SOIL GAS AND SOIL SAMPLING GULF STATE CREOSOTE-HATTIESBURG, MISSISSIPPI PREPARED BY US EPA/ERT-MAY 1990

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SOIL GAS AND SOIL SAMPLING

GULF STATES CREOSOTE HATTIESBURG, MISSISSIPPI

May, 1990

EPA Work Assignment No.: 1-335

Weston Work Order No.: 3347-11-01-2335 EPA Contract No.: 68-03-3482

FINAL REPORT

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rd:eh/ONEILL/FR-2335.R1

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1.1 Background

The former Gulf States Creosote Site is located in a commercial area of Hattiesburg. Mississippi (Figure 1). The site was an active wood preserving facility which operated from approximately 1920 to 1960. The property is currently owned by the city of Hattiesburg and subleased to several automobile dealerships, car-parts stores, a beverage distributer, a food store, and a furniture store. The process areas and wood drying/drip areas have been regraded, covered with asphalt, and are no longer evident. The former site encompasses approximately 20 acres, and is bordered on the east by railroad tracks, on the west by Gordons Creek, on the south by a drainage swale which feeds into Gordons creek, and on the north by Timothy Street.

The increase in surface runoff as a result of development and paving in the immediate area of the former site has significantly effected the flow into Gordons Creek. As a result, the U.S. Army Corps of Engineers (Corps) has been requested to rechannel Gordons Creek. In preparation for the rechannelization project, the Corps requested assistance from the US EPA in characterizing the nature and extent of contamination which may be present in the area as a result of former wood treating operations at the Gulf State Creosote site. Creosote and coal tar seeps are evident along the eastern bank of Gordons Creek.

1.2 Purpose of Investigation

The purpose of the ERT/REAC investigation was to identify the extent of contamination in the area of the former creosote plant. Specifically, areas adjacent to Gordons Creek were to be sampled in order to determine the nature and extent of contamination. If possible, an estimate of the volume of contaminated material/soils in the immediate area surrounding Gordons Creek was to be calculated.

During the soils investigation, ambient air monitoring/sampling was planned in order to identify any local air quality degradation which may have resulted from the presence of creosote residuals or during intrusive activities.

1.3 Summary of Activities

From January 20 to January 25, 1990, ERT/REAC completed a soil gas survey and preliminary soil sampling effort in the area surrounding the former plant site. The activities conducted during that investigation were summarized in a trip report dated February 16, 1990. Approximately 65 soil gas sampling stations were monitored and/or sampled. Analysis of soil gas samples was completed on-site using the ERT TAGA mobile tandem mass spectrometer (MS/MS). Fifteen soil borings were installed, from which ten (10) soil samples were collected and analyzed for Polynuclear Aromatic Hydrocarbons (PNA). Sampling activities had to be suspended due to an unusual amount of rain and subsequent rise in the water table.

On March 19 and 20, 1990, ERT/REAC returned to Hattiesburg to complete the soil borings and subsurface soil sampling investigation. The activities completed during this site visit are summarized in a trip report dated March 30, 1990. A total of fifteen soil borings were installed, from which nine (9) samples were collected and analyzed for PNA's.

Air sampling and monitoring was also conducted on January 23rd. A total of three air samples representing static conditions (i.e. prior to soil disturbance) were collected and analyzed for PNA compounds.

Appendix A includes copies of the two trip reports.



1.4 Methods of Investigation

Soil gas sampling activities followed procedures outlined in ERT/REAC Standard Operating Procedure (SOP) #2149. The installation of soil borings was facilitated through the use of a "Little Beaver" power auger and procedures defined in ERT/REAC SOP #2122. Samples were collected using a stainless steel hand auger and followed procedures defined in ERT/REAC SOP #2012 and 2127.

Air sampling procedures were conducted according to ERT/REAC SOP# 2066, and air monitoring procedures followed ERT/REAC SOP# 2060, "RAM-1". Sample collection protocols and analytical techniques for PNA's adhered to NIOSH air sampling procedure #5515.

2.0 RESULTS

2.1 Soil Gas Survey

The soil gas survey produced mixed results. A summary of real-time monitoring data collected with the HNU Photoionization Detector and Foxboro Organic Vapor Analyzer is provided in Table 1. Preliminary soil gas sample results indicated naphthalene, the target compound of interest, in the 10 to 100 parts per billion (ppb) range in numerous samples. Benzene, toluene, and xylene (BTX) were also identified in some samples in the low ppb range. After additional quantification, the TAGA results were finalized and the contaminants were found to be below the detection limit in all soil gas samples. The detection limit for naphthalene was calculated to be 40 ppb. Appendix B contains a copy of the final TAGA data.

2.2 Soil Borings/Soil Sample Collection

A total of nineteen (19) soil samples were analyzed for PNA compounds by GC/MS. Those samples were collected from fourteen (14) different borings. Depth of sample collection waried between five (5) and fifteen (15) feet below surface. This range corresponds to the contaminated soil horizons. Of the 19 samples analyzed, twelve (12) can be considered contaminated with various PNA compounds. Table 2 presents a summary of soil results for those samples collected in January, 1990. Table 3 presents a summary of soil results for those samples collected in March, 1990.

2.3 Air Sampling/Monitoring

Air sampling consisted of collecting 900 liters of air through an XAD tube/filter using a personal air sampling pump. Three locations were sampled prior to initiating site activities. All samples indicated non-detectable levels of contaminants. Air monitoring consisted of using an MIE RAM-1 with data logger. Total particulate concentrations were integrated over a 2.5 hour period. Average concentration was .008 mg/m³ with a maximum measurement of 1.09 mg/m³.

3.0 DISCUSSION OF RESULTS

3.1 Soil Gas Results

There appears to be no relationship between real-time screening results and TAGA (MS/MS) soil gas analysis. Furthermore, there does not appear to be any spatial relationship between screening results and the former plant site location. Some inconsistency and variation in screening results could be due to a combination of equipment failure, weather conditions (high humidity), and soil moisture.

The soil gas concentrations, which proved to be lower than TAGA detection limits, could in part be due to the time span between site activity and sampling (30 years). Either the creosote compounds have naturally decayed to a point where volatilization is minimal or the material has migrated and collected to downgradient locations.

3.2 Soil Sampling Results

Of the soil samples identified as contaminated, those collected from the area bordered by West Pine Street and Gordons Creek, south of the drainage ditch which runs underneath West Pine Street, appear to contain the highest concentrations (Figure 2). Specifically, samples collected from this area include D-00, D01, E-24, E-25, and E-27. Sample E-20, located on the northeast side of the drainage swale also had significant contamination. The contamination identified in B-25 is significantly less than that identified in other samples and may have been influenced by surface conditions (i.e., adjacent asphalt parking lots). This sample was collected from the 0 to 12 inch depth. Likewise, the minor contamination found in sample D-03A may have also been influenced by surface conditions.

3.3 Air Sampling/Monitoring Results

Because of the extreme precipitation encountered, the air sampling and ambient monitoring effort was abandoned. Static air quality conditions do not appear to reflect any effect from coal tar residuals. The sampling results are not representative of ambient conditions which may result during intrusive soil disturbance activities in contaminated areas.

4.0 CONCLUSIONS AND VOLUME ESTIMATES

4.1 Conclusions

The findings of this investigation indicate that there is no spatial relationship between the former plant site lay-out and the residual contamination (Figure 3). This investigation did not characterize conditions east of Timothy Street where the former process area and storage vessels were located. Due to natural surface drainage conditions and topographical relief, one would expect to find the bulk of contamination west of Timothy Street. The focus of this investigation was west of Timothy, and specifically the area(s) just east of Gordons Creek.

The fact that significant contamination was not found in areas removed from Gordons Creek could indicate that contaminants have migrated to that downgradient location over the years following plant closure. Another explanation may be that during shut down of the plant or construction of West Pine Street, the bulk of surface materials was dumped or bull-dozed into that area. Another explanation may be that contamination is randomly dispersed and so low in concentration that the soil gas sampling was not able to detect the contamination.

4.2 Volume Estimates

An estimated volume of soil that is contaminated with PNA compounds from the presence of creosote was computed based on ERT/REAC field observations. The soil borings and the visual assessment made along Gordons Creek provided enough information to approximate an area of contamination, which is designated on Figures 2 and 3. Based on an estimated thickness of the contamination (three feet), the volume calculation yielded approximately 7,200 yd³ of contaminated soil.

A second calculation was performed using a thickness of five feet as a worst case scenario. Creosote outcroppings approximately five feet in thickness were visible along the banks of Gordons Creek, and due to the thick underlying clay layer, the water table fluctuates quite a bit between the surface and twenty feet. These two observations support using five feet in the calculation which yielded a volume of 12,000 yd³.

These estimates will be used in planning remedial measures based on treatability studies presently being performed by ERT/REAC.

TABLE 1. SOIL GAS FIELD SCREENING DATA

GULF STATES CREOSOTE SITE HATTIESBURG, MISSISSIPPI JANUARY 22-26, 1990

Sample Location	Sample Number	Instrument OVA	Reading HNU	
•			·	
A01	01521	0.5	0.0	
A02	01522	3.5	0.0	
A03	01523	1.8	0.0	
A04	01524	2.0	0.0	
A05	01525	5.0	0.0	
A06	01526	1.0	0.0	
A07	01527	18.5	NR	
A08	01528	0.5	NR	
A09	01529	3.5	NR	
A10	01530	400.0	NR	
A11	00611	0.0	NR	
413 A12	00612	18.0	NR	
B01	01538	0.6	0.0	
B02	01539	0.4	0.0	
B03	01540	0.4	0.0	
⇒ B04	NS	NR	NR	
B05	NS	NR	NR	
B06	00621	1000.0	0.0	
B07	00622	2.0	NR	
908 ₹	00625	1.5	NR	
B09	00624	0.6	NR	
B10	NS	NR	NR	
J. B11	NS	NR	NR	
B12	00623	1.0	NR	
C01	01491	NR	0.0	
C02	NS	NR	0.0	
C03	01492	NR	0.0	
C04	NS	NR	0.0	
C05	01493	NR	0.0	
C06	01494	NR	0.5	
C07	NS	NR	0.0	
C08	NS	NR	0.0	
C09	01497	NR	0.5	
C10	NS	NR	NR	
C11	01498	NR	2.0	
C12	NS	NR	0.0	
Ou. C13	01500	NR	0.0	
C14	NS	NR	0.0	
De - C15	01501	NR	0.0	
Des (C13 C14 C15 C16	NS	NR	0.0	
.4 /		• • • •		

NS - No sample collected. NR - Reading not taken.

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TABLE 1 (CONT'D). SOIL GAS FIELD SCREENING DATA

GULF STATES CREOSOTE SITE HATTIESBURG, MISSISSIPPI JANUARY 22-26, 1990

Sample	Sample	Instrument Reading		
Location	Number	OVA	HNŪ	
C17	NS	NR	NR	
E01	01531	80.0	NR	
E02	01522	880.0	NR	
E03	01533	2.0	NR	
E04	01534	3.2	NR	
·E05	01535	2.4	NR	
£06	01536	2.0	NR	
E 07	01537	NR	NR	
E07	01502	0.8	0.0	
E08	NS	0.2	NR	
-E09	01503	0.6	NR	
-E10	01505	1.0	NR	
E11	01506	0.2	NR	
E12	NS	NR	NR	
E21	NS	NR	NR	
-E22	01509	NEG	NR	
.E23	01508	30.0	NR	
E24	01507	NR	NR	

NS - No sample collected. NR - Reading not taken. NEG - Negative reading.



TABLE 2. SUMMARY OF SOILS ANALYSIS

GULF STATES CREOSOTE SITE HATTIESBURG, MISSISSIPPI JANUARY, 1990

Parts per million (ppm)

	Sample Location	B0 2.5	D00	D00	D01	D01	E20
Compound Name	Sample Depth	0-12 in.	5 ft.	8 ft.	5 ft.	8 ft.	4 ft.
Naphthalene		•	178	354	280	148	4.1J
2-Methylnaphthalene		*	99	197	460	82	3.6J
1-Methylnaphthalene		*	72	104	340	45	*
Biphenyl		*	22J	55	9J	24	*
2,6-Dimethylnaphthalene		•	72	66	53	28	*
Acenaphthylene		*	4.4J	4.2J	2.3J	. *	*
Acenaphthene		*	259	156	225	81	14J
Dibenzofuran		•	158	125	114	78	4.73
Fluorene		*	245	140	219	90	9.4J
Phenanthrene		6.5J	718	325	715	229	26
Anthracene		•	465	210	521	114	69
Carbazole		*	173	96	157	38	15J
Fluoranthene		3J	844	215	763	188	138
ene		1.1J	181	64	266	65	98
nzo(a)anthracene		1.6J	181	54	259	62	104
Chrysene		2.9J	230	61	318	73	160
Benzo(b)fluoranthene		3.8J	*	78	143	127	248
Benzo(k)fluoranthene		*	231	74	135	121	236
Benzo(e)pyrene		2.5J	83	25	97	52	83
Benzo(a)pyrene	0	2.5J	125	35	133	55	116
Indeno(1,2,3-cd)pyrene		1.8J	51	15J	54	26	53
Dibenzo(a,h)anthracene		.5J	23	5J	19J	12J	17J
Benzo(g,h,i)perylene		1.5J	41	11J	42	22	42
75/~1			4,455		5322		

^{* -} Non-detectable levels.

J - Data indicates the presence of a compound that meets the identification criteria. The result is less than the lowest linear detection limit of 10.0 ug/ml, but greater than zero and the concentration is given as an approximate value.

TABLE 3. SUMMARY OF SOILS ANALYSIS

OULF STATES CREOSOTE SITE HATTIESBURG, MISSISSIPPI MARCH, 1990

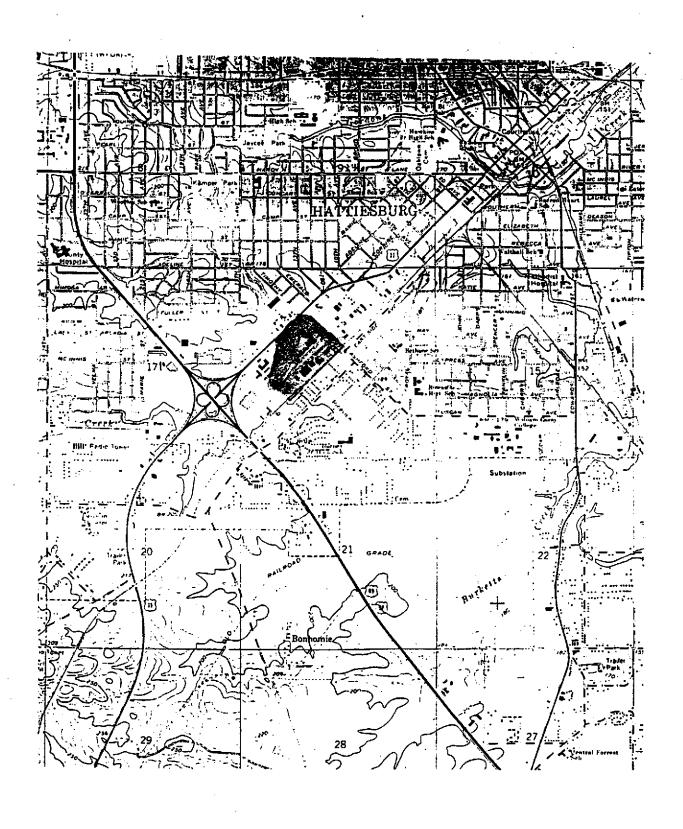
Paris per million (ppm)

Compound Name	Sample Location Sample Depth	D03A 10 ft. Top of Auger	D03A Bottom of Augor	E19 11 ft.	E24 8 ft.	E25 8 n.	B27 8 ft.
Naphthalone		0.53	7.3	2.5	544	48	753
2-Methylnaphthalene		•	.13 +	.9	224	26	293
1-Methylnaphthalene		•	.063	.6	107	26	193
Biphonyl		•	.023	.3J	\$5	3.51	140
2.6.Dimethylnaphthalene		• •	•	.43	71	13	160
Acenaphthylene		•	•	.043	7.33	241	20
Acenaphinene		•	.13	1.5	254	86	213
Dibenzolutan		•	.053	.7	159	37	125
Pluorene		•	.05J	.9	194	66	129
Phenanthrene		•	.043	2.7	420	136	425
Anthrucene		•	•	1.7	87	41	126
Carbazolo		•	.07	3	48	5.51	59
Fluoranthene		.13	.03J	2.9	224	144	288
Pyrene		.23	.043	3.4	180	126	296
Benzo(a)anthracene		.073	1	1.1	52	34	100
Chrysene		.083	•	1.2	. 42	37	86
Benzo(b)fluoranthene		•	•	1.0	•	•	86
Benzo(k)fluoranthene		•	•	.4	273	30	•
Benzo(e)pyrene		•	•	.5	•	9.73	31
Benzo(a) pyrene		•	•	.6	*●	11	42
Indeno(1,2,3-od)pyrene		•	•	•	•	•	•
Dibenzo(a,h)anthracene		•	•	•	•	•	. •
Benzo(g,h,l)perylene		•	•	•	•	•	•

^{• -} Non-detectable levels.

