

IMC FERTILIZER GROUP • MINERALS DIVISION
P.O. Box 867 • Bartow, Florida 33830
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PM
14 Dec. 1987
Bartow, FL
CM: # P387-961-036

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IMC FERTILIZER, INC.

INTERNATIONAL MINERALS & CHEMICAL CORPORATION

December 11, 1987

Mr. John Reynolds
Bureau of Air Quality Management
Department of Environmental Regulation
2600 Blair Stone Road
Twin Towers Office Building
Tallahassee, Florida 32301

Dear Mr. Reynolds:

Re: IMC Lonesome Dryer Application

Based on recent discussion, I am enclosing the following items for your reference in the consideration of the review of this permit application.

- 1. Copy of the Supplement to Emission Tests done in June.
- 2. The most recent test report for 11/5/87.
- 3. Waiver request until 1/13/88.

original copies on loan to John Reynolds.

The operation of this facility has only been in 1987. During this time, the only material dried has been 73 BPL Concentrate rock. This is the material for all testing that has been done.

Thank you for your attention in this matter. If you have further requirements, please contact me.

Very truly yours,

C. D. Turley
C. D. Turley

cc: M. Brandon - USEPA. w/ attachments.
R. S. Hearon

Copied: John Reynolds

BT/CHF

Bill Thomas - SWD.st

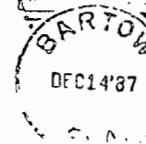
Jerry Campbell - HC/PC

12-18-87

Per John Reynolds - 12-18-87

no attachments (1/2) were mailed to the Dist. cc's

happy
holidays



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IMC FERTILIZER, INC.
INTERNATIONAL MINERALS & CHEMICAL CORPORATION

P.O. BOX 867 @ BARTOW, FLORIDA 33830

C. D. TURLEY - ENVIRON. SERVICES

TO:

MR. JOHN REYNOLDS
BUREAU OF AIR QUALITY MGT.
DEPT. OF ENVIRON. REG.
2600 BLAIR STONE ROAD
TWIN TOWERS OFFICE BLDG.
TALLAHASSEE, FL 32301

FORM 5-422 PRINTED IN U.S.A.

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of the return address*

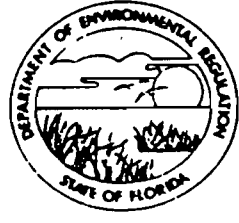
CERTIFIED

P 387 961 036

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STATE OF FLORIDA
DEPARTMENT OF ENVIRONMENTAL REGULATION



SOUTHWEST DISTRICT

4520 OAK FAIR BLVD.
TAMPA, FLORIDA 33610-9544
813-623-5561
SunCom—552-7612

BOB MARTINEZ
GOVERNOR

DALE TWACHTMANN
SECRETARY

DR. RICHARD D. GARRITY
DISTRICT MANAGER

WAIVER OF 90 DAY TIME LIMIT
UNDER SECTIONS 120.60(2) AND 403.0876, FLORIDA STATUTES

License (Permit, Certification) Application No. A029-111119
A029-111120

Applicant's Name: IMC Fertilizer, Inc.

The undersigned has read Sections 120.60(2) and 403.0876, Florida Statutes, and fully understands the applicant's rights under that section.

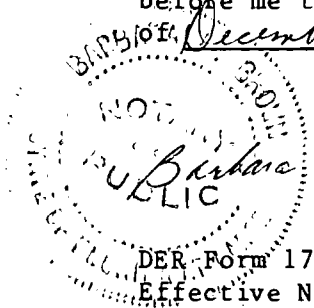
With regard to the above reference license (permit, certification) application, the applicant hereby with full knowledge and understanding of (his) (her) (its) rights under Sections 120.60(2) and 403.0876, Florida Statutes, waives the right under Sections 120.60(2) and 403.0876, Florida Statutes, to have the application approved or denied by the State of Florida Department of Environmental Regulation within the 90 day time period prescribed in Sections 120.60(2) and 403.0876, Florida Statutes. Said waiver is made freely and voluntarily by the applicant, is in (his) (her) (its) self-interest, and without any pressure or coercion by anyone employed by the State of Florida Department of Environmental Regulation.

This waiver shall expire on the 13th day of January 1988.

The undersigned is authorized to make this waiver on behalf of the applicant.

W. C. Cross
Signature
Vice Pres. & Gen. Mgr.
W. C. Cross-Fla. Minerals Operations
Please Type Name of Signee
12/11/87
Date

Sworn to and subscribed
before me this 14th day
of December 1987.



Notary Public, State of Florida at Large
My Commission Expires Feb. 13, 1991

Section 120.60, Florida Statutes

(2) When an application for a license is made as required by law, the agency shall conduct the proceedings required with reasonable dispatch and with due regard to the rights and privileges of all affected parties or aggrieved persons. Within 30 days after receipt of an application for a license, the agency shall examine the application, notify the applicant of any apparent errors or omissions, and request any additional information the agency is permitted by law to require. Failure to correct an error or omission or to supply additional information shall not be grounds for denial of the license unless the agency timely notified the applicant within this 30 day period. The agency shall notify the applicant if the activity for which he seeks a license is exempt from the licensing requirement and return any tendered application fee within 30 days after receipt of the original application or within 10 days after receipt of the timely requested additional information or correction of errors or omissions. Every application for license shall be approved or denied within 90 days after receipt of the original application or receipt of the timely requested additional information or correction of errors or omissions unless a shorter period of time for agency action is provided by law. The 90-day or shorter time period shall be tolled by the initiation of a proceeding under Section 120.57 and shall resume 10 days after the recommended order is submitted to the agency and the parties. Any application for a license not approved or denied within the 90-day period or shorter time period, within 15 days after conclusion of a public hearing held on the application, or within 45 days after the recommended order is submitted to the agency and the parties, whichever is latest, shall be deemed approved and, subject to the satisfactory completion of an examination, if required as prerequisite to licensure, the license shall be issued. The Public Service Commission, when issuing a license, and any other agency, if specifically exempted by law, shall be exempt from the time limitations within this subsection. Each agency, upon issuing or denying a license, shall state with particularity the grounds or basis for the issuance or denial of same, except where issuance is a ministerial act. On denial of a license application on which there has been no hearing, the denying agency shall inform the applicant of any right to a hearing pursuant to Section 120.57.

Section 403.0876, Florida Statutes

Permits; processing. ---Within 30 days after receipt of an application for a permit under this chapter, the department shall review the application and shall request submittal of all additional information the department is permitted by law to require. If the applicant believes any departmental request for additional information is not authorized by law or departmental rule, the applicant may request a hearing pursuant to s. 120.57. Within 30 days after receipt of such additional information, the department shall review it and may request only that information needed to clarify such additional information or to answer new questions raised by or directly related to such additional information. If the applicant believes the request of the department for such additional information is not authorized by law or departmental rule, the department, at the applicant's request, shall proceed to process the permit application. Permits shall be approved or denied within 90 days after receipt of the original application, the last item of timely requested additional material, or the applicant's written request to begin processing the permit application.

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DER
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EMISSION TESTS

FOR

PARTICULATE
OPACITY
SULFUR DIOXIDE
NITROGEN OXIDES

OF

IMC LONESOME
NO 1 DRYER SCRUBBER

PERMIT NOS AO29-111119
AND PSD-FL-088

CONDUCTED BY:

ENVIRONMENTAL SERVICES
IMC FERTILIZER, INC.
MINERALS OPERATIONS

AND

AIR CONSULTING AND ENGINEERING

Prepared by:

C. D. Turley

C. D. Turley

December 11, 1987

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- C-2 PROCESS CALIBRATION

I. INTRODUCTION

A. SUMMARY RESULTS

This report summarizes the emission tests conducted for the No 1 Dryer located at the IMC Lonesome Mine in Hillsborough County. The test was conducted on November 5, 1987 according to applicable USEPA and FDER sampling procedures. The purpose of the test was to establish the compliance status of this unit with the emission limitations contained in FDER Permit No A029-111119 and USEPA Permit No PSD-FL-088.

TABLE 1 RESULTS

Parameter	Unit	Run 1	Run 2	Run 3	Average	Limit
Temperature	deg F	155.0	155.0	154.0	154.7	
Pressure	in Hg	29.9	29.9	29.9		
Moisture	%V	28.0%	28.8%	28.0%	28.2%	
Volumetric Flow	acfm	100747	98619	96845	98737	182100
	scfm	62237	60202	59909	60783	
Process Wt. Rate	TPH	503	505	507	505	513
	%HOH wt	12.0%	11.8%	11.8%		
	TPH dry	443	446	447	445	450
	GPH	771	630	606	669	
	MMBtu/hr	115	95	90	100	190
Particulate	gr/scf	0.0088	0.0059	0.0050	0.0066	
	lb/hr	4.7	3.0	2.6	3.4	25
	lb/ton	0.011	0.007	0.006	0.008	0.06
Opacity	%	0%				10%
Sulfur Dioxide	ppm	27.9	36.9	25.0	29.9	
	lb/hr	16.4	20.3	13.9	16.9	110
	lb/MBtu	0.14	0.21	0.15	0.17	1.1
Nitrogen Oxides (as NO2)	ppm	77.0	82.0	82.0	80.3	81
	lb/hr	32.5	32.4	32.8	32.6	35.3

B. PARTICIPANTS

The Particulate and Opacity testing were conducted by personnel of the IMC Environmental Services Department. The gaseous emissions were determined by personnel of Air Consulting and Engineering.

Testers	Title	Affiliation
L. W. Bradley	Environmental Technician	IMC
D. A. Gilley	Environmental Technician	IMC
J. D. Norman	Environmental Technician	IMC
J. F. Tanner	Environmental Technician	IMC
C. D. Turley	Environmental Engineer	IMC
S. L. Neck	Environmental Engineer	ACE See Section IV
S. J. Carter	Environmental Technician	ACE See Section IV

Observer

M. D. Silcott	HCEPC
C. Gonzalez	HCEPC

C. TEST PROTOCOL

The test consisted of three individual runs which were organized in the following fashion:

Run 1 Length: 1 hour.

- Additional Method 2 Traverse.
- Particulate and opacity.
- Two 20 minute SO₂ samples.
- Operation of continuous analyzer for NO_x.

Run 2 Length: 1 hour.

- Additional Method 2 Traverse.
- Particulate and opacity.
- Two 20 minute SO₂ samples.
- Operation of continuous analyzer for NO_x.

Run 3 Length: 1 hour.

- Additional Method 2 Traverse.
- Particulate and opacity.
- Two 20 minute SO₂ samples.
- Operation of continuous analyzer for NO_x.

II. SOURCE DESCRIPTION

A. PROCESS DESCRIPTION

The IMC Lonesome No 1 Dryer is used to dry phosphate rock prior to shipping. The rock moves from wet rock storage piles located southwest of the dryer by conveyor belt to a feed hopper located above the dryer. The dryer is a fluid bed type fired with No 6 Oil. The exhaust gases from the dryer pass through four parallel cyclones, the I.D. fan and then to the scrubber where they vent to the atmosphere through the scrubber stack after final cleaning. The dried rock is transported by conveyor to the dry rock storage bins until shipment. The dryer is shown schematically in the following figure.

Dust is generated as the hot gases fluidize the feed rock within the dryer during operation. The cyclones and scrubber then remove the particulate matter generated. During the emission test, the dryer was operating at a rate of 505 TPH wet, 445 TPH dry (see following Section B).

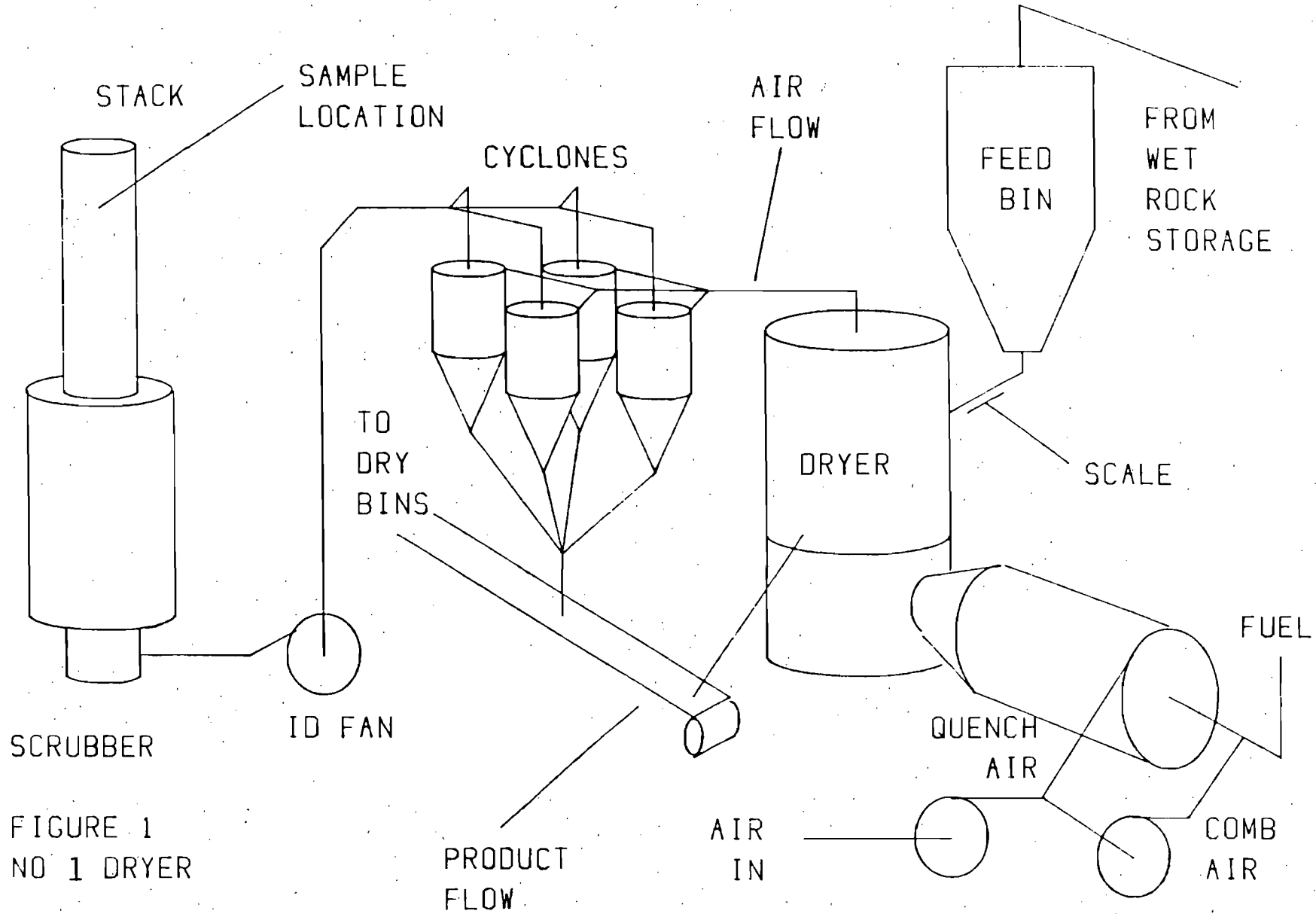


FIGURE 1
 NO 1 DRYER

B. PRODUCTION DETERMINATIONS AND PROCESS STATEMENT

1. TONNAGE

The dryer is equipped with a variable speed conveyor used to control the rock feed rate to the dryer. The feed rate is automatically controlled based on a required temperature of the rock exiting the dryer. The conveyor is also equipped with a belt scale used to determine the dryer wet feed tonnage. This measurement is not used in the operating control system of the dryer. The procedure and calibration are included in Appendix C-2. The calibration was within the required 5% limit.

TABLE 2 PRODUCTION TONNAGE

SCALE READINGS

	TIME		SCALE TONNAGE		TPH	%HOH	dry
	Start	End	Start	End			TPH
RUN 1	09:48	12:06	2306	3464	503	12.0%	443
RUN 2	12:06	13:53	3464	4365	505	11.8%	446
RUN 3	13:53	15:22	4365	5117	507	11.8%	447
OVERALL	09:48	15:22	2306	5117	505	11.9%	445

2. FUEL USAGE

The fuel used during this test was No 6 Oil. Analysis is included in Appendix B-2. One sample of the oil was collected during each run of the test. The dryer is equipped with a combustion burner designed to fire either oil or a coal-oil-mixture (COM). The dryer has two fuel supply systems for that burner. The original system for oil, prior to the COM conversion, incorporated an oil meter in the supply line to the burner. The new system for the COM or oil did not contain a meter. The system with the meter was used for the test. This meter is a positive displacement type. The readings and calculations are presented in the following table:

TABLE 3 FUEL OIL USAGE

	TIME		METER READINGS		GPH
	Start	End	Start	End	
RUN 1	09:40	12:07	312829	313018	771
RUN 2	12:07	13:28	313018	313103	630
RUN 3	13:28	15:14	313103	313210	606
OVERALL	09:40	15:14	312829	313210	684

3. SCRUBBER

The following scrubber parameters were measured during the test. The scrubber pressure drop was measured using a pressure transduced measurement. The readings are included in the following table:

TABLE 4 SCRUBBER MEASUREMENTS

	PSI	GPM	Del P	pH
RUN 1	92	525	22.0	6.2
RUN 2	92	525	22.5	6.4
RUN 3	92	525	22.5	6.4

4. OPERATIONAL PROBLEM

During this test, the primary fan damper for the dryer could not be properly opened. This condition apparently deteriorated during the test of the dryer scrubber. The result of this condition meant that the amount of quench air supplied to the dryer was restricted during the test. This damper has been replaced since this test. The effect of this is demonstrated by comparing the average dscfm of 88,000 in the initial 3/4 test with the average of 61,000 of this test. Because of the intermittent operation of this dryer, this problem had not been obvious prior to the scheduled test.

INTERNATIONAL MINERALS AND CHEMICAL CORPORATION

PROCESS INFORMATION AND CALCULATION

VERSION

PLANT: FT LONESOME PREPARATION STACK: NO 2 DRYER SCRUBBER

TYPE OF CONTROL DEVICE: WET SCRUBBER

TYPE OF EMISSION TEST: PARTICULATE TESTER: J.D. NORMAN, L BRADLEY
I. FANNING, D. JURLEY (ENG)

DATE: 11/5/87 SAMPLING TIME: 5.5 HOURS

PERMIT NO.: A029-11119 EXEMPTION
ASP ORDER NO.: PSD-FL-088

THE OPERATION OF THIS SYSTEM HAS BEEN VERIFIED DURING THE TIME OF TESTING AS PER THE SUBMITTED PROCEDURE RELATIVE TO THE ABOVE ORDER.

SIGNED: John D. Norman

PROCESS CALCULATION

SOURCE OF INFORMATION: DRYER SCALE

TIME	SCALE READING (TONS)
9:45 -	2306
3:15 -	5017
<u>5.50 HR</u>	2811 TONS DRIED

511 TPH
5.5 HRS 2811 TONS

AVG PROCESS RATE: 511 TPH MATERIAL: PHOSPHATE ROCK

EMISSION STANDARD: PERMIT CONDITIONS ALLOWABLE: 25 LB/HR

CALCULATED BY: J.D. Norman DATE: 11/12/87

PROCESS STATEMENT SIGNATURE: John D. Norman

I CERTIFY THAT THE ABOVE STATEMENT IS TRUE AND CORRECT TO THE BEST OF MY KNOWLEDGE. TITLE: Foreman

DATE: 11-13-87

III. PARTICULATE EMISSIONS

A. SAMPLE METHODS

The particulate test consisted of the three sample runs conducted according to the Methods:

Method 1:

Section 2.2.1. 12 point traverse shown in Figure 2. These points were also used for the intermediate traverses between runs.
Section 2.4. Absence of Cyclonic flow verified.

Method 2:

Section 2.3. Thermocouple used. Calibrations in Appendix C-1.
Section 4.1.2.1. Calibration duct has 12 in. diameter.
Section 4.1.2.3. Calibration system not capable of 5000 fpm, current maximum = 3800 fpm, for interval calibration.
Section 4.3. Laboratory grade thermometer used for calibration. Pre-test calibration procedures from EPA/600/4-77/027b vol. III used.

Method 3:

Section 1.1. Fyrite O₂ and CO₂ Gas Analyzers used.
Section 2.1. Figure 3-1, Grab sampling train used.
Section 2.1.1. Stainless steel probe length = 38 inches.

Method 4:

Section 1.2. Lange's equation used for calculation of HOH saturation content based on actual stack pressure in stead of Saturation Table. Traverse point temperatures used to calculate saturation at each point in calculations. In all final calculations, the saturation %HOH was used instead of measured values.

Method 5:

Section 2.1. Lear-Siegler Method 5 particulate train with IMC design modifications. The train is shown in Figure 3.
Section 2.1.2. Stainless steel probe liners used.
Section 2.1.5. Filter holder material unknown, frit is stainless steel, and gasket is hi-temp neoprene.
Section 2.1.7. Condenser used in lieu of glass impingers. Approved per letter dated 4/30/87, JTW to CDT.
Section 2.2.3. 250 ml glass containers with glass stoppers used.
Section 2.2.4. Filter petri dishes are plastic -- 150 x 15 mm.
Section 2.2.6. Silica gel stored and transported in drying tube.
Section 2.3.1. Disposal aluminum weighing dish used, tare = 2.5 g.
Section 4.1.4.2. The train remained connected as it was moved between sampling ports.
Section 4.1.5. Isokinetic conditions were maintained during sampling using a TI CC40 portable computer.
Section 4.3. Container No 1. Filters were dried at 104 °C for final weighing. Cooled in desiccator for ten minutes prior final weighing.
Section 4.3. Container No 2. Liquid equalized to etched volume mark, volume is reduced in container and then transferred to weighing dish.

Section 4.3. Container No 3. Not used. Drying tube used to transport silica gel.

Section 5. The equipment calibrations are contained in Appendix C-1.

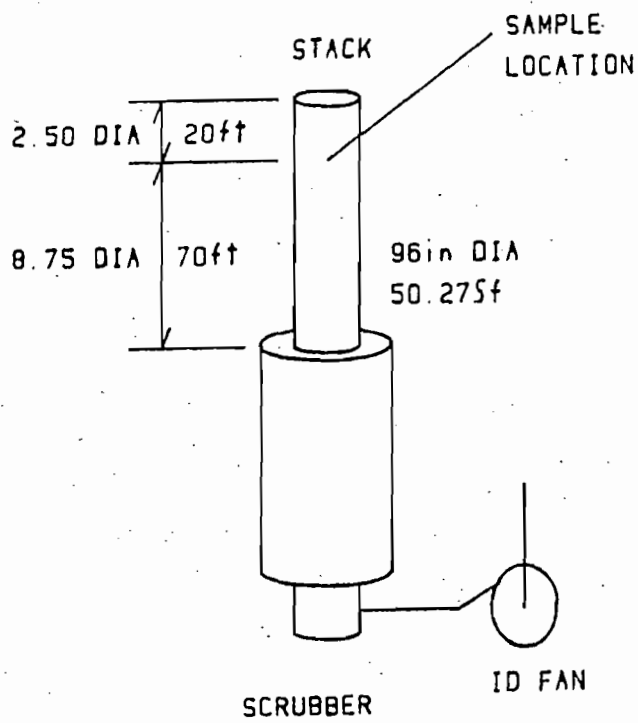
Section 5.4. Probe temperature measured directly behind the filter frit.

Section 6.5. The stack gas moisture content was verified against that of saturation calculated based on stack temperature and pressure.

Section 6.6. Acetone blank concentration not calculated. Wash weights not adjusted for positive blank determinations.

Method 9:

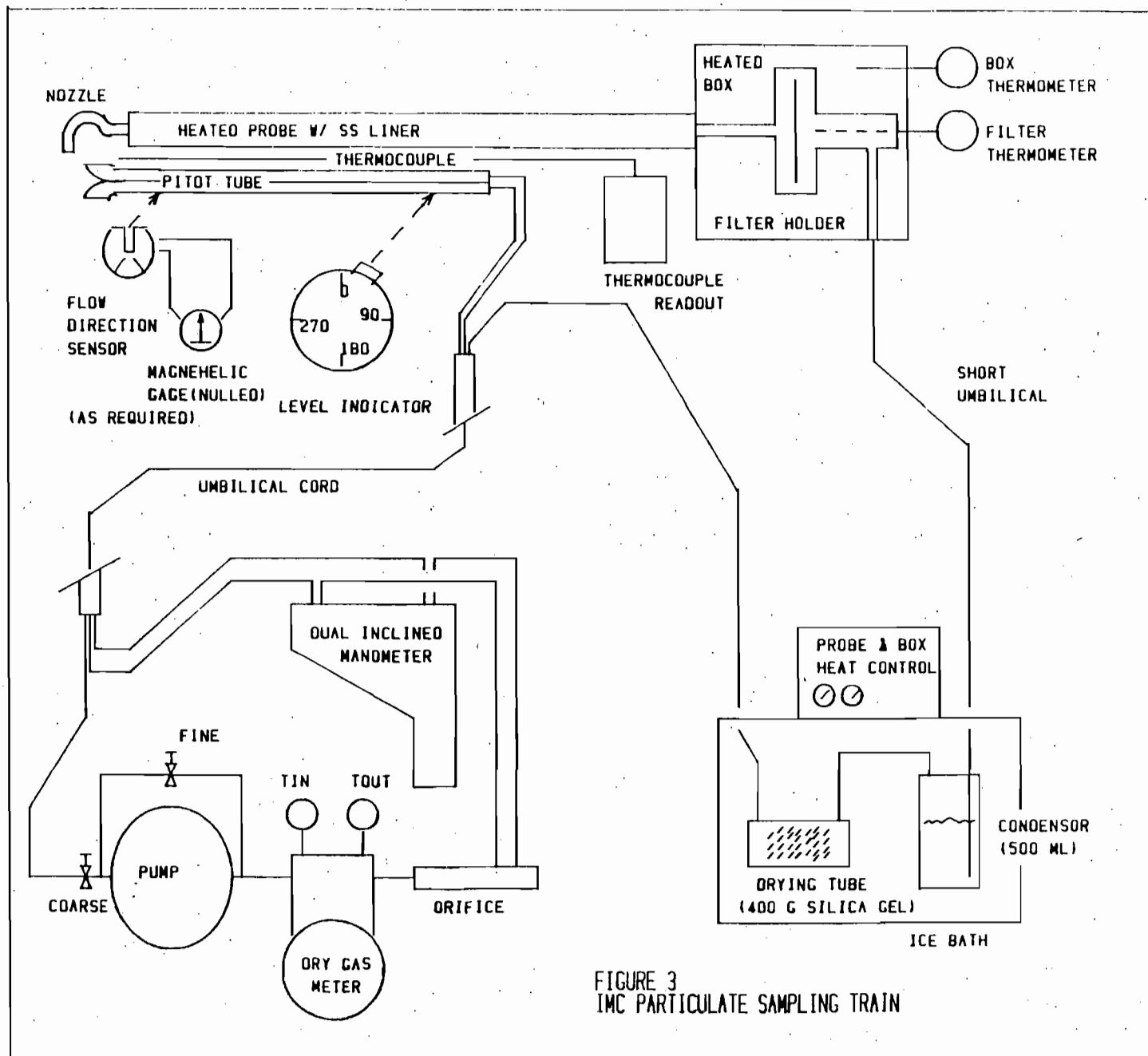
Section 2.4. Reading time of one hour used.



TRAVERSE POINTS

W-1	S-1	4 1/8 in
W-2	S-2	14
W-3	S-3	28 3/8
W-4	S-4	67 9/16
W-5	S-5	81 15/16
W-6	S-6	91 13/16

FIGURE 2 NO 1 DRYER TRAVERSE



B. SAMPLE RESULTS

1. VELOCITY TRAVERSE RESULTS

The test plan included three velocity traverses of the scrubber as means of confirming the particulate test traverses. The results of those determinations are presented for comparison in the following table. The volumetric flow is shown based on saturation at the stack temperature and as measured for comparison of the test run results.

TABLE 5 VELOCITY TRAVERSE RESULTS

No	1F	RUN 1	2F	RUN 2	3F	RUN 3	4F
TIME	09:20	10:35	11:35	12:00	13:21	14:05	15:18
Pbar	29.9	29.9	29.9	29.9	29.9	29.9	29.9
Pstk	29.88	29.88	29.88	29.88	29.88	29.88	29.88
Tdry	155	155	156	155	154	154	150
Twet	155	155	156	155	154	154	150
Cp	0.868	0.834	0.868	0.834	0.868	0.834	0.868
OO	14.5%	14.5%	14.5%	14.5%	14.5%	14.5%	14.5%
OCO	4.5%	4.5%	4.5%	4.5%	4.5%	4.5%	4.6%
%HOH sat	28.6%	28.0%	29.4%	28.8%	28.0%	28.0%	25.3%
SCFM sat	70412	62895	61844	61145	61026	60680	61813
%HOH msd		32.5%		35.7%		33.8%	
SCFM msd		62237		60202		59909	
NOx ppm	75	77	81	82	82	82	82

VELOCITY TRAVERSE CALCULATIONS

The following pages are the calculation of each individual traverse conducted for this test. The particulate test traverses were measured in a plane parallel to the stack wall. For the auxiliary traverses, an IMC design sensor was used to determine the cyclonic angles at each point. These angles are included in each of the Traverse Calculation pages and used in comparative calculation of the various emission rates. These pages also contain a comparison calculation of the individual traverse point velocity based on moisture content determined from the point stack temperature and velocities based on Method 2 calculations. The results were not significantly different. Summary results are based on the Method 2 equations.

Individual Velocity Traverse Calculations

No: B 1F
 Date: 11/05
 Time: 09:20
 Pbar: 29.9
 Pst: -0.3
 Ps: 29.88 = Pbar+Pst/13.6
 Cp: 0.868
 Fv: 13.58 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.20%
 O: 0.64%
 C: 87.39% %EA: 2.518 based on fuel analysis and %OO

Usages
 Rock: 509 TPH
 0.12 HOH
 448 Dry
 Oil: 771 GPH
 8.138 lb/gal
 18291 Btu/lb
 115 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		155.0			
		sat %HOH:		28.65%			
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.06			
Avg:	18.3	155.0	28.65%				
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.32	22	155	28.65%	37.3	37.3	34.6
2	0.30	20	155	28.65%	36.1	36.1	33.9
3	0.32	10	155	28.65%	37.3	37.3	36.7
4	0.35	18	155	28.65%	39.0	39.0	37.1
5	0.36	12	155	28.65%	39.6	39.6	38.7
6	0.32	12	155	28.65%	37.3	37.3	36.5
SW-1	0.27	28	155	28.65%	34.3	34.3	30.3
2	0.35	19	155	28.65%	39.0	39.0	36.9
3	0.32	15	155	28.65%	37.3	37.3	36.0
4	0.34	19	155	28.65%	38.5	38.5	36.4
5	0.40	24	155	28.65%	41.7	41.7	38.1
6	0.38	20	155	28.65%	40.7	40.7	38.2
				avg	38.2	38.2	36.1
				fps:	115111	115111	108926
				acfm:	28.65%	28.65%	
				%HOH:	70412	70412	66629
				scfm:	lb/hr		
				Emission:	std	point	cyc
NOx	ppm	Mgas	pVM/RT*60	37.8	37.8	35.8	
CO	0	28	4.3610	0.0	0.0	0.0	
VOC	0	44	6.8531	0.0	0.0	0.0	
SO2	0.0	64	9.9681	0.0	0.0	0.0	
PART	0.009 gr/scf		0.0001	5.3	5.3	5.0	

Individual Velocity Traverse Calculations

No: B 2F
 Date: 11/05
 Time: 11:35
 Pbar: 29.9
 Pst: -0.3
 Ps: 29.88 = Pbar+Pst/13.6
 Cp: 0.868
 Fv: 13.58 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.20%
 O: 0.64%
 C: 87.39% %EA: 2.518 based on fuel analysis and %OO

Rock: Usages
 505 TPH
 0.118 HOH
 446 Dry

Oil: 630 GPH
 8.138 lb/gal
 18291 Btu/lb
 94 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		156.1			
		sat %HOH:		29.41%			
Ms = Mdry*(1-%HOH)+18*%HOH				25.98			
Avg:	13.0	156.1	Ms:	29.41%			
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.22	20	156	29.35%	31.0	31.0	29.1
2	0.26	16	156	29.35%	33.7	33.7	32.4
3	0.25	14	157	30.07%	33.1	33.1	32.1
4	0.23	0	156	29.35%	31.7	31.7	31.7
5	0.26	10	156	29.35%	33.7	33.7	33.2
6	0.30	16	156	29.35%	36.2	36.2	34.8
SW-1	0.26	24	156	29.35%	33.7	33.7	30.8
2	0.25	18	156	29.35%	33.1	33.1	31.4
3	0.28	12	156	29.35%	35.0	35.0	34.2
4	0.28	6	156	29.35%	35.0	35.0	34.8
5	0.30	12	156	29.35%	36.2	36.2	35.4
6	0.28	8	156	29.35%	35.0	35.0	34.6
				avg			
				fps:	33.9	33.9	32.9
				acfm:	102376	102375	99185
				%HOH:	29.41%	29.41%	
				scfm:	61844	61843	59916
				Emission:	lb/hr		
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	81	46	7.1646	35.9	35.9	34.8	
CO	0	28	4.3610	0.0	0.0	0.0	
VOC	0	44	6.8531	0.0	0.0	0.0	
SO2	0.0	64	9.9681	0.0	0.0	0.0	
PART	0.009	gr/scf	0.0001	4.7	4.7	4.5	

Individual Velocity Traverse Calculations

No: B 3F
 Date: 11/05
 Time: 13:21
 Pbar: 29.9
 Pst: -0.3
 Ps: 29.88 = Pbar+Pst/13.6
 Cp: 0.868
 Fv: 13.58 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.19%
 O: 0.81%
 C: 87.32% %EA: 2.520 based on fuel analysis and %OO

Rock: 505 TPH
 0.118 HOH
 446 Dry

Oil: 630 GPH
 8.138 lb/gal
 18462 Btu/lb
 95 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:	154.0				
		sat %HOH:	27.96%				
Ms = Mdry*(1-%HOH)+18*%HOH			Ms:	26.14			
Avg:		14.2	154.0	27.96%			
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.24	18	154	27.96%	32.2	32.2	30.7
2	0.23	14	154	27.96%	31.6	31.6	30.6
3	0.26	8	154	27.96%	33.5	33.5	33.2
4	0.28	20	154	27.96%	34.8	34.8	32.7
5	0.24	14	154	27.96%	32.2	32.2	31.3
6	0.26	10	154	27.96%	33.5	33.5	33.0
SW-1	0.23	18	154	27.96%	31.6	31.6	30.0
2	0.22	14	154	27.96%	30.9	30.9	29.9
3	0.24	10	154	27.96%	32.2	32.2	31.7
4	0.24	18	154	27.96%	32.2	32.2	30.7
5	0.28	12	154	27.96%	34.8	34.8	34.1
6	0.25	14	154	27.96%	32.9	32.9	31.9
				avg			
				fps:	32.7	32.7	31.7
				acfm:	98651	98651	95466
				%HOH:	27.96%	27.96%	
				scfm:	61026	61026	59055
				lb/hr			
			Emission:				
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	82	46	7.1646	35.9	35.9	34.7	
CO	0	28	4.3610	0.0	0.0	0.0	
VOC	0	44	6.8531	0.0	0.0	0.0	
SO2	0.0	64	9.9681	0.0	0.0	0.0	
PART	0.006	gr/scf	0.0001	3.1	3.1	3.0	

Individual Velocity Traverse Calculations

No: B 4F			Usages
Date: 11/05		Rock: 507	TPH
Time: 15:18		0.118	HOH
Pbar: 29.9		447	Dry
Pst: -0.3			
Ps: 29.88 = Pbar+Pst/13.6		Oil: 606	GPH
Cp: 0.868		8.138	lb/gal
Fv: 13.58 = 85.49*Cp*sqrt(1/Ps)		18354	Btu/lb
Dia: 96	Area: 50.27	90	mmBtu/hr
%OO: 14.5%			
%OCO: 4.6%			
Mdry: 29.31			
H: 10.48%			
O: 0.67%			
C: 87.41%	%EA: 2.442 based on fuel analysis and %OO		

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

		Assumed Twb:	150.0				
		sat %HOH:	25.34%				
Ms = Mdry*(1-%HOH)+18*%HOH			26.44	Ms:			
Avg:	17.9		150.0	25.34%			
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.21	18	150	25.34%	29.9	29.9	28.4
2	0.24	15	150	25.34%	31.9	31.9	30.9
3	0.22	6	150	25.34%	30.6	30.6	30.4
4	0.24	20	150	25.34%	31.9	31.9	30.0
5	0.25	22	150	25.34%	32.6	32.6	30.2
6	0.24	18	150	25.34%	31.9	31.9	30.4
SW-1	0.24	16	150	25.34%	31.9	31.9	30.7
2	0.25	17	150	25.34%	32.6	32.6	31.2
3	0.24	20	150	25.34%	31.9	31.9	30.0
4	0.22	21	150	25.34%	30.6	30.6	28.6
5	0.24	20	150	25.34%	31.9	31.9	30.0
6	0.26	22	150	25.34%	33.2	33.2	30.8
				avg			
				fps:	31.8	31.8	30.1
				acfm:	95791	95791	90878
				%HOH:	25.34%	25.34%	
				scfm:	61813	61813	58643
				Emission:	lb/hr		
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	82	46	7.1646	36.3	36.3	34.5	
CO	0	28	4.3610	0.0	0.0	0.0	
VOC	0	44	6.8531	0.0	0.0	0.0	
SO2	0.0	64	9.9681	0.0	0.0	0.0	
PART	0.005	gr/scf	0.0000	2.6	2.6	2.5	

2. PARTICULATE RESULTS

The following pages are the completed particulate test results. These results are calculated by the IMC Main computer. The equations used by the program are presented in the fashion by which they are executed by the program. Note that the summary results in this report have been independently calculated as a means to double check results.

The equations are those contained in USEPA Methods 2 and 5. The Isokinetic Rate equation for the test has been changed. The equivalency of the program formula used to that of USEPA Method 5 is also shown. The formula used adjusts for the moisture content used in calculation in the case of a saturated wet stack with entrained droplets. The Isokinetic Rate is also calculated as the individual traverse point Isokinetic Rates which are included. The Isokinetic Rates for the test are indicated on the calculation summary sheet.



JOB: BA680
DATE: 11/13/87

INTERNATIONAL MINERALS & CHEMICAL CORPORATION
SOURCE SAMPLING CALCULATION REPORT

IMC SOURCE NO. 07100
OPTION NO. 000

TEST OF STACK.....NO 1 DRYER SCRUBBER.

AT PLANT.....LONESOME PREPARATION

CONDUCTED ON..... 11/05/87

D A T A S U M M A T I O N S

PARAMETER	UNITS	RUN 1	RUN 2	RUN 3
BAROMETRIC PRESSURE	IN OF HG	29.90	29.90	29.90
STATIC PRESSURE	IN OF HOH	0.30-	0.27-	0.30-
STACK PRESSURE	IN OF HG	29.88	29.88	29.88
AVERAGE SQRT DELTA-P	IN HOH1/2	0.528	0.516	0.508
AVERAGE DELTA-H	IN OF HOH	0.510	0.480	0.465
AVERAGE METER TEMP	DEGREES R	554.5	560.7	556.8
AVERAGE STACK TEMP	DEGREES R	614.1	615.2	614.0
METERED SAMPLE VOL	CUBIC FT	23.20	22.10	22.80
VOLUME OF WATER	CUBIC FT	10.750	11.693	11.175
TOTAL PARTICULATE	MG	12.7	8.0	7.1
AVERAGE % O-O	%	14.5	14.4	14.5
AVERAGE % C-O-O	%	4.5	4.5	4.5
AVERAGE % N-N + C-O	%	81.0	81.0	80.9
PITOT COEFFICIENT	UNITY	0.83	0.83	0.83
NOZZLE DIAMETER	IN	0.238	0.238	0.238
STACK AREA	SQUARE FT	50.270	50.270	50.270
M.C.F.	UNITY	1.01	1.01	1.01
TRAVERSE POINTS	UNITY	12	12	12
SAMPLING TIME	MINUTES	60	60	60

E. P. A. C A L C U L A T I O N S

VARIABLE	UNITS	RUN 1	RUN 2	RUN 3
STANDARD SAMPLE VOLUME	CUBIC FT	22.329	21.035	21.850
PROPORTION OF WATER	UNITY	0.325	0.357	0.338
PERCENT WATER VAPOR	%	28.02*	28.76*	27.96*
MOLECULAR WT OF DRY GAS	LB/LB-MOLE	29.430	29.434	29.440
MOLECULAR WT OF STK GAS	LB/LB-MOLE	26.227	26.145	26.241
ACTUAL STK GAS VELOCITY	FT/SEC	33.35	32.64	32.05
ACTUAL STK GAS FLOW	CU-FT/MIN	100,594	98,441	96,656
STANDARD STK GAS FLOW	CU-FT/MIN	62,182	60,119	59,804
STD STK GAS VELOCITY	FT/MIN	1,237	1,196	1,190
PART. CONCENTRATION.	GRAIN/CU-FT	.0088	.0059	.0050
PART. STACK EMISSIONS	LB/HOUR	4.68	3.02	2.57
ISOKINETIC RATE	%	97.4	94.9	99.1

*CALCULATED MOISTURE EXCEEDS SATURATION.
SATURATION VALUE ASSUMED.



