on of such flags	tnotes explaining
Comments: CV - cold vapor		

EPA Sample	No.	1
SAMPLE 4		1

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO.	88-1287
CONTRACT NUMBER	LAB RECEIPT DAT	TE <u>8/17/88</u>
LAB SAMPLE ID. NO. 4028	QC REPORT NO.	88-1287QC
Elements Identified a	nd Measured	
Concentration: Low	Medium	
Matrix: Water Soil X S	ludge	Other
ug/L or mg/kg dry weight	(Circle One)	
	Magnesium 4,610	
	Manganese 2,330	
	Mercury 0.19	
4. <u>Barium 77 A</u> 16.	Nickel 17	* P
	Potassium 1,440	
	Selenium 0.62	U N F
7. Calcium 37,500 P 19.	Silver 1.3	UNP
	Sodium 1,380	A
9. Cobalt 7.0 P 21.	Thallium 0.74	N F
	Vanadium 23	P
11. Iron 42,300 A 23.		N A
	nt Solids (%)	
Cyanide NR		
Footnote: For reporting results to EPA, sta	ndard result mia	lifiers are used
as defined on Cover Page. Additi	onal flags or for	otrotos orplainin
as defined on cover rage. Additi	on of such floor	Junes explaining
results are encouraged. Definiti		must be explicit
and contained on Cover Page, howe	ver.	
Comments of the sold		
Comments: CV - cold vapor		<u> </u>

EPA Sample	No.	
SAMPLE 5	1	

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO.	88-1287
CONTRACT NUMBER	LAB RECEIPT DATE	8/17/88
LAB SAMPLE ID. NO. 4029	QC REPORT NO	88-1287QC
Elements Identified ar	nd Measured	
Concentration: Low	Medium	and delication recovers
Matrix: Water Soil X SI	.udge	Other
ug/L or mg/kg dry weight	(Circle One)	
	Magnesium 3,620	<u>p</u> .
	Manganese 1,340	N A
	Mercury 0.17 Nickel 13	U CV
	Potassium 1,020	
	Selenium 0.59	U N F
7. Calcium 34,100 P 19.	Silver 1.2	U N P
	Sodium 1,390	A
	Challium 0.59	
	Manadium 15	P
	Zinc 133	N A
	nt Solids (%) 85	
Cyanide NR		
Footnote: For reporting results to EPA, star as defined on Cover Page. Addition results are encouraged. Definition and contained on Cover Page, however	onal flags or foot on of such flags m	notes explaining
Comments: CV - cold vapor		

EPA Sample No.	1
SAMPLE 6	 -

INORGANIC ANALYSIS DATA SHEET

THOROTAL TERMINIO	AIIII OIIIIII
AB NAME RECRA ENVIRONMENTAL, INC.	CASE NO. 88-1287
CONTRACT NUMBER	LAB RECEIPT DATE 8/17/88
AB SAMPLE ID. NO. 4030	QC REPORT NO. 88-1287QC
Elements Identified	and Measured
Concentration: Low	Medium
Matrix: Water Soil X	Sludge Other
2. Antimony 0.55 U N F 14. 3. Arsenic 7.7 N F 15. 4. Barium 105 A 16. 5. Beryllium 0.76 P 17. 6. Cadmium 0.66 U A 18. 7. Calcium 7,820 P 19. 8. Chromium 38 A 20. 9. Cobalt 7.4 P 21. 10. Copper 51 * A 22. 11. Iron 52,100 A 23. 12. Lead 761 A Perc Cyanide NR	Magnesium 2,120 P Manganese 1,610 N A Mercury 0.17 U CV Nickel 19 * P Potassium 1,480 P Selenium 0.55 U N F Silver 1.1 U N P Sodium 1,220 A A Thallium 0.55 U N F Vanadium 21 P P Zinc 361 N A ent Solids (%) 92.2 92.2 andard result qualifiers are used ional flags or footnotes explaining ion of such flags must be explicit
Comments: <u>CV - cold vapor</u>	

1	EPA Sample No.	1
1	SAMPLE 7	1

INORGANIC ANALYSIS DATA SHEET

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO. 88-1287
CONTRACT NUMBER	LAB RECEIPT DATE 8/17/88
LAB SAMPLE ID. NO4031	QC REPORT NO88-1287QC
Flowerts Identified an	nd Mongraphed
Elements Identified an	
Concentration: Low	Medium
Matrix: Water Soil X Sl	udge Other
ug/L or mg/kg dry weight	(Circle One)
1. Aluminum 15,500 P 13. M	Magnesium 1,810 P
2. Antimony 0.68 U N F 14. M	Manganese 83 N A
	Mercury 0.20 U CV
	Nickel 11 * P
	Potassium 1,840 P
	Selenium 0.68 UNF
7. Calcium 3,810 P 19. S	
8. Chromium 16 A 20. S	Sodium 324 A
	Challium 0.68 UNF
	Manadium 23 P
11. <u>Iron</u> 7,570 A 23. Z	
	nt Solids (%) 73.5
Cyanide <u>NR</u>	
Footnote: For reporting results to EPA, stan	dard result qualifiers are used
as defined on Cover Page. Addition	onal flags or footnotes explaining
results are encouraged. Definition	on of such flags must be explicit
and contained on Cover Page, howev	er.
Comments: <u>CV</u> - cold vapor	•

Q. C. Report No. <u>88-1287QC</u>

INITIAL AND CONTINUING CALIBRATION VERIFICATION 3

LAB	NAME RECE	RA ENVIRONME	ENTAL, IN	<u> 1C.</u>			CASE N	ю	88-12	87
							SOW NO)		
CAT	Ε						UNITS	ug/L		
Com	cound	Initia	al Calib.	1	Con	tinuing (Calibra	ation ²	,	
Meta	als:	True Value	Found	<u>%R</u>	True Value	Found	<u>%R</u>	Found	<u>%R</u>	Method ⁴
1.	Aluminum	1,000	1,040	104	1,000	1,020	102		 	ll P
2.	Antimony	100	103	103	100	100	100			F
	Arsenic	80	78	98	80	75	94		1	!! F
	Barium		9,880	99	10,000	10,050	101		1	H A
	Beryllium	500	492	98	500	477	95			P
	Cadmium	500	518	104	500	514	103			H A
	Calcium	20,000	21,200	106	20,000	20,600	103	<u> </u>		P
	Chromium	500	479	96	500	487	97	<u> </u>	1	A
	Cobalt	500	504	101	500	468	94	<u> </u>		ii P
	Copper	500	498	99	500	510	102			· A
	Iron	800	830	104	800	760	95		1	A
	Lead	2,500	2,500	100	2,500	2,590	104	<u> </u>	1	H A
	Magnesium		5,400	108	5,000	5,200	104			P
	Manganese	500	508	102	500	512	103		1	ii A
	Mercury	8.0	7.7	96	8.0	7.6	95		1	CV
16.	Nickel	300	330	107	300	300	100			A
	Potassium	5,000	5,200	104	5,000	4,900	98			A
18.	Selenium	50	50	100	50	46	92			F
	Silver	300	271	90	300	268	89		1	P
	Sodium	20,000	21,300	107	20,000	20,200	101			II A
	Thallium	80	82	103	80	84	105			F
	Vanadium	1,000	1,000	100	1,000	950	95			P
	Zinc	500	500	100	500	502	101		1	A
	er:		1						1	
									1	
Cya	nide	_	<u> </u>	-	_	T -				
3 _C	ontrol Lim		y and Ti	n 80-1	² Cc 20; Ot - ICP; A -	her Meta	ls 90-		nide 8	
1	imicate Wi	aryurar Me	man nac	ع ت	ICE; A -	TTOTIE IN		r dringe	* W	

^{*}US EPA or commercially purchased standard

Q.C. Report No. <u>88-1287QC</u>

BLANKS

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO	88-1287
DATE	UNITS	UG/L

Compound	1	Continuing (Blank V 2	Calibration Value 3	4	Preparation Blank Matrix: Matrix: SOIL 1 2	
Metals:		 			 	
1. Aluminum 2. Antimony 3. Arsenic 4. Barium 5. Beryllium 6. Cadmium 7. Calcium 8. Chromium 9. Cobalt 10. Copper 11. Iron 12. Lead 13. Magnesium 14. Manganese 15. Mercury 16. Nickel 17. Potassium 18. Selenium 19. Silver 20. Sodium 21. Thallium 22. Vanadium 23. Zinc Other:	40U 5U 5U 20U 7U 6U 500U 10U 30U 6U 30U 160U 100U 6U 0.2U 20U 500U 5U 10U 500U 5U 30U	40U 5U 5U 20U 7U 6U 500U 10U 30U 6U 100U 6U 0.2U 20U 500U 5U 10U 30U 5U 5U	20U 7U 6U 500U 10U 30U 6U 100U 6U 20U 500U 10U 500U 5U 30U			40U 5U 5U 20U 7U 6U 500U 10U 30U 6U 30U 160U 100U 6U 0.2U 20U 500U 5U 10U 500U 5U 30U
Cyanide						

 $^{^{1}}$ Reporting Units: aqueous, ug/L; solid, mg/kg

Q. C. Report No. 88-1287QC SPIKE SAMPLE RECOVERY

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO88-1287
	EPA Sample No. SAMPLE 1
DATE	Lab Sample ID No. 4025
	Units UG
Matrix	SOIL

		Control Limit	Chilend Complet	C1-		
			Spiked Sample	<u>Sample</u>	<u>Spiked</u>	
Com	oound	%R	Result (SSR)	Result (SR)	Added (SA)	§R ¹
Meta	als:					
1.	Aluminum	75-125	-	_	_	NR
2.	Antimony	**	5U	5U	50	0 N
3.	Arsenic	11 '	45	49	20	0 N
4.	Barium	11	2,428	676	2,000	88
5.	Beryllium	11	51	9.5	50	83
6.	Cadmium	!!	53	6U	50	105
7.	Calcium	11	_	-	_	NR
8.	Chromium	!!	324	133	200	96
9.	Cobalt	, 11	467	30	500	87
10.	Copper	"	308	99	250	84
11.	Iron	"	_	-	-	NR
12.	Lead	*1	990	573	500	83
13.	Magnesium	"	<u> </u>	-	-	NR
14.	Manganese	"	487	324	200	68 N
15.	Mercury	11	0.392	0.2U	0.4	98
16.	Nickel	11	430	75	400	89
17.	Potassium	"			_	NR
18.	Selenium	"	5U	5U	10	0 N
19.	Silver	**	10U	10U	50	0 N
20.	Sodium	11			-	NR
21.	Thallium	"1	29	5U	50	58 N
22.	Vanadium	11	601	194	500	81
	Zinc	"	768	689	200	40 N
Othe	er:					j
						1
Cyar	uide]	11				

1	₹R	=	[(SSR	_	SR)	/SA]	X	100
---	----	---	-------	---	-----	------	---	-----

"N" - Out of control

"NR" - Not required

Comments:	

Q. C. Report No. 88-1287QC

DUPLICATES

LAB NAME RECRA ENVIRONMENTAL, INC.	CASE NO88-1287
	EPA Sample No. SAMPLE 1
DATE	Lab Sample ID No. 4023,4024
	UnitsMG/KG DRY
Matrix	SOIL

]
Compour	nd	Control Limit ¹	Sample(S)	Duplicate(D)	RPD ²
Metals:	ļ				
1. Al 2. Ar 3. Ar 4. Ba 5. Be 6. Ca 7. Ca 8. Ch 9. Cc 10. Cc 11. Ir 12. Le 13. Ma 14. Ma 15. Me 16. Ni 17. Po 18. Se 19. Si	eminum entimony esenic eryllium edmium elcium encomium elcium encomium elcium encomium elcium encomium elcium encomium elcium el		33,500 0.79U 7.48 109 1.56 0.94U 3,990 20.7 4.52 19.2 12,000 88.7 3,070 53.6 0.24U 14 3,020 0.79U 1.6U	30,900 0.79U 7.78 102 1.40 0.94U 3,890 20.7 4.98 11.8 12,100 90.3 3,080 47.7 0.24U 9.3 2,960 0.79U 1.6U	6.9 NC 3.9 6.6 11 NC 2.4 0 9.7 46 * 0.82 1.8 0.32 12 NC 39 * 3.7 NC
21. Th	xdium mallium		582 0.79U	576 0.79U	0.94 NC
	nadium nc		28.8 112	31.8 103	9.9
Other:					
Cyanide	-				

