DEC 2 1977

Mr. John Reinhardt Solid Waste Section Wisconsin Department of Natural Resources Post Office Box 7921 Madison, Wisconsin 53707

Dear Mr. Reinhardt:

Enclosed is the information concerning contaminated sediments and soils at the Kerr-McGee plant site in Milwaukee that I spoke to you about. I would appreciate you or your staff reviewing this data and evaluating some of the alternatives of disposal we discussed over the telephone on November 28, 1977. If you have any questions or comments do not hesitate to call me at (312) 353-2110.

Yours truly,

fillian A William H. Miner

ENVIRONMENTAL PROTECTION AGENCY Office of Enforcement EPA-330/2-77-022

THE POTENTIAL FOR POLLUTION OF THE LITTLE MENOMONEE RIVER FROM THE KERR-McGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN

(September-October 1977)

November 1977

National Enforcement Investigations Center - Denver

and

Region V - Chicago

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CONTENTS

The Average

Ĩ	INTRODUCTION 1	
II	SUMMARY AND CONCLUSIONS	
III	RECOMMENDATIONS 5	
IV	STUDY METHODS	
- V -	GEOLOGIC OBSERVATIONS AND ANALYTICAL RESULTS	
- .	•	
• •	TABLES	
1	Location of Wells Installed -	
1 2	Location of Wells Installed - Sept. 19-20, 1977	
•	Location of Wells Installed - Sept. 19-20, 1977	
2	Location of Wells Installed - Sept. 19-20, 1977	
2 3	Location of Wells Installed - Sept. 19-20, 1977 8 Transects Used in Locating Soil Sampling and Well Sites - Sept. 1977 . 9 Water Level Measurements in Wells Drilled - Sept. 19-20, 1977 12 Water Level Measurements in Wells and a Spring - Oct. 19, 1977	
2 3 4	Location of Wells Installed - Sept. 19-20, 1977	

FIGURES

Aerial View of Kerr-McGee/Moss-American Site . . 1 . 7

APPENDICES

A	Well Logs,	Casing	and	Perforation	Record
	and Field	l Notes	- 9,	/19-20/77	

B Chain-of-Custody Procedures C Soil and Water Sample Analyses

I. INTRODUCTION

Creosote contamination of the Little Menomonee River from the operation of a Kerr-McGee/Moss-American creosote plant near Milwaukee, Wisconsin has been a long-standing problem in the area. Severe contamination occurred in the past as a result of the direct discharge of wastewater and now, even though the plant has ceased operations, continued contamination from residual creosote deposits is of concern.

The National Enforcement Investigations Center (NEIC) of the Environmental Protection Agency (EPA) was requested by EPA's Region V to conduct an investigation of the plant site and recommend measures to eliminate further pollution of the River. This investigation was conducted in September and October of 1977. A review of the history of the site and the resulting pollution is presented in an earlier report by NEIC.*

Due to the highly technical and complex nature of the subject matter concerning creosote contamination of the Little Menomonee River, the Honorable Myron L. Gordon, U.S. District Judge (Eastern District, Wisconsin) appointed David A. Saichek, of Gaines and Saichek, to serve as Special Master in the matter of United States v. Kerr-McGee/Moss-American, Inc.^{**} All drilling, sampling and other field observations conducted by NEIC were done in accordance with requirements specified in a letter from the Special Master dated April 29, 1977 to Mr. Frank J. Daily, Attorney for the defendant, and Mr. Charles H. Bohl, U.S.

 National Enforcement Investigations Center, June 1977. Impact of Creosote Deposits in the Little Menomonee River, Wisconsin (April 1977). Denver: Environmental Protection Agency, 330/2-77-016, 47 p.

** Case No. 75-C-277.

Attorney. In addition, on-site field operations including well drilling, sampling, water level measurements, well casing removal, well backfilling and abandonment procedures were observed by the Special Master, the EPA Region V Attorney, and representatives of the defendant.

This report describes the field methods used during the September and October surveys. It also presents the results of the investigation and recommends measures to reduce or eliminate movement of creosote from the plant site into the Little Menomonee River.

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II. SUMMARY AND CONCLUSIONS

As requested by EPA, Region V, the National Enforcement Investigations Center conducted an investigation of the abandoned Kerr-McGee/Moss-American plant site near Milwaukee, Wisconsin. The objectives of the investigation were to determine the extent of creosote contamination in soil and groundwater, to evaluate the potential for continued pollution of the Little Menomonee River from this site, and to determine the measures necessary to eliminate the source of pollution.

During a site visit on September 19 and 20, 1977, six wells were drilled, casings were installed, groundwater levels were determined and water samples were collected from four of the wells for analysis [Appendix A]. In addition, core samples were collected at depths over 4.6 m (15 ft) during well drilling. Transects were laid out in relation to the wells and to structures in the area, and more than 100 soil samples were collected at depths up to 90 cm (3 ft) for later analysis.

Another site visit was made on October 19, 1977 to collect a second series of water samples from the wells and to remove the well casings. The wells were backfilled and the property was restored to its presurvey condition to the satisfaction of Company representatives present on the site.

All soil and water samples were returned to the NEIC under chainof-custody procedures [Appendix B] and analyzed for methylene chloride extractables [Appendix C]. The moisture content of the soil samples was also determined. The presence of creosote in the extracts was confirmed by gas chromatography on all of the extracts and the individual components of creosote were confirmed by mass spectrometry analyses of selected samples. Groundwater level at each well was measured during both visits. These measurements plus general observations of the area and a knowledge of geology and hydrology indicate that the groundwater flows toward the Little Menomonee River.

The presence of creosote in water samples from two wells during both site visits and from a third well during the October visit shows that the groundwater is contaminated with creosote.

Analyses of core samples and soil samples taken from the transects show that the entire area is contaminated with creosote, in some cases to a depth of at least 4.6 m (15 ft) below ground level. In one case, the concentration of methylene chloride extractables in the soil was greater than 27% and it was greater than 10% in several more samples.

Such widespread and high concentrations of creosote at the Kerr-McGee/Moss-American plant site make it evident that contamination of the Little Menomonee River will continue unabated for years if no efforts are made to control surface runoff and mobility of the creosote-contaminated groundwater.