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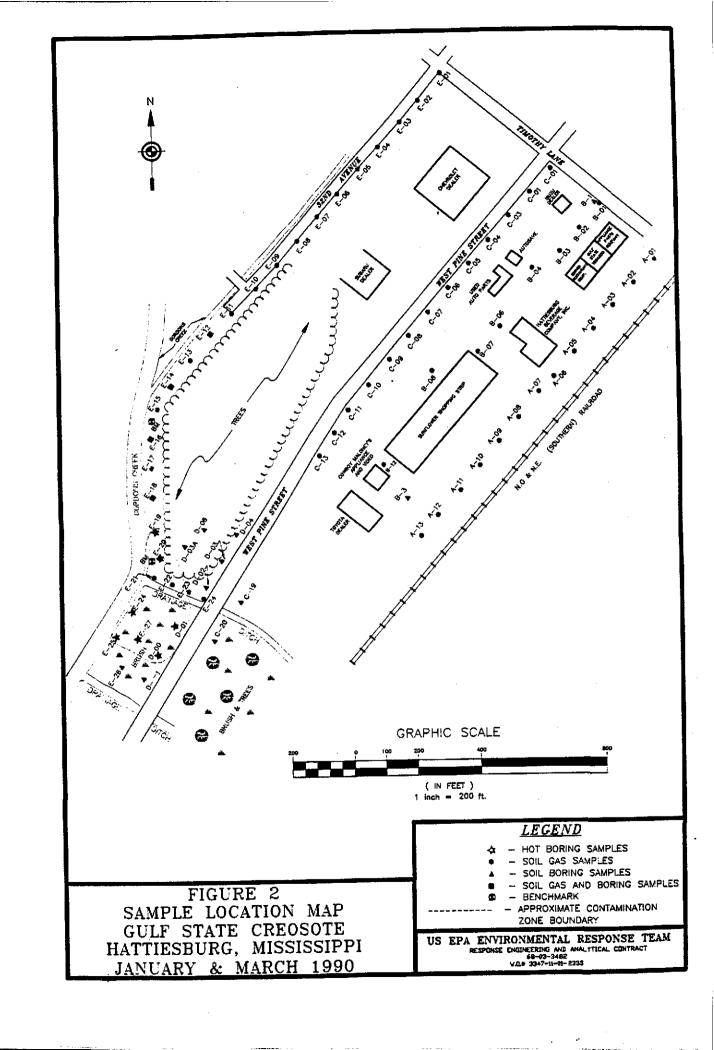


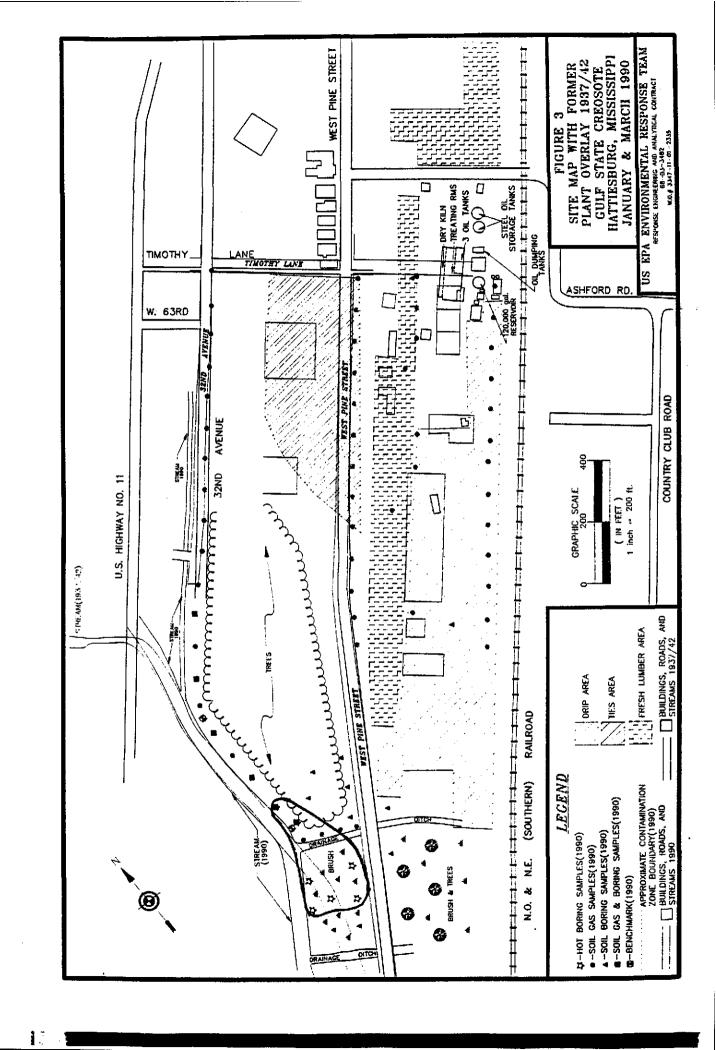
US EPA ENVIRONMENTAL RESPONSE TEAM

RESPONSE ENGINEERING AND ANALYTICAL CONTRACT 68-03-3482

GULF STATES CREOSOTE

Figure 1. Site Location Map





APPENDIX A GULF STATES CREOSOTE TRIP REPORTS APRIL, 1990



REAC SUPPORT ORGANIZATION GSA RARITAN DEPOT WOODBRIDGE AVENUE BUILDING 209, BAY F EDISON, NJ 08837 PHONE: 201-632-9200

DATE:

February 16, 1990

TO:

Harry Compton, US EPA-ERT Work Assignment Manager

FROM:

Martin O'Neill, REAC Task Leader With

THRU:

Craig Moylan, REAC O&A Section Chief WOB/fel

SUBJECT:

GULF STATES CREOSOTE SOIL GAS AND SOIL SAMPLING SURVEY:

WA 3347-11-01-2335 - TRIP REPORT

BACKGROUND

The former Gulf States Creosote Site is located in a commercial area of Hattiesburg, Mississippi. The site was an active wood preserving facility from approximately 1920 to 1960. It is currently owned by the city of Hattiesburg and subleased to several automobile dealerships, car-parts stores, a beverage distributer, a food store, and a furniture store. The process areas and wood drying/drip areas have been regraded, covered with asphalt, and are no longer evident. The former site encompasses approximately 20 acres, and is bordered on the east by railroad tracks, on the west by Gordens Creek, on the south by a drainage swale which feeds into Gordens Creek, and on the north by Timothy Street.

The increase in surface runoff as a result of development and paving in the immediate area of the former site has significantly effected the flow into Gordens Creek. As a result, the U.S. Army Corps. of Engineers (Corps) has been requested to re-channel Gordens Creek. In preparation for the re-channelization project, the Corps requested assistance from the U.S. EPA in characterizing the nature and extent of contamination which may be present in the area as a result of the former Gulf States wood treating operation.

The purpose of the ERT/REAC sampling effort was to identify the extent of contamination by completing a soil gas survey in the area, and to determine the nature of soil and/or groundwater contamination through the installation of well points. In addition, air sampling was planned in an attempt to determine if contaminants associated with the wood treating process, namely polynuclear aromatic hydrocarbons (PAHs) were present in ambient air as a result of soil disturbance activities (i.e., soil borings).

OBSERVATIONS

The ERT/REAC TAGA arrived in Hattiesburg on Saturday, January 20, 1990, at which time TAGA operators Dave Mickunas, Mark Bernick, Joe Gorski, and Gmae Loy commenced with instrument calibration and preparation. Additional ERT/REAC personnel including Harry Compton, Mark Sprenger, Greg Powell, Martin O'Neill, Akos Fekete, and Mark Ellis arrived

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in Hattiesburg on Sunday. ERT/REAC performed a site walk-through, and area familiarization survey on Sunday.

On Monday, January 22, ERT/REAC personnel met with the U.S. EPA On-Scene Coordinator (OSC), Don Rigor, to discuss planned activities and the general sampling/survey approach. Following the meeting, REAC personnel commenced with equipment pick-up and logistical setup at the site. The TAGA was mobilized to the site and began preparing for analysis of tedlar bags in conjunction with the soil gas sampling effort.

On Monday afternoon, ERT/REAC commenced with the soil gas sampling program. A total of four (4) transects were established running north to south, parallel to Pine Street. Transect A was established adjacent to the railroad tracks along the eastern border of the site. Transect C was placed along the eastern curb of Pine Street. Transect B was located approximately equidistant, and between, transect A and C, and transect E was placed along the eastern curb of 32nd Street. Where possible, sampling points were placed at 100 foot intervals along each of the transects. Because of building locations, paved areas and lack of access, the location of sampling stations along transect B varied from the expected straight line. Figure 1 provides the location of soil gas sample stations and soil borings.

The soil gas sampling continued into and was completed on Tuesday, January 23. A total of 49 stations were sampled. Sampling stations were surveyed using the HnU PI 101 photoionization detector, and Foxboro Organic Vapor Analyzer (OVA). In general, the HnU's were not responsive to the contaminants of concern whereas the OVA's were. Readings averaged 2 to 5 units and ranged from 1 to 400 units (at location A-07). A Tedlar bag was collected at only those stations where positive readings were obtained on the OVA. Bag samples were analyzed using the TAGA tandem mass spectrometer (MS/MS). Holding times for soil gas samples were less than 2 hours. The TAGA field report is presented in the Appendix of this trip report.

On Tuesday afternoon, a hand auger team commenced with the soil boring program in the area adjacent to the railroad tracks and along the tree-line west of Pine Street. The purpose of the hand auger points was to gather preliminary information regarding local surficial geology, and provide for a "quick and dirty" screening for the presence of creosote compounds. Another team began gathering and preparing equipment as required for the well point installations planned for Wednesday and Thursday.

A series of background samples were also collected on Tuesday. Two sampling stations were established in up-wind locations, and one station was located down-wind from the site. The background sample results were to be used in evaluating the possible effects of soil disturbance activities on ambient air quality.

Following review of the TAGA soil gas data on Wednesday, ERT in conjunction with the OSC decided that the investigation would focus on the areas adjacent to Gordens Creek. Additionally, it was decided that samples would be collected from each of the soil borings and sent to the ERT/REAC laboratory in Edison, NJ. The TAGA was no longer needed and could be de-mobilized. TAGA personnel spent Wednesday completing analyses and readying the instrument for the trip back to Edison. TAGA personnel departed Hattiesburg on Wednesday and the TAGA departed early on Thursday, January 25.

Preparation of equipment and establishment of a temporary decontamination facility was completed by Wednesday at noon. Two soil boring teams, both equipped with hydraulic power augers (Mini-Beaver), started sampling shortly after noon. Fourteen soil borings were drilled

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prior to a torrential rain-storm that arrived at approximately 1500. This severe downpour, together with the 2+ inches of rain that the Hattiesburg area received during Tuesday night, made working conditions challenging and sampling of distinct depths difficult. Shortly after the rain commenced on Wednesday, the entire soil column at 1 to 2 feet below ground surface became saturated. ERT/REAC sampling efforts were abandoned at 1530, and on-site operations ceased. Equipment was decontaminated and some of the unnecessary pieces were shipped back to REAC in Edison, NJ. A total of ten (10) soil samples were also shipped to the ERT/REAC laboratory.

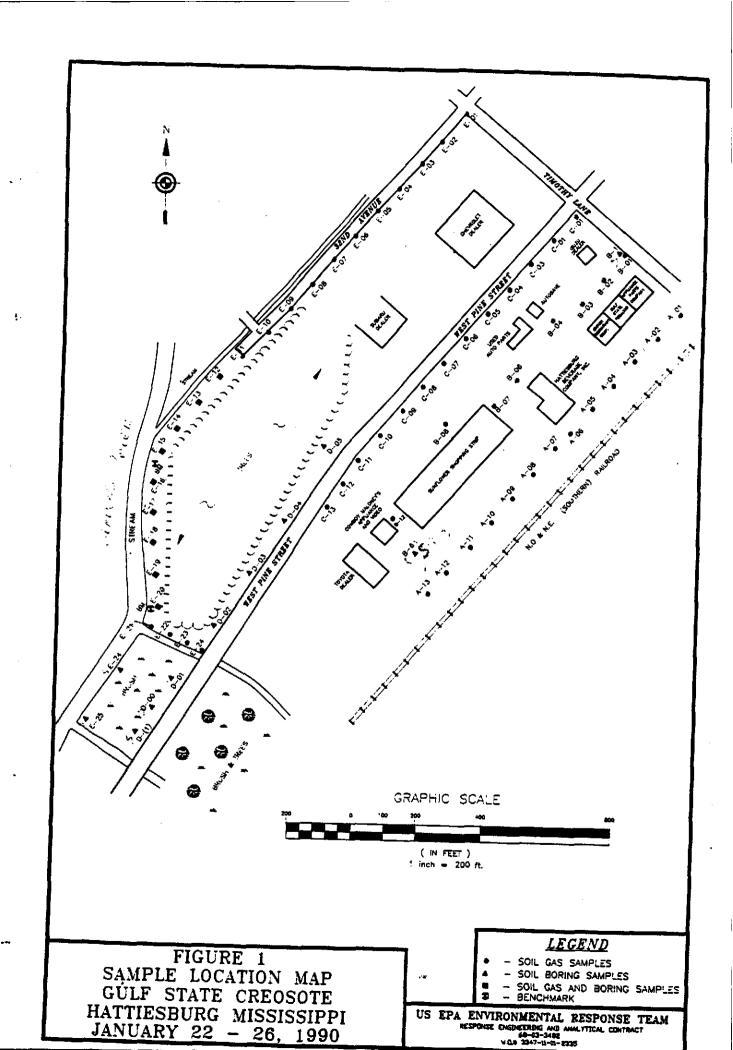
The rain continued throughout the night on Wednesday and into Thursday. Since characterization of distinct subsurface soil horizons would be compromised by the extremely elevated water table, ERT, in conjunction with the OSC, decided to postpone additional boring samples until dryer conditions prevailed. In addition, because of the rain and extreme wet conditions, the air sampling/monitoring program had to be suspended.

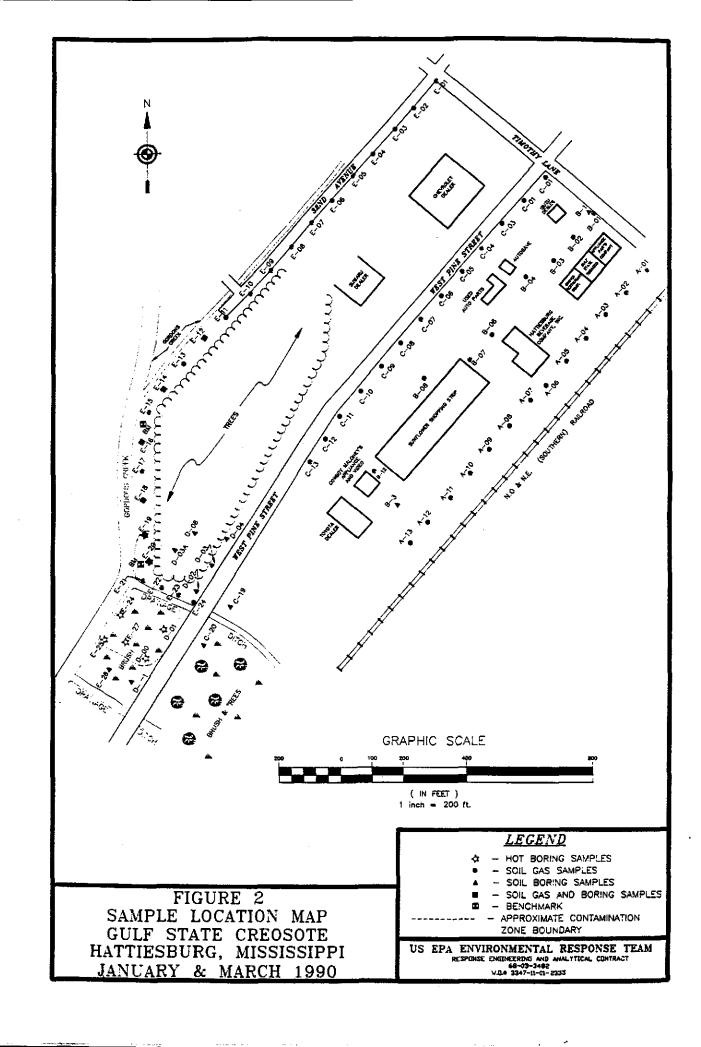
On Thursday, a team consisting of M. Ellis, G. Prince, G. Powell, and D. Rigor (OSC), commenced with the topographic survey of soil gas sampling locations, soil borings and significant features. The rest of the team began packaging the equipment for shipment back to Edison, and returning the rental equipment in preparation of site demobilization. A. Fekete, M. O'Neill, W. Batz, H. Compton, and M. Sprenger returned to NJ on Thursday night. The topographic survey was completed on Thursday which allowed the remaining crew to return to NJ on Friday.

FUTURE ACTIVITIES

The ten (10) soil samples and three (3) ambient air samples are currently being analyzed for PAH compounds by the S&A section of REAC. Draft analytical data are expected to be available in late February. Final analytical data are to be delivered to ERT on March 1, 1990.

REAC is currently preparing ACAD presentations of the land survey, soil gas contours and the extent of the former site operations. Preparation and delivery of a final summary report is planned for April 1, 1990.







REAC SUPPORT ORGANIZATION GSA RARITAN DEPOT WOODBRIDGE AVENUE BUILDING 209, BAY F EDISON, NJ 08837 PHONE, 201-632-9200

DATE:

March 30, 1990

TO:

Harry Compton, Work Assignment Manager

FROM:

Mark Ellis, REAC Geologist Me

Martin O'Neill, REAC Task Leader

THRU:

Craig Moylan, O&A Section Chief truchille Cin

SUBJECT:

GULF STATE CREOSOTE SOIL SAMPLING SURVEY:

W.A. # 3347-11-01-2335 - TRIP REPORT

BACKGROUND

The former Gulf States Creosote Site is located in a commercial area of Hattiesburg, Mississippi. The site was an active wood preserving facility which operated from approximately 1920 to 1960. It is currently owned by the city of Hattiesburg and subleased to several automobile dealerships, car-parts stores, a beverage distributer, a food store, and a furniture store. The process areas and wood drying/drip areas have been regraded, covered with asphalt, and are no longer evident. The former site encompasses approximately 20 acres, and is bordered on the east by railroad tracks, on the west by Gordons Creek, on the south by a drainage swale which feeds into Gordons creek, and on the north by Timothy Street.

The increase in surface runoff as a result of development and paving in the immediate area of the former site has significantly effected the flow into Gordons Creek. As a result, the U.S. Army Corps of Engineers (Corps) has been requested to rechannel Gordons Creek. In preparation for the rechannelization project, the Corps requested assistance from the US EPA in characterizing the nature and extent of contamination, which may be present in the area as a result of former wood treating operations at the Gulf State Creosote site.