^{*} Out of Control

NC - Non calculable RPD due to value(s) less than CRDL

¹ To be added at a later date

 $^{^{2}}$ RPD = [|S - D|/((S + D)/2)] x 100

Q.C. Report No. <u>88-1287QC</u>

INSTRUMENT DETECTION LIMITS AND

LABORATORY CONTROL SAMPLE

LAB NAME	RECRA ENVIRONMENTAL, INC.	CASE NO. <u>88-1287</u>	DATE
			LCS NO. MDS-976

<u>Compound</u>		Required Detection Limits (CRDL)-ug/L	Instrument Detection Limits (IDL)-ug/L ICP/AA Furnace ID#PLASMA40 ID# PE 500 PE 5000		Lab Control Samule ug/L mg/kg (circle one) True Found %R		
META	ALS:						
. 1.	Aluminum	200	40	!	!	4,610	*
	Antimony	60		5		-	
	Arsenic	10		5	†	1.89	
4.		200	20		 	-	
5.	Beryllium	5	5			0.70	
6.	Cadmium	5	5			18.8	
7.	Calcium	5000	500		l I	_	
8.	Chromium	10	10			189	
9.	Cobalt	50	30		i i	_	
10.		25	6			1,020	1
11.	Iron	100	30			13,700	
12.	Lead	5	160	5		489	i
	Magnesium	5000	100			_	
	Manganese	15	6			213	1
	Mercury	0.2	0.2			6.5	
	Nickel	40	20			159	
17.	Potassium	5000	500			_	
		5		5			
19.		10	10	<u> </u>	<u> </u>	1.1	!
		5000	500		<u> </u>		
21.	Thallium	10		5		-	
	Vanadium	50	30	-	<u> </u>	9.6	
	Zinc	20	5			1,350	
Othe	er:i		 	 	 		
Cyrar	nide	10	<u> </u>	 	 	· ·	
Cyan	<u> </u>	10 i	NR	NR			i

NR - Not Required

* - See Attached

SRM CONFIDENCE LIMITS

ELEMENT	MDS-976
	S - 1
	(ug/g)
Al	2,010-7,110
As	0-88.9
Be	0-2.99
Cd	10.5-27.8
Cr	150-246
Cu	882-1280
Fe	11,200-21,700
Pb	372-680
Mn	182-223
Hg	0-36.1
Ni	164-225
٧	1.7-24.4
Zn	1,190-1,450
Ag	0-203
Ti	0-4860



QC Report No. <u>88-1287QC</u>

HOLDING TIMES

LAB NAME	RECRA ENVIRONMENTAL,	INC.		
DATE			CASE NO	88-1287

EPA		Date	Mercury	Mercury	CN Prep	CN
Sample No.	Matrix	Received	Prep Date	Holding Time ¹ (Days)	- Date	Holding Time ¹ (Days)
SAMPLE 1	SOIL	8/17/88	8/23/88	6	 - -	1
SAMPLE 2	SOIL	8/17/88	8/23/88	6	_	
SAMPLE 3	SOIL	8/17/88	8/23/88	6	_	-
SAMPLE 4	SOIL	8/17/88	8/23/88	6	_	
SAMPLE 5	SOIL	8/17/88	8/23/88	6	_	
SAMPLE 6	SOIL	8/17/88	8/23/88	6	_	
SAMPLE 7	SOIL	8/17/88	8/23/88	6		

					i 	
					 	!
				1	1	
					! !	
					! 	! !
				[[]	 	1
			! ! !	ŧ 	1 1 1 1	
·			! ! !	 	f f 1	1
					1	
	 				1	
			 		† ‡ t	

¹Holding time is defined as number of days between the date received and the sample preparation date.

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND



VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION SAMPLE 1	MATRIX: (soil/water)	Soil
LABORATORY NUMBER 8801271	SAMPLE WT/VOLUME:	_5g
DATE COLLECTED/RECEIVED 8-18-88	UNIT (ug/l or ug/kg):	ne/ke
DATE OF ANALYSIS 8-25-88	DILUTION FACTOR:	1

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	85	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane Trans-1,2-Dichloroethene	ND	5
Trans-1,2-Dichloroethene	סא	5
Chloroform	ND	5
1,2-Dichloroethane	ON	5
1,1,1-Trichloroethane	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1.2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	112	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	16	5
1,1,2,2-TetrachTorethane	ND	5
Toluene	ND	5
Chlorobenzene	ND	5
Ethylbenzene	ND	5

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIM	SOIT	PERCENT RECOVERY
Toluene-d8 Bromofluorobenzene 1,2-Dichloroethane-d4	88-110	81-117	132
	86-115	74-121	76
	76-114	70-121	123

ND - NOT DETECTED

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION SA	MPLE 2	MATRIX: (soil/water)	<u>Soil</u>
LABORATORY NUMBER 88012	272	SAMPLE WT/YOLUME:	5g
DATE COLLECTED/RECEIVED _	8-18-88	UNIT (ug/l or ug/kg):	ug/kg
DATE OF ANALYSIS	8 25-88	DILUTION FACTOR:	1

		DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	55	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	65	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3 Dichloropropene	ND	5
2-Chloroethylvinylether	D	10
Bromoform	עא	5
Tetrachloroethene	15	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	NO	5
Chlorobenzene Ethylbenzene	ND ND	5 5

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIMITS		PERCENT RECOVERY	
	Water	Soil		
Toluene-d8	88-110	81-117	123	
Bromofluorobenzene	86-115	74-121	86	
1,2-Dichloroethane-d4	76-114	70-121	123	

ND - NOT DETECTED

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Sample 3	MATRIX: (soil/water)	Soil
LABORATORY NUMBER 8801273	SAMPLE WT/VOLUME:	58
DATE COLLECTED/RECEIVED 8/18/88	UNIT (ug/l or ug/kg):	ug/kg
DATE OF ANALYSIS 8/26/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	l ND	10
Chloroethane	D	10
Methylene Chloride	45	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	מא	5
Chloroform	מא	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	100	5
Dibromochloromethane	ND ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3 Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	15	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	ND	5
Chlorobenzene	ND	5
Ethylbenzene	ND	5

SURROGATE COMPOUND	RROGATE COMPOUND QC LIMITS		PERCENT RECOVERY	
	Water	Soil		
Toluene-d8	88 110	81-117	76	
Bromofluorobenzene	86-115	74-121	54	
1,2-Dichloroethane-d4	76-114	70-121	73	

ND - NOT DETECTED *: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Sample 4	MATRIX: (soil/water)	Soil
LABORATORY NUMBER 8801274	SAMPLE WT/VOLUME:	5g
DATE COLLECTED/RECEIVED 8/18/88	UNIT (ug/l or ug/kg):	ng/kg
DATE OF ANALYSIS 8/25/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	170	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	90	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	ND_	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	ND	5
Chlorobenzene	ND ND	5
Ethylbenzene	ND	5

SURROGATE COMPOUND			PERCENT RECOVERY	
	Water	Soil		
Toluene-d8	88-110		101	
Bromofluorobenzene 1,2-Dichloroethane-d4	86-115 76-114	74-121 70-121	65 175	

ND - NOT DETECTED *: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Sample 5	MATRIX: (soil/water)	<u>Soil</u>
LABORATORY NUMBER 8801275	SAMPLE WT/YOLUME:	<u> 5</u> g
DATE COLLECTED/RECEIVED 8/18/88	UNIT (ug/1 or ug/kg):	ug/kg
DATE OF ANALYSIS 8/26/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	110	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	75	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	40	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	9	5
1,2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND ND	5
Trichloroethene	230	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	50	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	ND	5
Chlorobenzene	ND	5
Ethylbenzene	ND	5

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIMITS	PERCENT RECOVERY
	Water Soil	
Toluene-d8	88 110 81-117	110
Bromofluorobenzene	86-115 74-121	41
1.2-Dichloroethane-d4	76-114 70-121	67

ND - NOT DETECTED *: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Sample 6	MATRIX: (soil/water)	soil
LABORATORY NUMBER 8801276	SAMPLE WT/VOLUME:	5g
DATE COLLECTED/RECEIVED 8/18/88	UNIT (ug/l or ug/kg):	ig/kg
DATE OF ANALYSIS 8/26/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl_Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	85	5
Trichlorofluoromethane	NĎ	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	18	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	9	5
1,2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	130	5
Dibromochloromethane	ND	5
1,1,2 Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	NO	10
Bromoform	ND	5
Tetrachloroethene	23	5
1,1,2,2-TetrachTorethane	ON D	5
Toluene	ND	5
Chlorobenzene	ND	5
Ethylbenzene	NO	5

SURROGATE COMPOUND	QC LIMI		PERCENT RECOVERY	
	Water	Soil		
Toluene-d8	88-110	81-117	101	
Bromofluorobenzene	86-115	74-121	46	
1,2-Dichloroethane-d4	76-114	70-121	73	

ND - NOT DETECTED
*: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Sample 7	MATRIX: (soil/water)	soil
LABORATORY NUMBER 8801277	SAMPLE WT/VOLUME:	58
DATE COLLECTED/RECEIVED 8/18/88	UNIT (ug/l or ug/kg):	na/ka
DATE OF ANALYSIS 8/26/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane .	DND	10
Methylene Chloride	43	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	30	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	- ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	180	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	D	5
Tetrachloroethene	30	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	ND	5
Chlorobenzene	ND	5
Ethylbenzene	ND	5

SURROGATE COMPOUND	QC LIMITS		PERCENT RECOVERY	
	Water	Soil		
Toluene-d8	88-110	81-117	73	
Bromofluorobenzene	86-115	74-121	55	
1,2-Dichloroethane-d4	76-114	70-121	63	

ND - NOT DETECTED *: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION _	Sample 1 MS	MATRIX: (soil/water)	Soil
LABORATORY NUMBER 880	01271	SAMPLE WT/VOLUME:	5g
DATE COLLECTED/RECEIVED	8/18/88	UNIT (ug/l or ug/kg):	ug/kg
DATE OF ANALYSIS 8/26	5/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	60 .	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	51	5
1.1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1.1.1-Trichloroethane	25	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	180	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	50	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	16	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	52	5
Chlorobenzene	39	5
Ethylbenzene	ND	5

SURROGATE COMPOUND	QC LIMITS Water Soil	PERCENT RECOVERY
Toluene-d8 Bromofluorobenzene 1 2-Dichloroethane-d4	88 110 81-117 86-115 74-121 76-114 70-121	86 48 72

ND - NOT DETECTED *: PROBABLE CONTAMINATION

SEP 27 '88 10:10

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICAT	TION Sample 1 MSD	MATRIX: (soil/water)	Soil
LABORATORY NUMBER	8801271	SAMPLE WT/VOLUME:	5 <u>g</u>
DATE COLLECTED/RE	CEIVED 8/18/88	UNIT (ug/l or ug/kg):	ug/kg
DATE OF ANALYSIS	8/26/88	DILUTION FACTOR: 1	

YOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	65	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	19	5
1,1-Dichloroethane	ND	5
1,1-Dichloroethane Trans-1,2-Dichloroethene	ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1.1.1-Trichloroethane	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	ND	5
1,2-Dichloropropane Trans-1,3 Dichloropropene	ND	5
Trichloroethene	110	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	38	5
cis-1,3 Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	ND	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	57	5
Chlorobenzene	38	5
Ethylbenzene	ND	5

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIMITS		PERCENT RECOVERY
	Water	Soil	
Toluene-d8 Bromofluorobenzene 1,2-Dichloroethane-d4	88-110 86-115 76-114	81-117 74-121 70-121	93 21 81

ND - NOT DETECTED *: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Method Blank	MATRIX: (soil/water)	Soil
LABORATORY NUMBER Method Blank	SAMPLE WT/VOLUME:	5g
DATE COLLECTED/RECEIVED N/A	UNIT (ug/l or ug/kg):	ug/kg
DATE OF ANALYSIS 8/25/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	24	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1,1-Dichloroethane Trans-1,2-Dichloroethene	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	מא	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane Trans-1,3-Dichloropropene	ND	5
Trans-1,3-Dichloropropene	ND	5
Trichloroethene	ND	5
Dibromochloromethane	ΝĎ	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	, ND	5
Tetrachloroethene	ND	5
1,1,2,2-Tetrachlorethane	D	5
Toluene	NO	5
Chlorobenzene	D	5
Ethylbenzene	ND	5

SURROGATE COMPOUND	QC LIMITS		PERCENT RECOVERY	
	Water	Soil		
Toluene-d8 Bromofluorobenzene 1,2-Dichloroethane-d4	86-115	81-117 74-121 70-121	92 96 95	

ND - NOT DETECTED *: PROBABLE CONTAMINATION

RECRA ENVIRONMENTAL, INC. COLUMBIA, MARYLAND

VOLATILE ORGANICS ANALYSIS DATA SHEET

SAMPLE IDENTIFICATION Method Blank	MATRIX: (soil/water)	Sc11
LABORATORY NUMBER Method Blank	SAMPLE WT/VOLUME:	5g
DATE COLLECTED/RECEIVED N/A	UNIT (ug/l or ug/kg):	ng/kg
DATE OF ANALYSIS 8/26/88	DILUTION FACTOR: 1	

