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Bainbridge Island, Washington**

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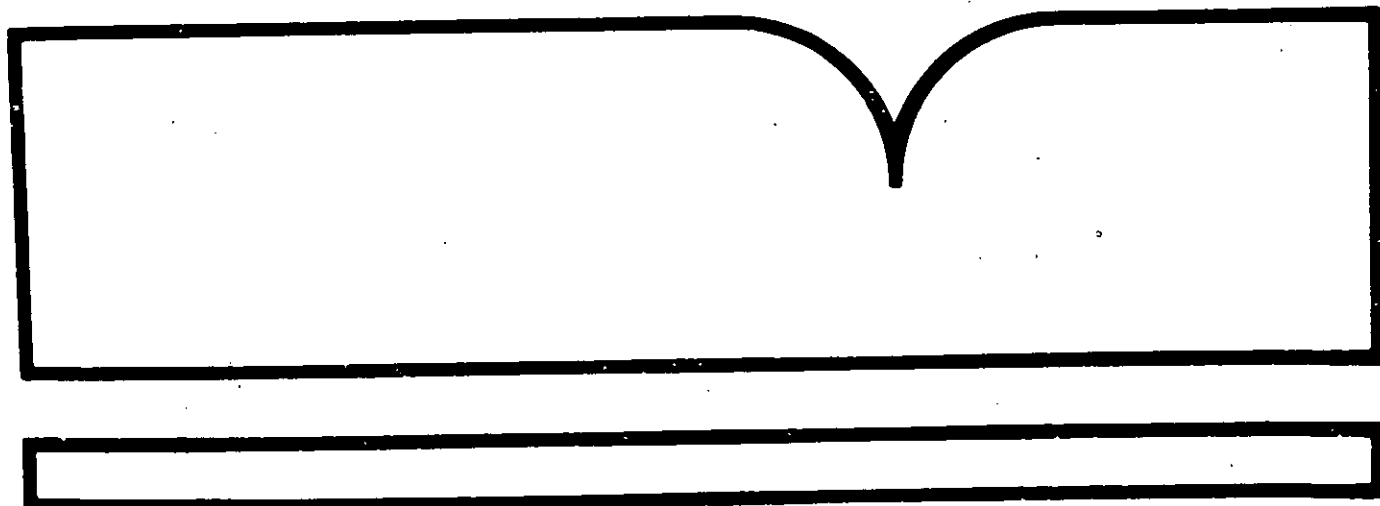
Emissions Test Report Air Toxics
Sampling at Wyckoff, Inc.
Bainbridge Island, Washington

Engineering-Science, Fairfax, VA

Prepared for

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AIR TOXICS SAMPLING AT WYCKOFF, INC.
BAINBRIDGE ISLAND, WASHINGTON

Submitted to

U.S. Environmental Protection Agency
Region X
1200 Sixth Avenue
Seattle, Washington 98101

Property of the U.S. Environmental Protection Agency

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March 1986
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Fairfax, Virginia 22030

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ENGINEERING-SCIENCE

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CHAPTER 1

INTRODUCTION AND SUMMARY

From July 22 to August 2, 1985, ES sampled five sources at two plants in the Seattle, Washington area to collect data on emission of toxic compounds. This report discusses the results of sampling two vacuum exhausts and the fugitive emissions from two retorts at Wyckoff, Inc.'s, creosote treating plant.

CHAPTER 2

PROCESS DESCRIPTION

Wyckoff, Inc., operates a log-treating plant on Bainbridge Island which is devoted to pentachlorophenol and creosote pressure-treating of logs for utility poles and marine pilings. The equipment for treating the logs is the same regardless of the preservative being used, although operating parameters are somewhat different.

Wyckoff has six retorts (five of which are operable) connected to two vacuum systems. In one system, one retort is used solely for penta treating, and one for either penta or creosote. The other system (three retorts) is dedicated to creosote treating. Each retort is a steam-jacketed insulated cylinder 130 feet long with a 90-inch inside diameter. Logs are debarked, loaded onto trams, and placed inside a retort. The door to the end of the retort is closed and bolted shut and the treating cycle commenced.

The treating process consists of several major phases and some minor ones, and there are general operating standards established by the Douglas Fir Association. The actual operating conditions and inclusion or exclusion of certain steps are determined by the operating personnel. The sampling plan was predicated on an incomplete understanding of the standard cycle and was not appropriate for the way the process is operated, at least at this facility. The following paragraph discusses the process as operated at Wyckoff.

The first step after bolting the door is evacuating the retort to 22" Hg which takes 45-60 minutes. The retort is then filled, via vacuum, with preservative and heated to 210°F with the steam jacket. The contents are held at 210° and 22" vacuum for about 48 hours. This conditioning phase removes water from the logs and the phase is ended when all the water has been extracted, as measured by periodic gauging of the collected water. After conditioning, the vacuum is released and the retort emptied, refilled with cold preservative, and then emptied again. This phase takes about three hours, and then the retort is pressurized with 30 psig air for about an hour. The retort is again filled (1-1/2 - 2 hours) with preservative and the pressure-treating begun. The bath is heated to 240°F and kept under pressure for about two hours. An optional expansion bath, lasting 1-2 hours if used, follows the pressure-treating. The retort is again pumped out (one hour) and live steam injected for two hours. The last step is called final vacuum and removes residual preservative and allows the load to cool down. When the temperature drops below 175°F, the door can be opened and the load removed.

Normally, the door is open for about 30 minutes while the load is pulled out and a new load put in. Copies of the process records for the loads handled while sampling was being performed are included in Appendix A.

The initial plan for sampling envisioned that vacuum vent emissions occurred during several phases because the retort was placed under vacuum during these times. However, this vacuum was used as the motive force for filling the retort with liquid and thus there was no discharge to the atmosphere as a result. Observations of the process and additional discussions with plant management and operating personnel confirmed that significant emissions occurred throughout the entire conditioning time and during final vacuum and not during other phases of the heating process. These two phases were the ones sampled.

The pollutants of concern are pentachlorophenol (C_6HCl_5), creosote compounds and PNAs. The specific compounds that comprise creosote are PNAs and typically include 2-4% each of acenaphthene, fluorene, diphenylene oxide, anthracene, carbazole, 12-14% phenanthrene and naphthalene. The remainder is unspecified PNAs. Penta consists of 5% pentachlorophenol in a light fuel oil (equivalent to #2 diesel fuel).

CHAPTER 3

TEST SCHEDULE AND SAMPLING LOCATIONS

As previously discussed, emission samples were taken from the vacuum vents when emissions were being vented, and from the open retort during load changes. Separate samples were taken from creosote and from penta treating. Each test involved collecting duplicate samples with paired sampling trains. Figures 3.1 and 3.2 portray the general arrangements for vacuum vent and retort fugitive sampling.

Table 3.1 provides a chronological summary of the test runs. As mentioned in Chapter 2, initially, samples were to be collected during other phases of the treating cycle. Observation of the process and further clarifications of the process by plant management led to the conclusion that the emissions during these other phases were negligible and therefore were not sampled. Copies of the field data are included as Appendix B.

TABLE 3.1

SAMPLING CHRONOLOGY

Date	Run Code	Sampling Time	Sample Type	Retort	Process Cycle
7/26/85	WCF-X-1A	1248-1319	NIOSH-XAD	D	Creosote fugitives
	WCF-X-1B	1248-1319	NIOSH-XAD	D	Creosote fugitives
7/29/85	WCV-WMS-1A	1235-1435	WMS-XAD	E	Penta-conditioning
	WCV-WMS-1B	1230-1430	WMS-XAD	E	Initial vacuum
7/31/85	WCV-WMS-2A	0927-1106	WMS-XAD	E	Penta-cool down
	WCV-WMS-2B	0922-1102	WMS-XAD	E	Final vacuum
7/31/85	WPF-X-1A	1116-1209	NIOSH-XAD	E	Penta fugitives
	WPF-X-1B	1116-1209	NIOSH-XAD	E	
7/31/85	WCF-X-2A	1337-1410	NIOSH-XAD	D	Creosote fugitives
	WCF-X-2B	1337-1417	NIOSH-XAD	D	
7/31/85	WCV-WMS-1A	1700-1830	WMS-XAD	C	Creosote-conditioning
	WCV-WMS-1B	1705-1825	WMS-XAD	C	Initial vacuum
8/1/85	WCV-X-2	0808-0828	NIOSH-XAD	C	Creosote-conditioning End of vacuum
8/1/85	WCV-WMS-2A	1800-1900	WMS-XAD	C	Creosote-cool down
	WCV-WMS-2B	1805-1900	WMS-XAD	C	Final vacuum

CHAPTER 4

SAMPLING AND ANALYTICAL PROCEDURES

The sampling and analytical procedures for this project are derived from EPA's "Test Methods for Evaluating Solid Waste" (SW-846, July 1982); "Sampling and Analysis Methods for Hazardous Waste Combustion (Draft)" (A.D. Little, Inc., February 1983); "NIOSH Manual of Analytical Methods", U.S. DHEW, NIOSH, Publication #75-121; and Title 40, Part 60, Appendix A, Code of Federal Regulations. The laboratory results are contained in Appendix C of this report.

SAMPLING PROCEDURES

Modified Method 5 - General

The MMS sampling train was used to collect organic compounds with boiling points between 100°C and 300°C such as products of incomplete combustion (PICs), PNAs, PCP and creosote compounds.

Extracts, rinses, and related materials from the sampling system for each sampling period were analyzed to quantify the specific compounds of interest for each test. A total of 13 runs were conducted.

Sampling Train

The gas stream was directed to a modified EPA Method 5 train, depicted in Figure 4-1. This system consists of the following components in series: nozzle, probe, heated particle filter, one sorbent module, a bank of impingers, and a meter box. A detailed discussion of the physical construction of the unmodified Method 5 train and its assembly is given by Martin; maintenance is given by Rom (APTD-0581, APTD-0576).

A glass fiber (particle) filter was enclosed in a glass housing and supported by a stainless steel mesh. This section of the train was contained in an electrically heated box with manual temperature regulation to maintain a sample gas temperature of about 120°C.

A typical sorbent module/condenser is also depicted in Figure 4-1. There are separate sections for cooling the incoming gas stream and for trapping organic gas constituents. Cooling was accomplished by routing the gas through a coil of glass tubing surrounded by water circulated from an ice and water bath. Organic constituents of the gas are removed by 20/40 mesh XAD-2 resin contained in an all-glass trap. High collection efficiencies (90-100%) are typical for vapor phase organic species

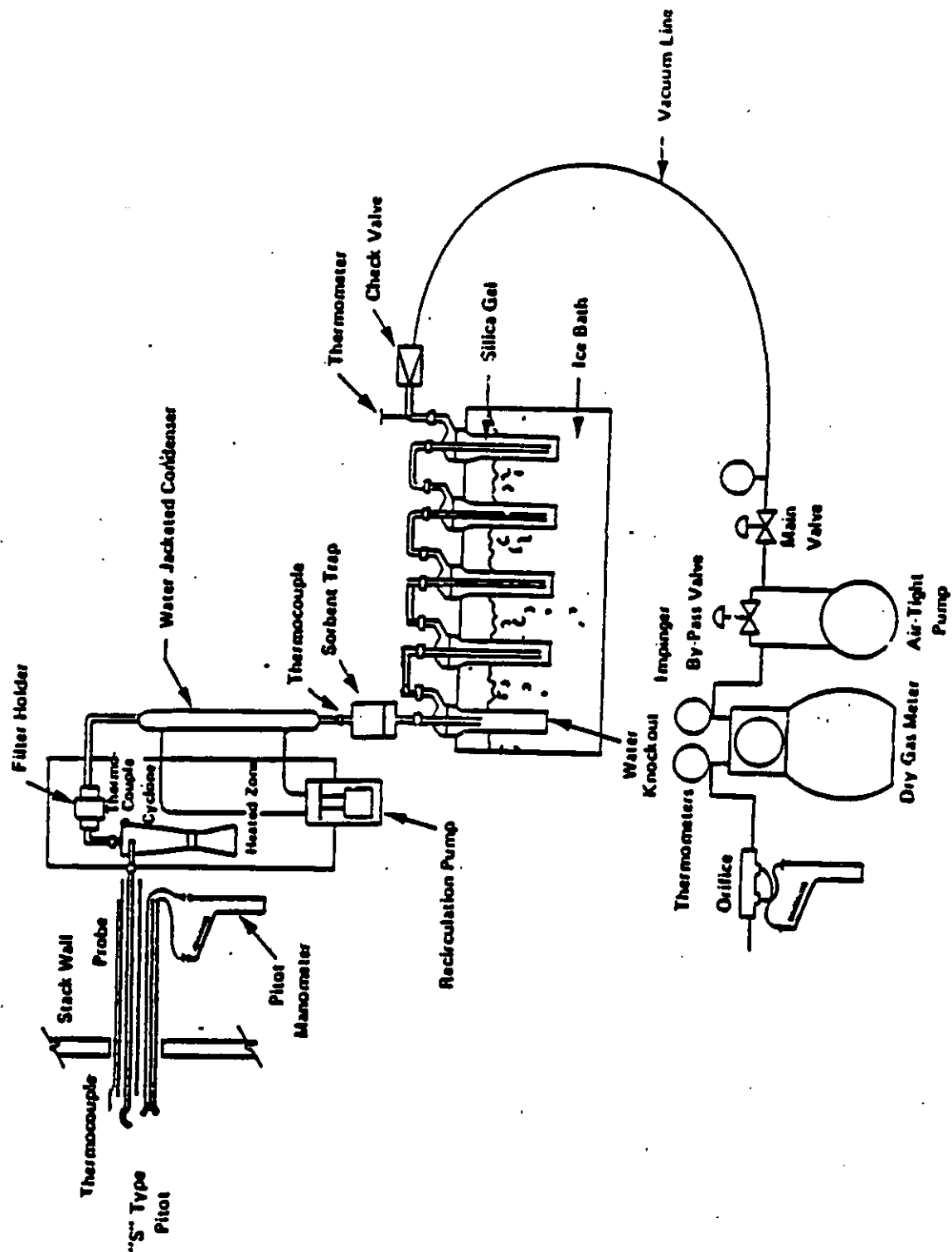


Figure 4.1 HHV Drain schematic diagram

The temperature of the gas at the outlet of the sorbent was measured and maintained between the limits of 0° and 20°C. Cooling of the gas condenses part of the sample gas moisture which may entrain and/or dissolve certain organic species. To insure that this material is collected, the sorbent module was positioned vertically such that the condensate percolated through the resin bed and into the bank of impingers.

During sampling of Wyckoff's vacuum vents, the mass of organic matter condensed on the surfaces of the cooling coil was large enough to markedly reduce the heat transfer rate and prevented proper cooling of the sample gas. The detailed results section of this report contains additional discussion of the effects of the organic compound condensation.

The impinger bank consisted of four impingers connected in series. The first impinger was initially empty to collect the condensate from the preceding sorbent module. The Greenberg-Smith nozzle was replaced by a very short stem such that the sampled gas does not bubble through the collected condensate.

The second and third impingers were included to remove acid species from the gas, and each contained 100 ml chromatographic-grade distilled water. The fourth impinger was filled with silica gel to absorb moisture remaining in the gas, thereby assuring accurate gas flow measurements and preventing damage to the pumping system.

A standard Method 5 meter box containing a variety of gas handling and metering devices followed the impinger bank.

All components of the sampling train were joined either by stainless steel Swagelok® fittings or by ground glass ball-and-socket joints capable of leak-free seals under sampling conditions. Stopcock grease was not used at any point in the system.

All components of the train which contact the sample gas were vigorously cleaned prior to transport to the field. The cleaning consisted of soaking in chromerge®, rinsing with tap water, distilled water, and chromatographic-grade methanol, successively. All components were sealed with methylene chloride-rinsed aluminum foil. Preparation of the XAD modules and resin is discussed in the analytical section.

Prior to each run, the individual components of the sampling train were assembled, and the probe/sampling train leak-tested. The heating and cooling systems are started and allowed to reach equilibrium. Then, the nozzle was plugged and the system evacuated to about 200 mm Hg (15"). If the leakage exceeded the lesser of 4% of the sampling rate or 0.02 cfm, the leak was corrected.

Sampling Procedure

The sampling plan called for multiple point isokinetic sampling for 3-4 hours. The condensation of organic material discussed above was great enough to prevent sampling isokinetically for the duration of the sampling period. Further, the exhaust gas velocity was very low across the cross section of the stack and, within the accuracy of the pitot tube

the cross section of the stack and, within the accuracy of the pitot tube system, essentially constant so lack of multiple point traversing and isokinetic sampling was judged to have minimal consequences. The samples were collected at a point of average velocity at a rate as near to isokinetic as possible.

Post-Test Operations

At the conclusion of the test, the train was leak checked and allowed to cool. The probe/nozzle unit was removed and capped, and the filter holder inlet capped, and all components taken to the cleanup area for sample recovery. The basic procedures of EPA Method 5 were followed, modified as necessary for recovery of samples for organic analysis. The probe, nozzle and filter holder were each rinsed three times with acetone and then three times with methylene chloride. All liquid samples were transferred to glass bottles with Teflon®-lined caps which had been cleaned as previously described for the glassware. The XAD module was removed from the train and sealed with grease-free glass balls and sockets. The impingers were removed and the volume of condensed water collected measured. The percolated condensate from the first impinger was transferred to a sample bottle along with rinses from the condenser, the contents of the second and third impingers and acetone and methylene chloride rinses of all three impingers and connecting glassware. All bottles were tightly capped, sealed with Teflon®-lined caps and Teflon® tape, and the liquid level marked on the bottles.