ERT/REAC completed a soil gas survey and a series of preliminary soil borings in January, 1990. Because of an unexpected amount of rainfall, completion of the soil sampling effort had to be postponed.

The purpose of this ERT/REAC sampling event was to complete the subsurface soil investigations in the area of the former Gulf States Creosote Plant. Samples were to be collected at borings which appeared to be contaminated and analyzed for Poly Aromatic Hydrocarbon (PAH) compounds by the S&A Section of REAC.

OBSERVATIONS

The ERT/REAC team consisting of George Prince and Mark Ellis arrived in Hattiesburg, MS at approximately 1430 on Monday, March 19, 1990. A meeting was held with the On-Scene Coordinator (OSC), Don Rigger, Richard Ball from the state of Mississippi, Department of Environmental Quality.

rd/ELLIS/TR-2335

Greg Powell of ERT Cincinnati, and JoAnna Cole from Region IV TAT to discuss the planned scope of work.

Following the meeting, the sampling teams met on site. One team began soil borings using the "Little Beaver" two-man power auger, while the other team surveyed new sampling locations from existing transects. Three borings were completed at locations D-1, E-26 and E-27, as depicted on Figure 1, Sample Location Map. Field data sheets are included in Appendix A. They include a rough geologic log and final depths for each boring. Boring locations E-26 and E-27 were measured 100 feet perpendicular from transect D. Boring E-27 had a distinct creosote odor, and a sample was collected using a stainless steel bucket auger. Site activities were completed at approximately 1745.

On Tuesday morning, the sampling teams met on site at 0730 and commenced boring at location E-25. The equipment was decontaminated after the completion of this boring using a high-pressure steam cleaner. A boring was then advanced at location E-24. Samples were collected at each of these locations from the bottom of the boreholes, or approximately nine feet.

A series of borings were drilled along transect D (D-02, D-03, D-03A, D-04, D-06). The first two locations. D-02 and D-03, were abandoned after the auger flights met refusal between three and six feet, due to an unknown thickness of fill material. Two samples were collected at location D-03A. One was representative of the wet sands just above a thick white clay layer and the other was a sample of the clay material. The clay layer was sampled to determine if the downward movement of the creosote is being retarded by the clay layer. Borings D-04 and D-06 were drilled to 10 feet and 14 feet, respectively. Both of these holes appeared to be clean (no odor), so no samples were collected.

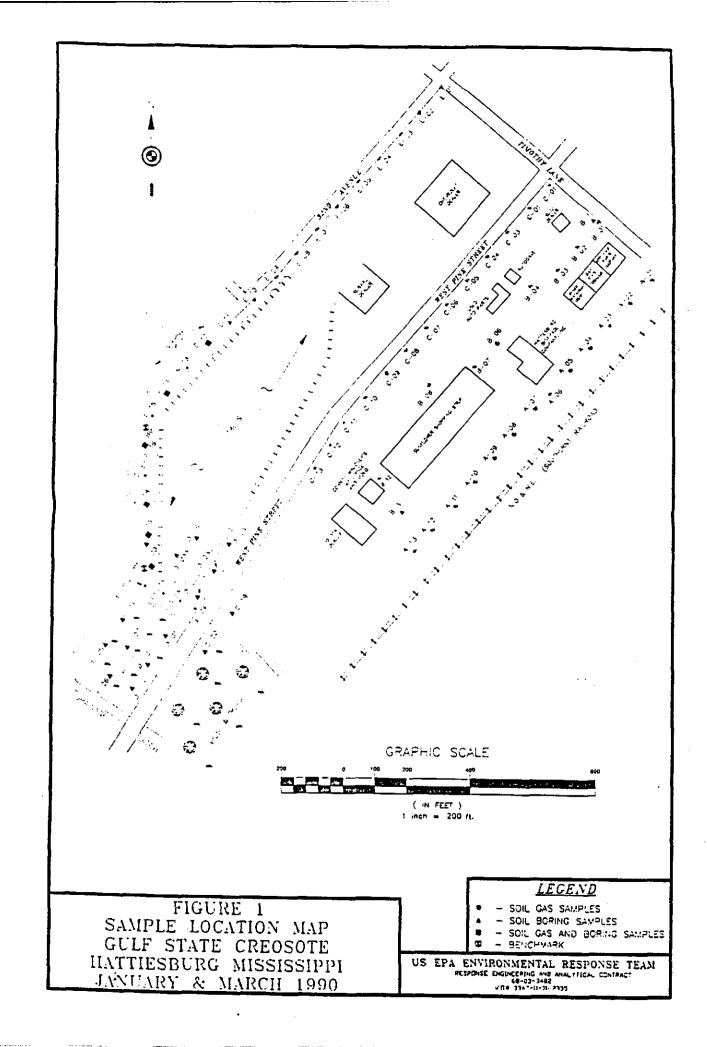
Borings were completed at locations E-19, C-19 and C-20. Two samples were collected from boring E-19, again, one representing the sands above the clay layer and one of the clay itself. Samples were also collected at C-19 and C-20. These samples were collected in order to verify the presence/absence of any creosote compounds in the area east of West Pine Street.

After visual observation, it was decided that it was not necessary to drill or sample areas west of Gordons Creek. The crew mobilized to the decontamination area, cleaned all of the equipment, and began packaging it for shipment back to Edison. Site activities were completed at 1645 and the equipment was shipped back to Edison, NJ via Federal Express.

FUTURE ACTIVITIES

REAC is currently preparing to perform some preliminary treatability studies. Shaker tests will be performed to determine the likelihood of using microbial treatment as a means of remediation at the site.

The ACAD section at REAC is presently updating the sample location map which will be included in a final summary report.



APPENDIX B

TAGA SOIL GAS ANALYSIS GULF STATES CREOSOTE APRIL, 1990

TAGA FIELD REPORT

Prepared by Roy F. Weston, Inc.

Gulf States Creosote Site Hattiesburg, Mississippi

February 13, 1990

EPA Work Assignment No. 1-335 Project No. 3347-11-01-2335 EPA Contract No. 68-03-3482

> Analysis by: Mark Bernick Gmae Loy Dave Mickunas

Prepared by: Mark Bernick Gmae Loy

Reviewed by: Dave Mickunas

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APPENDIX

A	Gas	Cylinder	Certifications
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B Methane Analysis Data

C Calibration Curves For Sampling Periods

INTRODUCTION

This report presents the results of EPA Work Assignment No. 0-335, Weston Order No. 3347-11-01-2335, EPA Contract No. 68-03-3482.

The Response Engineering and Analytical Contract (REAC) was tasked by the USEPA/ERT to mobilize the U.S. EPA Trace Atmospheric Gas Analyzer (TAGA) to Hattiesburg, Mississippi to analyze soil gas samples from the Gulf States Creosote Plant Site. The analysis commenced on January 22, 1990, and concluded one day earlier than anticipated on January 23, 1990, due to the meteorological forecast calling for rain that would impair soil gas sampling.

The goal of the investigation was to analyze soil gas and soil head space samples for target compounds to identify creosote, gas and oil contaminated soils. Naphthalene was used as the target compound for identification of creosote contaminated soil; benzene, toluene, and xylene were monitored as indicators of gasoline and/or oil contaminated soil.

TABLE 1

TARGET COMPOUND LIST

benzene naphthalene toluene xylene

PROCEDURE

TAGA PROCEDURE

The following operating procedures were performed during each analysis day using the TAGA 6000E:

- (1) The first and third quandrupoles were scanned for 15 minutes each; this readied the instrument electronically.
- (2) A gas mixture containing trichloroethene and tetrachloroethene was introduced by a mass flow controller into the sample air stream to optimize the first and third quadrupoles for sensitivity and mass assignment.
- (3) The instrument was calibrated before and after each sampling period to generate target compound response factors ion counts per second/parts per billion by volume (ICPS/ppbv).

Gas Calibration System

The gas calibration system consisted of a regulated gas cylinder with a mass flow controller. This calibration system was used to generate analytes' response factors (ICPS/ppbv), which were then used to quantify trace components in soil gas samples. The following is a list of target compounds, for which the instrument was calibrated, using the gas cylinder method:

benzene toluene xylene

A gas cylinder, containing a known mixture of target compounds as certified by Scott Specialty Gases (see Appendix A), was regulated at preset flow rates and diluted with ambient air. This dilution of the gas cylinder gave known analyte concentrations. Software, as described in the Plessey Interface Manual, utilized the analyte's cylinder concentration, the gas flow rate, the air sampling flow rate, and the atmospheric pressure to calculate the analyte's response factors (RFs). These response factors were obtained for the ion pairs of each compound of interest in the cylinder:

Cylinder ALM-001166

benzene toluene xylene

Tedlar Bag Calibration System

The Tedlar bag calibration system consisted of Tedlar bags, a 500 milliliter (ml) gas tight syringe, 5 and 60 ml sterile Becton-Dickinson B-D type syringes, and a temperature probe. The 5 and 60 ml syringes were filled with approximately 5 grams (gm) of the analyte and allowed to equilibrate for 15 minutes at room temperature. The temperature of the syringes was measured, and the concentration of the analyte in the gas phase of the syringe was calculated using the vapor pressure of the analyte, atmospheric pressure and the temperature of the syringe. The analyte was then diluted in the Tedlar bag to the target calibration concentration using ambient air and a volume of the syringe gas phase. The Tedlar bag was then attached directly to the TAGA sampling line and analyzed undiluted for the analyte's ion pairs.

A linear regression was run using the analyte's known Tedlar bag concentration and the respective ion pair's ion counts. The slope of the regression was equal to the ion pair's response factor. The target compound calibrated using the Tedlar bag calibration method was naphthalene.

Tedlar Bag Analysis

The Tedlar bags were received and stored inside an opaque plastic bag to prevent analyte degradation from by light. The bag sample number, location, sampling date and time were logged into the Taga computer prior to analysis. The Tedlar bag was attached directly to the TAGA sampling line and analyzed undiluted for the target compounds. Once the sample's target ion response equilibrated, a two minute file was collected and saved for each bag analyzed.

After the intensity data was downloaded off the hard disk; it was processed as described in the PC-Plessey Interface Manual. The PC-Plessey calculates an average concentration for each target compound. This concentration is arrived at by taking the signal intensity of selected parent/daughter ion pairs of a compound, dividing them by the appropriate response factor, and averaging the resultant concentrations of the ion pairs.

Soil Head Space Analysis

Soil samples were received and stored at room temperature in 40 ml glass Volatile Organic Analysis (VOA) bottles. These bottles contained about 35 ml of soil with about 5 ml of head space remaining. The samples were allowed to equilibrate at room temperature for one hour prior to analysis. The sample head space was bleed directly into the TAGA sampling line and analyzed undiluted for the target compounds. Once the sample's target ion response equilibrated, a two minute file was collected and saved for each sample analyzed.

After the intensity data was downloaded off the hard disk, it was processed as described in the PC-Plessey Interface Manual. Basically, this calculates an average concentration for each target compound. This concentration is arrived at by taking the signal intensity of selected parent/daughter ion pairs of a compound, dividing them by the appropriate response factor, and averaging the resultant concentrations of the ion pairs. The results of the soil head space target compound analysis are reported in ppbv.

CENTURY ORGANIC VAPOR ANALYZER PROCEDURE

Methane Analysis

A Model Century Organic Vapor Analyzer 128 Gas Chromatograph (OVA 128GC) was set up in the "GC Mode", as described in the instrument manual, using a 12 inch Poropak T 60/80 Mesh Stainless Steel OVA column (PT-12 column). An injection port was added to the front end of the column to allow gas-tight syringe injections of Tedlar bag samples, and a strip chart recorder was attached to the OVA 128GC to record the signal response of the flame ionization detector (FID).

A gas cylinder containing 889 parts per million by volume (ppmv) of methane as certified by Scott Specialty Gases (see Appendix A) was used as the methane calibration standard. Injections of 5 and 10 microliters (ul) of the calibration standard were recorded with the OVA 128GC CALIBRATE Switch set to X10. This gave a methane peak response equivalent to 45.0 ppmv and 89.9 ppmv methane respectively, for a sample injection of 100 ul.

A sample injection of 100 ul was used for analyzing the Tedlar bags for methane. The sample injection size was reduced to 10 or 5 ul if the signal response for a sample went off scale. The results of the methane analysis are present in ppmv methane.

RESULTS AND DISCUSSION

The TAGA 6000E laboratory performed both Tedlar bag and soil head space analyses for target compounds of samples from the Gulf States Creosote Site in Hattiesburg, Mississippi. The study was designed to monitor soil gas and head space using naphthalene as an indicator of creosote contamination. Naphthalene was chosen as an indicator as it was found to be present in a creosote contaminated soil sample from this site analyzed prior to mobilization. Additionally, benzene, toluene, and xylene were used as indicators of gas and/or oil contaminated soil.

The results of these analyses are broken down into five sampling periods which correspond to five calibration periods each having their own set of response factors, detection and quantitation limits, and maximum percent deviations. These are presented in the Quality Assurance/Quality Control section of this report. These sampling periods were:

Sampling Period I	January 22, 1990	14:30 - 17:00
Sampling Period II	January 22, 1990	17:00 - 19:14
Sampling Period III	January 22, 1990	19:14 - 21:00
Sampling Period IV	January 23, 1990	10:18 - 15:51
Sampling Period V	January 23, 1990	15:51 - 19:26

The results of this study are broken down by sampling period and are presented in Tables 2 - 6 respectively. These tables list the TAGA file, target compound results, Tedlar bag identification number (ID NO.), sampling date and time, and sample location. The tables present results of Tedlar bag soil gas analysis except TAGA file numbers AHT063, 64, 68 - 71, soil head space analysis results.

Additionally, selected Tedlar bags were analyzed for methane. The results of this analysis are presented in Table 7, including a list of the Tedlar bag ID No. and the methane results.

TABLE 2

SAMPLING PERIOD I TAGA RESULTS

CONCENTRATIONS IN PPBV

TAGA					•	SAMPL	ING	SAMPLE
FILE	BEN	TOL	XYL	NAP	ID NO.	DATE	TIME	LOCATION
AHT007	DL=5	DL=15	DL=10	DL=35	SG01492	1-22-90	14:40	C03
800THA	DL=5	DL=15	DL=10	DL=35	SG01521	1-22-90	14:32	AD1
AHT009	DL=5	DL=15	DL=10	DL=35	SG01497	1-22-90	15:40	C09
AHT010	10-J	17 - J	14 <i>-J</i>	DL=35	SG01491	1-22-90	14:20	C01
AHT011	DL=5	DL=15	DL=10	DL=35	SG01522	1-22-90	14:49	A02
AHT012	DL=5	DL=15	DL=10	DL=35	SG01526	1-22-90	15:28	A06
AHT013	DL=5	DL=15	DL=10	DL=35	SG01530	1-22-90	16:10	A010
AHT014	12-J	19 - J	16-J	DL=35	SG01493	1-22-90	14:55	C05
AHT015	DL=5	DL=15	DL=10	DL=35	SG01523	1-22-90	14:58	A03
AHT016	DL=5	DL=15	DL=10	DL=35	SG01524	1-22-90	15:10	A04

DL = DETECTION LIMIT

J - VALUE ABOVE DETECTION LIMIT BUT BELOW QUANTITATION LIMIT

TABLE 3

SAMPLING PERIOD II TAGA RESULTS

CONCENTRATIONS IN PPBV

TAGA FILE	BEN	TOL	XYL	NAP	ID NO.	SAMPL: DATE	ING SAMPLE TIME LOCATION
AHT019	11 - J	DL=20	DL=12	DL=40	SG01494	1-22-90	15:00 C06
AHT020	DL=6	DL=20	DL=12	DL=40	SG01525	1-22-90	15:19 A05
AHT021	DL=6	DL=20	DL=12	DL=40	SG01527	1-22-90	15:43 A07
AHT022	DL=6	DL=20	DL=12	DL=40	SG01528	1-22-90	15:53 AO8
AHT023	DL=6	DL=20	DL=12	DL=40	SG01529	1-22-90	16:01 A09
AHT024	31 - J	31 -J	18-J	DL=40	SG01498	1-22-90	16:10 C-11
AHT025	DL=6	DL=20	DL=12	DL=40	SG01500	1-22-90	16:20 CO13
AHT026	DL=6	DL=20	DL=12	DL=40	SG00611	1-22-90	16:25 A011
AHT027	DL=6	DL=20	DL=12	DL=40	SG00612	1-22-90	16:33 A012
AHT028	DL=6	DL=20	DL=12	DL=40	SG01501	1-22-90	16:42 CO15

DL = DETECTION LIMIT

J = VALUE ABOVE DETECTION LIMIT BUT BELOW QUANTITATION LIMIT

TABLE 4

SAMPLING PERIOD III TAGA RESULTS

CONCENTRATIONS IN PPBV

TAGA						SAMPL	ING	SAMPLE
FILE	BEN	TOL	XXL	NAP	ID NO.	DATE	TIME	LOCATION
AHT030	DL=8	DL=25	DL=16	DL=38	SG01531	1-22-90	16:4	4
AHT031	DL=8	DL=25	DL=16	DL=38		1-22-90		=
AHT032	DL=8	30-J	DL=16	DL=38	SG00613	1-22-90	16:0	6 E015
AHT033	DL=8	DL=25	DL=16	DL=38	SG00614	1-22-90	17:1	5 E014
AHT034	DL=8	DL=25	DL-16	DL=38	SG01534	1-22-90	17:14	4 E04
AHT035	DL=8	DL=25	DL=16	DL=38	SG01533	1-22-90	17:1	0· E03
AHT036	DL=8	DL=25	DL=16	DL=38	SG01535	1-22-90	17:2	5 E05
AHT037	DL=8	DL=25	DL=16	DL=38	SG01536	1-22-90	17:3	0 E06