The provision of an interceptor ditch, however, would prevent the contaminated runoff, groundwater and soil from entering the River. A granular carbon adsorption column may be necessary if the ditch effluent contains dissolved contaminants.

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III. RECOMMENDATIONS

The inflow to the Little Menomonee River of creosote-contaminated surface runoff, groundwater, and soil could be prevented by installing an interceptor ditch along the railroad track north of the site and between the contaminated area and the Little Menomonee River [Figure 1, Section IV].

The ditch should intercept the groundwater and surface runoff from the Kerr-McGee/Moss-American property. A system of skimmers and adsorption media (straw), placed in the ditch, would remove creosote from the interceptor discharge before it enters the Little Menomonee River. At 2.4 m (8 ft) deep, the ditch would collect any creosote floating on the water table and any soil washed off during a storm. what an Cart

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The ditch would have to be maintained regularly to remove accumulated skimmed material and solids that have settled.

If the water leaving the ditch is contaminated with soluble or emulsified components of creosote, it may be necessary to install a granular activated carbon column to treat the discharge from the ditch.

IV. STUDY METHODS

WELL AND SOIL SAMPLING SITE LOCATIONS

The investigated site consists of approximately $44,500 \text{ m}^2$ (11 acres) located in the NW 1/4, Section 8, T.8N., R.21E. near Brown Deer, Milwaukee County, Wisconsin [Figure 1]. From an aerial photograph of the site and an on-site inspection, the decision was made that six observation wells would be drilled to characterize the degree of contamination of the soil and the groundwater with creosote. One well was drilled near the northeast corner of a green shed close to the former retort area, and where it was assumed that the concentration of creosote would be the highest. Five other wells were drilled in relation to this "green shed" well, as described in Table 1; Figure 1 shows the approximate location of these wells. Because of surface characteristics (water, muck, etc.), it was not practicable to drill all of the wells at what was considered optimum locations.

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Once the wells were located, transects were established in relation to the wells and to structures in the area [Table 2]. The transects enabled the location of sites for collection of samples of soil from the surface and down to depths of 90 cm (3 ft) below land surface. The analysis of the samples collected allowed a more complete evaluation of the extent of contamination in the area.

The six observation wells were drilled on the eastern half of the Kerr-McGee/Moss-American plant site on September 19 and 20, 1977 using a truck-mounted continuous flight auger drill rig. In the field, a geologic log was prepared of each hole, including observations made by studying cuttings brought from the wells, noting rig behavior and taking drive samples.

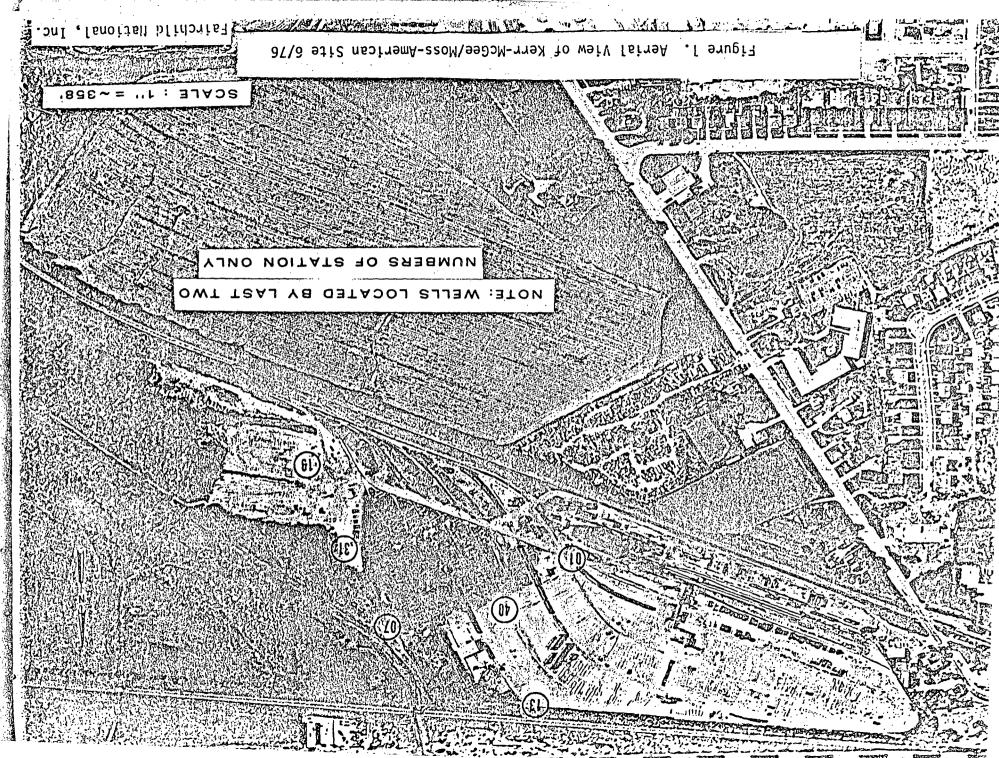


Table 1

MILWAUKEE, WISCONSIN September 19-20, 1977					
 Well	Location Description [†]				
772401	<pre>2.4 m (8 ft) from NE corner of the green shed, bearing 045°</pre>				
772407	226 m (740 ft) from well 772401, bearing 045°				
772419	315 m (1,035 ft) from well 772401, bearing 080°				
772413	177 m (582 ft) from well 772401, bearing 335°				
772440	About 91 m (298 ft) from well 772401 and 29 m ⁻ (96 ft) north of transect A				
772431	Halfway between wells 772407 and 772419 on transect F				

LOCATION OF WELLS INSTALLED AT THE KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN September 19-20, 1977

+ All bearings were taken with magnetic compass

Table 2

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TRANSECTS USED IN LOCATING SOIL SAMPLING AND WELL SITES KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN September, 1977

Transect	Description
A	Line connecting wells 772401 and 772407
В	Line connecting wells 772401 and 772413
Ċ	Line connecting wells 772407 and 772413
D	Line east from well 772413 and parallel to railroad tracks
E	Line connecting wells 772401 and 772419
F	Line connecting wells 772407 and 772419
G	Line parallel to and 1 m (3 ft) south of retorts
H	Line parallel to and 0.3 m (1 ft) north of north wall of retort building

9

A drive sample was collected by removing the auger from the well and replacing it with a "split-spoon sampler" on the bottom end of the drill rod. The split spoon was driven into the undisturbed earth at the bottom of the well by dropping a 59 kg (130 lb) weight. The number of blows required to drive split-spoon sampler 46 cm (18 in) deep was recorded at the time of sampling. These data are recorded in the drill logs.