JOB: BA680
DATE: 11/13/87
TEST OF STACK.....NO 1 DRYER SCRUBBER

INTERNATIONAL MINERALS & CHEMICAL CORPORATION
VELOCITY TRAVERSE CALCULATIONS
AT PLANT.....LONESOME PREPARATION

IMC SOURCE NO. 07100
OPTION NO. 000
CONDUCTED ON..... 11/05/87

-----RUN 1-----					-----RUN 2-----					-----RUN 3-----				
ELAP TIME (MIN)	TRV PT ID	STACK VELOCITY (FT/SEC)	NOZZLE FLOW (CU-FT/MIN)	ISOKINETIC RATE (PERCENT)	TRV PT ID	STACK VELOCITY (FT/SEC)	NOZZLE FLOW (CU-FT/MIN)	ISOKINETIC RATE (PERCENT)	TRV PT ID	STACK VELOCITY (FT/SEC)	NOZZLE FLOW (CU-FT/MIN)	ISOKINETIC RATE (PERCENT)		
05	S-1	31.58	0.60	103.	W-1	31.65	0.22	38.	S-1	31.54	0.60	102.		
10	S-2	31.50	0.54	92.	W-2	31.65	0.60	102.	S-2	31.54	0.62	106.		
15	S-3	31.50	0.56	96.	W-3	31.65	0.60	102.	S-3	31.54	0.56	96.		
20	S-4	31.50	0.53	91.	W-4	31.65	0.56	96.	S-4	31.54	0.59	101.		
25	S-5	42.23	0.81	103.	W-5	34.67	0.62	97.	S-5	34.55	0.62	97.		
30	S-6	42.23	0.74	95.	W-6	34.67	0.65	102.	S-6	34.55	0.65	101.		
35	W-1	31.60	0.56	96.	S-1	31.65	0.60	102.	W-1	31.54	0.59	102.		
40	W-2	31.60	0.56	96.	S-2	31.65	0.59	101.	W-2	31.54	0.56	95.		
45	W-3	31.60	0.59	101.	S-3	31.65	0.56	95.	W-3	31.54	0.59	100.		
50	W-4	31.60	0.59	101.	S-4	31.60	0.59	100.	W-4	31.54	0.55	95.		
55	W-5	31.60	0.59	100.	S-5	34.56	0.65	101.	W-5	31.54	0.58	100.		
60	W-6	31.60	0.56	95.	S-6	34.56	0.65	101.	W-6	31.54	0.55	94.		
**AVERAGES		33.35	0.60	97.		32.64	0.57	95.		32.05	0.59	99.		

RESULT EQUATIONS AND CALCULATIONS

Average Meter Temperature

$$T_m = \frac{1}{n} \sum_{i=1}^n (T_{in_i} + T_{out_i}) + 440$$

- T_m = Average Meter Temperature, °R
- T_{in_i} = Inlet Temperature at each Traverse Point, °F
- T_{out_i} = Outlet Temperature at each Traverse Point, °F
- n = Number of Traverse Point Readings

Average Meter Pressure Drop

$$\Delta H = \frac{1}{n} \sum_{i=1}^n \Delta H_i$$

- ΔH = Average Meter ΔH , Inch H₂O
- ΔH_i = Traverse Point ΔH Reading, Inch H₂O

Standard Sample Volume

$$V_s(Std) = V_m \cdot \frac{P_{bar} + 29.92}{19.92}$$

- $V_s(Std)$ = Standard Sample Volume, ft³ @ 14.7 psia, 70°F
- V_m = Metered Volume ($V_{load} - V_{leak}$), ft³
- P_{bar} = Barometric Pressure, Inch Hg

Standard Meter Volume Sampled

$$V_m(Std) = 0.0474 V_{li}$$

- $V_m(Std)$ = Volume of Meter Collected, ft³ @ 14.7 psia, 70°F
- V_{li} = Liquid Volume collected, ml

Proportion of Water Vapor

$$W_{H_2O} = V_m(Std) / (V_m(Std) + V_s(Std))$$

Average Stack Temperature

$$T_s = \frac{1}{n} \sum_{i=1}^n T_{s_i} + 460$$

- T_s = Average Stack Temperature, °R
- T_{s_i} = Stack Temperature at each Traverse Point, °F

Percent Moisture Vapor

- $\% H_2O$ = minimum ($\Delta H_p \times 100$); (Saturation)
- $\% H_2O$ = Moisture Content of Stack Gas
- Saturation = Moisture Content of Gas at Saturation for the Calculated Stack Temperature based on the following formulas:
 1. From Lange's Handbook of Chemistry, pg. 1436
 - $\log_{10} P_s = 4.10765 + 1730.180 / (233.0 + T)$ T = 60°C
 - $\log_{10} P_s = 7.96661 - 1668.21 / (228.0 + T)$ 60 + 2 = 130°C
 - T = Stack Temperature, °C
 - P_s = Saturation Pressure of Water, mmHg

- 2. Solving for P_s
 - Saturation = $100 \times P_s / (P_{bar} + P_{stc})$
 - all Pressure in equivalent units
 - Average Oxygen and Carbon Dioxide Content

$$\% O_2 = \frac{1}{n} \sum_{i=1}^n \% O_{2_i} \quad 21.0\% \text{ if } n = 0$$

$$\% CO_2 = \frac{1}{n} \sum_{i=1}^n \% CO_{2_i} \quad 1.0\% \text{ if } n = 0$$

- $\% O_2$ = Average Oxygen Content, %. This defaults to 21.0% if no readings are made.
- $\% CO_2$ = Average Carbon Dioxide Content, %. This defaults to 1.0% if no readings are made.
- n = Number of Readings made of Individual Constituent.
- $\% O_{2_i}$ = Individual Oxygen Reading
- $\% CO_{2_i}$ = Individual Carbon Dioxide Reading

Dry Molecular Weight of Stack Gas

$$M_d = 0.441 \% CO_2 + 0.322 \% O_2 + 0.28(2 \% H_2 + \% CO)$$

- M_d = Dry Molecular Weight, lb/lb-mole, 29.9 lb/lb-mole, etc. is calculated by default values of % CO₂ and % O₂. If M_d is calculated, it is assumed for remaining runs when no value given.

$$(\% H_2 + \% CO) = 100 - (\% O_2 + \% CO_2)$$

Molecular Weight of Stack Gas

$$M_s = M_d(1 - \% H_2O/100) + 18.12 \% H_2O/100$$

- M_s = Stack gas molecular weight, lb/lb-mole.

Average Square-root of Stack Velocity Reading

$$(\sqrt{SP})_{ave} = \frac{1}{n} \sum_{i=1}^n \sqrt{SP_i}$$

$$(\sqrt{SP})_{ave} = \text{Average square root of } SP, \sqrt{\text{Inch H}_2\text{O}}$$

- n = Number of Positive Traverse Point Readings
- SP_i = Traverse Point SP Reading, Inch H₂O

Stack Pressure

$$P_s = P_{bar} + P_{stc} / 13.6$$

- P_s = Absolute Stack Pressure, Inch Hg
- P_{stc} = Static Stack Pressure, Inch H₂O

Average Stack Velocity

$$V_s = K_s C_p (\sqrt{SP})_{ave} \sqrt{\frac{T_s}{P_s M_s}}$$

- V_s = Stack Velocity, Eps.
- C_p = Pitot Coefficient, unitless.
- $K_s = 85.48 \text{ ft/sec} (\text{lb/lb-mole})^{-1/2}$

Actual Stack Flow Rate

$$Q_s = 80 A_s V_s$$

- Q_s = Stack Flow Rate, acfm
- A_s = Area of Stack, ft²

Standard Stack Flow Rate

$$Q_s(Std) = Q_s (530/T_s) (P_s / 29.92) (1 - \% H_2O / 100)$$

$$Q_s(Std) = \text{Standard Stack Flow Rate, acfm}$$

Particulate Concentration

$$C_p = (M_s / 1000) (7700 / 453.6) / V_s(Std)$$

- C_p = Particulate Concentration, g/ft³
- M_s = Total Particulate Catch, Probe Wash + Filter Catch, mg

Emission Rate

$$pmr = C_p Q_s(Std) (80/7000)$$

- pmr = Particulate Emission Rate, lb/hr.

Isokinetic Rate

$$I = \frac{100 V_m(\text{std})}{A_m \bar{v}_m(\text{std})}$$

I - Average Isokinetic Rate, I. This equation is equivalent to that used in USEPA Method 3. This equation automatically compensates for the moisture content determined.

T - Total Sampling Time, min.

A_m - Area of Nozzle, ft²

v_m(std) - Standard Stack Velocity, Q_s(std)/A_s, fpm.

Traverse Point Calculation

Nozzle Flow

$$Q_{n_i} = (V_{n_i} - v_{n_i-1}) \left(\frac{T_{n_i}}{T_{n_i-1} - \text{moist}_i} \right) \left(\frac{P_{\text{bar}} + 14.7/13.6}{P_{n_i}} \right) \sqrt{1 - \frac{\% \text{H}_2\text{O}}{100}} c_i$$

Q_{n_i} - Nozzle Flow Rate, acfm.

t_i - Traverse Point Time, min.
Stack Velocity

$$V_{n_i} = 87.44 C_p \sqrt{\frac{v_{n_i} - P_{n_i}}{P_{n_i} A_n}}$$

v_{n_i} - Traverse Point Velocity, fpm.

Isokinetic Rate

$$I_i = (Q_{n_i} / 60 A_n) 100 / v_{n_i}$$

I_i - Traverse Point Isokinetic Rate, I.

Isokinetic Rate Equation Equivalency

$$I = 100 v_m(\text{std}) / A_m \bar{v}_m(\text{std})$$

$$= 100 v_m(\text{std}) / A_m \bar{Q}_s(\text{std}) / A_s$$

$$= \frac{100 \bar{v}_m(\text{std})}{A_m \bar{A}_s \bar{v}_s \left(\frac{P_{\text{bar}}}{P_s} \right) \frac{1}{29.32} \left(1 - \frac{\% \text{H}_2\text{O}}{100} \right) / c_i}$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{510}{29.32}$$

$$A_m \bar{v}_s \left(\frac{P_s}{P_m} \right) \frac{1}{29.32} \left(1 - \frac{\% \text{H}_2\text{O}}{100} \right)$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \left(\frac{P_s}{P_m} \right) \left(1 - \frac{\% \text{H}_2\text{O}}{100} \right)$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} \left(1 - \frac{\% \text{H}_2\text{O}}{100} \right) \frac{v_m(\text{std})}{v_m(\text{std}) - v_s(\text{std})}$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} \left(\frac{v_m(\text{std})}{v_m(\text{std}) - v_s(\text{std})} \right)$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} (v_m(\text{std}) - v_s(\text{std})) / v_m(\text{std})$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} \frac{v_m(\text{std})}{v_m(\text{std})} + 100 v_s \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} \frac{v_s(\text{std})}{v_m(\text{std})}$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} + 100 v_s \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} \left(\frac{v_s(\text{std})}{v_m(\text{std})} \right) \left(\frac{510}{29.32} \right) / v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} \left(\frac{1}{29.32} \right)$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= 100 v_m \left(\frac{P_s}{P_m} \right) \frac{P_s}{P_m} + \frac{P_s}{P_m} \left(\frac{v_s(\text{std})}{v_m(\text{std})} \right) 100$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$= \frac{100 \left(\frac{P_s}{P_m} \right) \left(v_m \frac{P_s}{P_m} + \frac{v_s(\text{std})}{v_m(\text{std})} \right)}$$

$$A_m \bar{v}_s \bar{Q}_s$$

$$I = 1.447 \frac{P_s}{P_m} \left(\frac{v_s(\text{std})}{v_m(\text{std})} + \frac{v_m}{P_{\text{bar}} + 14.7/13.6} \right)$$

$$A_m \bar{v}_s \bar{Q}_s$$

NOTE:

These two equations are equivalent in the case when the measured moisture content, based on T_g, does not exceed saturation moisture content. In the case when it does exceed it, the final formula does not produce the true isokinetic rate of sampling.

3. OPACITY RESULTS

The Visible Emissions were determined during the first particulate test run. The field sheet follows.

VISIBLE EMISSION OBSERVATION					START	DOWN FROM					READING				
					END	TIME TO									
MIN	00	15	30	45	MIN	00	15	30	45	MIN	00	15	30	45	
0:	0	0	0	0	20:	0	0	0	0	40:	0	0	0	0	
1:	0	0	0	0	21:	0	0	0	0	41:	0	0	0	0	
2:	0	0	0	0	22:	0	0	0	0	42:	0	0	0	0	
3:	0	0	0	0	23:	0	0	0	0	43:	0	0	0	0	
4:	0	0	0	0	24:	0	0	0	0	44:	0	0	0	0	
5:	0	0	0	0	25:	0	0	0	0	45:	0	0	0	0	
6:	0	0	0	0	26:	0	0	0	0	46:	0	0	0	0	
7:	0	0	0	0	27:	0	0	0	0	47:	0	0	0	0	
8:	0	0	0	0	28:	0	0	0	0	48:	0	0	0	0	
9:	0	0	0	0	29:	0	0	0	0	49:	0	0	0	0	
10:	0	0	0	0	30:	0	0	0	0	50:	0	0	0	0	
11:	0	0	0	0	31:	0	0	0	0	51:	0	0	0	0	
12:	0	0	0	0	32:	0	0	0	0	52:	0	0	0	0	
13:	0	0	0	0	33:	0	0	0	0	53:	0	0	0	0	
14:	0	0	0	0	34:	0	0	0	0	54:	0	0	0	0	
15:	0	0	0	0	35:	0	0	0	0	55:	0	0	0	0	
16:	0	0	0	0	36:	0	0	0	0	56:	0	0	0	0	
17:	0	0	0	0	37:	0	0	0	0	57:	0	0	0	0	
18:	0	0	0	0	38:	0	0	0	0	58:	0	0	0	0	
19:	0	0	0	0	39:	0	0	0	0	59:	0	0	0	0	

PLANT: FT. LONESOME PREP STACK: NO. 7 DRYER SCRUBBER

PERMIT NO: A029-11119 STACK HT: 125' STACK DIA: 96"

PLUME COLOR: WHITE PLUME TYPE: TRAILING

BACKGROUND COLOR: GREY SKY CONDITIONS: OVERCAST 1/951

WIND DIRECTION: NW WIND SPEED: 5-10 AMBIENT TEMPERATURE: 80°F

DISTANCE TO STACK: 200 yds DIRECTION TO STACK: NW

OBSERVER: J. Tanner DATE: 11/5/87

STACK LAYOUT SKETCH



EMISSION POINT

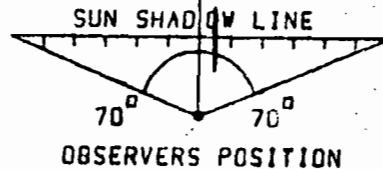
DRAW NORTH ARROW

STATE OF FLORIDA DEPARTMENT OF ENVIRONMENTAL REGULATION

THIS IS TO CERTIFY THAT

JERRY F. TANNER HAS COMPLETED THE STATE OF FLORIDA VISIBLE EMISSIONS EVALUATION TRAINING EPA REFERENCE METHOD 9. THIS CERTIFICATE EXPIRES Mar 9, 1988

J. F. Tanner BEARER'S SIGNATURE



OBSERVERS POSITION

INTERNATIONAL MINERALS AND CHEMICAL CORPORATION

PROCESS INFORMATION AND CALCULATION

VERSION 1

PLANT: FT LONESOME PREP STACK: NO 7 DRYER SCRUBBER

TYPE OF CONTROL DEVICE: WET SCRUBBER

TYPE OF EMISSION TEST: VISIBLE TESTER: J. TANNER

DATE: 11/5/87 SAMPLING TIME: 60 MINUTES

PERMIT NO.: A029-11119 EXEMPTION ASP ORDER NO.: PSD-FL-088

THE OPERATION OF THIS SYSTEM HAS BEEN VERIFIED DURING THE TIME OF TESTING AS PER THE SUBMITTED PROCEDURE RELATIVE TO THE ABOVE ASP ORDER.

SIGNED: J. O. Norman

PROCESS CALCULATION

SOURCE OF INFORMATION: DRYER SCALE

TIME	SCALE READING (TONS)
9 ⁴⁵	2306
3 ¹⁵	5117
5.5 HR	2811 TONS DRIED

5.5 HOURS ^{511 T.P.H.} 2811 TONS DRIED

AVG PROCESS RATE: 511 TPH MATERIAL: PHOSPHATE ROCK

EMISSION STANDARD: PERMIT CONDITIONAL ALLOWABLE: 10% OPACITY

CALCULATED BY: J. O. Norman DATE: 11/12/87

PROCESS STATEMENT

SIGNATURE: Marvin E. Driggs

I CERTIFY THAT THE ABOVE STATEMENT IS TRUE AND CORRECT TO THE BEST OF MY KNOWLEDGE.

TITLE: Supervisor

DATE: 11-20-87

C. ANALYSES

Three analyses are performed as a laboratory operation for each test. Type A/E fiberglass filters are desiccated and pre-weighed. The silica gel is pre-weighed and placed in the train drying tubes. These items are then placed in the IMC sampling trailer for the next sample to be conducted. Once the sampling has been completed, the recovered filters, the silica gel tubes, and the acetone washings are removed from the trailer for final processing. The acetone wash includes dust recovered from the nozzle, the probe liner, and the front half of the filter holder and any material on the filter seal. The filters are dried at 104 °C and the acetone washes are evaporated at room temperature prior to final weighing. Along with the acetone wash, a blank is weighed. The specific procedures followed are listed on the lab sheet in Appendix B-1.

IV. GASEOUS EMISSION RESULTS

The complete report received from Air Consulting and Engineering is included as this section.

SOURCE TEST REPORT

for

OXIDES OF NITROGEN
AND
SULFUR DIOXIDE

at

INTERNATIONAL MINERALS AND CHEMICAL CORPORATION
NUMBER 1 ROCK DRYER
FT. LONESOME MINE
BRADLEY, FLORIDA

FDER PERMIT NO. A029-111119
PSD-FL-088

NOVEMBER 5, 1987

Prepared for:

INTERNATIONNAL MINERALS AND CHEMICAL CORPORATION
POST OFFICE BOX 867
BARTOW, FLORIDA 33830

Prepared by:

AIR CONSULTING AND ENGINEERING
2106 N.W. 67th PLACE, SUITE 4
GAINESVILLE, FLORIDA 32606
(904) 335-1889

191-87-06

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2.0	SUMMARY AND DISCUSSION OF RESULTS.....	2
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APPENDICES

APPENDIX A--SO₂. H₂O EMISSION DATA

APPENDIX B--FIELD DATA SHEETS

SO₂. H₂O

SO₂ LABORATORY ANALYSIS

APPENDIX C--C.E.M. REFERENCE METHODS

APPENDIX D--NO_x STRIP CHART RECORDS

APPENDIX E--C.E.M. DATA REDUCTIONS

APPENDIX F--QUALITY ASSURANCE DATA

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To the best of my knowledge, all applicable field and analytical procedures comply with Florida Department of Environmental Regulation requirements and all test data and plant operating data are true and correct.

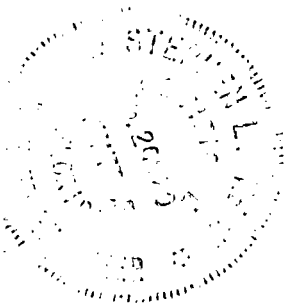
Stephen L. Neck

Stephen L. Neck, P.E.

State of Florida
Registration No. 20020

December 10, 1987

Date



SEAL

1.0 INTRODUCTION

Personnel from Air Consulting and Engineering (ACE) performed Environmental Protection Agency (EPA) reference Method 7E continuous emission monitoring (CEM) for oxides of nitrogen (NO_x), as well as EPA Method 6 SO_2 emission measurements at International Minerals and Chemical Corporation's (IMC) Lonesome Mine Number 1 rock dryer. This work was performed on November 5, 1987, under contract to IMC.

Mssrs. Carlos Gonzalez and Mike Silcott of the Hillsborough County Environmental Protection Commission observed testing.

Mr. Dave Turlee of IMC served as test coordinator as well as test director for EPA Methods 1-5 and 9 as performed by IMC field crews.

2.0 SUMMARY AND DISCUSSION OF RESULTS

Results of the testing are summarized in Table 1. Emission values are reported in terms of concentration.

Throughout the day, a problem with zero drift of the NO_x analyzer was experienced. Although all drifts were within Method 7E specifications (either 2% or 5% depending on interpretation), additional recalibrations were conducted at the analyzer to ensure that the upward drift did not prevent the source from demonstrating compliance. Following each recalibration at the analyzer, a sample bias test was performed at the sample interface.

Complete documentation of all strip chart records are provided in the report appendices.

Because the 50.75 ppm NO/N₂ and 86.2 ppm NO/N₂ calibration gases were a little over 6 months old, Method 7 sampling and analysis were performed for recertification purposes. The results (with rechecks) yielded values within 5% of the tag value. Tag values were therefore taken to remain accurate (Appendix F--Calibration Gas Certification) as specified in Method 7E.

Table 1 Emission Summary
 IMC Lonesome Mine
 Rock Dryer No. 1
 November 5, 1987

Time	Run Number	NO _x		SO ₂ *	
		ppmv _d	lb/SCF (10 ⁻⁶)	ppmvd	lb/SCF (10 ⁻⁶)
1048-1246	1	77	9.19	27.9	4.63
1304-1404	2	82	9.79	36.9	6.13
1419-1519	3	82	9.79	25.0	4.15
-----	AVERAGE	80	9.59	29.9	4.97

NOTE: $(\text{ppmv}_d)(2.595 \times 10^{-6})(M) = \text{lb/SCF}$

Where M = Molecular Weight

= 46 NO₂ (NO_x)

= 64 SO₂

* EACH RUN IS AVERAGE OF TWO 20 MINUTE SAMPLES

3.0 FIELD AND ANALYTICAL PROCEDURES

3.1 Sulfur Dioxide (SO₂) Sampling and Analysis--EPA Method 6

Sulfur Dioxide (SO₂) samples were collected by the emission measurement method specified by the United States Environmental Protection Agency (EPA). Hydrogen peroxide (H₂O₂) was used as the collection media for SO₂. A schematic diagram of the sampling train used is shown in Figure 1.

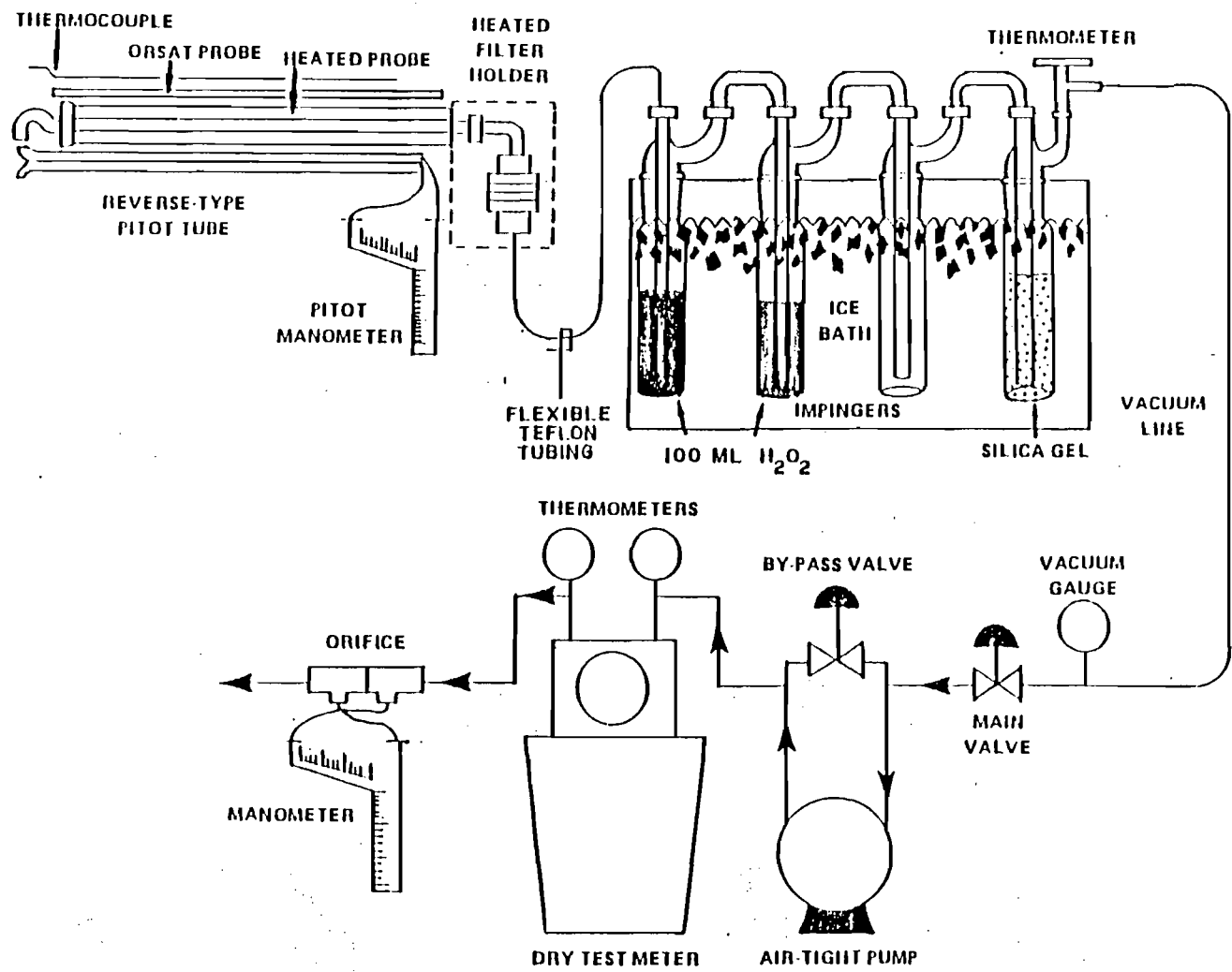
PREPARATION OF EQUIPMENT

1. FILTERS - Gelman type "A" filters.
2. FILTER HOLDER, FLEXIBLE TEFLON[®] TUBING, AND SAMPLING PROBE - The filter holder, flexible Teflon[®] tubing, and sampling probe were washed vigorously with soapy water and brushes, rinsed with acetone and distilled water, and dried prior to the test program. All openings on the sampling equipment were sealed while in transit to the test site.
3. IMPINGERS - The Greenburg-Smith impingers were cleaned with a warm soapy water solution and brushes, rinsed with distilled water and acetone, and dried. The impingers were sealed tightly during transit.

TEST PROCEDURE

Prior to performing the actual sample runs, certain stack and stack gas parameters were measured. These preliminary measurements included the average gas temperature, the stack gas velocity head, the stack gas moisture content, and the stack dimensions at the point where the tests were being performed. The stack gas temperature was determined by using a bi-metallic thermocouple and calibrated pyrometer.

A single sampling point was selected so that a representative sample could be extracted from the gas stream. This point was located in the stack centroid.



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FIGURE 1
EPA METHOD 5/6 SAMPLING TRAIN

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and
ENGINEERING

The gases sampled passed through the following components: a pyrex glass probe; a glass fiber filter; flexible Teflon[®] tubing; two impingers each with 100 ml of three percent hydrogen peroxide; one impinger dry; one impinger with 200 grams of silica gel; a flexible sample line; an air-tight pump; a dry test meter; and a calibrated orifice. The second impinger had a standard tip, while the first, third, and fourth impingers had modified tips with a 0.5 inch I.D. opening. Sample recovery was accomplished by the following procedures:

1. The volume of condensate from the first three impingers was measured for calculating stack gas moisture and then placed in Container 3. All sample exposed surfaces from the rear half of the filter holder through the dry trap impinger were thoroughly washed with distilled-deionized water and the washings were also placed in a glass container. The container was sealed and the liquid level marked. Although polyethylene sample bottles are specified, sulfuric acid reaction with glass is extremely negligible at these concentrations.
2. The used silica gel from the fourth impinger was transferred to its original container and sealed.

LABORATORY ANALYSIS

The four sample containers from each sample run were analyzed according to the following procedures:

1. A sample aliquot was taken and added to isopropanol in a 20/80 ratio. Thorin indicator was added and the solution was titrated to a pink endpoint using nominal 0.0100 normal barium perchlorate. Replicate titrations were made until results agreed within 1% or 0.2 ml. Blanks were titrated in the same fashion.
2. The used silica gel in its tared container was weighed to the nearest gram.

DATA

The field and laboratory data sheets, calculation sheets, and nomenclature definitions are included in the appendix of this report.

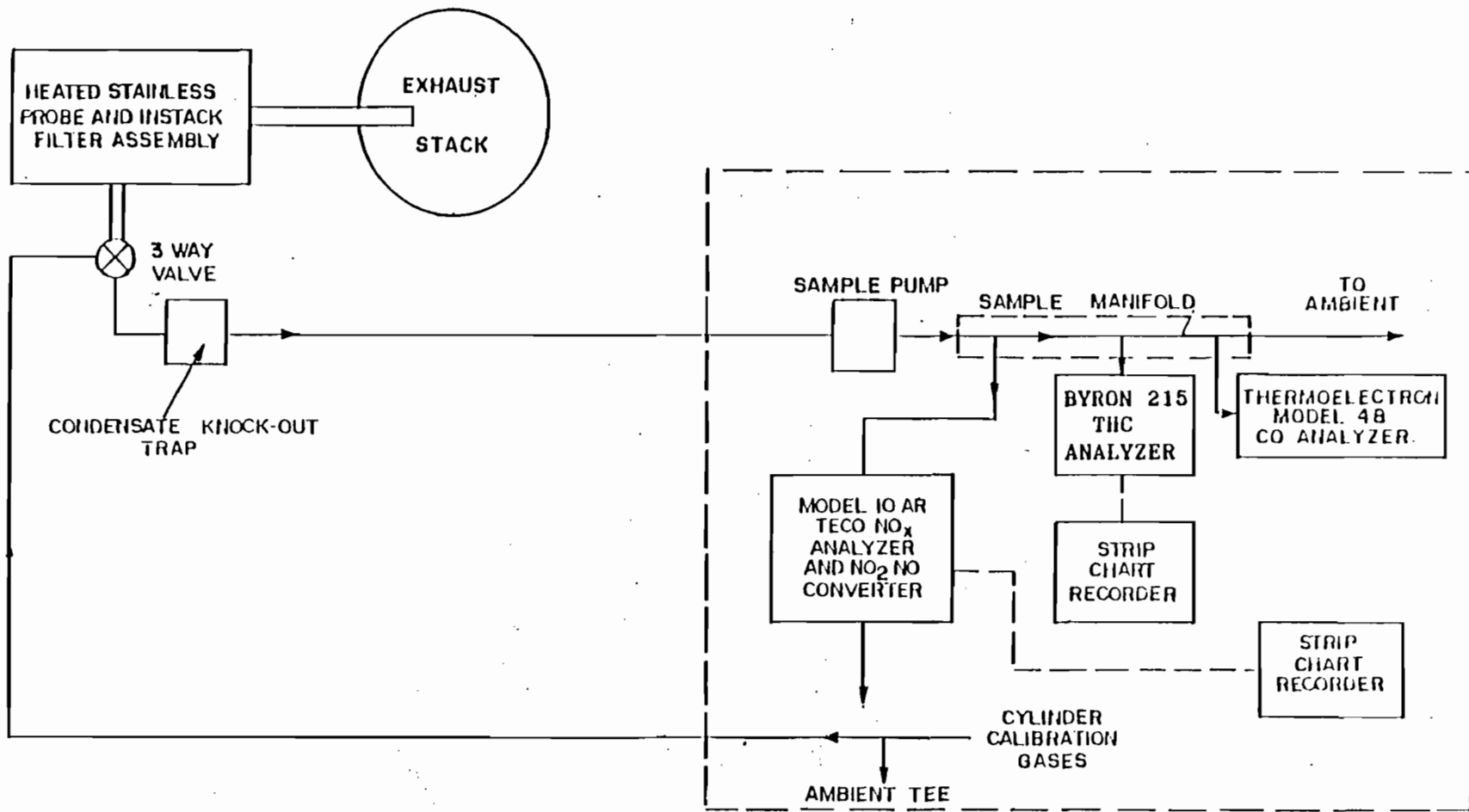
3.2 Continuous Emission Monitoring Methods

Sampling System

The sampling system was set up as shown in Figure 2. A description of all analyzers is provided in Appendix C. Teflon[®] sample lines were used throughout the system. In addition to the CEMs shown, a Teledyne 320 O₂ analyzer was also used with O₂ values ranging from 14.0 to 15.5% during testing.

Test Procedures

EPA reference Methods 7E, 10, and 25A were employed in the testing. These reference methods are provided in Appendix D. All procedures were strictly adhered to.



8

FIGURE 2

EPA METHODS 7E, 10, 25A
SAMPLING TRAIN

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ENGINEERING

APPENDIX A
SO₂, H₂O EMISSION DATA

AIR CONSULTING AND ENGINEERING

Complete Emission Results

 Plant: IMC
 Location: LONESOME MINE Date: 11/5/87
 Stack: UNIT 1 Run 1 From 1023 - 1045

Y Factor	0.999	Nozzle Diameter	0.250 In
Total Time	22.00 Min	Nozzle Area	0.000341 Sq Ft
Stack Area	1.000 Sq Ft	Barometric Pressure	29.85 ''Hg
Stack Temperature	152 Deg F	Meter Temperature	78 Deg F
Stack Pressure	29.85 ''Hg	Meter Orifice Diff	1.100 ''H2O
Stack SQR Vel Head	0.702 ''H2O	Meter Volume	11.982 cf
		Condensate Vol	92.00 ml

1. Volume Water Vapor Sampled	4.342	SCF
2. Volume Standard Dry Gas Sampled	11.757	SCF
3. Total Standard Sample Volume	16.100	SCF
4. Percent Water	26.525	*
5. Percent Dry Air	73.475	
6. Molecular Weight of Dry Stack Gas	29.184	
7. Molecular Weight of Wet Stack Gas	26.217	
8. Specific Gravity Stack Gas	0.91	
9. Percent Oxygen [0.2]	15.20	
10. Percent Carbon Dioxide [CO.2]	3.60	
11. Percent Excess Air	243.715	
12. Velocity of Stack	44.557	FPS
13. Actual Volumetric Flow	2673	ACFM
14. Dry Volumetric Flow	1964	ACFMD
15. Standard Volumetric Flow	1691	SCFMD
16. Emission Rate	0.0000	gr/SCF
17. Emission Rate	0.0000	gr/ACF
18. Emission Rate	0.00	Lbs/Hr
19. Percent Isokinetic	92.7	
20. SO2 Concentration	0.00000533	Lbs/SCF
21. SO2 Emission Rate	0.54	Lbs/Hr
22. SO2 Parts per Million [Dry Basis]	32.09	
23. SO2 Parts per Million [Wet Basis]	23.58	

 Probe/Nozzle Wash 0 Mg
 Filter 0 Mg
 Total 0.000 Mg

* Moisture Calculated from Saturation Vapor Pressure

AIR CONSULTING AND ENGINEERING

Complete Emission Results

Plant: IMC
 Location: LONESOME MINE Date: 11/5/87
 Stack: UNIT 1 Run 2 From 1138 - 1158

Y Factor	0.999	Nozzle Diameter	0.250 In
Total Time	20.00 Min	Nozzle Area	0.000341 Sq Ft
Stack Area	1.000 Sq Ft	Barometric Pressure	29.85 ''Hg
Stack Temperature	158 Deg F	Meter Temperature	84 Deg F
Stack Pressure	29.85 ''Hg	Meter Orifice Diff	1.100 ''H2O
Stack SQR Vel Head	0.548 ''H2O	Meter Volume	10.963 cf
		Condensate Vol	107.40 ml

1. Volume Water Vapor Sampled	5.069	SCF
2. Volume Standard Dry Gas Sampled	10.639	SCF
3. Total Standard Sample Volume	15.708	SCF
4. Percent Water	30.706	*
5. Percent Dry Air	69.294	
6. Molecular Weight of Dry Stack Gas	29.184	
7. Molecular Weight of Wet Stack Gas	25.750	
8. Specific Gravity Stack Gas	0.89	
9. Percent Oxygen [0.2]	15.20	
10. Percent Carbon Dioxide [CO.2]	3.60	
11. Percent Excess Air	243.715	
12. Velocity of Stack	35.262	FPS
13. Actual Volumetric Flow	2116	ACFM
14. Dry Volumetric Flow	1466	ACFMD
15. Standard Volumetric Flow	1250	SCFMD
16. Emission Rate	0.0000	gr/SCF
17. Emission Rate	0.0000	gr/ACF
18. Emission Rate	0.00	Lbs/Hr
19. Percent Isokinetic	124.8	
20. SO2 Concentration	0.00000394	Lbs/SCF
21. SO2 Emission Rate	0.30	Lbs/Hr
22. SO2 Parts per Million [Dry Basis]	23.75	
23. SO2 Parts per Million [Wet Basis]	16.45	

Probe/Nozzle Wash	0	Mg
Filter	0	Mg
Total	0.000	Mg

* Moisture Calculated from Saturation Vapor Pressure

AIR CONSULTING AND ENGINEERING

Complete Emission Results

Plant:	IMC	Date:	11/5/87
Location:	LONESOME MINE	Run 3 From	1213 - 1233
Stack:	UNIT 1		

Y Factor	0.999	Nozzle Diameter	0.250 In
Total Time	20.00 Min	Nozzle Area	0.000341 Sq Ft
Stack Area	1.000 Sq Ft	Barometric Pressure	29.85 ''Hg
Stack Temperature	160 Deg F	Meter Temperature	89 Deg F
Stack Pressure	29.85 ''Hg	Meter Orifice Diff	1.100 ''H2O
Stack SOR Vel Head	0.541 ''H2O	Meter Volume	11.072 cf
		Condensate Vol	88.30 ml

1. Volume Water Vapor Sampled	4.168	SCF
2. Volume Standard Dry Gas Sampled	10.642	SCF
3. Total Standard Sample Volume	14.809	SCF
4. Percent Water	28.143	
5. Percent Dry Air	71.857	
6. Molecular Weight of Dry Stack Gas	29.184	
7. Molecular Weight of Wet Stack Gas	26.037	
8. Specific Gravity Stack Gas	0.90	
9. Percent Oxygen [0.2]	15.20	
10. Percent Carbon Dioxide [CO.2]	3.60	
11. Percent Excess Air	243.715	
12. Velocity of Stack	34.687	FPS
13. Actual Volumetric Flow	2081	ACFM
14. Dry Volumetric Flow	1496	ACFMD
15. Standard Volumetric Flow	1270	SCFMD
16. Emission Rate	0.0000	gr/SCF
17. Emission Rate	0.0000	gr/ACF
18. Emission Rate	0.00	Lbs/Hr
19. Percent Isokinetic	122.9	
20. SO2 Concentration	0.00000589	Lbs/SCF
21. SO2 Emission Rate	0.45	Lbs/Hr
22. SO2 Parts per Million [Dry Basis]	35.49	
23. SO2 Parts per Million [Wet Basis]	25.50	

Probe/Nozzle Wash	0	Mg
Filter	0	Mg
Total	0.000	Mg

AIR CONSULTING AND ENGINEERING

Complete Emission Results

 Plant: IMC
 Location: LONESOME MINE Date: 11/5/87
 Stack: UNIT 1 Run 4 From 1248 - 1308