For these tests, one XAD module was exposed as a field blank to the lab. There was one trip blank included in the shipment. The field blanks were installed into a sample train, leak checked, heated to 120°C for 2 hours, and then recovered along with the other sample train fractions.

NIOSH Sorbent Tube Sampling

Retort fugitive emissions were to be sampled for 6 hours (3 runs @ 2 hours/run) with a MM5 train. This approach became impractical when we discovered that the door was left open for only half an hour rather than 8 hours. A sampling system using NIOSH-type sorbent tubes was devised to collect a sample in a short time (approximately 30 minutes) and retain the analytical sensitivity of MM5.

Sampling Train

Figure 4.2 displays a typical NIOSH design sorbent tube and the sampling train used to extract the sample gas from the stack. An EPA Method 6 type metering/flow control system was used to pull the sample gas and measure the sampled gas volume. The sample train consisted of one or two sorbent tubes containing 600 mg of XAD-2, a moisture removal trap, sample line, and control console containing pump, rotameter and dry gas meter. Flow rates and total gas volume were set and measured to match the capacity of the sorbent and the analytical sensitivity necessary. Two separate systems were used to collect duplicate samples.

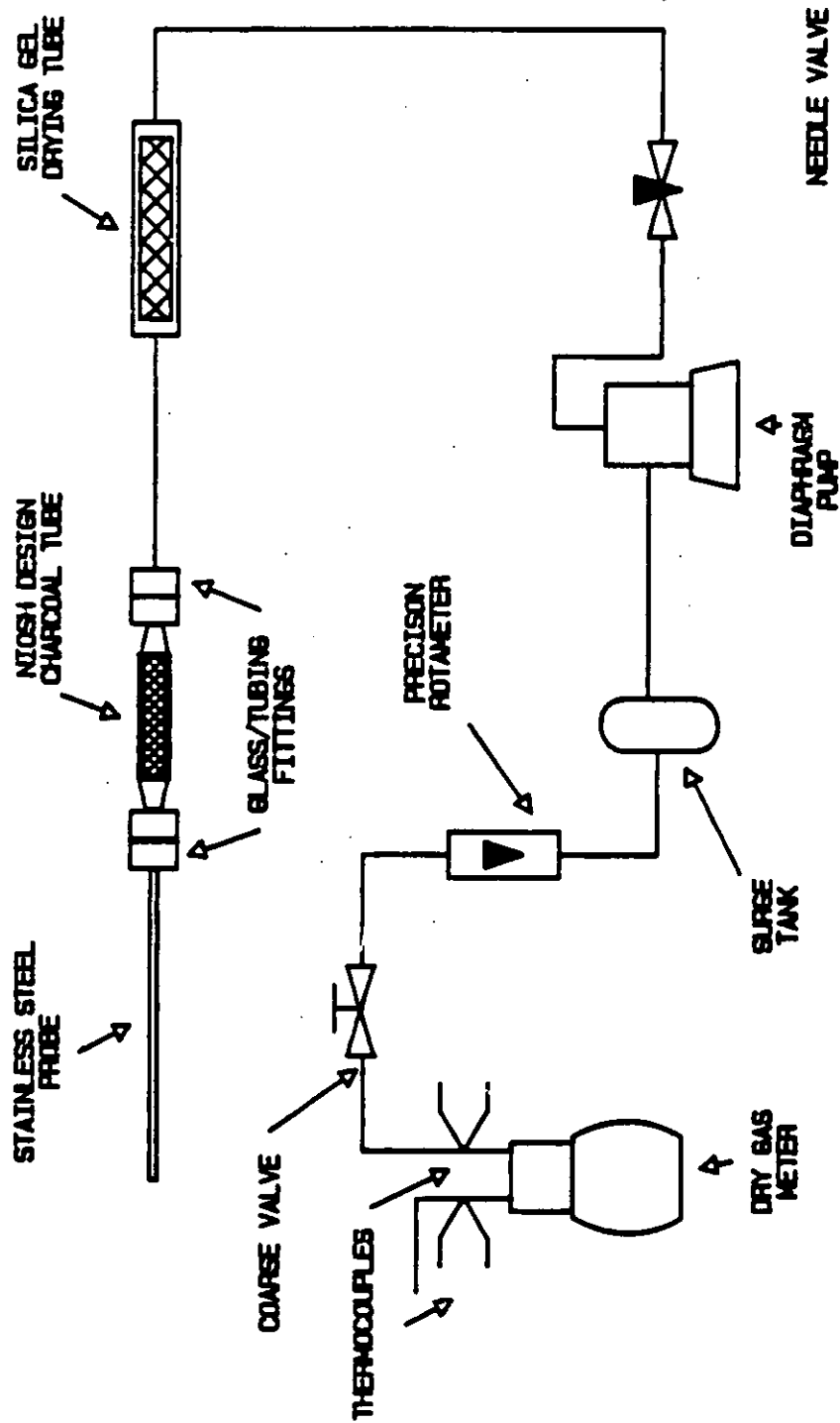


FIGURE 4.2

CHARCOAL TUBE SAMPLING TRAIN

Figure 4.3 shows how the tubes, sample lines and anemometer were assembled. The anemometer was used to obtain an average velocity for the sampling time as well as intermediate velocity readings. The area of the retort opening which had gas outflowing or inflowing was estimated by observing a set of wind sock "directional flags" placed in front of the opening.

Sampling Procedure

Samples were collected by breaking off the ends of the flame sealed sorbent tubes and pulling sample gas through the tube at 0.25 liter/minute for about 30 minutes. At the conclusion of sampling, the tubes were removed from the trains and capped with caps supplied by the manufacturer, and then placed in an envelope, pre-labeled with the date, run number, and sample ID number.

Post-Test Operations

Field blanks were taken by breaking the ends of a tube and exposing to ambient air for a time period equal to sample collection time. A trip blank (a tube taken to the field but not opened) was stored and shipped with the sample tubes.

ANALYTICAL PROCEDURES

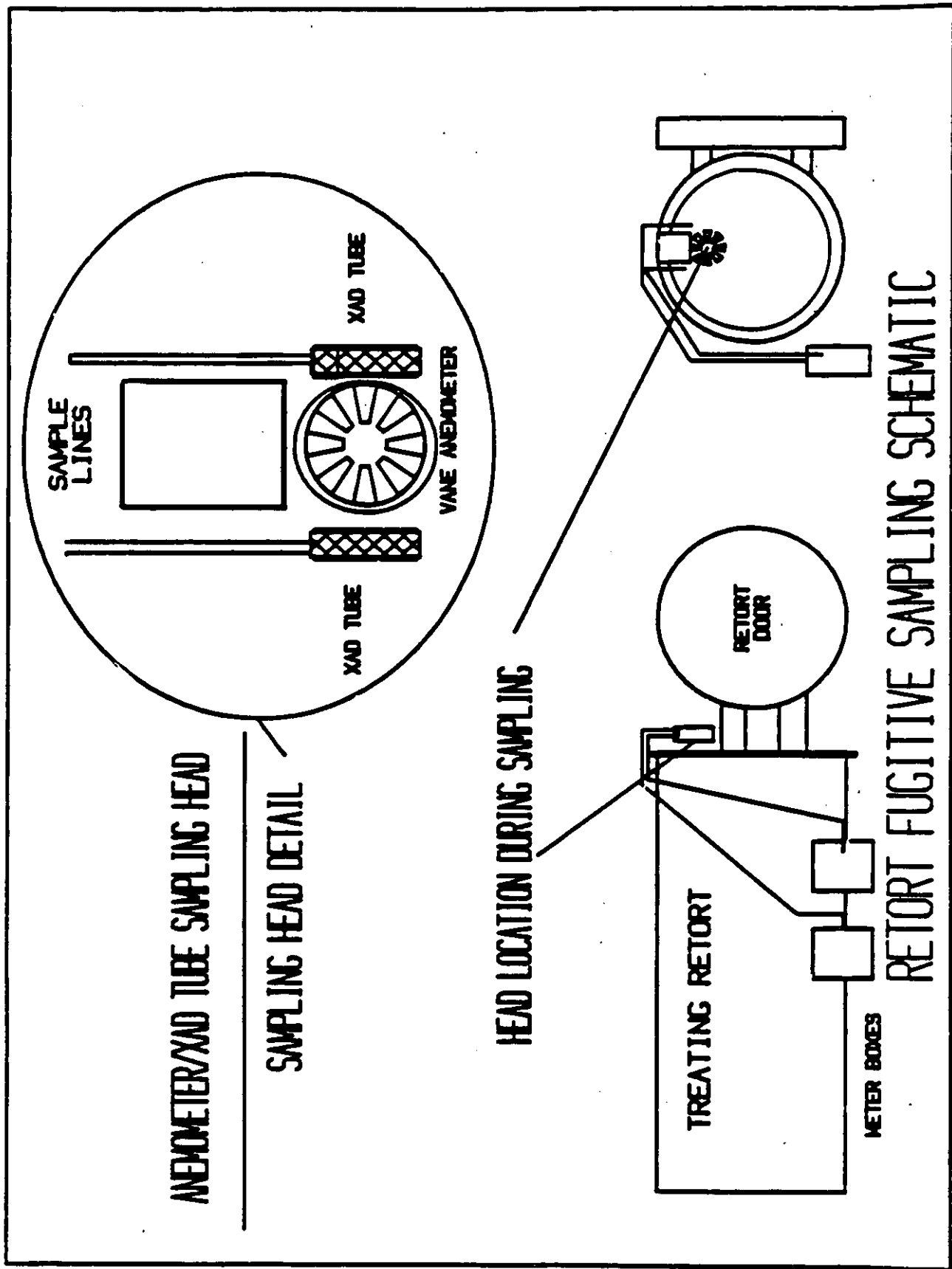
Modified Method 5 Analyses

Preparation and analysis of the MM5 samples is described below.

1. Container No. 1 (contents of the first, second, and third impingers from the MM5 sample train) - Noted the physical properties of the sample as to color, consistency, presence of solids, and measured the volume. Spiked the sample as necessary with the required surrogate compounds for QC purposes. Without adjusting the sample pH, transferred the impinger solution to a 1000-ml separatory funnel. Rinsed the sample container with 20 ml of acetone, followed by two 20 ml portions of methylene chloride, adding the rinses to the separatory funnel. Extracted the sample with three separate aliquots of methylene chloride, transferring to a K-D evaporator, after filtering through pre-extracted, dried Na_2SO_4 .

2. Container No. 2 (methylene chloride, acetone rinse of probe, filter housing, and any miscellaneous glassware) - Noted the physical properties of the sample as to color, consistency, presence of solids, and measure the volume. Spiked the original solvent sample as necessary for QC purposes. Added to the K-D evaporator as in Step A.

3. Container No. 3 (filter) - Noted the physical properties of the filter as to color, consistency, and presence of particles. Placed the filter into the thimble containing the XAD-2[®] resin and proceed as described in 4 below.



4. XAD-2[®] Adsorbent - Observed and noted the physical properties of the sample. Spiked the sample as necessary with compounds directly into the adsorbents before their removal from the glass sorbent trap. Expel the entire contents of the sorbent trap into a glass extraction thimble with a coarse-fritted bottom.

The resin in the thimble was covered with glass wool to prevent the resin from floating out into the Soxhlet extractor. The sorbent trap was rinsed with 10 ml acetone and then three 10 ml portions of methylene chloride, and these rinses added to the receiver. The Soxhlet extractor was charged with 250 ml methylene chloride, and this resin extracted for 24 hours with a cycle time of 10 to 14 times per hour. The extractate was then added to the K-D evaporator as described above for Container No. 1.

6. After all components of the sample were combined in the K-D evaporator, the volume was reduced to a known, small volume (5 ml). This sample was further evaporated or rediluted as required to obtain analytical responses within the ranges of the calibration curves for the various analytes.

The samples were injected into a temperature-programmed capillary column gas chromatograph with mass spectrometer (GC/MS) for identification and quantification of analytes. The samples were first analyzed on GC/FID to obtain approximate analyte concentrations to establish appropriate dilutions, if any, for mass spec analysis. The general procedure described in EPA publication SW 846 was used for set up and calibration of the GC/MS and compound identification performed by the data system with verification by the analyst. Quantification of analytes was made by response factor calculation relative to phenanthrene for the PHAs and to specific calibration standards for pentachlorophenol and naphthalene.

CHAPTER 5

RESULTS

The emission rates for a variety of organic compounds have been estimated and are summarized in this section. The major parameters determining the calculated emission rates are volumetric flow rate of exhaust gas and pollutant concentration, calculated from various measurements. The major elements of the measurement and their precision are discussed below.

Exhaust Gas Flow Rate - Vacuum Vents

The vacuum vent stack discharges the vapor from a condensate knockout tank which receives the exhaust from the process vacuum pumps. These tanks are covered with a steel deck with a steel stack over the middle of the tank. The rate of gas discharged is determined by the pressure in tank and gas exits through stack and through cracks and holes in the deck plating. The pulsations of the vacuum are somewhat damped by the volume of the tanks, but the exhaust gas velocity varies in a pulsing manner. A preliminary velocity traverse was conducted in each of the stacks, the average velocity calculated, and a point of average velocity selected. During sampling, the velocity head at this point was measured and assumed to be average velocity. The velocity data herein reported represent the average value of the velocity at the point over the sample period, and were calculated with Equation 2-9 of EPA Method 2. (40 CFR Part 60, Appendix A).

Normal operations at Wyckoff has three retorts operating at different phases of the treating cycle and the vapor discharges from all three venting to one knockout tank. For these tests, however, only 1 retort was operated in order to obtain samples from 1 phase of the cycle and thus the total gas flow was lower than normal. The resulting increase in uncertainty as to flow rates was considered less important than obtaining a sample representative of the emission from one and only one phase of the treating cycle. Table 5.1 summarizes the calculated stack parameters. These flow rates reflect the system exhaust in the first hours of vacuum on a charge, whether the final vacuum/cool-down or the beginning of conditioning. Limited data taken with a vane anemometer at the end of conditioning (i.e., after 36 hours of vacuum) on the creosote system yield a gas velocity of 4.8 ft/sec or approximately 360 acfm, which is slightly lower than the initial hours. This is expected as most of the water and air have been removed from the logs and the gas discharge probably represents the continuing volatilization of treating oil and air leakage caused by the system vacuum (22" Hg).

Exhaust Gas Flow Rate - Retort Fugitives

The method employed to estimate the volume of gas discharged from the retort caused by opening the doors yielded surprisingly reproducible results. The flow rates were calculated by measuring the velocity of the outflowing gas near the top center of the retort opening with a vane anemometer, estimating the fraction of retort opening in each of three regions (in-flowing, out-flowing, and stagnant) and calculating a flow rate as the product of velocity and discharge area. Table 5.2 summarizes the measurements and resulting flow rates. These data represent the flow rate while the retort is empty with door wide open, i.e., after the treated load has been pulled, but before a new load is put in. When a treated load is pulled, the volume occupied by the logs and trams is replaced with ambient air and when a new charge is loaded, a volume of air is displaced by the new load. The total retort volume is about 5,700 ft³ and a typical full load probably occupies 50-75% of the total volume. The air drawn in with removal of a load can be considered as displaced when a new load is charged. This amounts to 3,000-4,500 ft³ during a 2-5 minute period. If the retort is open and empty for half an hour and is discharging at a rate of 4,000 acfm, the total gas exhausted is 120,000 ft³ and the volume displaced by changing loads is relatively insignificant (only 3-4% of the total).

The major areas of uncertainty associated with these measurements and results are estimation of the area of discharge and the uniformity of the velocity across that area. The area estimation is probably within +25% of the true value. The velocity measurements taken are reasonably accurate (+5%) but an uncertainty derives from whether the velocity at the measured point is in fact the average bulk velocity of the exhaust gas. Some basic heat transfer calculations suggest that 1×10^6 ft³ of air are required to cool the retort from 180° to 80°F. If the retort cools down in 2 hours, the flow rate would be 8,300 acfm which compares favorably, given the nature and extent of the variables involved, with measured values of 4,000-5,000 acfm.

Sampling Data

Tables 5.3 and 5.4 summarize the measured and calculated values for the MM5 and fugitive sample collection. The calculation procedures are routine and the equations of Method 5 are used. Example calculations are included in Appendix D. Table 5.5a and 5.5b present the results of GC/MS analysis of the samples. The results are expressed as total ug of the analytes in the sample and reflect the instrumental value, any dilutions or concentrations and calculation of total mass of analytes in the sample. Data are presented for PNA compounds found in the sample from EPA's priority pollutant list and includes those specific compounds which were specifically targeted for analysis (pentachlorophenol, naphthalene).

Emission Estimates

The data in tables 5.6 -5.9 are derived from the analysis results, sampling parameters, and gas flow rate estimations. The mass emission

rates should not be considered as representative of overall facility emission; they are supplied to provide a reference for time rate of discharge.

The fugitive emission rates are representative, within the limits previously discussed, of short-term emission rate; process schedules, actual work practices regarding open door and recharging, and treating oil variation would need to be considered to establish overall emission rates. The vacuum vent rates are more reliable and representative of emission from the treating process; again, however, process operating schedule, and work practices such time of conditioning need to be considered in extrapolating single retort data to the whole facility. Some estimate of the overall emission rate can be made for the available data. A typical treating cycle is listed below showing the duration of each phase of this cycle.

Phase	Duration (hrs)	Pressure (P) or Vacuum (V)	Discharge to Atmosphere	Comments
Conditioning	48	V	Yes	
Pressure Treating	1	P	No	
Expansion	3	V	No	
Steam Cleaning	2	V	Yes	Negligible gas flow
Final Vacuum	1	V	Yes	
Fugitive	1/2	—	Yes	

Table 5.10 presents estimated emissions for one complete cycle of penta and creosote treating. The rate given are based on the flow measurements and concentration data previously discussed.