After the split-spoon sampler was removed, the auger was reinstalled in the hole, and drilling continued until the desired depth was reached. Upon reaching this depth, the auger was removed from the hole and a section of 5-cm (2-in) diameter PVC^{*} pipe, capped on the end and perforated with hacksaw slots, was installed in the hole. Backfill around the casing was obtained from cuttings which had been withdrawn from the hole.

During the drilling of the green-shed well (772401), it was noted that cores obtained from 3.0 to 3.5 m (10 to 11.5 ft) and from 4.6 to 5.0 m (15 to 16.5 ft) had layers of creosote ^{**} filling fractures and partings. As drilling progressed to about 6 m (20 ft), the auger began to lift water to the surface. This water had globules of creosote floating on the surface. As the auger was withdrawn, it was noted that the bottom 4.6 m (15 ft) was not coated with clay as had been the experience in other holes on the site, but instead was coated with creosote. This apparently was the result of drilling through a layer of creosote which coated the auger and prevented any clay from adhering to the auger.

* Polyvinyl Chloride

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* Identified on the basis of odor and appearance

SAMPLING AND ANALYTICAL PROCEDURES

Well Samples

Upon completing the six observation wells, the depth to groundwater was determined, and water sampling proceeded. Depth to water, obtained with a Powers' electric well sounder, was recorded as depth below the undisturbed land surface [Table 3].

After the depth was recorded, a 4.6 m (15 ft) long, 2.5 cm (1 in) diameter suction hose was installed in the casing and well was pumped until dry or until suction was broken. This was done twice at each well, then water samples were collected in glass containers prerinsed with water from each well using a Homelight gasoline-powered water pump. Water could not be obtained from two of the wells (772407 and 772419) due to the tight clay which comprises the bulk of the formation.

During each pumping of the green-shed well (772401), the water was noticed to contain globules of creosote, and as the pump broke suction a creosote froth was produced.

The soil samples from the wells were taken by opening the splitspoon sampler and placing the lower portions of the core in chemically clean glass jars capped with Teflon-lined lids. The upper portion of the core, which included material that fell into the hole as the auger was being removed, was not saved as a part of the sample.

All samples, soil and water, were taken to the NEIC Laboratory for analysis. Chain-of-custody procedures were followed throughout.

On October 19, the site was again visited by NEIC to re-sample the wells and return the site to pre-survey conditions. Several of

WATER LEVEL MEASUREMENTS IN WELLS DRILLED AT THE
KERR-MCGEE/MOSS-AMERICAN PLANT SITE
MILWAUKEE, WISCONSIN
September 19 - 20, 1977

Table 3

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Well	Date (Sept			Depth to water bo (cm)	elow land surface (ft)
772401	20		#	• 142	4.7
772407	19	-		27	0.9
772419	20			.143	4.7
772413	20			33.5	1.1
772440	20			42.7	1.4
772431	20		`	79.2	2.6

the wells, all of which had been capped with a plastic cover, had been vandalized and the well casing was broken off at or just above the ground surface. As before, the depth to water level at each of the wells was measured, this time using a steel tape [Table 4]. Then the wells were purged to a depth of 4.6 m (15 ft) and left to recover before sampling.

Also as noted during the previous visit, the well nearest the green shed produced a black, greasy liquid during purging. Well 772407, which produced no water during the earlier attempt at sampling, produced a very muddy, thick liquid during purging.

During the October visit, samples were obtained by use of a handpowered diaphragm pump which was rinsed with acetone between sampling at each well. Well 772407 was difficult to sample, probably because of mud and clay blocking the slots in the casing. To get enough sample for analysis, it was finally necessary to use a hand-operated bailer. The sample contained about 1,000 ml of water and 900 g of clayey sediment.

The sample from the well nearest the green shed was brownish with an iridescent sheen on the surface. The sample was not as dark or greasy as the material removed when the well had been purged earler in the day.

At sampling site 772452, about 31 m (100 ft) southwest of well 772440, the discharge from a small spring flowed through a small clump of vegetation and then toward the Little Menomonee River. A small pond had formed which was partially covered by an oily, iridescent sheen. As the water came up out of the ground, a sample was taken.

Table 4

WATER	<i>LEVEL</i>	MEASUREMENTS	IN	WELLS	AND	A	SPRING	AT	THE	
	KEI	R-MCGEE/MOSS-	AMI	ERICAN	PLAI	٧T	SITE			
		MILWAUKEE	Z, 1	VISCONS	SIN					
		October	1	9, 1977	7					

Well	Depth to wate	r below land surface
	Cm	ft
772401	101.60	3.3
772407	36.48	1.0
772419	93.98	3.1
772413	36.83	1.2
772440	34.29	1.1
772431	76.20	2.5
772452	Sur	face
	-	

After sampling, all casings were pulled from the wells and removed from the site. The holes were backfilled with a mixture of sand and bentonite clay, and the survey stakes placed during the earlier site visit were removed.

Soil Samples

Soil samples were collected from sites on the transects described in Table 2. At each of the sites except the wells, a small coring device was used to collect a surface sample. An auger was then used to excavate a hole to an intermediate depth of 30 to 50 cm (1 to 1.6 ft) where another core sample was collected. The holes were then deepened with the auger to final depths varying between about 55 and 90 cm (1.8 to 3.0 ft) and again a small core sample was collected from the bottom of each hole.

Analytical Procedures

The soil and water samples were analyzed for methylene chloride extractables. In addition, the moisture content of the soils was determined. The methylene chloride extracts were also analyzed for the presence of creosote compounds by gas chromatography.