Y Factor	0.999	Nozzle Diameter	0.250 In
Total Time	20.00 Min	Nozzle Area	0.000341 Sq Ft
Stack Area	1.000 Sq Ft	Barometric Pressure	29.85 ''Hg
Stack Temperature	153 Deg F	Meter Temperature	93 Deg F
Stack Pressure	29.85 ''Hg	Meter Orifice Diff	1.100 ''H2O
Stack SQR Vel Head	0.539 ''H2O	Meter Volume	11.302 cf
		Condensate Vol	88.10 ml

1. Volume Water Vapor Sampled	4.158	SCF
2. Volume Standard Dry Gas Sampled	10.789	SCF
3. Total Standard Sample Volume	14.947	SCF
4. Percent Water	27.522	*
5. Percent Dry Air	72.478	
6. Molecular Weight of Dry Stack Gas	29.184	
7. Molecular Weight of Wet Stack Gas	26.106	
8. Specific Gravity Stack Gas	0.91	
9. Percent Oxygen [0.2]	15.20	
10. Percent Carbon Dioxide [CO.2]	3.60	
11. Percent Excess Air	243.715	
12. Velocity of Stack	34.306	FPS
13. Actual Volumetric Flow	2058	ACFM
14. Dry Volumetric Flow	1492	ACFMD
15. Standard Volumetric Flow	1281	SCFMD
16. Emission Rate	0.0000	gr/SCF
17. Emission Rate	0.0000	gr/ACF
18. Emission Rate	0.00	Lbs/Hr
19. Percent Isokinetic	123.5	
20. SO2 Concentration	0.00000636	Lbs/SCF
21. SO2 Emission Rate	0.49	Lbs/Hr
22. SO2 Parts per Million [Dry Basis]	38.32	
23. SO2 Parts per Million [Wet Basis]	27.77	

 Probe/Nozzle Wash 0 Mg
 Filter 0 Mg
 Total 0.000 Mg

AIR CONSULTING AND ENGINEERING

Complete Emission Results

Plant:	IMC	Date:	11/5/87
Location:	LONESOME MINE	Run 5	From 1405 - 1425
Stack:	UNIT 1		

Y Factor	0.999	Nozzle Diameter	0.250 In
Total Time	20.00 Min	Nozzle Area	0.000341 Sq Ft
Stack Area	1.000 Sq Ft	Barometric Pressure	29.85 ''Hg
Stack Temperature	157 Deg F	Meter Temperature	95 Deg F
Stack Pressure	29.85 ''Hg	Meter Orifice Diff	1.100 ''H2O
Stack SQR Vel Head	0.529 ''H2O	Meter Volume	11.290 cf
		Condensate Vol	89.30 ml

1. Volume Water Vapor Sampled	4.215	SCF
2. Volume Standard Dry Gas Sampled	10.739	SCF
3. Total Standard Sample Volume	14.954	SCF
4. Percent Water	28.187	
5. Percent Dry Air	71.813	
6. Molecular Weight of Dry Stack Gas	29.184	
7. Molecular Weight of Wet Stack Gas	26.032	
8. Specific Gravity Stack Gas	0.90	
9. Percent Oxygen [0.2]	15.20	
10. Percent Carbon Dioxide [CO.2]	3.60	
11. Percent Excess Air	243.715	
12. Velocity of Stack	33.867	FPS
13. Actual Volumetric Flow	2032	ACFM
14. Dry Volumetric Flow	1459	ACFMD
15. Standard Volumetric Flow	1245	SCFMD
16. Emission Rate	0.0000	gr/SCF
17. Emission Rate	0.0000	gr/ACF
18. Emission Rate	0.00	Lbs/Hr
19. Percent Isokinetic	126.5	
20. SO2 Concentration	0.00000398	Lbs/SCF
21. SO2 Emission Rate	0.30	Lbs/Hr
22. SO2 Parts per Million [Dry Basis]	23.95	
23. SO2 Parts per Million [Wet Basis]	17.20	

Probe/Nozzle Wash	0		Mg
Filter	0		Mg
Total	0.000		Mg

AIR CONSULTING AND ENGINEERING

Complete Emission Results

Plant:	IMC	Date:	11/5/87
Location:	LONESOME MINE	Run 6	From 1439 - 1459
Stack:	UNIT 1		

Y Factor	0.999	Nozzle Diameter	0.250 In
Total Time	20.00 Min	Nozzle Area	0.000341 Sq Ft
Stack Area	1.000 Sq Ft	Barometric Pressure	29.85 ''Hg
Stack Temperature	160 Deg F	Meter Temperature	100 Deg F
Stack Pressure	29.85 ''Hg	Meter Orifice Diff	1.100 ''H2O
Stack SQR Vel Head	0.529 ''H2O	Meter Volume	10.936 cf
		Condensate Vol	91.90 ml

- | | | |
|---|------------|---------|
| 1. Volume Water Vapor Sampled | 4.338 | SCF |
| 2. Volume Standard Dry Gas Sampled | 10.309 | SCF |
| 3. Total Standard Sample Volume | 14.647 | SCF |
| 4. Percent Water | 29.615 | |
| 5. Percent Dry Air | 70.385 | |
| 6. Molecular Weight of Dry Stack Gas | 29.184 | |
| 7. Molecular Weight of Wet Stack Gas | 25.872 | |
| 8. Specific Gravity Stack Gas | 0.90 | |
| 9. Percent Oxygen [0.2] | 15.20 | |
| 10. Percent Carbon Dioxide [CO.2] | 3.60 | |
| 11. Percent Excess Air | 243.715 | |
| 12. Velocity of Stack | 34.054 | FPS |
| 13. Actual Volumetric Flow | 2043 | ACFM |
| 14. Dry Volumetric Flow | 1438 | ACFMD |
| 15. Standard Volumetric Flow | 1221 | SCFMD |
| 16. Emission Rate | 0.0000 | gr/SCF |
| 17. Emission Rate | 0.0000 | gr/ACF |
| 18. Emission Rate | 0.00 | Lbs/Hr |
| 19. Percent Isokinetic | 123.8 | |
| 20. SO2 Concentration | 0.00000434 | Lbs/SCF |
| 21. SO2 Emission Rate | 0.32 | Lbs/Hr |
| 22. SO2 Parts per Million [Dry Basis] | 26.14 | |
| 23. SO2 Parts per Million [Wet Basis] | 18.40 | |

Probe/Nozzle Wash	0	Mg
Filter	0	Mg
Total	0.000	Mg

APPENDIX B

FIELD DATA SHEETS
SO₂, H₂O

SO₂ LABORATORY ANALYSIS

STACK SAMPLING FIELD DATA SHEET



2106 N.W. 67th PLACE, SUITE 4
GAINESVILLE, FLORIDA 32606

TEST ID _____

PAGE _____ OF _____

PLANT IMC - LONESOME MINE SOURCE UNIT 1

PLANT LOCATION _____

TYPE OF SAMPLING TRAIN METHOD 6

TYPE OF SAMPLES SO₂

DATE 11-5-87 RUN NO. 1

TIME START 1023 TIME END 1043

SAMPLE TIME _____ min/pt 22 Total min

BAR PRESS. _____ "Hg STACK PRESS. _____ "Hg

ASSUMED MOISTURE 35 % FDA _____

WEATHER OVERCAST TEMP. _____ °F

METER BOX NO. 1 ΔH _____ γ 0.999

NOMOGRAPH C_f _____ PITOT CORR. FACTOR _____

NOZZLE CALIBRATION _____ = _____

STACK DIMENSIONS _____

STACK AREA _____ (EFFECTIVE _____ ft²)

STACK HEIGHT _____ ft.

STACK DIAMETER: UPSTREAM _____ DOWNSTREAM _____

PORT SIZE _____ in. NIPPLE LENGTH _____

U CORD LENGTH: _____

REMARKS: _____

11.76

MAT'L PROCESSING RATE _____

GAS METER READINGS: FINAL 908.682 ft.³

INITIAL 896.700 ft.³

NET 11.982 ft.³

IMPINGERS VOL. GAIN 86 ml.

SILICA GEL NO. 84 WT. GAIN 6.0

FILTER NO. _____ TOTAL CONDENSATE 92.0 ml.

ORSAT

	1	2	3	4	AVG
% CO ₂	<u>15.2</u>				<u>3.6</u>
% O ₂	<u>15.2</u>				<u>15.2</u>
% CO					
% N ₂					

F₀ _____ F₀ RANGE _____

LEAK CHECKS: METER BOX/PUMP _____

ORSAT BAG _____ GAS SAMPLE SYSTEM _____

ORSAT ANALYZER _____

PRE-TEST 0.15 CFM 16" Hg POST-TEST 0.012 4.5" Hg

BOX OPERATOR Neck PROBE HOLDER CANON

PYROMETER NO. 1 PITOT TUBE NO. _____

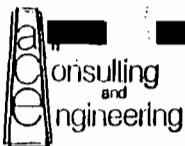
PITOT TUBE LEAK CHECK: PRETEST _____

POST-TEST(+) _____ H₂O _____ SEC

POST-TEST(-) _____ H₂O _____ SEC

PORT AND TRAVERSE POINT NUMBER	DISTANCE FROM INSIDE STACK WALL (IN.)	CLOCK TIME	GAS METER READING (FT ³)	STACK VELOCITY HEAD	METER ORIFICE PRESS. DIFF. ("H ₂ O)		STACK GAS TEMP. (°F)	SAMPLE BOX TEMP. (°F)	LAST IMPINGER TEMP. (°F)	DRY GAS METER TEMP. (°F)	VACUUM ON SAMPLE TRAIN ("Hg)
					CALC.	ACTUAL					
		<u>1028</u>	<u>899</u>	<u>.50</u>	<u>1.1</u>	<u>1.1</u>	<u>149</u>	<u>308</u>	<u>77</u>	<u>76</u>	<u>2.5</u>
		<u>1033</u>	<u>902.0</u>	<u>.49</u>	<u>1.1</u>	<u>1.1</u>	<u>151</u>	<u>308</u>	<u>70</u>	<u>77</u>	<u>2.5</u>
		<u>1038</u>	<u>904.59</u>	<u>.49</u>	<u>1.1</u>	<u>1.1</u>	<u>152</u>	<u>309</u>	<u>63</u>	<u>78</u>	<u>2.5</u>
		<u>1045</u>	<u>908.682</u>	<u>.49</u>	<u>1.1</u>	<u>1.1</u>	<u>155</u>	<u>312</u>	<u>58</u>	<u>80</u>	<u>2.5</u>

STACK SAMPLING FIELD DATA SHEET

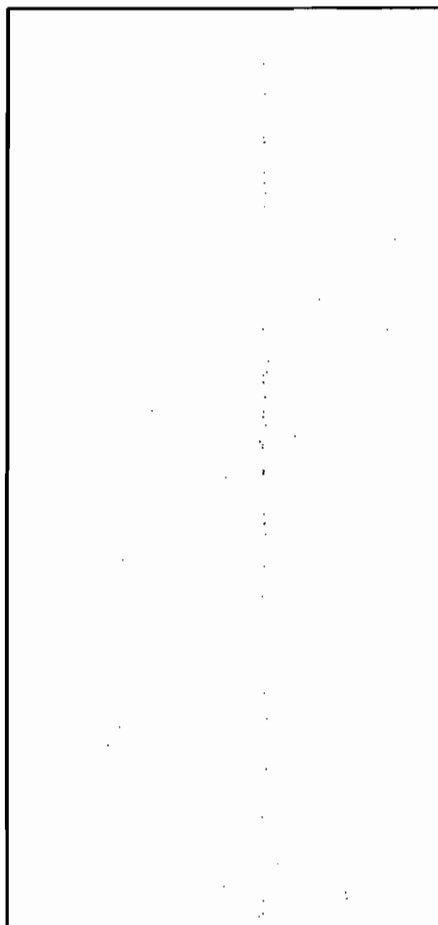


2106 N.W. 67th PLACE, SUITE 4
GAINESVILLE, FLORIDA 32606

TEST ID _____

PAGE _____ OF _____

PLANT Imc SOURCE UNIT 1
 PLANT LOCATION LONESOME MINE
 TYPE OF SAMPLING TRAIN EPA-6
 TYPE OF SAMPLES SO₂
 DATE 11-5-87 RUN NO. 5
 TIME START 1405 TIME END 1425
 SAMPLE TIME 2 min/pt 20 Total min
 BAR PRESS. 29.85 "Hg STACK PRESS. 29.85 "Hg
 ASSUMED MOISTURE _____ % FDA _____
 WEATHER CLEARING TEMP. _____ °F
 METER BOX NO. 1 ΔH _____ Y _____
 NOMOGRAPH C₁ _____ PITOT CORR. FACTOR _____
 NOZZLE CALIBRATION _____ = _____
 STACK DIMENSIONS _____
 STACK AREA _____ (EFFECTIVE _____ ft²)
 STACK HEIGHT _____ ft.
 STACK DIAMETER: UPSTREAM _____ DOWNSTREAM _____
 PORT SIZE _____ in. NIPPLE LENGTH _____
 U CORD LENGTH: _____
 REMARKS: _____



MAT'L PROCESSING RATE _____
 GAS METER READINGS: FINAL 953.710 in.³
 INITIAL 942.500 in.³
 NET 11.210 in.³
 IMPINGERS VOL. GAIN 8.1 ml.
 SILICA GEL NO. 36 WT. GAIN 5.3
 FILTER NO. _____ TOTAL CONDENSATE 87.3 ml

ORSAT	1	2	3	4	AVG
% CO ₂					3.6
% O ₂					15.2
% CO					
% N ₂					

F₀ _____ F₀ RANGE _____
 LEAK CHECKS: METER BOX/PUMP _____
 ORSAT BAG _____ GAS SAMPLE SYSTEM _____
 ORSAT ANALYZER _____
 PRE-TEST 0.005 CFM 10"Hg POST-TEST 0.005 CFM 5"Hg
 BOX OPERATOR Neel PROBE HOLDER CARTER
 PYROMETER NO. _____ PITOT TUBE NO. _____
 PITOT TUBE LEAK CHECK: PRETEST _____
 POST-TEST(+) _____ H₂O _____ SEC
 POST-TEST(-) _____ H₂O _____ SEC

PORT AND TRAVERSE POINT NUMBER	DISTANCE FROM INSIDE STACK WALL (IN.)	CLOCK TIME	GAS METER READING (FT ³)	STACK VELOCITY HEAD	METER ORIFICE PRESS. DIFF. ("H ₂ O)		STACK GAS TEMP (°F)	SAMPLE BOX TEMP (°F)	LAST IMPINGER TEMP (°F)	DRY GAS METER TEMP (°F)	VACUUM ON SAMPLE TRAIN ("Hg)
					CALC.	ACTUAL					
		1410	945.37	.28	1.01	1.1	149	297	51	93	2.5
		1415	948.18	.28	1.1	1.1	156	258	42	94	2.5
		1420	950.07	.28	1.1	1.1	161	252	45	95	2.5
		1425	953.790	.28	1.1	1.1	163	259	48	97	2.5
							164 288				

AIR CONSULTING AND ENGINEERING

SO₂
LAB DATA

Plant Name IMC Date Analyzed 11-24-87

Analyzed By NECK

Stack	Sample No.	V.T.	V.T.B.	N.	V.Soln.	V.A.
UNIT 1	RUN 1	4.0/3.9	0.1	0.0096	480	20
	RUN 2	2.9/2.8			450	20
	RUN 3	3.8/3.8			500	20
	RUN 4	3.9/3.8			540	20
	RUN 5	1.9/1.9			700	20
	RUN 6	1.8/1.7			800	20

- V.T. = Volume of Barium perchlorate titrant used for sample (ml)
- V.T.B. = Volume of Barium perchlorate titrant used for blank (ml)
- N. = Normality of Barium perchlorate
- V.Soln. = Total solution volume
- V.A. = Volume of sample aliquot titrated (ml)

APPENDIX C
C.E.M. REFERENCE METHODS

Method 7E—Determination of Nitrogen Oxides Emissions From Stationary Sources (Instrumental Analyzer Procedure)

1. Applicability and Principle

1.1 **Applicability.** This method is applicable to the determination of nitrogen oxides (NO_x) concentrations in emissions from stationary sources only when specified within the regulations.

1.2 **Principle.** A gas sample is continuously extracted from a stack, and a portion of the sample is conveyed to an instrumental chemiluminescent analyzer for determination of NO_x concentration. Performance specifications and test procedures are provided to ensure reliable data.

2. Range and Sensitivity

Same as Method 6C, Sections 2.1 and 2.2.

3. Definitions

3.1 **Measurement System.** The total equipment required for the determination of NO_x concentration. The measurement system consists of the following major subsystems:

3.1.1 **Sample Interface, Gas Analyzer, and Data Recorder.** Same as Method 6C, Sections 3.1.1, 3.1.2, and 3.1.3.

3.1.2 **NO₂ to NO Converter.** A device that converts the nitrogen dioxide (NO₂) in the sample gas to nitrogen oxide (NO).

3.2 **Span, Calibration Gas, Analyzer Calibration Error, Sampling System Bias, Zero Drift, Calibration Drift, and Response Time.** Same as Method 6C, Sections 3.2 through 3.8.

3.3 **Interference Response.** The output response of the measurement system to a component in the sample gas, other than the gas component being measured.

4. Measurement System Performance Specifications

Same as Method 6C, Sections 4.1 through 4.4.

5. Apparatus and Reagents

5.1 **Measurement System.** Any measurement system for NO_x that meets the specifications of this method. A schematic of an acceptable measurement system is shown in Figure 6C-1 of Method 6C. The essential components of the measurement system are described below:

5.1.1 **Sample Probe, Sample Line, Calibration Valve Assembly, Moisture Removal System, Particulate Filter, Sample Pump, Sample Flow Rate Control, Sample Gas Manifold, and Data Recorder.** Same as Method 6C, Sections 5.1.1 through 5.1.9, and 5.1.11.

5.1.2 **NO₂ to NO Converter.** That portion of the system that converts the nitrogen dioxide (NO₂) in the sample gas to nitrogen oxide (NO). An NO₂ to NO converter is not necessary if data are presented to

demonstrate that the NO₂ portion of the exhaust gas is less than 5 percent of the total NO_x concentration.

5.1.3 **NO_x Analyzer.** An analyzer based on the principles of chemiluminescence, to determine continuously the NO_x concentration in the sample gas stream. The analyzer shall meet the applicable performance specifications of Section 4. A means of controlling the analyzer flow rate and a device for determining proper sample flow rate (e.g., precision rotameter, pressure gauge downstream of all flow controls, etc.) shall be provided at the analyzer.

5.2 **NO_x Calibration Gases.** The calibration gases for the NO_x analyzer shall be NO in N₂. Three calibration gases, as specified in Sections 5.3.1 through 5.3.3, of Method 6C, shall be used. Ambient air may be used for the zero gas.

6. Measurement System Performance Test Procedures

Perform the following procedures before measurement of emissions (Section 7).

6.1 **Calibration Gas Concentration Verification.** Follow Section 6.1 of Method 6C, except if calibration gas analysis is required, use Method 7, and change all 5 percent performance values to 10 percent (or 10 ppm, whichever is greater).

6.2 **Interference Response.** Conduct an interference response test of the analyzer prior to its initial use in the field. Thereafter, recheck the measurement system if changes are made in the instrumentation that could alter the interference response (e.g., changes in the gas detector). Conduct the interference response in accordance with Section 5.4 of Method 20.

6.3 **Measurement System Preparation, Analyzer Calibration Error, and Sample System Bias Check.** Follow Sections 6.2 through 6.4 of Method 6C.

6.4 **NO₂ to NO Conversion Efficiency.** Unless data are presented to demonstrate that the NO₂ concentration within the sample stream is not greater than 5 percent of the NO_x concentration, conduct an NO₂ to NO conversion efficiency test in accordance with Section 5.6 of Method 20.

7. Emission Test Procedure

7.1 **Selection of Sampling Site and Sampling Points.** Select a measurement site and sampling points using the same criteria that are applicable to tests performed using Method 7.

7.2 **Sample Collection.** Position the sampling probe at the first measurement point, and begin sampling at the same rate used during the system calibration drift test. Maintain constant rate sampling (i.e., ± 10 percent) during the entire run. The sampling time per run shall be the same as the total time required to perform a run using Method 7, plus twice the system response time. For each run, use only those measurements obtained after twice the response time of the measurement system has elapsed, to determine the average effluent concentration.

7.3 **Zero and Calibration Drift Test.** Follow Section 7.4 of Method 6C.

8. Emission Calculation

Follow Section 8 of Method 6C.

9. Bibliography

Same as bibliography of Method 6C.

Method 6C—Determination of Sulfur Dioxide Emissions From Stationary Sources (Instrumental Analyzer Procedure)

1. Applicability and Principle.

1.1 Applicability. This method is applicable to the determination of sulfur dioxide (SO₂) concentrations in controlled and uncontrolled emissions from stationary sources only when specified within the regulations.

1.2 Principle. A gas sample is continuously extracted from a stack, and a portion of the sample is conveyed to an instrumental analyzer for determination of SO₂ gas concentration using an ultraviolet (UV), nondispersive infrared (NDIR), or fluorescence analyzer. Performance specifications and test procedures are provided to ensure reliable data.

2. Range and Sensitivity.

2.1 Analytical Range. The analytical range is determined by the instrumental design. For this method, a portion of the analytical range is selected by choosing the span of the monitoring system. The span of the monitoring system shall be selected such that the pollutant gas concentration equivalent to the emission standard is not less than 30 percent of the span. If at any time during a run the measured gas concentration exceeds the span, the run shall be considered invalid.

2.2 Sensitivity. The minimum detectable limit depends on the analytical range, span, and signal-to-noise ratio of the measurement system. For a well designed system, the minimum detectable limit should be less than 2 percent of the span.

3. Definitions.

3.1 Measurement System. The total equipment required for the determination of gas concentration. The measurement system consists of the following major subsystems:

3.1.1 Sample Interface. That portion of a system used for one or more of the following: sample acquisition, sample transport, sample conditioning, or protection of the analyzers from the effects of the stack effluent.

3.1.2 Gas Analyzer. That portion of the system that senses the gas to be measured and generates an output proportional to its concentration.

3.1.3 Data Recorder. A strip chart recorder, analog computer, or digital recorder for recording measurement data from the analyzer output.

3.2 Span. The upper limit of the gas concentration measurement range displayed on the data recorder.

3.3 Calibration Gas. A known concentration of a gas in an appropriate diluent gas.

3.4 Analyzer Calibration Error. The difference between the gas concentration exhibited by the gas analyzer and the known concentration of the calibration gas when the calibration gas is introduced directly to the analyzer.

3.5 Sampling System Bias. The difference between the gas concentrations exhibited by the measurement system when a known concentration gas is introduced at the outlet of the sampling probe and when the same gas is introduced directly to the analyzer.

3.6 Zero Drift. The difference in the measurement system output reading from the initial calibration response at the zero concentration level after a stated period of operation during which no unscheduled maintenance, repair, or adjustment took place.

3.7 Calibration Drift. The difference in the measurement system output reading from the initial calibration response at a mid-range calibration value after a stated period of operation during which no unscheduled maintenance, repair, or adjustment took place.

3.8 Response Time. The amount of time required for the measurement system to display 95 percent of a step change in gas concentration on the data recorder.

3.9 Interference Check. A method for detecting analytical interferences and excessive biases through direct comparison of gas concentrations provided by the measurement system and by a modified Method 6 procedure. For this check, the modified Method 6 samples are acquired at the sample by-pass discharge vent.

3.10 Calibration Curve. A graph or other systematic method of establishing the relationship between the analyzer response and the actual gas concentration introduced to the analyzer.

4. Measurement System Performance Specifications.

4.1 Analyzer Calibration Error. Less than ± 2 percent of the span for the zero, mid-range, and high-range calibration gases.

4.2 Sampling System Bias. Less than ± 5 percent of the span for the zero, and mid- or high-range calibration gases.

4.3 Zero Drift. Less than ± 3 percent of the span over the period of each run.

4.4 Calibration Drift. Less than ± 3 percent of the span over the period of each run.

4.5 Interference Check. Less than ± 7 percent of the modified Method 6 result for each run.

5. Apparatus and Reagents.

5.1 Measurement System. Any measurement system for SO₂ that meets the specifications of this method. A schematic of an acceptable measurement system is shown in Figure 6C-1. The essential components of the measurement system are described below:

5.1.1 Sample Probe. Glass, stainless steel, or equivalent, of sufficient length to traverse the sample points. The sampling probe shall be heated to prevent condensation.

5.1.2 Sample Line. Heated (sufficient to prevent condensation) stainless steel or Teflon tubing, to transport the sample gas to the moisture removal system.

5.1.3 Sample Transport Lines. Stainless steel or Teflon tubing, to transport the sample from the moisture removal system to the sample pump, sample flow rate control, and sample gas manifold.

5.1.4 Calibration Valve Assembly. A three-way valve assembly, or equivalent, for blocking the sample gas flow and introducing calibration gases to the measurement system at the outlet of the sampling probe when in the calibration mode.

5.1.5 Moisture Removal System. A refrigerator-type condenser or similar device (e.g., permeation dryer), to remove condensate continuously from the sample gas while maintaining minimal contact between the condensate and the sample gas. The moisture removal system is not necessary for analyzers that can measure gas concentrations on a wet basis; for these analyzers, (1) heat the sample line and all interface components up to the inlet of the analyzer sufficiently to prevent condensation and (2) determine the moisture content and correct the measured gas concentrations to a dry basis using appropriate methods, subject to the approval of the Administrator. The determination of sample moisture content is not necessary for pollutant analyzers that measure concentrations on a wet basis when (1) a wet basis CO₂ analyzer operated according to Method 3A is used to obtain simultaneous measurements, and (2) the pollutant/CO₂ measurements are used to determine emissions in units of the standard.

5.1.6 Particulate Filter. An in-stack or heated (sufficient to prevent water condensation) out-of-stack filter. The filter glass fiber mat. Additional filters at the inlet or outlet of the moisture removal system and inlet of the analyzer may be used to prevent accumulation of particulate material in the measurement system and extend the useful life of the components. All filters shall be fabricated of materials that are nonreactive to the gas being sampled.

5.1.7 Sample Pump. A leak-free pump, to pull the sample gas through the system at a flow rate sufficient to minimize the response time of the measurement system. The pump may be constructed of any material that is nonreactive to the gas being sampled.

5.1.8 Sample Flow Rate Control. A sample flow rate control valve and rotameter, or equivalent, to maintain a constant sampling rate within 10 percent.

(Note.—The tester may elect to install a back-pressure regulator to maintain the sample gas manifold at a constant pressure in order to protect the analyzer(s) from overpressurization, and to minimize the need for flow rate adjustments.)

5.1.9 Sample Gas Manifold. A sample gas manifold, to divert a portion of the sample gas stream to the analyzer, and the remainder to the by-pass discharge vent. The sample gas manifold should also include provisions for introducing calibration gases directly to the analyzer. The manifold may be constructed of any material that is nonreactive to the gas being sampled.

5.1.10 Gas Analyzer. A UV or NDIR absorption or fluorescence analyzer, to determine continuously the SO₂ concentration in the sample gas stream. The analyzer shall meet the applicable performance specifications of Section 4. A means of controlling the analyzer flow rate and a device for determining proper sample flow rate (e.g., precision rotameter, pressure gauge downstream of all flow controls, etc.) shall be provided at the analyzer.

(Note.—Housing the analyzer(s) in a clean, thermally-stable, vibration-free environment will minimize drift in the analyzer calibration.)

Method 6C—Determination of Sulfur Dioxide Emissions From Stationary Sources (Instrumental Analyzer Procedure)

1. Applicability and Principle.

1.1 Applicability. This method is applicable to the determination of sulfur dioxide (SO₂) concentrations in controlled and uncontrolled emissions from stationary sources only when specified within the regulations.

1.2 Principle. A gas sample is continuously extracted from a stack, and a portion of the sample is conveyed to an instrumental analyzer for determination of SO₂ gas concentration using an ultraviolet (UV), nondispersive infrared (NDIR), or fluorescence analyzer. Performance specifications and test procedures are provided to ensure reliable data.

2. Range and Sensitivity.

2.1 Analytical Range. The analytical range is determined by the instrumental design. For this method, a portion of the analytical range is selected by choosing the span of the monitoring system. The span of the monitoring system shall be selected such that the pollutant gas concentration equivalent to the emission standard is not less than 30 percent of the span. If at any time during a run the measured gas concentration exceeds the span, the run shall be considered invalid.

2.2 Sensitivity. The minimum detectable limit depends on the analytical range, span, and signal-to-noise ratio of the measurement system. For a well designed system, the minimum detectable limit should be less than 2 percent of the span.

3. Definitions.

3.1 Measurement System. The total equipment required for the determination of gas concentration. The measurement system consists of the following major subsystems:

3.1.1 Sample Interface. That portion of a system used for one or more of the following: sample acquisition, sample transport, sample conditioning, or protection of the analyzers from the effects of the stack effluent.

3.1.2 Gas Analyzer. That portion of the system that senses the gas to be measured and generates an output proportional to its concentration.

3.1.3 Data Recorder. A strip chart recorder, analog computer, or digital recorder for recording measurement data from the analyzer output.

3.2 Span. The upper limit of the gas concentration measurement range displayed on the data recorder.

3.3 Calibration Gas. A known concentration of a gas in an appropriate diluent gas.

3.4 Analyzer Calibration Error. The difference between the gas concentration exhibited by the gas analyzer and the known concentration of the calibration gas when the calibration gas is introduced directly to the analyzer.

3.5 Sampling System Bias. The difference between the gas concentrations exhibited by the measurement system when a known concentration gas is introduced at the outlet of the sampling probe and when the same gas is introduced directly to the analyzer.

3.6 Zero Drift. The difference in the measurement system output reading from the initial calibration response at the zero concentration level after a stated period of operation during which no unscheduled maintenance, repair, or adjustment took place.

3.7 Calibration Drift. The difference in the measurement system output reading from the initial calibration response at a mid-range calibration value after a stated period of operation during which no unscheduled maintenance, repair, or adjustment took place.

3.8 Response Time. The amount of time required for the measurement system to display 95 percent of a step change in gas concentration on the data recorder.

3.9 Interference Check. A method for detecting analytical interferences and excessive biases through direct comparison of gas concentrations provided by the measurement system and by a modified Method 6 procedure. For this check, the modified Method 6 samples are acquired at the sample by-pass discharge vent.

3.10 Calibration Curve. A graph or other systematic method of establishing the relationship between the analyzer response and the actual gas concentration introduced to the analyzer.

4. Measurement System Performance Specifications.

4.1 Analyzer Calibration Error. Less than ± 2 percent of the span for the zero, mid-range, and high-range calibration gases.

4.2 Sampling System Bias. Less than ± 5 percent of the span for the zero, and mid- or high-range calibration gases.

4.3 Zero Drift. Less than ± 3 percent of the span over the period of each run.

4.4 Calibration Drift. Less than ± 3 percent of the span over the period of each run.

4.5 Interference Check. Less than ± 7 percent of the modified Method 6 result for each run.

5. Apparatus and Reagents.

5.1 Measurement System. Any measurement system for SO₂ that meets the specifications of this method. A schematic of an acceptable measurement system is shown in Figure 6C-1. The essential components of the measurement system are described below:

5.1.1 Sample Probs. Glass, stainless steel, or equivalent, of sufficient length to traverse the sample points. The sampling probe shall be heated to prevent condensation.

5.1.2 Sample Line. Heated (sufficient to prevent condensation) stainless steel or Teflon tubing, to transport the sample gas to the moisture removal system.

5.1.3 Sample Transport Lines. Stainless steel or Teflon tubing, to transport the sample from the moisture removal system to the sample pump, sample flow rate control, and sample gas manifold.

5.1.4 Calibration Valve Assembly. A three-way valve assembly, or equivalent, for blocking the sample gas flow and introducing calibration gases to the measurement system at the outlet of the sampling probe when in the calibration mode.

5.1.5 Moisture Removal System. A refrigerator-type condenser or similar device (e.g., permeation dryer), to remove condensate continuously from the sample gas while maintaining minimal contact between the condensate and the sample gas. The moisture removal system is not necessary for analyzers that can measure gas concentrations on a wet basis; for these analyzers, (1) heat the sample line and all interface components up to the inlet of the analyzer sufficiently to prevent condensation and (2) determine the moisture content and correct the measured gas concentrations to a dry basis using appropriate methods, subject to the approval of the Administrator. The determination of sample moisture content is not necessary for pollutant analyzers that measure concentrations on a wet basis when (1) a wet basis CO₂ analyzer operated according to Method 3A is used to obtain simultaneous measurements, and (2) the pollutant/CO₂ measurements are used to determine emissions in units of the standard.

5.1.6 Particulate Filter. An in-stack or heated (sufficient to prevent water condensation) out-of-stack filter. The filter glass fiber mat. Additional filters at the inlet or outlet of the moisture removal system and inlet of the analyzer may be used to prevent accumulation of particulate material in the measurement system and extend the useful life of the components. All filters shall be fabricated of materials that are nonreactive to the gas being sampled.

5.1.7 Sample Pump. A leak-free pump, to pull the sample gas through the system at a flow rate sufficient to minimize the response time of the measurement system. The pump may be constructed of any material that is nonreactive to the gas being sampled.

5.1.8 Sample Flow Rate Control. A sample flow rate control valve and rotameter, or equivalent, to maintain a constant sampling rate within 10 percent.

(Note.—The tester may elect to install a back-pressure regulator to maintain the sample gas manifold at a constant pressure in order to protect the analyzer(s) from overpressurization, and to minimize the need for flow rate adjustments.)

5.1.9 Sample Gas Manifold. A sample gas manifold, to divert a portion of the sample gas stream to the analyzer, and the remainder to the by-pass discharge vent. The sample gas manifold should also include provisions for introducing calibration gases directly to the analyzer. The manifold may be constructed of any material that is nonreactive to the gas being sampled.

5.1.10 Gas Analyzer. A UV or NDIR absorption or fluorescence analyzer, to determine continuously the SO₂ concentration in the sample gas stream. The analyzer shall meet the applicable performance specifications of Section 4. A means of controlling the analyzer flow rate and a device for determining proper sample flow rate (e.g., precision rotameter, pressure gauge downstream of all flow controls, etc.) shall be provided at the analyzer.

(Note.—Housing the analyzer(s) in a clean, thermally-stable, vibration-free environment will minimize drift in the analyzer calibration.)

5.1.11 Data Recorder. A strip chart recorder, analog computer, or digital recorder, for recording measurement data. The data recorder resolution (i.e., readability) shall be 0.5 percent of span. Alternatively, a digital or analog meter having a resolution of 0.5 percent of span may be used to obtain the analyzer responses and the readings may be recorded manually. If this alternative is used, the readings shall be obtained at equally spaced intervals over the duration of the sampling run. For sampling run durations of less than 1 hour, measurements at 1-minute intervals or a minimum of 30 measurements, whichever is less restrictive, shall be obtained. For sampling run durations greater than 1 hour, measurements at 2-minute intervals or a minimum of 96 measurements, whichever is less restrictive, shall be obtained.

5.2 Method 6 Apparatus and Reagents. The apparatus and reagents described in Method 6, and shown by the schematic of the sampling train in Figure 6C-2, to conduct the interference check.

5.3 SO₂ Calibration Gases. The calibration gases for the gas analyzer shall be SO₂ in N₂ or SO₂ in air. Alternatively, SO₂/CO₂, SO₂/O₂, or SO₂/CO₂/O₂ gas mixtures in N₂ may be used. For fluorescence-based analyzers, the O₂ and CO₂ concentrations of the calibration gases as introduced to the analyzer shall be within 1 percent (absolute) O₂ and 1 percent (absolute) CO₂ of the O₂ and CO₂ concentrations of the effluent samples as introduced to the analyzer. Alternatively, for fluorescence-based analyzers, use calibration blends of SO₂ in air and the nomographs provided by the vendor to determine the quenching correction factor (the effluent O₂ and CO₂ concentrations must be known). Use three calibration gases as specified below:

5.3.1 High-Range Gas. Concentration equivalent to 80 to 90 percent of the span.

5.3.2 Mid-Range Gas. Concentration equivalent to 50 to 60 percent of the span.

5.3.3 Zero Gas. Concentration of less than 0.25 percent of the span. Purified ambient air may be used for the zero gas by passing air through a charcoal filter, or through one or more impingers containing a solution of 3 percent H₂O₂.

6. Measurement System Performance Test Procedures.

Perform the following procedures before measurement of emissions (Section 7).

6.1 Calibration Gas Concentration Verification. There are two alternatives for establishing the concentrations of calibration gases. Alternative Number 1 is preferred.

6.1.1 Alternative Number 1—Use of calibration gases that are analyzed following the Environmental Protection Agency Traceability Protocol Number 1 (see Citation 1 in the Bibliography). Obtain a certification from the gas manufacturer that Protocol Number 1 was followed.

6.1.2 Alternative Number 2—Use of calibration gases not prepared according to Protocol Number 1. If this alternative is chosen, obtain gas mixtures with a manufacturer's tolerance not to exceed ± 2 percent of the tag value. Within 6 months before the emission test, analyze each of the calibration gases in triplicate using Method 6. Citation 2 in the Bibliography describes procedures and techniques that may be used for this analysis. Record the results on a data sheet (example is shown in Figure 6C-3).

Each of the individual SO₂ analytical results for each calibration gas shall be within 5 percent (or 5 ppm, whichever is greater) of the triplicate set average; otherwise, discard the entire set, and repeat the triplicate analyses. If the average of the triplicate analyses is within 5 percent of the calibration gas manufacturer's cylinder tag value, use the tag value; otherwise, conduct at least three additional analyses until the results of six consecutive runs agree with 5 percent (or 5 ppm, whichever is greater) of their average. Then use this average for the cylinder value.

6.2 Measurement System Preparation. Assemble the measurement system by following the manufacturer's written instructions for preparing and preconditioning the gas analyzer and, as applicable, the other system components. Introduce the calibration gases in any sequence, and make all necessary adjustments to calibrate the analyzer and the data recorder. Adjust system components to achieve correct sampling rates.

6.3 Analyzer Calibration Error. Conduct the analyzer calibration error check by introducing calibration gases to the measurement system at any point upstream of the gas analyzer as follows:

6.3.1 After the measurement system has been prepared for use, introduce the zero, mid-range, and high-range gases to the analyzer. During this check, make no adjustments to the system except those necessary to achieve the correct calibration gas flow rate at the analyzer. Record the analyzer responses to each calibration gas on a form similar to Figure 6C-4.

Note.—A calibration curve established prior to the analyzer calibration error check may be used to convert the analyzer response to the equivalent gas concentration introduced to the analyzer. However, the same correction procedure shall be used for all effluent and calibration measurements obtained during the test.

6.3.2 The analyzer calibration error check shall be considered invalid if the gas concentration displayed by the analyzer exceeds ± 2 percent of the span for any of the calibration gases. If an invalid calibration is exhibited, take corrective action, and repeat the analyzer calibration error check until acceptable performance is achieved.

6.4 Sampling System Bias Check. Perform the sampling system bias check by introducing calibration gases at the calibration valve installed at the outlet of the sampling probe. A zero gas and either the mid-range or high-range gas, whichever most closely approximates the effluent concentrations, shall be used for this check as follows:

6.4.1 Introduce the upscale calibration gas, and record the gas concentration displayed by the analyzer on a form similar to Figure 6C-6. Then introduce zero gas, and record the gas concentration displayed by the analyzer. During the sampling system bias check, operate the system at the normal sampling rate, and make no adjustments to the measurement system other than those necessary to achieve proper calibration gas flow rates at the analyzer. Alternately introduce the zero and upscale gases until a stable response is achieved. The tester shall determine the measurement system response time by observing the times required to achieve a stable response for both the zero

and upscale gases. Note the longer of the two times as the response time.

6.4.2 The sampling system bias check shall be considered invalid if the difference between the gas concentrations displayed by the measurement system for the analyzer calibration error check and for the sampling system bias check exceeds ± 5 percent of the span for either the zero or upscale calibration gas. If an invalid calibration is exhibited, take corrective action, and repeat the sampling system bias check until acceptable performance is achieved. If adjustment to the analyzer is required, first repeat the analyzer calibration error check, then repeat the sampling system bias check.