VOLATILE COMPOUND	CONCENTRATION	DETECTION LIMIT
Chloromethane	ND	10
Bromomethane	ND	10
Vinyl Chloride	ND	10
Chloroethane	ND	10
Methylene Chloride	23	5
Trichlorofluoromethane	ND	5
1,1-Dichloroethene	ND	5
1.1-Dichloroethane	ND	5
Trans-1,2-Dichloroethene	ND	5
Chloroform	ND	5
1,2-Dichloroethane	ND	5
1,1,1-Trichloroethane Carbon Tetrachloride	ND	5
Carbon Tetrachloride	ND	5
Bromodichloromethane	ND	5
1,2-Dichloropropane	ND	5
Trans-1.3-Dichloropropene	ND	5
Trichloroethene	ND	5
Dibromochloromethane	ND	5
1,1,2-Trichloroethane	ND	5
Benzene	ND	5
cis-1,3-Dichloropropene	ND	5
2-Chloroethylvinylether	ND	10
Bromoform	ND	5
Tetrachloroethene	ND	5
1,1,2,2-Tetrachlorethane	ND	5
Toluene	ND	5
Chlorobenzene	ЙЙ	5
Ethylbenzene	ND	5

SURROGATE RECOVERY DATA

SURROGATE COMPOUND	QC LIMITS		PERCENT RECOVERY	
	Water	Soil		
Toluene-d8 Bromofluorobenzene 1,2-Dichloroethane-d4	88 110 86-115 76-114	81-117 74-121 70-121	87 104 100	

ND - NOT DETECTED *: PROBABLE CONTAMINATION

MEMO

To:

Bruce Clegg

Reference No. 2366

From:

Tony Misercola

Date:

3/30/89

Re:

Analytical Data - Ernst Steel/Pyramid Company

Revised 10-19-89 Memo

This memo details an assessment of analytical results reported by Recra Environmental for seven soil samples collected at the Ernst Steel/Pyramid Site on August 17, 1988. The samples submitted for analysis consisted of the following:

<u>Matrix</u>	Investigative Samples	Field Duplicates	<u>Total</u>
Soil/Fill	6	1	7

All samples were submitted for Target Compound List (TCL) VOAs, BNAs, Pesticides/PCBs and Target Analyte List (TAL) metals. Methods employed for the following parameters were taken from USEPA SW-846 "Test Methods for Evaluating Solid Waste", third edition, September 1986.

<u>Parameter</u>	<u>Method</u>
VOA	8240
BNA	8270
Pest/PCBs	8080
Metals	6010/7000 Series

The QA/QC by which these data have been assessed are outlined in the aforementioned methods and the documents entitled "Laboratory Functional Guidelines for Evaluating Organic and Inorganic Analyses", written by the USEPA.

Based on review of this data set and other related quality control data, the following are noted:

1. <u>Sample Holding Times</u>

Based on criteria in USEPA SW-846, the following holding time requirements have been established:

VOA (Solids) 14 days as long as stored at 4°C

BNA (Solids) 14 days prior to extraction, as long as stored at 4°C

40 days from extraction to analysis

Metals (Solids) 6 months prior to analyses

Mercury (Solids) 28 days prior to analyses

By comparing the sampling dates in the project field book and subsequent Chain of Custody forms to the actual dates of extraction and/or analysis, it is noted that all samples were extracted and/or analyzed prior to the expiration of their prescribed holding times.

2. <u>Surrogate Spike Recoveries</u>

Laboratory performance on individual samples is established by means of spiking activities. All samples submitted for the TCL VOAs, BNAs and Pest/PCBs were spiked with surrogates prior to sample preparation.

All samples submitted for TCL, BNAs and Pest/PCBs yielded surrogate recoveries within the control limits established in USEPA SW-846. Therefore, no qualification of these data is required on this basis.

However, problems did occur in surrogate recoveries for VOAs. Table 1 shows that seven of the 10 samples analyzed (1 MS/MSD and 1 method blank) obtained surrogate recoveries that were outside recommended control limits.

In my memo to Dr. Mohan Khare of Recra Environmental, all poor surrogate recoveries were duly noted. His response was that all calculations were correct as well, all instrumental performance criteria were met. The samples should be re-analyzed to establish whether matrix interference or laboratory error was responsible for the poor surrogate performance.

Upon receiving results of the reanalysis on September 22, 1988, it was noted that surrogates for the VOAs were again outside recommended control limits except for the method blanks. Therefore, based on the results of the reanalysis, it is best assumed that the apparent complexity of the sample matrix not only caused poor recoveries of the surrogates but also caused several detectable compound concentrations which were not evident in the

original analyses. Therefore, the following qualifications shall be made for the results of the VOAs in all seven samples:

- a) All positive VOA results are flagged as estimates (J) due to out-of-control surrogate spike recoveries.
- b) Sample quantitation limits are flagged as estimated (UJ) due to out-of-control surrogate spike recoveries.

3. <u>Laboratory Reagent/Blank Analyses</u>

The assessment of results on blank analysis is for the purpose of determining the existence and magnitude of contamination problems. Solid laboratory blank samples were prepared from 30g of sodium sulfate while liquid blanks were prepared by deionized water.

All laboratory reagent blanks submitted for TCL, BNAs, Pest/PCBs and TAL metals yielded non-detectable concentrations of the analytes of interest. It should be noted that methylene chloride was detected in the VOA blanks at the method quantitation limit (5 ppb) and in the reanalysis at 23 ppb. This analyte, however, is a laboratory artifact and should have no effect on sample data.

Matrix Spike/Matrix Spike Duplicate (MS/MSD) Analyses

In general, no action is taken on MS/MSD data to qualify an entire case. This data alone cannot be used to evaluate the precision and accuracy of individual samples.

The precision of an analytical method is demonstrated by the reproducibility of an analytical data. Presented in Table 2A are the relative percent differences between duplicate matrix spike analyses.

All RPD results for VOAs, BNAs, Pest/PCBs were within their control limits established by USEPA SW-846 demonstrating adequate precision among the analyses.

All spike recoveries for MS/MSD analyses for VOAs were within their respective control limits indicating satisfactory accuracy, of the analyses performed. Table 2A shows no more than two spike recoveries in the BNAs and one spike recovery for Pest/PCBs to be outside recommended control limits. This situation is still acceptable in terms of quality control, however, since results for these spike recoveries were lower than method control limit criteria, a slight low bias in 1,2,4-trichlorobenzene and g-bhc results for the soils analyzed may be implied.

5. <u>Matrix Spike and Duplicate Results for TAL Metals</u>

As previously noted, reasoning behind spiking and duplicating sample analyses is to generate accuracy and precision information of a particular matrix. Presented in Table 2B are the results of spike recoveries for the specific metals analyzed for. Problems existed for spike recoveries of seven metals, those being: antimony; arsenic; manganese; selenium; silver; thallium and zinc. As a result of these low recoveries, the qualifications as shown on Table 3A must be made regarding the results of these metals in the seven soil samples.

Duplicate analysis was performed for metals on Sample No. 1 and are presented in Table 2C. Satisfactory precision was demonstrated with low RPD values for all metals except for copper and nickel which exhibited higher RPD values than the recommended control limit of 35± percent. Therefore, all associated values of copper and nickel shall be qualified as presented in Table 3B.

6. Field OA/OC

In accordance with proper QA/QC, one field duplicate was taken at location 19N22E. The two samples were submitted as No. 4 and No. 5 respectively to the laboratory. The results of the field duplicates yielded satisfactory results except two semi-volatiles detected slightly above the detection limit in Sample No.5, showed less than detectable quantities in Sample No. 4. The discrepancies in this data may be attributed to inhomogenity in the sample matrix. Organic analyses are highly dependent on the sample matrix. Slight changes in the matrix may have a substantial effect on the reproducibility of analytical data.

7. Summary

In summary, there were a number of problems that arose that made it necessary for some for the data to be qualified those being:

- a) Poor surrogate recoveries in the VOA analyses due to matrix interference. On this basis, all VOC results associated with the soil samples had to be qualified as estimated (data qualifer).
- b) Due to poor recoveries on sample spikes for metallic analytes antimony, arsenic, manganese, selenium, silver, thallium and zinc had to be qualified as either estimated values or unusable. The laboratory has been notified of these results in an attempt to find the reason that QC criteria was not met for these analytes.

c) On the basis of poor precision established for duplicate analyses of copper and nickel, all associated values for these two analytes had to be qualified as estimated.

References:

- 1. "Laboratory Data Validation Functional Guidelines for Evaluating Organics Analyses", Technical Directive Document, Prepared by the USEPA Data Validation Work Group, NUS Corporation SuperFund Division.
- 2. "Laboratory Data Validation Functional Guidelines for Evaluating Inorganics Analyses", prepared by the USEPA Office of Emergency and Remedial Response.

TM:jd Attachments

Table 1 Summary of Surrogates Outside Quality Control Limits on First Analysis

<u>I.D. No.</u>	Surrogate	Percent Recovery	Percent Control <u>Limit</u>
1	Toluene-d ₈	126	81-117
4	Toluene-d8	134	81-117
	BFB	73	74-121
5	Toluene-d8	128	81-117
6	Toluene-d8	119	81-117
7	Toluene-d8	124	81-117
1 (M/S)	Toluene-d8	123	81-117
Method Blank	Toluene-d ₈	112	88-110

Table 2A Relative Percent Differences Between Duplicate Matrix Spike Analysis Organic Parameters

Lab ID No.	<u>Parameter</u>	RPD ¹	RPD Control <u>Limit</u> ²	Spike Percent <u>Recovery</u>	Spike/Dup. Percent <u>Recovery</u>	Spik <u>Matı</u>
	GC/MS Volatile Compounds					Soil
	1,1-Dichloroethene	4	22	156	262	
	Trichloroethene	0	24	88	88	
	Benzene	0	21	126	126	
•	Toluene	3	21	136	132	
•	Chlorobenzene	0	21	100	100	
	Base/Neutral Acid Compounds	5				Soil
	Phenol	2	35	47	48	,
:	2-Chlorophenol	8	50	38	41	
	1,4-Dichlorobenzene	12	27	40	45	
	N-Nitroso-di-n-Propylamine	7	38	53	57	
	1,2,4-Trichlorobenzene	8	23	34*	37*	
	Chloro-3-Methylphenol	2	33	40	41	
	Acenapthene	2	19	46	47	
	4-Nitrophenol	8	50	13	12	
	2,4-Dinitrotoluene	3	47	· 33	32	
	Pentachlorophenol	7	47	44	41	
	Pyrene	13	36	58	66	
	Pesticides/PCBs					Soil
7	g-BHC	17	50	51	43*	
	Heptachlor	10	31	92	83	
	Aldrin	31	43	66	48	
•	Dieldrin	12	38	64	57	
	Endrin	13	45	88	77	
	4,4-DDT	17	50	62	53	

¹ Relative percent difference between spike duplicate percent recoveries.

² Control limits for RPD analyses detailed in USEPA SW-846 (September 1986).

^{*} Outside spike recovery control limits established in USEPA SW-846 (September 1986).

Table 2B Spike Recovery Results for Metal Analyses

I.D. No.	<u>Metal</u>	Control Limit	Percent Recovery
Sample 1	Antimony	75-125	0*
•	Arsenic	11	0*
	Barium	**	88
	Berylium	Ħ	83
	Cadmium	u ·	105
	Chromium	11	96
	Cobalt	H	87
	Copper	п	84
	Manganese	H	68*
	Mercury	н	98
	Nickel	n	89
	Selenium	11	0*
	Silver	II	0*
	Thallium	11	58*
	Vanadium	***	81
	Zinc	11	40*
	Lead	11	83

^{*}Outside control limits established in USEPA SW-846 (September 1986).

Table 2C RPD Values for Duplicate Metal Analyses

I.D. No.	<u>Metal</u>	RPD ¹	Control RPD <u>Limit²</u>	Initial <u>Result</u> (mg/Kg)	Duplicate <u>Result</u>
Sample 1	Aluminum Arsenic Barium Berylium Calcium Chromium Cobalt Copper Iron Lead	6.9 3.9 6.6 11 2.4 0 9.7 46* 0.82		33,500 7.48 109 1.56 3,990 20.7 4.52 19.2 12,000 88.7	30,960 7.78 102 1.40 3,890 20.7 4.98 11.8 12,100
	Magnesium Manganese Nickel Potassium Sodium Vanadium Zinc	1.8 0.32 12 39* 3.7 0.94 9.9 8.3		3,070 53.6 14 3,020 582 28.8 112	90.3 3,080 47.7 9.3 2,960 576 31.8 103

 $^{^1\}mathrm{Relative}$ Percent Difference between duplicate sample analyses.