TABLE 5.1
FLOW RATE DATA

Run	Date	Average Stack Temp. °F	Moisture %	Velocity ft/sec	Volumetric Flow Rates	
					acfm	dscfm
WPV-MM5-1A	7/29	152.4	26.9	12.2		
-1B	7/29	<u>152.3</u>	<u>27.9</u>	<u>12.2</u>		
Avg		152.4	27.2	12.2	780	490
WPV-MM5-2A	7/31	145.8	21.4	11.5		
-2B	7/31	<u>145.3</u>	<u>22.6</u>	<u>11.8</u>		
Avg		145.6	22.0	11.7	750	510
WCV-MM5-1A	7/31	131.5	17.2	6.2		
-1B	7/31	<u>131.2</u>	<u>16.7</u>	<u>6.3</u>		
Avg		131.4	17.0	6.3	460	340
WCV-MM5-2A	8/1	145.4	20.5	6.3		
-2B	8/1	<u>145.3</u>	<u>21.7</u>	<u>6.3</u>		
Avg		145.4	21.1	6.3	460	320

TABLE 5.2

RETORT FUGITIVE FLOW RATES

Retort	C	D	E
Treating Oil	Creosote	Creosote	Penta
Date	7/26/85	7/31/85	7/31/85
Time	1255-1355	1350-1420	1157-1227
Average Velocity ft/min	302	313	372
(Range of Intermediate Velocities)	(316-375)	(286-333)	(400)
Average Discharge Area, % (Range)	37 (29-41)	30 (13-35)	30 (22-35)
Average Discharge Area, ft ²	16.4	13.3	13.3
Average Discharge Gas Flow Rate acfm	4960	4160	4950

TABLE 5.3
MODIFIED METHOD 5 SAMPLING

Run ID	WCV 1A	WCV 2A	HPV 1A	HPV 2A
<u>SAMPLING DATA</u>				
Meter Volume, acf	52.666	36.822	59.403	73.376
Meter Pressure (DH), in H ₂ O	1.02	1.27	0.60	1.56
Meter Temp, °F	90.3	94.2	96.7	85.2
Barometric Pressure, in. Hg	29.9	29.9	29.9	29.9
Gamma	1.013	1.013	1.007	1.007
Condensate, ml	227.9	196.1	445.8	416.4
Area of Stack, ft ²	1.227	1.227	1.069	1.069
<u>DERIVED DATA</u>				
Meter Volume, dscf	51.477	35.759	56.996	72.057
Water Volume, scf	10.727	9.230	20.984	19.600
Moisture Content, %	17.2	20.5	26.9	21.4
Molecular Weight, Dry	28.84	28.84	28.84	28.84
Molecular Weight, Wet	26.98	26.62	25.92	26.52
Stack Velocity, ft/sec	6.25	6.27	12.20	11.62
Stack Temp, °F	131.3	145.4	152.4	145.6
Volumetric Flow Rate				
- acfm	460	462	783	745
- dscfm	342	319	493	509

TABLE 5.4

FUGITIVE EMISSION SAMPLING DATA

Run ID	WPF-X 1A	WPF-X 1B	WCF-X 1A	WCF-X 2A	WCF-X 2B
Meter Volume, liters	15.240	14.23	9.835	10.22	8.18
Meter Volume, acf	0.539	0.520	0.348	0.361	0.289
Meter Pressure, in. H ₂ O	1.00	1.20	1.00	0.97	1.21
Meter Temp, °F	83.6	79.8	86.1	81.3	77.9
Barometric Pressure, in. Hg	29.9	29.9	29.9	29.9	29.9
Gamma	0.951	0.955	0.951	0.951	0.955
Meter Volume, dscf	0.500	0.489	0.321	0.321	0.273

TABLE 5.5a

VACUUM VENT SAMPLING ANALYSIS RESULTS
Total ug of Analyte

Compound	Creosote		XCV X-2	Penta	
	WCV	WCV		WPV	WPV
	1A	2A		1A	1A
Naphthalene	2,500,000	900,000	7200	2,000,000	1,000,000
Acenaphthylene	2,400	8,200	ND	ND	ND
Acenaphthene	3,600	40,000	20	30,000	25,000
Phenanthrene	1,400	3,400	ND	2,100	1,700
Fluoranthene	ND	ND	ND	ND	ND
Fluorene	4,700	25,000	ND	8,400	9,800
Pyrene	ND	ND	ND	ND	ND
Pentachlorophenol	ND	ND	ND	ND	ND
Anthracene	ND	ND	ND	ND	ND
Detection Limit	1,000	1,000	10	1,000	1,000

TABLE 5.5b

FUGITIVE SAMPLING ANALYSIS RESULTS
Total ug of Analyte

Compound	WPF 1A	WPF 1B	WCF 1A	WCF 2A	WCV 2B
Naphthalene	50	200	3,000	11,070	1,200
Acenaphthylene	ND	ND	ND	90	60
Acenaphthene	ND	10	2,200	1,900	1,500
Phenanthrene	ND	10	640	300	130
Fluoranthene	ND	ND	300	40	ND
Fluorene	ND	20	740	580	340
Pyrene	ND	ND	ND	20	ND
Pentachlorophenol	ND	600	ND	ND	ND
Anthracene	ND	ND	100	100	20
Detection Limit	10	10	10	10	10

TABLE 5.6a

EMISSION RATES
PROCESS CYCLE - CREOSOTE CONDITIONING (INITIAL VACUUM)
EXHAUST GAS FLOW RATE - 342 dscfm

Compound	Emissions		
	\$/dscf	ppmv	\$/hour
Naphthalene	107×10^{-6}	323	2.20
Acenaphthylene	0.103×10^{-6}	0.26	0.0021
Acenaphthene	0.154×10^{-6}	0.39	0.0032
Phenanthrene	0.060×10^{-6}	0.13	0.0012
Fluoranthene	$<0.043 \times 10^{-6}$	<0.08	<0.0009
Fluorene	0.201×10^{-6}	0.47	0.0041
Pyrene	$<0.043 \times 10^{-6}$	<0.08	<0.0009
Pentachlorophenol	$<0.043 \times 10^{-6}$	<0.06	<0.0009
Anthracene	$<0.043 \times 10^{-6}$	<0.09	<0.0009

TABLE 5.6b

EMISSION RATES
PROCESS CYCLE - CREOSOTE COOL-DOWN (FINAL VACUUM)
EXHAUST GAS FLOW RATE, 319 dscfm

Compound	#/dscf	Emissions	
		ppmv	#/hour
Naphthalene	55.5×10^{-6}	168	1.06
Acenaphthylene	0.506×10^{-6}	1.29	0.0097
Acenaphthene	2.47×10^{-6}	6.19	0.0472
Phenanthrene	0.210×10^{-6}	0.46	0.0040
Fluoranthene	$<0.062 \times 10^{-6}$	<0.12	<0.0012
Fluorene	1.54×10^{-6}	3.59	0.0295
Pyrene	$<0.062 \times 10^{-6}$	<0.12	<0.0012
Pentachlorophenol	$<0.062 \times 10^{-6}$	<0.09	<0.0012
Anthracene	$<0.062 \times 10^{-6}$	<0.13	<0.0012

TABLE 5.7a

EMISSION RATES
PROCESS CYCLE - PENTA CONDITIONING (INITIAL VACUUM)
EXHAUST GAS FLOW RATE, 493 dscfm

Compound	Emissions		
	#/dscf	ppmv	#/hour
Naphthalene	77.4×10^{-6}	234	2.29
Acenaphthylene	$<0.039 \times 10^{-6}$	<0.10	<0.0011
Acenaphthene	1.16×10^{-6}	2.91	0.0343
Phenanthrene	0.081×10^{-6}	0.18	0.0024
Fluoranthene	$<0.039 \times 10^{-6}$	<0.07	<0.0011
Fluorene	0.325×10^{-6}	0.76	0.0096
Pyrene	$<0.039 \times 10^{-6}$	<0.07	<0.0011
Pentachlorophenol	$<0.039 \times 10^{-6}$	<0.06	<0.0011
Anthracene	$<0.039 \times 10^{-6}$	<0.08	<0.0011

TABLE 5.7b

EMISSION RATES
PROCESS CYCLE - PENTA COOL DOWN (FINAL VACUUM)
EXHAUST GAS FLOW RATE, 493 dscfm

Compound	Emissions		
	#/dscf	ppmv	#/hour
Naphthalene	30.6×10^{-6}	92.4	0.93
Acenaphthalene	$<0.031 \times 10^{-6}$	<0.08	<0.0009
Acenaphthene	0.765×10^{-6}	1.92	0.0234
Phenanthrene	0.052×10^{-6}	0.11	0.0016
Fluoranthene	$<0.031 \times 10^{-6}$	<0.06	<0.0009
Fluorene	0.300×10^{-6}	0.70	0.0092
Pyrene	$<0.031 \times 10^{-6}$	<0.06	<0.0009
Pentachlorophenol	$<0.031 \times 10^{-6}$	<0.04	<0.0009
Anthracene	$<0.031 \times 10^{-6}$	<0.07	<0.0009

TABLE 5.8

PROCESS CYCLE - CREOSOTE FUGITIVE
EXHAUST GAS FLOW RATE, approximately 3800 dscfm

Compound	Sample	Emissions		
		g/dscf	ppmv	g/hour
Naphthalene	A	72.4 x 10 ⁻⁶	219	16.5
	B	9.69 x 10 ⁻⁶	29.3	2.21
Acenaphthalene	A	0.59 x 10 ⁻⁶	1.5	0.134
	B	0.49 x 10 ⁻⁶	1.2	0.111
Acenaphthene	A	12.4 x 10 ⁻⁶	31.2	2.83
	B	12.1 x 10 ⁻⁶	30.4	2.76
Phenanthrene	A	1.96 x 10 ⁻⁶	4.3	0.447
	B	1.05 x 10 ⁻⁶	2.3	0.239
Fluoranthene	A	0.26 x 10 ⁻⁶	0.50	0.060
	B	<0.08 x 10 ⁻⁶	<0.15	<0.018
Fluorene	A	3.79 x 10 ⁻⁶	8.8	0.865
	B	2.75 x 10 ⁻⁶	6.4	0.626
Pyrene	A	0.13 x 10 ⁻⁶	0.25	0.030
	B	<0.081 x 10 ⁻⁶	<0.15	<0.018
Pentachlorophenol	A	<0.065 x 10 ⁻⁶	<0.10	<0.015
	B	<0.081 x 10 ⁻⁶	<0.12	<0.018
Anthracene	A	<0.65 x 10 ⁻⁶	1.4	0.149
	B	0.16 x 10 ⁻⁶	0.35	0.037

TABLE 5.9

EMISSION RATES
PROCESS CYCLE - PENTA FUGITIVE
EXHAUST GAS FLOW RATE, approximately 4500 dscfm

Compound	Emissions		
	g/dscf	ppmv	g/hour
Naphthalene	0.900×10^{-6}	2.7	0.24
Acenaphthalene	$<0.045 \times 10^{-6}$	<0.11	<0.012
Acenaphthene	0.045×10^{-6}	0.11	0.012
Phenanthrene	0.045×10^{-6}	0.10	0.012
Fluoranthene	$<0.045 \times 10^{-6}$	<0.09	<0.012
Fluorene	0.090×10^{-6}	0.21	0.024
Pyrene	$<0.045 \times 10^{-6}$	<0.09	<0.012
Pentachlorophenol	2.71×10^{-6}	3.9	0.72
Anthracene	$<0.045 \times 10^{-6}$	<0.1	<0.012

TABLE 5.10a

TOTAL CREOSOTE CYCLE EMISSIONS
(LB/HR)

COMPOUND	EMISSIONS (LB/HR)			CYCLE TOTAL (56 HR)
	CONDITIONING (48 HR)	FINAL (1 HR)	FUGITIVE (1/2 HR)	
Naphthalene	1.06	1.06	4.68	111
Acenaphthalene	0.101	0.0097	0.613	0.172
Acenaphthene	0.154	0.0472	1.40	1.60
Phenanthrene	0.0576	0.004	0.172	0.233
Fluoranthene	<0.0432 ^a	<0.0012	0.0195	<0.0639
Fluorene	0.197	0.0295	0.373	0.599
Pyrene	<0.0432	<0.0012	0.012	<0.0564
Pentachlorophenol	<0.0432	<0.0012	0.0083	<0.0527
Anthracene	<0.0432	<0.0012	<0.0465	<0.0477

$$\frac{111.5}{56 \text{ hr}} = 1.99 \frac{\text{lb}}{\text{hr}} \times 24 \text{ hr} \times 2 = 111.5 \text{ lb/day}$$

$$\times 412 \text{ lb/day} = 84,815 \text{ lb/day}$$

$$43.4 \text{ tons/day}$$

TABLE 5.10b.

TOTAL PENTA CYCLE EMISSIONS
(LB/HR)

COMPOUND	EMISSIONS (LB/HR)			CYCLE TOTAL (56 HR)
	CONDITIONING (48 HR)	FINAL (1 HR)	FUGITIVE (1/2 HR)	
Naphthalene	110	0.930	0.120	111
Acenaphthalene	<0.0528	<0.0009	<0.006	<0.0597
Acenaphthene	1.65	1.920	0.006	3.57
Phenanthrene	0.115	0.110	0.006	0.231
Fluoranthene	<0.0528	<0.0009	<0.006	<0.597
Fluorene	0.461	0.0092	0.012	0.482
Pyrene	<0.0528	<0.0009	<0.006	<0.0597
Pentachlorophenol	<0.0528	<0.0009	0.360	<0.414
Anthracene	<0.0528	<0.0009	<0.006	<0.0597

* < indicates values calculated lower limit of detection.

$$\frac{2.91 \text{ lb}}{\text{hr}} \times \frac{24 \text{ hr}}{\text{day}} \times \frac{365 \text{ days}}{\text{yr}} = 455 \text{ lbs/yr of Nap}$$

56 hr/cycle

$$\frac{4.68}{0.12} = 39 \text{ times. One notation}$$

APPENDIX A

PROCESS DATA RECORDS

A-1

PENTA LOAD

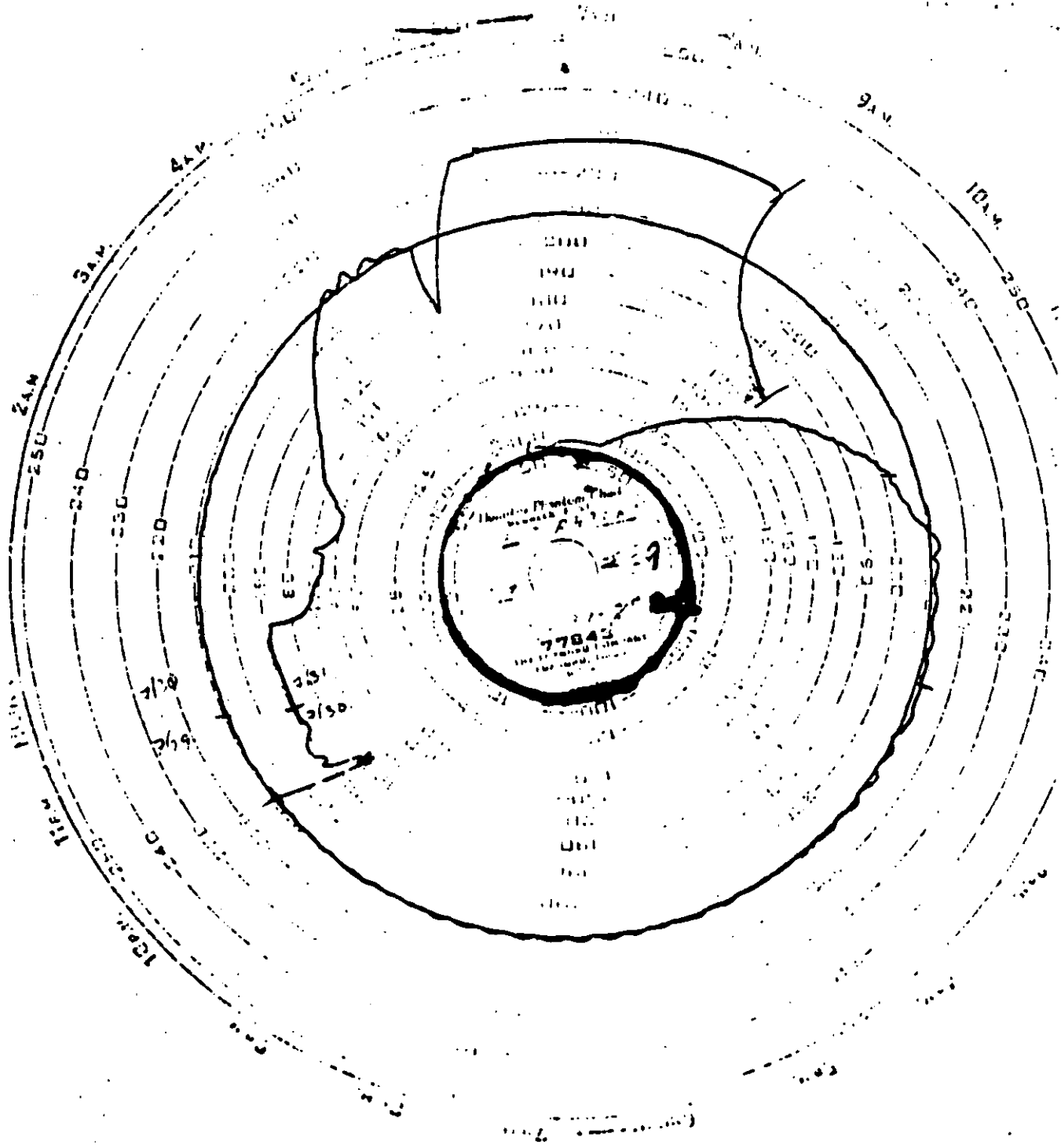
E- RETORT

		APPROX HRS <u>7-30-85</u>
START LOAD	11:00 PM	
PUMP-OUT + COLD OIL		2 $\frac{3}{4}$
30" I AIR		1 $\frac{1}{4}$
PUMP BACK		1 $\frac{3}{4}$
PRESS		1 $\frac{1}{2}$
X-BATH		1 $\frac{1}{2}$
PUMP-OUT		1 HR
L-STEAM		2 HRS
FINAL UAC		2 HRS
		<u>13 $\frac{3}{4}$ TOTAL HRS</u>