7. Emission Test Procedure.

7.1 Selection of Sampling Site and Sampling Points. Select a measurement site and sampling points using the same criteria that are applicable to Method 6.

7.2 Interference Check Preparation. For each individual analyzer, conduct an interference check for at least three runs during the initial field test on a particular source category. Retain the results, and report them with each test performed on that source category.

If an interference check is being performed, assemble the modified Method 6 train (flow control valve, two midjet impingers containing 3 percent H₂O₂, and dry gas meter) as shown in Figure 6C-2. Install the sampling train to obtain a sample at the measurement system sample by-pass discharge vent. Record the initial dry gas meter reading.

7.3 Sample Collection. Position the sampling probe at the first measurement point, and begin sampling at the same rate as used during the sampling system bias check. Maintain constant rate sampling (i.e., ± 10 percent) during the entire run. The sampling time per run shall be the same as for Method 6 plus twice the system response time. For each run, use only those measurements obtained after twice response time of the measurement system has elapsed, to determine the average effluent concentration. If an interference check is being performed, open the flow control valve on the modified Method 6 train concurrent with the initiation of the sampling period, and adjust the flow to 1 liter per minute (± 10 percent).

(Note.—If a pump is not used in the modified Method 6 train, caution should be exercised in adjusting the flow rate since overpressurization of the impingers may cause leakage in the impinger train, resulting in positively biased results).

7.4 Zero and Calibration Drift Tests. Immediately preceding and following each run, or if adjustments are necessary for the measurement system during the run, repeat the sampling system bias check procedure described in Section 6.4 (Make no adjustments to the measurement system until after the drift checks are completed.) Record and analyzer's responses on a form similar to Figure 6C-5.

7.4.1 If either the zero or upscale calibration value exceeds the sampling system bias specification, then the run is considered invalid. Repeat both the analyzer calibration error check procedure (Section 6.3) and the sampling system bias check procedure (Section 6.4) before repeating the run.

5.1.11 Data Recorder. A strip chart recorder, analog computer, or digital recorder, for recording measurement data. The data recorder resolution (i.e., readability) shall be 0.5 percent of span. Alternatively, a digital or analog meter having a resolution of 0.5 percent of span may be used to obtain the analyzer responses and the readings may be recorded manually. If this alternative is used, the readings shall be obtained at equally spaced intervals over the duration of the sampling run. For sampling run durations of less than 1 hour, measurements at 1-minute intervals or a minimum of 30 measurements, whichever is less restrictive, shall be obtained. For sampling run durations greater than 1 hour, measurements at 2-minute intervals or a minimum of 96 measurements, whichever is less restrictive, shall be obtained.

5.2 Method 6 Apparatus and Reagents. The apparatus and reagents described in Method 6, and shown by the schematic of the sampling train in Figure 6C-2, to conduct the interference check.

5.3 SO₂ Calibration Gases. The calibration gases for the gas analyzer shall be SO₂ in N₂ or SO₂ in air. Alternatively, SO₂/CO₂, SO₂/O₂, or SO₂/CO₂/O₂ gas mixtures in N₂ may be used. For fluorescence-based analyzers, the O₂ and CO₂ concentrations of the calibration gases as introduced to the analyzer shall be within 1 percent (absolute) O₂ and 1 percent (absolute) CO₂ of the O₂ and CO₂ concentrations of the effluent samples as introduced to the analyzer. Alternatively, for fluorescence-based analyzers, use calibration blends of SO₂ in air and the nomographs provided by the vendor to determine the quenching correction factor (the effluent O₂ and CO₂ concentrations must be known). Use three calibration gases as specified below:

5.3.1 High-Range Gas. Concentration equivalent to 80 to 90 percent of the span.

5.3.2 Mid-Range Gas. Concentration equivalent to 50 to 60 percent of the span.

5.3.3 Zero Gas. Concentration of less than 0.25 percent of the span. Purified ambient air may be used for the zero gas by passing air through a charcoal filter, or through one or more impingers containing a solution of 3 percent H₂O₂.

6. Measurement System Performance Test Procedures.

Perform the following procedures before measurement of emissions (Section 7).

6.1 Calibration Gas Concentration Verification. There are two alternatives for establishing the concentrations of calibration gases. Alternative Number 1 is preferred.

6.1.1 Alternative Number 1—Use of calibration gases that are analyzed following the Environmental Protection Agency Traceability Protocol Number 1 (see Citation 1 in the Bibliography). Obtain a certification from the gas manufacturer that Protocol Number 1 was followed.

6.1.2 Alternative Number 2—Use of calibration gases not prepared according to Protocol Number 1. If this alternative is chosen, obtain gas mixtures with a manufacturer's tolerance not to exceed ± 2 percent of the tag value. Within 6 months before the emission test, analyze each of the calibration gases in triplicate using Method 6. Citation 2 in the Bibliography describes procedures and techniques that may be used for this analysis. Record the results on a data sheet (example is shown in Figure 6C-3).

Each of the individual SO₂ analytical results for each calibration gas shall be within 5 percent (or 5 ppm, whichever is greater) of the triplicate set average; otherwise, discard the entire set, and repeat the triplicate analyses. If the average of the triplicate analyses is within 5 percent of the calibration gas manufacturer's cylinder tag value, use the tag value; otherwise, conduct at least three additional analyses until the results of six consecutive runs agree with 5 percent (or 5 ppm, whichever is greater) of their average. Then use this average for the cylinder value.

6.2 Measurement System Preparation. Assemble the measurement system by following the manufacturer's written instructions for preparing and preconditioning the gas analyzer and, as applicable, the other system components. Introduce the calibration gases in any sequence, and make all necessary adjustments to calibrate the analyzer and the data recorder. Adjust system components to achieve correct sampling rates.

6.3 Analyzer Calibration Error. Conduct the analyzer calibration error check by introducing calibration gases to the measurement system at any point upstream of the gas analyzer as follows:

6.3.1 After the measurement system has been prepared for use, introduce the zero, mid-range, and high-range gases to the analyzer. During this check, make no adjustments to the system except those necessary to achieve the correct calibration gas flow rate at the analyzer. Record the analyzer responses to each calibration gas on a form similar to Figure 6C-4.

Note.—A calibration curve established prior to the analyzer calibration error check may be used to convert the analyzer response to the equivalent gas concentration introduced to the analyzer. However, the same correction procedure shall be used for all effluent and calibration measurements obtained during the test.

6.3.2 The analyzer calibration error check shall be considered invalid if the gas concentration displayed by the analyzer exceeds ± 2 percent of the span for any of the calibration gases. If an invalid calibration is exhibited, take corrective action, and repeat the analyzer calibration error check until acceptable performance is achieved.

6.4 Sampling System Bias Check. Perform the sampling system bias check by introducing calibration gases at the calibration valve installed at the outlet of the sampling probe. A zero gas and either the mid-range or high-range gas, whichever most closely approximates the effluent concentrations, shall be used for this check as follows:

6.4.1 Introduce the upscale calibration gas, and record the gas concentration displayed by the analyzer on a form similar to Figure 6C-6. Then introduce zero gas, and record the gas concentration displayed by the analyzer. During the sampling system bias check, operate the system at the normal sampling rate, and make no adjustments to the measurement system other than those necessary to achieve proper calibration gas flow rates at the analyzer. Alternately introduce the zero and upscale gases until a stable response is achieved. The tester shall determine the measurement system response time by observing the times required to achieve a stable response for both the zero

and upscale gases. Note the longer of the two times as the response time.

6.4.2 The sampling system bias check shall be considered invalid if the difference between the gas concentrations displayed by the measurement system for the analyzer calibration error check and for the sampling system bias check exceeds ± 5 percent of the span for either the zero or upscale calibration gas. If an invalid calibration is exhibited, take corrective action, and repeat the sampling system bias check until acceptable performance is achieved. If adjustment to the analyzer is required, first repeat the analyzer calibration error check, then repeat the sampling system bias check.

7. Emission Test Procedure.

7.1 Selection of Sampling Site and Sampling Points. Select a measurement site and sampling points using the same criteria that are applicable to Method 6.

7.2 Interference Check Preparation. For each individual analyzer, conduct an interference check for at least three runs during the initial field test on a particular source category. Retain the results, and report them with each test performed on that source category.

If an interference check is being performed, assemble the modified Method 6 train (flow control valve, two midjet impingers containing 3 percent H₂O₂, and dry gas meter) as shown in Figure 6C-2. Install the sampling train to obtain a sample at the measurement system sample by-pass discharge vent. Record the initial dry gas meter reading.

7.3 Sample Collection. Position the sampling probe at the first measurement point, and begin sampling at the same rate as used during the sampling system bias check. Maintain constant rate sampling (i.e., ± 10 percent) during the entire run. The sampling time per run shall be the same as for Method 6 plus twice the system response time. For each run, use only those measurements obtained after twice response time of the measurement system has elapsed, to determine the average effluent concentration. If an interference check is being performed, open the flow control valve on the modified Method 6 train concurrent with the initiation of the sampling period, and adjust the flow to 1 liter per minute (± 10 percent).

(Note.—If a pump is not used in the modified Method 6 train, caution should be exercised in adjusting the flow rate since overpressurization of the impingers may cause leakage in the impinger train, resulting in positively biased results).

7.4 Zero and Calibration Drift Tests. Immediately preceding and following each run, or if adjustments are necessary for the measurement system during the run, repeat the sampling system bias check procedure described in Section 6.4 (Make no adjustments to the measurement system until after the drift checks are completed.) Record and analyzer's responses on a form similar to Figure 6C-5.

7.4.1 If either the zero or upscale calibration value exceeds the sampling system bias specification, then the run is considered invalid. Repeat both the analyzer calibration error check procedure (Section 6.3) and the sampling system bias check procedure (Section 6.4) before repeating the run.

7.4.2 If both the zero and upscale calibration values are within the sampling system bias specification, then use the average of the initial and final bias check values to calculate the gas concentration for the run. If the zero or upscale calibration drift value exceeds the drift limits, based on the difference between the sampling system bias check responses immediately before and after the run, repeat both the analyzer calibration error check procedure (Section 6.3) and the sampling system bias check procedure (Section 6.4) before conducting additional runs.

7.5 Interference Check (if performed). After completing the run, record the final dry gas meter reading, meter temperature, and barometric pressure. Recover and analyze the contents of the midget impingers, and determine the SO₂ gas concentration using the procedures of Method 6. (It is not necessary to analyze EPA performance audit samples for Method 6.) Determine the average gas concentration exhibited by the analyzer for the run. If the gas concentrations provided by the analyzer and the modified Method 6 differ by more than 7 percent of the modified Method 6 result, the run is invalidated.

8. Emission Calculation.

The average gas effluent concentration is determined from the average gas concentration displayed by the gas analyzer, and is adjusted for the zero and upscale sampling system bias checks, as determined in accordance with Section 7.4. The average gas concentration displayed by the analyzer may be determined by integration of the area under the curve for chart recorders, or by averaging all of the effluent measurements. Alternatively, the average may be calculated from measurements recorded at equally spaced intervals over the entire duration of the run. For sampling run durations of less than 1 hour, measurements at 1-minute intervals or a minimum of 30 measurements, whichever is less restrictive, shall be used. For sampling run durations greater than 1 hour, measurements at 2-minute intervals or a minimum of 96 measurements, whichever is less restrictive, shall be used. Calculate the effluent gas concentration using Equation 6C-1.

$$C_{\text{gas}} = (\bar{C} - C_0) \frac{C_{\text{MS}}}{C_{\text{MS}} - C_0}$$

Eq. 6C-1

where:

- C_{gas} = Effluent gas concentration, dry basis, ppm.
- \bar{C} = Average gas concentration indicated by gas analyzer, dry basis, ppm.
- C₀ = Average of initial and final system calibration bias check responses for the zero gas, ppm.
- C_{MS} = Average of initial and final system calibration bias check responses for the upscale calibration gas, ppm.
- C_{MS} = Actual concentration of the upscale calibration gas, ppm.

9. Bibliography.

1. Traceability Protocol for Establishing True Concentrations of Gases Used for Calibrations and Audits of Continuous Source Emission Monitors: Protocol Number 1. U.S. Environmental Protection Agency, Quality Assurance Division, Research Triangle Park, NC, June 1978.
2. Westlin, Peter R. and J. W. Brown. Methods for Collecting and Analyzing Gas Cylinder Samples. Source Evaluation Society Newsletter, 3(3):5-15, September 1978.

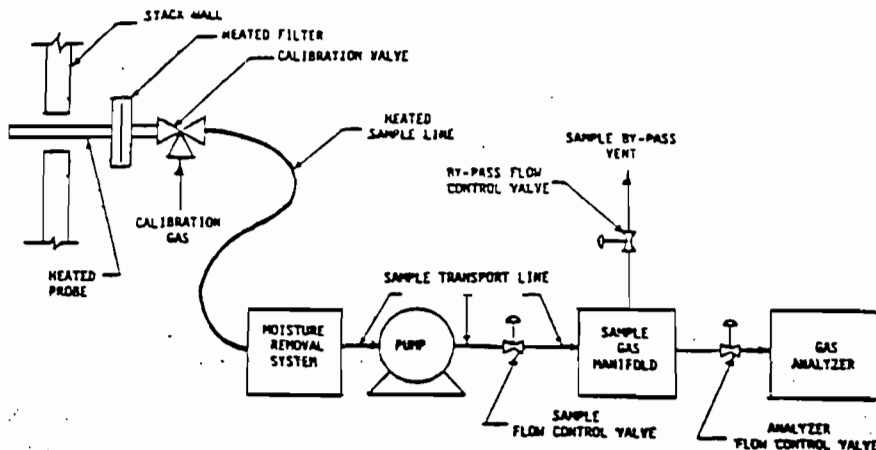


Figure 6C-1. Measurement System Schematic

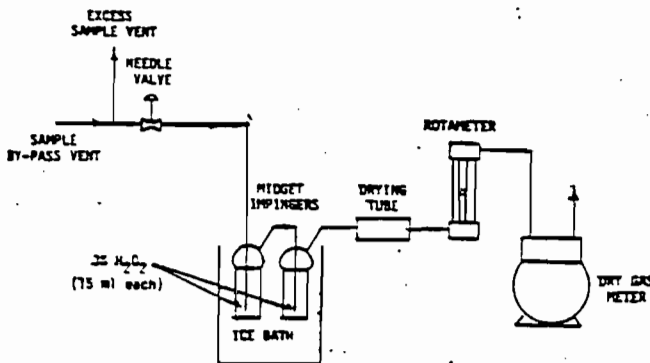


Figure 6C-2. Interference Check Sampling Train

7.4.2 If both the zero and upscale calibration values are within the sampling system bias specification, then use the average of the initial and final bias check values to calculate the gas concentration for the run. If the zero or upscale calibration drift value exceeds the drift limits, based on the difference between the sampling system bias check responses immediately before and after the run, repeat both the analyzer calibration error check procedure (Section 6.3) and the sampling system bias check procedure (Section 6.4) before conducting additional runs.

7.5 Interference Check (if performed). After completing the run, record the final dry gas meter reading, meter temperature, and barometric pressure. Recover and analyze the contents of the midget impingers, and determine the SO₂ gas concentration using the procedures of Method 6. (It is not necessary to analyze EPA performance audit samples for Method 6.) Determine the average gas concentration exhibited by the analyzer for the run. If the gas concentrations provided by the analyzer and the modified Method 6 differ by more than 7 percent of the modified Method 6 result, the run is invalidated.

8. Emission Calculation.

The average gas effluent concentration is determined from the average gas concentration displayed by the gas analyzer, and is adjusted for the zero and upscale sampling system bias checks, as determined in accordance with Section 7.4. The average gas concentration displayed by the analyzer may be determined by integration of the area under the curve for chart recorders, or by averaging all of the effluent measurements. Alternatively, the average may be calculated from measurements recorded at equally spaced intervals over the entire duration of the run. For sampling run durations of less than 1 hour, measurements at 1-minute intervals or a minimum of 30 measurements, whichever is less restrictive, shall be used. For sampling run durations greater than 1 hour, measurements at 2-minute intervals or a minimum of 96 measurements, whichever is less restrictive, shall be used. Calculate the effluent gas concentration using Equation 6C-1.

$$C_{\text{gas}} = (\bar{C} - C_0) \frac{C_{M_6}}{C_m - C_0}$$

Eq. 6C-1

where:

- C_{gas} = Effluent gas concentration, dry basis, ppm.
- \bar{C} = Average gas concentration indicated by gas analyzer, dry basis, ppm.
- C₀ = Average of initial and final system calibration bias check responses for the zero gas, ppm.
- C_m = Average of initial and final system calibration bias check responses for the upscale calibration gas, ppm.
- C_{M₆} = Actual concentration of the upscale calibration gas, ppm.

9. Bibliography.

1. Traceability Protocol for Establishing True Concentrations of Gases Used for Calibrations and Audits of Continuous Source Emission Monitors: Protocol Number 1. U.S. Environmental Protection Agency, Quality Assurance Division, Research Triangle Park, NC, June 1978.
2. Westlin, Peter R. and J. W. Brown. Methods for Collecting and Analyzing Gas Cylinder Samples. Source Evaluation Society Newsletter, 3(3):5-15, September 1978.

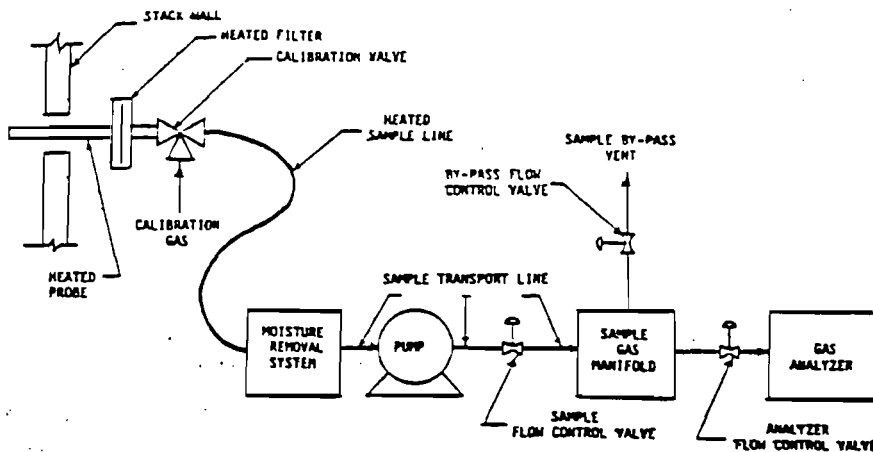


Figure 6C-1. Measurement System Schematic

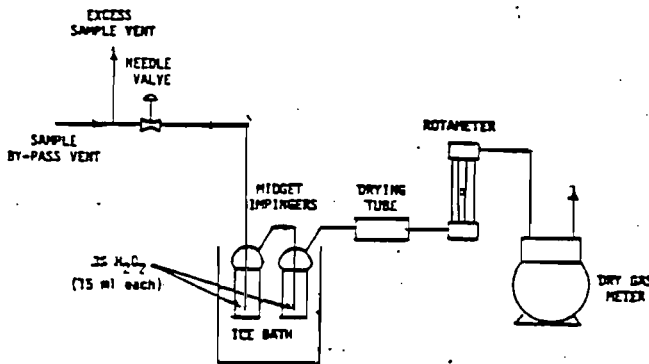


Figure 6C-2. Interference Check Sampling Train

Date _____
 Analytic method used _____

Date: _____
 Analyzer calibration data _____
 for sampling runs: _____
 Span: _____

FIGURE 6C-3.—ANALYSIS OF CALIBRATION GASES

	Gas concentration (indicate units)		
	Zero *	Mid-range *	High-range *
Sample run:			
1.....			
2.....			
3.....			
Average.....			
Maximum percent deviation.....			

- * Average must be less than 0.25 percent of span.
- * Average must be 50 to 60 percent of span.
- * Average must be 80 to 90 percent of span.

Source identification: _____
 Test personnel: _____

FIGURE 6C-4.—ANALYZER CALIBRATION DATA

	Cylinder value	Analyzer calibration response (indicate units)	Absolute difference	Difference (percent of span)
Zero gas.....				
Mid-range gas.....				
High-range gas.....				

Source identification: _____
 Test personnel: _____
 Date: _____
 Run number: _____
 Span: _____

	Analyzer calibration response	Initial values		Final values	
		System calibration response	System cal. bias (percent of span)	(percent of span)	
				System cal. bias	Drift
Zero gas.....					
Upscale gas.....					

$$\text{System Calibration Bias} = \frac{\text{System Cal. Response} - \text{Analyzer Cal. Response}}{\text{Span}} \times 100$$

$$\text{Drift} = \frac{\text{Final System Cal. Response} - \text{Initial System Cal. Response}}{\text{Span}} \times 100$$

Figure 6C-5. System calibration bias and drift data.

Date _____
 Analytic method used _____

Date: _____
 Analyzer calibration data _____
 for sampling runs: _____
 Span: _____

FIGURE 6C-3.—ANALYSIS OF CALIBRATION GASES

	Gas concentration (indicate units)		
	Zero *	Mid-range *	High-range *
Sample run:			
1			
2			
3			
Average			
Maximum percent deviation			

* Average must be less than 0.25 percent of span.
 * Average must be 50 to 60 percent of span.
 * Average must be 80 to 90 percent of span.

Source identification: _____
 Test personnel: _____

FIGURE 6C-4.—ANALYZER CALIBRATION DATA

	Cylinder value	Analyzer calibration response (indicate units)	Absolute difference	Difference (percent of span)
Zero gas				
Mid-range gas				
High-range gas				

Source identification: _____
 Test personnel: _____
 Date: _____
 Run number: _____
 Span: _____

	Analyzer calibration response	Initial values		Final values	
		System calibration response	System cal. bias (percent of span)	System cal. bias (percent of span)	Drift
Zero gas					
Upscale gas					

$$\text{System Calibration Bias} = \frac{\text{System Cal. Response} - \text{Analyzer Cal. Response}}{\text{Span}} \times 100$$

$$\text{Drift} = \frac{\text{Final System Cal. Response} - \text{Initial System Cal. Response}}{\text{Span}} \times 100$$

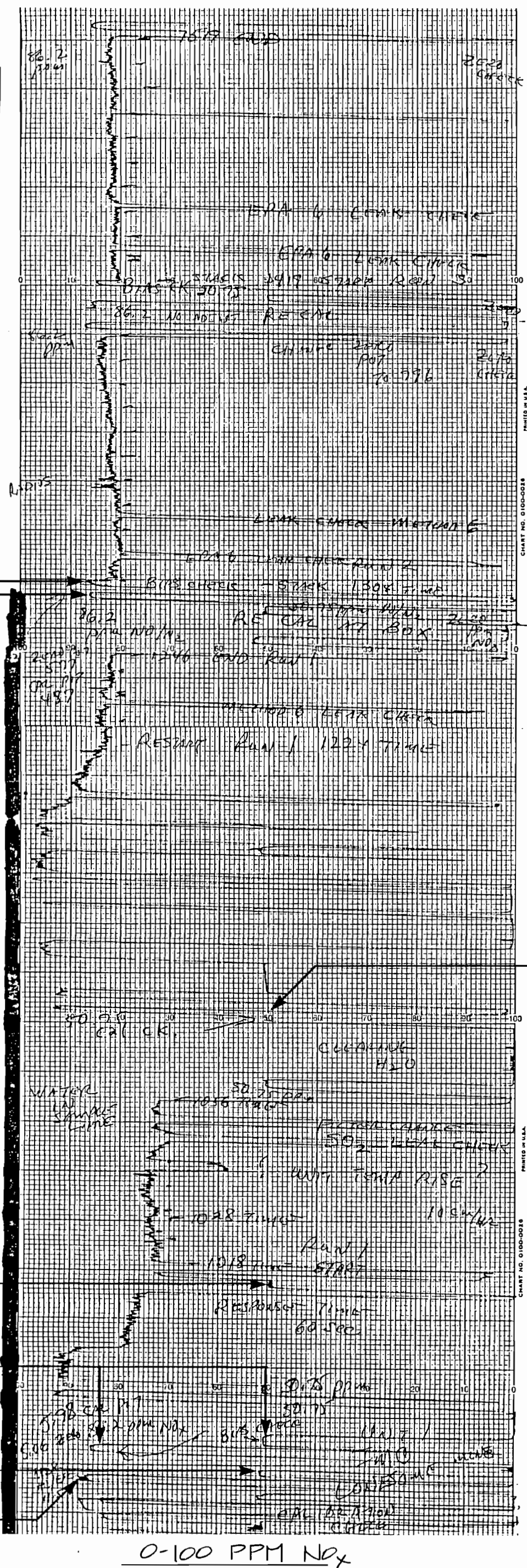
Figure 6C-5. System calibration bias and drift data.

APPENDIX D

NO_x STRIP CHART RECORDS

NOx STRIP CHART
 IMC UNIT - 1
 NOVEMBER 5, 1987

86.2 CAL. CK.
 86.2 BIAS CK.
 86.2 BIAS CK. & LINEARITY CK.
 RESPONSE TIME
 86.2 & 50.75 PPM BIAS
 50.75 PPM LINEARITY
 86.2 PPM CAL. SET



RUN 3

50.75 CAL @ BOX
 86.2 CAL. CK. & RE-SET

RUN 2

RE-CAL @ BOX
 50.75 CAL. CK.
 END RUN 1

↑ CONT. RUN 1

EXCESS FUEL PROBLEM
 50.75 CAL. CK.

STOP TO CLEAR H2O

↑ RUN 1

86.2 PPM CAL. SET

AIR CONSULTING & ENGINEERING



APPENDIX E
C.E.M. DATA REDUCTIONS

EMISSION SUMMARY

FACILITY: Imc - Lonesome mine
 SOURCE: UNIT 1
 DATE: 11-5-87

PAGE 1 OF 2

TIME	MAXIMUM	MINIMUM	AVERAGE
1018 - 1024	75	71	73
1024 - 1030	75	71	74
1030 - 1036	76	72	74
1036 - 1042	74	75	74
1042 - 1048	74	70	72
1048 - 1056	74	72	73
1224 - 1230	85	82	83
1230 - 1236	85	82	82
1236 - 1242	83	81	82
1242 - 1246	82	82	82
1304 - 1310	82	80	81
1310 - 1316	80 81	80	81
1316 - 1322	81	80	81
1322 - 1328	81	80	81
1328 - 1334	83	81	82
1334 - 1340	83	81	82
1340 - 1346	83	81	82
1346 - 1352	84	82	83
1352 - 1358	84	83	83
1358 - 1404	85	83	84

77

82

15 MINUTE SUMMARY

TIME	MAXIMUM	MINIMUM	AVERAGE
1409 - 1425	81	80	81
1425 - 1431	82	80	81
1431 - 1437	81	81	81
1437 - 1443	83	81	82
1443 - 1449	82	81	81
1449 - 1455	82	81	82
1455 - 1500	83	82	82
1500 - 1507	82	81	81
1507 - 1512	82	81	82
1512 - 1519	83	82	82

82 ppm

APPENDIX F
QUALITY ASSURANCE DATA

C.E.M. CALIBRATION ERROR

CONTINUOUS MONITOR ACCURACY CERTIFICATION

PLANT: IMC

LOCATION: Lonesome Mine

SOURCE ID: Unit 1

DATE: November 5. 1987

Calibration Gas	NO _x --Run 1		% Span
	Monitor Value ppm	Difference ppm	
ZERO	0.00	0.00	0.00
50.75 NO/N ₂	51.25	+0.50	+0.50
86.2 NO/N ₂	86.20	0.00	0.00
50.75 BIAS CHECK	50.50	-0.25	-0.25

Calibration Gas	NO _x --Run 2		% Span
	Monitor Value ppm	Difference ppm	
ZERO	0.00	0.00	0.00
50.75 NO/N ₂	50.75	0.00	0.00
86.2 NO/N ₂	86.20	0.00	0.00
86.2 BIAS CHECK	86.20	0.00	0.00

Calibration Gas	NO _x --Run 3		% Span
	Monitor Value ppm	Difference ppm	
ZERO	0.00	0.00	0.00
86.2 NO/N ₂	86.20	86.20	0.00
50.75 NO/N ₂	50.00	0.75	0.75
86.2 BIAS CHECK	85.80	0.40	0.40

C.E.M. ZERO AND SPAN DRIFTS

CONTINUOUS MONITOR DRIFT CERTIFICATION

PLANT: IMC

LOCATION: Lonesome Mine

SOURCE ID: Unit 1

DATE: November 5, 1987

GAS ID: 50.75 NO_x. 86.2 NO_x

Run Number	Time	Span Drift				Zero Drift			
		Initial	Final	Drift	% Span	Initial	Final	Drift	% Span
1	1100	50.75	50.75	0.0	0.0	0.0	1.0	1.0	1.0
		50.75	52.75	2.0	2.0	0.0	2.0	2.0	2.0
2	1406	86.20	87.00	0.8	0.8	0.0	2.0	2.0	2.0
3	1522	86.20	86.00	0.2	0.2	0.0	1.0	1.0	1.0

NO. - NO

CONVERTER CHECK

2400 ppm NO_x (82.6 DIVISIONS)

25% O₂ RANGE 300 ppm NO_x RANGE

10 20 30 40 50 60 70 80 90 100

82.6 DIVISIONS
20% O₂

DRAFT = 1.0%

CONVERTED

UNITS

20 0 = 1.0%

2 25 27

90 80 70 60 50 40 30 20 10 0

90
80
70
60
50
40
30
20
10
0

BEST AVAILABLE COPY

240 ppm NOx (82.7 DIVISIONS)

25% O2 RANGE 300 ppm NOx RANGE

0 10 20 30 40 50 60 70 80 90 100

82.6 DIVISIONS
20% O2

DRAFT = 1.0%

CONVERTED

CHIEF

2.0

2.25-2.7

NOx
ppm
20
10
5
2.5

90 80 70 60 50 40 30 20 10 0

NO_x INTERFERENCE

Interference Response

Procedure: Introduce the gaseous components into the measurement system.
Record the interference response.

Specification: Interference \leq 2 percent of span.

Compliance Test Result of January 18, 1980, Thermo Electron, Inc.

<u>Gas Type</u>	<u>ppm</u>	<u>Analyzer Response</u>	<u>Percent of Span</u>
CO	500	<.1 ppm	<.1%
O2	201	<.1 ppm	<.1%
CO2	10%	<.1 ppm	<.1%
H2	20.9%	<.1 ppm	<.1%

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ANALYZER TYPE

10AR RANGE 0-25 PPM

SERIAL NO.

10AR-014B-80TEST GAS TYPECONCENTRATION PPMANALYZER
OUTPUT RESPONSE% OF SPANCO500< .1 PPM< .1%SO₂201< .1 PPM< .1%CO₂10%< .1 PPM< .1%O₂20.9%< .1 PPM< .1%

**CALIBRATION GAS
CERTIFICATIONS**



Scott Specialty Gases

PLUMSTEADVILLE, PA. 18949

PHONE: (215) 766-8861

TWX: 510-665-9344

AIR CONSULTING AND ENGINEERING
ATTN: STEVE NECK
2106 NW 67TH PLACE
SUITE 4
GAINESVILLE, FL 32606

Date: 3-27-87
Our Project No.: 338458
Your P.O. No.: 87102

Gentlemen:

Thank you for choosing Scott for your Specialty Gas needs. The analyses for the gases ordered, as reported by our laboratory, are listed below. Results are in volume percent, unless otherwise indicated.

ANALYTICAL REPORT

Cyl. No. <u>AAL-1988</u>	Analytical Accuracy <u>±1%</u>
Component	Concentration
<u>CARBON MONOXIDE</u>	<u>14.99 PPM</u>
<u>NITROGEN</u>	<u>BALANCE</u>
<u>NBS TRACEABLE</u>	

Cyl. No. <u>AAL-771</u>	Analytical Accuracy <u>±1%</u>
Component	Concentration
<u>CARBON MONOXIDE</u>	<u>6.114 PPM</u>
<u>NITROGEN</u>	<u>BALANCE</u>
<u>NBS TRACEABLE</u>	

Cyl. No. <u>AAL-2618</u>	Analytical Accuracy <u>±1%</u>
Component	Concentration
<u>NITRIC OXIDE</u>	<u>86.17 PPM</u>
<u>NITROGEN</u>	<u>BALANCE</u>
<u>NBS TRACEABLE</u>	

Cyl. No. <u>AAL-12822</u>	Analytical Accuracy <u>±1%</u>
Component	Concentration
<u>NITRIC OXIDE</u>	<u>50.75 PPM</u>
<u>NITROGEN</u>	<u>BALANCE</u>
<u>NBS TRACEABLE</u>	

Analyst John E. Sanson
JOHN SANSON

Approved By Francis E. Nevill
FRANCIS E. NEVILL

The only liability of this Company for gas which fails to comply with this analysis shall be replacement thereof by the Company without extra cost.

CERTIFIED REFERENCE MATERIALS EPA PROTOCOL GASES
ACUBLEND® CALIBRATION & SPECIALTY GAS MIXTURES PURE GASES
ACCESSORY PRODUCTS CUSTOM ANALYTICAL SERVICES

TROY, MICHIGAN / SAN BERNARDINO, CALIFORNIA / HOUSTON, TEXAS

ACE

3016-B NE 20th Way
Gainesville, Florida 32601

NOx EMISSION TEST DATA

Job No. _____

Plant Name _____

Date 10-31-87

Source Tested 86.17 ppm NO/NOx

Run No. _____

Sample No. _____
 Time _____
 Flask No. _____
 VF--Flask and Stopcock Volume, ml _____
 BI--Initial Barometric Pressure, in Hg _____
 MI--Initial Manometer Reading, in Hg* _____
 PI--Initial Flask Pressure, in Hg Abs. _____
 TI--Initial Flask Temperature, OF _____
 BF--Final Barometric Pressure in Hg _____
 MF--Final Manometer Reading, in Hg* _____
 PF--Final Flask Pressure, in Hg Abs. _____
 TF--Final Flask Temperature, OF _____
 VS--Volume of Gas Sampled at STP, ml _____
 TS--Stack Temperature, OF _____
 PS--Stack Pressure, in Hg _____
 P1--Percent Moisture of Stack Gas, by Vol. _____
 CO2--Percent Carbon Dioxide, Dry basis _____
 O2--Percent Oxygen, Dry Basis _____
 VS--Stack Volumetric Flow Rate at STP, SCFMD _____
 MNO2--Mass of NO2 in Sample, ug _____
 PPM NO2--Parts per Million NO2 at STP, ppm _____
 LBS NO2--Pounds NO2 per Cubic Foot, lbs/dscf (10⁻⁵) _____
 LBS NO2--Pounds NO2 per hour, lbs/hr _____
 F--Factor of Fuel Fired, dscf/MM BTU _____
 LBS NO2--Pounds NO2 per million BTU, lbs/MM BTU _____
 PPM--Parts per Hour 100% Acid _____
 LBS NO2--Pounds NO2 per Ton 100% Acid _____

	1	2	3
Flask No.	16/16	20/20	5/5
VF	2025	2045	1961
BI	30.27	30.27	30.27
MI	3.02	2.97	2.97
PI	27.25	27.3	27.3
TI	76	76	76
BF	30.0	30.0	30.0
MF	-.15	-.25	-.25
PF	29.85	29.75	29.75
TF	80	80	80
VS	175	176	169
TS			
PS			
P1			
CO2			
O2			
VS			
MNO2	284	297	348
PPM NO2	84.8	87.4	107.5
LBS NO2			
LBS NO2/hr			
F			
LBS NO2/MM BTU			
PPM			
LBS NO2/Ton			

FUE CHECK → 297.7
92 88.2

Manometer reading is a negative value when the mercury backs up toward the flask.
 STP--Standard Temperature and Pressure: 68OF, 29.92 in Hg.

MI = PBI + MI
 MF = PBF + MF

$$S = (17.65) \times (VF - 25) \times \left[\left(\frac{PF}{TF + 460} \right) - \left(\frac{PI}{TI + 460} \right) \right]$$

PM NO2 = (523) x (MNO2) ÷ (VS)
 LBS NO2/dscf = (6.23 x 10⁻⁵) x (MNO2) ÷ (VS)
 LBS NO2/hr = (60) x (QS) x (lbs/NO2/dscf)
 LBS NO2/MM BTU = (lbs NO2/dscf) x (F) x $\left(\frac{20.9}{20.9 - \%O_2} \right)$

Tech. Initials _____
 Charged By _____
 Sampled By _____
 Final Pressure & Temp by _____

ACE

3016-B NE 20th Way
Gainesville, Florida 32601

NOx EMISSION TEST DATA

Job No. _____

Plant Name _____

Date 10-31-87

Source Tested 50.75 PPM
NO/NOx

Run No. _____

change to NOx vacuum pump

Sample No.	1	2	3	
Time				
Flask No.	21/21	24/24	3/3	
VF--Flask and Stopcock Volume, ml	2071	1954	1968	
PBI--Initial Barometric Pressure, in Hg	30.27	30.27	30.27	
MI--Initial Manometer Reading, in Hg*	29.40	29.40	29.40	
PI--Initial Flask Pressure, in Hg Abs.	0.87	0.87	0.87	
TI--Initial Flask Temperature, °F	76	76	76	
PBF--Final Barometric Pressure in Hg	30.0	30.0	30.0	
MF--Final Manometer Reading, in Hg*	+1.10	+1.10	+1.20	
PF--Final Flask Pressure, in Hg Abs.	30.10	30.10	30.20	
TF--Final Flask Temperature, °F	80	80	80	
VS--Volume of Gas Sampled at STP, ml	1954	1842	1864	
TS--Stack Temperature, °F				
PS--Stack Pressure, in Hg				
%M--Percent Moisture of Stack Gas, by Vol.				
%CO ₂ --Percent Carbon Dioxide, Dry basis				
%O ₂ --Percent Oxygen, Dry Basis				
QS--Stack Volumetric Flow Rate at STP, SCFMD				
MNO ₂ --Mass of NO ₂ in Sample, ug	183.9	193.8	195.4	
Parts per Million NO ₂ at STP, ppm	49.2	55.0	54.8	53.0
Pounds NO ₂ per Cubic Foot, lbs/dscf (10 ⁻⁵)				
Pounds NO ₂ per hour, lbs/hr		199.5	199.5	
F-Factor of Fuel Fired, dscf/MM BTU		56.6	56.0	
Pounds NO ₂ per million BTU, lbs/MM BTU				
Tons per Hour 100% Acid				
Pounds NO ₂ per Ton 100% Acid				

*R. ECHERK
VALUES*

*Manometer reading is a negative value when the mercury backs up toward the flask.
STP-Standard Temperature and Pressure: 68°F, 29.92 in Hg.

PI = PBI + MI
PF = PBF + MF

$$VS = (17.65) \times (VF - 25) \times \left[\left(\frac{PF}{TF + 460} \right) - \left(\frac{PI}{TI + 460} \right) \right]$$

PPM NO₂ = (523) x (MNO₂) ÷ (VS)
 Lbs NO₂/dscf = (6.23 x 10⁻⁵) x (MNO₂) ÷ (VS)
 Lbs NO₂/hr = (60) x (QS) x (lbs/NO₂/dscf)
 Lbs NO₂/MM BTU = (lbs NO₂/dscf) x (F) x $\left(\frac{20.9}{20.9 - \%O_2} \right)$

Tech. Initials _____
 Charged By _____
 Sampled By _____
 Final Pressure & Temp by _____

ESE ENVIRONMENTAL SCIENCE AND ENGINEERING, INC.

NITROGEN OXIDE LABORATORY DATA

PLANT ACE Calibration gases
 ANALYZED BY Debra Nede

DATE 11/4/82

STACK	RUN/SAMPLE NO.	FLASK NO.	ABSORBANCE (A)	MASS OF NOX (m)
	20		.181	297.3
	16		.173	284.1
	5		.212	348.2
	3		.119	195.4
	21		.112	183.9
	24		.118	193.8
	100		.119	
	200		.243	
	300		.385	
	400		.472	

CALIBRATION FACTOR

$$K_c = \frac{100 (A_1 + 2A_2 + 3A_3 + 4A_4)}{(A_1^2 + A_2^2 + A_3^2 + A_4^2)}$$

$$K_c = \frac{100 (.119 + 2(.243) + 3(.385) + 4(.472))}{.119^2 + .243^2 + .385^2 + .472^2}$$

$$K_c = 821.2$$

MASS OF NO_x

$$m = 2K_c AF$$

A = _____

F = Dilution Factor = _____

11/19/87

Review of Gas Calibration STD

	Abs.	Mean of ND
Flask # 3	.130	199.5
5	.194	297.7
24	.130	199.5
100 μ	.138	
200	.254	
300	.393	
400	.521	corr coef .9997

$R_e = 767.3$

APPENDIX G
PROJECT PARTICIPANTS

PROJECT PARTICIPANTS

ACE

Stephen L. Neck. P.E.