DL = DETECTION LIMIT

J = VALUE ABOVE DETECTION LIMIT BUT BELOW QUANTITATION LIMIT

TABLE 5

SAMPLING PERIOD IV TAGA RESULTS

CONCENTRATIONS IN PPBV

					*	SAMPL	TMC	SAMPLE
TAGA								
FILE	BEN	TOL	XYL	NAP	ID NO.	DATE	TIME	LOCATION
AHT043	7-J	40-J	30-J	82-J	SG01538	1-23-90	09:07	7 B01
AHT044	フーJ	30-J	24-J	57 - J	SG01502	1-23-90	09:12	2 E07
AHT045	9-J	51 - J	41 - J	122 - J	SG01540	1-23-90	09:30	B03
AHT046	7 - J	36 - J	27 - J	73ーゴ	SG01505	1-23-90	09:45	E10
AHT047	7 - J	49 - J	25-J	60 - J	SG01503	1-23-90	09:30	E09
AHT049	DL=6	30 − J	24-J	75 - J	SG01504	1-23-90	09:39	E10
AHT050	10 - J	31 - J	23-J	75 - J	SG01506	1-23-90	09:53	E11
AHT051	DL=6	28 - J	20-J	53 - J	SG00621	1-23-90	10:22	B06
AHT052	16 - J	33 - J	22 - J	50-J	SG00622	1-23-90	10:35	5 B07
AHT055	14-J	40-J	26-J	53-J	SG00623	1-23-90	10:47	7 B012
AHT056	14-J	661	42-J	80~J	SG01508	1-23-90	10:5	7 E23
AHT057	DL=6	41-J	15-J	37 - J	SG01509	1-23-90	11:10	5 E22
AHT058	7 - J	37 - J	23 - J	52 - J	SG01510	1-23-90	11:4	5 F1
AHT059	10 - J	DL=25	DL=14	DL=29	SG00624	1-23-90		B09
AHT060	DL=6	33 - J	22 - J	47 - J	SG01507	1-23-90		E24
AHT062	18-J	48-J	111	51 - J	SG00625	1-23-90	12:0	B B08
AHT063	11-J	51 - J	102	938	BORE HO	LE 9' DE	EP	A01
AHT064	22	131	457	9709	E05321	SOIL H	EAD S	PACE
			·					

DL = DETECTION LIMIT

J = VALUE ABOVE DETECTION LIMIT BUT BELOW QUANTITATION LIMIT

TABLE 6

SAMPLING PERIOD V TAGA RESULTS CONCENTRATIONS IN PPBV

TAGA FILE	BEN	TOL	XYL	NAP	ID NO.	TITLE	·
AHT068 AHT069 AHT070 AHT071	24 27 23 24	35-J 37-J 34-J 32-J	41-J 46-J 39-J 34-J	DL=38 DL=38 DL=38 DL=38	D05309 K05322	SOIL HEAD SOIL HEAD SOIL HEAD	SPACE SPACE

DL = DETECTION LIMIT

J - VALUE ABOVE DETECTION LIMIT BUT BELOW QUANTITATION LIMIT

TABLE 7

METHANE ANALYSIS RESULTS CONCENTRATION IN PPMV

DATE - January 23, 1990 SAMPLE ID NO.METHANE

SG01538 ND SG01540 ND SG01505 ND SG01506 ND SG01502 ND SG01503 ND SG01503 ND SG00622 ND SG00621 800 SG00623 ND SG01507 ND SG01507 ND SG01509 ND SG01510 ND SG01510 ND SG01508 ND SG00625 ND SG00624 ND

ND = NOT DETECTED

QUALITY ASSURANCE/QUALITY CONTROL

The compound parent/daughter ion pairs used for ion profile quantitation and detection are listed below:

Compound	ID	Parent Mass/Daughter Mass
benzene	BNZ	78/39
benzene	BNZ	78/52
toluene	TOL	92/39
toluene	TOL	92/51
xylene	XYL	106/65
xylene	XYL	106/91
naphthalene	NAP	128/78
naphthalene	NAP	128/128

Additional ion pairs had been calibrated and monitored for, but due to background interferences or insensitivity, they were not used.

Calculations for the Summary of Actual and Intermediate Response Factors for the Target Compounds' Ion Pairs During the Sampling Period

Response factors (RF) were generated from the final and initial calibration events, as described 1 the procedure. Tables 8 - 12 contain the RFs in units of ion counts per second/parts per billion by volume (ICPS/ppbv). The actual RFs are used to calculate the intermediate RFs, which are used to calculate the concentrations reported in the results. The following is a list of the target compounds and the identification "ID" used in Tables 8 - 12.

ID	COMPOUND
BNZ	benzene
TOL	toluene
XYL	xylene
NAP	naphthalene

The following equation was used to calculate the intermediate response factors (IRF) found in Tables 8 - 12.

$$IRF = 2 (RF1 \times RF2)$$

 $(RF1 + RF2)$

where:

IRF = intermediate response factor (ICPS/ppbv)

RF1 = the response factor for an ion pair measured during the initial calibration event (ICPS/ppbv)

RF2 = the response factor for the same ion pair measured during the final calibration event (ICPS/ppbv)

For example, the entry for the 78/39 ion pair of benzene from Table 8 is:

RF1 = 31.14 (ICPS/ppbv) RF2 = 37.13 (ICPS/ppbv)

and then

$$\frac{2(31.14 \times 37.13)}{(31.14 + 37.13)} = \frac{2312.45}{68.27} = 33.87 \text{ ICPS/ppbv}$$

TABLE 8

THE SUMMARY OF ACTUAL AND INTERMEDIATE RESPONSE FACTORS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD I

	CALIBRATION TIME	01/22/90 14:30	01/22/90 17:00		
ID	PM/DM	RESPONSE FACTOR	RESPONSE FACTOR	INTERMEDIATE RESPONSE FACTOR	
BEN	78/ 39	31.14	37.13	33.87	
BEN	78/ 52	44.63	53.21	48.54	
TOL	92/ 39	6.23	8.19	7.08	
TOL	92/ 51	17.11	22.49	19.43	
XYL*	106/ 39	2.59	2.50	2.54	
XYL*	106/ 51	2.49	2.42	2.45	
XYL	106/ 65	9.87	9.56	9.71	
XYL	106/ 91	65.90	63.84	64.85	
NAP	128/ 78	2.12	2.29	2.20	
NAP	128/128	106.56	115.42	110.81	

ID = Identification Code

PM = Parent Mass

DM = Daughter Mass

^{*} Ion Pairs Not Used in Quantitation.

TABLE 9

THE SUMMARY OF ACTUAL AND INTERMEDIATE RESPONSE FACTORS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD II

	CALIBRATION	, ,	01/22/90 19:14		
	TIME	14:30	19:14	INTERMEDIATE	
.+		RESPONSE	RESPONSE	RESPONSE	
ID	PM/DM	FACTOR	FACTOR	FACTOR	
BEN	78/ 39	31.14	28.63	29.83	
BEN	78/ 52	44.63	41.02	42.75	
TOL	92/ 39	6.23	5.83	6.03	
TOL	92/ 51	17.11	16.02	16.55	
XYL*	106/ 39	2.59	1.78	2.11	
XYL*	106/ 51	2.49	1.72	2.04	
XYL	106/ 65	9.87	6.81	8.06	
XYL	106/ 91	65.90	45.50	53.83	
NAP	128/ 78	2.12	1.85	1.98	
NAP	128/128	106.56	93.28	99.48	

ID = Identification Code

PM = Parent Mass DM = Daughter Mass

* Ion Pairs Not Used in Quantitation.

TABLE 10

THE SUMMARY OF ACTUAL AND INTERMEDIATE RESPONSE FACTORS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD III

	CALIBRATION TIME	01/22/90 14:30	01/22/90 21:00		
ID	PM/DM	RESPONSE FACTOR	RESPONSE FACTOR	INTERMEDIATE RESPONSE FACTOR	
	Elly Del	FACION	FACIOR	PACIOR	
BEN	78/ 39	31.14	23.24	26.61	
BEN	78/ 52	44.63	33.56	38.31	
TOL	92/ 39	6.23	4.09	4.94	
TOL	92/ 51	17.11	10.07	12.68	
XYL*	106/ 39	2.59	1.41	1.82	
XYL*	106/ 51	2.49	1.16	1.58	
XYL	106/ 65	9.87	4.82	6.48	
XYL	106/ 91	65.90	34.10	44.94	
NAP	128/ 78	2.12	2.10	2.11	
NAP	128/128	106.56	94.70	100.28	

ID = Identification Code

PM = Parent Mass

DM = Daughter Mass

^{*} Ion Pairs Not Used in Quantitation.

TABLE 11

THE SUMMARY OF ACTUAL AND INTERMEDIATE RESPONSE FACTORS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD IV

	CALIBRATION	01/23/ 9 0	01/23/90	
	TIME	10:18	15:51	
				INTERMEDIATE
		RESPONSE	RESPONSE	RESPONSE
ID	PM/DM	FACTOR	FACTOR	FACTOR
BEN	78/ 39	30.30	24.09	26.840
BEN	78/ 52	42.72	31.25	36.094
TOL	92/ 39	6.10	3.38	4.349
TOL	92/ 51	16.34	11.72	13.653
XYL*	106/ 39	3.14	1.78	2.268
XYL*	106/ 51	2.75	0.74	1.165
XYL	106/ 65	11.50	5.70	7.621
XXT	106/ 91	73.31	33.37	45.863
NAP	128/ 78	4.54	2.23	2.991
NAP	128/128	222.17	109.31	146.527

ID = Identification Code

PM = Parent Mass DM = Daughter Mass

^{*} Ion Pairs Not Used in Quantitation.

TABLE 12

THE SUMMARY OF ACTUAL AND INTERMEDIATE RESPONSE FACTORS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD V

	CALIBRATION TIME	01/23/90 10:18	01/23/90 19:26			
ID	PM/DM	RESPONSE FACTOR	RESPONSE FACTOR	INTERMEDIATE RESPONSE FACTOR		
BEN	78/ 39	30.30	21.33	25.04		
BEN	78/ 52	42.72	31.01	35.93		
TOL	92/ 39	6.10	3.03	4.05		
TOL	92/ 51	16.34	9.72	12.19		
XYL*	106/ 39	3.14	1.13	1.66		
XYL*	106/ 51	2.75	1.08	1.55		
XYL	106/ 65	11.50	4.92	6.89		
XYL	106/ 91	73.31	31.16	43.73		
NAP	128/ 78	4.54	1.65	2.42		
NAP	128/128	222.17	49.50	80.96		

ID = Identification Code

PM = Parent Mass
DM = Daughter Mass

^{*} Ion Pairs Not Used in Quantitation.

Calculations for the Summary of the Detection and Quantitation Concentration Limits for the Target Compounds' Ion Pairs During the Sampling Periods

The ion pairs' detection concentration limits (DL) and quantitation concentration limits (QL) were generated from the ion pairs' intensity of the standard deviation (SD) of the measurement of an ambient air Tedlar bag; as well as its intermediate response factor. Tables 13 - 17 contain these IRFs that are in units of ion counts per second/part per billion by volume (ICPS/ppbv).

The following equation was used to calculate the detection concentration limits found in Table 13 to 17.

$$DL = \underbrace{3 \times SD}_{IRF}$$

where:

DL = detection limit concentration for an ion pair (ppbv)

SD = Standard deviation of the ion intensity for an ion pair of the measurement of an ambient air Tedlar bag (ICPS)

IRF = intermediate response factor for an ion pair (ICPS/ppbv)

For example, the entry for the 78/39 ion pair of benzene from Table 13 is:

SD = 59 ICPS

IRF = 33.87 ICPS/ppbv

$$DL = 3 \times 59 = 5.2 \text{ ppbv}$$

33.87

The following equation was used to calculated the quantitation limit concentrations found in Table 13 to 17:

$$QL = 10 X SD$$
IRF

where:

QL = quantitation limit concentration for an ion pair (ppbv)

SD = ion intensity for an ion pair of the measurement of an ambient air Tediar bag (ICPS)

IRF = intermediate response factor for an ion pair (ICPS/ppbv)

For example, the entry for the 78/39 ion pair of benzene from Table 13 is:

SD = 59 ICPS

IRF = 33.87 ICPS/ppbv

$$QL = \frac{10 \times 59}{33.87} = 17.4 \text{ ppbv}$$

TABLE 13

THE SUMMARY OF DETECTION AND QUANTITIATION CONCENTRATION LIMITS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD [

ID	PM/DM	IRF (ICPS)	ERROR BAR	DL (ICPS)	QL (ICPS)	(ppbv)	(bbpa)	INTENSITY (ICPS)	SD (ICPS)
BEN	78/ 39	33.87	0.088	177	590	5.2	17.4	• 71	59
BEN	78/ 52	48.54	0.088	162	540	3.3	11.1	92	54
TOL	92/ 39	7.08	0.136	150	500	21.2	70.7	42	50
TOL	92/ 51	19.43	0.136	153	510	7.9	26.2	71	51
XYL*	106/ 39	2.54	0.016	138	460	54.2	180.8	29	46
XYL*	106/ 51	2.45	0.016	153	510	62.4	207.8	58	51
XYL	106/ 65	9.71	0.016	156	520	16.1	53.5	48	52
XYL	106/ 91	64.85	0.016	228	760	3.5	11.7	292	76
NAP	128/ 78	2.20	0.040	141	470	64.0	213.5	32	47
NAP	128/128	110.81	0.040	630	2100	5.7	19.0	1379	210

ID = Identification Code

PM = Parent Mass

DM = Daughter Mass

IRF = Intermediate Response Factors

DL = Detection Limit

QL = Quantitation Limit

SD = Standard Deviation

^{*} Ion Pairs Not Used in Quantitation

TABLE 14

THE SUMMARY OF DETECTION AND QUANITATION CONCENTRATION LIMITS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD II

ID	PM/DM	IRF (ICPS)	ERROR BAR	DL (ICPS)	QL (ICPS)	(ppbv)	(bbpv)	INTENSITY (ICPS)	SD (ICPS)
BEN	78/ 39	29.83	0.042	210	700	7.0	23.5	131	70
BEN	78/ 52	42.75	0.042	204	680	4.8	15.9	195	68
TOL	92/ 39	6.03	0.033	177	590	29.4	97.9	61	59
TOL	92/ 51	16.55	0.033	171	570	10.3	34.4	125	57
XYL*	106/ 39	2.11	0.183	150	500	71.0	236.8	46	50
XYL*	106/ 51	2.04	0.183	174	580	85.4	284.8	63	58
XYL	106/ 65	8.06	0.183	147	490	18.2	60.8	103	49
XYL	106/ 91	53.83	0.183	279	930	5.2	17.3	439	93
NAP	128/ 78	1.98	0.066	147	490	74.4	247.9	38	49
NAP	128/128	99.48	0.066	591	1970	5.9	19.8	1561	197

ID = Identification Code

PM = Parent Mass
DM = Daughter Mass

IRF = Intermediate Response Factors

DL = Detection Limit
QL = Quantitation Limit
SD = Standard Deviation

* Ion Pairs Not Used in Quantitation

TABLE 15

THE SUMMARY OF DETECTION AND QUANITATION CONCENTRATION LIMITS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD III

ID	PM/DM	IRF (ICPS)	ERROR BAR	DL (ICPS) (QL ICPS)	(ppbv)	(ppbv)	INTENSITY (ICPS)	SD (ICPS)
BEN	78/ 39	26.61	0.145	207	690	7.8	25.9	205	69
BEN	78/ 52	38.31	0.142	243	810	6.3	21.1	291	81
TOL	92/ 39	4.94	0.207	171	570	34.6	115.4	94	57
TOL	92/ 51	12.68	0.259	189	630	14.9	49.7	191	63
XYL*	106/ 39	1.82	0.294	171	570	93.7	312.5	59	57
XYL*	106/ 51	1.58	0.365	165	550	104.3	347.6	65	55
XYL	106/ 65	6.48	0.344	147	490	22.7	75.7	120	49
XXT	106/ 91	44.94	0.318	345	1150	7.7	25.6	677	115
NAP	128/ 78	2.11	0.004	147	490	69.7	232.4	39	49
NAP	128/128	100.28	0.059	618	2060	6.2	20.5	1696	206

ID = Identification Code

PM = Parent Mass

DM = Daughter Mass

IRF = Intermediate Response Factors

DL = Detection Limit

QL = Quantitation Limit

SD = Standard Deviation

^{*} Ion Pairs Not Used in Quantitation

TABLE 16

THE SUMMARY OF DETECTION AND QUANITATION CONCENTRATION LIMITS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD IV

ID	PM/DM	IRF (ICPS)	ERROR BAR	DL (ICPS)	QL (ICPS)	(ppbv)	(ppbv)	INTENSITY (ICPS)	SD (ICPS)
BEN	78/ 39	26.84	0.114	168	560	6.3	20.9	91	56
BEN	78/ 52	36.09	0.155	183	610	5.1	16.9	113	61
TOL	92/ 39	4.35	0.287	159	530	36.6	121.9	51	53-
TOL	92/ 51	13.65	0.164	162	540	11.9	39.6	84	54
XYL*	106/ 39	2.27	0.276	150	500	66.1	220.4	51	50
XXT*	106/ 51	1.17	0.576	177	590	151.9	506.2	82	59
XYL	106/ 65	7.62	0.337	165	550	21.6	72.2	8.5	55
XYL	106/ 91	45.86	0.374	249	830	5.4	18.1	. 427	83
NAP	128/ 78	2.99	0.341	150	500	50.2	167.2	77	50
NAP	128/128	146.53	0.340	876	2920	6.0	19.9	5209	292

ID = Identification Code

PM = Parent Mass

DM = Daughter Mass

IRF = Intermediate Response Factors

= Detection Limit
= Quantitation Limit DL

QL

SD = Standard Deviation

^{*} Ion Pairs Not Used in Quantitation

TABLE 17

THE SUMMARY OF DETECTION AND QUANITATION CONCENTRATION LIMITS FOR THE TARGET COMPOUNDS' ION PAIRS DURING SAMPLING PERIOD V

ID	PM/DM (I	IRF (CPS)	ERROR BAR (DL (ICPS) (QL ICPS)	DL (ppbů)	-	INTENSITY (ICPS) (SD (ICPS)
BEN	78/ 39 2	5.04	0.174	174	580	6.9	23.2	78	58
BEN	78/ 52 3	5.93	0.159	168	560	4.7	15.6	96	56
TOL	92/ 39	4.05	0.336	159	530	39.3	130.9	51	53
TOL	92/ 51 1	12.19	0.254	153	510	12.6	41.9	50	51
XYL*	106/ 39	1.66	0.471	258	860	155.6	518.6	46	86
XYL*	106/ 51	1.55	0.435	159	530	102.4	341.3	76	53
XYL	106/ 65	6.89	0.401	153	510	22.2	74.0	51	51
XYL	106/ 91 4	3.73	0.403	216	720	4.9	16.5	199	72
NAP	128/ 78	2.42	0.466	159	530	65.6	218.8	53	53
NAP	128/128 8	0.96	0.636	816	2720	10.1	33.6	2809	272

ID = Identification Code

PM = Parent Mass

DM = Daughter Mass

IRF = Intermediate Response Factors

DL = Detection Limit

QL = Quantitation Limit

SD = Standard Deviation

^{*} Ion Pairs Not Used in Quantitation

Calculations for the Detection and Quantitation Concentration Limits for the Target Compounds During the Sampling Periods

The detection concentration limits (DL) and quantitation concentration limits (QL) for compound were generated by averaging the respective DLs and QLs of the target compounds' ion pairs. Only the designated ion pairs in Tables 13-17 were used to determine the DLs and Qls, because others have background interferences or were insensitive.