LOAD SHOULD BE DONE
BY 2:00 PM WED. 7-31-85

BUTCH - HOME PHONE
842-3341

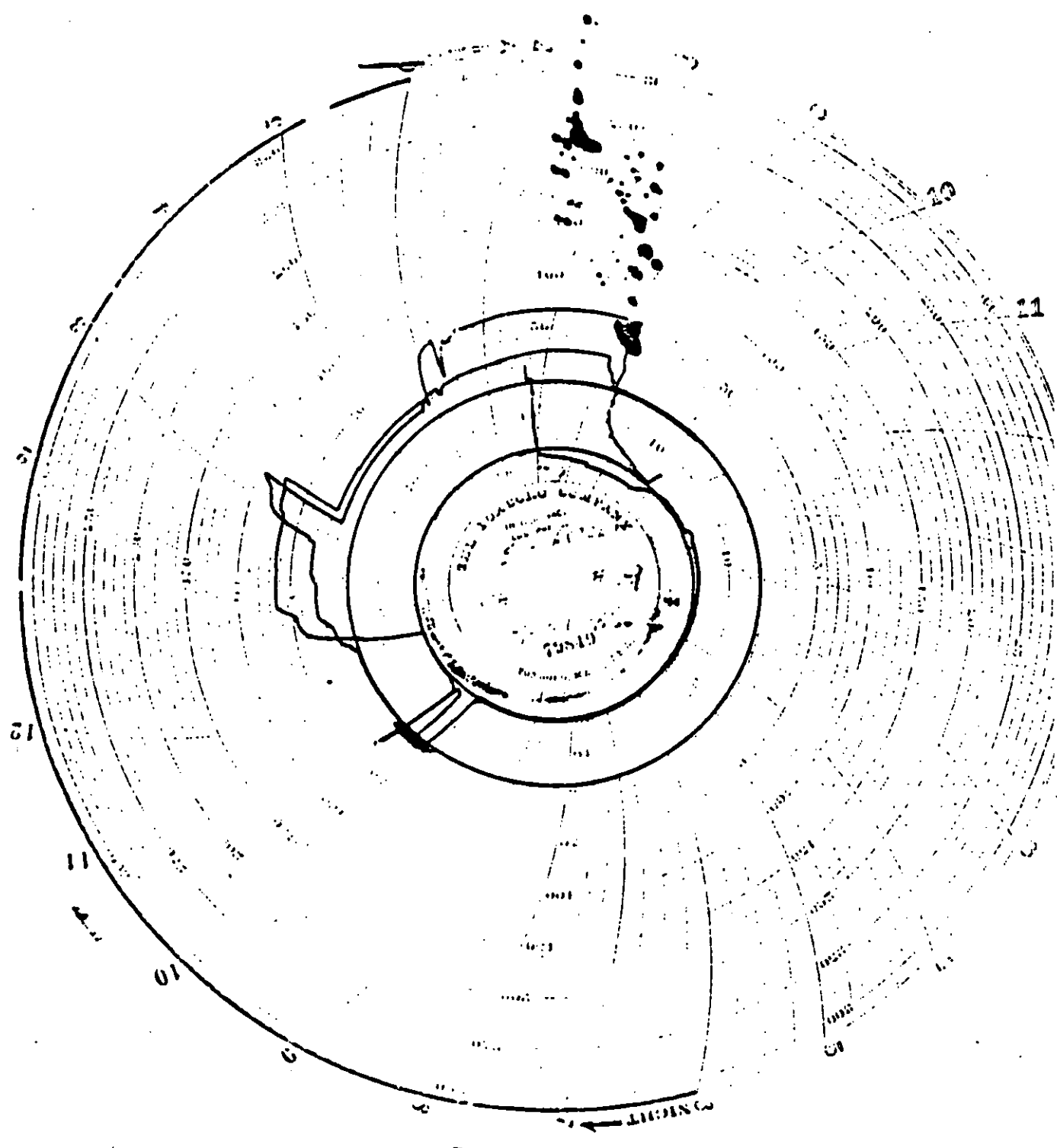
OR PAGER 997-0497



Target E (1000 ft)
 1000 ft
 1000 ft

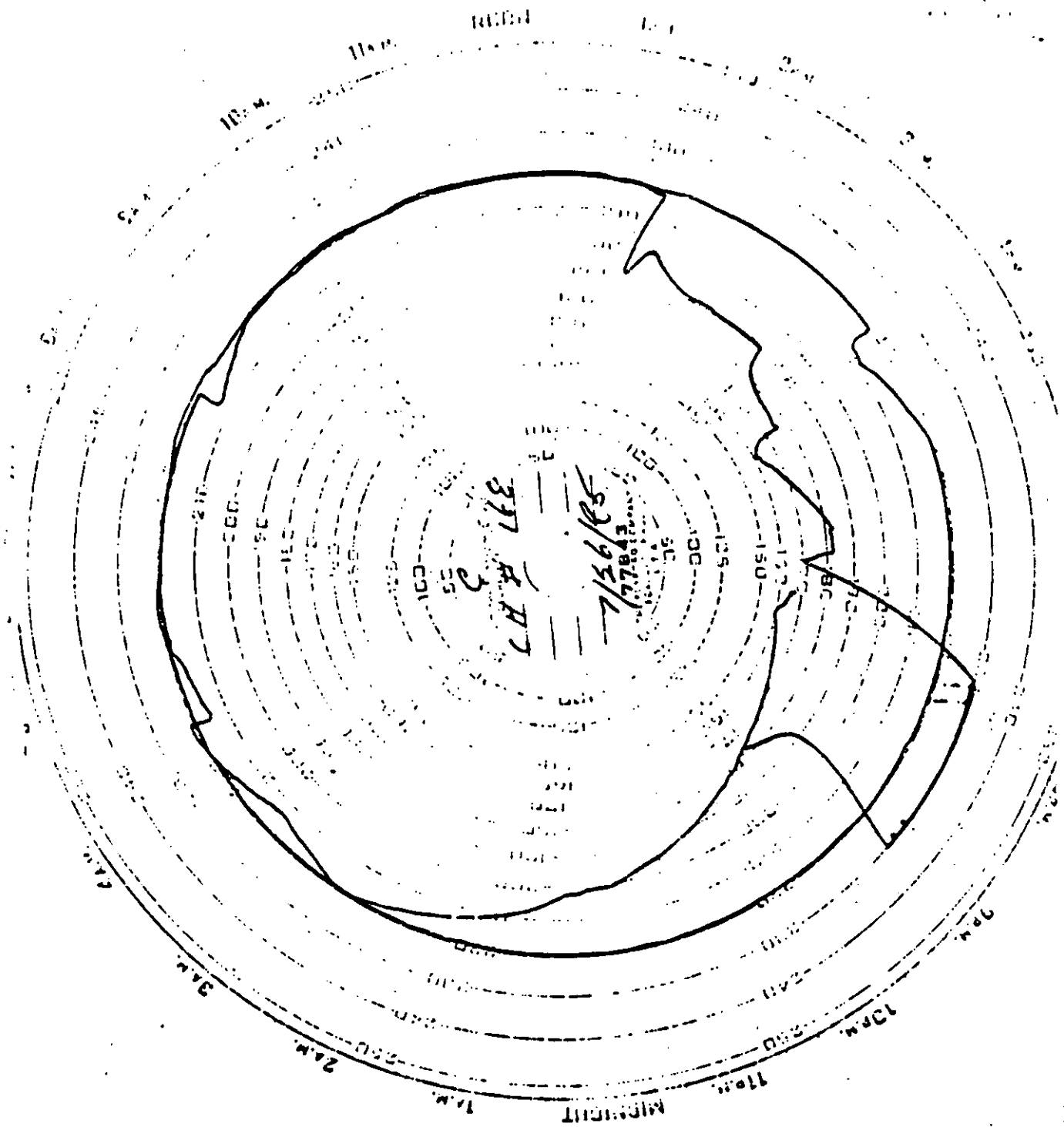
Target E (1000 ft)