²RPD Control Limit Assumed to be 35± Percent

Table 3A Metals Qualified as a Result of Poor Spike Recoveries

Sample No.	Analyte	Conc. (mg/Kg)	<u>Qualifier</u>
1	Antimony	<0.79	R
	Arsenic	7.6	Ţ
	Manganese	51	J
	Selenium	<0.79	R
	Silver	<1.6	R
	Thallium	<0.79	R
	Zinc	108	J
2	Antimony	< 0.57	R
	Arsenic	8.4	J
	Manganese	3010	J
	Selenium	< 0.57	R
	Silver	<1.2	R
	Thallium	<0.57	R
	Zinc	155	. J
3	Antimony	<0.53	R
	Arsenic	4.9	J
	Manganese	2590	Ĵ
	Selenium	< 0.53	Ř
	Silver	<1.1	R
	Thallium	< 0.53	R
	Zinc	88	J
4	Antimony	<0.62	R
•	Arsenic	12	Ĵ
	Manganese	2330	Ĵ
	Selenium	< 0.62	R
	Silver	<1.3	R
	Thallium	0.74	J
	Zinc	211	J

Table 3A Metals Qualified as a Result of Poor Spike Recoveries

Sample No.	Analyte	Conc. (mg/Kg)	<u>Qualifier</u>
5	Antimony Arsenic Manganese Selenium Silver Thallium Zinc	<0.59 8.9 1340 <0.59 <1.2 <0.59 133	R J R R R
6	Antimony	<0.55	R
	Arsenic	7.7	J
	Manganese	1610	J
	Selenium	<0.55	R
	Silver	<1.1	R
	Thallium	<0.55	R
	Zinc	361	J
7	Antimony	<0.68	R
	Arsenic	9.5	J
	Manganese	83	J
	Selenium	<0.68	R
	Silver	<1.4	R
	Thallium	<0.68	R
	Zinc	89	J

NOTES:

R - unusable

J - Values should be viewed as estimates

Table 3B Metals Qualified as a Result of Poor Precision

Sample No.	<u>Metal</u>	<u>Results</u> (mg/Kg)	Qualifier
1	Copper Nickel	16 12	J
2	Copper Nickel	36 13	J
3	Copper Nickel	42 9.5	J
4	Copper Nickel	12 17	J
5	Copper Nickel	35 13	J
6	Copper Nickel	51 19	. J J
7	Copper Nickel	15 11	J

NOTES:

J = the associated value is an estimated quantity because quality control criteria were not met.



651 Colby Drive, Waterloo, Ontario, Canada N2V 1C2 (519) 884-0510

August 24, 1988

Reference No. 2366

Dr. Kare
RECRA ENVIRONMENTAL INC.
8320 Guilford Road, Building F
Columbia, MD
21048

Dear Dr. Kare:

Re: Surrogate Recoveries for Volatiles

On Thursday, August 18, 1988, Conestoga-Rovers & Associates (CRA) delivered seven soil samples for the Target Compound List of VOAs, BNAs, PCBs/Pesticides. Upon initial review of results sent by telefax on Monday, August 22, 1988, some deficiencies have been identified with surrogate recoveries for five of the soil samples and the additional matrix spike sample. Following is a list of the surrogates outside control limits for the individual samples:

Number	Surrogate	Percent Recovery	Control <u>Limits</u>
1	Toluene d ₈	126	81-117
4	Toluene d ₈	134	81-117
_	BFB	73	74-121
5	Toluene d ₈	128	81-117
6	Toluene d ₈	119	81-117
7	Toluene d ₈	124	81-117
1 (MS)	Toluene dg	123	81-117

The evaluation of the results is not necessarily straight forward, the sample itself may produce effects due to such factors as interference and high concentrations of analytes.

USEPA-SW-846 states that the following procedures be adhered to when recovery for surrogates is out of control:

- 1. Check to be sure there are no errors in calculations, check instrument performance.
- 2. Recalculate the data and/or re-analyze the extract if any of the above checks reveal a problem.

August 24, 1988 Dr. Kare Page. 2. #2366

3. If none of the above are a problem, the effected fraction should be repurged to establish whether non-conformance was due to the sample matrix.

If you have any questions pertaining to this letter, please do not hesitate to contact me at 716-283-6720.

Yours very truly,

CONESTOGA-ROVERS & ASSOCIATES

anthony Misercola

Anthony Misercola

Per Ja

Environmental Chemist

AM:jd

cc: B. Clegg



CONESTOGA-NOVERS & ASSOCIATES LIMITED

651 Colby Drive, Waterloo, Ontario, Canada N2V 1C2 (519) 884-0510

September 22, 1988

Reference No. 2366

Mr. Jim Miller Recra Environmental Inc. 10 Hazelwood Drive Suite 106 Amherst, NY 14150

Dear Mr. Miller:

Re: Deficiencies in QC Results - Ernst Steel/Pyramid Co.

In reviewing the analytical results for TCL metals on the seven soil samples submitted on August 17, 1988 for the above mentioned, some deficiencies were noted. We would like your comments on problems associated with these deficiencies and your corrective actions.

1. Duplicate Analysis

Duplicate results for copper and zinc on soil sample #1 fell outside the recommended RPD control limits of 35 percent. Because of this, all associated results for copper and zinc in the remaining soil samples had to be qualified as estimates.

2. Matrix Spike Data

Data obtained on spike recoveries for the following metals were outside the quality control limits established in USEPA SW-846:

Metal	% Recovery
Antimony	0
Arsenic	0
Manganese	68
Selenium	0
Silver	0
Thallium	5 8
Zinc	40

Why was there such poor recovery for these metals, particularly since the sample itself contained very low concentrations of the above mentioned metals? Was the sample re-extracted or reanalyzed?

All results for the metals that had 0 percent recovery are unusable for quantitative purposes.

3. Form 7, P14, "Laboratory Control Sample Sheet", does not include true values and percent recoveries for the TCL metals.

If you have any questions or need further clarification, please do not hesitate to contact me at 716-283-6720.

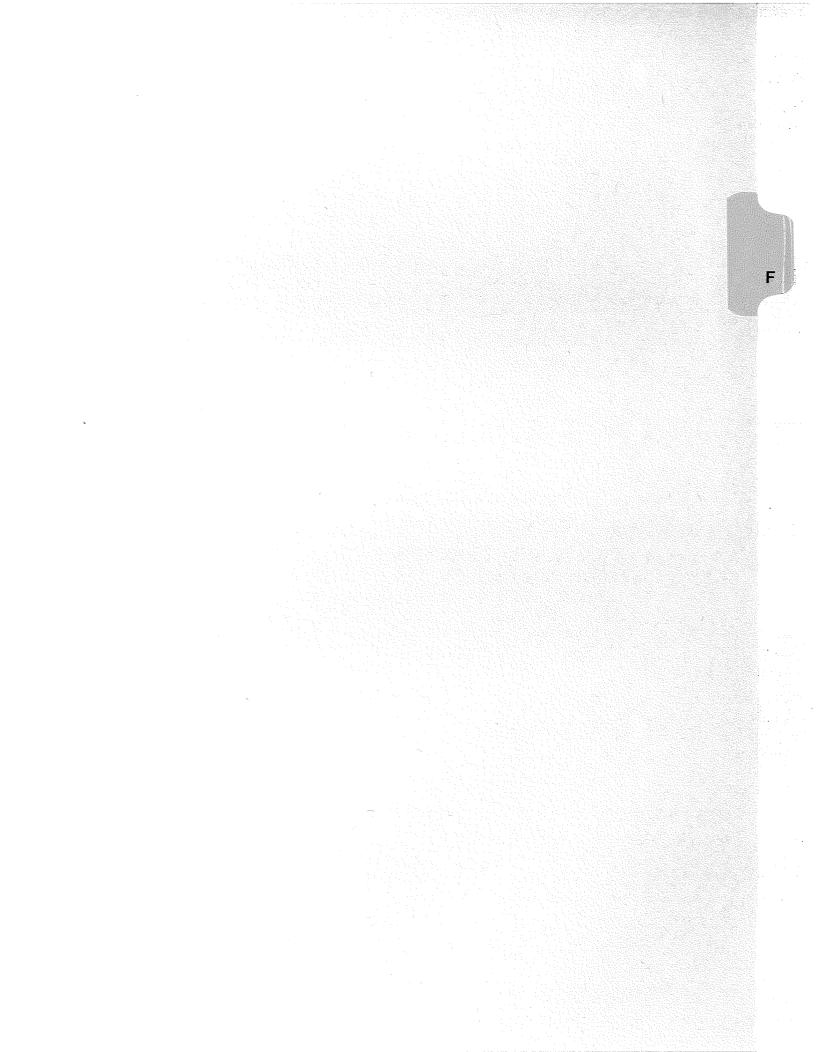
Yours very truly,

CONESTOGA-ROVERS & ASSOCIATES

Anthony Misercola Project Chemist

AM:jd

cc: B. Clegg



APPENDIX F

GROUNDWATER SAMPLING RESULTS



HUNTINGDON ANALYTICAL SERVICES
Division of EMPIRE SOILS INVESTIGATIONS INC.
PO Box 250 Middleport New York 14105
Telephone (716) 735-3400 Telex 131246

ENVIRONMENTAL ANALYTICAL REPORT

HAS REF. #30-616, #30-618, #30-622, and #30-629

Study Nos. E046.029 - E046.031 and E046.033

September 27, 1988

REPORT CODE LEGEND:

<DL = LESS THAN DETECTION LIMIT</pre>

ND = NOT DETECTED

NA = NOT APPLICABLE



HAS Reference Numbers: #30-616, #30-618, #30-622, and #30-629

September 27, 1988

Statement of Work Performed

I hereby declare that the work was performed under my supervision according to the procedures outlined by the following U.S. Environmental Protection Agency references and that this report provides a correct and faithful record of the results obtained.

- 40 CFR Part 136, "Guidelines Establishing Test Procedures for the Analysis of Pollutants Under the Clean Water Act," October 26, 1984 (Federal Register) U.S. Environmental Protection Agency.
- U.S. Environmental Protection Agency, "Test Methods of Evaluating Solid Waste Physical/Chemical Methods," Office of Solid Waste and Emergency Response, SW-846, 2nd Edition and 3rd Edition.

Richard D. Fitzpatrick

Environmental Laboratory Director

MODIFIED METHOD 8020

				Method		l	
Sample Identification:		OW3-88	OW103-881	Blank	SW-1	SW-2	SW-102
HAS Sample #30-		616-001	616-002		618-015	618-0161	017
Date Sampled:		9/12/88	9/12/88		9/13/88	9/13/88	9/13/88
Date Received:		9/13/88	9/13/881		9/13/88	9/13/88	9/13/88
Date Analyzed:	j	9/14/88	9/14/88	9/14/881	9/14/881	9/14/88	9/14/88
Holding Time, Days:		2	2 1		11	1 1	1
Dilution Factor:		1	1 1	1 1	1	1 1	1
İ	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/l	μg/l	μg/l	μg/l	μg/l	μg/l	μg/l
1	1		1	1			
_Toluene	i 0.50 i	<dl< td=""><td><dl td="" <=""><td><dl i<="" td=""><td><dl i<="" td=""><td><dl i<="" td=""><td><dl< td=""></dl<></td></dl></td></dl></td></dl></td></dl></td></dl<>	<dl td="" <=""><td><dl i<="" td=""><td><dl i<="" td=""><td><dl i<="" td=""><td><dl< td=""></dl<></td></dl></td></dl></td></dl></td></dl>	<dl i<="" td=""><td><dl i<="" td=""><td><dl i<="" td=""><td><dl< td=""></dl<></td></dl></td></dl></td></dl>	<dl i<="" td=""><td><dl i<="" td=""><td><dl< td=""></dl<></td></dl></td></dl>	<dl i<="" td=""><td><dl< td=""></dl<></td></dl>	<dl< td=""></dl<>

Quality Control

Surrogate	% Rec.					
İ						1
α,α,α-Trifluorotoluene	96	91	97	92	93	95

i		1	Method			Method	
Sample Identification:		SW-3	Blank	OW1-88	OW2-88	Blank	
HAS Sample #30-		618-018		622-0011	622-0021	1	
Date Sampled:		9/13/88		9/15/88	9/15/88	1	
Date Received:		9/13/88		9/15/881	9/15/88]	
Date Analyzed:		9/14/88	9/14/881	9/15/881	9/15/88	9/15/88	
Holding Time, Days:		1	1	0 1	0 [1	
Dilution Factor:	1	1 1	1	1	1 1	1 [
	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/l	μg/l	μg/l	μg/1	μg/l	μg/l	$\mu g/1$
	1 1	1	l	1	1	1	
Toluene	0.5	<dl td="" <=""><td><dl td="" <=""><td>0.53</td><td><dl< td=""><td><dl td="" <=""><td></td></dl></td></dl<></td></dl></td></dl>	<dl td="" <=""><td>0.53</td><td><dl< td=""><td><dl td="" <=""><td></td></dl></td></dl<></td></dl>	0.53	<dl< td=""><td><dl td="" <=""><td></td></dl></td></dl<>	<dl td="" <=""><td></td></dl>	