C.E.M. Field Testing
Report Preparation
SO₂ Analysis

Sidney J. Carter

Field Testing

Karie L. Philman

Document Production
Report Preparation

ESE. INC.

Dagmar Neck

NO_x Calibration Gas Analyses

IMC. INC.

Dave Turlee

Project Coordinator

HCEPC

Carlos Gonzalez

Test Observer

Mike Silcott

Test Observer

APPENDIX

A. FIELD SHEETS

A-1 PARTICULATE FIELD SHEETS

Contents: Velocity Traverses.

Particulate Field Sheets (3)

Computer Verification Sheets (2)

Field Check Sheet



SOURCE IDENTIFICATION (PLANT/STACK)																														
1 PIT LOM SOME PREPARATION																														
2 NO. 1 DRYER SCRUBBER																														
3 3 12 1			5 50.27			10 10 5 1.1			2.5			PHOS. ROCK																		
RUN TIME (MILITARY)										NOZZLE DIAMETER (")					PITOT CP COEFFICIENT					ORSAT ANALYSIS										
START					FINISH					CO ₂ READINGS					O ₂ READINGS															
HR		MIN			HR		MIN			RDG 1		RDG 2			RDG 3			RDG 1		RDG 2			RDG 3							
4 12		00			13		05			23.80		83.4			4.5			4.6		4.5			14.4		14.5			14.5		
5 (-27)			29.90			133.5			248			37			4.3															

65	66						
RUN 2	OF 3						
67	68	69	70	71	72		
TEST DATE							
11		588					
73	74	75	76	77	78	79	80
OPTION				SOURCE ID #			
71.00							

OPERATOR: (1) D. NORMAN (2) L. BRADLEY (3) J. JANNER (4) D. CILLEY (5) D. TORLEY (ENG)

METER NO. 1	2	PROBE LENGTH 8'	9'	FILTER NO. 901	901	NOMOGRAPH INFORMATION	
BOB NO. 2	2	PROBE HEATER T. C	C	PROBE WASH NO. 5	5	H ₂ O 3000	TS REF. 155°F
METER ORIF. CH. 1.855		AMBIENT TEMP. 70'±	70'±	SILICA GEL NO. 99C	99C	FACTORY 1.807201	AP REF. 110°F

COMMENTS: 740 ml
-500
240 ml
+8-SILICA
248

S	T	EL. TIME (MIN.)	TRAVERSE POINT	METER VOLUME (FT ³)	ΔP (H ₂ O)	ΔH (H ₂ O)	STACK TEMP. (°F)	METER TEMP. (°F)		IMP. TEMP.	S BOX TEMP.	PROBE TEMP.	PUMP VAC.
								INLET	OUTLET				
6		5	W-1	134.2	25	45	156	96	90	48	240		0.0
6		10	2	136.1	25	45	156	102	92				0.0
6		15	3	138.0	25	45	156	104	92				0.0
6		20	4	139.8	25	45	156	106	94				0.0
6		25	5	141.8	30	54	156	108	94				0.0
6		30	6	143.9	30	54	156	108	96	48	242		0.0
6		35	S-1	145.8	25	45	156	104	92	50	232		0.0
6		40	2	147.7	25	45	156	110	96				0.0
6		45	3	149.5	25	45	156	110	98				0.0
6		50	4	151.4	25	45	154	110	98				0.0
6		55	5	153.5	30	54	152	110	98	52	242		0.0
6		60	6	155.6	30	54	152	110	98				0.0



SOURCE IDENTIFICATION (PLANT/STACK)																															
1 FT. LOMF SOME PREPARATION																															
2 NO. 1 DRYER SCRUBBER																															
PTS. NO.		ID. NO.		MINS. NO.		STACK AREA (FT ²)				M.C.F.				FEED RATE (TPH)				ALLOWED EMISSIONS LB/DAY				FEED DESCRIPTION									
3		12		1		550.27				1010511				25				PHOS ROCK													
RUN TIME (MILITARY)												NOZZLE DIAMETER (")				PITOT Cp COEFFICIENT				ORGAN ANALYSIS											
START						FINISH						CO ₂ READINGS						O ₂ READINGS													
HR		MIN		HR		MIN		RDG 1		RDG 2		RDG 3		RDG 1		RDG 2		RDG 3													
12		00		13		05		238.0		83.4		4.5		4.6		4.5		14.4		14.5		14.5									
STATIC PRESSURE (MM HG)		BAROMETRIC PRESSURE (MM HG)		INITIAL METER VOLUME (FT ³)		LEAK CHECK VOLUME (FT ³)		EL. TIME AT LEAK CHECK		NET H ₂ O COLLECTED (MG)		NET FILTER WEIGHT (MG)		NET PROBEWASH WEIGHT (MG)																	
(-27)		29.90		133.5						248		37		43																	

PAGE 2 OF 3					
TEST DATE					
11		588			
SOURCE ID #					
7100					

OPERATORS: (1) D. NORMAN (2) L. BRADLEY (3) J. TANNER (4) D. GILLEY (5) D. TORLEY (ENG)

METER BODY NO.		PROBE LENGTH		FILTER NO.		NOMOGRAPH INFORMATION			
1		81		901		T _{H2O}		TS REF.	
SNOX NO. SET		PROBE HEATER SET.		PROBE WASH NO.		3000		155°F	
2		C		5		C FACTOR		1.807201	
METER ORIFICE #		AMBIENT TEMP.		SILICA GEL NO.		1.807201		110°F	
1.PSS		70.2		986					

COMMENTS: 740ml
-500
240ml
+8-SMICA
248

S. T.	EL. TIME (MIN.)	TRAVERSE POINT	METER VOLUME (FT ³)	ΔP (H ₂ O)	ΔH (H ₂ O)	STACK TEMP. (°F)	METER TEMP. (°F)		IMP. TEMP.	S BOX TEMP.	PROBE TEMP.	PUMP VAC.
							INLET	OUTLET				
6	5	W-1	134.2	25	45	156	96	90	48	240		0.0
6	10	2	136.7	25	45	156	102	92				0.0
6	15	3	138.0	25	45	156	104	92				0.0
6	20	4	139.8	25	45	156	106	94				0.0
6	25	5	141.8	30	54	156	108	94				0.0
6	30	6	143.9	30	54	156	108	96	48	242		0.0
6	35	5-1	145.8	25	45	156	104	92	50	232		0.0
6	40	2	147.7	25	45	156	110	96				0.0
6	45	3	149.5	25	45	156	110	98				0.0
6	50	4	151.4	25	45	154	110	98				0.0
6	55	5	153.5	30	54	152	110	98	52	242		0.0
6	60	6	155.6	30	54	152	110	98				0.0



SOURCE IDENTIFICATION (PLANT/STACK)																																			
1 FT. LONESOME PREPARATION																																			
2 NO. 1 DRYER SCRUBBER																																			
3			12			1			550.27			1010			511			25			PHAS ROCK														
PUMP TIME (MIN:SEC)																		NOZZLE DIAMETER (")						PITOT Cp COEFFICIENT			ORSAT ANALYSIS								
START						FINISH						CO ₂ READINGS						O ₂ READINGS																	
HR		MIN		HR		MIN		RDG 1		RDG 2		RDG 3		RDG 1		RDG 2		RDG 3																	
4		1405		1510		2380		834		45		46		46		145		145		145															
5			STATIC PRESSURE (PSIG)			BAROMETRIC PRESSURE (IN HG)			INITIAL METER VOLUME (FT ³)			LEAK CHECK VOLUME (FT ³)			EL. TIME AT LEAK CHECK			NET H ₂ O COLLECTED (MM)			NET FILTER WEIGHT (MM)			NET PROBLWASH WEIGHT (MG)											
5			(-30)			29.90			155.9						237			43			28														

65		66													
RUN		3 OF 3													
67		68		69		70		71		72					
TEST DATE:															
11 5 87															
73		74		75		76		77		78		79		80	
OPTION						SOURCE ID #									
7100															

OPERATORS: (1) D. NORMAN (2) P. CULLEY (3) J. TANNER (4) L. BRADLEY (5) D. TURLEY (ENGINEER)

METER BOX NO		PROBE LENGTH		FILTER NO.		MONOGRAPH INFORMATION					
2		8'		902		H ₂ O		3090		TS REF.	
NO 2		6		6		L. FACTOR		1.807201		155°F	
METER OFFICE #		AMBIENT TEMP		SILICA GEL NO.						110°F	
1.855		70.0		987							

COMMENTS: 728 MI
-500 MI
228 MI
+9 SILICA
237 MI

ST	EL. TIME (MIN.)	TRAVERSE POINT	METER VOLUME (FT ³)	ΔP (IN H ₂ O)	ΔH (IN H ₂ O)	STACK TEMP. (°F)	METER TEMP. (°F)		IMP. TEMP.	S. BOX TEMP.	PROBE TEMP.	PUMP VAC.
							INLET	OUTLET				
6	5	1	157.8	25	45	154	96	84	50	240		0.0
6	10	2	159.8	25	45	154	100	88				0.0
6	15	3	161.6	25	45	154	100	90				0.0
6	20	4	163.5	25	45	154	102	90	52	242		0.0
6	25	5	165.5	30	54	154	102	90				0.0
6	30	6	167.6	30	54	154	104	92				0.0
6												
6	35	W 1	169.5	25	45	154	94	88	52	242		0.0
6	40	2	171.3	25	45	154	104	92				0.0
6	45	3	173.2	25	45	154	106	94				0.0
6	50	4	175.0	25	45	154	106	94				0.0
6	55	5	176.9	25	45	154	108	96				0.0
6	60	6	178.7	25	45	154	108	96	50	244		0.0
6												
6												
6												
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6												
6												
6												
6												
6												
6												



JOB: BA680
DATE: 11/13/87

INTERNATIONAL MINERALS & CHEMICAL CORPORATION
SOURCE SAMPLING DATA VERIFICATION REPORT

IMC SOURCE NO. 07100
OPTION NO. 000

TEST OF STACK.....NO 1 DRYER SCRUBBER

AT PLANT.....LONESOME PREPARATION

CONDUCTED ON..... 11/05/87

-----CONSTANT SAMPLING PARAMETERS-----

TRAVERSE POINTS PER RUN.....12	NUMBER OF RUNS.....3	PROCESS FEED DESCRIPTION..... PHOS ROCK
READINGS PER TRAVERSE POINT..... 1	STACK AREA (SQ. FEET).....50.2700	PROCESS FEED RATE (TONS PER HR)... 511.0
MINUTES PER READING..... 5	METER CORRECTION FACTOR..... 1.010	EMISSIONS ALLOWED (LBS PER HR).... 25.00

-----SAMPLE READINGS BY RUN-----

-----I T E M N A M E-----	-----UNITS-----	---RUN 1---	---RUN 2---	---RUN 3---	
TIME OF RUN.....TIME AT START	(24-HR CLOCK)	10:35	12:00	14:05	
	TIME AT FINISH	(24-HR CLOCK)	11:45	13:05	15:10
EQUIPMENT PARAMETERS.....NOZZLE DIAMETER	INCHES	0.2380	0.2380	0.2380	
	PITOT COEFFICIENT	UNITY	0.834	0.834	0.834
ORSAT ANALYSIS READINGS.....%CO2 READING 1	PERCENT	4.5	4.5	4.5	
	READING 2	PERCENT	4.5	4.6	4.6
	READING 3	PERCENT	4.5	4.5	4.6
	%O2 READING 1	PERCENT	14.5	14.4	14.5
	READING 2	PERCENT	14.5	14.5	14.5
	READING 3	PERCENT	14.5	14.5	14.5
PRESSURE MEASUREMENTS.....STATIC STACK PRESSURE	INCHES OF H2O	0.30-	0.27-	0.30-	
	BAROMETRIC PRESSURE	INCHES OF HG	29.90	29.90	29.90
METER MEASUREMENTS.....INITIAL READING	CUBIC FEET	109.10	133.50	155.90	
	LEAK CHECK VOLUME ADJ	CUBIC FEET	0.00	0.00	0.00
	ELAPSED TIME AT LEAK CHK	MINUTES	00.	00.	00.
MOISTURE AND PARTICULATE.....TOTAL LIQUID COLLECTED	MILLILITERS	228.0	248.0	237.0	
	NET FILTER WEIGHT	MILLIGRAMS	8.2	3.7	4.3
	NET PROBEGASH WEIGHT	MILLIGRAMS	4.5	4.3	2.8



JOB: BA680
DATE: 11/13/87

INTERNATIONAL MINERALS & CHEMICAL CORPORATION
SOURCE SAMPLING DATA VERIFICATION REPORT

IMC SOURCE NO. 07100
OPTION NO. 000

TEST OF STACK.....NO 1 DRYER SCRUBBER

AT PLANT.....LONESOME PREPARATION

CONDUCTED ON..... 11/05/87

-----RUN 1-----							
ELAP TIME	TRV PT	METER VOLUME	DEL P	DEL H	STK T	METER T IN	METER T OUT
0		109.10					
5	S-1	111.00	.25	.45	155	84	84
10	S-2	112.70	.25	.45	152	88	84
15	S-3	114.50	.25	.45	152	98	86
20	S-4	116.20	.25	.45	152	100	86
25	S-5	118.80	.45	.81	151	102	88
30	S-6	121.20	.45	.81	151	104	90
35	W-1	123.00	.25	.45	156	100	90
40	W-2	124.80	.25	.45	156	102	92
45	W-3	126.70	.25	.45	156	104	92
50	W-4	128.60	.25	.45	156	104	92
55	W-5	130.50	.25	.45	156	106	92
60	W-6	132.30	.25	.45	156	106	94

-----RUN 2-----							
TRV PT	METER VOLUME	DEL P	DEL H	STK T	METER T IN	METER T OUT	
	133.50						
W-1	134.20	.25	.45	156	96	90	
W-2	136.10	.25	.45	156	102	92	
W-3	138.00	.25	.45	156	104	92	
W-4	139.80	.25	.45	156	106	94	
W-5	141.80	.30	.54	156	108	94	
W-6	143.90	.30	.54	156	108	96	
S-1	145.80	.25	.45	156	104	92	
S-2	147.70	.25	.45	156	110	96	
S-3	149.50	.25	.45	156	110	98	
S-4	151.40	.25	.45	154	110	98	
S-5	153.50	.30	.54	152	110	98	
S-6	155.60	.30	.54	152	110	98	

-----RUN 3-----							
TRV PT	METER VOLUME	DEL P	DEL H	STK T	METER T IN	METER T OUT	
	155.90						
S-1	157.80	.25	.45	154	96	84	
S-2	159.80	.25	.45	154	100	88	
S-3	161.60	.25	.45	154	100	90	
S-4	163.50	.25	.45	154	102	90	
S-5	165.50	.30	.54	154	102	90	
S-6	167.60	.30	.54	154	104	92	
W-1	169.50	.25	.45	154	94	88	
W-2	171.30	.25	.45	154	104	92	
W-3	173.20	.25	.45	154	106	94	
W-4	175.00	.25	.45	154	106	94	
W-5	176.90	.25	.45	154	108	96	
W-6	178.70	.25	.45	154	108	96	

SAMPLING CHECK LIST

DATE 11/5/87 BY JOA

STACK 7100

STAINLESS STEEL PROBE NOZZLES

RUN	MEASUREMENT	DEVIATION	AVERAGE
1	12380	0	12380
2	12380	0	12380
3	12380	0	12380

STAINLESS STEEL PROBE LINERS

RUN	VISUAL CHECK OK	CONNECTED OK	CHANGED DURING RUN
1	✓	✓	NO
2	✓	✓	NO
3	✓	✓	NO

"S" TYPE PITOT

RUN	CALIBRATION DATE	VISUAL INSPECTION	LEAK CHECK*
1	6/7/87	OK	✓
2	6/7/87	OK	✓
3	6/7/87	OK	✓

SAMPLE FILTER CHECK

RUN	VISUAL CHECK OK	CHANGED DURING RUN
1	✓	NO
2	✓	NO
3	✓	NO

* 15 sec @ 3" HOH

SYSTEM LEAK CHECKS:
PRE-TEST

RUN	CF LEAKAGE	INCH HG	SECONDS
1	0.00	15	60
2	0.00	10	60
3	0.00	15	60

POST-TEST

RUN	CF LEAKAGE	INCH HG	SECONDS
1	0.00	3.0	60
2	0.00	3.0	60
3	0.00	3.0	60

SPECIFIC CHECK ITEMS:

- Sample points selected per Method 1?
- Dual inclined manometer leveled and zeroed?
- Sample box thermometer calibration date.
- Nozzle and pitot tube parallel to stack wall during sampling?
- Was stack static pressure measured?
- Required DH found using calculator? or Nomograph?
- Were sample recovery procedures followed?
- Date of last impinger thermometer calibration.
- Stack thermocouple installed from pitot tube end?
- Were 3 grab OCO and OO readings taken?
< 0.3 lb/lb-mole difference in average of three?
- Single point integrated sample for OCO and OO, Tedlar bag leak check for 10 minutes at 2-4 in HOH?
- Condenser: 500 ml distilled HOH?
- Silica gel completely spent?
- Date of last filter thermometer calibration.
- Dates, last dry gas meter thremometer calibrations.
- Date of last stack thermocouple calibration series.

RUN 1	RUN 2	RUN 3
✓	✓	✓
✓	✓	✓
6/6/87	6/6/87	6/6/87
✓	✓	✓
✓	✓	✓
✓	✓	✓
6/6/87	6/6/87	6/6/87
✓	✓	✓
✓	✓	✓
✓	✓	✓
NO	NO	NO
✓	✓	✓
NO	NO	NO
6/6/87	6/6/87	6/6/87
6/6/87	6/6/87	6/6/87
6/6/87	6/6/87	6/6/87

IMC PROCEDURE FOR SAMPLE RECOVERY OF METHOD 5 SAMPLING

SAMPLE RECOVERY:

1. Probe is removed from stack and allowed to cool until it can be handled safely.
2. The external particulate matter is wiped from around the probe nozzle and the nozzle is sealed tightly to prevent loss or gain of particulate matter. The probe is then removed from the sampling train and the probe outlet is cleaned and sealed in the same manner as the probe tip.
3. The probe is then transferred to the air-conditioned IMC sampling trailer for cleaning or transport back to the environmental laboratory for analysis.
4. The probe nozzle and probe connection fitting are removed from the probe, cleaned with a nylon bristle brush and rinsed with acetone from a spray wash bottle into a clean identified glass storage bottle. The probe liner is also cleaned with a nylon bristle brush and rinsed with acetone into the same glass storage bottle. The nozzle, probe connection fitting and probe liner are cleaned until the acetone rinse shows no visible particles. Bottles are filled to equal marked levels and volume is noted on lab analysis sheet.
5. The glass connection from the probe outlet to the filter holder inlet is brushed and rinsed into the identified acetone storage bottle until the acetone rinse shows no visible particles.
6. The filter holder is then disassembled using care not to lose any of the captured particulate. The top half of the glass filter holder is brushed and rinsed into the same identified acetone storage bottle until the acetone rinse shows no visible particles. The acetone storage bottle is sealed tightly to prevent any loss of sample.
7. A blank acetone sample is taken from the same wash bottle used for the sample acetone rinse. (All sample bottles, including the acetone blank, are filled to the same marked level.) The blank sample storage bottle is sealed tightly to prevent any spillage and marked with an identifying number.

FILTER RECOVERY:

1. The filter holder rubber "O" ring is removed from the disassembled filter holder with a pair of tweezers. Any particles that adhere to the "O" ring are transferred into a transparent petri dish using a nylon bristle brush.
2. The filter is then transferred to the petri dish taking care not to spill any of the particles captured on the filter. Check the filter for any visible damage that may have occurred during sampling. The petri dish is sealed tightly to prevent any sample loss and is marked with an identifying number.

SILICA GEL RECOVERY:

1. Silica gel tube is removed from sampling train. Note the condition of the gel color. The gel tube is marked with an identifying number.

IMPINGER RECOVERY:

1. The distilled H₂O condenser impinger is disassembled from the sampling train and the contents are poured into a graduated cylinder. The water volume is on the field data sheet to the nearest ml. (± 1 ml.) Notice any color or film in the liquid.
2. The impinger catch is then discarded unless other analysis is desired or required.

Were there any deviations from the above procedures? None

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VELOCITY TRANSDUCER
 ENVIRONMENTAL SERVICES
 ELECTRICAL ENGINEERS
 & CHEMICAL CORPORATION

PROJECT: #1 RUN

9:20

DATE: 11-5-87

S.W. side

N.W. side

Point	Velocity Pressure	Point	Velocity Pressure
0g	CO2	1	CC-22 132
14.5'	4.5'	2	CC-20 130
14.5'	4.5'	3	CC-10 132
14.5'	4.5'	4	C-18 135
		5	C-19 132
		6	C-24 34
		7	C-20 40
		8	C-12 32
		9	B
		10	↑
		11	9:20
		12	
		13	
		14	
		15	
		16	
		17	
		18	
		19	
		20	
		21	
		22	
		23	
		24	
		25	
		26	

VELOCITY TRANSDUCER: In the STATIC: In HOSE
 ENVIRONMENTAL SERVICES: In the HOSE: In HOSE
 ELECTRICAL ENGINEERS: In the HOSE: In HOSE
 & CHEMICAL CORPORATION: In the HOSE: In HOSE

IMC Velocity Traverse

Stack: #1 Low Same

Date: 11-5-87

point	standard		cyclonic	
	dP std	temp	alpha	dP cyc

Names: BRADLEY - Gilly

S.W ¹	.22	156	CC20	.23
2	.26	156	CC16	.26
3	.25	157	CC14	.26
4	.23	156	0	.23
5	.26	156	C10	.25
6	.30	156	C16	.32
7				
8				
9				
10				

Time Start: 11:35 Finish: 11:47

Barometric Pressure:

Static Pressure: -.3

Stack Temperature Dry: 156 Wet:

% Moisture:

Pitot Tube Cp:

Stack Diameter: Area:

11				
12				

	time	read1	read2	read3	avg
--	------	-------	-------	-------	-----

00 11:50 14.5 14.5 14.5

000 11:58 4.5 4.5 4.5

N.W ¹	.26	156	CC24	.25
2	.25	156	CC18	.26
3	.27	156	CC12	.26
4	.28	156	C6	.27
5	.30	156	C12	.31
6	.28	156	C8	.29
7				

Mdry: Mstk:

	standard	cyclonic
--	----------	----------

alpha: XXXXX

fpm:

acfm:

scfm:

8				
9				
10				
11				
12				

Comments: Run # 1 AFTER

IMC Velocity Traverse					Stack: #1 Longshore	Date: 11-5-87
point	standard		cyclonic		Names: BRADLEY : Billy	
	dP std	temp	alpha	dP cyc		
NW 1	.24	154	CC18	.23	Time Start: 1 : 21	Finish: 1 : 28
2	.23	154	CC14	.22	Barometric Pressure:	
3	.26	154	CC8	.27	Static Pressure: .3	
4	.28	154	C20	.29	Stack Temperature Dry:	Wet: 154
5	.24	154	C14	.22	% Moisture:	
6	.26	154	C10	.24	Pitot Tube Cp:	
7					Stack Diameter: Area:	
8						
9						
10						
11					time	read1 read2 read3 avg
12					00 1 : 32	14.4 14.5 14.5
					000 1 : 45	4.5 4.6 4.5
SW 1	.23	154	CC18	.24	Mdry:	Mstk:
2	.22	154	CC14	.24	standard	cyclonic
3	.24	154	CC10	.25	alpha: XXXXX	
4	.24	154	C18	.24	fpm:	
5	.28	154	C12	.27	acfm:	
6	.25	154	C14	.24	scfm:	
7						
8					Comments: Run #2 AFTER	
9						
10						
11						
12						

IMC Velocity Traverse

Stack: #1 Lanesane

Date: 11-5-87

point	standard dP std	temp	cyclonic alpha	dP cyc
-------	--------------------	------	-------------------	--------

Names: BRADLEY, GILLY

1 NW .21 150 CC18 .22

Time Start: 3:18 Finish: 3:32

2 .24 150 CC15 .25

Barometric Pressure:

3 .22 150 CC6 .24

Static Pressure: -.3

4 .24 150 C20 .24

Stack Temperature Dry: Wet:

5 .25 150 C22 .26

% Moisture:

6 .24 150 C18 .25

Pitot Tube Cp:

7

Stack Diameter: Area:

8

9

10

	time	read1	read2	read3	avg
--	------	-------	-------	-------	-----

11

OO 3:40 14.5 14.5 14.5

12

OCO 3:50 4.5 4.6 4.6

SW 1 .24 150 CC16 .25

Mdry: Mstk:

2 .25 150 CC17 .25

	standard	cyclonic
--	----------	----------

3 .24 150 CC20 .25

alpha: XXXXX

4 .22 150 C-21 .23

fpm:

5 .24 150 C-20 .25

acfm:

6 .26 150 C22 .26

scfm:

7

8

Comments: RUN #3 AFTER

9

10

11

12

A-2 OPERATING LOG
Contents: Measurement Sheet

TESTING PARAMETER LOG for TEST: DATE 11/5/87 DRYER NO 1 BY J. Miller
 MARK STRIP CHARTS in CONTROL ROOM (time and for runs)

TIME	EVENT LOG	TIME	NOx DIVs	RUN	TIME	START RUN	SCF	TIT VOL.
:		:		1 START	:	1		
:		:		1 MID	:	2		
:		:		1 END	:	3		
:		:		2 START	:	4		
:		:		2 MID	:	5		
:		:		2 END	:	6		
:		:		3 START				
:		:		3 MID	TIME	SAMPLE	TIME	SAMPLE
:		:		3 END	11:50	OIL	11:15	U/F
:		:			?	OIL	1:15	U/F
:		TIME	CO DIVs	RUN	3:15	OIL	3:15	U/F
:		:		1 START	:	FEED	:	IN
:		:		1 MID	:	FEED	:	IN
:		:		1 END	:	FEED	:	IN
:		:		2 START	TIME	OO 1	OO 2	OO 3
:		:		2 MID	:			
:		:		2 END	:			
:		:		3 START	:			
:		:		3 MID	:			
:		:		3 END	:			
:		TIME	VOC DIVs	RUN	TIME	OCO 1	OCO 2	OCO 3
:		:		1 START	:			
:		:		1 MID	:			
:		:		1 END	:			
:		:		2 START	:			
:		:		2 MID	:			
:		:		2 END	:			
:		:		3 START	:			
:		:		3 MID	:			
:		:		3 END	:			
:		TIME	BELT	METER	TIME	T-AMB	TIME	T ROCK IN
:		8:00		175°F	8:00	70°C	8:00	26°C
:		10:00		172°F	11:55	70°C	11:55	27°C
:		12:04		166°F	1:45	67°C	1:45	27°C
:		2:00		162°F	3:25	71°C	3:25	26°C
:		3:21		162°F				

B. ANALYSES

B-1 LABORATORY

Contents: Lab Analysis Sheet

FILTER PAN WEIGHTS

PAN NO.	1	2	3
FINAL WTS	3.4369	3.4442	3.4346
TARE WTS	2.6320	2.6371	2.6311
NET WTS	.8049	.8071	.8035
INITIAL WEIGHING	11/9/87		FINAL WEIGHING
HRS DRIED	1 HR.		HRS DRIED 38 CT: 100° C
DATE	11/9/87	WEIGHED BY JDT	DATE 11/12/87 WEIGHED BY JDT
COMMENTS			

FILTER WEIGHTS

FILTER NO	900	901	902
FINAL WT	.8049	.8071	.8035
TARE WT	.7967	.8034	.7992
NET WT	.0082	.0037	.0043
INITIAL WEIGHING	CONTINUOUS		FINAL WEIGHING
HRS DESICCATED	CONTINUOUS		HRS DRIED 38 HRS
DATE	8/26/87	WEIGHED BY JDT	DATE 11/12/87 WEIGHED BY JDT
COMMENTS			

ACETONE WASH WEIGHTS

BOTTLE NO	4	5	6	7
FINAL WT	2.6281	2.6451	2.6407	2.6240
TARE WT	2.6236	2.6429	2.6379	2.6235
NET WT	.0045	.0022	.0028	.0005
SPILLAGE				

WASH VOLUME (mls)

WASH VOLUME (mls)	140 ml
DATE	10/12/87
WEIGHED BY	JDT
COMMENTS	

SILICA GEL WEIGHTS

TUBE NO	985	986	987
FINAL WT	4.08	4.08	4.09
TARE WT	2.00	4.00	4.00
NET WT	8	8	9
INITIAL WEIGHING			FINAL WEIGHING 11/6/87
HRS DRIED	48		DATE 11/6/87
DATE	11/4/87	WEIGHED BY JDT	WEIGHED BY JDT
COMMENTS			

PRE-SAMPLING FILTER & SILICA GEL WEIGHING PROCEDURES

FILTER:

1. The filters are checked visually against light for irregularities, flaws, and pin hole leaks.
2. Filters are oven dried for two to three hours, desiccated for two hours and weighed to the nearest 0.1 mg.
3. Filters are stored in sealed plastic petri containers.

SILICA GEL:

1. Six to sixteen mesh, indicating type silica gel is dried at 175 degrees C for two hours. (New silica gel is used as received.)
2. Four hundred grams of silica gel is weighed to the nearest 0.5 grams.

IMC PROCEDURE FOR FILTER, ACETONE PROBE WASH SILICA GEL ANALYSIS

FILTER ANALYSIS:

1. Sample filter is carefully transferred from the petri storage container to a tared aluminum dish and oven dried at 105 degrees C from two to twenty-four hours. The sample is removed from the oven desiccated ten to fifteen minutes and weighed. This is recorded as the final weight.

ACETONE PROBE WASH ANALYSIS:

1. The lid is removed from the glass acetone was storage bottle and placed under a closed ventilation hood to evaporate the acetone.
2. The acetone wash sample is then carefully transferred to a tared aluminum dish using acetone and the aid of a rubber policeman.
3. The sample is again placed under a ventilation hood to evaporate the acetone.
4. After the acetone has completely evaporated from the sample the aluminum dish is oven dried at 105 degrees C from two to three hours. The sample is removed from the oven desiccated ten to fifteen minutes and weighed. This is recorded as the final weight.
5. The blank acetone sample is analyzed in the same manner as the particulate acetone wash sample.

SILICA GEL ANALYSIS:

1. The used silica gel is weighed to the nearest 0.5 gram and recorded.

Were there any deviations from the above procedures?

(John D. Brown) None

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Date: 11/5/87 Source: FT LONESOME NO 2 DRYER SCRUBBER

Chain of Custody Procedures

- 1) Sample filters are kept in a locked environmental laboratory in a desiccator until their initial weighing.
- 2) Initial filter weights are recorded in sampling log book which includes : a) weight
b) date weighed
c) weighers initials
- 3) Silica gel weighed and placed in drying tubes in the environmental laboratory.
- 4) Filters, in sealed nonreactive containers, and drying tubes are locked in sampling trailer for field use.
- 5) After sampling, filters are sealed in the their holders. The probes and drying tubes are also sealed. All are then locked in the sampling trailer and transported to the environmental laboratory for analysis.
- 6) Filters, silica gel, and probe washes handled according to analysis procedures. These weights are recorded on lab analysis sheets. Precautions taken during analysis procedure are as follows:
 - a) initial weighing date of aluminum dishes
 - b) initial desiccation of aluminum dishes
 - c) no. of hrs sample is dried and at what temperature
 - d) final weighing date of aluminum dishes
 - e) initials of person performing analysis
- 7) Environmental laboratory is routinely patrolled by IMC Security personnel.

Above procedures followed?

Name	Date
John D. Nam	11/18/87
John D. Nam	11/18/87
R. W. Buckley	11/18/87
R. A. Collier	11/18/87
J. E. Lammie	11/18/87

B-2 FUEL ANALYSES

Contents: All results for all testing on 11/5, 9A-11A. Results 8A was collected during an attempted VOC test on 8/28.

PEMBROKE LABORATORIES, INC.

(813) 285-8742

(813) 533-0969

Mailing address: 528 Gooch Rd., Ft. Meade, FL. 33841

IMC Noralyn
P. O. Box 867
Bartow, FL 33830

Date Received: 11-09-87

Date Reported: 12-01-87

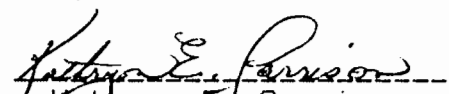
Attn: Mr. J. N. Allen

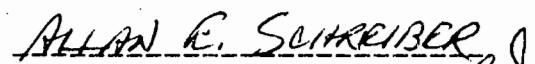
P.O. #UCP7543

Lab Numbers	MC-4953	MC-4954	MC-4955	MC-4956
Sample I.D.	8A	9A	10A	11A
BTU/lb	18357.	18291.	18462.	18354.
Density	0.980	0.977	0.977	0.977
Carbon %	87.49	87.39	87.32	87.41
Hydrogen %	10.48	10.20	10.19	10.48
Nitrogen %	0.11	<0.10	0.30	<0.10
Oxygen %	0.84	0.64	0.81	0.67
Sulfur %	1.33	1.55	1.49	1.26
Ash %	0.058	0.060	0.048	0.064

Thank you for this opportunity to serve you!

Respectfully submitted,


Kathryn E. Garrison
Laboratory Supervisor


Allan E. Schreiber
Laboratory Director

Laboratory ID 84172

C. CALIBRATION DATA

C-1 PARTICULATE TEST CALIBRATIONS

Contents: Meter Calibration
 Meter Calibration - Post test
 Pitot Calibrations
 Temperature Calibrations

METER BOX CALIBRATIONS

DIAPHRAGM PUMP LEAK CHECK @ 0.02 cfm	6/4/87	BAROMETER CALIBRATION	6/4/87
0.565 Vwet	10 mins	MERCURY BAROMETER READING	30.10
0.558 Vdry		ANEROID BAROMETER READING	30.00
0.007 diff	0.0007 leak cfm	ADJUSTED	0.10

PRE-TEST METER BOX CALIBRATION

CONTROL BOX NO. 1 DATE: 6/4/87 Pbar: 30.08 "Hg CALIBRATED BY: *J.D. Norma*

ORIFICE SETTING DELTA H in HOH	WET TEST METER VOLUME Vw cf	DRY GAS METER VOLUME Vd cf	TEMPERATURE degree F				TIME theta MINS	Y	DELTA H@
			WET TEST METER tw	DRY GAS METER		AVERAGE td			
				INLET tdi	OUTLET tdo				
0.50	6.00	F 883.23	76	106	92	559	min 15 sec 33 15.55	1.003	1.819
		I 877.00							
		V 6.23							
1.00	15.00	F 899.55	76	110	92	561	min 27 sec 40 27.67	1.007	1.836
		I 884.00							
		V 15.55							
2.00	12.00	F 917.16	76	100	80	550	min 15 sec 34 15.57	1.008	1.853
		I 905.00							
		V 12.16							
3.00	32.00	F 952.86	76	112	92	562	min 34 sec 22 34.37	1.014	1.864
		I 920.00							
		V 32.86							
4.00	45.00	F 1000.06	76	114	94	564	min 42 sec 6 42.10	1.018	1.879
		I 954.00							
		V 46.06							
5.00	45.00	F 50.56	76	116	96	566	min 37 sec 44 37.73	1.008	1.881
		I 4.00							
		V 46.56							
AVERAGE							1.010	1.855	

POST-TEST METER BOX CALIBRATION

CONTROL BOX NO. 1 DATE: 11/17/87 Pbar: 30.24 "Hg CALIBRATED BY: *J.D. Norma*

ORIFICE SETTING DELTA H in HOH	WET TEST METER VOLUME Vw cf	DRY GAS METER VOLUME Vd cf	TEMPERATURE degree F				TIME theta MINS	Y	DELTA H@
			WET TEST METER tw	DRY GAS METER		AVERAGE td			
				INLET tdi	OUTLET tdo				
0.50	30.00	F 219.39	76	110	96	563	min 78 sec 55 78.92	1.003	1.851
		I 188.00							
		V 31.39							
0.50	6.00	F 242.60	75	114	98	566	min 15 sec 51 15.85	1.006	1.850
		I 236.30							
		V 6.30							
0.50	10.00	F 257.52	75	114	98	566	min 26 sec 25 26.42	1.004	1.850
		I 247.00							
		V 10.52							
AVERAGE							1.004	1.850	

CALIBRATION EQUATIONS

$$Y = \frac{V_w \times P_{bar} (td + 460)}{V_d (P_{bar} + \Delta H / 13.6) (tw + 460)}$$

$$\Delta H @ = \frac{0.0317 \Delta H (tw + 460) \theta^2}{P_{bar} (td + 460) V_w}$$

DATE: 6/7/87

PITOT TUBE CALIBRATION

CALIBRATED BY: JD Norma

PITOT LENGTH: 8 FT

NO. 1 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
1	1.10	.78	.834	0
2	1.10	.78	.834	0
3	1.10	.78	.834	0
AVERAGE Cp			.834	

NO. 1 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
1	.24	.18	.857	0
2	.24	.18	.857	0
3	.24	.18	.857	0
AVERAGE Cp			.857	

NO. 2 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
1	1.10	.78	.834	0
2	1.10	.78	.834	0
3	1.10	.78	.834	0
AVERAGE Cp			.834	

NO. 2 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
1	.24	.18	.857	0
2	.24	.18	.857	0
3	.24	.18	.857	0
AVERAGE Cp			.857	

NO. 3 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
1	1.10	.78	.834	0
2	1.10	.78	.834	0
3	1.10	.78	.834	0
AVERAGE Cp			.834	

NO. 3 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
1	.24	.18	.857	0
2	.24	.18	.857	0
3	.24	.18	.857	0
AVERAGE Cp			.857	

NO. 4 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
AVERAGE Cp				

NO. 4 PROBE				
RUN NO.	DELTA P "S"	DELTA P "STD"	Cp "S"	DEV.
AVERAGE Cp				

PITOT TUBE CALIBRATION PROCEDURE

MANOMETER:

A permanently mounted Dwyer dual incline manometer is used for calibrating the pitot tubes. This eliminates the disconnecting and reconnecting of pitot tube lines which could cause bias calibration results. The manometer is filled with clean fluid of the proper density.

PITOT TUBES:

The pitot tubes to be calibrated are checked for nicks, dents or any misalignments that could cause bias calibration results. Permanent numbers are affixed to the pitot tubes.

CALIBRATION:

1. Level and zero dual inclined manometer.
2. Turn on fan and allow flow in calibration duct to stabilize.
3. Connect the standard pitot tube and the S type pitot tube to the manometer.
4. Seal the type S entry part with duct tape.
5. Position the standard pitot tube near the center of the duct. Check to be sure that the pitot tube is properly aligned and perpendicular to the duct.
6. The intake area is adjusted to give the desired flow rate, as measured by the standard pitot tube, and is recorded.
7. The standard pitot tube is withdrawn from the duct and the standard port is sealed with duct tape.
8. The "S" type pitot tube is inserted into the calibration duct at the same point measured by the standard pitot tube. Check the pitot tube for proper alignment in the duct. Wait for the manometer fluid to stabilize and record the measured flow rate. Remove the pitot tube from the duct.
9. Repeat steps five through eight above until three sets of velocity measurements are obtained.

CALCULATION:

1. Calculate the "S" type pitot tube coefficient for each set of measurements using the following equation:
$$C_p(S) = C_p(\text{Std}) \text{SQRT}(\Delta P \text{ Std} / \Delta P \text{ S})$$
2. The average of the three readings shall be used as the "S" type pitot tube coefficient.
3. Deviation of the three sets of readings can not exceed ± 0.02 inches.