Individual components of creosote were also confirmed in selected samples by mass spectrometry. In all cases, unless otherwise indicated, most of the methylene chloride extractable material was creosote.

V. GEOLOGIC OBSERVATIONS AND ANALYTICAL RESULTS

The entire Kerr-McGee/Moss-American plant site is underlain by a thick section [at least 17 m (55 ft)] of plastic, clayey glacial till with a greater percentage of gravel material in the till section along the eastern side of the plant site near the Little Menomonee River. The till is locally overlain by a man-made fill and/or by a section of organic peat. In the Kerr-McGee/Moss-American plant site, the till showed evidence of horizontal openings filled with creosote to a depth of at least 4.6 m (15 ft) in the undisturbed core samples. Groundwater occurs in the till section throughout the entire area of the Kerr-McGee/Moss-American plant site investigated. In humid areas such as in the Milwaukee vicinity, the groundwater table configuration is characteristically a subdued replica of the land surface which, at the subject site, slopes north and east across the Kerr-McGee/Moss-American site toward the Little Menomonee River. Groundwater and soil samples obtained from the plant site indicated evidence of severe creosote contamination, especially in the area of the former retort building. As the groundwater and soil are known to contain creosote and the water table slopes toward the Little Menomonee River, any movement will be in this direction.

As reported in Table 5, the water samples collected from the observation well (772401) near the retort contained 4,910 mg/l of methylene chloride extractables in the September sample and 99 mg/l during the October sampling. Even higher concentrations of creosote were noted in the water standing in the well. The suction line from the well to the pump was coated with creosote adhering to the inside of the pipe. Cores obtained from the surface 0.5 m (0 to 1.5 ft) and from depths within the hole -- 3.0 to 3.5 m and 4.6 to 5.0 m (10 to 11.5 ft)

Table 5

ANALYTICAL RESULTS - WELL WATER SAMPLES COLLECTED AT THE KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN

Well		ethylene Chloride ept. 20, 1977 O mg/l	
772401	NE corner of green shed	4,190	99
772407	NE corner of area near RR	NS ⁺	600 ⁺⁺
772413	Northside center	ND ⁺⁺⁺	<2 ^{†††}
772419	SE corner of area	NS [†]	<2 ^{†††}
772431	Eastside center	ND ⁺⁺⁺	<2 ⁺⁺⁺
772440	Center of area	20	9
772452	Spring sample, ∿100 ft sout of 772440	hwest	<2 ⁺⁺⁺

* No sample obtained due to slow water level recovery after well development and purging.

- ++ Value rounded off to nearest 100 mg/l. This sample contained a considerable amount of very finely divided clay-like mud. An 1,800 ml slurry of this material was extracted. The slurry consisted of about 1,100 ml of water and 900 g of mud. The moisture content of the mud was 45%. A small amount of the methylene chloride extractables in this sample was creosote. The majority of the extractable material resembled a highly weathered heavy fuel oil.
- *** None detected above the detection limit of methylene chloride extractables or no identification as creosote by gas chromatography.

and 15 to 16.5 ft) -- contained large amounts (2,400 to 7,900 mg/kg) of methylene chloride extractables [Table 6].

It appears that there is a creosote layer, probably several inches thick, at shallow depth in the retort area. In addition, heavier fractions of creosote are apparently sinking through the groundwater to greater depths.

Much of the creosote may be temporarily sorbed on the soil material but a finite portion does move toward the Little Menomonee River. A sample of groundwater obtained from well 772440, near the center of the storage area, was found to contain 20 mg/l of methylene chloride extractables in the September sample and 9 mg/l during the October sample. A core sample, taken at a depth of 3.0 to 3.5 m (10 to 11.5 ft) from this well showed no detectable level of creosote. Creosote was not detected in water samples from any of the other wells except that during the October sampling, the mud-water mixture from well 772407 contained a high concentration of methylene chloride extractables which resembled highly weathered heavy fuel oil. A fraction of this extract was found to contain creosote components. This may be a result of the strong tendency for creosote to adhere to the well casing and to the pump suction line.

Analysis of the well core samples [Table 6] showed creosote at depths more than 4.6 m (15 ft) below the surface and concentrations of methylene chloride extractables as high as 23,700 mg/kg (2.37%). As previously discussed, this creosote was visible and detected by appearance and odor as the cores were removed from the split-spoon sampler. The creosote had evidently percolated through the soil and the groundwater surface to fill voids in the soil to at least the 4.6 m (15 ft) depth.

The presence of creosote from soil samples [Table 6] is randomly distributed throughout the area investigated. It reached a high of

Table 6

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ANALYTICAL RESULTS - SOIL AND WELL CORE SAMPLES AT THE KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN September 19-20, 1977

	Description	· · · · · · · · · · · · · · · · · · ·	Depth	% Moisture	Methylene Chloride Extractable (Dry Weight) - mg/kg
		STATION NO.			
Green shed,	well		0-1.5 ft 5-6.5 ft	8.7 24.2	2,50 <u>0</u> ND
•		*	10-11.5 ft	16.5	7,900
· · ·			15-16.5 ft	10.6	2,400
· ·	· ·	STATION NO.			
Transect.A,	105 ft from S	Station 772401		11.9	9,800
•		•.	62 cm	17.5	15,400
•	-		90 cm	14.9	1,700
Thomsont A	220 ft from (STATION NO. Station 772401		13.8	9,200
ITansect A,	220 FU FUN 3	Station //2401	Surface 35 cm	27.4	2,100
•	•		50 cm	35.5	11,100
		STATION NO	772404		t 1
Transect A	351 ft from 9	STATION NO. Station 772401		27.0	8,500
frunseet ny			32 cm	53.5	4,300
		•	82 cm	30.0	1,000
	, ·	STATION NO.	772405		6 ³
Transect A.	474 ft from S	Station 772401		39.0	5,900
			35 cm	51.3	13,100
		· ·	65 cm -	45.5	4,200
		STATION NO.			х •
Transect A,	597 ft from S	Station 772401		40.0	9,000
			Mid-Depth	64.5	31,800
	. ·		73 cm	60.4	3,800
		STATION NO.	772407	-	
NE corner o	f site, well		0-1.5 ft	20.1	23,700
-		•	5-6.5 ft	22.1	ND ⁺
			10-11.5 ft	12.1	NU