The following equation was used to calculate the compound's detection limit concentration found in Table 18:

$$DL = \frac{DL1 + DL2 + ... + DLn}{n}$$

where:

DL = detection limit for a compound (ppbv)

DL1 = detection limit for the first ion pair (ppbv)

DL2 = detection limit for the second ion pair (ppbv)

DL3 = detection limit for the nth ion pair (ppbv)

n = number of ion pairs to be averaged

For example, the entry for the 78/39 and 78/52 ion pairs of benzene from Table 12 is:

$$DL = \frac{5.2 + 3.3}{2} = \frac{8.5}{2} = 4.25 \text{ ppbv}$$

This number was rounded up to the next whole number resulting in the detection limit equal to 5 ppbv.

The following equation was used to calculate the compound's quantitation limit concentration found in Table 18:

$$QL = \frac{QL1 + QL2 + ... + QLn}{n}$$

where:

QL = quantitation limit for a compound (ppbv)

QL1 = quantitation limit for the first ion pair (ppbv)

QL2 = quantitation limit for the second ion pair (ppbv)

QL3 = quantitation limit for the nth ion pair (ppbv)

n = number of ion pairs to be averaged

For example, the entry for the 78/39 and 78/52 ion pairs of benzene from Table 13 is:

$$QL = \frac{17.4 + 11.1}{2} = \frac{28.5}{2} = 14.25 \text{ ppbv}$$

This number was rounded up to the next whole number resulting in the quantitation limit equal to 15 ppbv.

TABLE 18

SUMMARY OF DETECTION AND QUANTITATION CONCERTRATION LIMITS FOR TARGET COMPOUNDS DURING THE SAMPLING PERIODS

CONCENTRATIONS IN PPBV

COMPOUND		SAMI	PLING	SAMI	PLING	SAMP	LING	SAME	LING	SAM	PLING
	PERI	OD I	PERI	OD II	PERI	OD III	PERI	OD IV	PERI	OD V	
	DL	QL	DL	QL	DL	QL	DL	QL	DL	QL	
benzene		5	15	6	20	8	24	6	19	6	20
toluene		15	49	20	67	25	83	25	81	26	87
xylene		10	33	12	40	16	51	14	46	14	46
naphthalene		35	116	40	134	38	127	29	94	38	127

DL = Detection Limit

QL = Quantitation Limit

Calculations for the Potential Maximum Concentration Percent Deviations for Target Compounds During the Sampling Periods

The potential maximum concentration percent deviations presented in Table 19 are called "error bars" for simplicity. They represent the potential bias in the concentration due to changes in the sensitivity of the TAGA. "Error bars" were calculated using the following equation:

"error bars" =
$$\frac{|RF1 - RF2|}{(RF1 + RF2)}$$
 X 100

where:

error bars = maximum concentration percent deviation (unitless)

RF1 = the response factor for an ion pair measured during the initial calibration event (ICPS/ppbv)

RF2 = the response factor for the same ion pair measured during the final calibration event (ICPS/ppbv)

The above calculation was repeated for each ion pair. The "error bars" for each compound were averaged to give a single value for each compound between the two calibrations each sampling period.

For example, using the BNZ data from Table 8 for the 78/39 ion pair:

RF1 = 31.14

RF2 = 37.13

and then

$$\frac{|RF1 - RF2|}{(RF1 + RF2)}$$
 X 100 = $\frac{|31.14 - 37.13|}{31.14 + 37.13}$ X 100 = 8.8%

and the other BNZ ion pair: 78/51 the "error bar" is 8.8%. These ion pair "error bars" are averaged to give an "error bar" for BNZ equal to 8.8%, which is the entry in Table 19.

TABLE 19

THE SUMMARY OF THE POTENTIAL MAXIMUM CONCENTRATION PERCENT DEVIATIONS FOR THE TARGET COMPOUNDS DURING THE SAMPLING PERIOD

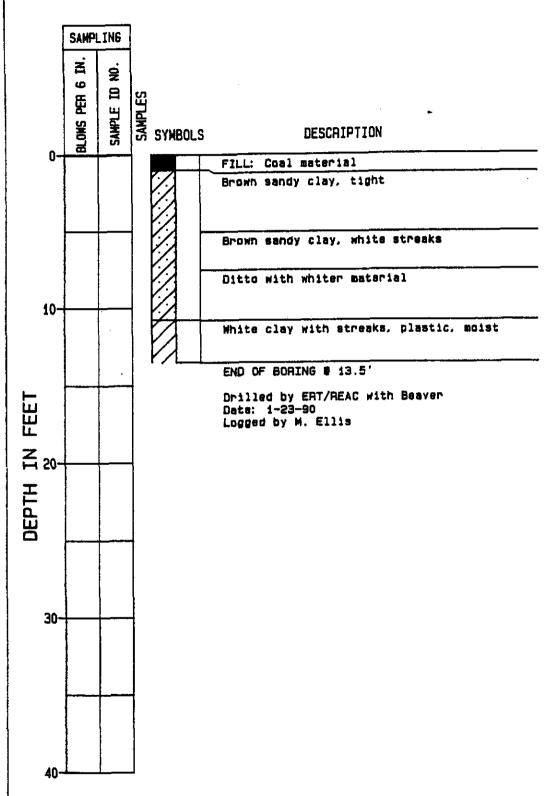
SAMPLING PER	toD I	II	III	ıv	v
СОМРОИИД	ERROR BAR	ERROR BAR	ERROR BAR	ERROR BAR	ERROR B.
	PERCENTAGE	PERCENTAGE	PERCENTAGE	PERCENTAGE	PERCENTA(
benzene	8.8	4.2	14.4	13.5	16.7
toluene	13.6	3.3	23.3	22.6	29.5
xylene	1.6	18.3	33.1	35.6	40.2
naphthalene	4.0	6.6	3.2	34.1	55.1

APPENDIX C

SOIL BORING LOGS GULF STATES CREOSOTE APRIL, 1990

BORING B-01

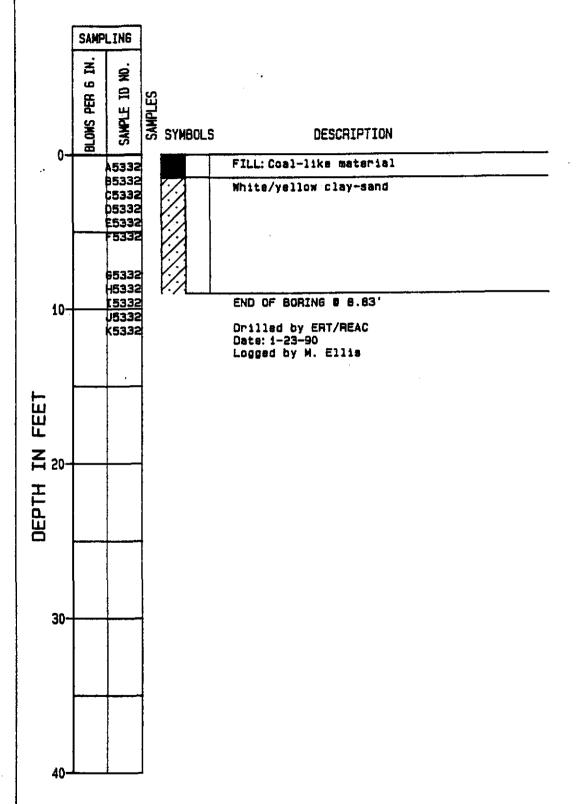
SULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

BORING B-02.5

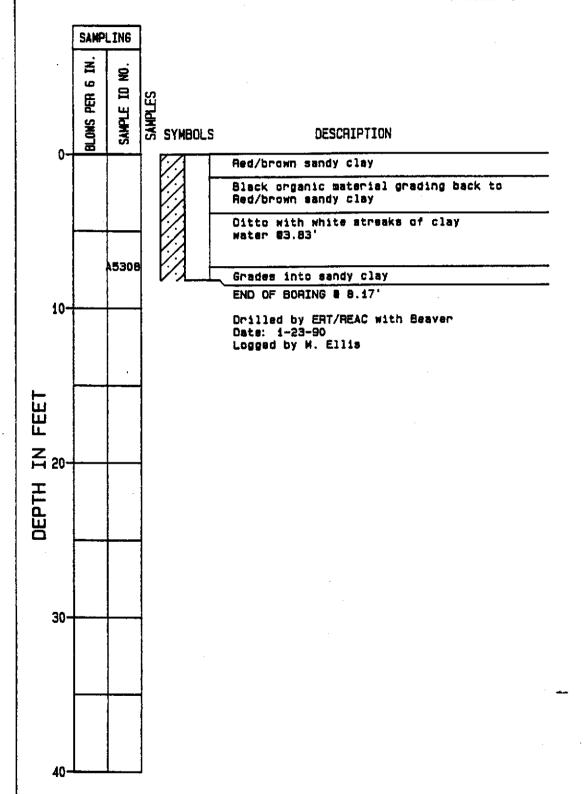
GULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

BORING B-3

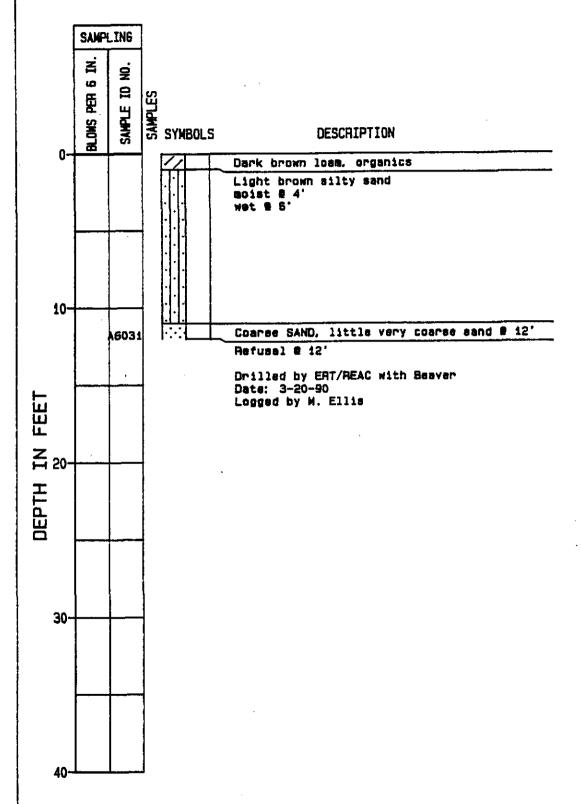
GULF STATES CREOSOTE HATTIESBURG, MS



BORING LOG

BORING C-19

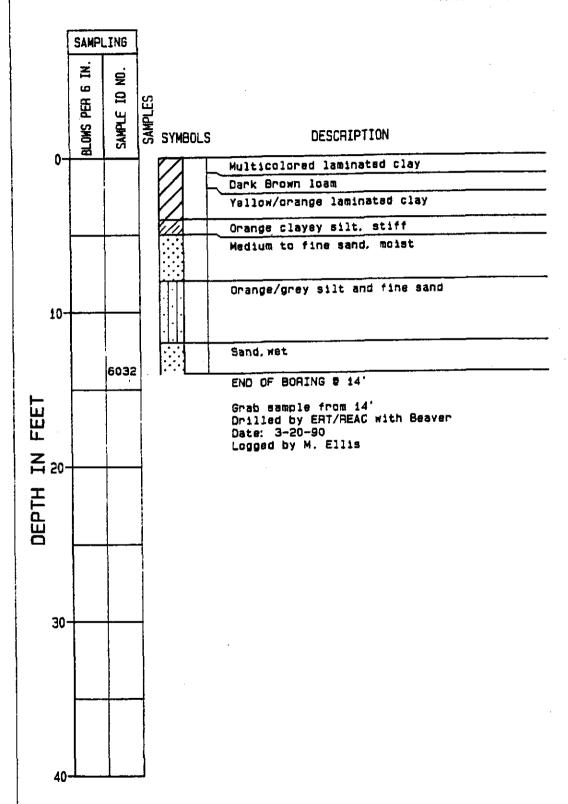
GULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

BORING C-20

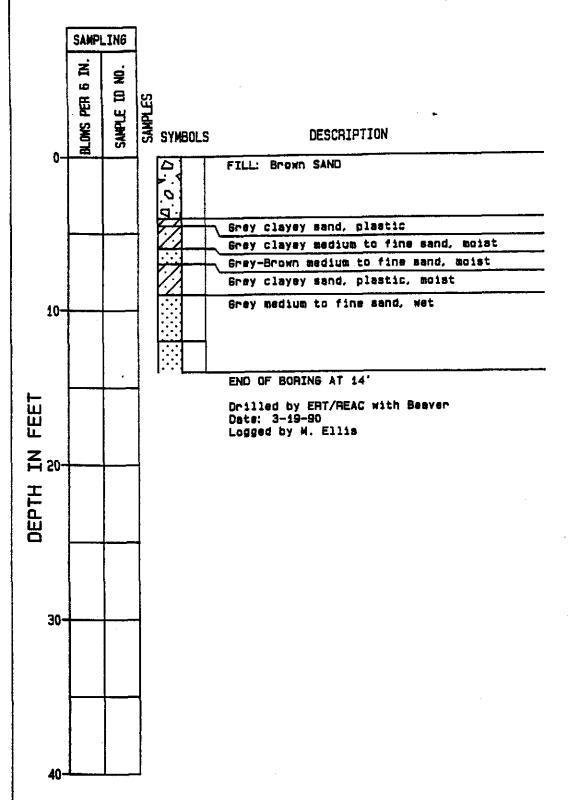
GULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

BORING D- -1

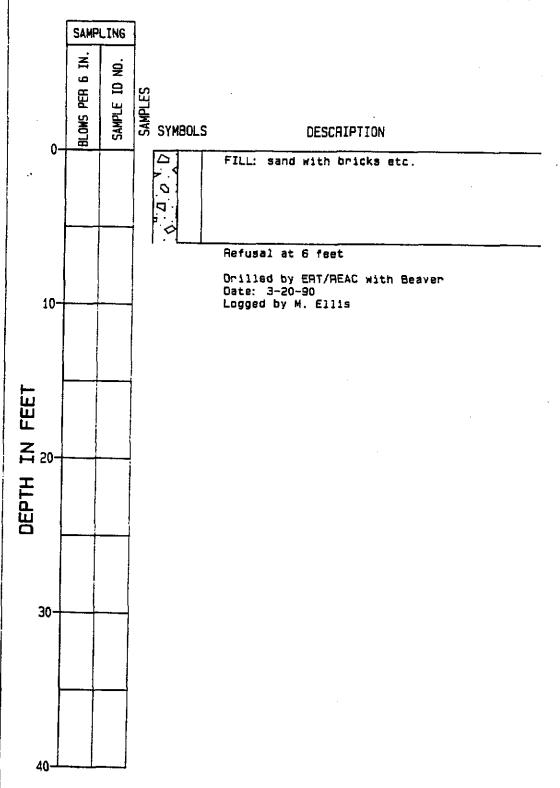
GULF STATE CREGSOTE HATTIESBURG, MS



BORING LOG

BORING D-02

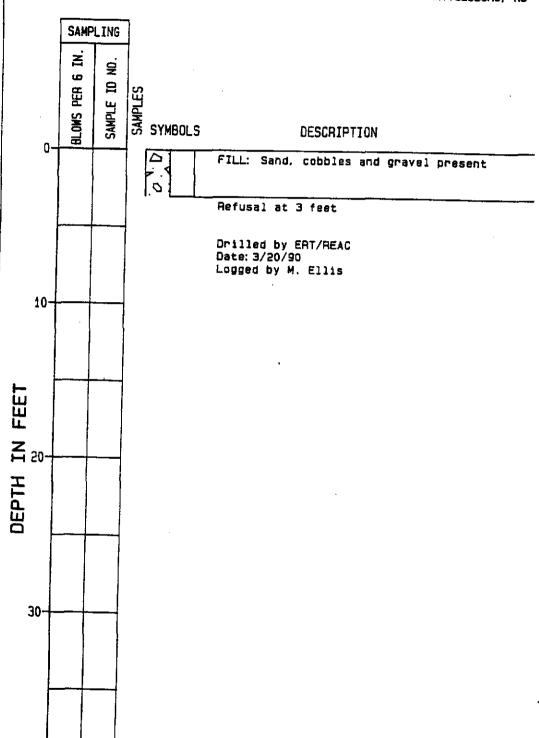
GULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