DIAL THERMOMETER CALIBRATION

DATE 6-6-87

CALIBRATED BY _____

AMBIENT TEMP 78° deg F

BAROMETRIC PRESSURE 30.09 in Hg

DIAL THERMOMETER		TEMPERATURE READINGS			
REF THERMOMETER <u>227-769</u>		ICE WATER	OIL	BOIL WATER/OIL	
METER BOX IN	UNIT <u>78°</u>	<u>32°</u>	<u>100°</u>	<u>212°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>100°</u>	<u>212°</u>	
AVG DEV	DEV <u>0</u>	<u>0</u>	<u>0</u>	<u>0</u>	
METER BOX OUT	UNIT <u>78°</u>	<u>32°</u>	<u>101°</u>	<u>212°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>100°</u>	<u>212°</u>	
AVG DEV	DEV <u>0</u>	<u>0</u>	<u>1°</u>	<u>0°</u>	
SAMPLE BOX BOX NO 1	UNIT <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>211°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>	
AVG DEV	DEV <u>0°</u>	<u>0</u>	<u>0</u>	<u>1°</u>	
SAMPLE BOX FILTER NO 1	UNIT <u>78°</u>	<u>32°</u>	<u>151°</u>	<u>212°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>0°</u>	<u>0°</u>	
SAMPLE BOX BOX NO 2	UNIT <u>78°</u>	<u>32°</u>	<u>151</u>	<u>211°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150</u>	<u>212°</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>1°</u>	<u>1°</u>	
SAMPLE BOX FILTER NO 2	UNIT <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>210</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>0°</u>	<u>2°</u>	
SAMPLE BOX BOX NO 3	UNIT <u>78°</u>	<u>32°</u>	<u>151</u>	<u>211°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150</u>	<u>212°</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>1°</u>	<u>1°</u>	
SAMPLE BOX FILTER NO 3	UNIT <u>78°</u>	<u>32°</u>	<u>151</u>	<u>212°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150</u>	<u>212°</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>1°</u>	<u>0°</u>	
IMPINGER	UNIT <u>78°</u>	<u>32°</u>	<u>100°</u>	<u>151°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>100°</u>	<u>150°</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>0°</u>	<u>1°</u>	
STACK DRY/WET BULB	UNIT <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>211°</u>	
	REF <u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>	
AVG DEV	DEV <u>0°</u>	<u>0°</u>	<u>0</u>	<u>1</u>	

DEV FOR INDIVIDUAL READING =
$$\frac{(T_{unit} + 460) - (T_{ref} + 460)}{(T_{ref} + 460)} \times 100$$

PITOT THERMOCOUPLE CALIBRATION

DATE 6-6-87 CALIBRATED BY _____
 AMBIENT TEMP 78° deg F BAROMETRIC PRESSURE 30.09 in Hg

PROBE THERMOCOUPLE LENGTH: <u>8'</u>			TEMPERATURE READINGS			
REF THERMOMETER <u>227-769</u>			ICE WATER	OIL	BOIL WATER/OIL	
PROBE 1	BOX 1	UNIT	<u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>
		REF	<u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>
		AVG DEV	<u>0</u>	<u>0°</u>	<u>0°</u>	<u>0°</u>
PROBE 1	BOX 2	UNIT				
		REF				
		AVG DEV				
PROBE 1	BOX 3	UNIT				
		REF				
		AVG DEV				
PROBE 2	BOX 1	UNIT	<u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>
		REF	<u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>
		AVG DEV	<u>0°</u>	<u>0°</u>	<u>0°</u>	<u>0°</u>
PROBE 2	BOX 2	UNIT				
		REF				
		AVG DEV				
PROBE 2	BOX 3	UNIT				
		REF				
		AVG DEV				
PROBE 3	BOX 1	UNIT	<u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>
		REF	<u>78°</u>	<u>32°</u>	<u>150°</u>	<u>212°</u>
		AVG DEV	<u>0°</u>	<u>0°</u>	<u>0°</u>	<u>0°</u>
PROBE 3	BOX 2	UNIT				
		REF				
		AVG DEV				
PROBE 3	BOX 3	UNIT				
		REF				
		AVG DEV				

DEV FOR INDIVIDUAL READING =
$$\frac{(T_{unit} + 460) - (T_{ref} + 460)}{(T_{ref} + 460)} \times 100$$

C-2 PROCESS CALIBRATION
Contents: Scale Calibration

ER & LOCATION: KONESOME PROP. BEST-AVAILABLE COPY
 TE: 11-2-87 CONVEYOR NO.: #1 WEIGH FEEDER TEST BY: MLW
 ST LENGTH: 465 FEET: 15 REVOLUTIONS: 7Min 18 Sec. TIME
 ST CHAIN CAL CON: NA TONS R-CAL: 100.27 TONS

FUNCTION	AS FOUND	AS LEFT		
TOTAL	NA	NA		
ZERO	1586	1500		
SPAN	35127	35107		
TEST DUR	4772	4772		
CAL CON	100.27	100.3		

ZERO BALANCE TEST

TEST NO.	1	2	3	4	5	6
READING AFTER	1514	1501	1499	1500		
READING START	1586	1514	1501	1499		
DIFFERENCE	72	13	2	1		
ERROR IN %	4.5%	.8%	.1%	.06%		
ZERO NO.	1514	1501	1499	1500		

SPAN TEST

TEST NO.	2	2	3	4	5	6
READING AFTER	35127	35117	35107			
READING START	35170	35127	35117			
REG cal con	74.90	100.30	100.30			
CAL CON	100.27	100.27	100.27			
DIFFERENCE	25.37	.03	.03			
ERROR IN %	25.3%	.03%	.03%			
SPAN NO.	35127	35117	35107			

COMMENTS:

DATE & LOCATION: LONESOME

BEST AVAILABLE COPY

DATE: 11-2-87

CONVEYOR NO.: #1 WEIGH FEEDER

TEST BY: MLW

TEST LENGTH: 465

FEET: 15

REVOLUTIONS: 7 min. 18 sec. TIME

TEST CHAIN CAL CON: NA

TONS R-CAL: NA

TONS

Static Wt. = 170.2

FUNCTION	AS FOUND	AS LEFT		
TOTAL	NA	NA		
ZERO	1586	1500		
SPAN	27160	27160		
TEST DUR	4772	4772		
CAL CON TEST TONS	26.38	26.40		

ZERO BALANCE TEST

TEST NO.	1	2	3	4	5	6
READING AFTER	1514	1501	1499	1500		
READING START	1586	1514	1501	1499		
DIFFERENCE	72	13	2	1		
ERROR IN %	4.5%	0.8%	.1%	.06%		
ZERO NO.	1514	1501	1499	1500		

SPAN TEST

TEST NO.	1	2	3	4	5	6
READING AFTER	27160	27140	27119	27201	27180	27160
READING START		27160	27140	27119	27201	27180
REG cal con TEST TONS	34.10	26.40	26.40	26.30	26.40	26.40
CAL CON Test TONS	26.38	26.38	26.38	26.38	26.38	26.38
DIFFERENCE	7.72	.02	.02	.08	.02	.02
ERROR IN %	29.3%	.07%	.07%	.3%	.07	.07
SPAN NO.	27160	27140	27119	27201	27180	27160

COMMENTS: USED static wts. on span Test

DER

DEC 17 1987

BAQM

OPERATION ANALYSIS
SUPPLEMENT TO EMISSION TESTS

FOR

IMC LONESOME

NO 1 DRYER

TESTS: 4/3/87
6/10/87

AND

NO 2 DRYER

TESTS: 6/8/87
6/9/87

Prepared by:

C. D. Turley

July 15, 1987

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A. TEST CALCULATIONS

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INTRODUCTION

This is a analysis of the various operating parameters that have been measured or calculated during the compliance testing done on the IMC Lonesome dryers. The purpose of this is to provide additional information to assist in the interpretation of the compliance test reports related to these dryers. This report is subitted separately and refers to the information in the referenced tests. The results from these tests are used in the summary calculation tables of this report. Individual run and traverse calculation pages from the reports are included in the Appendix. Also included for reference are the rejected results from the April 3, 1987 test of the No 1 Dryer.

Tests:	Date	Dryer	Test types
	04/03	1	Particulate, VE, SO ₂ , NO _x , CO, and VOC.
	06/08	2	NO _x only.
	06/09	2	Particulate, VE, SO ₂ , NO _x , CO, and VOC.
	06/10	1	Particulate, VE, SO ₂ , NO _x , CO, and VOC.

The permits that apply to these dryers have some basic inconsistencies with normal operating procedures for such units. This analysis was also done to begin to develop a consistent basis for describing the operation of these dryers. They are permitted below their capacity to operate. This type of analysis is necessary if estimates of the emissions and their ability to comply are to be made at their real capacity. It is also necessary if the basis of the permit limitations are to be evaluated.

The limitations in the permit application for the fuel conversion of these dryer which were based on estimates and assumptions. This type of analysis is necessary to evaluate the validity of those assumptions. Much of the fuel related estimates were based on emission factors that apply to combustion in steam generating boilers which are in not similiar to the method of operation of a phosphate rock dryer. A steam generator operates at the minimum excess air possible in order to maximize gas temperatures for the purpose of heat transfer. In a rock dryer, the purpose of combustion is to heat air that will evaporate moisture from the feed rock. Compared to a boiler, a dryer operates at large quantities of excess air.

SCALE VS BIN LEVEL COMPARISON

During tests of both dryers at Lonesome, measurements were made that enable the comparison of the dryer operating tonnage determined by scale reading and by bin level measurements. The rock moisture and density were also determined in order to produce a more reliable calculation. These results were also calculated in the same manner that the previous 4/3/87 test tonnages were determined in order to estimate the possible error that had occurred.

date	04/03			06/08	06/09			06/10		
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	AVE	1R	2R	3R	1R	2R	3R
Scale calculations										
stph				507	492	499	499	507	506	508
HOHin				12.0%	12.0%	11.8%	8.8%	11.2%	11.2%	11.0%
dtpH				446	433	440	455	450	450	452
avg				446			443			450
Bin Level calculations										
Tin				84.2	83	82	81	78	79	79
density	95	95	95	96.5	95	93	92	94	97	96
HOHout				3.6%	4.0%	4.0%	4.2%	3.4%	3.0%	4.2%
den adj				94.4	92.7	90.7	89.6	92.0	95.2	93.5
btph	538	548	404	482	434	435	357	451	442	1006
dtpH				464	417	417	342	436	429	964
Comparison of dry production rate determination										
% diff				4.1%	-3.7%	-5.2%		-3.2%	-4.6%	
abs avg										4.2%
Bin Level prediction of scale reading based on permit assumptions										
pre ass	613	625	460	541	495	506	420	519	494	1135
% diff				6.2%	0.6%	1.4%		2.5%	-2.5%	
abs avg										2.6%

On the tests of 6/9 and 6/10, there were some obvious errors in the final bin measurements. These were ignored in the calculation of the average of the percent difference in the determination of the dry TPH by the two methods which was within an 4.2%. The density of the rock was determined on a dry basis. In order to calculate based on the bin level measurements, this measurement was adjusted based on the final moisture content of the rock in the same ratio as the specific gravity with and without the moisture, $((1-\%HOH) \times 2.6 + \%HOH \times 1) / 2.6$. It should be noted that the density assumed for the 4/3 test of 95 lb/cf is equal to the average of the eight measurements made during the June tests. The apparent agreement of predicting the scale tonnage based on bin level measurements based on parameters assumed in the permit is 2.6% for the June tests.

In the future, it reasonable to expect that either measurement of operating tonnage could be determined by either method in compliance with 40 CFR 60.403(d). The bin level method would be considered as a contingency means in order to conduct a test if scale problems could not be rectified for a scheduled test. Compliance with the 5% limit would be maintained based on an assumption of 95 lb/cf density. In either case, in order to produce reliable calculations, rock moisture determinations should be conducted. Increased care must be taken with the measurement of the bin levels for such testing in the future.

This plan for a back-up method of determining feed rate is necessary because of the limited operation of the No 1 dryer. No 2 was operated only to conduct these tests. The current schedule requires operation only once every two to three weeks. If a test is scheduled for one of these operations, the status of the scale will not be known until the start-up on the previous night. The test could be postponed but the production cannot. This will mean that the test will be postponed until the next operation in two or three weeks if bin level readings cannot be used.

FUEL METER VS TANK LEVEL

During the Lonesome dryer testing, corresponding readings for the oil meters and the supply tank level were taken for comparison. Oil samples collected to correspond to the test runs in June testing. The April analyses cited are the average of four analyses for 2 split samples by two different labs. The analyses are included in Appendix B.

date	04/03			06/08	06/09	06/10				
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	AVE	1R	2R	3R	1R	2R	3R
Meter calculations (used for test result calculations)										
M-gph	906	856	880	950	920	943	923	940	919	795
lb/gal	8.176	8.176	8.176	8.097	8.105	8.097	8.097	8.088	8.072	8.105
btu/lb	18893	18893	18893	18493	18538	18521	18492	18497	18545	18509
MMbtu	140	132	136	142	138	141	138	141	138	119
Tank level calculations for comparison										
L-gph	867	752	786		841	894	888	834	814	729
Toil	170	170	170		196	192	196	195	192	190
Mark's Standard Handbook for Mechanical Engineers, Seventh Edition										
Table 13, pg 7-22: 0.00035 /F for <15 deg API										
expand	3.0%	3.0%	3.0%		3.7%	3.6%	4.1%	3.8%	3.6%	3.5%
L-exp	893	774	809		872	926	925	865	843	754
% diff	-1.5%	-9.5%	-8.1%		-5.2%	-1.8%	0.2%	-8.0%	-8.3%	-5.1%
			-6.4%				-2.3%			-7.1%

The level flow rate was corrected to the meter oil temperature by the cited coefficient of expansion. The fuel oil in the tank had been preheated and would have expanded to some degree also. The temperature difference for the expansion was based on meter temperature and the ambient temperature recorded. The overall temperature of the oil in the tank would be difficult to determine. The compliance calculation were based on the meter flow readings. This also affects the resulting emissions in that the fuel analyses will have a reference temperature other than the operating temperature recorded. This should bias fuel related calculations high.

The dryers have two fuel supply systems. One contains the oil meter and electric preheaters and the other has a boiler preheater with not oil meter in the system. For all testing, the system with the meter has been used. If it could not be, then the only measurement of fuel useage would be the tank level indication. There are not specific requirements regarding this determination that apply to this testing. In order to produce the most reliable results, an attempt to use the meter will be made.

STACK MOISTURE

The stack moisture content of the these dryers should always be that of saturation at the stack temperature. Several separate determinations of the moisture content were made to verify the saturation. The results were not conclusive. The earlier test measurements exceed saturation, while the later measured moisture contents are generally below saturation.

date	04/03			06/08	06/09			06/10		
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	FINAL	1R	2R	3R	1R	2R	3R
Pbar	29.59	29.59	29.59	30.22	30.15	30.15	30.15	30.14	30.14	30.14
Scrubber Inlet Duct										
in Tdry	178	178	175	188	189	188	182	176	176	176
in Twet				152	151	150	148	150	150	150
delta P	20.1	20.2	17.7	25.0	24.5	25.8	25.0	28.4	28.4	28.5
Pduct	31.07	31.08	30.89	32.06	31.95	32.04	31.99	32.23	32.23	32.24
%sat				24.8%	24.3%	23.6%	22.5%	23.5%	23.5%	23.5%
Stack conditions										
Tstk	153.0	153.0	150.0	156.4	155.8	155.8	153.6	155.9	155.8	151.4
Pstk	29.56	29.56	29.56	30.16	30.09	30.10	30.09	30.11	30.11	30.11
%satstk	27.6%	27.6%	25.6%	29.4%	29.0%	29.0%	27.5%	29.1%	29.0%	26.0%
Average of two %HOH determinations during SO2 runs by ACE										
c%HOH	25.3%	22.3%	21.1%		29.6%	30.6%	28.2%	28.8%	29.4%	25.7%
%HOH used in the emission calculation for compliance										
%meas	27.5%	19.5%	24.9%	29.4%	25.8%	26.7%	26.3%	24.1%	23.4%	23.6%

STACK MOLECULAR WEIGHT

For all the complete test series, two separate measurements of the stack oxygen and carbon dioxide contents were made for comparison. These results are presented below. The stack molecular weights used in compliance are included.

date	04/03			06/08	06/09			06/10		
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	FINAL	1R	2R	3R	1R	2R	3R
Readings determined by Fryite Analyzers (3 grab avg)										
OO	15.0%	14.5%	15.0%	13.0%	13.5%	13.8%	14.0%	14.5%	14.5%	14.5%
OCO	4.5%	3.5%	4.5%	4.0%	4.5%	4.5%	4.5%	4.5%	4.5%	4.5%
Mdry	29.32	29.14	29.32	29.16	29.27	29.27	29.28	29.30	29.30	29.30
Readings determined by Orsat (integrated) by ACE										
cOO	14.2%	14.2%	14.3%		13.8%	14.0%	14.4%	14.6%	14.6%	14.6%
cOCO	4.3%	4.2%	4.0%		4.6%	4.6%	3.6%	4.3%	4.3%	4.3%
cMdry	29.26	29.24	29.21		29.29	29.30	29.15	29.27	29.27	29.27
Ms based on Fryite and stack temperature saturation %HOH										
Ms sat	26.20	26.07	26.42	25.88	26.00	26.00	26.18	26.01	26.02	26.36
Ms based on ACE determinations										
cMs	26.41	26.73	26.85		25.95	25.84	26.01	26.02	25.96	26.37
Compliance calculation based on Fryite and measured %HOH										
Ms used	26.21	26.97	26.50	25.88	26.35	26.26	26.31	26.57	26.66	26.64

VOLUMETRIC FLOW RATE

The stack volumetric flow determinations from each of the runs of the complete test series are included. The values on 6/8 are the results of the final traverse for that test day. For the June tests, the cyclonic yaw angles were determined and used to calculate the cyclonic flow rates. The affect of the stack saturation and measured moisture contents are also compared. The bias shows the degree that the compliance emission are affected by ignoring the effects in the stack gases.

date	04/03			06/08	06/09			06/10		
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	FINAL	1R	2R	3R	1R	2R	3R
ACFM measured										
meas	136946	136765	132970	168744	155830	148541	154971	130781	130870	127612
						No 2 avg: 157022		No 1 avg: 132657		
cyc				161963	150120	143049	148112	126016	127475	122641
SCFM comparison (flow - meas) / flow										
meas	84487	93695	85543	102905	99639	93738	98811	85575	86520	84772
						No 2 avg: 98773		No 1 avg: 86766		
wet	116534	116392	113906	145700	134371	127857	134120	112818	112916	110900
sat	84391	84291	84728	102902	95372	90751	97267	80021	80163	82017
bias	0.1%	11.2%	1.0%	0.0%	4.5%	3.3%	1.6%	6.9%	7.9%	3.4%
cyc				98770	95988	90273	94438	82456	84276	81470
bias				4.2%	3.8%	3.8%	4.6%	3.8%	2.7%	4.1%

There is an apparent difference in the flow of the two systems. This was generally true in the previous testing done by Brewster. There is no obvious explanation for this difference.

SO2 CALCULATIONS

Samples of the water reporting to the scrubber and underflow were collected during the runs of the tests. On the last two days, these corresponded to the determination of the stack SO₂ emissions. The samples were analyzed for sulfate content. This was done as method to determine the apparent collection efficiency of the rock in the dryer and the scrubber. Based on the original application for the PSD permit for the conversion of the fuels for these two dryers, the removal efficiency for the rock should have been approximately 82%. In the following table, each of the two dryers seem to achieve 92%. The scrubbers seems to demonstrate a collection of approximately 50%.

date	04/03			06/08	06/09			06/10		
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	AVE	1R	2R	3R	1R	2R	3R
gpm	450	450	450	500	570	570	570	580	580	580
Water Sulfate content in mg/l										
in				94.6	114.4	107.4	116.5	100.4	108.6	109.3
out				154.3	166.2	145.3	158.8	145.3	138.3	134.9
fuel S	1.85%	1.85%	1.85%	1.28%	1.28%	1.26%	1.30%	1.26%	1.25%	1.24%
Sulfur dioxide (lb/hr)										
scrubber				9.9	9.8	7.2	8.0	8.7	5.7	4.9
fuel	275	259	267	197	191	192	194	192	186	160
stack	27.9	20.7	18.2		6.5	8.9	7.5	8.0	9.2	4.8
Apparent efficiency										
rock					91.4%	91.7%	92.0%	91.3%	92.0%	93.9%
scrubber					60.0%	44.8%	51.6%	52.1%	38.5%	50.7%
total	89.9%	92.0%	93.2%		96.6%	95.4%	96.1%	95.8%	95.1%	97.0%
Minimum efficiency required for compliance @ 1.1 lb/mmBtu										
min eff	43.9%	43.9%	43.9%		20.3%	19.2%	21.8%	19.3%	18.4%	17.9%

The collection of the SO₂ based on the fuel analysis into the systems confirm that there is a removal and that it is similar in each scrubber. It should be noted that these results are applicable to the rock being dried for these tests.

PARTICULATE COLLECTION

The scrubber water supply and underflow samples were also analyzed for filterable solids as means of determining the apparent particulate collection efficiency for the scrubbers. Each of the scrubbers seem to operate at approximately 97% collection efficiency.

date	04/03			06/08	06/09	06/10				
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	AVE	1R	2R	3R	1R	2R	3R
Suspended solid in scrubber water in mg/l										
in				9	4	59	107	332	74	26
out				2138	1571	1287	1130	1756	1887	1272
Particulate lb/hr										
underflow				532	446	350	291	413	526	361
stack	13.0	19.1	12.3		12.0	13.4	10.8	10.8	12.1	14.6
apparent eff					97.4%	96.3%	96.4%	97.4%	97.8%	96.1%
minimum eff @ 25 lb/hr				95.3%	94.4%	92.9%	91.4%	93.9%	95.2%	93.1%
Ash particulate reporting to dryer										
%Ash	0.09%	0.09%	0.09%	0.05%	0.04%	0.02%	0.02%	0.02%	0.05%	0.06%
lb/hr	6.4	6.0	6.2	4.2	3.1	1.6	1.7	1.4	3.7	3.6

NOX MODEL

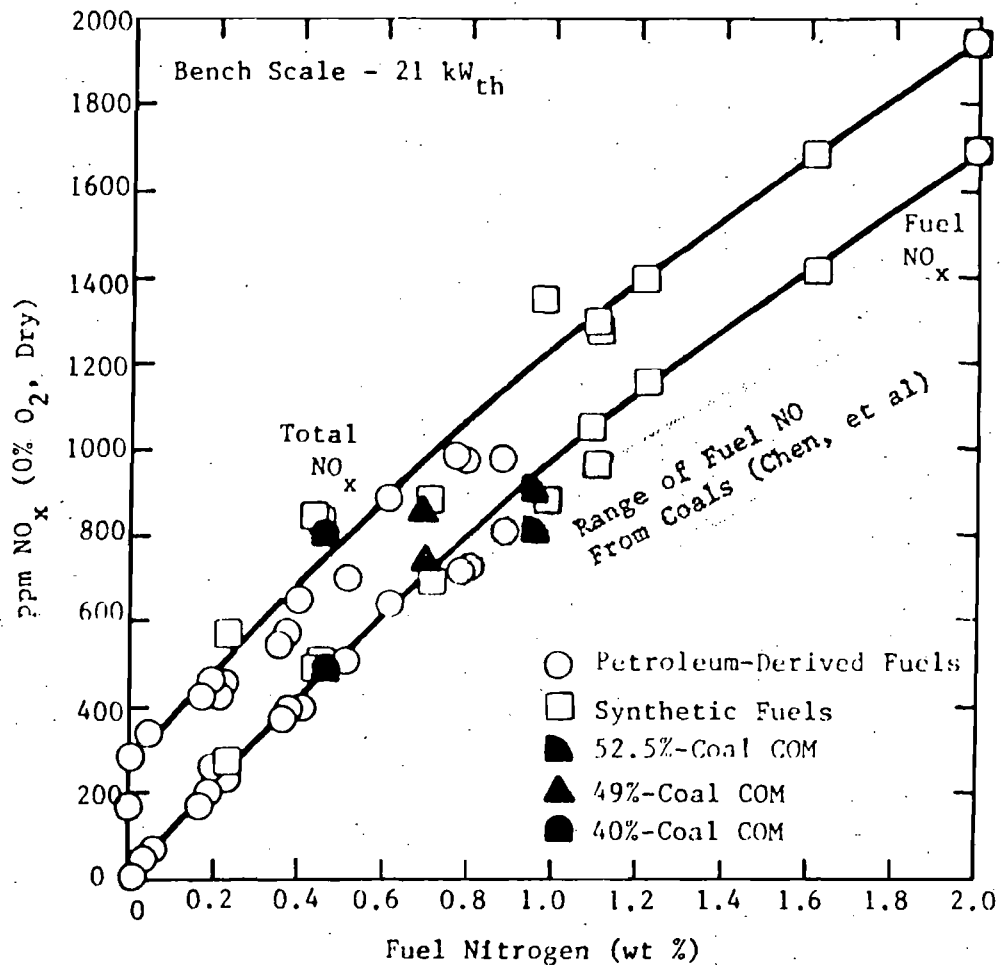
Prior to the testing of the dryers, an attempt was made to predict the nitrogen oxide emissions. The NO_x emissions will result from either oxidation of the nitrogen content in the fuel and disassociation of the nitrogen in the combustion air. These reactions are reversible depending on the conditions of the flame combustion. The fuel nitrogen can also produce diatomic nitrogen which would not contribute to the emissions. These tests provide a limited number of points to evaluate the modeling predictions. The models:

1. 100% conversion of fuel nitrogen to nitrogen dioxide.
2. The following figure from a paper by England, et al, presented at the 4th Symposium on Coal Slurry Combustion, May 10-12, 1982. The approximation of ppm @ 3% EA = 100 x %N fuel + 225 was used to represent lower portion of curve.
3. The equation from AP42, pg 1.3-2, lb/1000gal = 22 + 400 x (%N fuel)².

The data points used were determinations which had a specific fuel analysis associated with it. The model calculations were based on lb/lb fuel to eliminate the uncertainty associated with the saturation and cyclonic considerations in the determination of volumetric flow rate. The measured values used were those of the compliance determination. A simple linear regression was done for the eight data points. The models and results are compared in the graph following.

date	04/03			06/08	06/09	06/10				
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	2F	1R	2R	3R	1R	2R	3R
%N	0.30%	0.30%	0.30%	0.25%	0.14%	0.28%	0.23%	0.15%	0.23%	0.30%
meas	40.6	44.3	40.5	44.4	34.3	34.9	37.5	31.3	31.0	29.2
fuel	73.7	69.5	71.5	64.7	34.3	70.2	56.5	37.5	56.1	63.5
england	82.0	77.4	79.6	78.1	56.8	81.4	71.2	60.1	71.2	71.4
AP42	53.1	50.1	51.6	45.7	27.5	50.3	39.9	29.1	39.7	46.1

It can be concluded that the AP42 curve is more reliable than the England model. It is also apparent that, for this limited operation, none of the predictions were valid. The regression curve implies that the emissions seem to have limited dependence on fuel nitrogen. This could only be confirmed at several other operating conditions. The most interesting point would be at maximum fuel usage which is currently unattainable due to permit restriction on tonnage.

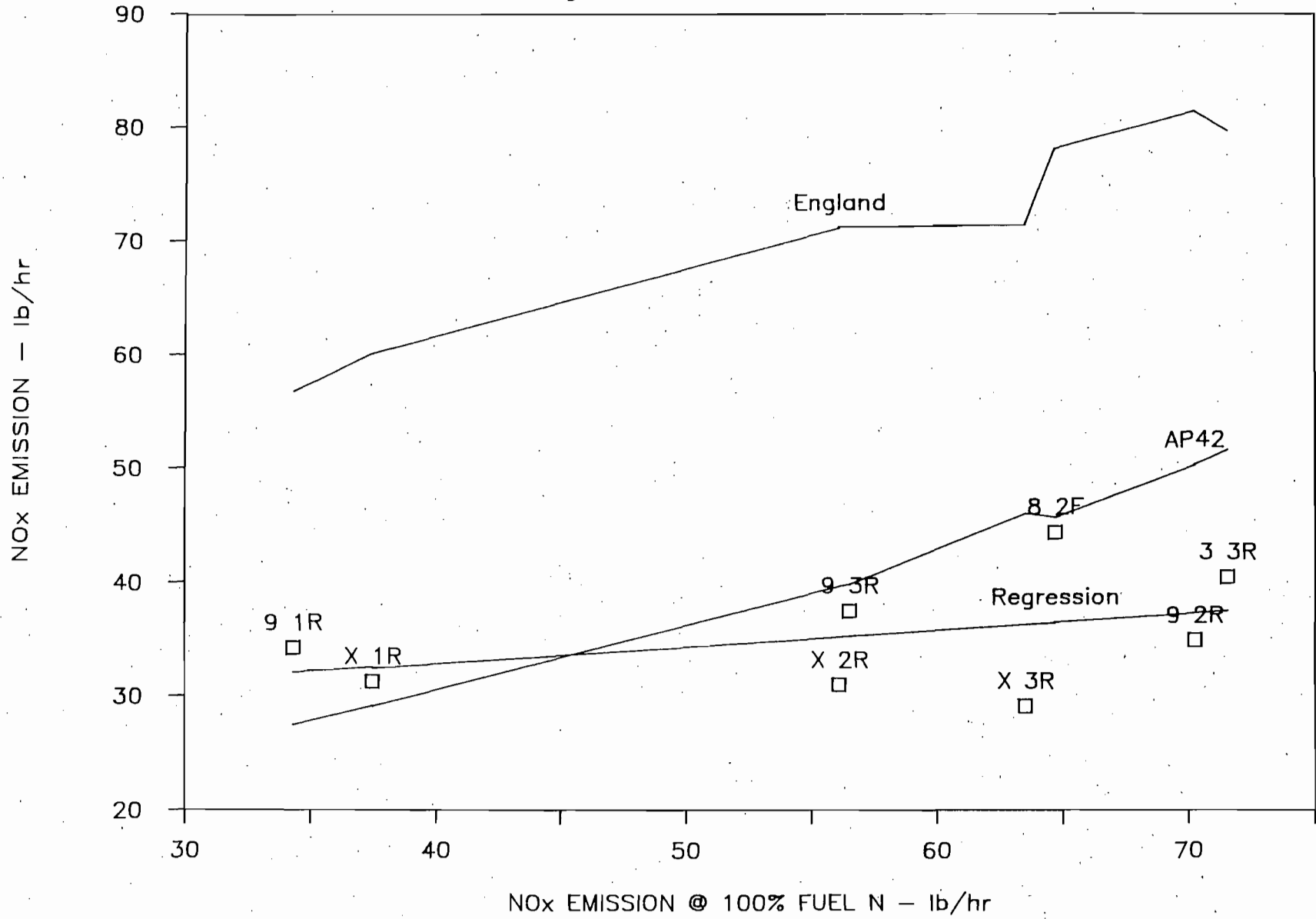


Influence of Fuel Nitrogen on NO_x Emissions
(Unstaged, 3% Excess O₂).

FIGURE 1

ENGLAND NO_x RELATION

Figure 2 NOx MODEL COMPARISON



COMBUSTION CALCULATIONS

The following are summary tables of the run calculations in Appendix A based on using the fuel analysis and the two measured volumetric flow rates for comparison. These should be considered on preliminary type calculations since the leakage of air in with the rock is not considered. This is usually balanced visually by the operator and should be minimal unless conditions have changed. The meter gph was used without an expansion correction. The radiant and convective heat loss have not used in the heat balance calculation.

The water balance for the run is based on the calculation of the amount of moisture evaporated from the rock, the combustion water, and the amount indicated in the stack gases. These are used to calculate the amount evaporated or condensed in the scrubber.

date	04/03			06/08	06/09	06/10				
dryer	1	1	1	2	2	2	2	1	1	1
run	1R	2R	3R	FINAL	1R	2R	3R	1R	2R	3R
STACK GAS COMPOSITION										
Fuel Analysis										
%C	86.86%	86.86%	86.86%	86.59%	86.79%	86.65%	86.55%	86.79%	86.67%	86.90%
%H	10.09%	10.09%	10.09%	10.16%	10.10%	10.58%	10.30%	10.54%	10.52%	10.47%
%O	1.25%	1.25%	1.25%	1.72%	1.68%	1.23%	1.62%	1.26%	1.33%	1.09%
Measured values										
OO	15.0%	14.5%	15.0%	13.0%	13.5%	13.8%	14.0%	14.5%	14.5%	14.5%
OCO	4.5%	3.5%	4.5%	4.0%	4.5%	4.5%	4.5%	4.5%	4.5%	4.5%
EA	2.53	3.52	2.53	2.96	2.51	2.48	2.49	2.49	2.49	2.49
Values based on standard volumetric flow										
OO				16.6%	16.6%	16.2%	16.5%	15.7%	15.9%	16.5%
OCO				3.4%	3.4%	3.7%	3.4%	4.1%	3.9%	3.5%
EA				3.61	3.61	3.19	3.54	2.84	2.98	3.48
Values based on volumetric flow										
OO				16.4%	16.2%	15.8%	16.3%	15.2%	15.4%	16.2%
OCO				3.5%	3.7%	4.0%	3.7%	4.5%	4.3%	3.7%
EA				3.43	3.25	2.86	3.28	2.47	2.60	3.17
HOH BALANCE lb/lb fuel										
products				0.914	0.909	0.952	0.927	0.949	0.947	0.942
rock				11.49	10.99	10.62	10.85	10.76	11.54	11.19
Scrubber evaporation or condensation (-)										
std flow				3.48	1.40	1.19	1.71	-1.48	-2.33	-0.55
cyc flow				2.84	2.49	2.17	1.67	0.16	-0.21	0.17
HEAT BALANCE Btu/lb fuel										
fuel				18493	18538	18521	18492	18497	18545	18509
std				23082	19869	19379	20531	16817	16932	19051
cyc				22411	20879	20293	20432	18369	18970	19710

These calculations indicate that the systems are reasonably well defined for calculation on a first approximation basis. Further refinement is possible in future permit testing. They are a basis for analysis of existing permit condition and refining them in terms of operating conditions.

APPENDIX

A. TEST CALCULATIONS

Contents: Individual Velocity Traverse Calculations
Comparison Calculations

Individual Velocity Traverse Calculations

No: 8 1F		Usages
Date: 06/08	Rock: 492 TPH	
Time: 13:23	0.12 HOH	
Pbar: 30.22	433 Dry	
Pst: -0.8		
Ps: 30.16 = Pbar+Pst/13.6	Oil: 990 GPH	
Cp: 0.834	8.097 lb/gal	
Fv: 12.98 = 85.49*Cp*sqrt(1/Ps)	18493 Btu/lb	
Dia: 96 Area: 50.27	148 mmBtu/hr	
%O: 13.0%		
%CO: 4.0%		
Mdry: 29.16		
H: 10.16%		
O: 1.72%		
C: 86.59%	%EA: 2.958 based on fuel analysis and %CO	

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

	Assumed Twb:	155.3					
	sat %HOH:	28.55%					
Ms = Mdry*(1-%HOH)+18*%HOH	Ms:	25.97					
Avg:	16.3	155.3	28.56%				
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.64	16	154	27.70%	50.5	50.4	48.5
2	0.85	16	156	29.08%	58.3	58.4	56.1
3	0.75	16	156	29.08%	54.7	54.8	52.7
4	0.65	16	156	29.08%	50.9	51.0	49.1
5	0.98	16	156	29.08%	62.6	62.7	60.2
6	0.90	16	156	29.08%	59.9	60.0	57.7
SW-1	0.60	18	154	27.70%	48.9	48.8	46.4
2	0.82	18	155	28.38%	57.2	57.2	54.4
3	0.74	16	155	28.38%	54.4	54.3	52.2
4	0.86	16	155	28.38%	58.6	58.6	56.3
5	0.95	16	155	28.38%	61.6	61.6	59.2
6	0.94	16	155	28.38%	61.3	61.2	58.9
				avg			
				fps:	56.6	56.6	54.3
				acfm:	170630	170642	163760
				%HOH:	28.55%	28.56%	
				scfm:	105461	105463	101209
				Emission:	lb/hr		
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	59	46	7.1646	44.6	44.6	42.8	

COMPARISON CALCULATIONS

RUN: 1F 06/08/87 time 13:23 ref: 1A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8659	0.0722	OCO	1 0.0722	OCO	0.21	Btu/lb
hydrogen	1	0.1016	0.1016	NO2	1 0.0002	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0025	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0128	0.0004	-fuel O	-0.0005	NN	0.25	RT/p
oxygen	16	0.0172	0.0011	HOH	0.25 0.0254	rock	0.25	378.20
ash		0.0005				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0976	cmb NN	0.3679	10.303	139.16	155	87.8	174
cmb NN	28	0.3679	OCO	44 0.0722	3.175	27.29	155	87.8	45
total		176.07	NO2	46 0.0002	0.008	0.07			
			OSO	64 0.0004	0.026	0.15			
			total dry			166.67			
			HOH	18 0.0508	0.914	19.21	155	87.8	956
			total products						1175

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18493 btu/lb
Fuel GPH	990	0	148 mmbtu/HR
lb/gal	8.097	-100.0%	1269 GPH @190 mmBtu/hr
lb/hr	8016		

SCRUBBER	gpm Inlet Twb	In hoh sat
	500 152	24.81%

ROCK	%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	492	12.0% in						
Dry TPH	433							
Calc Bin TPH	449	3.6% out						
Meas Bin TPH	485		Rock dried	107.93		194	84.2	2963
(Meas-Calc)/Meas	7.5%		Rock HOH	4.03		194	84.2	443
Evaporated TPH	42.8		Evap HOH	0.5937 10.69	224.5	155	84.2	11189

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	250 225	405 stoichometric	0.0082	0.0675
	AP42	152 134	475 @ 3% Excess Air	0.0099	0.0815
			286 stoichometric	0.0058	0.0477

Standard Flow at measured %HOH	wSCFM	measured %HOH	NOx emission model	NOx meas	fuel	eng	ap42		
	147608	28.55%		59	86	103	60		
Dry Flow	789.40			45.4	65.8	79.5	46.5		
Stk HOH	15.015	315.48							
Net HOH 0.1896	3.414	71.73	155	88	3568	54.73	evap gpm		
Scrubber HOH	27.774		148	88	1672				
Excess Air		622.73							
EA OO 0.3452	11.046	130.55	155	88	171				
EA NN 1.3014	36.438	492.18	155	88	614				
			total	21795					
				p + rk	23.59%	16.54%	3.46%	29.21	EA 3.537
				inlet	24.81%				
				meas	28.55%	13.00%	4.00%	29.16	2.958
				ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	saturation %HOH	NOx emission model	NOx meas	fuel	eng	ap42		
	141664	28.55%		59	89	108	63		
Dry Flow	757.61			43.6	65.8	79.5	46.5		
Stk HOH	14.410	302.78							
Net HOH 0.1561	2.809	59.02	155	88	2936	45.04	evap gpm		
Scrubber HOH	28.378		148	88	1708				
Excess Air		590.94							
EA OO 0.3276	10.482	123.89	155	88	163				
EA NN 1.2349	34.578	467.06	155	88	583				
			total	21159					
				sat	28.55%	16.35%	3.60%	29.23	3.356
				cyc	28.55%				
				p + rk	24.34%				

Individual Velocity Traverse Calculations

No:	8 2F	Usages
Date:	06/08	Rock: 512 TPH
Time:	13:49	0.12 HOH
Pbar:	30.22	451 Dry
Pst:	-0.8	
Ps:	30.16 = Pbar+Pst/13.6	Oil: 972 GPH
Cp:	0.834	8.097 lb/gal
Fv:	12.98 = 85.49*Cp*sqrt(1/Ps)	18493 Btu/lb
Dia:	96 Area: 50.27	146 mmBtu/hr
%OO:	13.0%	
%OCO:	4.0%	
Mdry:	29.16	
H:	10.16%	
O:	1.72%	
C:	86.59%	%EA: 2.958 based on fuel analysis and %OCO

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

	Assumed Twb:	155.8					
	sat %HOH:	28.96%					
Ms = Mdry*(1-%HOH)+18*%HOH	Ms:	25.93					
Avg:	16.3	155.8	28.97%				
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.65	16	154	27.70%	51.0	50.8	48.8
2	0.84	16	156	29.08%	58.0	58.0	55.8
3	0.80	16	157	29.79%	56.6	56.7	54.5
4	0.58	16	157	29.79%	48.2	48.3	46.4
5	0.87	16	156	29.08%	59.0	59.0	56.8
6	0.92	16	156	29.08%	60.7	60.7	58.4
SW-1	0.65	18	154	27.70%	51.0	50.8	48.3
2	0.85	18	156	29.08%	58.3	58.4	55.5
3	0.86	16	156	29.08%	58.7	58.7	56.4
4	0.76	16	156	29.08%	55.2	55.2	53.0
5	0.95	16	156	29.08%	61.7	61.7	59.3
6	0.98	16	156	29.08%	62.6	62.7	60.2
				avg			
				fps:	56.7	56.8	54.5
				acfm:	171144	171154	164244
				%HOH:	28.96%	28.97%	
				scfm:	105077	105075	100832
				Emission:	lb/hr		
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	59	46	7.1646	44.4	44.4	42.6	