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Description Station 772408 begin mark Methylene Chloride Extractable (Dry Weight) mg/kg Transect B, 145 ft from Station 772401 Surface 50 cm 7.8 12.6 4,400 69 cm 4;500 4,400 69 cm Transect B, 291 ft from Station 772401 Surface 9 cm 7.5 800 4,800 4,800 4,800 4,800 4,800 69 cm 800 4,800 4,800 4,800 4,800 69 cm Transect B, 436.5 ft from Station 772401 Surface 8 cm 7.5 800 4,800 68 cm 800 4,800 4,300 Transect B, 436.5 ft from Station 772401 Surface 8 cm 16.0 5,300 3,100 68 cm 5,300 3,100 10-11.5 ft 16.0 1,500 North side, well STATION NO. 772410 10-11.5 ft 16.6 16.6 1,700 ND 12 ft north of Transect E, 207 ft from Station 772401 Surface 5.5 cm 18.2 34,300 12 ft north of Transect E, 414 ft from Station 772401 Surface 6.0 cm 12.2 12,900 12 ft north of Transect E, 611 ft from Station 772401 Surface 6.0 cm 12.0 20,000 12 ft north of Transect E, 611 ft from Station 772401 Surface 8.0 12.0 20,000 40 cm 15.1 5,300 53.00	•	· · · ·		
Transect B, 145 ft from Station 772401Surface 7.84;50050 cm 12.64;40050 cm 12.64;500STATION NO. 772401Surface 7.5800Transect B, 291 ft from Station 772401Surface 7.580040 cm 22.76,300G cm 21.65,300Transect B, 436.5 ft from Station 772401Surface 16.05,300Transect B, 436.5 ft from Station 772401Surface 16.05,300STATION NO. 772410Surface 16.05,300STATION NO. 772413STATION NO. 772413STATION NO. 772413STATION NO. 77241412 ft north of Transect E, 207 ftSTATION NO. 772414Station 772401Surface 13.371,00038 cm 20.510,40055 cm 18.234,300STATION NO. 772415ft north of Transect E, 414 ftSurface 12.212,90050 cm 24.4ND ⁺ 60 cm 22.6ND ⁺ Station 772401Surface 12.020,000	Description	Depth		Chloride Extractable (Dry Weight)
Transect B, 291 ft from Station 772401Surface7.580040 cm22.74,80069 cm21.64,300Transect B, 436.5 ft fromSTATION NO. 772410Surface16.0Transect B, 436.5 ft fromSTATION NO. 772401Surface16.042 cm29.53,10068 cm29.81,500North side, wel1STATION NO. 7724135-6.5 ft21.012 ft north of Transect E, 207 ftSurface13.371,00012 ft north of Transect E, 207 ftSurface13.371,00012 ft north of Transect E, 414 ftSurface12.212,90050 cm24.4ND [†] 60 cm22.6ND [†] 12 ft north of Transect E, 611 ftSurface12.020,000		Surface 50 cm	12.6	4,400
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North side, well $\frac{5-6.5 \text{ ft}}{10-11.5 \text{ ft}} 21.0 \\ 10-11.5 \text{ ft}} 16.6 \\ \frac{\text{STATION NO. 772414}}{12 \text{ ft north of Transect E, 207 ft}}$ 12 ft north of Transect E, 207 ft $38 \text{ cm} = 20.5 \\ 10,400 \\ 55 \text{ cm} = 18.2 \\ 34,300 \\ 12 \text{ ft north of Transect E, 414 ft}}$ 12 ft north of Transect E, 414 ft $50 \text{ cm} = 24.4 \\ 60 \text{ cm} = 22.6 \\ ND^{\dagger} \\ 60 \text{ cm} = 22.6 \\ ND^{\dagger} \\ 12 \text{ ft north of Transect E, 611 ft}}$ 12 ft north of Transect E, 611 ft $50 \text{ cm} = 12.0 \\ 20,000 \\ 12 \text{ ft north of Transect E, 611 ft}}$		01 Surface 42 cm	29.5	3,100
12 ft north of Transect E, 207 ft from Station 772401Surface 13.371,00038 cm 20.510,40055 cm 18.234,30012 ft north of Transect E, 414 ft from Station 772401Surface 12.212,90050 cm 24.4ND ⁺ 60 cm 22.6ND ⁺ 12 ft north of Transect E, 611 ft from Station 772401Surface 12.020,000	North side, well	5-6.5 ft 10-11.5 ft		
55 cm 18.2 34,300 STATION NO. 772415 12 ft north of Transect E, 414 ft Surface 12.2 12,900 12 ft north of Transect E, 414 ft Surface 12.2 12,900 50 cm 24.4 ND ⁺ 60 cm 22.6 ND ⁺ STATION NO. 772416 12 ft north of Transect E, 611 ft Surface 12.0 20,000	12 ft north of Transect E, 207 ft		13.3	71,000
STATION NO. 772415 12 ft north of Transect E, 414 ft Surface 12.2 12,900 50 cm 24.4 ND ⁺ 60 cm 22.6 ND ⁺ STATION NO. 772416 12 ft north of Transect E, 611 ft from Station 772401 Surface 12.0 20,000		38 cm	20.5	10,400
12 ft north of Transect E, 414 ft Surface 12.2 12,900 50 cm 24.4 ND ⁺ 60 cm 22.6 ND ⁺ STATION NO. 772416 12 ft north of Transect E, 611 ft Surface 12.0 20,000		55 cm	18.2	34,300
from Station 772401 Surface 12.2 12,900 50 cm 24.4 ND ⁺ 60 cm 22.6 ND ⁺ STATION NO. 772416 12 ft north of Transect E, 611 ft Surface 12.0 20,000		772415		
60 cm 22.6 ND [†] STATION NO. 772416 12 ft north of Transect E, 611 ft from Station 772401 Surface 12.0 20,000	12 ft north of Transect E, 414 ft from Station 772401	Surface	12.2	12,900
STATION NO. 772416 12 ft north of Transect E, 611 ft from Station 772401 Surface 12.0 20,000		50 cm	- 24.4	ND
12 ft north of Transect E, 611 ftfrom Station 772401Surface12.020,000		60 cm	22.6	ND [†]
from Station 772401 Surface 12.0 20,000		772416		
40 cm 15.1 5,300		Surface	12.0	20,000
	·	40 cm	15.1	5,300