BORING D-03

GULF STATE CREOSOTE HATTIESBURG, MS

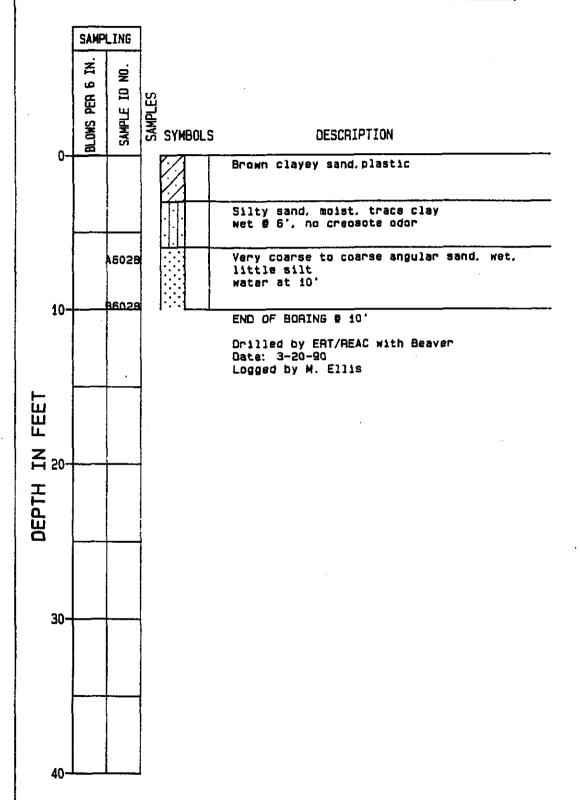


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BORING LOG

BORING D-03A

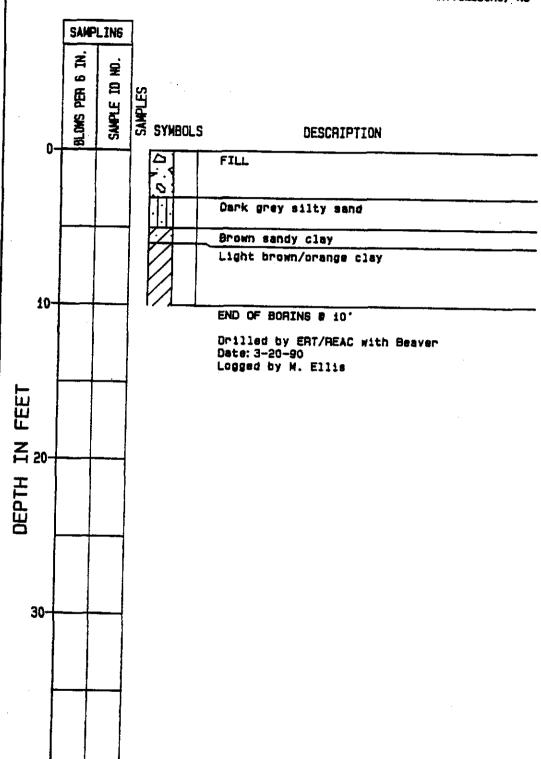
GULF STATE CREDSOTE HATTIESBURG, MS



BORING LOG

BORING D-04

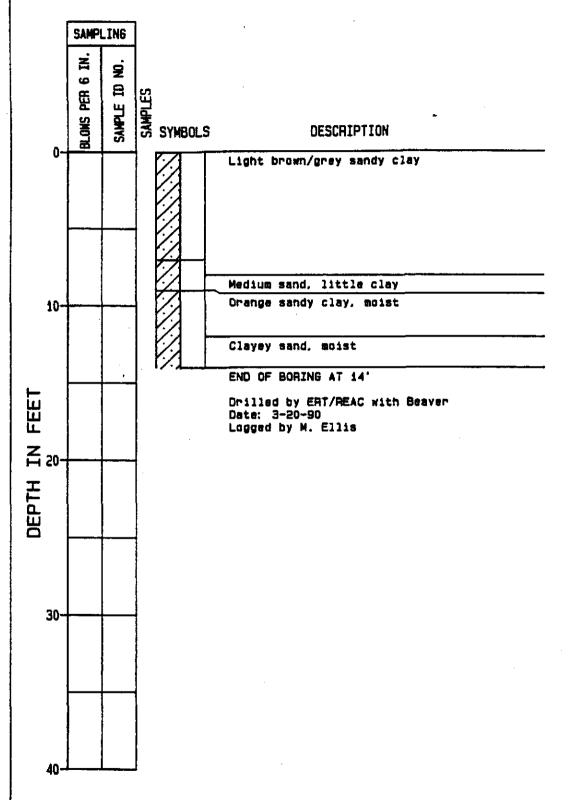
GULF STATE CREUSOTE HATTIESBURG, MS



BORING LOG

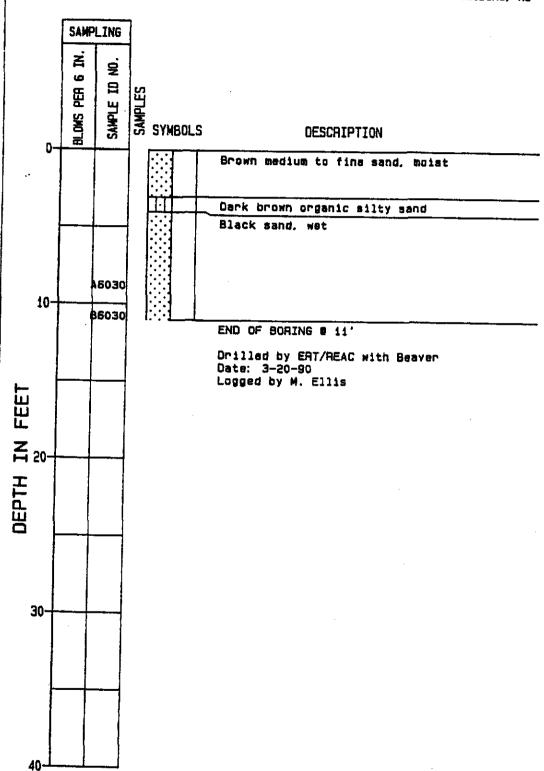
BORING D-06

GULF STATE CREOSOTE HATTIESBURG, MS



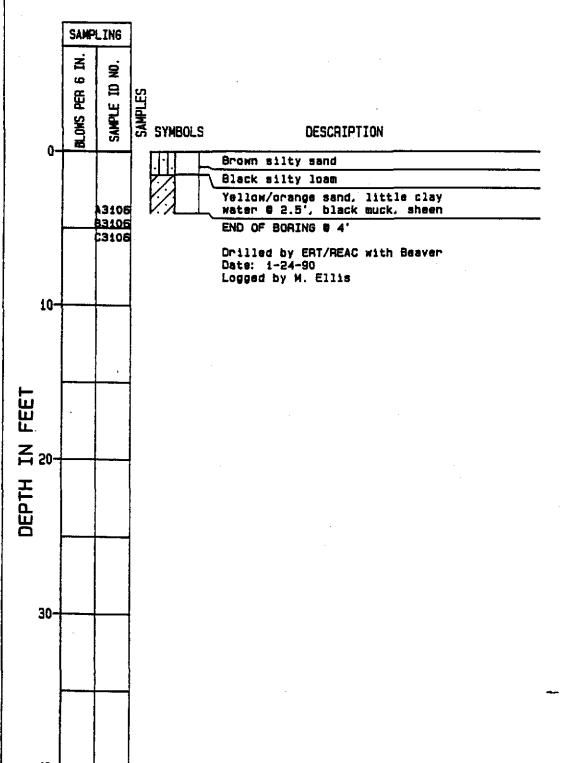
BORING LOG

GULF STATE CREOSOTE HATTIESBURG, MS



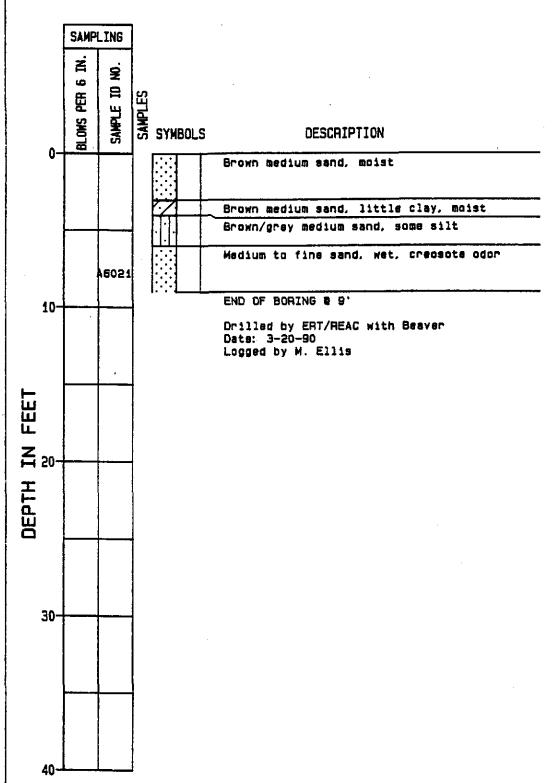
BORING LOG

GULF STATE CREOSOTE HATTIESBURG, MS



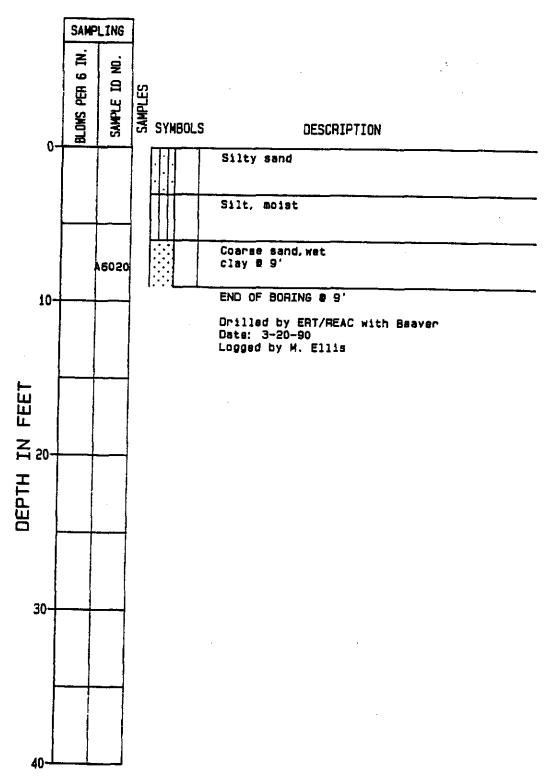
BORING LOG

GULF STATE CREDSOTE HATTIESBURG, MS



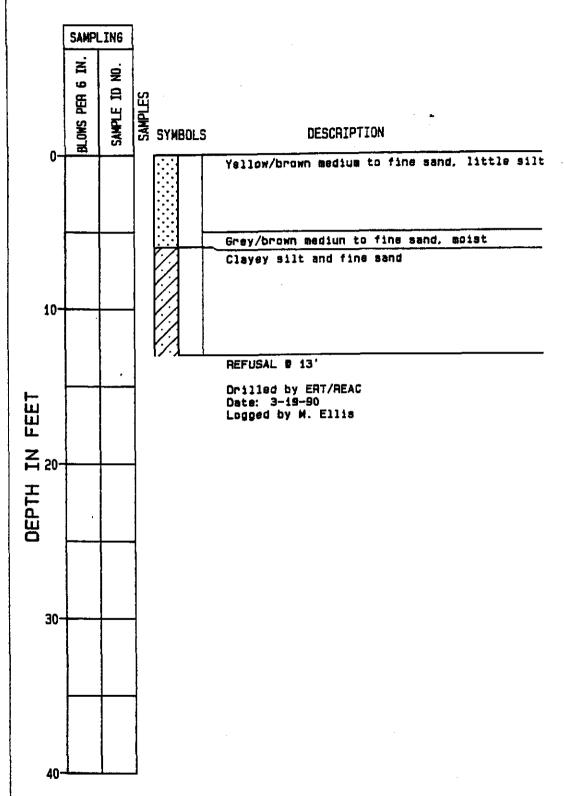
BORING LOG

GULF STATE CREOSOTE HATTIESBURG, NS



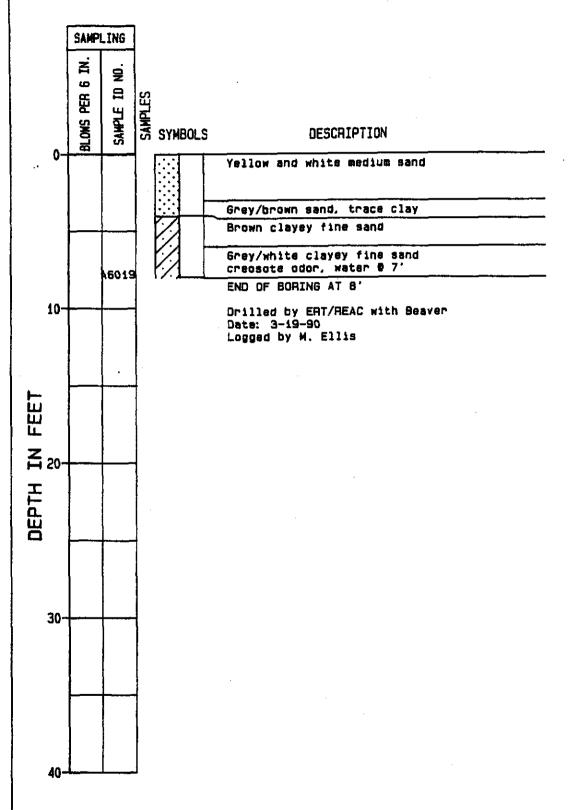
BORING LOG

GULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

GULF STATE CREOSOTE HATTIESBURG, MS



BORING LOG

APPENDIX D

FIELD DATA SHEETS GULF STATES CREOSOTE APRIL, 1990

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rd/BATZ/WORKSHEET

ENVIRONMENTAL RESPONSE TEAM AIR SAMPLING WORKSHEET

Roy F. Weston, Inc. REAC Project, Edison, NJ EPA Contract No. 68-03-3482

SITE GUIF SH	te Creos	sole w.		3-1011-74	
DATE 123190		EP		· ONEIN	
SAMPLE NO.	3203F	3204F	3206F	Field Blanke	
Sample Location	Hacks.	Neur cire K on Aned	pehins umr stæk	Field blank	
Remarks					
Pump No.	3404	7325	7342		
Collection Media	XAD+filter	· VADEGILET	YAD-Filte	XADIGIKI	
Analysis Requested	<u>5515</u>	<u>5515</u>	5515	<u>5515</u>	
Time of Day	9:45	10:00	10:10	9:00	
Time/Counter (Start)	0				<u></u>
Time/Counter (Stop)	480 Mun	<u>454min</u>	453mm		 .
Total Sampling Time	480 min	454min	453min.	<u> </u>	-M*
Pump Fault	Y. (N)	Y/N	Y/N	Y/N	Y/N
Flow Rate (Start)	2 l/m	21/m	<u>2l/m</u>		
Flow Rate (Stop)	7 Um	alfin	24m		
Flow Rate (Average)	alla	- a l/m	_2e/m		
Volume Sampled	960li.	908	906		
Air Monitoring Data HNU OVA LEL/RAM					
WEATHER PARAMETERS Weather Conditions Wind direction See Si	Sun ry K mypPressur	Tempo	erature <u>27</u> Humidity <u>2</u>	Windspeed	2 m/h
GENERAL COMMENTS:		<i></i>			



ENVIRONMENTAL RESPONSE TEAM AIR SAMPLING WORKSHEET

Roy F. Weston, Inc. REAC Project, Edison, NJ EPA Contract No. 68-03-3482

SITE Gulf Str. SAMPLERS, WB DATE 1123190	te	EF		2335 1284 (cmp	ton
SAMPLE NO.	Realtin				
Sample Location	Do	EDIZ hendspace	E014 Neadspace	H6 headspace	KI8 handspace
Remarks	above Bor	e Above bo	OUA DOTE	Eva 5	apore pro
Pump No.	HUUI	Hau i ova 5	HITE.	7447VI	Thuc i
Collection Media	HXYA	DVK COM	B OVA	<u>wa</u>	80a
Analysis Requested	none-				>
Time of Day	410	415	420	445pm	5:00 pm
Time/Counter (Start)					
Time/Counter (Stop)					
Total Sampling Time		_			
Pump Fault	Y/N	Y/N	Y/N	Y/N	Y/N
Flow Rate (Start)					
Flow Rate (Stop)					
Flow Rate (Average)					
Volume Sampled					_
Air Monitoring Data HNU OVA LEL/RAM	3.5 3.5	<u> </u>	0		6
WEATHER PARAMETERS Weather Conditions Wind direction	Sunny Pressi	Temp ure <u>30,30</u>	erature <u>27</u> Humidity <u>a</u>	<u>C</u> Windspo 고기 Met II	eed
GENERAL COMMENTS:	· • •				

MIE INC. Model PDL-1 SN 2151 V2.8 12/87

CURRENT DATE: 1/23/90 CURRENT TIME: 22:35:59

1BRATION

J.014 V = 0.000 mgm3 2.223 V = 4.500 mgm3

LOWER ALARM:

0.000

 ϵ_{mgm}

UPPER ALARM:

0.000

mgm3

UNITS: mgm3

INPUT READS: 0.046 mgm3
TEST STARTING DATE: 1/23/90
TEST STARTING TIME: 14:40:31
ELAPSED TIME: 0 DAYS 2:22:47

OVERALL AVG: 0.008 mgm3 OVERALL MIN: - 0.014 mgm3

MIN OCCURRED 1/23/90 @ 15:21:08

OVERALL MAX: 1.086 mgm3

MAX OCCURRED 1/23/90 @ 16:59:24

STEL: 0.058 mgm3

STEL OCCURRED 1/23/90 @ 16:48:11

TIME HISTORY
PERIOD LENGTH: 0:15:00
#_OF PERIODS COMBINED: 10

MIN AVG MAX IO

DATE: 1/23/90 TIME: 14:40:31 TAG #: 1 0-0.014 0.012 1.086 * AMP DIST

SAMPLES LOGGED:

8567

Emgm	SAMPLES	.	%
- 0.374	8203	************	****095.75
0.028 0.430	356 4	**** •	004.15 000.04
0.833	4	•	000.04

Lab No.: Date: 1123 90 Time:	Samplers: PC: Site Name: Sample Location:		lowel	ومدمل	Chain of Custody No. REAC Task Leader: EPA Task Monitor: Project No.:	Comptor &
	silt	color muck oder loam flow	FACE WATER	STREAM width depth velocity pools nifles	rubble clay cm/s gravel org: % shell other	anic er
SAMPLE TYPE stream/surface groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer ponar trowl other _ oucke) sugar ekman	SAMPLE color odor temp DO cond	ORP salinity _	epth 32	WEATHER PARAME ambient temp barometric pressure relative humidity weather conditions	
TOC required?Yes If No, explainYes If No, explain Grain size analysis required? If No, explain ORGANICS A. halogenated & aromatic volume aroma	_No YesNo		LIMITED CHEMII A. total cyanid B. total phenoi C. petroleum i D. pH E. alkalinity F. hardness G. total dissolv H. total susper I. suifate OTHER ANALYS	s sydrocarbons red solids nded solids	CONTAINER plastic jar plastic jar acetate core plastic bag plastic bucket 4L plastic STORAGE wet ice dry ice ambient BIOASSESSME See attached See comment	PRESERVATIVES HNO3 NaOH Zn Acetate HCL Na2SO4 other
8. metals scan (ICP) C. metals, other RCRA A. EP toxicitymetals _ B. ignitability C. corrosivitypH D. reactivity COMMENTS:		herbicides	AIR SAMPLING Sampling Method Sample Flow Rat Sampling Time Volume Collected Cool N SAMP N SAM	S	Collection Media	ctions
	10.8	'- nois-(white pla	stic cla	d 1/5trea	ke

Lab No.:		States creosote.	Chain of Custody No. REAC Task Leader: O'NRLL EPA Task Monitor: COM PTO Project No.:
industrial wooded lowland committed farmland lacustr residential gully hedgerows floodplain	SOIL TYPE paluatrine rock clay driverrine gravel muck ine auti loam silt peat color	odor depth flow velocity_	BOTTOM rock stit rubble clay cm/s gravel organic % shell other
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment siudge ANALYSES TO BE PERFORME	DEVICE kernmerer ponar trowl other bucket sugar ekman	SAMPLE INFORMATION colo PAP oder ORP temp salinity DO sample depth cond tide stage	WEATHER PARAMETERS ambient temp barometric pressure relative humidity
TOC required?Yes If No. explain Grain size analysis required? If No. explain	No Yes No	— U. pri	SAMPLE PREPARATION CONTAINER PRESERVATIVES HNO3 plastic jar NaOH s acetate core Zn Acetate plastic bag HCL
ORGANICS A. halogenated & aromatic vota B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extractable	Jacs at	E. alkalinity F. hardness G. total dissolved solids H. total suspended solids I. suifate	plastic bucket Na2SO4 4L plastic other STORAGE wet ice
G. pesticides, drinking water H. herbicides, drinking water NORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other		OTHER ANALYSES (specify)	BICASSESSMENT See attached data sheet See comments
A. EP toxicitymetals B. ignitability C. corrosivitypH D. reactivity	pesticidesherbicides	AIR SAMPLING Sampling Method Sample Flow Rate Sampling Time Volume Collected	Collection Media Special Shipping Instructions #Field Blanks#Sample Blanks
OMMENTS: D-C" 12" 12"	ance coal no	Heral / sample	Taken Bo was + Jac

BI 10" - > GHIJIK JAK OVA = NP

Lab No.:	Samplers: COMPTO Site Name: GUKS Sample Location: B-	itates Ca	PEU SOTE 1-12	Chain of Custody No. REAC Task Leader: _ EPA Task Monitor: _ Project No.:	O'NEAL
	palustrine rock clay friverrine gravel muc sand loar silt pea	n flow	width depth velocity	rubble cla cm/s gravel org % shell oth	y Janic
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer ponar trowd other bucket sugar ekman	SAMPLE INFO color oder temp DO cond	ORPsalinity	barometric pressure	
ANALYSES TO BE PERFORME TOC required?Yes If No. explain Grain size analysis required? If No. explain ORGANICS A. halogenated & aromatic voi B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extractable G. pesticides, drinking water H. herbicides, drinking water INORGANICS A. metals, priority pollutant B. metals scan (ICP)	_No YesNo atiles	A. (B. (C. (D. (F.) G. (H.)	ED CHEMISTRY total cyanide total phenol petroleum hydrocarbons ph alkalinity nardness total dissolved solids total suspended solids ulfate ER ANALYSES (specify)	CONTAINER glass jar plastic jar acetate core plastic bucket 4L plastic STORAGE wet ice dry ice ambient BIOASSESSME See attached See comment	PRESERVATIVES HN03 NaOH Zn Acetate HCL Na2SO4 other
C. metals, other RCRA A. EP toxicitymetals _ B. ignitability C. corrosivitypH D. reactivity	pesticidesherbio	cides Samp Samp Samp	AMPLING ling Method le Flow Rate ling Time le Collected	Collection Media Special Shipping Instru #Field Blanks#S	ctions
COMMENTS: 1135 Augus	1.5 - 24" 310" W 7'4 R	Black or	N SANDY CIA govic Materia apy RBSC dingila Kao	lyradinsbackte	

Nº 006031

Lab No.:	Samplers:				
Date: 3/2-0/20 Time: /4/5	Site Name:	C-\$35 19		EPA Task Monitor:	335
SITE DESCRIPTION landfill old field uplar industrial wooded lowla	SOIL TYPI nd palustrine rock cland riverrine gravel matrine sand to		width depth velocity	rubble cia cm/s gravel or % shell ott	
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer poner trowl other bucket sugar ekman	odor ORP temp salinii DO samp	ON ty le depth stage	barometric pressure relative humidity	·
ANALYSES TO BE PERFORM	MED			SAMPLE PRE	PARATION
TOC required?Yes If No, explain Grain size analysis required? If No, explain ORGANICS A. hatogenated & aromatic B. volatiles-USEPA 524 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extract G. pesticides, drinking wate H. herbicides, drinking wate H. herbicides, drinking wate INORGANICS A. metals, priority pollutant B. metals scan (iCP) C. metals, other	YesNo voistiles ables-USEPA 625	B. total ph C. petrolet D. pH E. alkalinit F. hardnet G. total dii H. total su I. suifate	anide ienol um hydrocarbons	plastic bag plastic bucket 4L plastic STORAGE wet ice dry ice ambient BIOASSESSM	d data sheet
1-2' H	brown loan brown kilty he ditto	ml 4-8	ethod v Rate me ected ditto	(Sample Blanks A: Btm Samp Samb

Nº 006032

Lab No.:	_ Samplers: _			·	Chain of Custody No. REAC Task Leader:	
Date: 3/20/90	_ Site Name:			·	EPA Task Monitor:	
Time: 1440	Sample Loc	ation:	<u>C-20</u>		Project No.:	<i>33<u>5</u></i>
industrial wooded	upland palustrine. lowland riverrine lacustrine	SOIL TYPE rock clay gravel muck sand loam silt peat color	SURFACE WATER color odor flow direction	STREAM width depth velocity pools riffles	rubble clay cm/s gravel org	
SAMPLE TYPE stream/surface soil groundwater pond/ brackish river ocean/saline efflue sediment sludg	bucket nt sugar	ponar other	odor ORP _ temp salinity	depth	WEATHER PARAME ambient temp barometric pressure relative humidity weather conditions_	
ANALYSES TO BE PERF	OAMED				SAMPLE PREP	PARATION
TOC required?Yes			LIMITED CHEM A. total cyan B. total phen	ide	CONTAINER glass jar plastic jar	PRESERVATIVES HNO3 NaOH
Grain size analysis required in No. explain			C. petroleum D. pH E. alkalinity	hydrocarbons	acetate core plastic bag plastic bucket	Zn Acetate HCL Na2SO4
ORGANICS A. halogenated & arom B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid ex G. pesticides, drinking	(tractables-USEPA 6	25	F. hardness G. total disso	endød solids	STORAGE wet ice dry ice ambient	other
H. herbicides, drinking INORGANICS A. metals, priority pollu B. metals scan (ICP) C. metals, other	-				— BIOASSESSME See attached See commen	data sheet
ACRA A. EP toxicitym B. ignitability C. corrosivitypH D. reactivity	etalspesticid	esherbici	AIR SAMPLING Sampling Meth Sample Flow R Sampling Time Volume Collect	late	Collection Media Special Shipping Instru #Field Blanks#S	octions
of the	-1' milli -1 -2' dk bron -3' yllow	un loan	util chays	7-	8' detto	wilt-time

No 006017

Roy F. Weston, Inc. REAC, Edison, N.J. EPA Contract 68-03-3482

Chain of Custody No. Lab No: REAC Task Leader: Date: Site Name: EPA Task Monitor: Time: Sample Location: Project No.: SITE DESCRIPTION SOIL TYPE SURFACE WATER STREAM BOTTOM landfil) Old field upland palustrine rock clay color . width_ rock industrial baboow lowland rivernne gravei muck odor depth_ rubble clay commercial farmland lacustrine sand loam gravel organic residential guily Silt peat direction HACE other hedgerows Copepiato color sand SAMPLE TYPE DEVICE SAMPLE INFORMATION WEATHER PARAMETERS 3)ream/surface kemmerer DODAY color pH. ambient temp groundwater pond/lake ORP trowi other odor barometric pressure brackish river bucket temo salinity . relative humidity. ocean/saline effluent sugar DO sample depth weather conditions. sediment sludge ekman cond tide stage ANALYSES TO BE PERFORMED SAMPLE PREPARATION TOC required? ____ ___Yes ____ No LIMITED CHEMISTRY CONTAINER PRESERVATIVES If No. explain __ A. total cyanide glass jar HN_O3 B. total phenoi plastic jar NaOH Grain size analysis required? ______Yes _____No C. petroleum hydrocarbons acetate core Zn Acetate If No, explain D. pH plastic bad HCL E. alkalinity plastic bucket Na2SO4 **ORGANICS** F. hardness 4L plastic other A. halogenated & aromatic volatiles G. total dissolved solids B. volatiles-USEPA 624 H. total suspended solids C. trihalomethanes I. suifate STORAGE D. pesticides/PCB wet ice E. PCB dry ice F. base neutral/acid extractables-USEPA 625 OTHER ANALYSES (specify) ambient G. pesticides, drinking water H. herbicides, drinking water BIOASSESSMENT INORGANICS See attached data sheet A. metals, priority pollutant See comments B. metals scan (ICP) C. metals, other **RCRA** AIR SAMPLING A. EP toxicity _ __metals ____pesticides ____herbicides Sampling Method _ Collection Media 8. ignitability Sample Flow Rate_ Special Shipping Instructions

Sampling Time

Valume Collected

COMMENTS: grey clayer sand, plastic grey clayer med-fine sands worst grey times and brown, alpents grey clayer sands, worst plastic

C. corresivity_

D. reactivity

Nº 006023

Roy F. Weston, Inc. REAC, Edison, N.J. EPA Contract 68-03-3482

Lab No.: Samplers: Samplers: Site Name: Sample Location		- <i>o</i> a		•	0
landfill old field upland palustrine rot industrial wooded lowland riverrine gracommercial farmland lacustrine sail residential gully	ivel muck	SURFACE WATER color odor flow direction	STREAM width depth velocity pools riffles	rubble c cm/s gravel o % shell o	lit lay rganic ther
SAMPLE TYPE DEVICE stream/surface soil kemmerer pon groundwater pond/lake trowl othe brackish river bucket ocean/saline effluent sugar sediment sludge ekman	ar er	temp salinity		barometric pressurelative humidity_	METERS re
ANALYSES TO BE PERFORMED	· · · · · · · · · · · · · · · · · · ·			SAMPLE PR	EPARATION
TOC required?YesNo If No. explainYesNo Grain size analysis required?YesNo		B. total pher	nide	CONTAINER glass jar plastic jar acetate core	PRESERVATIVES HNO3 NaOH Zn Acetate
If No, explain		D. pH E. alkalinity	,	plastic bag plastic bucke	HCL
ORGANICS A. halogenated & aromatic volatiles B. volatiles-USEPA 624		F. hardness G. total diss		4L plastic	other
C. trihalomethanes D. pesticides/PCB E. PCB		i. sulfate		STORAGE wet ice dry ice	
F. base neutral/acid extractables-USEPA 625 G. pesticides, drinking water H. herbicides, drinking water		OTHER ANAL	YSES (specify)	ambient .	
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other				BIOASSESSA See attach See commo	ed data sheet
RCRA A. EP toxicitymetalspesticides _ B. ignitability C. corrosivitypH D. reactivity		AIR SAMPLING Sampling Mett Sample Flow F Sampling Time Volume Collect	Rate	Collection Media Special Shipping Inst	ructions

COMMENTS

0-3 fell material
3-6' more fill, brichaste.
drill augus are bending
Stopped dulling

N: 005510

Lab No.: Samplers Date: 1-24 Site Name Time: 33 Sample L	· Creff	tote Creas	Il de	Chain of Custody No REAC Task Leader: EPA Task Monitor: Project No.:	<u> </u>
SITE DESCRIPTION landfill old field upland palustrine industrial wooded lowland riverrine commercial farmland lacustrine residential gully hedgerows floodplain	SOIL TYPE rock clay gravel muck sand loam silt peat color	SURFACE WATER color odor flow direction	STREAM width depth velocity pools riffles	rubble cia cm/s gravel org	
SAMPLE TYPE DEVICE stream/surface soil kemmere groundwater pond/lake trowl brackish river bucket ocean/saline effluent sugar sediment sludge ekman	er ponar other	odor ORP _ temp satinity DO sample		_ barometric pressure	1CC %
ANALYSES TO SE PERFORMED TOC required?	No	LIMITED CHEI A. total cyan B. total pher C. petroleum D. pH E. alkalinity	iide	plastic bag	PRESERVATIVES HNO3 NaOH Zn Acetate HCL
ORGANICS A. halogenated & aromatic volatiles 5. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extractables-USEPA G. pesticides, drinking water H. herbicides, drinking water	625	F. hardness G. total disse	ended solids	plastic bucket 4L plastic STORAGE wet ice dry ice ambient	Na2SO4 other
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other		·		BIOASSESSME See attached See comment	data sheet
RCRA A. EP toxicitymetalspestici B. ignitability C. corrosivitypH D. reactivity COMMENTS:	desherbicide	AIR SAMPLING Sampling Meth Sample Flow R Sampling Time Volume Collect	od	Collection Media Special Shipping Instru- #Field Blanks#S	ctions

Nº 005511

Roy F. Weston, Inc. REAC, Edison, N.J. EPA Contract 68-03-3482

Date: 1279 Site	nplers: QDP Name: C74 H S nple Location: DX	tate Creasate 20 - 5 ft.	Chain of Custody No. REAC Task Leader: EPA Task Monitor: Project No.:
SITE DESCRIPTION landfill old field upland pale industrial wooded lowland rive commercial farmland lacustrine residential gully hedgerows floodplain	/	SURFACE WATER STREAM color width odor depth flow velocity direction pools riffles	rubble claycm/s gravel organic% shell other
stream/surface soil ker groundwater pond/lake tro brackish river bu- ocean/saline effluent sur	EVICE mmerer ponar ow) other icket gar man	SAMPLE INFORMATION color pH odor ORP temp satinity DO sample depth cond tide stage	relative humidity 1(1/67)
ANALYSES TO BE PERFORMED TOC required?	YesNo	LIMITED CHEMISTRY A. total cyanide B. total phenol C. petroleum hydrocarbons D. pH E. alkalinity F. hardness G. total dissolved solids H. total suspended solids i. sulfate OTHER ANALYSES (specify)	plastic bag HCL plastic bucket Na2SO4 4L plastic other STORAGE wet ica dry ice ambient
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other RCRA A. EP toxicitymetals B. ignitability C. corrosivitypH D. reactivity	pesticidesherbicide	AIR SAMPLING Sampling Method Sample Flow Rate Sampling Time Volume Collected	BIOASSESSMENT See attached data sheet See comments Collection Media Special Shipping Instructions #Field Blanks #Sample Blanks

OVA- De ppin

N: 005321

Roy F. Weston, Inc.