COMPARISON CALCULATIONS

RUN: 2F 06/08/87 time 13:49 ref: LA 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8659	0.0722	OCO	1 0.0722	OCO	0.21	Btu/lb
hydrogen	1	0.1016	0.1016	NO2	1 0.0002	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0025	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0128	0.0004	-fuel O	-0.0005	NN	0.25	RT/p
oxygen	16	0.0172	0.0011	HOH	0.25 0.0254	rock	0.25	378.20
ash		0.0005				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0976	cmb NN	0.3679	10.303	139.16	156	87.8	175
cmb NN	28	0.3679	OCO	44 0.0722	3.175	27.29	156	87.8	45
total		176.07	NO2	46 0.0002	0.008	0.07			
			OSO	64 0.0004	0.026	0.15			
			total dry			166.67			
			HOH	18 0.0508	0.914	19.21	156	87.8	956
			total products						1177

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18493 btu/lb
Fuel GPH	972	0	146 mmbtu/HR
lb/gal	8.097	-100.0%	1269 GPH @190 mmBtu/hr
lb/hr	7873		

SCRUBBER	gpm	Inlet Twb	In hoh sat
	500	152	24.81%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	512	12.0%	in						
Dry TPH	451								
Calc Bin TPH	468	3.6%	out						
Meas Bin TPH	485			Rock dried	114.52		194	84.2	3144
(Meas-Calc)/Meas	3.6%			Rock HOH	4.28		194	84.2	470
Evaporated TPH	44.6			Evap HOH	0.6300	11.34	156	84.2	11875

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	250	225	405 stoichometric	0.0082 0.0675
	AP42	152	134	475 @ 3% Excess Air	0.0099 0.0815
				286 stoichometric	0.0058 0.0477

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	147912	28.96%							NOx ppm	59	84	102	60	
Dry Flow					800.75				lb/hr	45.2	64.7	78.1	45.7	
Stk HOH	15.536		326.43						evap gpm					
Net HOH	0.1823	3.282	68.96	156	88	3432	51.68							
Scrubber HOH	28.469			148	88	1714								
Excess Air			634.08						Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.3515	11.247	132.93	156	88	176			std	28.96%	16.60%	3.41%	29.21	3.601
EA NN	1.3251	37.102	501.15	156	88	631			p + rk	24.33%				
						22618			inlet	24.81%				
									meas	28.96%	13.00%	4.00%	29.16	2.958
									ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	141949	28.96%							NOx ppm	59	88	106	62	
Dry Flow					768.47				lb/hr	43.4	64.7	78.1	45.7	
Stk HOH	14.910		313.27						evap gpm					
Net HOH	0.1475	2.656	55.80	156	88	2777	41.82							
Scrubber HOH	29.096			148	88	1752								
Excess Air			601.80						Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.3336	10.675	126.16	156	88	167			std	28.96%				
EA NN	1.2576	35.213	475.63	156	88	599			p + rk	28.96%	16.42%	3.55%	29.22	3.418
						21959			sat	28.96%				
									cyc	28.96%				
									p + rk	25.10%				

Individual Velocity Traverse Calculations

No: 8 3F		Usages
Date: 06/08	Rock: 516 TPH	
Time: 14:16	0.12 HOH	
Pbar: 30.22	454 Dry	
Pst: -0.8		
Ps: 30.16 = Pbar+Pst/13.6	Oil: 950 GPH	
Cp: 0.834	8.097 lb/gal	
Fv: 12.98 = 85.49*Cp*sqrt(1/Ps)	18493 Btu/lb	
Dia: 96 Area: 50.27	142 mmBtu/hr	
%OO: 13.0%		
%OCO: 4.0%		
Mdry: 29.16		
H: 10.16%		
O: 1.72%		
C: 86.59%	%EA: 2.958 based on fuel analysis and %OCO	

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

		Assumed Twb:	156.1				
		sat %HOH:	29.14%				
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:	25.91				
Avg:		16.3	156.1	29.14%			
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.55	16	154	27.70%	47.0	46.7	44.9
2	0.82	16	156	29.08%	57.3	57.3	55.1
3	0.80	16	156	29.08%	56.6	56.6	54.4
4	0.60	16	157	29.79%	49.0	49.1	47.2
5	0.92	16	157	29.79%	60.7	60.9	58.5
6	0.98	16	156	29.08%	62.7	62.7	60.2
SW-1	0.55	18	154	27.70%	47.0	46.7	44.4
2	0.82	18	156	29.08%	57.3	57.3	54.5
3	0.80	16	157	29.79%	56.6	56.7	54.5
4	0.68	16	157	29.79%	52.2	52.3	50.3
5	0.95	16	157	29.79%	61.7	61.8	59.4
6	0.88	16	156	29.08%	59.4	59.4	57.1
				avg			
				fps:	55.6	55.6	53.4
				acfm:	167769	167794	161027
				%HOH:	29.14%	29.14%	
				scfm:	102708	102712	98570
				Emission:	lb/hr		
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	56	46	7.1646	41.2	41.2	39.5	

COMPARISON CALCULATIONS

RUN: 3F 06/08/87 time 14:16 ref: LA 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		Btu/lb
carbon	12	0.8659	0.0722	OCO	1 0.0722	OCO	0.21	
hydrogen	1	0.1016	0.1016	NO2	1 0.0002	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0025	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0128	0.0004	-fuel O	-0.0005	NN	0.25	RT/p
oxygen	16	0.0172	0.0011	HOH	0.25 0.0254	rock	0.25	378.20
ash		0.0005				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0976	cmb NN	0.3679	10.303	139.16	156	87.8	176
cmb NN	28	0.3679	OCO	44 0.0722	3.175	27.29	156	87.8	46
total		176.07	NO2	46 0.0002	0.008	0.07			
			OSO	64 0.0004	0.026	0.15			
			total dry			166.67			
			HOH	18 0.0508	0.914	19.21	156	87.8	956
			total products						1178

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18493 btu/lb
Fuel GPH	950	0	142 mmbtu/HR
lb/gal	8.097	-100.0%	1269 GPH @190 mmBtu/hr
lb/hr	7692		

SCRUBBER	gpm Inlet Twb	In hoh sat
	500	152 24.81%

ROCK	%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	516	12.0% in						
Dry TPH	454							
Calc Bin TPH	471	3.6% out						
Meas Bin TPH	485		Rock dried	117.98		194	84.2	3239
(Meas-Calc)/Meas	3.0%		Rock HOH	4.41		194	84.2	484
Evaporated TPH	44.9		Evap HOH	0.6490	11.68	245.5	156	84.2 12235

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	405	405	stoichometric	0.0082	0.0675
	England	250 225	@ 3% Excess Air	0.0099	0.0815
	AP42	152 134	stoichometric	0.0058	0.0477

Standard Flow at measured %HOH	wSCFM	144937	NOx emission model					
	mhoh	29.14%	NOx meas	fuel	eng	ap42		
	mole/lb	lb/lb	ppm	56	102	60		
Dry Flow		801.16	lb/hr	42.0	63.2	76.3	44.6	
Stk HOH	15.677	329.40						
Net HOH 0.1711	3.081	64.73	47.39 evap gpm					
Scrubber HOH	29.420	148						
Excess Air		634.49						
EA OO 0.3517	11.255	133.02	Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA NN 1.3259	37.126	501.48		29.14%	16.60%	3.41%	29.21	3.604
			total	22938	p + rk	24.83%		
					inlet	24.81%		
					meas	29.14%	13.00%	4.00% 29.16 2.958
					ACE	0.00%	0.00%	0.00% 28.00 ERR

Cyclonic Flow @ saturation %HOH	wSCFM	139112	NOx emission model					
	shoh	29.14%	NOx meas	fuel	eng	ap42		
	mole/lb	lb/lb	ppm	56	106	62		
Dry Flow		768.97	lb/hr	40.3	63.2	76.3	44.6	
Stk HOH	15.047	316.16						
Net HOH 0.1361	2.451	51.49	37.70 evap gpm					
Scrubber HOH	30.050	148						
Excess Air		602.30						
EA OO 0.3339	10.684	126.27	Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA NN 1.2587	35.243	476.03		29.14%	16.42%	3.55%	29.22	3.421
			total	22276	sat	29.14%		
					cyc	29.14%		
					p + rk	25.61%		

Individual Velocity Traverse Calculations

No: 8 4F		Usages
Date: 06/08	Rock:	507 TPH
Time: 14:30		0.12 HOH
Pbar: 30.22		446 Dry
Pst: -0.8		
Ps: 30.16 = Pbar+Pst/13.6	Oil:	950 GPH
Cp: 0.834		8.097 lb/gal
Fv: 12.98 = 85.49*Cp*sqrt(1/Ps)		18493 Btu/lb
Dia: 96 Area: 50.27		142 mmBtu/hr
%OO: 13.0%		
%OCO: 4.0%		
Mdry: 29.16		
H: 10.16%		
O: 1.72%		
C: 86.59%	%EA: 2.958 based on fuel analysis and %OCO	

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts ave + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts pt + 460) / (Mdry * (1 - \%HOH pt) + 18 * \%HOH pt)}$
 $cyc = fps pt * \cos(\alpha)$
 $sat \%HOH = eqns from Lange's Handbook of Chemistry, pg 1436$

		Assumed Twb:	156.4				
		sat %HOH:	29.37%				
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:	25.88				
Avg:		16.3	156.4	29.38%			
point	dp	alpha	t	%HOH	std	point	cyc
SE-1	0.60	16	154	27.70%	49.1	48.8	46.9
2	0.80	16	157	29.79%	56.7	56.7	54.5
3	0.80	16	157	29.79%	56.7	56.7	54.5
4	0.62	16	157	29.79%	49.9	50.0	48.0
5	0.95	16	157	29.79%	61.8	61.8	59.4
6	0.95	16	157	29.79%	61.8	61.8	59.4
SW-1	0.60	18	154	27.70%	49.1	48.8	46.4
2	0.80	18	156	29.08%	56.7	56.6	53.8
3	0.78	16	157	29.79%	56.0	56.0	53.9
4	0.65	16	157	29.79%	51.1	51.2	49.2
5	0.98	16	157	29.79%	62.7	62.8	60.4
6	0.90	16	157	29.79%	60.1	60.2	57.9
			avg				
			fps:	56.0	56.0	53.7	
			acfm:	168744	168772	161963	
			%HOH:	29.37%	29.38%		
			scfm:	102905	102909	98758	
			Emission:	lb/hr			
	ppm	Mgas	pVM/RT*60	std	point	cyc	
NOx	56	46	7.1646	41.3	41.3	39.6	

COMPARISON CALCULATIONS

RUN: 4F 06/08/87 time 14:30 ref: 1A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8659	0.0722	OCO	1 0.0722	OCO	0.21	Btu/lb
hydrogen	1	0.1016	0.1016	NO2	1 0.0002	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0025	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0128	0.0004	-fuel O	-0.0005	NN	0.25	RT/p
oxygen	16	0.0172	0.0011	HOH	0.25 0.0254	rock	0.25	378.20
ash		0.0005				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0976	cmb NN	0.3679	10.303	139.16	156	87.8	177
cmb NN	28	0.3679	OCO	44 0.0722	3.175	27.29	156	87.8	46
total		176.07	NO2	46 0.0002	0.008	0.07			
			OSO	64 0.0004	0.026	0.15			
			total dry			166.67			
			HOH	18 0.0508	0.914	19.21	156	87.8	956
			total products						1179

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18493 btu/lb
Fuel GPH	950	0	142 mmbtu/HR
lb/gal	8.097	-100.0%	1269 GPH @190 mmBtu/hr
lb/hr	7692		

SCRUBBER	gpm	Inlet Twb	In hoh sat
	500	152	24.81%

ROCK	%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	507	12.0% in						
Dry TPH	446							
Calc Bin TPH	463	3.6% out						
Meas Bin TPH	485		Rock dried	116.04		194	84.2	3185
(Meas-Calc)/Meas	4.6%		Rock HOH	4.33		194	84.2	476
Evaporated TPH	44.2		Evap HOH	0.6383	11.49	241.4	156	84.2
								12036

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	405	405 stoichometric	0.0082	0.0675
	AP42	250	475 @ 3% Excess Air	0.0099	0.0815
		152	286 stoichometric	0.0058	0.0477

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	145700	29.37%							NOx	56	84	102	59	
Dry Flow					802.70				ppm	42.1	63.2	76.3	44.6	
Stk HOH	15.887				333.82				lb/hr					
Net HOH	0.1935	3.483	73.18	156	88	3643	53.58	evap gpm						
Scrubber HOH	29.017			148	88	1747								
Excess Air			636.03						Stack gas composition					
EA OO	0.3526	11.282	133.34	156	88	178			%HOH	OO	OCO	Mdry	EA	
EA NN	1.3292	37.216	502.69	156	88	638			std	29.37%	16.61%	3.40%	29.21	3.612
					total	23082			p + rk	24.51%				
									inlet	24.81%				
									meas	29.37%	13.00%	4.00%	29.16	2.958
									ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	139846	29.37%							NOx	56	88	106	62	
Dry Flow					770.45				lb/hr	40.4	63.2	76.3	44.6	
Stk HOH	15.249				320.40									
Net HOH	0.1580	2.844	59.77	156	88	2975	43.76	evap gpm						
Scrubber HOH	29.656			148	88	1785								
Excess Air			603.78						Stack gas composition					
EA OO	0.3347	10.710	126.58	156	88	169			%HOH	OO	OCO	Mdry	EA	
EA NN	1.2618	35.329	477.20	156	88	606			sat	29.37%				
					total	22411			cyc	29.37%	16.43%	3.54%	29.22	3.429
									p + rk	25.28%				

Individual Velocity Traverse Calculations

No: 9 1F
 Date: 06/09
 Time: 14:00
 Pbar: 30.15
 Pst: -0.78
 Ps: 30.09 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 13.00 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 13.5%
 %OCO: 4.5%
 Mdry: 29.27
 H: 10.10%
 O: 1.68%
 C: 86.79% ZEA: 2.508 based on fuel analysis and %OO

Usages
 Rock: 489 TPH
 0.12 HOH
 431 Dry
 Oil: 929 GPH
 8.105 lb/gal
 18538 Btu/lb
 140 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		155.3			151							
		sat %HOH:		28.62%			25.79%		28.62%					
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.04			26.36		26.04					
Avg:	point	dp	alpha	t	%HOH	std	point	cyc	inlet	std	std	cyc	meas	meas
SE-1	0.52	18	155	28.45%	45.6	45.5	43.3	45.3	45.6	43.3				
2	0.54	18	155	28.45%	46.4	46.4	44.1	46.1	46.4	44.2				
3	0.56	18	155	28.45%	47.3	47.2	44.9	47.0	47.3	45.0				
4	0.56	16	155	28.45%	47.3	47.2	45.4	47.0	47.3	45.4				
5	0.92	16	155	28.45%	60.6	60.6	58.2	60.2	60.6	58.2				
6	0.94	16	155	28.45%	61.2	61.2	58.8	60.9	61.2	58.9				
SW-1	0.52	18	155	28.45%	45.6	45.5	43.3	45.3	45.6	43.3				
2	0.56	18	155	28.45%	47.3	47.2	44.9	47.0	47.3	45.0				
3	0.56	18	155	28.45%	47.3	47.2	44.9	47.0	47.3	45.0				
4	0.78	16	156	29.14%	55.8	55.9	53.7	55.5	55.8	53.6				
5	0.84	16	156	29.14%	57.9	58.0	55.8	57.5	57.9	55.7				
6	0.96	16	156	29.14%	61.9	62.0	59.6	61.5	61.9	59.5				
avg														
				fps:	52.0	52.0	49.8	51.7	52.0	49.8				
				acfm:	156847	156859	150066	155896	156847	150054				
				%HOH:	28.62%	28.62%		25.79%	28.62%	28.62%				
				scfm:	96634	96639	92454	99852	96634	92449				
Emission: lb/hr														
	ppm	Mgas	pVM/RT*60	std	point	cyc								
NOx	48	46	7.1646	33.2	33.2	31.8	34.3	33.2	31.8					
CO	4.1	28	4.3610	1.7	1.7	1.7	1.8	1.7	1.7					
VOC	0.8	44	6.8531	0.5	0.5	0.5	0.5	0.5	0.5					
SO2	5.8	64	9.9681	5.6	5.6	5.3	5.8	5.6	5.6					
PART	0.014	gr/scf	0.0001	11.6	11.6	11.1	12.0	11.6	11.1					

COMPARISON CALCULATIONS

RUN: 1F 06/09/87 time 14:00 ref: 2A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	fuel	oxygen required	mole/lb fuel	Btu/°F/#		Btu/lb
carbon	12	0.8679	0.0723		OCO	1 0.0723	OCO	0.21	
hydrogen	1	0.1010	0.1010		NO2	1 0.0001	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0014	0.0001		OSO	1 0.0004	OO	0.23	
sulfur	32	0.0128	0.0004		-fuel O	-0.0005	NN	0.25	RT/p
oxygen	16	0.0168	0.0011		HOH	0.25 0.0253	rock	0.25	378.20
ash		0.0004					liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0976	cmb NN	0.3678	10.297	139.09	155	90	168
cmb NN	28	0.3678	OCO	44 0.0723	3.182	27.35	155	90	44
total		175.98	NO2	46 0.0001	0.005	0.04			
			OSO	64 0.0004	0.026	0.15			
			total dry			166.63			
			HOH	18 0.0505	0.909	19.10	155	90	949
			total products						1161

FUEL	Meter	Tank	
Expansion @ 60	95.2%	99.0%	18538 btu/lb
Fuel GPH	929	874	140 mmbtu/HR
lb/gal	8.105	-6.0%	1265 GPH @190 mmBtu/hr
lb/hr	7533		

SCRUBBER	gpm Inlet Twb	In hoh sat
	570 151	24.29%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	489	12.0% in							
Dry TPH	431								
Calc Bin TPH	448	4.0% out							
Meas Bin TPH	445		Rock dried		114.30		179	83	2743
(Meas-Calc)/Meas	-0.8%		Rock HOH		4.76		179	83	457
Evaporated TPH	40.8		Evap HOH	0.6013	10.82	227.4	155	83	11339

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	140 225	227 stoichometric	0.0046	0.0378
	AP42	48 134	365 @ 3% Excess Air	0.0076	0.0626
			182 stoichometric	0.0037	0.0303

Standard Flow at measured %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	135376	28.62%							NOx	48	49	81	39	
Dry Flow					769.69				ppm	33.8	34.7	57.4	27.7	
Stk HOH	14.686				308.58				lb/hr					
Net HOH	0.1641	2.953	62.05	155	90	3084	44.50	evap gpm						
Scrubber HOH	34.879			136	90	1604								
Excess Air			603.05						Stack gas composition					
EA OO	0.3343	10.697	126.43	155	90	161			%HOH	OO	OCO	Mdry	EA	
EA NN	1.2602	35.287	476.63	155	90	576			std	28.62%	16.43%	3.55%	29.23	3.427
						total	21125	p + rk		24.26%				
								inlet		24.29%				
								meas		28.62%	13.53%	4.53%	29.27	2.508
								ACE		0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	129524	28.62%							NOx	48	51	85	41	
Dry Flow					736.41				ppm	32.4	34.7	57.4	27.7	
Stk HOH	14.052				295.24				lb/hr					
Net HOH	0.1288	2.318	48.71	155	90	2421	34.93	evap gpm						
Scrubber HOH	35.514			136	90	1634								
Excess Air			569.78						Stack gas composition					
EA OO	0.3158	10.107	119.45	155	90	152			%HOH	OO	OCO	Mdry	EA	
EA NN	1.1907	33.340	450.33	155	90	544			sat	28.62%				
						total	20450	cyc		28.62%	16.22%	3.71%	29.24	3.238
								p + rk		25.08%				

Individual Velocity Traverse Calculations

No: 9 IR	Usages
Date: 06/09	Rock: 492 TPH
Time: 14:15	0.12 HOH
Pbar: 30.15	433 Dry
Pst: -0.78	
Ps: 30.09 = Pbar+Pst/13.6	Oil: 920 GPH
Cp: 0.834	8.105 lb/gal
Fv: 13.00 = 85.49*Cp*sqrt(1/Ps)	18538 Btu/lb
Dia: 96 Area: 50.27	138 mmBtu/hr
ZOO: 13.5%	
ZOCO: 4.5%	
Mdry: 29.27	
H: 10.10%	
O: 1.68%	
C: 86.79% ZEA: 2.508 based on fuel analysis and ZOO	

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

		Assumed Twb:	155.8			151				
		sat %HOH:	29.03%			25.79%	25.85%			
Ms = Mdry*(1-%HOH)+18*%HOH	Ms:	26.00				26.36	26.35			
Avg:	17.0	155.8	29.03%			std	std	cyc		
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas
SE-1	0.65	18	155	28.45%	51.0	50.9	48.4	50.6	50.7	48.2
2	0.45	18	155	28.45%	42.4	42.4	40.3	42.1	42.1	40.1
3	0.55	18	155	28.45%	46.9	46.8	44.5	46.6	46.6	44.3
4	0.55	16	155	28.45%	46.9	46.8	45.0	46.6	46.6	44.8
5	0.94	16	155	28.45%	61.3	61.2	58.8	60.9	60.9	58.6
6	0.94	16	155	28.45%	61.3	61.2	58.8	60.9	60.9	58.6
SW-1	0.50	18	155	28.45%	44.7	44.6	42.5	44.4	44.4	42.3
2	0.55	18	157	29.86%	46.9	47.0	44.7	46.6	46.6	44.3
3	0.55	18	157	29.86%	46.9	47.0	44.7	46.6	46.6	44.3
4	0.80	16	157	29.86%	56.6	56.7	54.5	56.2	56.2	54.0
5	0.80	16	157	29.86%	56.6	56.7	54.5	56.2	56.2	54.0
6	0.98	16	157	29.86%	62.6	62.8	60.4	62.2	62.2	59.8
			avg							
			fps:	52.0	52.0	49.8	51.7	51.7	49.4	
			acfm:	156899	156914	150120	155811	155830	149083	
			%HOH:	29.03%	29.03%		25.79%	25.85%	25.85%	
			scfm:	96023	96022	91865	99703	99639	95325	
			Emission:	lb/hr						
	ppm	Mgas	pVM/RT*60	std	point	cyc				
NOx	48	46	7.1646	33.0	33.0	31.6	34.3	34.3	32.8	
CO	4.1	28	4.3610	1.7	1.7	1.6	1.8	1.8	1.7	
VOC	0.8	44	6.8531	0.5	0.5	0.5	0.5	0.5	0.5	
SO2	6.6	64	9.9681	6.3	6.3	6.0	6.6	6.5	6.3	
PART	0.014	gr/scf	0.0001	11.6	11.6	11.1	12.0	12.0	11.5	

COMPARISON CALCULATIONS

RUN: 1R 06/09/87 time 14:15 ref: 2A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#				
carbon	12	0.8679	0.0723	OCO	1 0.0723	OCO	0.21		Btu/lb	
hydrogen	1	0.1010	0.1010	NO2	1 0.0001	HOH	0.45	Huv	HOH 1015	
nitrogen	14	0.0014	0.0001	OSO	1 0.0004	OO	0.23			
sulfur	32	0.0128	0.0004	-fuel O	-0.0005	NN	0.25		RT/p	
oxygen	16	0.0168	0.0011	HOH	0.25 0.0253	rock	0.25		378.20	
ash		0.0004				liq HOH	1			
Combustion Air		mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0976	36.89	cmb NN	0.3678	10.297	139.09	156	90	169
cmb NN	28	0.3678	139.09	OCO	44 0.0723	3.182	27.35	156	90	44
total			175.98	NO2	46 0.0001	0.005	0.04			
				OSO	64 0.0004	0.026	0.15			
				total dry			166.63			
				HOH	18 0.0505	0.909	19.10	156	90	950
				total products						1163

FUEL	Meter	Tank	
Expansion @ 60	95.2%	99.0%	18538 btu/lb
Fuel GPH	920	874	138 mmbtu/HR
lb/gal	8.105	-5.0%	1265 GPH @190 mmBtu/hr
lb/hr	7457		

SCRUBBER	gpm Inlet Twb	In hoh sat
	570 151	24.29%

ROCK	%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	492	12.0% in						
Dry TPH	433							
Calc Bin TPH	451	4.0% out						
Meas Bin TPH	445		Rock dried	116.09		179	83	2786
(Meas-Calc)/Meas	-1.3%		Rock HOH	4.84		179	83	464
Evaporated TPH	41.0		Evap HOH	0.6107	10.99	231.0	156	83 11519

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	140 225	227 stoichometric	0.0046	0.0378
	AP42	48 134	365 @ 3% Excess Air	0.0076	0.0626
			182 stoichometric	0.0037	0.0303

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx	meas	fuel	eng	ap42
	134371	25.85%							NOx	meas	fuel	eng	ap42	
Dry Flow					801.74				ppm	48	47	78	38	
Stk HOH					279.47				lb/hr	34.9	34.3	56.8	27.5	
Net HOH	0.0777	1.399	29.39	156	90	1461	20.86	evap gpm						
Scrubber HOH	36.820			136	90	1694								
Excess Air		635.11							Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.3520	11.266	133.15	156	90	171			std	25.85%	16.61%	3.41%	29.21	3.609
EA NN	1.3272	37.162	501.96	156	90	612			p + rk	23.78%				
						19869			inlet	24.29%				
									meas	25.85%	13.53%	4.53%	29.27	2.508
									ACE	29.60%	13.80%	4.60%	29.29	2.456

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx	meas	fuel	eng	ap42
	129448	29.03%							NOx	meas	fuel	eng	ap42	
Dry Flow					739.26				ppm	48	51	85	41	
Stk HOH					302.33				lb/hr	32.2	34.3	56.8	27.5	
Net HOH	0.1382	2.487	52.25	156	90	2598	37.09	evap gpm						
Scrubber HOH	35.732			136	90	1644								
Excess Air		572.63							Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.3174	10.157	120.05	156	90	154			sat	29.03%				
EA NN	1.1967	33.506	452.58	156	90	551			cyc	29.03%	16.24%	3.70%	29.24	3.254
						20879			p + rk	25.28%				

Individual Velocity Traverse Calculations

No: 9 2F
 Date: 06/09
 Time: 15:30
 Pbar: 30.15
 Pst: -0.74
 Ps: 30.10 = Pbar + Pst / 13.6
 Cp: 0.834
 Fv: 13.00 = 85.49 * Cp * sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 13.8%
 %OCO: 4.5%
 Mdry: 29.27
 H: 10.58%
 O: 1.23%
 C: 86.65% %EA: 2.485 based on fuel analysis and %OO

Usages
 Rock: 496 TPH
 0.118 HOH
 438 Dry
 Oil: 937 GPH
 8.097 lb/gal
 18521 Btu/lb
 140 mmBtu/hr

Calculation equation summary

fps: std = $Fv \cdot \sqrt{dp \cdot (Ts_{ave} + 460) / Ms}$
 point = $Fv \cdot \sqrt{dp \cdot (Ts_{pt} + 460) / (Mdry \cdot (1 - \%HOH_{pt}) + 18 \cdot \%HOH_{pt})}$
 cyc = $fps_{pt} \cdot \cos(\alpha)$
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		156.9	150					
		sat %HOH:		29.79%	25.16%	29.79%				
Ms = $Mdry \cdot (1 - \%HOH) + 18 \cdot \%HOH$		Ms:		25.91	26.44	25.91				
Avg:	17.0	156.9	29.79%	std		std		cyc		
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas
SE-1	0.54	18	157	29.85%	46.6	46.6	44.3	46.1	46.6	44.3
2	0.54	16	157	29.85%	46.6	46.6	44.8	46.1	46.6	44.8
3	0.58	16	157	29.85%	48.3	48.3	46.4	47.8	48.3	46.4
4	0.60	18	157	29.85%	49.1	49.1	46.7	48.6	49.1	46.7
5	0.90	18	157	29.85%	60.2	60.2	57.2	59.6	60.2	57.2
6	0.96	16	157	29.85%	62.1	62.1	59.7	61.5	62.1	59.7
SW-1	0.64	16	156	29.14%	50.7	50.6	48.7	50.2	50.7	48.8
2	0.58	18	157	29.85%	48.3	48.3	45.9	47.8	48.3	45.9
3	0.58	18	157	29.85%	48.3	48.3	45.9	47.8	48.3	45.9
4	0.76	18	157	29.85%	55.3	55.3	52.6	54.7	55.3	52.6
5	0.86	16	157	29.85%	58.8	58.8	56.5	58.2	58.8	56.5
6	0.94	16	157	29.85%	61.5	61.5	59.1	60.9	61.5	59.1
avg										
		fps:		53.0	53.0	50.7	52.5	53.0	50.7	
		acfm:		159792	159793	152814	158205	159792	152812	
		%HOH:		29.79%	29.79%		25.16%	29.79%	29.79%	
		scfm:		96575	96575	92357	101928	96575	92357	
Emission: lb/hr										
	ppm	Mgas	pVM/RT*60	std	point	cyc				
NOx	48	46	7.1646	33.2	33.2	31.8	35.1	33.2	31.8	
CO	4.1	28	4.3610	1.7	1.7	1.7	1.8	1.7	1.7	
VOC	0.8	44	6.8531	0.5	0.5	0.5	0.6	0.5	0.5	
SO2	12.0	64	9.9681	11.5	11.5	11.0	12.2	11.5	11.0	
PART	0.017	gr/scf	0.0001	13.8	13.8	13.2	14.5	13.8	13.2	

COMPARISON CALCULATIONS

RUN: 2F 06/09/87 time 15:30 ref: 3A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#	
carbon	12	0.8665	0.0722	OCO	1 0.0722	OCO	0.21 Btu/lb
hydrogen	1	0.1058	0.1058	NO2	1 0.0002	HOH	0.45 Huv HOH 1015
nitrogen	14	0.0028	0.0002	OSO	1 0.0004	OO	0.23
sulfur	32	0.0126	0.0004	-fuel O	-0.0004	NN	0.25 RT/p
oxygen	16	0.0123	0.0008	HOH	0.25 0.0265	rock	0.25 378.20
ash		0.0002				liq HOH	1

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0989	cmb NN	0.3727	10.436	140.97	157	90	175
cmb NN	28	0.3727	OCO	44 0.0722	3.177	27.31	157	90	45
total		178.36	NO2	46 0.0002	0.009	0.08			
			OSO	64 0.0004	0.025	0.15			
			total dry			168.50			
			HOH	18 0.0529	0.952	20.01	157	90	995
			total products						1214

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18521 btu/lb
Fuel GPH	937	927	140 mmbtu/HR
lb/gal	8.097	-1.0%	1267 GPH @190 mmBtu/hr
lb/hr	7585		

SCRUBBER	gpm	Inlet Twb	In hoh sat
	570	150	23.63%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	496	11.8% in							
Dry TPH	438								
Calc Bin TPH	456	4.0% out							
Meas Bin TPH	455		Rock dried		115.46		184	82	2944
(Meas-Calc)/Meas	-0.2%		Rock HOH		4.81		184	82	491
Evaporated TPH	40.3		Evap HOH	0.5909	10.64	223.5	157	82	11154

Nox CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	280	449 stoichometric	0.0092	0.0756
	AP42	189	505 @ 3% Excess Air	0.0107	0.0876
			322 stoichometric	0.0066	0.0542

Standard Flow at measured %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	Nox emission model	meas	fuel	eng	ap42	
	137559	29.79%							NOx	48	99	115	71	
Dry Flow					763.91				ppm	33.8	69.8	80.9	50.0	
Stk HOH			15.429		324.18				lb/hr					
Net HOH	0.2134		3.841		80.70	157	90	4014	evap gpm					
Scrubber HOH			33.730			136	90	1552						
Excess Air					595.40				Stack gas composition					
EA OO	0.3300		10.561		124.82	157	90	163	%HOH	OO	OCO	Mdry	EA	
EA NN	1.2443		34.839		470.58	157	90	583	std	29.79%	16.34%	3.57%	29.23	3.338
								22114	p + rk	24.17%				
									inlet	23.63%				
									meas	29.79%	13.80%	4.50%	29.27	2.485
									ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	Nox emission model	meas	fuel	eng	ap42	
	131552	29.79%							NOx	48	104	120	74	
Dry Flow					730.55				ppm	32.4	69.8	80.9	50.0	
Stk HOH			14.755		310.02				lb/hr					
Net HOH	0.1759		3.167		66.54	157	90	3310	evap gpm					
Scrubber HOH			34.404			136	90	1583						
Excess Air					562.04				Stack gas composition					
EA OO	0.3115		9.970		117.83	157	90	153	%HOH	OO	OCO	Mdry	EA	
EA NN	1.1745		32.887		444.21	157	90	550	sat	29.79%				
								21399	cyc	29.79%	16.13%	3.74%	29.24	3.151
									p + rk	25.00%				

Individual Velocity Traverse Calculations

No: 9 2R	Usages
Date: 06/09	Rock: 499 TPH
Time: 15:40	0.118 HOH
Pbar: 30.15	440 Dry
Pst: -0.74	
Ps: 30.10 =Pbar+Pst/13.6	Oil: 943 GPH
Cp: 0.834	8.097 lb/gal
Fv: 13.00 = 85.49*Cp*sqrt(1/Ps)	18521 Btu/lb
Dia: 96 Area: 50.27	141 mmBtu/hr
%O: 13.8%	
%CO: 4.5%	
Mdry: 29.27	
H: 10.58%	
O: 1.23%	
C: 86.65%	%EA: 2.485 based on fuel analysis and %O

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

		Assumed Twb:	157.0			150				
		sat %HOH:	29.85%			25.16%	26.68%			
Ms = Mdry*(1-%HOH)+18*%HOH	Ms:		25.91			26.44	26.26			
Avg:		17.0	157.0	29.85%				std	std	cyc
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas
SE-1	0.50	18	157	29.85%	44.8	44.8	42.7	44.4	44.5	42.4
2	0.50	16	157	29.85%	44.8	44.8	43.1	44.4	44.5	42.8
3	0.78	16	157	29.85%	56.0	56.0	53.8	55.5	55.6	53.5
4	0.78	18	157	29.85%	56.0	56.0	53.3	55.5	55.6	52.9
5	0.50	18	157	29.85%	44.8	44.8	42.7	44.4	44.5	42.4
6	0.50	16	157	29.85%	44.8	44.8	43.1	44.4	44.5	42.8
SW-1	0.80	16	157	29.85%	56.7	56.7	54.5	56.2	56.3	54.2
2	0.50	18	157	29.85%	44.8	44.8	42.7	44.4	44.5	42.4
3	0.50	18	157	29.85%	44.8	44.8	42.7	44.4	44.5	42.4
4	0.50	18	157	29.85%	44.8	44.8	42.7	44.4	44.5	42.4
5	0.77	16	157	29.85%	55.7	55.7	53.5	55.1	55.3	53.1
6	0.80	16	157	29.85%	56.7	56.7	54.5	56.2	56.3	54.2
				avg						
				fps:	49.6	49.6	47.4	49.1	49.3	47.1
				acfm:	149562	149562	143049	148057	148541	142073
				%HOH:	29.85%	29.85%		25.16%	26.68%	26.68%
				scfm:	90303	90303	86371	95377	93738	89657
				Emission:	lb/hr					
	ppm	Mgas	pVM/RT*60	std	point	cyc				
NOx	52	46	7.1646	33.6	33.6	32.2	35.5	34.9	33.4	
CO	4.2	28	4.3610	1.7	1.7	1.6	1.7	1.7	1.6	
VOC	1	44	6.8531	0.6	0.6	0.6	0.7	0.6	0.6	
SO2	9.5	64	9.9681	8.5	8.5	8.2	9.0	8.9	8.5	
PART	0.017	gr/scf	0.0001	12.9	12.9	12.3	13.6	13.4	12.8	

COMPARISON CALCULATIONS

RUN: 2R 06/09/87 time 15:40 ref: 3A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#	
carbon	12	0.8665	0.0722	OCO	1 0.0722	OCO	0.21
hydrogen	1	0.1058	0.1058	NO2	1 0.0002	HOH	0.45 Huv HOH
nitrogen	14	0.0028	0.0002	OSO	1 0.0004	OO	0.23
sulfur	32	0.0126	0.0004	-fuel O	-0.0004	NN	0.25
oxygen	16	0.0123	0.0008	HOH	0.25 0.0265	rock	0.25
ash		0.0002				liq HOH	1

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0989	cmb NN	0.3727	10.436	140.97	156	90	172
cmb NN	28	0.3727	OCO	44	0.0722	3.177	156	90	44
total		178.36	NO2	46	0.0002	0.009			
			OSO	64	0.0004	0.025			
			total dry			168.50			
			HOH	18	0.0529	0.952	156	90	995
			total products						1210

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18521 btu/lb
Fuel GPH	943	927	141 mmbtu/HR
lb/gal	8.097	-1.7%	1267 GPH @190 mmBtu/hr
lb/hr	7634		

SCRUBBER	gpm Inlet Twb	In hoh sat
	570 150	23.63%

ROCK	%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	499	11.8% in						
Dry TPH	440							
Calc Bin TPH	458	4.0% out						
Meas Bin TPH	455		Rock dried	115.31		184	82	2940
(Meas-Calc)/Meas	-0.7%		Rock HOH	4.80		184	82	490
Evaporated TPH	40.5		Evap HOH	0.5901	10.62	223.2	156	82

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	280 225	449 stoichometric	0.0092	0.0756
	AP42	189 133	505 @ 3% Excess Air	0.0107	0.0876
			322 stoichometric	0.0066	0.0542

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	127857	26.68%							NOx ppm	52	103	119	74	
Dry Flow					736.74				lb/hr	35.6	70.2	81.4	50.3	
Stk HOH	12.762	268.15							evap gpm					
Net HOH	0.0660	1.187	24.95	156	90	1240	18.13							
Scrubber HOH	36.144			136	90	1663								
Excess Air		568.23							Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.3150	10.079	119.13	156	90	153			std	26.68%	16.17%	3.71%	29.24	3.186
EA NN	1.1875	33.249	449.11	156	90	547			p + rk	24.82%				
					total	19379			inlet	23.63%				
									meas	26.68%	13.80%	4.50%	29.27	2.485
									ACE	30.57%	14.00%	4.60%	29.30	2.410

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	123129	29.85%							NOx ppm	52	111	129	80	
Dry Flow					678.83				lb/hr	32.8	70.2	81.4	50.3	
Stk HOH	13.750	288.90							evap gpm					
Net HOH	0.1208	2.175	45.70	156	90	2272	33.21							
Scrubber HOH	35.156			136	90	1617								
Excess Air		510.33							Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.2829	9.052	106.99	156	90	137			sat	29.85%				
EA NN	1.0665	29.861	403.34	156	90	491			cyc	29.85%	15.76%	4.02%	29.27	2.861
					total	20293			p + rk	26.38%				

Individual Velocity Traverse Calculations

No: 9 3F
 Date: 06/09
 Time: 17:10
 Pbar: 30.15
 Pst: -0.78
 Ps: 30.09 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 13.00 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.0%
 %OCO: 4.5%
 Mdry: 29.28
 H: 10.30%
 O: 1.62%
 C: 86.55% %EA: 2.488 based on fuel analysis and %OO