H-2



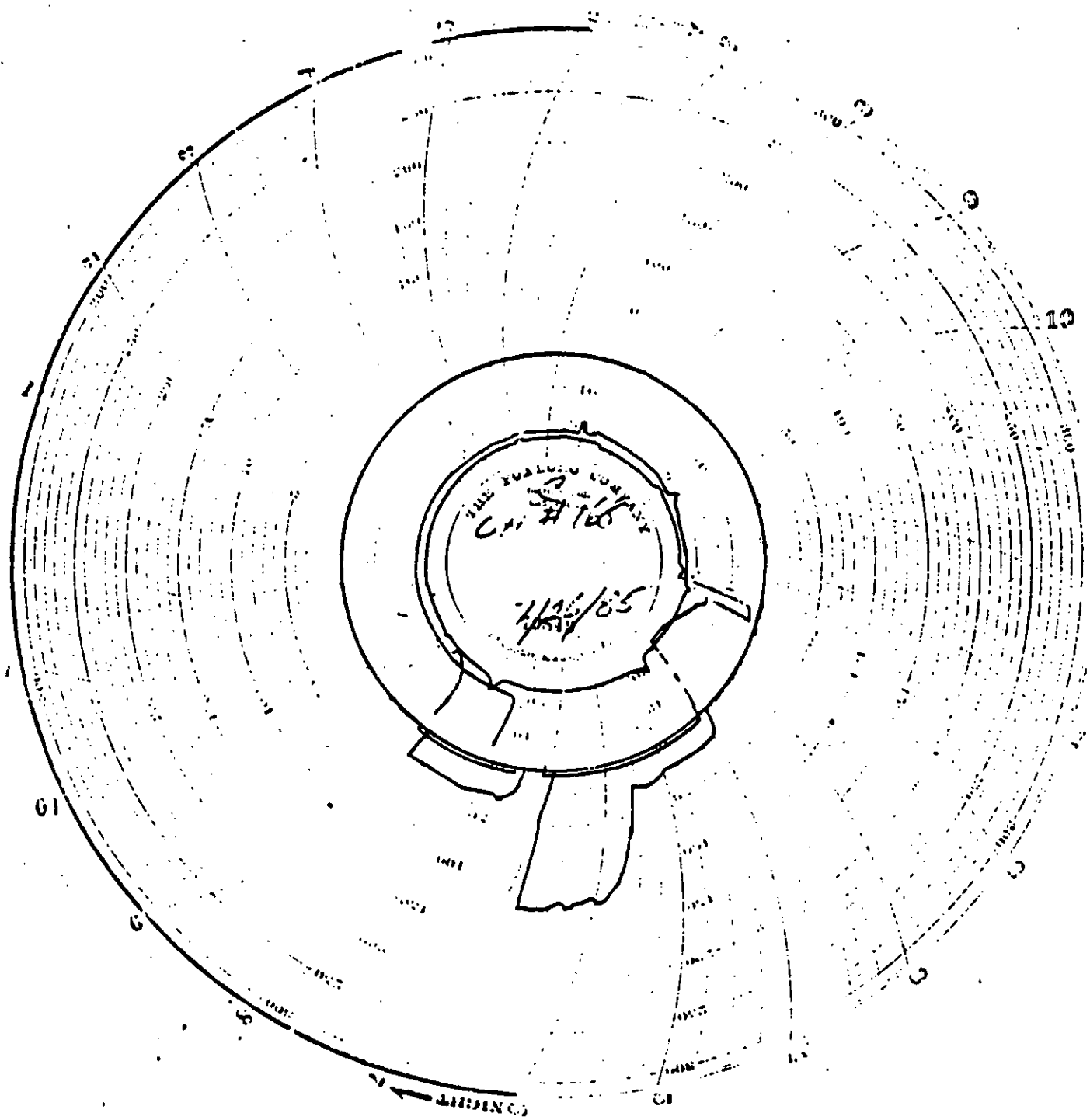
VACUUM, " Hg. PRESSURE CHART

A-3

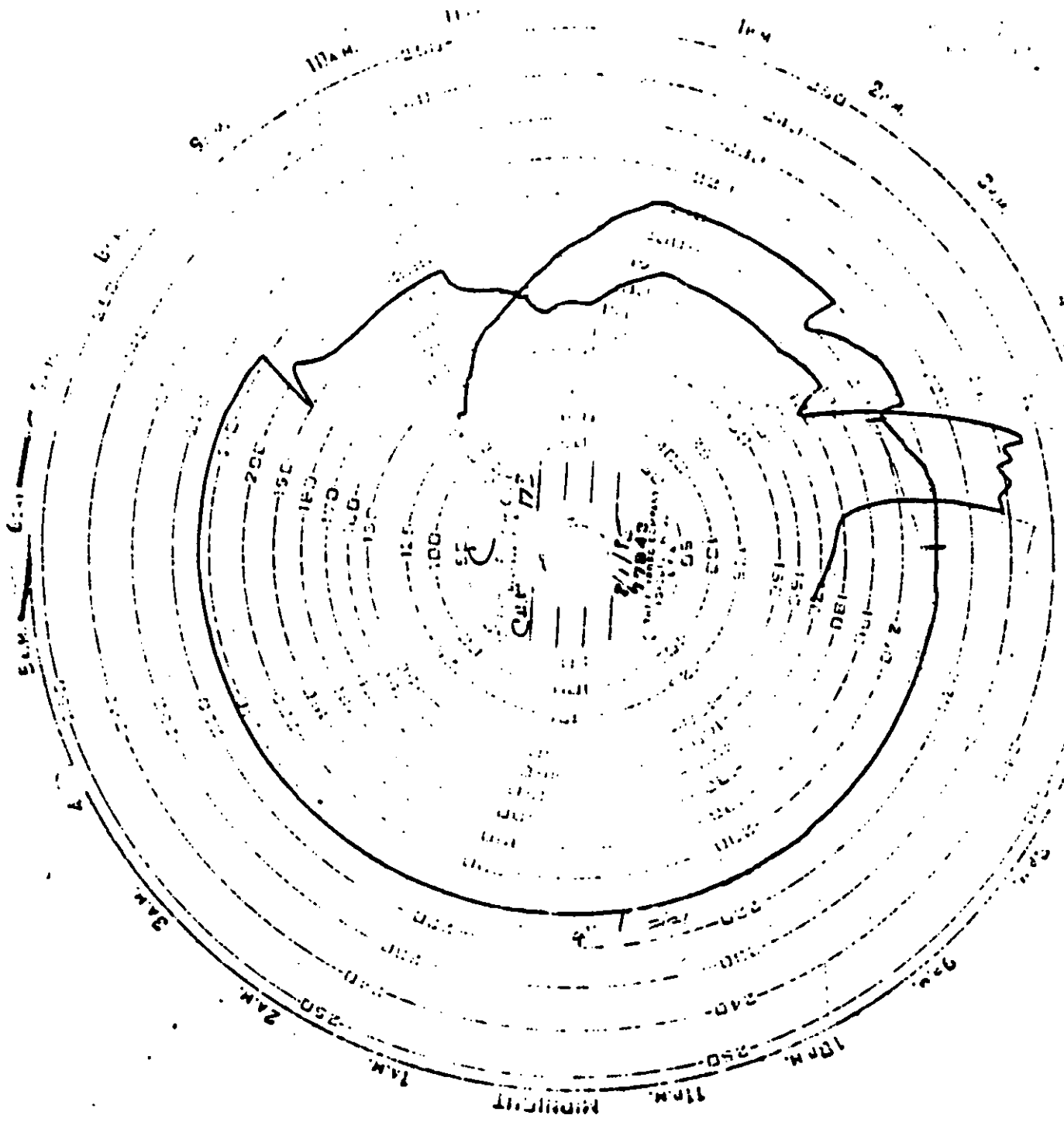


REPORT ()
 START: 10:00
 END: 10:59

A-4

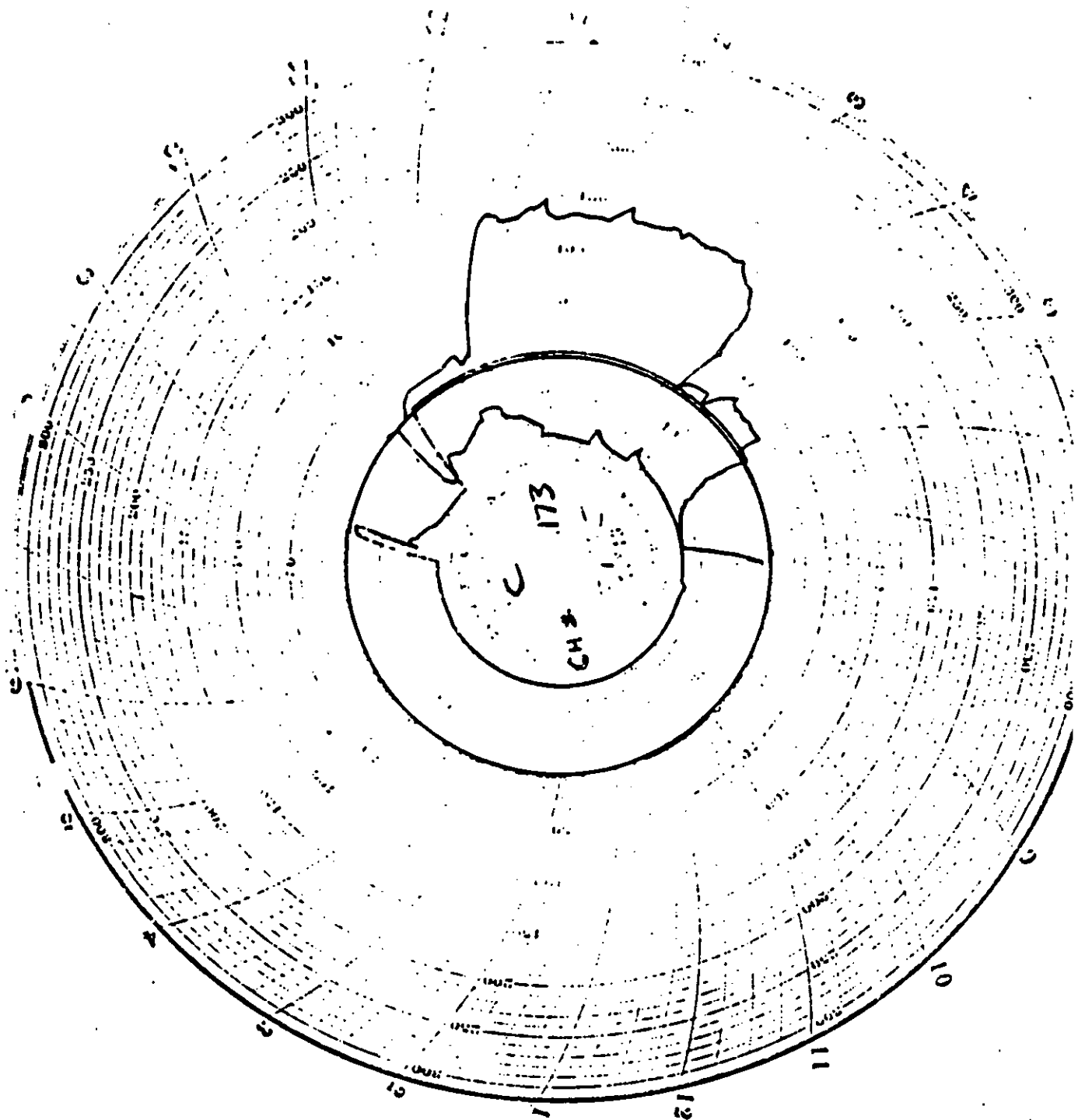


A-5

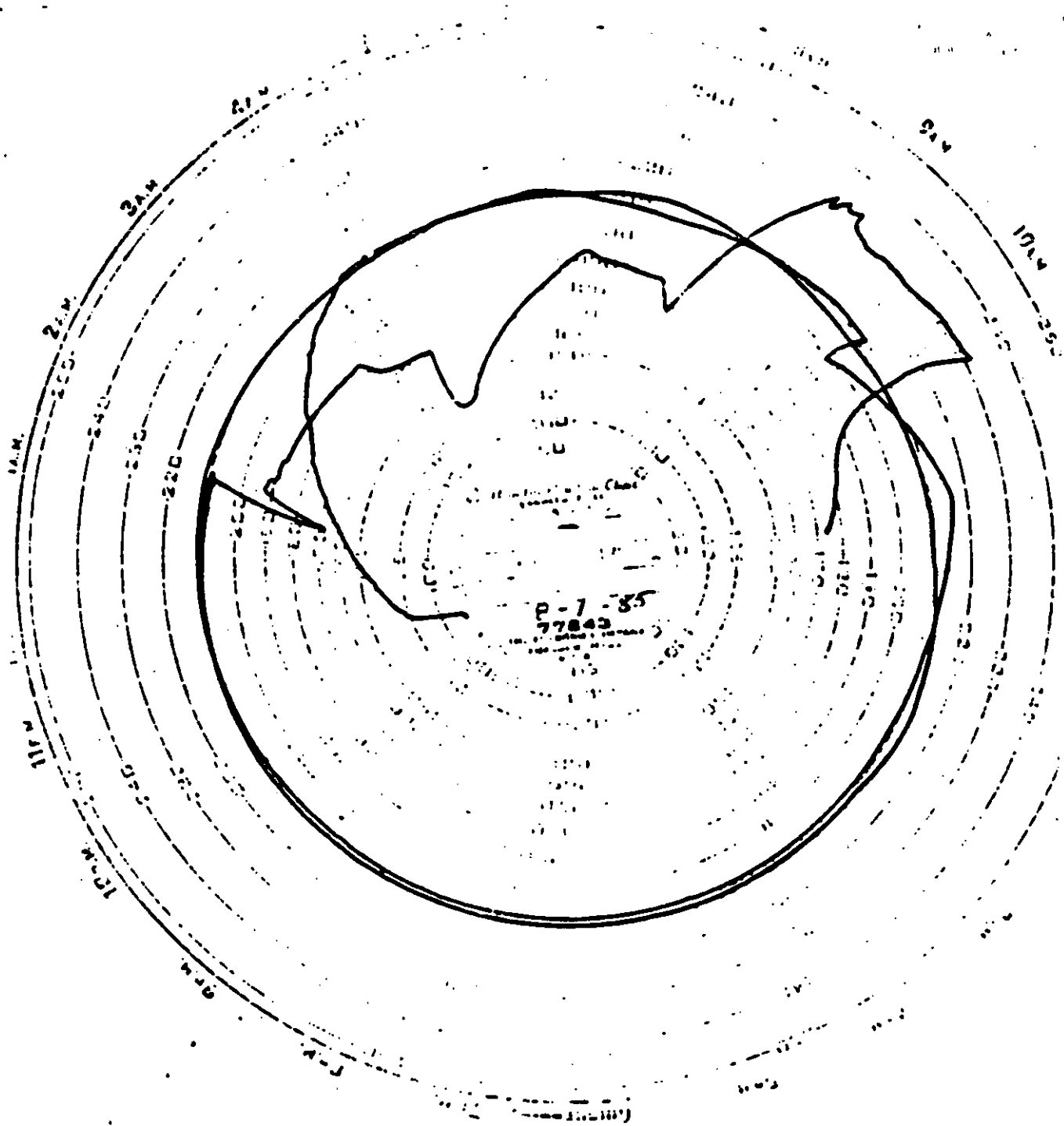


REPORT C. GREENE,
 START 7/2/85
 STOP 8/1/85

A-6

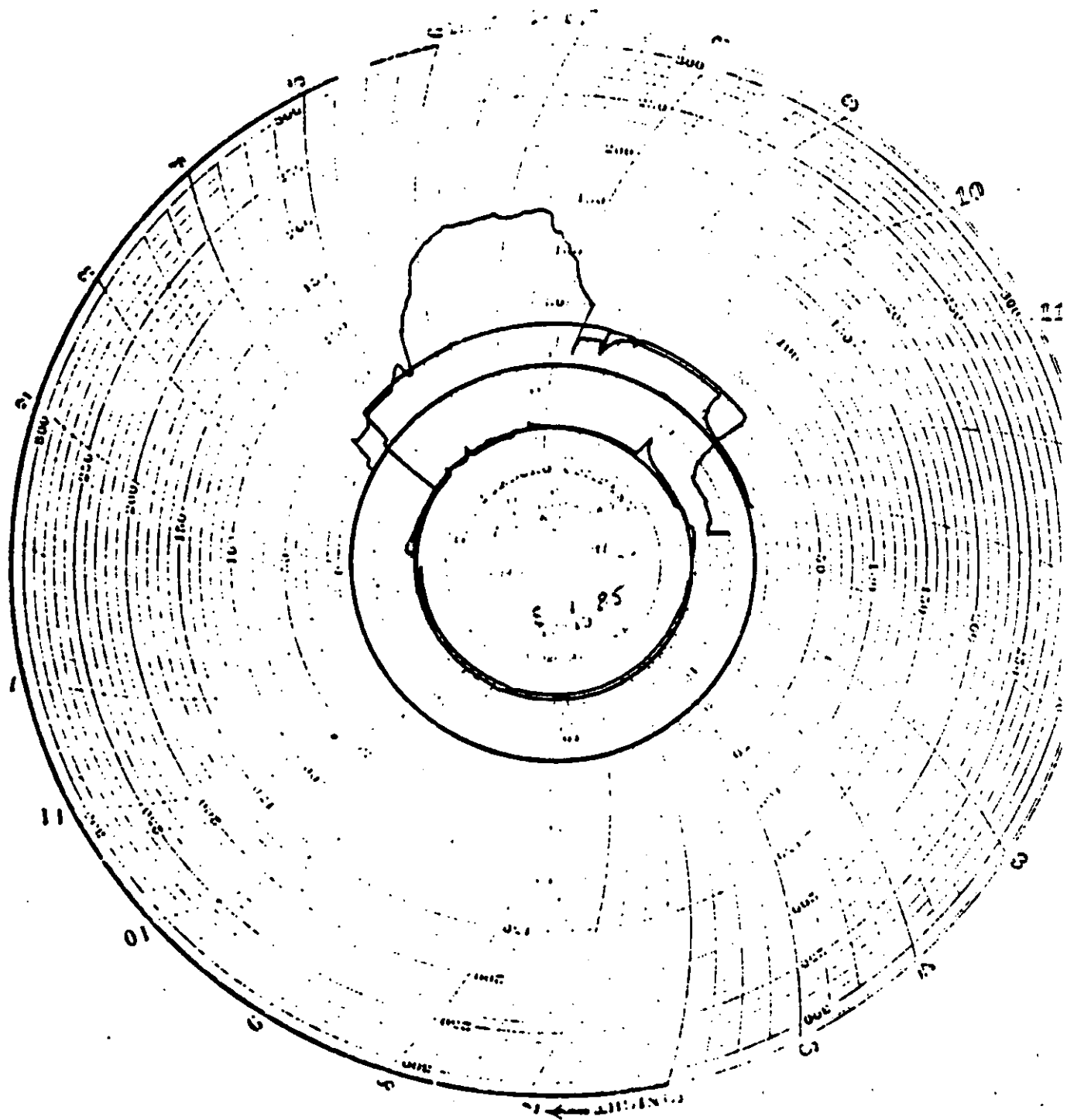


A-7



Track of P-7-85
Start: 7:30 PM
End: 8:10 PM

A-3



A-9

APPENDIX B

FIELD DATA

B-1

FIELD DATA

Plant WYCCORP
 Date 2/21/83
 Sampling Location PORT LAMAR FLS
 Sample Type XAD
 Run Number WPF-X1A
 Operator _____
 Ambient Temperature _____
 Barometric Pressure _____
 Static Pressure (P_s) _____
 Filter Number(s) _____
 Pretest Leak Rate = ____ ofm @ ____ in. Hg
 Pretest Pitot Leak Check _____
 Pretest Orsat Leak Check _____
 Read and Record all Data Every ____ Minutes

Probe Length and Type _____
 Pitot Tube I.D. No. _____
 Nozzle I.D. _____
 Assumed Moisture, % _____
 Temp. Readout S/N _____
 Meter Box Number VOST 0 F
 Meter Hg _____
 C Factor _____
 Meter Gamma -951
 Heater Box Setting _____
 Reference ΔP _____
 Post Test Leak Rate = ____ ofm @ ____ in. Hg
 Post Test Pitot Leak Check _____
 Post Test Orsat Leak Check _____

Schematic of

Traverse Point Layout

VOST PORT LAMAR

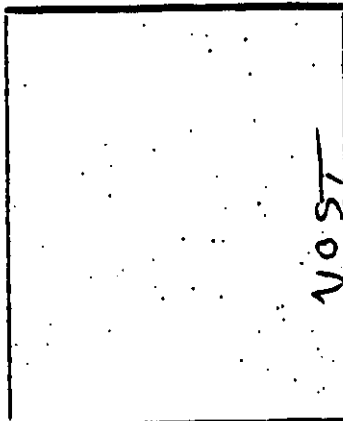
Pig 10

Traverse Point Number	Sampling Time, (min) / (24-hour clock)	Gas Meter Reading (V_g) ft ³	Velocity Head (P_v) in. H ₂ O	Orifice Prod. Differential (H) in. H ₂ O	Stack Temp. (T_s) °F	Dry Gas Meter Inlet ($T_{m, in}$) °F	Dry Gas Meter Outlet ($T_{m, out}$) °F	Pump Box Temp. Vacuum Filter in. Hg	Sample In-linger Temp. °F
0	1116	83.15		48	73	78	X	1.0	
5	1121	83.33		48		79		0.9	
10	1126	84.61	5000	48	74	CLOSE		1.0	
15	1139	84.61		48	74	82		0.9	
20	1145	86.2		48	74	83		0.9	
25	1154	88.40		48	76	85		0.9	
30	1159	89.8		48	78	86		0.9	
40	1209	92.00		48	78	87		1.0	
50	1219	94.8		48	78	88		1.0	
60	1229	97.39	5000	48	78	88		1.0	
1				48				1.0	
2		15.29		48		83.06		1.0	
3									
4		14.13							
5									
6		14.19	12500						

FIELD DATA

Plant WYCKOFF
 Date 2/23/55
 Sampling Location CRACK D
 Sample Type XAD
 Run Number WQ-FX-2B
 Operator JB
 Ambient Temperature _____
 Barometric Pressure _____
 Static Pressure (P_g) _____
 Filter Number(s) _____
 Pretest Leak Rate - 0 cfm @ _____ in. Hg
 Pretest Pitot Leak Check _____
 Pretest Orsat Leak Check _____
 Read and Record all Data Every _____ Minutes

Probe Length and Type _____
 Pitot Tube I.D. No. _____
 Nozzle I.D. _____
 Assumed Moisture, % _____
 Temp. Readout S/N _____
 Meter Box Number 1057 °C
 Meter Hg _____
 C Factor _____
 Meter Gamma 0.55
 Meter Box Setting _____
 Reference ΔP _____
 Post Test Leak Rate - 0 cfm @ _____ in. Hg
 Post Test Pitot Leak Check _____
 Post Test Orsat Leak Check _____



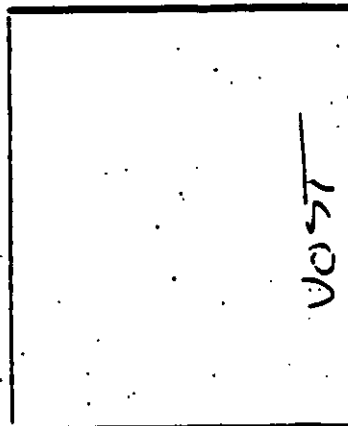
Schematic of
 Traverse Point Layout

Traverse Point Number	Sampling / Time (min) / (clock)	Gas Meter Reading (V _g) ft	Velocity Head (P _g) in. H ₂ O	Orifice Pres. Differential (H ₂ O) Desired Actual	Stack Temp. (T _g)	Dry Gas Meter Temp. Inlet (T _{in}) °C Outlet (T _{out}) °F	Pump In. Hg	Sample Box Temp. Filter Temp. °F	In-Player Temp. °F
1	10 / 1333	481.49		48	72	25	24	1-2	
2	10 / 1343	82.9		48	72	25	25	1-2	
3	10 / 1347	84.05		48	92	25	25	1-2	
4	10 / 1353	85.7		48	92	26	26	1-2	
5	20 / 1357	86.44		48	92	26	26	1-2	
6	28 / 1402	87.75		48	92	27	27	1-2	
7	10 / 1407	88.01		48	92	27	27	1-2	
8	32-37 / 1410	4179.17		48	92	27	27	1-2	
9	1			48	92	27	27	1-2	
10	1			48	92	27	27	1-2	
11	1			48	92	27	27	1-2	
12	1			48	92	27	27	1-2	
13	1			48	92	27	27	1-2	
14	1			48	92	27	27	1-2	
15	1			48	92	27	27	1-2	
16	1			48	92	27	27	1-2	
17	1			48	92	27	27	1-2	
18	1			48	92	27	27	1-2	
19	1			48	92	27	27	1-2	
20	1			48	92	27	27	1-2	

FIELD DATA

Plant Wyesoff
 Date 7/3/85
 Sampling Location CAGE D
 Sample Type CO2
 Run Number WGE-X-24
 Operator AB
 Ambient Temperature 71°F
 Barometric Pressure 30
 Static Pressure (P_g) —
 Filter Number(s) —
 Pretest Leak Rate = — cfm @ — in. Hg
 Pretest Pitot Leak Check —
 Pretest Orsat Leak Check —
 Read and Record all Data Every — Minutes

Probe Length and Type —
 Pitot Tube I.D. No. —
 Nozzle I.D. —
 Assumed Moisture, % —
 Temp. Readout S/M —
 Meter Box Number V057 °F
 Meter No. —
 C Factor —
 Meter Gamma 0.751
 Heater Box Setting —
 Reference ΔP —
 Post Test Leak Rate = — cfm @ — in. Hg
 Post Test Pitot Leak Check —
 Post Test Orsat Leak Check —



Schematic of
 Traverse Point Layout

B-7

Traverse Point Number	Sampling Time, (min) / (clock)	Gas Meter Reading (V _g) ft ³	Velocity Head (P _g) in. H ₂ O	Orifice Pres. Differential (H) in. H ₂ O	Stack Temp. (T _g) °F	Dry Gas Meter Temp. Inlet (T _m) °F Outlet (T _m) °F	Pump Vacuum in. Hg	Sample Box Temp. Filter Temp. °F	In-Plunger Temp. °F
0	11332	2122.37		48	48	78	78	1.0	
6	11343	90.8		48	132	78	78	1.0	
10	11347	200.97		48	77	78	80		
16	11353	01.6		48	77	78	80		
20	11357	02.37		48	83	78	80		
25	11402	03.76		48	86	78	80	0.75	
30	11407	05.07		48	86	78	80	0.75	
35	11412	06.30		48	87	78	80	1.0	
40	11413	2107.61	4800	48	93 (94)	78	80	0.95	
1	1								
1	1	10.22		48		78	80	0.95	
1	1								
1	1	9.51							
1	1								
1	1	2.76							
1	1	0							
1	1								
1	1								