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21

•	Table 6	(Continue	ed)	
KERR_M	GEE/MOSS	-AMERICAN	PLANT	SITE

STATION NO. 77241 12 ft north of Transect E, 808 ft from Station 772401	1 <u>7</u> Surface 30 cm	4.9	- 1,300
			- 1,300
	30 ст		1,000
. · · · · · · · · · · · · · · · · · · ·		11.5	2,400
	60 cm	13.9	2,200
STATION NO. 77241	9		· +
)-1.5 ft	17.7	
	5-6.5 ft		ND ND
	D-11.5 f	τ 14.6	ND
STATION NO. 77242	20		
	Surface	18.3	40,200
	50 cm	22.7	19,300
	75 cm	18.5	13,600
STATION NO 77240	10		
STATION NO. 77242 Transect H, Midpoint of retort bldg			
71 ft from Station 772401	Surface	17.1	10,500
	47 cm	21.3	3,800
	70	04 4	· .
	72 cm	24.4	2,200
STATION NO. 77242	22		
	Surface	8.1	4,900
	33 cm	21.1	6,200
	73 cm	-22.9	2,200
STATION NO. 77242) 2		
STATION NO. 77242 Transect G, Midpoint between 5th &	23		
	Surface	13.0	41,900
	40 cm	19.7	2,900
	70	00 F	12 000
	72 cm	23.5	13,900

•••

Description	Depth	% Moisture	Methylene Chloride Extractable (Dry Weight) mg/kg
STATION NO. 7	772424		
Transect G, Midpoint of 4th retort from left on south facing side	Surface	11.9	- 92,700
	38 cm	25.8	127,000
	63 cm	25.9	36,400
STATION NO. 7 Transect G, Midpoint of 2nd retort	772425		•
from left on south facing side	Surface	2.4	104,000
	38 cm	43.2	133,000
	. 70 cm	38.7	89,900
STATION NO.	772426		
<pre>Inside retort room, 40.3 ft from NE corner, 2 ft from north wall</pre>	Surface	5.0	72,900
	70 cm	26.0	, 7,100
STATION NO.	772427		
Inside retort room, 20 ft from NW corner, 4 ft in from north wall	Surface	3.3	279,000
	60 cm	20.2	5,600
STATION NO. Transect F, 168 ft from Station 772419		8.4	2,400
	52 cm . 60 cm		4,100 ND
STATION NO. Transect F, 336 ft from Station 772419		9.9 41.0 38.5	1,900 2,8 <u>0</u> 0 ND

Description	Depth	% Moisture	Methylene Chloride Extractable (Dry Weight) mg/kg
Eastside center, well	7 <u>2431</u> 5-6.5 ft 10-11.5 f		ND - ND
STATION NO. 77 Transect F, 504 ft from Station 772419	72432 Surface 36 cm 58 cm	27.0 50.8 47.2	11,200 35,8 <u>0</u> 0 ND
STATION NO. 7 Transect D, 75 ft from Station 772413	72433 Surface 40 cm	24.2 31.8 41.8	8,8 <u>0</u> 0 ND <u>:</u> ND
STATION NO. 7 Transect D, 168 ft from Station 772413	72434 Surtace 40 cm 70 cm	23.3 32.2 50.0	9,100 1,700 3,700
STATION NO. 7 Transect D, 261 ft from Station 772413	72435 Surface 40 cm 70 cm	33.3 50.8 55.3	19,700 1,000 1,000
STATION NO. 7 Transect D, 354 ft from Station 772413	72436 Surface 50 cm 80 cm	30.2 51.0 58.5	10,200 1,100 15,600
STATION NO. 7 Transect C, 152 ft from Station 772407	72437 Surface 32 cm 65 cm	26.0 74.5 73.6	8,600 11,000 2,800
STATION NO. 7 Transect C, 304 ft from Station 772407	72438 Surface 52 cm 70 cm	25.0 59.9 58.5	1,900 3,900 3,100

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23

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Description	Depth	% Moisture	Methylene Chloride Extractable (Dry Weight) mg/kg
STATION NO.	772439	······································	-
Transect C, 456 ft from Station 77240		14.9	900
	38 cm	38.9	25,300
	70 cm	60.4	21,100
STATION NO.	772440		
Center of site, well	10-11.5	ft 13.5	ND ⁺
STATION NO.	772441		
50 ft at 280° magnetic from a point in the RR ditch 23 ft	Surface	21.9	1,600
downstream of the confluence with the dock ditch	37 cm	24.2	1,100
	72 cm	34.6	1,400

 ND - None detected above the detection limit of 500 mg/kg or no identification as creosote by gas chromatography. 279,000 mg/kg (27.9%) in a sample taken inside the retort room and was greater than 100,000 mg/kg (10%) at depths up to 38 cm (15 in) below the ground surface at sites outside the buildings.

The absence of creosote in some of the water samples collected and the relatively low concentrations found in wells other than the one nearest the green shed may be an indication of the low solubility of creosote or the strong tendency for creosote to adhere to the well casing and to the pump suction line. It may also be an indication that the wells were placed at the edge of a plane of higher concentration of contaminants. This latter possibility is thought most likely because the wells could not be drilled in optimum locations due to poor surface conditions which precluded ready access.

The high concentrations of creosote found in the groundwater and in the soil at all levels, particularly at the surface, indicate that the pollution of the Little Menomonee River will continue if corrective measures are not taken. It seems likely that a ditch dug along the north and east boundaries of the study area would intercept both groundwater and surface runoff. The creosote thus intercepted could be removed by skimming and sediment removal prior to release of the water to the River. If soluble materials in the water orove to be a problem, it may be necessary to treat the water with activated carbon prior to discharge.