Quality Control

		Ì				
Surrogate	% Rec.	8 Rec.	% Rec.	% Rec.	8 Rec.	Rec.
		[]	1	l	<u> </u>
α, α, α -Trifluorotoluene	93	97	89	91	93	<u> </u>

METHOD 8080 POLYCHLORINATED BIPHENYLS

			1	Method		[
Sample Identification:		OW3-88	OW103-88	Blank		L	i
HAS Sample #30-616		001	002	• •		[
Date Sampled:		9/12/88	9/12/88]	
Date Received:		9/13/88	9/13/88			l	
Date Prepared:		9/13/88	9/13/88	9/13/88		l	
Date Analyzed:		9/14/88	9/14/88	9/14/88		ſ	
Dilution Factor:		11	1 1	1		1	
	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/l	μg/l	μg/1	μg/1	μg/l	μg/l	μg/l
			1			l	
AC-1254	1.0	ND	ND I	ND			
	1			1			
Hexabromobenzene (Surrog	ate) % Rec.	53	55	75			

1						Method	
Sample Identification:	SW-1	SW-2	SW-102	SW-3	Blank		
HAS Sample #30-618		015	016	017	018		
Date Sampled:		9/13/88	9/13/88	9/13/88	9/13/88		
Date Received:		9/13/88	9/13/88	9/13/88	9/13/88		
Date Prepared:		9/13/88	9/13/88	9/13/88	9/13/88	9/13/88	
Date Analyzed:	9/14/881	9/14/88	9/14/88	9/14/88	9/14/881		
Dilution Factor:		1 1	11	11	11	1 1	
	Detection		Result	Result	Result	Result	Result
COMPOUND	Limit µg/1	μg/l	μg/l	μg/l	μg/l	μg/l	μg/l
			_	1	_ ·		
AC-1254	1.0	ND	ND I	ND	ND I	ND 1	
			1		ĺ	1	I
Hexabromobenzene (Surrog	ate) % Rec.	77	<u>71</u>	69	95	124	

				Method			
Sample Identification:		OW1-88	OW2-88	Blank			<u> </u>
HAS Sample #30-622		001	002				<u> </u>
Date Sampled:		9/15/88	9/15/88				<u></u>
Date Received;		9/15/881	9/15/88				
Date Prepared:		9/15/881	9/15/88	9/15/88			
Date Analyzed:		9/15/881	9/15/88	9/15/881			<u> </u>
Dilution Factor:		1	1 1	1 1			<u></u>
	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/l	μg/l	μg/l	μg/1	μg/l	μg/l	. μg/l
			1				
AC-1254	1.0	ND I	ND 1	ND			
				I			
Hexabromobenzene (Surrog	ate) % Rec.	54	60	78 i			

METHOD 8080 POLYCHLORINATED BIPHENYLS

	İ				Method		
Sample Identification:		SS1-88	SS2-88	SS3-88	Blank		
HAS Sample #30-616		003 [004	005			
Date Sampled:		9/12/88	9/12/88	9/12/88			
Date Received:		9/13/881	9/13/88	9/13/881			
Date Prepared:		9/13/88	9/13/88	9/13/88	9/13/88		
Date Analyzed:		9/14/881	9/14/88	9/14/88	9/14/88		
Dilution Factor:	1	1	11	1 1	11		
	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/g	μg/g	μg/g	μg/g	μg/g	μg/g	μg/g
AC-1254	0.20	 DM	ND	I D I	ND		
Hexabromobenzene (Surrog	 ate) % Rec.	51 <u> </u>	*	125	124		

^{*}Surrogate omitted during preparation.

	[]		1	
Sample Identification:		SS-4	SS-5	SS-6	SS-7	SS-107	<u> SS-8</u>
HAS Sample #30-618		001	002	003	004	005 [006
Date Sampled:		9/13/88	9/13/88	9/13/88	9/13/88	9/13/88	9/13/88
Date Received:		9/13/88	9/13/88	9/13/88	9/13/88	9/13/88	9/13/88
Date Prepared:	1	9/13/88	9/13/88	9/13/88	9/13/88	9/13/88	9/13/88
Date Analyzed:	1	9/15/88	9/15/88	9/15/88	9/15/88	9/15/88	9/15/88
Dilution Factor:		1	1 1	1 1	11	1	1
	Detection		Result	Result	Result	Result	Result
COMPOUND	Limit μg/g	μg/g	μg/g	μg/g	ug/g	ug/g	ug/g
	1		1			1	
AC-1254	i 0.20 i	1.2 i	ND I	0.60 j	1.6 j	2.2	ND
	İ	1	1	I	1	ĺ	
Hexabromobenzene (Surrog	ate) % Rec.	100 j	106 j	73 i	67 j	87 j	<u>59</u>

				l			
Sample Identification:	i	SS-9	SS-10	Sed 1	Sed 2	Sed 3	<u>Sed 103</u>
HAS Sample #30-618 -	007 [008 1	009	010	011	012	
Date Sampled:	9/13/88	9/13/881	9/13/88	9/13/88	9/13/881	9/13/88	
Date Received:	1	9/13/88	9/13/88	9/13/88	9/13/88	9/13/88	9/13/88
Date Prepared:	1	9/13/88	9/13/881	9/13/881	9/13/88	9/13/88	9/13/88
Date Analyzed:	1	9/15/88	9/15/881	9/15/881	9/15/88	9/15/881	9/15/88
Dilution Factor:	İ	1 1	1	1 i	11	1	1
	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/g	μg/g	ug/g	ug/g	μg/g	μg/g	ug/g
	1						
AC-1254	i 0.20 i	ND i	ND I	ND I	ND	ND I	ND
						1	
Hexabromobenzene (Surrog	ate) % Rec.	57 i	75 i	50 i	54	48	76

METHOD 8080 POLYCHLORINATED BIPHENYLS

				Method		l	
Sample Identification:		Sed 4	Sed 5	Blank			
HAS Sample #30-618		013	014	/			
Date Sampled:	1	9/13/88	9/13/88			1	
Date Received:		9/13/881	9/13/881				
Date Prepared:		9/13/88	9/13/881	9/13/88			
Date Analyzed:	J	9/15/88	9/15/88	9/15/881			
Dilution Factor:		1 1	1 1	1 1			
	Detection	Result	Result	Result	Result	Result	Result
COMPOUND	Limit µg/g	μg/g	μg/g	μg/g	μg/g	μg/g	μg/g
AC-1254	0.20	ND I	ND	ND [
Hexabromobenzene (Surrog	ate) % Rec.	41	67 <u> </u>	78 i	:		

PREPARED FOR: CONESTOGA ROVERS

Metals Analysis for: TOTAL LEAD

SAMPLE ID		EPA	DATE	DATE	DETECTION	RESULT	l QC
CUST.	HAS #	METHOD	PREPARED	ANALYZED	LIMIT	ppm	%SPKR/RPI
	1676 001						
W3-88	616-001	•	9-13-88	9-15-88	0.002	<dl< td=""><td>*95</td></dl<>	* 95
W103-88	616-002	1	•	9-15-88	0.002	<dl< td=""><td>104/24.8</td></dl<>	104/24.8
SS1-88	616-003	•	•	9-15-88	2.27	219	*95
SS2-88	616-004	•	•	9-15-88	1.47	507	* 95
SS3-88	616-005	16010	9-13-88	9-15-88	1.99	181	170/6.1
SS-4	 618-001	 6010	 9-13-88	 9-14-88	 2.29	29600	¦ * 95
SS-5	1618-002	•	,	9-14-88	2.05	8980	* 95
SS-6	618-003	6010	,	9-14-88	1.6	5570	* 95
SS-7	618-004	6010		9-14-88	2.15	14200	* 95
SS-107	618-005	6010		9-14-88	2.28	14200	*95
SS-8	1618-006	•		9-14-88	2.19	563	* 95
SS-9	618-007	•	1	9-14-88	2.24	312	* 95
SS-10	618-008	6010		9-14-88	2.06	4370	*95
ED 1	618-009	•		9-14-88	2.14	286	*95
SED 2	618-010	•		9-14-88	1.57	.118	*95
SED 3	618-011			9-14-88	1.99	38.1	* 95
ED 103	618-012			9-14-88	1.51	31.5	* 95
ED 4	618-013	6010		9-14-88	2.04	255	* 95
ED 5	618-014			9-14-88	2.34	35	* 95
W-1	618-015			9-15-88	0.002	<dl td="" <=""><td>*95</td></dl>	* 95
W-2	618-016	7421		9-15-88	0.002	<dl td="" <=""><td>*95</td></dl>	* 95
W-102	618-017	7421		9-15-88	0.002	0.002	* 95
W-3	618-018	7421		9-15-88	0.002	<dl td="" <=""><td>*95</td></dl>	* 95
					1.002		
		1		İ	Ī		

*THIS INDICATES A 95% CONFIDENCE LIMIT ACHIEVED WITH AN EPA QUALITY CONTROL SOLUTION ANALYZED ALONG WITH YOUR SAMPLE.

PREPARED FOR: CONESTOGA ROVERS

Metals Analysis for: TOTAL LEAD

,	SAMPLE ID	HAS #	EPA METHOD	DATE PREPARED	DATE ANALYZED	DETECTION LIMIT	RESULT ppm	QC %SPKR/RPD	
									1
	I IOW1-88	 622-001	1 7421	 9-15-88	9-15-88	0.002	0.003	l *95	
	OW2-88	622-002	7421	9-15-88	9-15-88	0.002	<dl< td=""><td>108/1.8</td><td></td></dl<>	108/1.8	
	TRIP BLK	622-003	7421	9-15-88	9-15-88	0.002	0.024	* 95	
	ĺ					İ			:
	OW-3A	629-001	7421	9-21-88	9-22-88	0.002	<dl< td=""><td>*95</td><td></td></dl<>	* 95	
	OW-2A	629-002	7421	9-21-88	9-22-88	0.002	<dl< td=""><td>*95</td><td></td></dl<>	* 95	
	OW-1A	629-003	7421	9-21-88	9-22-88	0.002	<dl< td=""><td>*95</td><td></td></dl<>	* 95	
	OW-4A	629-004	7421	9-21-88	9-22-88	0.002	<dl< td=""><td>*95</td><td></td></dl<>	* 95	
	SW-3A	629-005	7421	9-21-88	9-22-88	0.002	0.002	* 95	
	SW-1A	629-006	7421	9-21-88	9-22-88	0.002	0.007	113/<1.0	
	SW-2A	629-007	7421	9-21-88	9-22-88	i 0.002 i	0.008	*95	
					· 	İ			
						<u>.</u>			

*THIS INDICATES A 95% CONFIDENCE LIMIT ACHIEVED WITH AN EPA QUALITY CONTROL SOLUTION ANALYZED ALONG WITH YOUR SAMPLE.

PREPARED FOR: CONESTOGA ROVERS

Metals Analysis for: TOTAL CHROMIUM

SAMPLE ID CUST.	HAS #	EPA METHOD	DATE PREPARED	DATE ANALYZED	DETECTION LIMIT	RESULT ppm	QC %SPKR/RPD
 OW3-88	 616-001	 7191	 9-13-88	 9-15-88	l 0.003	l <dl< td=""><td> *95</td></dl<>	 * 95
OW103-88		7191		19-15-88	0.003	⟨DL	126/<1.0
SS1-88		6010		9-15-88	2.27	16	+95
ISS2-88	•	•	•	9-15-88	1.47	19.3	*95
ISS3-88	616-005	6010		9-15-88	1.99	22.5	85/1.3
1	}		1	1			1
SS-4	618-001	6010	9-13-88	9-14-88	2.29	188	*95
iss-5	618-002		•	9-14-88	2.05	189	* 95
ISS-6	618-003	6010	9-13-88	9-14-88	1.6	479	* 95
SS-7	618-004	6010	9-13-88	9-29-88	2.15	165	* 95
SS-107	618-005	6010	9-13-88	9-29-88	2.28	134	*95
SS-8	618-006	6010	9-13-88	9-14-88	2.19	30.1	* 95
SS-9	618-007	6010	9-13-88	9-14-88	2.24	19.2	* 95
SS-10	618-008	6010	9-13-88	9-14-88	2.06	27.8	* 95
SED 1	618-009	6010	9-13-88	9-14-88	2.14	25.1	* 95
SED 2	618-010	6010	9-13-88	9-14-88	1.57	19.8	* 95
SED 3	618-011		,	9-14-88	1.99	18.8	* 95
SED 103	618-012	, ,		9-14-88	1.51	19.1	* 95
SED-4		•		9-14-88	2.04	21.2	* 95
SED-5	1			9-14-88	2.34	16.7	* 95
SW-1	•	, ,		9-15-88	0.003	0.005	*95
SW-2	618-016			9-15-88	0.003	<dl< td=""><td> *95 </td></dl<>	*95
SW-102	618-017	, ,		9-15-88	0.003	<dl< td=""><td>*95 </td></dl<>	* 95
SW-3	618-018	7191	9-13-88	9-15-88	0.003	<dl< td=""><td>*95 </td></dl<>	* 95
	·						
							İ

*THIS INDICATES A 95% CONFIDENCE LIMIT ACHIEVED WITH AN EPA QUALITY CONTROL SOLUTION ANALYZED ALONG WITH YOUR SAMPLE.