1-
2-
3-
4-
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Plant WYCOFF
Date 7/28/85
Sampling Location PERMIT VAC VENT
Sample Type AA-5
Run Number UP-V-AMS-1-B-A (60)
Operator JB JMY
Ambient Temperature 69°
Barometric Pressure
Static Pressure (P_s)
Filter Number(s) AAA
Pretest Leak Rate 0.03 cfm @ 16 in. Hg
Pretest Pitot Leak Check
Pretest Orsat Leak Check
Read and Record all Data Every Minutes

Probe Length and Type 2'
Pitot Tube I.D. No.
Nozzle I.D.
Assumed Moisture, %
Temp. Readout S/M
Meter Box Number 2010-8407
Meter No. 1.552
C Factor
Meter Gamma 1.007
Heater Box Setting
Reference ΔP
Post Test Leak Rate 0.008 cfm @ 20 in. Hg
Post Test Pitot Leak Check
Post Test Orsat Leak Check

Schematic of
Traverse Point Layout

* Condenser Temp checked to 130°F for approx 2-3 minutes at approx 1340-1345

Traverse Point Number	Sampling Time, (min) / (clock)	Gas Meter Reading (V _m) ft.	Velocity Head (P _v) in. H ₂ O	Orifice Pres. Differential (H) in. H ₂ O	Stack Temp. (T _g) °F	Dry Gas Meter Inlet (T _m) °F	Dry Gas Meter Outlet (T _m) °F	Pump Vacuum in. Hg	Sample Box Temp. Filter Temp. °F	In-Plunger Temp. °F
372	0 / 1235	588.064	0.035	0.69	146	91	63	13.3	183	83
373	15 / 1250	595.3	0.040	0.69	152	93	65	13.3	183	83
374	30 / 1305	602.9	0.040	0.69	156	93	65	13.3	183	83
375	45 / 1320	610.4	0.040	0.69	159	94.5	65	13.3	183	83
376	60 / 1335	618.0	0.035	0.69	154	93.5	65	13.3	183	83
377	75 / 1350	625.6	0.035	0.69	151	94.5	65	13.3	183	83
378	90 / 1400	630.7	0.040	0.69	155	97.5	65	13.3	183	83
379	90 / 1405	633.1	0.035	0.69	154	97.5	65	13.3	183	83
380	95 / 1410	635.6	0.035	0.69	152	98	65	13.3	183	83
381	100 / 1415	638.0	0.035	0.69	152	98	65	13.3	183	83
382	105 / 1420	641.4	0.035	0.69	152	100	65	13.3	183	83
383	110 / 1425	642.7	0.035	0.69	154	100	65	13.3	183	83
384	115 / 1430	645.1	0.035	0.69	154	100	65	13.3	183	83
385	120 / 1435	645.4	0.035	0.69	151	99	65	13.3	183	83
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SAMPLE RECOVERY DATA

Plant: Wycoff
 Date: _____
 Sampling Location: Penia Vac. Vent
 Sample Type: MMS
 Run Number: WP-V-MMS1-A
 Sample Box Number: _____
 Clean-up Man: _____
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: N/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>615</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>100</u>	ml	<u>100</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>415</u>	

Silica Gel

Final Volume:	<u>466.5</u>	g	_____	g	_____	g
Initial Volume	<u>234.7</u>	g	<u>202.0</u>	g	_____	g
Net Volume	_____	g	_____	g	<u>29.8</u>	g
Total Moisture:	_____		<u>29.8</u>		<u>445.8</u>	

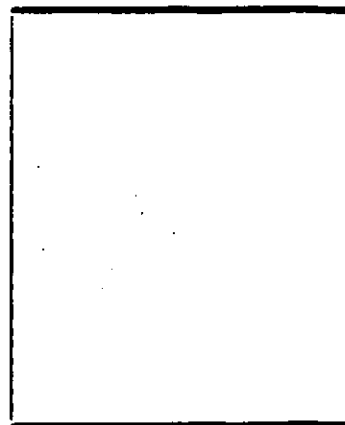
Description of Impinger Catch: _____

13.46

FIELD DATA

Plant Wycoff
 Date 7/29/83
 Sampling Location Pentac Vaa Vent
 Sample Type HMS
 Run Number WP-V-M-5-1-B341
 Operator dfb, jdy
 Ambient Temperature 68° 76°
 Barometric Pressure NA
 Static Pressure (P_g) NA
 Filter Number(s) NA
 Pretest Leak Rate - 0.05 cfm @ 15 in. Hg
 Pretest Pitot Leak Check Adm
 Pretest Orsat Leak Check Adm
 Read and Record all Data Every 1 Minute

Probe Length and Type 2'
 Pitot Tube I.D. No. 2'
 Nozzle I.D. 2'
 Assumed Moisture, % 2'
 Temp. Readout S/M 2'
 Meter Box Number 455
 Meter No. 1703
 C Factor 1.013
 Meter Gamma 1.013
 Heater Box Setting 1.013
 Reference ΔP 1.013
 Post Test Leak Rate - 0.05 cfm @ 15 in. Hg
 Post Test Pitot Leak Check Adm
 Post Test Orsat Leak Check Adm



Schematic of
 Traverse Point Layout

0.5 cfm

Traverse Point Number	Sampling Time, (min) / (clock)	Gas Meter Reading (V _g) ft	Velocity Head (P _g) in. H ₂ O	Orifice Pres. Differential (H) in. H ₂ O	Stack Temp. (T _g) °F	Dry Gas Meter Temp. Inlet (T _g) °F	Outlet (T _g) °F	Pump Vacuum in. Hg	Sample Box Temp. Filter °F	In- plunger Temp. °F
1	0 11230	095.455	0.035	0.76	1450	60	60	8	245	25
2	15 11625	22.38	0.035	0.76	1470	68	68	8	243	35
3	30 11300	111.0	0.040	0.76	155	94	94	9	260	70
4	45 11315	118.05	0.040	0.76	154	96	96	10 1/2	245	69
5	60 11330	125.67	0.035	0.76	153	96	96	12	241	65
6	75 11345	138.8	0.035	0.76	155	99	99	14 1/2	252	66
7	90 11400	141.0	0.035	0.76	155	102	102	16 1/2	252	62
8	105 11415	148.6	0.035	0.76	152	104	104	18	234	71
9	120 11430	156.334	0.035	0.76	152	104	104	(19)		
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SAMPLE RECOVERY DATA

Plant: Wycoff
 Date: _____
 Sampling Location: Pinto Vnc. Vent
 Sample Type: MMS
 Run Number: WP-V-MMS1-B
 Sample Box Number: _____
 Clean-up Man: _____
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: N/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>645</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>100</u>	ml	<u>100</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>445</u>	

Silica Gel

Final Volume:	<u>505.5</u>	g	_____	g	_____	g
Initial Volume:	<u>251.0</u>	g	<u>229.2</u>	g	_____	g
Net Volume:	_____	g	_____	g	<u>25.3</u>	g
Total Moisture:	_____		<u>25.3</u>		<u>470.3</u>	

Description of Impinger Catch: _____

P-11

SAMPLE RECOVERY DATA

Plant: Wyeoff
 Date: 7/31/85
 Sampling Location: Vac. Vent
 Sample Type: MMS
 Run Number: WP-V-MMS-2A
 Sample Box Number: _____
 Clean-up Man: ThJ
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: N/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>585</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>100</u>	ml	<u>00</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>385</u>	

Silica Gel

Final Volume:	<u>505.0</u>	g	_____	g	_____	g
Initial Volume:	<u>241.9</u>	g	<u>231.7</u>	g	_____	g
Net Volume:	_____	g	_____	g	<u>31.4</u>	g
Total Moisture:	_____		_____		<u>416.4</u>	

Description of Impinger Catch: _____

FIELD DATA

Plant	WYCKOFF
Date	7/31/85
Sampling Location	Porter Lake Venn
Sample Type	WGS
Run Number	WGV-775-2.03
Operator	JFB
Ambient Temperature	62°
Barometric Pressure	30.
Static Pressure (P _s)	
Filter Number(s)	
Pretreat Leak Rate -	0.007 cfm @ 1" in. Hg
Pretreat Pitot Leak Check	
Pretreat Orsat Leak Check	
Read and Record all Data Every	Minutes

Probe Length and Type	4
Pitot Tube I.D. No.	
Nozzle I.D.	.375
Assumed Moisture, %	22
Temp. Readout S/M	
Water Box Number	255
Water M _g	1.788
C Factor	
Water Gases	1.013
Water Box Setting	
Reference ΔP	
Post Test Leak Rate -	0.017 @ 20 in. Hg
Post Test Pitot Leak Check	
Post Test Orsat Leak Check	

Schematic of
Traverse Point Layout

[illegible]

SAMPLE RECOVERY DATA

Plant: Wycott
 Date: 7/31/85
 Sampling Location: Vac. Vard
 Sample Type: MMS
 Run Number: WP-V-MMS-28
 Sample Box Number: _____
 Clean-up Man: MS
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: N/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>620</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>100</u>	ml	<u>100</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>420</u>	

Silica Gel

Final Volume:	<u>500.0</u>	g	_____	g	_____	g
Initial Volume	<u>225.5</u>	g	<u>245.8</u>	g	_____	g
Net Volume	_____	g	_____	g	_____	g
Total Moisture:	_____		<u>28.7</u>		<u>28.7</u>	
					<u>448.7</u>	

Description of Impinger Catch: _____

SAMPLE RECOVERY DATA

Plant: Wycoff
 Date: 7/31/85
 Sampling Location: _____
 Sample Type: MMS
 Run Number: WC-V-MMS-1A
 Sample Box Number: _____
 Clean-up Man: 212
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: N/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>370</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>100</u>	ml	<u>100</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>190</u>	

Silica Gel

Final Volume:	<u>512.5</u>	g	_____	g	_____	g
Initial Volume	<u>240.9</u>	g	<u>233.7</u>	g	_____	g
Net Volume	_____	g	_____	g	<u>37.7</u>	g
Total Moisture:	_____		<u>37.9</u>		<u>227.9</u>	

Description of Impinger Catch: _____

[illegible]

Schematic of Traverse Point Layout

Read and Record all Data	Every	Minutes
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Carlsen

Q

[illegible]

B-18

SAMPLE RECOVERY DATA

Plant: Wycoff
 Date: 7/31/85
 Sampling Location: _____
 Sample Type: MM's
 Run Number: WC-U-MMS-1B
 Sample Box Number: _____
 Clean-up Man: HLB
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: Y/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>370</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>150</u>	ml	<u>100</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>170</u>	

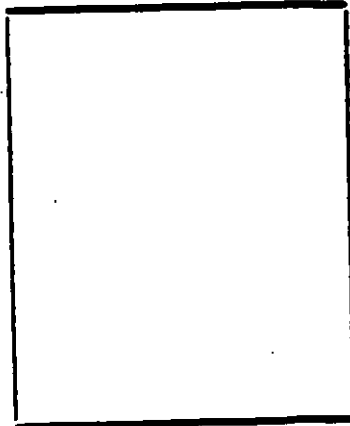
Silica Gel

Final Volume:	<u>513.5</u>	g	_____	g	_____	g
Initial Volume	<u>2443.</u>	g	<u>240.4</u>	g	_____	g
Net Volume	_____	g	_____	g	<u>28.8</u>	g
Total Moisture:	_____		<u>28.8</u>		<u>198.8</u>	

Description of Impinger Catch: _____

FIELD DATA

Plant U455555
 Date 8/1/95
 Sampling Location Site 1000
 Sample Type WV-X8
 Run Number WV-X8
 Operator WV-X8
 Ambient Temperature 62°C
 Barometric Pressure _____
 Static Pressure (P_0) _____
 Filter Number(s) _____
 Pretest Leak Rate = ____ ofm @ ____ in. Hg
 Pretest Pitot Leak Check _____
 Pretest Orsat Leak Check _____
 Read and Record all Data Every ____ Minutes



Schematic of
 Traverse Point Layout

Approved by WV-X8

Traverse Point Number	Sampling Time (min) / (clock)	Gas Meter Reading (V_g) ft ³	Match - Density Wedge (P_g) in. H ₂ O	Orifice Pressure Differential (in. H ₂ O) Desired Actual	Stack Temp. (T_g) °F	Dry Gas Meter Temp. Inlet ($T_{g,in}$) °F Outlet ($T_{g,out}$) °F	Pump Vacuum in. Hg	Sample Box Temp. Filter Temp. °F	In-plier Temp. °F
305	0 / 08:13	2102.42	1.0	4.5 4.8	104	60 60	0		
306	5 / 08:13	2109.0	1.0	4.8 4.8	104	60 60	0		
307	10 / 08:13	2116.4	2.4	4.8 4.8	104	60 60	0		
308	15 / 08:13	2111.6	1.6	4.8 4.8	104	60 60	0		
309	20 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
310	25 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
311	30 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
312	35 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
313	40 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
314	45 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
315	50 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
316	55 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
317	60 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
318	65 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
319	70 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
320	75 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
321	80 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
322	85 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
323	90 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
324	95 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		
325	100 / 08:13	2112.1	5.0	4.8 4.8	104	60 60	0		

Date	Time	Location	Weather	Wind	Temp	Humidity	Pressure	Visibility	Clouds	Sea	Current	Tide	Remarks

Date 4/1/85
Sampling Location CEED/HCMW
Sample Type MMS
Run Number WCV-HS-2B
Operator RESNER
Ambient Temperature 68
Barometric Pressure 30
Static Pressure (P_s) T. 6
Filter Number(s) _____
Pretest Leak Rate = 0.00 cfm @ 14 in. Hg

	Read	and Record	all Data	Every	Minutes
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Post Test, Grant Leak Check

London

[illegible]

SAMPLE RECOVERY DATA

Plant: Wycoff
 Date: 8/1/85
 Sampling Location: Van Unit
 Sample Type: MMS
 Run Number: WC-V-MMS-2A
 Sample Box Number: -
 Clean-up Man: ALT
 Job Number: _____
 Comments: _____

FRONT HALF

Filter Number: N/A
 Description of Filter: _____

MOISTURE

Impingers

Final Volume:	<u>390</u>	ml	_____	ml	_____	ml
Initial Volume:	<u>0</u>	ml	<u>0</u>	ml	_____	ml
Net Volume:	_____	ml	_____	ml	_____	ml
Total H ₂ O:	_____		_____		<u>190</u>	

Silica Gel

Final Volume:	<u>484.0</u>	g	_____	g	_____	g
Initial Volume	<u>230.0</u>	g	<u>230.4</u>	g	_____	g
Net Volume	_____	g	_____	g	<u>23.4</u>	g
Total Moisture:	_____		_____		<u>213.6</u>	

Description of Impinger Catch: _____

SAMPLE RECOVERY DATA

Plant: W. H. Hoff
 Date: 9/1/75
 Sampling Location: Vac. Vind
 Sample Type: MMS
 Run Number: WC-U-MMS-2B
 Sample Box Number: —
 Clean-up Man: N/A
 Job Number: —
 Comments: —

FRONT HALF

Filter Number: N/A
 Description of Filter: —

MOISTURE

Impingers

Final Volume:	<u>375</u>	ml	<u>—</u>	ml	<u>—</u>	ml
Initial Volume:	<u>100</u>	ml	<u>100</u>	ml	<u>—</u>	ml
Net Volume:	<u>—</u>	ml	<u>—</u>	ml	<u>—</u>	ml
Total H ₂ O:	<u>—</u>		<u>—</u>		<u>175</u>	