APPENDIX A

WELL LOGS, CASING AND PERFORATION RECORD AND FIELD NOTES 9/19-20/77

APPENDIX A

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WELL LOGS, CASING AND PERFORATION RECORD AND FIELD NOTES, 9/19-20/77

KEY TO WELL LOGS



Fill, Cinders, gravel, clay and silt Till, gray, clayey silt, peat, vari-colored clay Gravel and sand, dense, sandy Peat, dark gray Total Depth

Record of casing depth and perforations.

 $\mathbf{\Sigma}^{\mathsf{f}}$ Record of surface of water level. Other notes on field logs are self-explanatory.

A-2 EPA Na Jual Ellivico Drill Log Amer Project: K-M Moss Project Number: Location: Actor of 212 and of eastern rr tro Hole Musber: 772407 Drill Crew: Soil Testing Servi- Geologist: Rouse Drill Hethod: CF Auger Date: 9/19/1977 Depth Fr Water Description Log Fills Cinclers, Clay, dark gray 31/18" -3.5.7.cl.al $\Delta = \Delta V$ 2 Till; Groy, clugey, silty, gravelig inc. filled 24/i8" with grovel, round to subrud in Peat; durk gray, wet Samplit 10 Sample 3 E 0.0,0 Derise duilling 0000 20 0000 00 30 Pense, rocks 0 40 0. Z 'n 50 1110 TP 551 60 Note; Hole cured, wouldn't Csy Moved 3'South Fo hold for csy, Drilled 251, 49 Moved 3' forther seath, drill. 50+ 81 CS9. Water @ 0.91 Refurned in p.m 70 North, drilled Zl .

EPA Na Jnal Enforcement Investigations lenter A-3 Drill Log Project: Stan Korn Mc Gei Moss Amer t Number: Location: SECor, line w/ east r. r, & south rel. (Airchote imber: 7724 19 Geologist: Rouse Hethod: C.F. Auger Drill Crew: 375, 1-c i Date: 9/19/1977 Log Water Description F+ F. II, grouil, cloy Δ 1 Till, Tun, clayey, silty, growelly, densy. with occil dense ground leaves 10 Water was 4.7' below 1. son 15 4/20/170 00 1000 20 - a 000 25

EPA Na Snal Enforcement Investigation. enter A-5 Drill Log Project: Kerr McCee Location: Nof rorthern ect Number: Number: 772413 track rr Geologist: Rouse Method: CFA-yer Drill Crew: STS, Inc Date: 9/19/1977 1, F7. Log Water Description Fill, ruilroud grods Hip ロ Peut, dark gray, soft, with roots 11 5 Clay, gray, plug fic becomes Till, clugey, silts, slight grovel, plastic + - + + 0 95- 5 g 10 5 21 15 1. 20 20 25 Break in scale, 35 .45 Group loyer 1 deusers andy 0.0 2 501 Water is 1.10 below 1.5 on 0930 3 9/20/17 TD= 50' 68

EPA Na onal Enforcement Investigation. Jenter A-4 Drill Log Rerr-mebee Project Number: Project: Location: 81 Not NF Cor, gree - shed Hole Number: 772401 Geologist: Rovs, Date: 9/19/1977 Drill Method: CFAuger Drill Crew: 575,100 Depth, Ff Log Water Description Fill gravel, clay Sumple i × D Till, gruyto green, clurry, slightly gravelly, d 10 in-pie 21 Creasote 14 Sample 3 26 Blown Creosoft in spop inployer 30 Blow, 20 Auger brings up water up creosote globules Bottom 151 of auger cleaned of during refreat of auger, creosote coating 25 25 Water 4.65' below 1.5. 0" 9/20/77@ 0920 Collected 2 gill sample From wella

EPA Na onal Enforcement Investigation. ·A-6 Drill Log Kerv McGee Project Number: Project: Location: ACOrox 75'N of 773404 Drill Crew: 575, 14- Geologist Hole Number: 772440 Geologist: Rous Drill Method: GF Auger Date: 9/19/197 Depth, FF Log Water Description! Fill, grovel, Glay, soft Till, cluyeg, silty plustic, gray. jD Blow 15 Gravil, clayer, dease 00,00 000ð 25 20 TPZO Water was 1.4' below 1.5.04 9/20/77/0925 collected less then 1/2 gal @ ن[:] ز

EPA Na..onal Enforcement Investigation. Jenter A-7 Drill Log Project: Kerr - MicGee Moss Ame -Location: East give, craiter (Airph ect Number: Number: 772431 (Airphite) Method: CFAuge Drill Crew: 517 5, 142 Geologist: ROUST Date: 9/2011977 Description 1,Ff Log Water Fill grovely clayer 00 Feat, durk gray V Till, gray, clayeg, silty, plastic with rush yru - I luyers, as show ĺŨ Jar, 15 25 Druse drilling ingravel -25 3) TD. 30 Water wes 2.6 1 below 1.5. on 9/20 @ 0950 30 Collected water sur ple (2 gal) (a 1320

APPENDIX C

SOIL AND WATER SAMPLE ANALYSES

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CONFIRMATION OF CREOSOTE IN WATER AND SOIL SAMPLES FROM THE KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN BY GAS CHROMATOGRAPHY/MASS SPECTROMETRY

Analyses of the water extract and soil extract from Kerr-McGee confirming creosote were completed. The samples were screened initially on the flame ionization gas chromatograph, then analyzed utilizing combined gas chromatography/mass spectrometry (GC/MS). The gas chromatograms readily confirmed the presence of creosote when the chromatogram of the standard was compared to the water sample extract and the soil extract. The major sample components in samples and standard all occurred in the same ratio. The mass spectra of the individual compounds in the standard creosote mix and standard spectra of individual compounds run using the same instrumental conditions as the samples were perfect matches and all were confirmed by mass spectrometry and matching gas chromatography retention times. Table I gives the results for the samples and the standard mix.