REAC, Edison, N.J. EPA Contract 68-03-3482

Lab No.:	Samplers:	ell Stat	Compton	Chain of Custody No. REAC Task Leader: EPA Task Monitor: Project No.:	0'NEL
	SOIL T d palustrine rock nd riverrine gravel trine sand silt color	clay color odor loam flow	depth velocity_	rock slit	y Janic
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer ponar trowl other _ bucket sugar ekman	SAMPLE IN color odor temp DO cond	ORP	barometric pressure relative humidity	
ANALYSES TO BE PERFORM TOC required?Yes If No, explain ORGANICS A. halogenated & aromatic v	No YesNo		AITED CHEMISTRY A. total cyanide 3. total phenol C. petroleum hydrocarbo D. pH E. alkalinity F. hardness 3. total dissolved solids	CONTAINER glass tal plastic jar acetate core plastic bag plastic bucket 4L plastic	PRESERVATIVES HNO3 NaOH Zn Acetate HCL Na2SO4 other
B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extracts G. pesticides, drinking water H. herbicides, drinking water	7	I	total suspended solid sulfate HER ANALYSES (speci	STORAGE wet ice dry ice	
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other	() \$1 (Ac	ross fram	BIOASSESSMI See attached 860 commer on Pine S	data sheet
A. EP toxicitymetals . B. ignitability C. corrosivitypH D. reactivity	pesticides	herbicides Sa Sa Sa	R SAMPLING mpling Method mple Flow Rate mpling Time lume Collected	Collection Media Special Shipping Instru	Sample Blanks
COMMENTS: A AAB C & D	5' dup 8' deep	gray,	, odorous	, in water	r

Nº 006028

Roy F. Weston, Inc. REAC, Edison, N.J. EPA Contract 68-03-3482

SITE DESCRIPTION landfill old field upland palu industrial wooded lowland rive commercial farmland lacustrine	SOIL TYPE	0110710			Project No.:	2335
residential gully hedgerows floodplain	· ·	y color ck oder m flow	E WATER	STREAM width depth velocity pools riffles	rubble o cm/s gravel o % shell o	alit day organic other
stream/surface soil kee groundwater pond/lake tro brackish river bu- ocean/saline effluent sur	VICE mmerer poner wi other cket gar man	SAMPLE INI calor odor temp DO cond	ORP satinity sample di		barometric pressurelative humidity_	METERS Ire
ANALYSES TO BE PERFORMED TOC required?YesNo If No. explain	YesNo		AITED CHEMINAL total cyanida 3. total phenolonic, petroleum in 5. ph 5. alkalinity 6. hardness 6. total dissolv 6. total susper 6. sulfate HER ANALYS	ed solids	SAMPLE PRI CONTAINER glass jar plastic jar acetate core plastic bag plastic bucke 4L plastic STORAGE wet ice dry ice ambient	PRESERVATIVES HNO3 NaOH Zn Acetate HCL
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other RCRA A. EP toxicitymetals B. ignitability C. corrosivitypH	pesticidesherb	picides Sai Sai	R SAMPLING mpling Method mple Flow Rai mpling Time ,		BIOASSESSI See attach See comm Collection Media Special Shipping Ins	ed d <u>ata</u> sheet ents

In woods and week corr of D-03 0-2' brown clayer sand, platic
3' ditti
3-6 Selly sand, mist tracely.
6' cuttings up wet, some
sand, no cresses start

6-9 detto 16'- noter, no odor will take somple

D-0314

A: wetsendyen B: Btmargur

Lab No.:	20	Samplers: _ Site Name: Sample Loc	ation:		<u> </u>			REAC 1	ask Leader: sk Monitor:	0.
industrial w commercial fa residential g	id field upland	i palustrine d riverrine rine	SOIL T rock gravel sand silt color _	YPE clay muck loam peat	color _ odor _	CE WATER	STREAM width depth velocity pools riffles	cm/s	rubbie c	it ay rganic ther
SAMPLE TYPE stream/surface groundwater brackish ocean/saline sediment	soil pond/lake river effluent sludge	DEVICE kemmerer trowi bucket sugar ekman	ponar other		SAMPLE II color odor temp DO cond	ORP salinity sample	depth	ambie barom relativ		'e
ANALYSES TO	BE PERFORMI	D	···					s	AMPLE PRE	PARATION
If No, explain Grain size analy If No, explain ORGANICS A. halogenate B. volatiles-U C. trihalometl D. pesticides/ E. PCB F. base neutr G. pesticides/ H. herbicides	hanes	Yes				D. pH E. alkalinity F. hardness G. total disso H. total susp I. sulfate	ide iot hydrocarbons	g pi ai pi 4i S S w di		other
B. metals sca C. metals, oth RCRA A. EP toxicity B. ignitability C. corrosivity, D. reactivity	n (ICP) hermetals _	pesticide		herbicid	es Sa Sa Sa Vo	R SAMPLING mpling Meth imple Flow R impling Time lume Collect	od	Collection Special S	hipping Insti	
COMMENTS	0-3' 3' sc	fell lty sa	wat vde,	der	k gu	7 10	9' det	to,		

Nº 006026

Lab No.:	Samplers: Site Name: Sample Location:_	N-0	6		REAC Task Le	ader:
	palustrine rock i riverrine grave	ciay co muck od loam flo peat dir	JRFACE WATER lor or w	STREAM width depth velocity pools riffles	rubbi cm/s grave % shell	slit le clay el organic other
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge	bucket suger ekman	colar odor temp	ORP salinity _	lepth	barometric pri relative humic weather cond	essure
ANALYSES TO BE PERFORME TOC required?Yes If No. explain Grain size analysis required? If No. explain ORGANICS A. halogenated & aromatic vol. B. volatiles-USEPA 624	No		LIMITED CHEM! A. total cyanid B. total phenoi C. petroleum i D. pH E. alkalinity F. hardness G. total dissoit H. total suspei	le I hydrocarbons wed solids	CONTAII glass jar plastic ja acetate c plastic be plastic be 4L plastic	HNO3 IT NaOH CORE Zn Acetate ag HCL ucket Na2SO4
C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extractable G. pesticides, drinking water H. herbicides, drinking water	03-USEPA 625		I. sulfate		STORAG wet ide dry ide ambient	
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other					See att	ESSMENT tached data sheet imments
RCRA A. EP toxicitymetals B. ignitability C. corrosivitypH D. reactivity	pesticides	_herbicides	AIR SAMPLING Sampling Method Sample Flow Rat Sampling Time _ Volume Collected		Collection Media Special Shipping #Field Blanks	
COMMENTS: 0-3' B 3-6' C 7' C 8' 'n	t brown/gregetto litto, moi ned sand ye sand	st littled clay, wo	ly ist	17' ds	yy sand	? word

N: 005314

Roy F. Weston, Inc.

REAC, Edison, N.J. EPA Contract 68-03-3482

Lab No.: Samplers: Prince Date: Site Name: G./f Time: Sample Location: E	1 E//s States 0 14	Chain of Custody No
SITE DESCRIPTION landfill old field upland palustrine rock clay industrial wooded lowland riverrine gravel muck commercial farmland lacustrine silt peat color	SURFACE WATER STREAM color width depth depth velocity direction pools iffles	rubble claycm/s gravel organic% snell other
stream/surface soil kemmerer ponar col groundwater pond/lake trow other od brackish river bucket and ten ocean/saline effluent sugar DC	MPLE INFORMATION or pH	barometric pressure
ANALYSES TO BE PERFORMED TOC required?YesNo If No, explainYesNo If No, explainYesNo If No, explainYesNo ORGANICS A. halogenated & aromatic volatiles B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extractables-USEPA 625 G. pesticides, drinking water H. herbicides, drinking water	LIMITED CHEMISTRY A. total cyanide B. total phenol C. petroleum hydrocarbons D. pH E. alkalinity F. hardness G. total dissolved solids H. total suspended solids I. sulfate OTHER ANALYSES (specify)	SAMPLE PREPARATION CONTAINER PRESERVATIVES glass jar HNO3 plastic jar NaOH acetate core Zn Acetate plastic bag HCL plastic bucket Na2SO4 4L plastic other STORAGE wet ice dry ice ambient BIOASSESSMENT See attached data sheet
A. metals, priority pollutant B. metals scan (ICP) C. metals, other RCRA A. EP toxicitymetalspesticidesherbicides B. ignitability C. corrosivitypH D. reactivity COMMENTS: Seil SAMple Silf Comments	AIR SAMPLING Sampling Method Sample Flow Rate Sampling Time Volume Collected V SAND Serie A	Collection Media Special Shipping Instructions #Field Blanks #Sample Blanks W. C. @ 4.5 -

Nº 006030

Roy F. Weston, Inc.

REAC, Edison, N.J. EPA Contract 68-03-3482

Lab No.: 3/20/90 Time:	Samplers: Site Name: Sample Loca	tion:E	- 19 - 19			REAC T	ask Leader: _	
,	f palustrine d riverrine rine	SOIL TYPE rock clay gravel mucl sand loam sift peat color	color k odor i flow	E WATER	STREAM width depth velocity pools riffles	cm/s	•	
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer trowl bucket sugar ekman	ponar other	color odor temp DO	ORP satinity _	depth	ambie barom relativ	etric pressure e humidity	ETERS
ANALYSES TO BE PERFORMI TOC required?Yes If No, explain Grain size analysis required? If No, explain ORGANICS	No	_		D. pH E. alkalinity F. hardness	de bl hydrocarbons	9 p a p	AMPLE PREF CONTAINER lass jar lastic jar cetate core lastic bag lastic bucket L plastic	PARATION PRESERVATIVES HNO3 NaOH Zn Acetate HCL Na2SO4 other
A. halogenated & aromatic vi B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extracta G. pesticides, drinking water H. herbicides, drinking water	bles-USEPA 62	25		G. total disso H. total suspe I. suifate THER ANALY		w	TORAGE let ice lry ice mbient	
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other			_				IOASSESSMI See attached See commer	data sheet
RCRA A. EP toxicitymetals _ B. ignitability C. corrosivitypH D. reactivity		sherbi	cides S4 S4 S4	R SAMPLING Impling Methological Imple Flow Ri Impling Time Jume Callect	od		n Media_ Shipping Instri anks#	sections
veete 4'		on pilt	ned-fine Z Aband Ls		A: 8	ptv-9	\ sampli	at bellow

Roy F. Weston, inc.

REAC, Edison, N.J. EPA Contract 68-03-3482

Lab No.:/ Sampler Date:/250	1	St. Geneto E-20		Chain of Custody No. REAC Task Leader: EPA Task Monitor: Project No.:	M. O. Neel
SITE DESCRIPTION landfill old field upland palustrir industrial goden lowland riverring commercial farmland lacustrine residential gulty hedgerows floodplage		SURFACE WATER color odor flow direction	STREAM width depth velocity pools riffles	cm/s gravel org	•
SAMPLE TYPE DEVICE stream/surface soil kemme groundwater pond/lake trowl brackish river bucket ocean/saline effluent sugar sediment sludge ekman	_	temp satinity	depth	barometric pressure relative humidity weather conditions	100 west rainy
ANALYSES TO BE PERFORMED TOC required?YesNo If No. explainYes If No. explainYes If No. explainYes ORGANICS A. halogenated & aromatic volatiles B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PC8 E. PC8		D. pH E. alkalinity F. hardness G. total diss H. total susp I. sulfate	nide nol n hydrocarbons olved solids pended solids	CONTAINER glass jar plastic jar acetate core plastic bag plastic bucket 4L plastic STORAGE wet ice dry ice	PRESERVATIVES HNO3 NaOH Zn Acetate HCL
Descendental/acid extractables-USE/ G. pesticides, drinking water H. herbicides, drinking water INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other	-A 025	AIR SAMPLIN	YSES (specify)	BIOASSESSM See attached See commen	d data sheet
A. EP toxicitymetalspes B. ignitability C. corrosivitypH D. reactivity	ticidesherbic		nod Pate	Collection Media Special Shipping Instru	Sample Blanks
COMMENTS: 01.0' brown setty -1.5' black setty 5-2.0 yellow-range. 5 water (black much	loan mud, little	Ţ		'semple (blea Sample, hole	

Nº 006021

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Lab No.: 3/30/20 Date: 3/30/20 Time: 0840	Samplers: Site Name: Sample Location:	<u>E-3</u>	1			ody No	
	SOIL spalustrine rock gravel gravel sand silt color	ciay co muck oc loam flo peat di	URFACE WATER blor blor bw rection	STREAM width depth velocity pools riffles	rock rubb cm/s grave	le clay el organic other	
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer ponar trowl other _ bucket sugar ekman	color odor temp DO	ORP satinity sample	depth	ambient temp barometric pi relative hymi	PARAMETERS pressure dity ditions	
ANALYSES TO BE PERFORME TOC required?Yes If No. explain Grain size analysis required? If No, explain ORGANICS A. halogenated & aromatic vol. B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PCB F. base neutral/acid extractations.	No YesNo		D. pH E. alkalinity F. hardness G. total diss H. total susp i. suifate	nide nol n hydrocarbons	SAMPLE CONTAI glass jai plastic ji acetate plastic b plastic b 4L plasti STORAC wet ice dry ice ambient	HNO3 AT NAOH COTE Zn Acetate PAGE HCL PUCKET Na2SO4 AC OTHER	
G. pesticides, drinking water H. herbicides, drinking water INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other					BIOASS See a	ESSMENT ittached data sheet omments	
A. EP toxicitymetals _ B. ignitability C. corrosivitypH D. reactivity	pesticides	_herbicides	AIR SAMPLIN Sampling Meti Sample Flow I Sampling Time Volume Collect	nod	Collection Media Special Shipping		<u> </u>

Beck

6' five- put med sent wint create other

pample at 81

Nº 006020

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Date: 3/292	Samplers:	-35	Chain of Custody No REAC Task Leader: EPA Task Monitor: Project No.: 2 3 3 5
	SOIL TYPE palustrine rock clay riverine gravel muck sand loam silt peat color	SURFACE WATER STREAM color width odor depth flow velocity direction pools riffles	rubble claycm/s gravel organic% shell other
stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment studge	DEVICE kemmerer ponar trowi other bucket sugar ekman	SAMPLE INFORMATION color pH odor ORP temp salinity DO sample depth cond tide stage	barometric pressure relative humidity
ANALYSES TO BE PERFORMED TOC required?Yes! If No, explain Grain size analysis required?! If No, explain ORGANICS A. halogenated & aromatic volations. Just the properties of the pesticides of the pesticides of the pesticides of the pesticides. Just the pesticides of	YesNo	LIMITED CHEMISTRY A. total cyanide B. total phenol C. petroleum hydrocarbon: D. pH E. alkalinity F. hardness G. total dissolved solids H. total suspended solids I. sulfate OTHER ANALYSES (specify)	plastic bag HCL plastic bucket Na2SQ4 4L plastic other STORAGE wet ice dry ice
INORGANICS A. metals, priority pollutant B. metals scan (ICP) C. metals, other			BIOASSESSMENT See attached data sheet See comments
RCRA A. EP toxicitymetals B. ignitability C. corrosivitypH D. reactivity	pesticidesherbicid	AIR SAMPLING es Sampling Method Sample Flow Rate Sampling Time Volume Collected	Collection Media Special Shipping Instructions
COMMENTS: 0-3'	sitty sand	Totaline Constitution	#Field Blanks #Sample Blanks

3' moist sell

9' but cley layer
crenote present

E-25 A: Surge

Lab No.: 3/19/91 Date: 3/19/91	Samplers:Site Name:Sample Location:	Gulf S	de Cues	70		K Leader: _ Monitor: _	335
	SOIL To sailt color	clay color muck odor loam flow peat direc	FACE WATER	STREAM width depth velocity pools riffles		SOTTOM ock slit ubble clay gravel orguinell other iand	anic
SAMPLE TYPE stream/surface soil groundwater pond/lake brackish river ocean/saline effluent sediment sludge:	DEVICE kemmerer ponar trowl other _ bucket sugar ekman	color	ORP salinity sample	depth	ambient to barometri relative h	c pressure umidity	TEAS
ANALYSES TO BE PERFORM TOC required?	NoNoNo olatiles		D. pH E. alkalinity F. hardness G. total disso	ide ioi hydrocarbons blyed solids ended solids	CON glass acet plass 4L p STO wet dry i amb	tic jar ate core tic bag tic bucket elastic PRAGE ice	PRESERVATIVES HNO3 NaOH Zn Acetate HCL Na2SO4 other
RCRA A. EP toxicitymetals B. ignitability C. corrosrvitypH D. reactivity	posticides	_herbicides	AIR SAMPLING Sampling Meth Sample Flow F Sampling Time Volume Collect	nod	Collection M Special Ship #Field Blank	oping Instru	iampie Blanks

Nº 006019

Roy F. Weston, Inc.

REAC, Edison, N.J. EPA Contract 68-03-3482

=	Samplers: Site Name: Sample Loca d palustrine d rivernne rine		SURFA color _ odor _ flow _	ce water	STREAM width depth velocity pools riffles	REAC T. EPA Tas Project (ask Leader:sk Monitor: No.: BOTTOM rock slit rubble clay gravel orga	anic Br
SAMPLE TYPE stream/surface groundwater pond/lake brackish river ocean/saline effluent sediment sludge	DEVICE kemmerer trowl bucket sugar ekman	ponar other	SAMPLE I color odor temp DO cond	ORP salinity	depth	WEAT ambie barom relative weath	HER PARAME nt temp etric pressure e humidity	
ANALYSES TO BE PERFORM TOC required?Yes If No. explain Grain size analysis required? If No. explain ORGANICS A. halogenated & aromatic v B. volatiles-USEPA 624 C. trihalomethanes D. pesticides/PCB E. PC8 Dasa neutral/acid extracta G. pesticides, drinking water H. herbicides, drinking water INORGANICS	Yes	No		D. pH E. alkalinity F. hardness G. total disse H. total susp I. suifate	ide nol n hydrocarbons		AMPLE PREP. CONTAINER Lastic jar cetate core lastic bag lastic bucket L plastic TORAGE ret ice ry ice mbient HOASSESSME See attached	PRESERVATIVES HNO3 NaOH Zn Acetate HCL Na2SO4 other
A. metals, priority pollutant B. metals scan (ICP) C. metals, other RCRA A. EP toxicitymetals B. ignitability C. corrosivitypH D. reactivity COMMENTS: 6-3	posticid Yllow To	white pred.	ides S S	NIR SAMPLING Sampling Meth Sample Flow F Sampling Time Volume Collect	nod	Special S	n Media Shipping Instru	is
13 6-8': 6	rown cle gray-ste versote ster at	a clase of smell con	aand in Lad Whander	(u				

On drawing, but NOT in report A13 BI E13 E14 E 15 EII In report, but MOT on drawing 3 B-2 5 E012 E014

[EPA]

APPENDIX D SITE RECONNAISSANCE OF GORDON'S CREEK

- 5		
	S. Kirchoff Field Notes	
	5-25-94	5-25-94 Soll, vegetation
6th FMH	temp = 90°s, no rain for weeks clear > cloudy skies drought-like 2 weeks ago H2O in Gordons creek ~ 2 ft. higher; more leachate evident today compared to yesterday	Stratigraphic Column from Cut- Cank of Gordon's Creek See Photo No. 3 for Sand - clduinterface Widozu a creosate thickness arey compact varied is derise clay, some on v. fine grained edge of HzOtoo difficult to estimate a lift thick on Some outdrops to 2 tt.
	W. Pine St. →	

A CUM/INDI