PREPARED FOR: CONESTOGA ROVERS

Metals Analysis for: TOTAL CHROMIUM

SAMPLE ID	HAS #	EPA METHOD	DATE PREPARED	•	DETECTION LIMIT	RESULT ppm	QC %SPKR/RPD
W1-88	 622-001	 7191	l 19-15-88	l 19-15-88	l l 0.003	l 0.013	l l *95
)W2-88	622-002	•	•	9-15-88	0.003	0.006	124/<1.0
RIP BLK	622-003		•	9-15-88	0.003	0.009	*95
W-3A	 629-001	 7191	 9-21-88	 9-22-88	l 0.003	 <dl< td=""><td> *95</td></dl<>	 *95
W-2A	1629-002	•	1	9-22-88	0.003	0.003	* 95
W-1A	1629-003	•	•	9-22-88	0.003	0.007	* 95
W-4A	629-004	•	•	9-22-88	0.003	<dl< td=""><td>*95</td></dl<>	* 95
	629-005	•	•	9-22-88	0.003	<dl< td=""><td>*95</td></dl<>	* 95
W-1A	629-006	•	,	9-22-88	0.003		100/12.7
W-2A	1629-007	7191	•	9-22-88	0.003	0.003	+95
							İ
	i	i			Ì		
	i	İ		İ	İ		İ
	i	İ			ĺ		
	İ	İ					
	Ì	İ					•
	1						
	1						
]			
							f
	1						[
	ĺ						İ
	Ì	İ					
	1						ĺ
	l		İ				
	1				[
]		
]

*THIS INDICATES A-95% CONFIDENCE LIMIT ACHIEVED WITH AN EPA QUALITY CONTROL SOLUTION ANALYZED ALONG WITH YOUR SAMPLE.

PREPARED FOR: CONESTOGA ROVERS

Metals Analysis for: EP TOX LEAD

• • • • • • • • •						• • • • • • •	
SAMPLE ID		EPA	DATE	DATE	DETECTION	RESULT	QC
CUST.	HAS #	METHOD	PREPARED	ANALYZED	LIMIT	mg/L	*
1	1			1			1
SS1-88	616-003	6010	9-15-88	9-15-88	0.02	0.049	95
SS2-88	616-004	6010	9-15-88	9-15-88	0.02	0.048	95
SS3-88	616-005	6010	9-15-88	9-15-88	0.02	0.054	95
	1			İ			1
SS-4	618-001	6010	9-15-88	9-15-88	0.02	49.9	*95
SS-5	618-002	6010	9-15-88	9-15-88	0.02	15.6	*95
SS-6	618-003	6010	9-15-88	9-15-88	0.02	35.6	**<1.0
SS-7	618-004	6010	9-15-88	9-15-88	0.02	132	*95
SS-107	618-005	6010	9-15-88	9-15-88	0.02	229	*95
SS-8	618-006	6010	9-15-88	9-15-88	0.02	0.23	*95
SS-9	618-007	6010	9-15-88	9-15-88	0.02	0.066	*95
SS-10	618-008	6010	9-15-88	9-15-88	0.02	50.4	*95
SED 1	618-009	6010	9-15-88	9-15-88	0.02	0.12	*95
SED 2	618-010	6010	9-15-88	9-15-88	0.02	0.057	*95
SED 3	618-011	6010	9-15-88	9-15-88	0.02	0.035	*95
SED 103	618-012	6010	9-15-88	9-15-88	0.02	0.02	*95
SED 4	618-013	6010	9-15-88	9-15-88	0.02	0.043	*95
SED 5	618-014	6010	9-15-88	9-15-88	0.02	<dl< td=""><td> *95 </td></dl<>	*95
İ		1		1	l	1	
İ	İ	1					
İ	İ	ĺ	1	1			
i	İ	ĺ		•	Į		
i	İ	İ	ĺ	1	[1 1
i	İ		ĺ	1	l		
i	1	İ		1			
•	•	•	•	•	•		

*THIS INDICATES A 95% CONFIDENCE LIMIT ACHIEVED WITH AN EPA QUALITY CONTROL SOLUTION ANALYZED ALONG WITH YOUR SAMPLE.

**THIS SAMPLE WAS ANALYZED IN DUPLICATE WITH THE RPD INDICATED ABOVE.

THESE SAMPLES WERE EXTRACTED AND ANALYZED ACCORDING TO METHOD: CONTAINED IN "TEST METHODS FOR EVALUATING SOLID WASTE: "PHYSICAL/CHEMICAL METHODS" SW-846, 3RD EDITION.

PREPARED FOR: CONESTOGA ROVERS

Metals Analysis for: EP TOX CHROMIUM

SAMPLE ID		EPA	DATE	DATE	DETECTION	RESULT	l QC
CUST.	HAS #	•	•	ANALYZED	LIMIT	mg/L	*
							,
	ſ		! 	<u> </u>	! 		1
SS1-88	616-003	6010	9-15-88	9-15-88	0.02	<dl< td=""><td>95</td></dl<>	95
•	616-004			9-15-88	0.02	<dl< td=""><td>95</td></dl<>	95
ISS3-88	616-005	•	9-15-88	9-15-88	0.02	<dl< td=""><td>95</td></dl<>	95
ISS-4	618-001	6010	9-15-88	9-15-88	0.02	<dl< td=""><td>+95</td></dl<>	+95
ISS-5	618-002		9-15-88	9-15-88	0.02	0.12	* 95
ISS-6	618-003			9-15-88	0.02		94.0/1.3
•	618-004			9-15-88	0.02	0.038	* 95
ISS-107	618-005		'	9-15-88	0.02	0.024	* 95
SS-8	618-006	6010	9-15-88	9-15-88	0.02	<dl< td=""><td>*95</td></dl<>	* 95
•	618-007			9-15-88	0.02	<dl< td=""><td>*95</td></dl<>	*95
ISS-10	618-008		,	9-15-88	0.02	<dl< td=""><td>*95</td></dl<>	* 95
SED 1	618-009			9-15-88	0.02	<dl< td=""><td>×95</td></dl<>	×95
•	618-010			9-15-88	0.02	0.024	i *95
SED 3	618-011			9-15-88	0.02	<dl< td=""><td>+95</td></dl<>	+95
SED 103	618-012			9-15-88	0.02	<dl< td=""><td>*95</td></dl<>	*95
SED 4	618-013			9-15-88	0.02	<dl< td=""><td>*95</td></dl<>	*95
SED 5	618-014	6010	9-15-88	9-15-88	0.02	<dl< td=""><td> *95</td></dl<>	*95
							İ
							İ
•							
•					,		
[i		
					1		
1 	: 				; ; [[
1	;	i 1		ı	· •	1	1

*THIS INDICATES A 795% CONFIDENCE LIMIT ACHIEVED WITH AN EPA QUALITY CONTROL SOLUTION ANALYZED ALONG WITH YOUR SAMPLE.

THESE SAMPLES WERE EXTRACTED AND ANALYZED ACCORDING TO METHODS CONTAINED IN "TEST METHODS FOR EVALUATING SOLID WASTE: PHYSICAL/CHEMICAL METHODS" SW-846, 3RD EDITION.

Inorganic Wet Chemical Analyses

Analyte: TOX and *EOX

EPA Method No.: **9020

Date Received: 9/13/88 - 9/15/88

 Sample I.D.		Date Sampled	Date Analyzed	Detection	 Concentration	Units
<u>30-616-001</u>	OW3-88	9/12/88	9/15/88	10.0 	<dl< td=""><td>μg/l</td></dl<>	μg/l
30-616-002	OW103-88	9/12/88	9/15/88	10.0	18.6	μg/l
30-616-003	SS1-88	9/12/88	9/13/88	10	44	mg/kg
30-616-004	SS2-88	9/12/88	9/13/88	10	34	mg/kg
30-616-005	SS3-88	9/12/88	9/13/88	 10	15	mg/kg
30-618-001	SS-4	9/13/88	9/14/88	10	34	mg/kg
30-618-002	SS-5	9/13/88	9/14/88	 10	75 I	mg/kg
30-618-003	SS-6	9/13/88	9/14/88	 10	60	mg/kg
30-618-004	SS-7	9/13/88	9/14/88	 10	 26	mg/kg
30-618-005		9/13/88	9/14/88	 10	 34	mg/kg
30-618-006	SS-8	9/13/88	9/14/88	 10	35	mg/kg
 <u>30-618-007</u>	SS-9	9/13/88	9/14/88	 10	25	mg/kg
30-618-008	SS-10	9/13/88	9/14/88	10	17	mg/kg
 <u>30-618-009</u>	Sed 1	9/13/88 »	9/14/88	10	<dl< td=""><td>mg/kg</td></dl<>	mg/kg
30-618-010	Sed 2	9/13/88	9/14/88	10	<dl< td=""><td>mg/kg</td></dl<>	mg/kg
30-618-011	Sed 3	9/13/88	9/14/88	10	<dl< td=""><td>mg/kg</td></dl<>	mg/kg
30-618-012	Sed 103	9/13/88	9/14/88	10	<dl< td=""><td>mg/kg</td></dl<>	mg/kg
30-618-013	Sed 4	9/13/88	9/14/88	10	111	mg/kg
30-618-014	Sed 5	9/13/88	9/14/88	10	13	mg/kg
 <u>30-618-015</u>	SW1-88	9/13/88	9/15/88	10.0	15.4	μg/l

^{*}Extractable organic halides, modified EPA method 9020 for solids as received.

^{**}Subcontracted

Inorganic Wet Chemical Analyses

Analyte: TOX and *EOX

EPA Method No.: **9020

Date Received: 9/13/88 - 9/15/88

		Date	Date	Detection		
Sample I.D.	Client I.D.	Sampled	Analyzed	Limit	Concentration	Units
30-618-016	 SW-2	9/13/88	 9/15/88	10.0	60.4	<u>ид/1</u>
30-618-017	 SW-102	9/13/88	 9/15/88	10.0	 35.4	μg/l
30-618-018	 SW-3	9/13/88	 9/15/88	10.0	 49.0	μg/l
30-622-001	 OW1-88	9/15/88	 9/19/88	10.0	 28.6	μg/l
30-622-002	 OW2-88	9/15/88	 9/19/88	10.0	 <dl td="" <=""><td>μg/l</td></dl>	μg/l
30-622-004	 Trip Blank	9/15/88	 9/19/88	10.0		μg/l

^{*}Extractable organic halides, modified EPA method 9020 for solids as received.