TABLE I

	GC/MS Identification			
	Commercial	Sample	Sample	
	Creosote	2311	2240	
Compound	Local	Soil	Water	
Napthalene	CF(1)	CF	CF	
2-methylnaphthalene	CF	CF	CF	
<pre>l-methylnaphthalene</pre>	CF	CF	CF	
Biphenyl	CF	CF	CF	
1,3-dimethylnaphthalene	CF	CF	CF	
Acenaphthene	CF	CF	CF	
Dibenzofuran	CF	CF	CF	
Fluorene	CF	CF	CF	
Anthracene	CF	- CF	CF	
(Phenanthrene)	CF(2)	CF	CF	
Carbazole	CF	CF	CF	
2-methylphenanthrene	Good (3)	Good	Good	
Fluoranthene	CF	CF	CF	
Pyrene	CF	CF	CF	

Note: CF - confirmed

- These compounds confirmed by GC/MS identification, GC/MS of standards, and coincidence of retention times with those of standards.
- (2) Phenanthrene and anthracene have identical mass spectra and the gas chromatography retention times are only slightly different, with phenanthrene eluting later. It appears as a definite shoulder on the anthacene peak with the GC conditions used when more dilute solutions are run. Both compounds are obviously present.
- (3) This compound was identified by mass spectrometry only. A standard was not available.

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ANALYTICAL METHODOLOGY - SOIL, WELL CORE AND GROUNDWATER SAMPLES COLLECTED SEPTEMBER 19-20, 1977 AT THE KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN

On September 23, 1977, 6 water and 113 soil samples were received at the NEIC Laboratory for creosote analysis. All samples were handled according to chain-of-custody procedures. The waters and soils were analyzed for methylene chloride extractables and the soils for moisture content. The methylene chloride extracts were also analyzed for the presence of creosote compounds by gas chromatography. The results are summarized in Tables 5 and 7 in the body of the report.

Methodology:

A. Moisture

About 10 grams of thoroughly mixed sample was accurately weighed in a tared 50 ml beaker and dried overnight in an oven at 105°C. The water loss was determined by reweighing the cooled and desiccated beakers. Calculations:

Wt. of water loss x loo = % moisture
Wt. of sample wet

B. Methylene Chloride Extractables

Soils - 10 grams of thoroughly mixed soil was weighed into a 150 ml beaker. Large stones, twigs, leaves, etc. were avoided. 30 grams of granular anhydrous sodium sulfate was added to the beaker and the soil and sodium sulfate were mixed thoroughly to obtain a coarse granular consistency. The mixture was then transferred to a 33 x 80 mm cellulose extraction thimble and placed in a Soxhlet extractor. 200 ml of methylene chloride was placed in a 500 ml flat-bottomed flask and attached to the extractor. The extractor was allowed to cycle for 2 to 3 hours, with a rate of about 10 cycles per hour. Each flask was then placed on a rotary evaporator and the solution was concentrated to a volume of about 20 ml. The remaining solvent was quantitatively transferred to a tared 50 ml beaker and evaporated to dryness on a warm hot plate under a gentle stream of carbon-filtered air. Each beaker was reweighed and the residue determined. Results were calculated on a dry weight basis using % moisture values.

Wt. of residue in mg x 1000 =
Wt. of sediment extracted wet (gm) x wt fraction solids
mg/kg dry basis
wt fraction solids = 1.00 - wt fraction moisture

C-3

Waters - From 100 ml to 3 liters of the water samples were extracted twice with 100 ml of methylene chloride. The extracts were combined and evaporated to dryness in tared beakers on a warm hot plate under a gentle stream of carbon-filtered air. Each beaker was reweighed and the residue determined.

C. Creosote Compounds

Creosote is a complex mixture of compounds, some of which can be detected on a flame ionization gas chromatograph. These compounds give a distinctive pattern of peaks. This pattern was used to identify the presence of creosote.

The methylene chloride extracts were dissolved in 10 ml of acetone and an aliquot was injected on a Hewlett-Packard 5700A gas chromatograph equipped with an automatic sampler. Chromatograms were compared with a "standard" creosote sample that was purchased locally. The standard creosote sample was labelled as containing 60% creosote by weight. The sample was cut with a light refined petroleum-based oil.

Some samples with methylene chloride residues as high as 7,000 mg/kg showed no creosote pattern by gas chromatography. These residues could be caused by either "natural background" material, noncreosote compounds which have been extracted by methylene chloride, or creosote compounds which are not chromatographable. The samples which contained detectable amounts of methylene chloride residue, but which exhibited no creosote pattern, were reported as None Detected.

ANALYTICAL METHODOLOGY GROUNDWATER SAMPLES COLLECTED OCTOBER 19, 1977 AT THE KERR-MCGEE/MOSS-AMERICAN PLANT SITE MILWAUKEE, WISCONSIN

On October 21, 1977 seven plant site well samples were delivered to the laboratory in accordance with NEIC chain-of-custody procedures. The samples were analyzed for total methylene chloride extractables and the extracts screened by flame ionization detection - gas chromatography (FID-GC) to determine the presence of creosote material.

FID-GC analysis showd that samples from Stations 772413, 772419, 772431, and 772452 did not contain any observable creosote material. All other samples contained observable amounts of creosote. The results of these analyses are presented in Table 5 in the body of the report.

Methodology:

A. Moisture

About 10 grams of thoroughly mixed sample was weighed in a tared 50 ml beaker and dried overnight in an oven at 105°C. The water loss was determined by reweighing the cooled and desiccated beakers. Calculations:

Wt. of water loss x 100 = % moisture
Wt. of sample wet

B. Methylene Chloride Extractables

Waters - From 1,000 ml to 3.5 liters of the water samples were extracted twice with 100 ml of methylene chloride. The extracts were combined and evaporated to dryness in tared beakers on a warm hot plate under a gentle stream of carbon-filtered air. Each beaker was reweighed and the residue determined.

C. Creosote Compounds

Creosote is a complex mixture of compounds, some of which can be detected on a flame ionization gas chromatograph. These compounds give a distinctive pattern of peaks. This pattern was used to identify the presence of creosote.

The methylene chloride extracts were dissolved in 10 ml of acetone and an aliquot was injected on a Hewlett-Packard 5700A gas chromatograph equipped with an automatic sampler. Chromatograms were compared with a "standard" creosote sample that was purchased locally. The standard creosote sample was labelled as containing 60% creosote by weight. The sample was mixed with a light refined petroleum-based oil.