Silica Gel

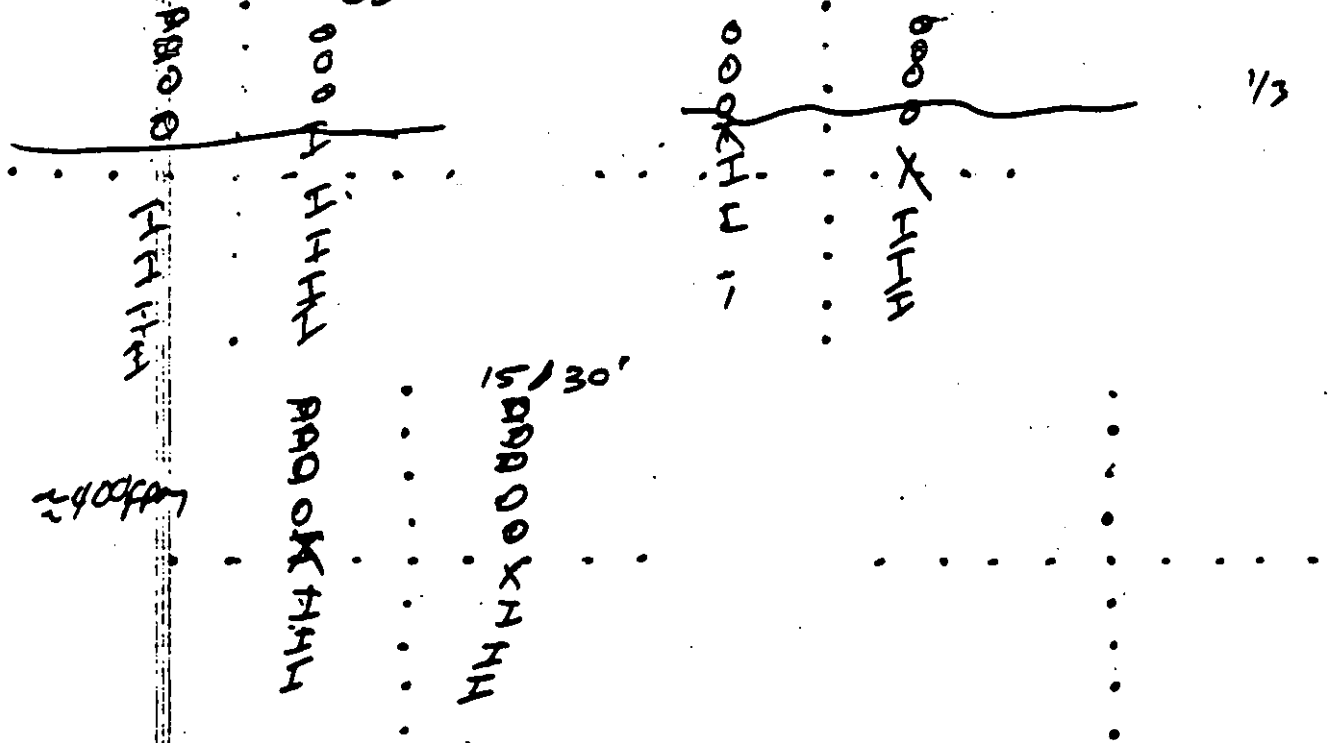
Final Volume:	<u>502.5</u>	g	<u>—</u>	g	<u>—</u>	g
Initial Volume	<u>244.7</u>	g	<u>236.6</u>	g	<u>—</u>	g
Net Volume	<u>—</u>	g	<u>—</u>	g	<u>265.9</u>	g
Total Moisture:	<u>—</u>		<u>—</u>		<u>234.7</u>	
					<u>196.1</u>	

Description of Impinger Catch: —

B-24

FLOW ESTIMATION CRO C 7/26/85

ANOMALOUS GND 6045 20mi
 05'00" 10'30" 2 400 fpm



B-2,5

CRSO

7

Stake

1350
1420

Anemometer 5 min
520

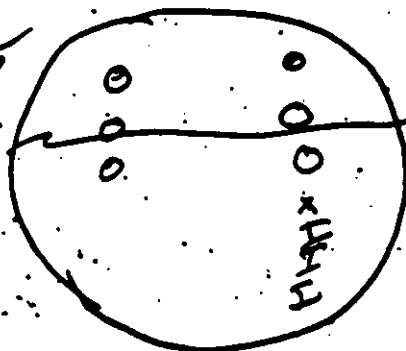
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min

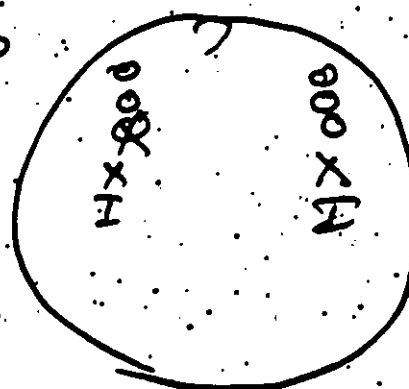
0 ft
9382 ft

22'00"

375
100 ft
15 sec



10

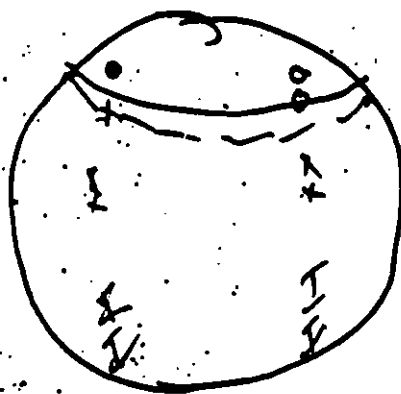


30

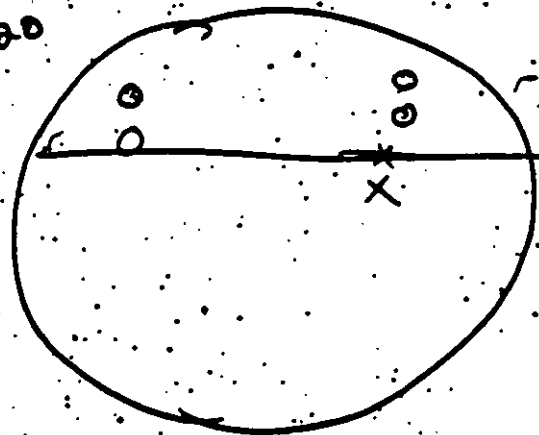
200 ft
36 sec

15

100 ft
21 sec



20



3

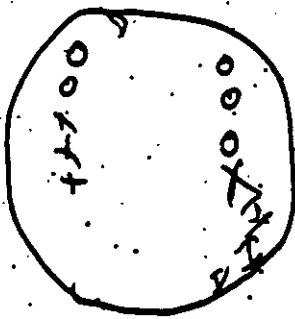
100 ft
18 sec

CRSO 0000

1350

23'00"

100 ft
18 sec

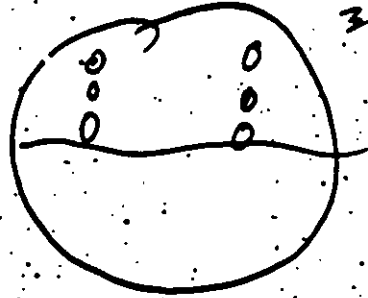


30

3/4

100
18 sec

B-2



30

6 Point Rotation Functions

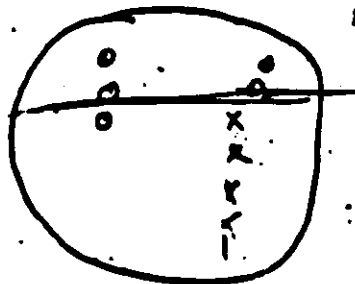
7/31/83

1152 Automation Start
END

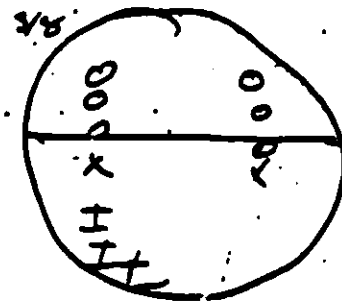
0 min
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50

0 ft
3
11135

0300
100 ft
19 sec

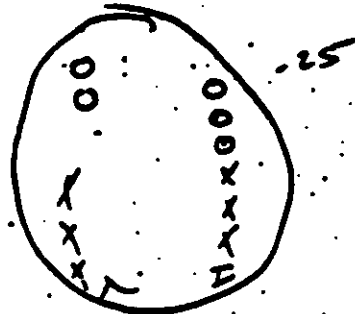


09

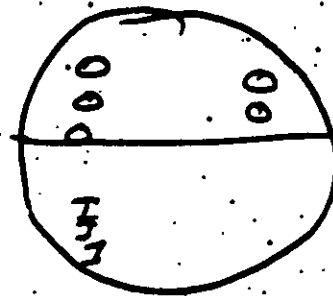


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17 sec

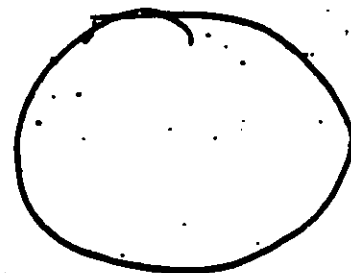
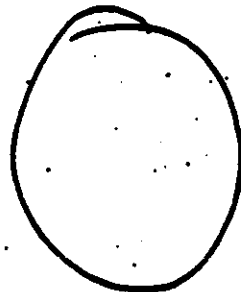
18'00"
100 ft
18 sec



25
14/3



100 ft
16 sec



B-27

PONTA
E

WYCKOPA

7/29/85

LOAD IN 0745

PILLOWS 0745 → 0845

START IN

0930

TIME

ROTOR TEMP

VAC

H₂O

GAUGES

1230

202

20

1245

204

20

1300

208

18

10"

1315

212

19

1330

212

17

11 1/2"

1345

211

19

1400

212

19

13

1415

212

19

1430

212

19

14

7/31

PONTA

ROTOR

E

P PSI

0900

239

35

0912

BLOWDOWN

0915

225

START

0920

212

0

VAC → 2

0935

206

17

(CHART + 4)

0945

201

19

0955

196

20

1005

191

21

1015

186

22

1025

1045

181

23

CROSS TO

E

1700

206

24

10 (CHART)

1715

210

24

1730

211

24

14 1/2

1745

210

24

1800

210

24

15

1815

210

24

15 1/2

1830

211

24

16 1/2

1328

WYCKOFF

2/20/85

REPORT P C2803078

START LINE 9244 0830

END OF LINE 9244 1030

FINAL VRC 229

T = 235 20ps1

B-29

4-10

8/1/85

CR50 VACUUM VENT

TIME		TEMP	VAC
1745		BLOWDOWN	STAY
1752	1800 INTO	VACUUM	
1800		200	14
1810		184	16
1820		181	20
1830		177	21
1840		176	22
1850		175	24
1900			

Run WCV-45 -2A, 2B

B-30

MASTER SAMPLE LOG

Plant W. C. C. Page of
 Location Bainbridge Island, WA Project Number

Sample Number	Run Code	Date	Sample Type	Container Type & Size	Disposition
WC-U-AAS 2A	-	8/1/85	FH AAS	450 L Bottle	
↓		↓	BH "	1 L Bottle	
↓		↓	Filter "	Petro. Dish	
↓		↓	XAD "	XAD Module	
WC-U-AAS 2B	-	8/1/85	FH AAS	450 L Bottle	
↓		↓	BH "	1 L Bottle	
↓		↓	F.H. "	Petro. Dish	
↓		↓	XAD "	XAD Module	
WC-U-AAS 3	-	8/1/85	FH ^{Reservoir} Blank	450 L Bottle	
↓		↓	BH	1 L Bottle	
↓		↓	Filter	Petro. Dish	
↓		↓	XAD	XAD Module	
WCF-XAD 1A	-		XAD Cartridge	Envelope	
WCF-XAD 1B	-		"	"	
WCF-X- 2A Front	-		"	"	
" Back	-		"	"	
WCF-X- 2B Front	-		"	"	
" Back	-		"	"	
WP-F-X 1A Front	-		"	"	
" Back	-		"	"	
WP-F-X 1B Front	-		"	"	
" Back	-		"	"	
WC-U-XZ Front	-		"	"	
" Back	-		"	"	

B-31

MASTER SAMPLE LOG

Plant Wycoff Page of
 Location Bainbridge Island, WA Project Number

Sample Number	Run Code	Date	Sample Type	Container Type & Size	Disposition
WP-V-MMS 1A	-	7/29/85	MMS F.H.	450 L Btl	
↓		↓	MMS B.H.	1 L Btl	
↓		↓	MMS Filter	Petri Dish	
↓		↓	MMS XAD	XAD Media	
WP-V-MMS 1B	-	7/29/85	MMS FH	450 L Btl	
↓		↓	" BH	1 L Btl	
↓		↓	" Filter	Petri Dish	
↓		↓	" XAD	XAD Media	
WP-V-MMS 2A	-	7/31/85	MMS F.H.	450 L Btl	
↓		↓	" B.H.	1 L Btl	
↓		↓	" Filter	Petri Dish	
↓		↓	" XAD	XAD Media	
WP-V-MMS 2B	-	7/31/85	MMS FH	450 L Btl	
↓		↓	" BH	1 L Btl	
↓		↓	" Filter	Petri Dish	
↓		↓	" XAD	XAD Media	
WC-V-MMS 1A	-	7/31/85	MMS FH	450 L Btl	
↓		↓	" BH	1 L Btl	
↓		↓	" Filter	Petri Dish	
↓		↓	" XAD	XAD Media	
WC-V-MMS 1B	-	7/31/85	MMS FH	450 L Btl	
↓		↓	" BH	1 L Btl	
↓		↓	" Filter	Petri Dish	
↓		↓	" XAD	XAD Media	

APPENDIX C
LABORATORY ANALYSIS

c-1

LABORATORY ANALYSIS REPORT

P.O. No. _____
Job No. 46921
Date Received 8-5-85
Date Reported _____

For Puget Sound / ES Fairfax Attention: John Bolstad

Address _____

Lab No.	<u>95207SE</u>	<u>952077F</u>	<u>952078E</u>	<u>952078 F+B</u>
Source of Sample	<u>WC-F-XAD</u>	<u>WC-F-XAD</u>	<u>WC-F-XAD</u>	<u>WC-F-XAD</u>
	<u>1A Front</u>	<u>2A Front</u>	<u>2A Back</u>	<u>2B Front + Back</u>
Date Collected:	<u>-</u>	<u>-</u>	<u>-</u>	<u>-</u>
Time Collected:	<u>-</u>	<u>-</u>	<u>-</u>	<u>-</u>

Analyses	Units	ANALYTICAL RESULTS			
Pentaabthophenol	ug	ND	ND	ND	ND
Acenaphthene		2200	1900		1500
Acenaphthylene		ND	90		60
Anthracene		100	180		20
Benz(a)anthracene		ND	ND		ND
Benz(a)fluoranthene					
Benz(b)fluoranthene					
Benz(e)pyrene					
Benz(a,b,i)perylene					
2-Chloronaphthalene					
Chrysene					
Dibenz(a,h)anthracene					
Fluoranthene		300	40		
Fluorene		240	580		340
Indeno(1,2,3-cd)pyrene		ND	ND		ND
Naphthalene		3000	11000	70	1200
Phenanthrene		640	200	ND	130
Pyrene		ND	20	ND	ND

COMMENTS:

THESE RESULTS WERE OBTAINED BY FOLLOWING ACCEPTED LABORATORY PROCEDURES.
THE LIABILITY OF THE CORPORATION SHALL NOT EXCEED THE AMOUNT PAID FOR THIS REPORT.

Laboratory Supervisor

C-1

LABORATORY ANALYSIS REPORT

P.O. No. _____
Job No. 46921
Date Received 8-5-85
Date Reported _____

For Puget Sound / ES Fairfax Attention: John Bolstad

Address _____

Lab No. _____

Source of Sample _____

Date Collected: _____

Time Collected: _____

<u>852079 F</u>	<u>852079 B</u>	<u>852080 FIB</u>	<u>852081 FIB</u>
<u>WP-V XAD</u>	<u>WP-V XAD</u>	<u>WP-V XAD</u>	<u>WC-V XAD</u>
<u>1A Front</u>	<u>1A Back</u>	<u>1B Front + Back</u>	<u>2 Front + Back</u>

Analyses	Units	ANALYTICAL RESULTS			
<u>Pentaabthophenol</u>	<u>ug</u>	<u>ND</u>	<u>ND</u>	<u>600</u>	<u>ND</u>
<u>Acanthophene</u>				<u>10</u>	<u>20</u>
<u>Acanthophene</u>				<u>ND</u>	<u>ND</u>
<u>Anthracene</u>					
<u>Benz(a) Anthracene</u>					
<u>Benz(a) Fluoranthene</u>					
<u>Benz(a) Fluoranthene</u>					
<u>Benz(a) pyrene</u>					
<u>Benz(a,h,i) pyrene</u>					
<u>2-Chloroanthracene</u>					
<u>Chrysene</u>					
<u>Dibenz(a,h)anthracene</u>					
<u>Fluoranthene</u>					
<u>Fluoranthene</u>				<u>20</u>	
<u>Indeno(1,2,3-cd) pyrene</u>				<u>ND</u>	
<u>Anthracene</u>			<u>50</u>	<u>200</u>	<u>7200</u>
<u>Phenanthrene</u>			<u>ND</u>	<u>10</u>	<u>ND</u>
<u>Pyrene</u>			<u>ND</u>	<u>ND</u>	<u>ND</u>

COMMENTS: _____

THESE RESULTS WERE OBTAINED BY FOLLOWING ACCEPTED LABORATORY PROCEDURES.
THE LIABILITY OF THE CORPORATION SHALL NOT EXCEED THE AMOUNT PAID FOR THIS REPORT.

Laboratory Supervisor _____



LABORATORY ANALYSIS REPORT

P.O. No. _____

Job No. 46921Date Received 8-5-85

Date Reported _____

For Puget Sound / ES FairfaxAttention John Bolstad

Address _____

Lab No. _____

Source of Sample _____

Date Collected: _____

Time Collected: _____

852065852067852069Blank 852074W-C-VW-C-VW-C-VBlankMMS-1AMMS-2AMMS-3

Analytes

Units detection
limits

ANALYTICAL RESULTS

Pentaabthiphenolug1000NDNDNDND (20)Acenaphthene3.6x10³8x10⁴Acenaphthylene24008200AnthraceneNDNDBenz(a)anthraceneBenz(a)fluorantheneBenz(a)phenanthreneBenz(a)pyreneBenz(a,b,i)perylene2-ChloroanthraceneChryseneDibenz(a,h)anthraceneFluorantheneFluoreneIndeno(1,2,3-cd)pyreneShanthalenePhenanthrenePyrene2.5x10⁶9.0x10⁵44014003400NDNDNDND

COMMENTS:

THESE RESULTS WERE OBTAINED BY FOLLOWING ACCEPTED LABORATORY PROCEDURES.
THE LIABILITY OF THE CORPORATION SHALL NOT EXCEED THE AMOUNT PAID FOR THIS REPORT.

Laboratory Supervisor

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LABORATORY ANALYSIS REPORT

P.O. No. _____
 Job No. 46921
 Date Received 8-5-85
 Date Reported _____

For Puget Sound / ES Fairfax Attention: John Bolstad

Address _____

Lab No. 852070 852072

Source of Sample W.P.V. W.P.V.