^{**}Subcontracted

SAMPLE CUSTODIAN REPORT

ELAP #10833

Analysis for: Conestoga-Rovers & Associates

Date: 9/13/88

7703 Niagara Falls Boulevard

P.O. #: 2366

Niagara Falls, NY 14304

Client #: E046.029

HAS Job #: 30-616

Sample Identification and Information

			Sample	1	pH Upon	HAS Preservation/
Seq. No.	Bottle #	Customer I.D.		Analysis Requested		
			Ground			
001	01	OW3-88	Water	8080	≈7	<u> </u>
	l			1		
	02 & 03	n	71	i tox	н	<u>1</u>
				1		
	04 & 05	п	17	8020	-	<u>1</u> 1
						2-filtered at HAS
	06	Ħ	17	Total Cr and Pb	≈ 7	with .45µ filter
	1					
002	01	OW103-88	11	8080	<u>≈</u> 7	11
	02 & 03	π	n	TOX	п	1 1
	[[
	04 & 05	n	"	8020	-	1
				Į į		2-filtered at HAS
	06	п	п	Total Cr and Pb	<u>≈7</u>	with .45µ filter
	[]			EP Tox, Total Cr		
003	01	SS1-88	Soil	and Pb. 8080	•	11
	1 1					
	02 & 03	П	п	TOX	-	<u> </u>
				EP Tox, Total Cr		
004	01	SS2-88	п	and Pb. 8080	-	11
	[]					1
	<u> 02 & 03 </u>	н	п	TOX	•	1 1
				EP Tox, Total Cr		
005	01	SS3-88	т.	and Pb. 8080	-	1
	1					
	02 & 03	Ħ	#	TOX	•	<u>l</u>
						!
		A STATE		!		
						<u> </u>
]	. ⊸		!		
	! !			!		1

PRESERVATION CODE:

- 1 Cool 4°C
- 2 Nitric Acid to pH<2
- 3 Sulfuric Acid to pH<2
- 4 Sodium Thiosulfate added for Residual Chlorine
- 5 Phosphoric Acid to pH<2

- 6 Sodium Hydroxide to pH>12
- 7 Phosphoric Acid to pH 4.0, 1 g/l of Copper Sulfate
- 8 pH 6-8 with Sodium Hydroxide or Sulfuric Acid
- 9 Hydrochloric Acid to pH<2

SAMPLE CUSTODIAN REPORT

ELAP #10833

Analysis for: Conestoga-Rovers & Associates

Date: 9/13/88

7703 Niagara Falls Boulevard

P.O. #: 2366

Niagara Falls, NY 14304

Client #: E046.030

HAS Job #: 30-618

Sample Identification and Information

			Sample	1	pH Upon	HAS Preservation
eq. No.	Bottle #	Customer I.D.	Type	Analysis Requested		
				EP Tox, Total Cr	1	
001	01	SS-4	Soil	and Pb. 8080	-	i 1
	1				1	1
	02 & 03	n	m	TOX	! ! -	1
	<u> </u>			EP Tox, Total Cr	<u> </u>	1
002	01	SS-5	n	and Pb, 8080	! ! -	1 1
002	1 01	33-3		I and ID, 0000	<u> </u>	<u> </u>
	02 & 03	п ,	п	TOX		1
·	102 & 03				•	<u> </u>
000	1 07		п	EP Tox, Total Cr		!
003	01	SS-6		and Pb. 8080	-	<u> </u>
	02 & 03	н	77	TOX	-	11
				EP Tox, Total Cr		
004	01	SS-7	п	and Pb. 8080	-	11
				[1
	02 & 03	77	п	TOX	-	<u>i</u> 1
	1	I		EP Tox, Total Cr		1
005	i 01 i	SS-107	, n	and Pb, 8080	-	i 1
	1					
	02 & 03	n	n	TOX	-	i 1
	1			EP Tox, Total Cr		1
006	01	SS-8	Ħ	and Pb. 8080	_	1 1
000	<u> </u>	33-0		1 and 10, 8080		<u> </u>
	 02 & 03	n i	н	l mov		ļ
	102 & 03			TOX	-	<u> </u>
			_	EP Tox, Total Cr		
007	01	SS-9	Ħ	and Pb. 8080	-	<u> </u>
	1			1		
	<u> 102 & 03 </u>	, , , , , , , , , , , , , , , , , , ,	n	TOX		1
	1 1			EP Tox, Total Cr		
800	01 1	SS-10	n	and Pb, 8080	•	11
	1			1	* 4	
	02 & 03		n	TOX		1
		•		EP Tox, Total Cr		
009	i 01 i	→ Sed 1	17	and Pb, 8080	-	1
	[1	***************************************	1		
	! 02 & 03	n 1	11	TOX	_	1 1
	ATION CODE		· · · · · · · · · · · · · · · · · · ·	LOX		

PRESERVATION CODE:

- 1 Cool 4°C
- 2 Nitric Acid to pH<2
- 3 Sulfuric Acid to pH<2
- 4 Sodium Thiosulfate added for Residual Chlorine
- 5 Phosphoric Acid to pH<2

- 6 Sodium Hydroxide to pH>12
- 7 Phosphoric Acid to pH 4.0, 1 g/l of Copper Sulfate
- 8 = pH 6-8 with Sodium Hydroxide or Sulfuric Acid
- 9 Hydrochloric Acid to pH<2

SAMPLE CUSTODIAN REPORT

ELAP #10833

_	• •	 ,	
A	lvsi	 ~~.	

for: Conestoga-Rovers & Associates

Date: 9/13/88

7703 Niagara Falls Boulevard

P.O. #: 2366

Niagara Falls, NY 14304

Client #: E046.030

HAS Job #: 30-618

Sample Identification and Information

]		Sample			HAS Preservation
Sea. No.	Bottle #	Customer I.D.	Type	Analysis Requested	Arrival	Comments
	1			EP Tox, Total Cr		
010	01 i	Sed 2	Soil	and Pb. 8080	.	i <u>1</u>
	1 1		1	1		
	02 & 03	n) } #	TOX	, ! -	1 1
	102 & 03 1		<u> </u>	EP Tox, Total Cr	! !	1
011	01	Sed 3	! ! "	and Pb. 8080	1	1
011	<u> </u>	Sed 2	<u> </u>	i and FB, 8080	<u> </u>	<u> </u>
		n	ļ j #	l may	!	1 7
	102 & 03	11	<u>''</u>	TOX	<u> </u>	<u> </u>
	1 1		l	EP Tox, Total Cr		!
012	01	Sed 103	<u> </u>	and Pb, 8080	<u> </u>	<u> </u>
			1	1		
	102 & 03	п	<u>"</u>	TOX	-	11
	Ī		ĺ	EP Tox, Total Cr		
013	i 01 i	Sed 4	j #	and Pb, 8080	-	1 1
	1 1		 			l
	02 & 03	n	! #	TOX	i -	i 1
	102 6 03 1		<u> </u> 	EP Tox, Total Cr	l	
014	01	Sed 5	! ! #	and Pb, 8080	i. İ -	1
014	<u> </u>		<u> </u>	and rb. 5000	1	1
		m	! ! "	l mov	{ !	! ! 1
	102 & 03		L	TOX		<u> </u>
	1		Ground		!	
015	01	SW-1	Water	1 8080	<u> ≈6</u>	
	1 1]			1
	02 1	Я	n	Total Cr and Pb	<2	1 2
	1		1	1		
	i03 & 04 i	n	,	TOX	≈6	<u> </u>
	1		1		[
	05 & 06	₩	, ,	8020	-	i 1
	103 4 00 1		1	1	<u> </u>	
016	01	SW-2	ł j n	8080	! ! ≈6	1
OTO	 		<u> </u>	1 0000		<u> </u>
	00	- n	 	I Toward Dh	! <2	1 2
	02	**	! "	Total Cr and Pb	<u> </u>	<u> </u>
		-4	! _		!	
	103 & 04	п	<u> </u>	TOX	<u> ≈6</u>	<u> </u>
	1 1		I	1		!
	105 & 06 1	Ħ	1 "	8020	-	1

- 1 Cool 4°C
- 2 Nitric Acid to pH<2
- 3 = Sulfuric Acid to pH<2
- 4 Sodium Thiosulfate added for Residual Chlorine
- 5 Phosphoric Acid to pH<2

- 6 Sodium Hydroxide to pH>12
- 7 = Phosphoric Acid to pH 4.0, 1 g/l of Copper Sulfate
- 8 pH 6-8 with Sodium Hydroxide or Sulfuric Acid
- 9 Hydrochloric Acid to pH<2

SAMPLE CUSTODIAN REPORT

ELAP #10833

Analysis for: Conestoga-Rovers & Associates

Date: 9/13/88

7703 Niagara Falls Boulevard

P.O. #: 2366

Niagara Falls, NY 14304

Client #: E046.030

HAS Job #: 30-618

Sample Identification and Information

Sea. No.	 Bottle #	Customer I.D.	Sample Type	 Analysis Requested		HAS Preservation/
017	 01	SW-102	Ground Water	8080	 ≈6	
	02	п	n	Total Cr and Pb	 <2	1 2
	 03 & 04	H	Ħ	 TOX	 ≈6	11
	 05 & 06	11	Ħ	8020	<u> </u> 	1
018	01	 SW-3	π	8080	[<u> </u> ≈6	1
	02	п	n	Total Cr and Pb	<2	2
	 03 & 04	п	н	TOX	 ≈6	1
	 05 & 06	π	TT TT	8020	 	1
						1
	<u> </u>					
		-		1	<u> </u>	
		,				

PRESERVATION CODE:

- 1 Cool 4°C
- 2 = Nitric Acid to pH<2
- 3 = Sulfuric Acid to pH<2</pre>
- 4 Sodium Thiosulfate added for Residual Chlorine
- 5 Phosphoric Acid to pH<2

- 6 Sodium Hydroxide to pH>12
- 7 Phosphoric Acid to pH 4.0, 1 g/l of Copper Sulfate
- 8 pH 6-8 with Sodium Hydroxide or Sulfuric Acid
- 9 Hydrochloric Acid to pH<2

SAMPLE CUSTODIAN REPORT

ELAP #10833

Analveie	for:	Conestoga-Rovers	۶.	Acc
Anaivsis	IOT:	Conescoga-Rovers	œ	ASS

Associates

Date: 9/15/88

7703 Niagara Falls Boulevard

P.O. #: 2366

Niagara Falls, NY 14304

Client #: E046.031

HAS Job #: 30-622

Sample Identification and Information

Seq. No.	 Bottle #	Customer I.D.	Sample Type	 Analysis Requested		HAS Preservation, Comments
001	01	OW1-88	Ground Water	 8080 PCB	 ≈6	1
	02	π	Ħ	Total Cr and Pb	<2	2
	 03 & 04	Ħ	n	 8020 (Toluene)	<u> </u>	1
	 05 & 06	π		TOX	L≈6	1
002	01	OW2-88	н	8080 PCB	 ≈6	1
	 02	11	Ħ	 Total Cr and Pb	<2	<u> </u> 2
	 03 & 04	Ħ	Ħ	 8020 (Toluene)	-	1
	 05 & 06	n	я	TOX	 ≈6	1 1
003		Trip Blank	n	Total Cr and Pb	<2	2
004		Trip Blank	10	TOX	≈ 7	1
		-				
						<u> </u>
	[<u> </u>
	ļ 1					<u> </u>
	[◄				[
						ļ

PRESERVATION CODE:

- 1 Cool 4°C
- 2 Nitric Acid to pH<2
- 3 = Sulfuric Acid to pH<2</pre>
- 4 Sodium Thiosulfate added for Residual Chlorine
- 5 = Phosphoric Acid to pH<2
- 6 Sodium Hydroxide to pH>12
- 7 = Phosphoric Acid to pH 4.0, 1 g/l of Copper Sulfate
- 8 pH 6-8 with Sodium Hydroxide or Sulfuric Acid
- 9 Hydrochloric Acid to pH<2

CLIENT: CRA

DEPARTMENT: ENVIRONMENTAL

9-20-88

BATCH: 30-629

PROJECT:

DATE:

CUSTODIAN: KAS

1=COOL 4 C

2=NITRIC ACID TO pH(2

3=SULFURIC ACID TO pH(2

4=SODIUM THIOSULFATE ADDED FOR RESIDUAL CHLORINE

5=PHOSPHORIC ACID TO pHC2

6=SODIUM HYDROXIDE TO pH>12

7=PHOSPHORIC ACID TO pH 4,1g/1 OF COPPER SULFATE

B=LAB PACK

9=SANITARY SEWER

10=APPROVED DISPOSAL FACILITY

11=STERILIZED FOR STERILE LAND FILL

					•	SAMPLE RECEIPT		•				
		BOTTLE	CLIENT	DATE SAMPLED	:DATE :RECEIVED	: :DESCRIPTION :	ANALYS REQUES			:PRESERVATION/ :PROCESS	ISAMPL ICONHE	E CONDITION NTS
10-6	29-001	; }	10N-3V	19-20-88	-; :9-20-88	: WATER	· 	Cr, Pb	i<2	1 #2	:	600D
	29-002		ION-2A		19-20-88		i	Cr, Pb	1(2	12	i	600D
	29-003		ION-IA		19-20-88		i	Cr, Pb	1(2	1 12	i	6000
	29-004		ION-4A		19-20-88		i	Cr, Pb	1(2	12	1	GOOD
	29-005		ISW-3A		19-20-88			Cr, Pb	1(2	1 12		6000
	29-006		ISW-IA		19-20-88			Cr, Pb	1(2	1 12	1	6000
	29-007		ISW-2A		19-20-88		•	Cr, Pb	1<2	1 12		600D
		:	1	1	;	;	:	•	:	:	:	
		:	1		1	!	ł		:	:	:	
		:	ŧ	:	1	:	!		:	1	:	
		!	:	;	:	1	;		:	ŧ	:	
		;	;	ł	1	:	1		:	:	:	
		!	:	;	1	:	:	•	:	•	:	
		:	:	:	1	:	!		:	ŧ	;	
		:	:	!	1	:	:		1	:	t	
		1	;	ŧ	:	1	ŧ		:	:	:	
		1	:	:	:	!	ŧ		:	:	:	
		1	:	1	:	1	†		:	å å	:	
		:	:	:	1	:	†		ŧ.	:	:	
		1	t	:	:	!	:		:	t	:	
		:	•	ŧ	:	;	1		:	:	:	
		;	:	i	1	:	1		1	:	:	
		;	:	ŧ	1	•	ŀ		:	i	1	
		:	;	:	:	:	ŧ		1	1	1	
				1	:	1	:		:	:	i	
		:	i	1		1	•		1	•	i	
		:	i	:		:			1	•	1	
		i	i			:	1		1	1	i	
		1	;	•	!	!	•			1	٠.	

HUN111	ngDun an	ALTITCAL	SERVILES		ANALY	ILAL KE	WUESI F	ORM	Pag	e / of	
Tracking Into Init 9-14 150	Addres	er Name 103 N.A.	GARA FALLS	5 BUD			23(66) J Address	Quote #	9-12 H	AS Ref.	30-616
	1 10M	Y MISE	RCOLL	283-676 Phone #	ල ල	Date Si	ubmitted	9/13/88	••	e VERBA	EN - 9-2 LS-9-15
Sampler(s) Oustomer	Signature	1	Sample	Sample	INo. of		of Samo	les 5	Client	No. EC	146.029
I.D.	_Date_	Time	Type	Location	•		nalysis R	equested	Comments	HAS No.	Billing
aw 3-881	9/13/88	3:25	lunter	Pyramid	15	8080,7	0x 802	2, Cx+Pb.		1001	825
8-E010x		13.25	<u> </u>) '	15		,			لانما	1 1
51-88	• •	Tup	5011		13	8080 7	OX EPÍC	X+Dbl C	+Pb	 ce'3	1050
55 2-88		I IFAP	<u> </u>	1	3		,, ,,			1004	1_1_
35 3-89	/1	Tnp		"	3_		4			1005	1
	-	 									
Relinqui	_	1	 	Received	by	Reling	guished by	/ Dat	e/Time	 Recei	l ved by
Relinqui	shed by	Date	e/Time	Received	by	Reling	pulshed by	/ Dat	e/Tima	Recei	ved by
Comments:		1						Wet Met. Chem.		70X Other	Final Total

Tracking	Ouston	HAS to ser Name	AES_		 	P.O. #Billing Addi		Quote #	·		
	_ _ Contac	<u> </u>		Phone #		Date Submit	ted /	9/13/88	Date Due	:	
	Signature	بروسان والمساور والمساور والمساور				Total # of	Samples		Client N	o .	
Oustomer I.D.	Date	Time	Sample Type	Sample Location		Analys	ls Requ	ested	Comments	IHAS No.	Billing
30-616-001	.02					TOX		i	210-ppb	1	
30-616-002	.02					/ '			18.6 ррь	<u> </u>	
				 	 			•			
Relingu	ished by	Date	/Time	Received		Relinquish	1 1	9/13/88	Pime 30	(1)	ved by
	lathed by		/Time	Received		Relipopulah		Date/1		Receiv	

HUNLING	<u> NA MUUE</u>	ALTIICAL	SERVICES	••••••••••••••••••••••••••••••••••••••	ANALYI	ICAL REQUE	.SI FURI	1	Page	cf_	-
Tracking		er Name	· AF	5		P.O. #	·	Quote #	HAS	S Ref. #	
ite Init.					_	Billing Ad	dress				e de la companya de l
	 					•					
					İ			/ /		•	
	Contac	t Name		Phone #		Date Submi	tted 9	/13/88	Date Due		
moler(s) S	ignature					Total # of	Samples		Client N	9.	
Austomer I.D.	Date	Time	Sample Type	Sample Location	No. of Cont.	Analy	sis Requ	ested	Comments	ihas yo.	Billing
) (0/10-cx)1	C.L	 		<u> </u>	<u> </u>	XX	VER	3ALS 9-15	1/Ong/l		
)-1316001-] 		L		١,			18.6 mg/e		
16160013	-0.	1			 	• ,			44 mg/kg		
1/1000/	0}	l !	•	<u> </u> 	<u> </u>	•(34 mg/kg		
5.6/6-09s									1 15 mg/kg	-	
									00		
		<u> </u>			<u> </u>						
		<u> </u>	artiferen errora errora errora errora errora errora errora errora errora errora errora errora errora errora er								
		l			 	<u></u>		•			/ // _ // // // // // // // // //
Relinquis	_	1	/Time	Received	_ /i	Relinquis	///	Date/1	rime (//. 0.0 (Receiv	ed by
Relinquis			/Time	Received		Relinquis		Date/		Receiv	ed by

Tracking to Ini -14 151	/ Address	n Name	IY	is BUN				3106_ ddress	_ Qux	ote #	HA	S Ref. #	<u>30-618</u>
	Contact	Y Mise	REALA	28367a Phone #	_ <u>20</u>	Date	Subm	itted	9/1-	3/88_		VERBALS 9-15	N-9-22 PM or 9-16
moler(s) ustomer I.D.	Signature Date	Time	Sample Type	Sample Location	No. of			f Samol Vais Re			Client N	1	16.030 Billing
54	9113158		Sail_	Rusaniel	3	80,800		/		C1+Pp	Southeartes	1001	1050
55 5	 	10.10	11	1	3	• • •	,,,	11	h			1007	
56	4,	\v:3¢	• • •		3	1	t 1	••	• •	1,		1003	
57	£,	10:40		1,	l L 3_	-1	١,	١,	١,			Looy	
<u> </u>	١,	10: 40	١,	"	3	1.	• 1	٠.	٨			1. 1005	
58	,	10:55	11		1 3	1,		٠,				1006	
59	١,	11:05			3	h	١,	`.	,,	•		1007	
310	١,	14:55	٠.	\	3	,l	١,	٠,	ĸ			8001	
seel 1		امر: ٥٥	• •	1 ,,	3	1.	11		Į.			1009	
ed 2	Lahed by	14:40			3			, ·	<i>(</i> *			100	V
_	-		Cust-rly	Received	y ya	Re.	inqui	shed by		Date/I	'1me	Receiv	ved by
Rel inqu	Lethed by	Date	/Time	Received	by	Rel	inquie	shed by		Date/T	'ime	Receiv	ved by

HUNT	INGDON AN	<u>ALYTICAL</u>	SERVICES		ANALY	TICAL REQU	JEST FO	DRM_		Page	a a of	<u></u>	
Tracking Date Ini	Custom	er Name				 P.O. # 2 Billing # 	•	Quo	te #	10	AS Ref. #	30.6	<u>18</u>
						, 					,		F
	Contac	t Name		Phone #		Date Subr	nitted	9113	88	Date Du	.		
Sampler(s) Customer I.D.	Signature Date	i I Time	Sample Type	Sample Location	[No. of		of Sampl Ysis Re	_	d i	Client N	l HAS No.	Bill	
	9/13/88	1	 \$0;	Rycamia	l	XOT, OB'CR	/	<u> </u>	1	Sallibalted	1011	105	_
Sect 103	1	112:15	4		3	1	'\	• ,	1		1012	1	
Sec 2 4	\ <u> </u>	77:30 	.		3	١, ١	•	۱.	١. ا		1013		1
5ecl 5		11:50	<u> </u>	<u> </u>	l L 3						014	1	
Sw 1	٠.	13:45	worker		6	NOT, OROR	Total	<u>.r +P</u> p	1000 N		· 015	82:	 5
ടധമ		14:30		<u> </u>	9	h k	•	• .	· .		ا ا <i>۵ا</i> ن	1	
20109		14:20	.,		16	۸ ۱.		41			017		
sw 3	,.	12:05	, \	<u> </u>	6	(1	• •	ţ.			018	<u>\partial \tau \tau \tau \tau \tau \tau \tau \tau</u>	
								·					
Relingu	lahed by	Date	/Time	Received	ру	Relinqui	shed by	1	Date/T	'ime	Receiv	ed by	
	Lethed by	Date	A Custon	Received	ру	Relinqui	shed by		Date/T	'ime	Receiv	ed by	,

HU	NLING	DUN AN	ALTICAL	SERVICES		ANALY	TICAL KEQUESI FOR	1	Page	of	1
Track	ing	Ouston	CiCiA er Name				P.O. # 23/66	Quote #	9-12 HA	S Ref. #	632
nte :	Init.		7703	NIAGARA	- FALLS BLUD	<u> </u>	Billing Address	***************************************			
9-14	TST	Addres	NF,	NY			1				
		——————————————————————————————————————	· Ú.		20.0						
		Contac		gcoud	<i>283 ⋅ 672</i> Phone #	0	Date Submitted	115/88	Date Due	WRATE.	N 9.22 9-19
ro franc	(a) Si	onature			A A A A A A A A A A A A A A A A A A A		 Total # of Samples	-3		6. FO ^L	
Custom	ar			Sample		No. 01			Client N	6. 1-0-	16.03
I.D.		Date	Time	Type	Location	i	Analysis Requ		Comments	IHAS No.	
	- ' /	115/88	0830	H20	001)	TOX (900			001	825
	_		<u> </u>	<u> </u>	<u> </u>	9	1 Toluene (80	(<u>0</u> 00	! 	<u> </u>	
			<u> </u> 	 		1	1 A ROCHIOC	1354 60	1 80)	<u> </u>	
			<u> </u>	l . L		11	Unetals To	•	Filk1/11.003		
	19	115/08	0730	H ₂ 0	1047	12	TO X (9020		 	003-	825
						10	1 Tolurge (80			1 00.3	Oix C
					<u> </u>						
<u></u>		·				<u> </u>	JAMCHOR 100		1 /	<u> </u>	
···					1	<u> </u>	1		F. 16/111003)
- 1	<u> </u>			H20	THIP BLANK	<u> </u>	TOX (9000			00.34	312 50
Ralin	mulsh	ed by	Date	e/Time	Received	1	metials Ta	-+ Pb)	Filk1/1/203	003	125
	Ji. C		· /_	(°0C)	Mecetived	υγ	Relinquished by	Date/1	l'IMe	Receiv	ed by
Relir		ed by		e/Time	Received	by	Relinquished by	Date/1	lime	Receiv	ed by
					<u> </u>		_ 	•	İ		
Coment	is:						Initial	Wet 60	C Trip	TOX	Final
	T			70 S. S.	r jan yr er as		1 TOCAL MET	IChem. I Or	g. Blanksi	Other 1	" tal

nun	ן רעסחטע יאני	IALTITUAL	SERVILES		ANALY	ILAL REWUESI FORM	Page of			
Tracking ata Ini	Custon	er Name	s DES			P.O. # Quote # Billing Address	43f	S Ref. #	<u>.</u>	
	Addres	18								
									to Principle and	
	Contrac	t Name		Phone #		Date Submitted 7/16/98	Date Die	}		
mpler(s) watomer I.D.	•	 Time	Sample Type	Sample Location		Total # of Samples 3 Analysis Requested	Client N Comments		 Billing	
6 2.2	 CO	1							DITTHM	
	109 1	1				'\n\				
	1	1. Cu								
· (v)	120.4	-			<u> </u>	TOY (TRIP 13km/K)	115mall 50mp	(1		
		 						<u> </u>		
	<u> </u>	<u> </u>		<u> </u>			<u> </u>	· 		
	! !	<u> </u>		<u>.</u>						
	1	1								
7				<u> </u>						
	<u> </u>			<u> </u>	<u></u> 			<u> </u>		
	<u> </u>	<u> </u>					<u> </u>	<u> </u>		
Relinou	ished by		/Time	Received	bu	Relinquished by Date	mi	اــــا		
Junio L	-	9/15/88		Ravyo	Teeles	Relinquished by Date,		Receive Ludy Of		
	lahed by		/Time	Received	by 0	Relinquished by Date	17:111.	Receiv		
Xmenta:		<u> </u>			1		Trip Trip Trip	Other	Final Total	

Tracking		CVA er Name		48 hr.	_TA	P.O. # 23/0/0			of of AS Ref. #	
te Init.	 Address	D	NIAGARA NY 14.	Faces Bed 304	<u>2.</u> 	Billing Address				
	10NU Contact	Y MISER	COLA	283-672 Phone #	<u>0</u>	Date Submitted 9/	20/88	Date Due	VERBAL WRITE	5-9-22 U-9-23
mpler(s) Si Nustomer			Sample		No. of	Total # of Samples		Client N	ю. E04 1	6-033
1	Date	Time	Type	Location	Cont.	Analysis Reque	sted	1	HAS No.	Billir
u-3A 9	120/88	1045	1t20		<u> </u>	Cr , fb		two3 pHC2	001	75
W-DA	(1	1115				11		"	002	
W-1A	(1	1/00				. ((1 4	003	
W-4A	11	1200	. 11			1/		11	004	
1-3A		1130	11			11		1)	005	
W-1A	II	1145	11		,	11		1.	004	
W- 2A	/·	1200	i,			li .		11	007	V
			·		`		وروار والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع والمراجع	1	<u> </u>	
Relinquish	ed by	0	Time	Received	by	Relinquished by	Date/	Time	Receiv	red by
Relinquish		Date	/Time	Received	ру	Relinquished by	Date/	Time	Receiv	red by
anments:	PYE	AMID	- 48	hr. Verba	10	Initial Met.	Wet Chem. O	Trip rg. Blanks	<u> </u>	Final