MMS-1A MMS-2A

Date Collected: _____

Time Collected: _____

Analysis	Units	Detection Limits	ANALYTICAL RESULTS		
<u>Pentachlorophenol</u>	<u>ug/g</u>	<u>ND</u>	<u>ND</u>		
<u>Arsenathione</u>		<u>3.0 x 10⁴</u>	<u>2.5 x 10⁴</u>		
<u>Arsenathylene</u>		<u>ND</u>	<u>ND</u>		
<u>Anthracene</u>					
<u>Benz(a) Anthracene</u>					
<u>Benz(b) Fluoranthene</u>					
<u>Benz(e) Fluoranthene</u>					
<u>Benz(a) Pyrene</u>					
<u>Benz(b,i) Pyrene</u>					
<u>2-Chloronaphthalene</u>					
<u>Chrysene</u>					
<u>Dibenz(a,h) Anthracene</u>					
<u>Fluoranthene</u>					
<u>Fluorene</u>		<u>8400</u>	<u>9800</u>		
<u>Indene(1,2,3-cd) Pyrene</u>		<u>ND</u>	<u>ND</u>		
<u>Naphthalene</u>		<u>2.0 x 10⁶</u>	<u>1.0 x 10⁶</u>		
<u>Phenanthrene</u>		<u>2.100</u>	<u>1700</u>		
<u>Pyrene</u>		<u>ND</u>	<u>ND</u>		

COMMENTS: _____

THESE RESULTS WERE OBTAINED BY FOLLOWING ACCEPTED LABORATORY PROCEDURES.
 THE LIABILITY OF THE CORPORATION SHALL NOT EXCEED THE AMOUNT PAID FOR THIS REPORT.

Laboratory Supervisor

C-4

APPENDIX D
CALCULATIONS

D-1

RUN NUMBER	MPV 1A	MPV 1B	MPV 2A	MPV 2B	WCV 1A	WCV 1B	WCV 2A	WCV 2B
METER VOLUME, ACF	59.403	60.779	73.376	74.512	52.666	47.046	36.822	36.326
METER PRESS(DH), IN	0.60	0.75	1.36	1.70	1.02	0.71	1.27	1.17
METER TEMP., DEG. F	96.7	95.9	85.2	94.8	90.3	78.0	94.2	74.6
BAROM PRESS, IN HG	29.9	29.9	29.9	29.9	29.9	29.9	29.9	29.9
GAMMA	1.007	1.013	1.007	1.013	1.013	1.007	1.013	1.007
METER VOL-STD, DSCF	56.996	58.770	72.057	72.360	51.477	46.721	35.759	36.346
CONDENSATE, ML	445.8	470.3	416.4	448.7	227.9	198.8	196.1	213.6
WATER VAPOR, SCF	20.984	22.137	19.600	21.120	10.727	9.358	9.230	10.054
MOISTURE CONTENT, %	26.9%	27.4%	21.4%	22.6%	17.2%	16.7%	20.5%	21.7%
MOLECULAR WT, DRY	28.84	28.84	28.84	28.84	28.84	28.84	28.84	28.84
MOLECULAR WT, WET	25.92	25.87	26.52	26.39	26.97	27.03	26.62	26.49

A5: 'RUN NUMBER
 B5: U ^MPV
 B6: (T) U ^1A
 A7: 'METER VOLUME, ACF
 B7: (F3) U 59.403
 A8: 'METER PRESS(DH), IN H2O
 B8: (F2) U 0.6
 A9: 'METER TEMP., DEG. F
 B9: (F1) U 96.7
 A10: 'BAROM PRESS, IN HG
 B10: U 29.9
 A11: 'GAMMA
 B11: (F3) U 1.007
 A14: 'METER VOL-STD, DSCF
 B14: (F3) +B7*B11*(530/(B9+460))*((B10+(B8/13.6))/(29.92)
 A15: 'CONDENSATE, ML
 B15: (F1) U 445.8
 A16: 'WATER VAPOR, SCF
 B16: (F3) +B15*0.04707
 A17: 'MOISTURE CONTENT, %
 B17: (P1) +B16/(B16+B14)
 A19: 'MOLECULAR WT, DRY
 B19: 28.84
 A20: 'MOLECULAR WT, WET
 B20: (F2) U 18*B17+((1-B17)*B19)

RUN NUMBER	WCF-X 1A	WCF-X 1B	WCF-X 1A	WCF-X 2A	WCF-X 2B
METER VOLUME, LITERS	9.835	7.575	15.240	10.22	0.18
METER VOLUME, ACF	0.348	0.268	0.539	0.361	0.289
METER PRESS(DH), IN H2O	1.00	1.18	1.00	0.97	1.21
METER TEMP., DEG. F	86.1	86.1	83.6	81.3	77.9
BAROM PRESS, IN HG	29.90	29.90	29.90	29.9	29.9
GAMMA	0.951	0.955	0.951	0.951	0.955
METER VOL-STD, DSCF	0.321	0.249	0.500	0.337	0.273

A5: 'RUN NUMBER
 C5: U 'WCF-X
 C6: (T) U '1A
 A7: 'METER VOLUME, LITERB
 C7: U 9.835
 A8: 'METER VOLUME, ACF
 C8: (F3) +C7/28.3
 A9: 'METER PRESS(DH), IN H2O
 C9: (F2) U 1
 A10: 'METER TEMP., DEG. F
 C10: (F1) U 86.1
 A11: 'BAROM PRESS, IN HG
 C11: (F2) U 29.9
 A12: 'GAMMA
 C12: (F3) U 0.951
 A15: 'METER VOL-STD, DSCF
 C15: (F3) +C8+C12*(530/(C10+460))*((C11+(C9/13.6))/29.92)

PROCESS CYCLE

WCV-1A

EXHAUST GAS FLOW
RATE, DSCFM

342

SAMPLE VOLUME

51.477

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	2,500,000	1.07E-04	323.20	2.20
ACENAPHTHYLENE	152.2	2,400	1.03E-07	0.26	0.00211
ACENAPHTHENE	154.2	3,600	1.54E-07	0.39	0.00316
PHENANTHRENE	178.2	1,400	6.06E-08	0.13	0.00123
FLUORANTHENE	202.3	(1,000)	-4.28E-08	-0.08	-0.00088
FLUORENE	166.2	4,700	2.01E-07	0.47	0.00413
PYRENE	202.1	(1,000)	-4.28E-08	-0.08	-0.00088
PENTACHLOROPHENOL	266.4	(1,000)	-4.28E-08	-0.08	-0.00088
ANTHRACENE	178.2	(1,000)	-4.28E-08	-0.08	-0.00088

PROCESS CYCLE

WCV-2A

EXHAUST GAS FLOW
RATE, DSCFM

319

SAMPLE VOLUME

35.759

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	900,000	5.55E-05	167.50	1.06
ACENAPHTHYLENE	152.2	0,200	5.06E-07	1.29	0.00968
ACENAPHTHENE	154.2	40,000	2.47E-06	6.19	0.04720
PHENANTHRENE	178.2	3,400	2.10E-07	0.46	0.00401
FLUORANTHENE	202.3	(1,000)	-6.17E-08	-0.12	-0.00118
FLUORENE	166.2	25,000	1.54E-06	3.59	0.02950
PYRENE	202.1	(1,000)	-6.17E-08	-0.12	-0.00118
PENTACHLOROPHENOL	266.4	(1,000)	-6.17E-08	-0.09	-0.00118

D-4

ANTHRACENE 178.2 (1,000) -6.17E-08 -0.13 -0.00118

PROCESS CYCLE

WPV-1A

EXHAUST GAS FLOW
RATE, DSCFM

493

SAMPLE VOLUME

56.996

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	2,000,000	7.74E-05	233.53	2.29
ACENAPHTHYLENE	152.2	(1,000)	-3.87E-08	-0.10	-0.00114
ACENAPHTHENE	154.2	30,000	1.16E-06	2.91	0.03432
PHENANTHRENE	178.2	2,100	8.12E-08	0.18	0.00240
FLUORANTHENE	202.3	(1,000)	-3.87E-08	-0.07	-0.00114
FLUORENE	166.2	8,400	3.25E-07	0.76	0.00961
PYRENE	202.1	(1,000)	-3.87E-08	-0.07	-0.00114
PENTACHLOROPHENOL	266.4	(1,000)	-3.87E-08	-0.06	-0.00114
ANTHRACENE	178.2	(1,000)	-3.87E-08	-0.08	-0.00114

PROCESS CYCLE

WPV-2A

EXHAUST GAS FLOW
RATE, DSCFM

509

SAMPLE VOLUME

72.057

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	1,000,000	3.06E-05	92.36	0.93
ACENAPHTHYLENE	152.2	(1,000)	-3.06E-08	-0.08	-0.00093
ACENAPHTHENE	154.2	25,000	7.65E-07	1.92	0.02336
PHENANTHRENE	178.2	1,700	5.20E-08	0.11	0.00159
FLUORANTHENE	202.3	(1,000)	-3.06E-08	-0.06	-0.00093
FLUORENE	166.2	9,600	3.00E-07	0.70	0.00916
PYRENE	202.1	(1,000)	-3.06E-08	-0.06	-0.00093

D-5

PENTACHLOROPHENOL	266.4	(1,000)	-3.06E-08	-0.04	-0.00093
ANTHRACENE	178.2	(1,000)	-3.06E-08	-0.07	-0.00093

PROCESS CYCLE WPF-X-1B

EXHAUST GAS FLOW
RATE, DSCFM 4478.04310

SAMPLE VOLUME 0.489

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	200	9.02E-07	2.72	0.24
ACENAPHTHYLENE	152.2	(10)	-4.51E-08	-0.11	-0.01211
ACENAPHTHENE	154.2	10	4.51E-08	0.11	0.01211
PHENANTHRENE	178.2	10	4.51E-08	0.10	0.01211
FLUORANTHENE	202.3	(10)	-4.51E-08	-0.09	-0.01211
FLUORENE	166.2	20	9.02E-08	0.21	0.02423
PYRENE	202.1	(10)	-4.51E-08	-0.09	-0.01211
PENTACHLOROPHENOL	266.4	600	2.71E-06	3.93	0.72679
ANTHRACENE	178.2	(10)	-4.51E-08	-0.10	-0.01211

PROCESS CYCLE WCF-X-2A

EXHAUST GAS FLOW
RATE, DSCFM 3763.36551

SAMPLE VOLUME 0.337

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	1,070	7.24E-05	218.61	16.35
ACENAPHTHYLENE	152.2	90	5.89E-07	1.50	0.13294
ACENAPHTHENE	154.2	1,900	1.24E-05	31.19	2.80659
PHENANTHRENE	178.2	300	1.96E-06	4.26	0.44312
FLUORANTHENE	202.3	40	2.62E-07	0.50	0.05909
FLUORENE	166.2	580	3.79E-06	8.83	0.85675

D-6

PYRENE	202.1	20	1.31E-07	0.25	0.02954
PENTACHLOROPHENOL	266.4	(10)	-6.54E-08	-0.10	-0.01477
ANTHRACENE	178.2	100	6.54E-07	1.42	0.14772

PROCESS CYCLE WCF-X-26

EXHAUST GAS FLOW
RATE, DSCFM 3763.36551

SAMPLE VOLUME 0.273

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	1,200	9.69E-06	29.25	2.19
ACENAPHTHYLENE	152.2	60	4.85E-07	1.23	0.10941
ACENAPHTHENE	154.2	1,500	1.21E-05	30.40	2.73516
PHENANTHRENE	178.2	130	1.05E-06	2.20	0.23705
FLUORANTHENE	202.3	(10)	-8.08E-08	-0.15	-0.01823
FLUORENE	166.2	340	2.75E-06	6.39	0.61997
PYRENE	202.1	(10)	-8.08E-08	-0.15	-0.01823
PENTACHLOROPHENOL	266.4	(10)	-8.08E-08	-0.12	-0.01823
ANTHRACENE	178.2	20	1.62E-07	0.35	0.03647

PROCESS CYCLE WCF-X-1AF

EXHAUST GAS FLOW
RATE, DSCFM 4487.08965

SAMPLE VOLUME 0.321

COMPOUND	MW	ug	#/DSCF	PPMV	#/HR
NAPHTHALENE	128.2	3,000	2.06E-05	62.20	5.55
ACENAPHTHYLENE	152.2	(10)	-6.87E-08	-0.17	-0.01843
ACENAPHTHENE	154.2	2,200	1.51E-05	37.92	4.06781
PHENANTHRENE	178.2	640	4.40E-06	9.55	1.10336
FLUORANTHENE	202.3	300	2.06E-06	3.94	0.55470

D-7

FLUORENE	166.2	740	5.00E-06	11.83	1.36426
PYRENE	202.1	(10)	-6.87E-08	-0.13	-0.01849
PENTACHLOROPHENOL	266.4	(10)	-6.87E-08	-0.10	-0.01849
ANTHRACENE	178.2	100	6.87E-07	1.49	0.18490

D-8

END

DATE

FILMED

2-12-87

NTIS

ESTIMATE OF VOLUME OF WOOD TREATED
DURING WYCKOFF TEST

$$\text{VOL. OF RESORT} = \pi r^2 h = \pi (3.75)^2 (130) = 5743$$

5740 ft³

$d = 90" = 7.5'$ $r = 3.75'$
 $h = 130'$

50% OF RESORT VOLUME = VOL. OF WOOD TREATED

$5740 \times 50\% = 2870 \text{ ft}^3 \text{ WOOD / BATCH}$

Ref
(2)

USE 50%
to get vol. wood
treated

calc. vol. of
resort

$\pi r^2 h$

Conditioning or vacuum control?

N/K report doesn't say

pressure - no control

Test Runs?

acc. - pug. - 4
 cond - 3
 final vac - 2

PCP - pug - 2
 cond - 2
 final vac - 2

CREOSOTE CYCLE (TABLE 5.10a)

REFERENCE 2
WYCKOFF, INC.

ESTIMATE OF TOTAL LBS. EMITTED DURING ENTIRE CYCLE

	COND. (48 HR)	FINAL VAC. (1 HR)	FUG. (1/2 HR)
Naphthalene	$\frac{1.06}{\sqrt{5088}}$	$\sqrt{1.06}$	$\frac{4.68}{\sqrt{2.34}}$
Acenaphthalene	$\frac{0.101}{\sqrt{4.85}}$	$\sqrt{0.0097}$	$\frac{0.613}{\sqrt{0.31}}$
Acenaphthene	$\frac{0.154}{\sqrt{7.39}}$	$\sqrt{0.0472}$	$\frac{1.40}{\sqrt{0.7}}$
Phenanthrene	$\frac{0.0576}{\sqrt{2.76}}$	$\sqrt{0.004}$	$\frac{0.172}{\sqrt{0.086}} \quad 0.09$
Fluoranthene	$\frac{<0.0432}{\sqrt{<2.07}}$	$\sqrt{<0.0012}$	$\frac{0.0195}{\sqrt{0.00975}} \quad 0.01$
Fluorene	$\frac{0.197}{\sqrt{9.46}}$	0.0295	$\frac{0.373}{\sqrt{0.19}}$
Pyrene	$\frac{<0.0432}{\sqrt{<2.07}}$	$\sqrt{<0.0012}$	$\frac{0.012}{\sqrt{0.006}}$
Pentachlorophenol	$\frac{<0.0432}{\sqrt{<2.07}}$	$\sqrt{<0.0012}$	$\frac{0.0083}{\sqrt{0.004}}$
Anthracene	$\frac{<0.0432}{\sqrt{<2.07}}$	$\sqrt{<0.0012}$	$\frac{<0.0465}{\sqrt{<0.02}}$

RSD = 1b/batch (Final vac = same #)

$$\frac{2870 \text{ ft}^3/\text{batch}}{1 \text{ ft}^3} = 1 \text{ b/ft}^3$$

Calculate 1b/batch

then

1b/ft³ wood

1 put in Table

6 TABLES - 3 creosote
3 PCP

(COND, VAC, FUG)

RCP Cycle (Table 5.10.b)
(lb/hr)

Compd	COND (48 HRS)	Final VAC. (1 HR)	FUG (1/2 HR)
Naphthalene	$\frac{110}{\sqrt{5280}}$	$\sqrt{0.930}$	$\sqrt{0.120}$ 0.00
Acenaphthalene	$\frac{<0.0528}{\sqrt{<2.53}}$	$\sqrt{<0.0009}$	$\frac{<0.006}{\sqrt{<0.0006}}$ <0.003
Acenaphthene	$\frac{1.65}{\sqrt{79.2}}$	$\sqrt{1.920}$	$\sqrt{0.006}$ 0.003
Phenanthrene	$\frac{0.115}{\sqrt{5.52}}$	$\sqrt{0.110}$	$\sqrt{0.006}$ 0.003
Fluoranthene	$\frac{<0.0528}{\sqrt{<2.53}}$	$\sqrt{<0.0009}$	$\sqrt{<0.006}$ <0.003
Fluorene	$\frac{0.461}{\sqrt{22.13}}$	$\sqrt{0.0092}$	$\sqrt{0.012}$ 0.006
Pyrene	$\frac{<0.0528}{\sqrt{<2.53}}$	$\sqrt{<0.0009}$	$\sqrt{<0.006}$ <0.003
Pentachlorophenol	$\frac{<0.0528}{\sqrt{<2.53}}$	$\sqrt{<0.0009}$	$\sqrt{0.360}$ 0.18
Anthracene	$\frac{<0.0528}{\sqrt{<2.53}}$	$\sqrt{<0.0009}$	$\sqrt{<0.006}$ <0.003

($<$ = below detection limit)

Ref = lb/batch (final vac. = same H)

÷ 2870 ft³/batch