Usages
 Rock: 494 TPH
 0.118 HOH
 435 Dry
 Oil: 914 GPH
 8.097 lb/gal
 18492 Btu/lb
 137 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		153.8			148				
		sat %HOH:		27.65%			23.94%	27.65%			
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.16			26.58	26.16			
Avg:		17.5	153.8	27.65%			std	std	cyc		
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas	
SE-1	0.56	18	154	27.76%	47.1	47.1	44.8	46.7	47.1	44.8	
2	0.58	18	152	26.43%	47.9	47.7	45.4	47.6	47.9	45.6	
3	0.60	16	154	27.76%	48.8	48.8	46.9	48.4	48.8	46.9	
4	0.60	16	154	27.76%	48.8	48.8	46.9	48.4	48.8	46.9	
5	0.88	16	154	27.76%	59.1	59.1	56.8	58.6	59.1	56.8	
6	0.94	18	154	27.76%	61.0	61.1	58.1	60.6	61.0	58.0	
SW-1	0.60	18	154	27.76%	48.8	48.8	46.4	48.4	48.8	46.4	
2	0.60	16	154	27.76%	48.8	48.8	46.9	48.4	48.8	46.9	
3	0.58	20	154	27.76%	47.9	48.0	45.1	47.6	47.9	45.1	
4	0.72	18	154	27.76%	53.4	53.4	50.8	53.0	53.4	50.8	
5	0.84	18	154	27.76%	57.7	57.7	54.9	57.2	57.7	54.9	
6	0.98	18	154	27.76%	62.3	62.3	59.3	61.8	62.3	59.3	
		avg									
		fps:		52.6	52.6	50.2	52.2	52.6	50.2		
		acfm:		158733	158739	151360	157479	158733	151354		
		%HOH:		27.65%	27.65%		23.94%	27.65%	27.65%		
		scfm:		99354	99355	94736	103619	99354	94735		
		Emission:		lb/hr							
	ppm	Mgas	pVM/RT*60	std	point	cyc					
NOx	52	46	7.1646	37.0	37.0	35.3	38.6	37.0	35.3		
CO	4.2	28	4.3610	1.8	1.8	1.7	1.9	1.8	1.7		
VOC	1	44	6.8531	0.7	0.7	0.6	0.7	0.7	0.6		
SO2	6.6	64	9.9681	6.6	6.6	6.3	6.8	6.6	6.3		
PART	0.013	gr/scf	0.0001	10.8	10.8	10.3	11.3	10.8	10.3		

Individual Velocity Traverse Calculations

No: 9 3R
 Date: 06/09
 Time: 17:20
 Pbar: 30.15
 Pst: -0.78
 Ps: 30.09 =Pbar+Pst/13.6
 Cp: 0.834
 Fv: 13.00 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.0%
 %OCO: 4.5%
 Mdry: 29.28
 H: 10.30%
 O: 1.62%
 C: 86.55% %EA: 2.488 based on fuel analysis and %OO

Usages
 Rock: 499 TPH
 0.118 HOH
 440 Dry
 Oil: 923 GPH
 8.097 lb/gal
 18492 Btu/lb
 138 mmBtu/hr

Calculation equation summary

fps: std = $Fv \cdot \sqrt{dp \cdot (Ts_{ave} + 460) / Ms}$
 point = $Fv \cdot \sqrt{dp \cdot (Ts_{pt} + 460) / (Mdry \cdot (1 - \%HOH_{pt}) + 18 \cdot \%HOH_{pt})}$
 cyc = $fps_{pt} \cdot \cos(\alpha)$
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		153.6			148			
		sat %HOH:		27.48%			23.94%	26.33%		
Ms = $Mdry \cdot (1 - \%HOH) + 18 \cdot \%HOH$		Ms:		26.18			26.58	26.31		
Avg:	17.5	153.6	27.49%			std	std	cyc		
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas
SE-1	0.60	18	156	29.14%	48.7	49.0	46.6	48.4	48.6	46.2
2	0.60	18	156	29.14%	48.7	49.0	46.6	48.4	48.6	46.2
3	0.60	16	152	26.43%	48.7	48.6	46.7	48.4	48.6	46.7
4	0.50	16	152	26.43%	44.5	44.3	42.6	44.2	44.4	42.7
5	0.75	16	152	26.43%	54.5	54.3	52.2	54.1	54.4	52.2
6	0.92	18	153	27.09%	60.3	60.3	57.3	59.9	60.2	57.3
SW-1	0.60	18	153	27.09%	48.7	48.7	46.3	48.4	48.6	46.2
2	0.65	16	153	27.09%	50.7	50.7	48.7	50.3	50.6	48.6
3	0.70	20	154	27.76%	52.6	52.7	49.5	52.2	52.5	49.3
4	0.50	18	154	27.76%	44.5	44.5	42.4	44.2	44.4	42.2
5	0.85	18	154	27.76%	58.0	58.1	55.2	57.6	57.9	55.0
6	0.85	18	154	27.76%	58.0	58.1	55.2	57.6	57.9	55.0
				avg						
				fps:	51.5	51.5	49.1	51.1	51.4	49.0
				acfm:	155355	155359	148112	154184	154971	147744
				%HOH:	27.48%	27.49%		23.94%	26.33%	26.33%
				scfm:	97505	97491	92944	101492	98811	94203
				Emission:	lb/hr					
	ppm	Mgas	pVM/RT*60	std	point	cyc				
NOx	53	46	7.1646	37.0	37.0	35.3	38.5	37.5	35.8	
CO	4	28	4.3610	1.7	1.7	1.6	1.8	1.7	1.6	
VOC	0.6	44	6.8531	0.4	0.4	0.4	0.4	0.4	0.4	
SO2	7.6	64	9.9681	7.4	7.4	7.1	7.7	7.5	7.2	
PART	0.013	gr/scf	0.0001	10.6	10.6	10.1	11.1	10.8	10.3	

COMPARISON CALCULATIONS

RUN: 3R 06/09/87 time 17:20 ref: 4A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#	
carbon	12	0.8655	0.0721	OCO	1 0.0721	OCO	0.21
hydrogen	1	0.1030	0.1030	NO2	1 0.0002	HOH	0.45 Huv HOH
nitrogen	14	0.0023	0.0002	OSO	1 0.0004	OO	0.23
sulfur	32	0.0130	0.0004	-fuel O	-0.0005	NN	0.25
oxygen	16	0.0162	0.0010	HOH	0.25 0.0257	rock	0.25
ash		0.0002				liq HOH	1

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0979	cmb NN	0.3692	10.338	139.64	154	78	195
cmb NN	28	0.3692	OCO	44	0.0721	3.173	154	78	50
total		176.69	NO2	46	0.0002	0.008			
			OSO	64	0.0004	0.026			
			total dry			167.14			
			HOH	18	0.0515	0.927	154	78	972
			total products						1218

FUEL	Meter	Tank	
Expansion @ 60	95.2%	99.4%	18492 btu/lb
Fuel GPH	923	927	138 mmbtu/HR
lb/gal	8.097	0.4%	1269 GPH @190 mmBtu/hr
lb/hr	7477		

SCRUBBER	gpm Inlet Twb	In hoh sat
	570 148	22.52%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	499	11.8% in							
Dry TPH	440								
Calc Bin TPH	459	4.0% out							
Meas Bin TPH	378		Rock dried		117.81		175	81	2769
(Meas-Calc)/Meas	-21.3%		Rock HOH		4.91		175	81	461
Evaporated TPH	40.6		Evap HOH	0.6029	10.85	228.0	154	81	11370

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	372 225	372 stoichometric	0.0076	0.0621
	AP42	129 134	455 @ 3% Excess Air	0.0095	0.0783
			262 stoichometric	0.0053	0.0438

Standard Flow at measured %HOH	wSCFM	134120	NOx emission model				
	mhoh	26.33%	NOx meas	fuel	eng	ap42	
	mole/lb	lb/lb	ppm	53	99	55	
Dry Flow		792.90	lb/hr	38.2	71.2	39.9	
Stk HOH	13.485	283.34	evap gpm				
Net HOH	0.0947	1.705					
Scrubber HOH	36.409	35.83					
Excess Air		625.76	Stack gas composition				
EA OO	0.3469	11.100	%HOH	OO	OCO	Mdry	
EA NN	1.3077	36.615	std	26.33%	16.55%	3.44%	
		494.57	total	20531		29.21	
			p + rk	23.79%		EA	
			inlet	22.52%			
			meas	26.33%	14.00%	4.53%	
			ACE	28.19%	14.40%	3.60%	

Cyclonic Flow @ saturation %HOH	wSCFM	128184	NOx emission model				
	shoh	27.48%	NOx meas	fuel	eng	ap42	
	mole/lb	lb/lb	ppm	53	105	59	
Dry Flow		745.94	lb/hr	36.0	71.2	39.9	
Stk HOH	13.453	282.66	evap gpm				
Net HOH	0.0929	1.673					
Scrubber HOH	36.442	35.15					
Excess Air		578.81	Stack gas composition				
EA OO	0.3208	10.267	%HOH	OO	OCO	Mdry	
EA NN	1.2096	33.868	sat	27.48%		EA	
		457.46	cyc	27.48%	16.27%	3.66%	
			total	20432		29.24	
			p + rk	24.91%		3.276	

Individual Velocity Traverse Calculations

No: 9 4F
 Date: 06/09
 Time: 19:20
 Pbar: 30.15
 Pst: -0.78
 Ps: 30.09 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 13.00 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 ZOO: 13.0%
 ZOCO: 4.0%
 Mdry: 29.16
 H: 10.30%
 O: 1.62%
 C: 86.55% ZEA: 2.943 based on fuel analysis and ZOO

Usages
 Rock: 504 TPH
 0.118 HOH
 445 Dry
 Oil: 935 GPH
 8.097 lb/gal
 18492 Btu/lb
 140 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-ZHOH pt)+18*ZHOH pt))
 cyc = fps pt*cos(alpha)
 sat ZHOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		155.0			148			
		sat ZHOH:		28.45%			23.94%	28.45%		
Ms = Mdry*(1-ZHOH)+18*ZHOH		Ms:		25.99			26.49	25.99		
Avg:	dp	alpha	t	ZHOH	std	point	cyc	inlet	meas	cyc
SE-1	0.52	18	154	27.76%	45.6	45.5	43.3	45.2	45.6	43.4
2	0.54	18	155	28.45%	46.5	46.5	44.2	46.0	46.5	44.2
3	0.56	18	155	28.45%	47.3	47.3	45.0	46.9	47.3	45.0
4	0.55	16	155	28.45%	46.9	46.9	45.1	46.4	46.9	45.1
5	0.92	16	155	28.45%	60.6	60.6	58.3	60.1	60.6	58.3
6	0.94	16	155	28.45%	61.3	61.3	58.9	60.7	61.3	58.9
SW-1	0.62	18	154	27.76%	49.8	49.7	47.2	49.3	49.8	47.4
2	0.84	18	156	29.14%	58.0	58.1	55.2	57.4	58.0	55.1
3	0.80	18	156	29.14%	56.6	56.7	53.9	56.0	56.6	53.8
4	0.68	16	155	28.45%	52.1	52.1	50.1	51.6	52.1	50.1
5	0.96	16	155	28.45%	62.0	62.0	59.6	61.4	62.0	59.6
6	0.98	16	155	28.45%	62.6	62.6	60.2	62.0	62.6	60.2
avg										
fps:				54.1	54.1	51.7	53.6	54.1	51.7	
acfm:				163162	163174	156074	161607	163162	156063	
ZHOH:				28.45%	28.45%		23.94%	28.45%	28.45%	
scfm:				100810	100814	96427	106133	100810	96423	
Emission: lb/hr										
	ppm	Mgas	pVM/RT*60	std	point	cyc				
NOx	53	46	7.1646	38.3	38.3	36.6	40.3	38.3	36.6	
CO	4	28	4.3610	1.8	1.8	1.7	1.9	1.8	1.7	
VOC	0.6	44	6.8531	0.4	0.4	0.4	0.4	0.4	0.4	
SO2	8.7	64	9.9681	8.7	8.7	8.3	9.2	8.7	8.3	
PART	0.013	gr/scf	0.0001	11.0	11.0	10.5	11.6	11.0	10.5	

COMPARISON CALCULATIONS

RUN: 4F 06/09/87 time 19:20 ref: 4A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		Btu/lb
carbon	12	0.8655	0.0721	OCO	1 0.0721	OCO	0.21	1015
hydrogen	1	0.1030	0.1030	NO2	1 0.0002	HOH	0.45 Huv	HOH
nitrogen	14	0.0023	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0130	0.0004	-fuel O	-0.0005	NN	0.25	RT/p
oxygen	16	0.0162	0.0010	HOH	0.25 0.0257	rock	0.25	378.20
ash		0.0002				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0979	cmb NN	0.3692	10.338	139.64	155	78	199
cmb NN	28	0.3692	OCO	44 0.0721	3.173	27.28	155	78	51
total		176.69	NO2	46 0.0002	0.008	0.06			
			OSO	64 0.0004	0.026	0.15			
			total dry			167.14			
			HOH	18 0.0515	0.927	19.48	155	78	973
			total products						1223

FUEL	Meter	Tank	
Expansion @ 60	95.2%	99.4%	18492 btu/lb
Fuel GPH	935	927	140 mmbtu/HR
lb/gal	8.097	-0.8%	1269 GPH @190 mmBtu/hr
lb/hr	7567		

SCRUBBER	gpm Inlet	Twb	In hoh sat
	570	148	22.52%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	504	11.8% in							
Dry TPH	445								
Calc Bin TPH	463	4.0% out							
Meas Bin TPH	378		Rock dried		117.54		175	81	2762
(Meas-Calc)/Meas	-22.5%		Rock HOH		4.90		175	81	460
Evaporated TPH	41.0		Evap HOH	0.6015	10.83	227.5	155	81	11351

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	372		372 stoichometric	0.0076	0.0621
	England	230 225	455 @ 3% Excess Air	0.0095	0.0783
	AP42	129 134	262 stoichometric	0.0053	0.0438

Standard Flow at measured %HOH	wSCFM	140884	NOx emission model					
	mhoh	28.45%	NOx meas	fuel	eng	ap42		
	mole/lb	lb/lb	ppm	53	98	55		
Dry Flow		799.30	lb/hr	39.0	72.1	40.3		
Stk HOH	15.123	317.74						
Net HOH	0.1871	3.368 70.76	50.97	evap gpm				
Scrubber HOH	34.293	134						
Excess Air		632.16						
EA OO	0.3504	11.213 132.53						
EA NN	1.3211	36.990 499.63						
			Stack gas composition					
			%HOH	OO	OCO	Mdry	EA	
			std	28.45%	16.58%	3.41%	29.21	3.578
			p + rk	23.61%				
			inlet	22.52%				
			meas	28.45%	13.00%	4.00%	29.16	2.943
			ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	134764	NOx emission model					
	shoh	28.45%	NOx meas	fuel	eng	ap42		
	mole/lb	lb/lb	ppm	53	102	57		
Dry Flow		764.58	lb/hr	37.3	72.1	40.3		
Stk HOH	14.466	303.94						
Net HOH	0.1506	2.711 56.96	41.03	evap gpm				
Scrubber HOH	34.949	134						
Excess Air		597.44						
EA OO	0.3312	10.597 125.25						
EA NN	1.2485	34.958 472.19						
			Stack gas composition					
			%HOH	OO	OCO	Mdry	EA	
			sat	28.45%				
			cyc	28.45%	16.38%	3.57%	29.23	3.381
			p + rk	24.42%				

Individual Velocity Traverse Calculations

No: X 1F
 Date: 06/10
 Time: 10:45
 Pbar: 30.14
 Pst: -0.42
 Ps: 30.11 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 12.99 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.54%
 O: 1.26%
 C: 86.79% %EA: 2.490 based on fuel analysis and %OO

Usages
 Rock: 492 TPH
 0.112 HOH
 436 Dry
 Oil: 946 GPH
 8.088 lb/gal
 18497 Btu/lb
 142 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:					150				
		sat %HOH:					25.15%		28.43%		
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:					26.46		26.09		
Avg:	point	dp	alpha	t	%HOH	std	point	cyc	std	std	cyc
									inlet	meas	meas
SE-1	0.45	18	154	27.75%	42.3	42.2	40.2	42.0	42.3	40.3	
2	0.52	18	155	28.43%	45.5	45.5	43.3	45.2	45.5	43.3	
3	0.48	16	155	28.43%	43.7	43.7	42.0	43.4	43.7	42.0	
4	0.50	16	155	28.43%	44.6	44.6	42.9	44.3	44.6	42.9	
5	0.78	8	155	28.43%	55.7	55.7	55.2	55.3	55.7	55.2	
6	0.48	8	155	28.43%	43.7	43.7	43.3	43.4	43.7	43.3	
SW-1	0.52	16	155	28.43%	45.5	45.5	43.7	45.2	45.5	43.7	
2	0.44	18	155	28.43%	41.8	41.8	39.8	41.6	41.8	39.8	
3	0.48	18	155	28.43%	43.7	43.7	41.6	43.4	43.7	41.6	
4	0.48	14	155	28.43%	43.7	43.7	42.4	43.4	43.7	42.4	
5	0.52	14	155	28.43%	45.5	45.5	44.1	45.2	45.5	44.1	
6	0.58	16	156	29.13%	48.0	48.2	46.3	47.7	48.0	46.2	
avg											
					fps:	45.3	45.3	43.7	45.0	45.3	43.7
					acfm:	136689	136692	131880	135727	136689	131876
					%HOH:	28.43%	28.43%		25.15%	28.43%	28.43%
					scfm:	84518	84518	81543	87771	84518	81542
Emission: lb/hr											
	ppm	Mgas	pVM/RT*60		std	point	cyc				
NOx	51	46	7.1646		30.9	30.9	29.8	32.1	30.9	29.8	
CO	2	28	4.3610		0.7	0.7	0.7	0.8	0.7	0.7	
VOC	0.7	44	6.8531		0.4	0.4	0.4	0.4	0.4	0.4	
SO2	9.4	64	9.9681		7.9	7.9	7.6	8.2	7.9	7.6	
PART	0.015	gr/scf	0.0001		10.7	10.7	10.3	11.1	10.7	10.3	

COMPARISON CALCULATIONS

RUN: 1F 06/10/87 time 10:45 ref: 5A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8679	0.0723	OCO	1 0.0723	OCO	0.21	Btu/lb
hydrogen	1	0.1054	0.1054	NO2	1 0.0001	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0015	0.0001	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0126	0.0004	-fuel O	-0.0004	NN	0.25	RT/p
oxygen	16	0.0126	0.0008	HOH	0.25 0.0264	rock	0.25	378.20
ash		0.0002				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0988	cmb NN	0.3724	10.427	140.85	155	87	177
cmb NN	28	0.3724	OCO	44	0.0723	3.182	155	87	45
total		178.21	NO2	46	0.0001	0.005			
			OSO	64	0.0004	0.025			
			total dry			168.39			
			HOH	18	0.0527	0.949	155	87	992
			total products						1215

FUEL	Meter	Tank	
Expansion @ 60	95.3%	99.1%	18497 btu/lb
Fuel GPH	946	867	142 mmbtu/HR
lb/gal	8.088	-8.4%	1270 GPH @190 mmBtu/hr
lb/hr	7653		

SCRUBBER	gpm	Inlet Twb	In hoh sat
	580	150	23.49%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	492	11.2%	in						
Dry TPH	436								
Calc Bin TPH	452	3.4%	out						
Meas Bin TPH	465			Rock dried	114.07		179	78	2880
(Meas-Calc)/Meas	2.9%			Rock HOH	4.01		179	78	406
Evaporated TPH	39.7			Evap HOH	0.5762	10.37	155	78	10887

NOX CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	150	225	241 stoichometric	0.0049
	AP42	54	133	375 @ 3% Excess Air	0.0079
				187 stoichometric	0.0038

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	118090	28.43%							NOx	51	61	98	48	
Dry Flow					662.63				ppm	31.5	37.7	60.5	29.3	
Stk HOH	12.527				263.21				lb/hr					
Net HOH	0.0670	1.206	25.34	155	87	1261	18.46	evap gpm						
Scrubber HOH	36.686			132	87	1651								
Excess Air		494.24							Stack gas composition					
EA OO	0.2740	8.767	103.61	155	87	137			%HOH	OO	OCO	Mdry	EA	
EA NN	1.0328	28.920	390.63	155	87	492			std	28.43%	15.64%	4.13%	29.29	2.773
					total	18929			p + rk	26.42%				
									inlet	23.49%				
									meas	28.43%	14.50%	4.50%	29.30	2.490
									ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	113936	28.43%							NOx	51	63	102	49	
Dry Flow					639.32				ppm	30.3	37.7	60.5	29.3	
Stk HOH	12.087				253.95				lb/hr					
Net HOH	0.0425	0.766	16.08	155	87	800	11.72	evap gpm						
Scrubber HOH	37.127			132	87	1671								
Excess Air		470.93							Stack gas composition					
EA OO	0.2610	8.353	98.73	155	87	131			%HOH	OO	OCO	Mdry	EA	
EA NN	0.9841	27.556	372.20	155	87	468			sat	28.43%				
					total	18458			cyc	28.43%	15.44%	4.28%	29.30	2.643
									p + rk	27.12%				

Individual Velocity Traverse Calculations

No: X 1R	Usages
Date: 06/10	Rock: 507 TPH
Time: 11:10	0.112 HOH
Pbar: 30.14	450 Dry
Pst: -0.42	
Ps: 30.11 = Pbar + Pst / 13.6	Oil: 940 GPH
Cp: 0.834	8.088 lb/gal
Fv: 12.99 = 85.49 * Cp * sqrt(1/Ps)	18497 Btu/lb
Dia: 96 Area: 50.27	141 mmBtu/hr
%O: 14.5%	
%CO: 4.5%	
Mdry: 29.30	
H: 10.54%	
O: 1.26%	
C: 86.79%	%EA: 2.490 based on fuel analysis and %O

Calculation equation summary

$fps: std = Fv * \sqrt{dp * (Ts_{ave} + 460) / Ms}$
 $point = Fv * \sqrt{dp * (Ts_{pt} + 460) / (Mdry * (1 - \%HOH_{pt}) + 18 * \%HOH_{pt})}$
 $cyc = fps_{pt} * \cos(\alpha)$
 $sat \%HOH = eqns \text{ from Lange's Handbook of Chemistry, pg 1436}$

		Assumed Twb:					150				
		sat %HOH:					25.15% 24.15%				
Ms = Mdry * (1 - %HOH) + 18 * %HOH		Ms:					26.46 26.57				
Avg:	dp	alpha	t	%HOH	std	point	cyc	inlet	std	std	cyc
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas	meas
SE-1	0.50	18	156	29.13%	44.7	44.7	42.5	44.3	44.2	42.1	
2	0.40	18	156	29.13%	40.0	40.0	38.0	39.6	39.6	37.6	
3	0.45	16	156	29.13%	42.4	42.4	40.8	42.1	42.0	40.3	
4	0.45	16	156	29.13%	42.4	42.4	40.8	42.1	42.0	40.3	
5	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.5	
6	0.60	18	156	29.13%	49.0	49.0	46.6	48.6	48.5	46.1	
SW-1	0.44	18	155	28.43%	41.9	41.8	39.8	41.6	41.5	39.5	
2	0.44	16	156	29.13%	41.9	41.9	40.3	41.6	41.5	39.9	
3	0.50	20	156	29.13%	44.7	44.7	42.0	44.3	44.2	41.6	
4	0.50	18	156	29.13%	44.7	44.7	42.5	44.3	44.2	42.1	
5	0.50	18	156	29.13%	44.7	44.7	42.5	44.3	44.2	42.1	
6	0.50	18	156	29.13%	44.7	44.7	42.5	44.3	44.2	42.1	
avg											
					fps:	43.8	43.8	41.8	43.5	43.4	41.3
					acfm:	132171	132173	126016	131060	130781	124689
					%HOH:	29.07%	29.07%		25.15%	24.15%	24.15%
					scfm:	80874	80874	77107	84627	85575	81588
Emission: lb/hr											
		ppm	Mgas	pVM/RT*60	std	point	cyc				
NOx	51	46	7.1646		29.6	29.6	28.2	30.9	31.3	29.8	
CO	2	28	4.3610		0.7	0.7	0.7	0.7	0.7	0.7	
VOC	0.7	44	6.8531		0.4	0.4	0.4	0.4	0.4	0.4	
SO2	9.4	64	9.9681		7.5	7.5	7.2	7.9	8.0	7.6	
PART	0.015	gr/scf	0.0001		10.2	10.2	9.7	10.7	10.8	10.3	

COMPARISON CALCULATIONS

RUN: 1R 06/10/87 time 11:10 ref: 5A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8679	0.0723	OCO	1 0.0723	OCO	0.21	Btu/lb
hydrogen	1	0.1054	0.1054	NO2	1 0.0001	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0015	0.0001	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0126	0.0004	-fuel O	-0.0004	NN	0.25	RT/p
oxygen	16	0.0126	0.0008	HOH	0.25 0.0264	rock	0.25	378.20
ash		0.0002				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0988	cmb NN	0.3724	10.427	140.85	156	87	180
cmb NN	28	0.3724	OCO	44 0.0723	3.182	27.35	156	87	46
total		178.21	NO2	46 0.0001	0.005	0.04			
			OSO	64 0.0004	0.025	0.15			
			total dry			168.39			
			HOH	18 0.0527	0.949	19.93	156	87	992
			total products						1218

FUEL	Meter	Tank	
Expansion @ 60	95.3%	99.1%	18497 btu/lb
Fuel GPH	940	867	141 mmbtu/HR
lb/gal	8.088	-7.8%	1270 GPH @190 mmBtu/hr
lb/hr	7603		

SCRUBBER	gpm Inlet Twb	In hoh sat
	580 150	23.49%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	507	11.2% in							
Dry TPH	450								
Calc Bin TPH	466	3.4% out							
Meas Bin TPH	465		Rock dried		118.33		179	78	2988
(Meas-Calc)/Meas	-0.1%		Rock HOH		4.16		179	78	421
Evaporated TPH	40.9		Evap HOH	0.5978	10.76	226.1	156	78	11299

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	150 225	241 stoichometric	0.0049	0.0405
	AP42	54 133	375 @ 3% Excess Air	0.0079	0.0650
			187 stoichometric	0.0038	0.0315

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	112818	24.15%							NOx ppm	51	60	96	47	
Dry Flow					675.31				lb/hr	31.8	37.5	60.1	29.1	
Stk HOH	10.232				214.99									
Net HOH	-0.0820				-31.02	156	87	-1544	evap gpm					
Scrubber HOH	39.617					132	87	1783						
Excess Air					506.92				Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.2810				106.27	156	87	143	std	24.15%	15.74%	4.05%	29.28	2.845
EA NN	1.0593				400.65	156	87	511	p + rk	26.70%				
						total	16817		inlet	23.49%				
									meas	24.15%	14.50%	4.50%	29.30	2.490
									ACE	28.81%	14.60%	4.30%	29.27	2.649

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	108707	29.07%							NOx ppm	51	67	107	52	
Dry Flow					608.49				lb/hr	28.7	37.5	60.1	29.1	
Stk HOH					249.37									
Net HOH	0.0089				3.36	156	87	167	evap gpm					
Scrubber HOH	37.981					132	87	1709						
Excess Air					440.10				Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.2440				92.26	156	87	124	sat	29.07%				
EA NN	0.9197				347.84	156	87	444	cyc	29.07%	15.16%	4.50%	29.33	2.470
						total	18369		p + rk	28.79%				

Individual Velocity Traverse Calculations

No: X 2F
 Date: 06/10
 Time: 12:30
 Pbar: 30.14
 Pst: -0.4
 Ps: 30.11 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 12.99 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.52%
 O: 1.33%
 C: 86.67% %EA: 2.491 based on fuel analysis and %OO

Usages
 Rock: 511 TPH
 0.112 HOH
 454 Dry
 Oil: 938 GPH
 8.072 lb/gal
 18545 Btu/lb
 140 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		154.9			150				
		sat %HOH:		28.37%			25.15%	28.37%			
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.09			26.46	26.09			
Avg:		15.7	154.9	28.37%			std	std		cyc	
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas	
SE-1	0.46	16	155	28.43%	42.8	42.8	41.1	42.5	42.8	41.1	
2	0.50	18	155	28.43%	44.6	44.6	42.4	44.3	44.6	42.4	
3	0.50	16	155	28.43%	44.6	44.6	42.9	44.3	44.6	42.9	
4	0.50	16	155	28.43%	44.6	44.6	42.9	44.3	44.6	42.9	
5	0.48	10	155	28.43%	43.7	43.7	43.0	43.4	43.7	43.0	
6	0.50	16	155	28.43%	44.6	44.6	42.9	44.3	44.6	42.9	
SW-1	0.50	18	154	27.74%	44.6	44.5	42.3	44.3	44.6	42.4	
2	0.46	16	155	28.43%	42.8	42.8	41.1	42.5	42.8	41.1	
3	0.48	16	155	28.43%	43.7	43.7	42.0	43.4	43.7	42.0	
4	0.48	16	155	28.43%	43.7	43.7	42.0	43.4	43.7	42.0	
5	0.50	16	155	28.43%	44.6	44.6	42.9	44.3	44.6	42.9	
6	0.56	14	155	28.43%	47.2	47.2	45.8	46.9	47.2	45.8	
		avg									
		fps:		44.3	44.3	42.6	44.0	44.3	42.6		
		acfm:		133572	133572	128535	132649	133572	128534		
		%HOH:		28.37%	28.37%		25.15%	28.37%	28.37%		
		scfm:		82674	82673	79555	85798	82674	79556		
		Emission:		lb/hr							
	ppm	Mgas	pVM/RT*60	std	point	cyc					
NOx	50	46	7.1646	29.6	29.6	28.5	30.7	29.6	28.5		
CO	2	28	4.3610	0.7	0.7	0.7	0.7	0.7	0.7		
VOC	1	44	6.8531	0.6	0.6	0.5	0.6	0.6	0.5		
SO2	10.6	64	9.9681	8.8	8.8	8.4	9.1	8.8	8.4		
PART	0.016	gr/scf	0.0001	11.5	11.5	11.1	12.0	11.5	11.1		

COMPARISON CALCULATIONS

RUN: 2F 06/10/87 time 12:30 ref: 6A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#	
carbon	12	0.8667	0.0722	OCO	1 0.0722	OCO	0.21
hydrogen	1	0.1052	0.1052	NO2	1 0.0002	HOH	0.45 Huv HOH
nitrogen	14	0.0023	0.0002	OSO	1 0.0004	OO	0.23
sulfur	32	0.0125	0.0004	-fuel O	-0.0004	NN	0.25
oxygen	16	0.0133	0.0008	HOH	0.25 0.0263	rock	0.25
ash		0.0005				liq HOH	1

Btu/lb
1015
RT/p
378.20

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0987	cmb NN	0.3720	10.415	140.68	155	90	169
cmb NN	28	0.3720	OCO	44 0.0722	3.178	27.32	155	90	43
total		177.99	NO2	46 0.0002	0.008	0.06			
			OSO	64 0.0004	0.025	0.15			
			total dry			168.20			
			HOH	18 0.0526	0.947	19.89	155	90	989
			total products						1201

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18545 btu/lb
Fuel GPH	938	845	140 mmbtu/HR
lb/gal	8.072	-10.0%	1269 GPH @190 mmbtu/hr
lb/hr	7575		

SCRUBBER	gpm	Inlet Twb	In hoh sat
	580	150	23.49%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	511	11.2% in							
Dry TPH	454								
Calc Bin TPH	468	3.0% out							
Meas Bin TPH	442			Rock dried	119.75		184	79	3143
(Meas-Calc)/Meas	-5.9%			Rock HOH	3.70		184	79	389
Evaporated TPH	43.2			Evap HOH	0.6333 11.40	239.5	155	79	11961

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	369 225	369 stoichometric	0.0076	0.0621
	AP42	128 133	455 @ 3% Excess Air	0.0096	0.0788
			261 stoichometric	0.0053	0.0440

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	115419	28.37%							NOx meas	50	95	120	67	
Dry Flow					654.84				lb/hr	30.2	57.2	72.6	40.5	
Stk HOH	12.344	259.36							evap gpm					
Net HOH	-0.0002	-0.003	-0.06	155	90	-3	-0.04							
Scrubber HOH	38.285	134	90	1685										
Excess Air			486.63						Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.2697	8.632	102.02	155	90	129			std	28.37%	15.58%	4.17%	29.29	2.734
EA NN	1.0169	28.474	384.61	155	90	462			p + rk	28.38%				
					total	18967			inlet	23.49%				
									meas	28.37%	14.50%	4.50%	29.30	2.491
									ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	NOx meas	fuel	eng	ap42	
	111066	28.37%							NOx meas	50	99	125	70	
Dry Flow					630.14				lb/hr	29.0	57.2	72.6	40.5	
Stk HOH	11.879	249.58							evap gpm					
Net HOH	-0.0260	-0.468	-9.84	155	90	-489	-7.10							
Scrubber HOH	38.751	134	90	1705										
Excess Air			461.94						Stack gas composition	%HOH	OO	OCO	Mdry	EA
EA OO	0.2561	8.194	96.84	155	90	122			sat	28.37%				
EA NN	0.9653	27.030	365.10	155	90	439			cyc	28.37%	15.37%	4.33%	29.31	2.595
					total	18471			p + rk	29.16%				

Individual Velocity Traverse Calculations

No: X 2R
 Date: 06/10
 Time: 12:30
 Pbar: 30.14
 Pst: -0.4
 Ps: 30.11 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 12.99 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.52%
 O: 1.33%
 C: 86.67% %EA: 2.491 based on fuel analysis and %OO

Usages
 Rock: 506 TPH
 0.112 HOH
 450 Dry
 Oil: 919 GPH
 8.072 lb/gal
 18545 Btu/lb
 138 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460))/Ms
 point = Fv*sqrt(dp*(Ts pt+460))/(Mdry*(1-%HOH pt)+18*%HOH pt)
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		155.8			150				
		sat %HOH:		29.01%			25.15%	23.38%			
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.02			26.46	26.66			
Avg:	point	dp	alpha	t	%HOH	std	point	cyc	std	std	cyc
SE-1	0.40	16	156	29.13%	40.0	40.0	38.4	39.6	39.5	38.0	
2	0.50	18	156	29.13%	44.7	44.7	42.5	44.3	44.2	42.0	
3	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.4	
4	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.4	
5	0.50	10	156	29.13%	44.7	44.7	44.0	44.3	44.2	43.5	
6	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.4	
SW-1	0.45	18	154	27.74%	42.4	42.2	40.2	42.1	41.9	39.8	
2	0.45	16	156	29.13%	42.4	42.4	40.8	42.1	41.9	40.3	
3	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.4	
4	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.4	
5	0.50	16	156	29.13%	44.7	44.7	43.0	44.3	44.2	42.4	
6	0.50	14	156	29.13%	44.7	44.7	43.4	44.3	44.2	42.8	
avg											
fps:				43.9	43.9	42.3	43.6	43.4	41.8		
acfm:				132461	132463	127475	131364	130870	125941		
%HOH:				29.01%	29.01%		25.15%	23.38%	23.38%		
scfm:				81135	81134	78078	84840	86520	83262		
Emission:		lb/hr									
	ppm	Mgas	pVM/RT*60	std	point	cyc					
NOx	50	46	7.1646	29.1	29.1	28.0	30.4	31.0	29.8		
CO	2	28	4.3610	0.7	0.7	0.7	0.7	0.8	0.7		
VOC	1	44	6.8531	0.6	0.6	0.5	0.6	0.6	0.6		
SO2	10.6	64	9.9681	8.6	8.6	8.3	9.0	9.2	8.8		
PART	0.016	gr/scf	0.0001	11.3	11.3	10.9	11.8	12.1	11.6		

COMPARISON CALCULATIONS

RUN: 2R 06/10/87 time 12:30 ref: 6A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8667	0.0722	OCO	1 0.0722	OCO	0.21	Btu/lb
hydrogen	1	0.1052	0.1052	NO2	1 0.0002	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0023	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0125	0.0004	-fuel 0	-0.0004	NN	0.25	RT/p
oxygen	16	0.0133	0.0008	HOH	0.25 0.0263	rock	0.25	378.20
ash		0.0005				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0987	cmb NN	0.3720	10.415	140.68	156	90	171
cmb NN	28	0.3720	OCO	44	0.0722	3.178	156	90	44
total		177.99	NO2	46	0.0002	0.008			
			OSO	64	0.0004	0.025			
			total dry			168.20			
			HOH	18	0.0526	0.947	156	90	989
			total products						1204

FUEL	Meter	Tank	
Expansion @ 60	95.4%	99.0%	18545 btu/lb
Fuel GPH	919	845	138 mmbtu/HR
lb/gal	8.072	-8.1%	1269 GPH @190 mmBtu/hr
lb/hr	7422		

SCRUBBER	gpm Inlet Twb	In hoh sat
	580 150	23.49%

ROCK		%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	506	11.2% in							
Dry TPH	450								
Calc Bin TPH	464	3.0% out							
Meas Bin TPH	442		Rock dried		121.20		184	79	3182
(Meas-Calc)/Meas	-5.0%		Rock HOH		3.75		184	79	394
Evaporated TPH	42.8		Evap HOH	0.6410	11.54	242.4	156	79	12110

NOx CALCULATION	Fuel	fuel thermal	ppm condition	lb/lb	scf/lb
	England	369 230 225	369 stoichometric	0.0076	0.0621
	AP42	128 133	455 @ 3% Excess Air	0.0096	0.0788
			261 stoichometric	0.0053	0.0440

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	112916	23.38%							NOx	50	89	113	63	
Dry Flow					699.45				ppm	31.6	56.1	71.2	39.7	
Stk HOH	10.156	213.39							lb/hr					
Net HOH -0.1294	-2.329	-48.94	156	90	-2433	-34.57	evap gpm							
Scrubber HOH	41.402		134	90	1822									
Excess Air		531.25							Stack gas composition					
EA OO 0.2945	9.423	111.37	156	90	143				%HOH	OO	OCO	Mdry	EA	
EA NN 1.1102	31.085	419.88	156	90	512				std	23.38%	15.92%	3.91%	29.26	2.985
					total 16932				p + rk	27.27%				
									inlet	23.49%				
									meas	23.38%	14.50%	4.50%	29.30	2.491
									ACE	29.37%	14.60%	4.30%	29.27	2.651

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	109987	29.01%							NOx	50	98	125	70	
Dry Flow					631.22				ppm	28.5	56.1	71.2	39.7	
Stk HOH	12.276	257.93							lb/hr					
Net HOH -0.0116	-0.209	-4.39	156	90	-218	-3.10	evap gpm							
Scrubber HOH	39.281		134	90	1728									
Excess Air		463.02							Stack gas composition					
EA OO 0.2567	8.213	97.07	156	90	124				%HOH	OO	OCO	Mdry	EA	
EA NN 0.9676	27.093	365.95	156	90	446				sat	29.01%				
					total 18970				cyc	29.01%	15.38%	4.33%	29.31	2.601
									p + rk	29.36%				

Individual Velocity Traverse Calculations

No: X 3F
 Date: 06/10
 Time: 13:45
 Pbar: 30.14
 Pst: -0.4
 Ps: 30.11 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 12.99 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.47%
 O: 1.09%
 C: 86.90% ZEA: 2.495 based on fuel analysis and %OO

Usages
 Rock: 515 TPH
 0.11 HOH
 458 Dry
 Oil: 777 GPH
 8.105 lb/gal
 18509 Btu/lb
 117 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		149.0			150				
		sat %HOH:		24.53%			25.15%	24.53%			
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.53			26.46	26.53			
Avg:		17.0	149.0	24.53%			std	std	cyc		
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas	
SE-1	0.45	16	149	24.53%	41.8	41.8	40.1	41.8	41.8	40.1	
2	0.48	16	149	24.53%	43.1	43.1	41.5	43.2	43.1	41.5	
3	0.50	18	149	24.53%	44.0	44.0	41.9	44.1	44.0	41.9	
4	0.50	18	149	24.53%	44.0	44.0	41.9	44.1	44.0	41.9	
5	0.50	18	149	24.53%	44.0	44.0	41.9	44.1	44.0	41.9	
6	0.48	16	149	24.53%	43.1	43.1	41.5	43.2	43.1	41.5	
SW-1	0.48	16	149	24.53%	43.1	43.1	41.5	43.2	43.1	41.5	
2	0.46	14	149	24.53%	42.2	42.2	41.0	42.3	42.2	41.0	
3	0.48	18	149	24.53%	43.1	43.1	41.0	43.2	43.1	41.0	
4	0.48	18	149	24.53%	43.1	43.1	41.0	43.2	43.1	41.0	
5	0.50	18	149	24.53%	44.0	44.0	41.9	44.1	44.0	41.9	
6	0.56	18	149	24.53%	46.6	46.6	44.3	46.6	46.6	44.3	
		avg									
		fps:		43.5	43.5	41.6	43.6	43.5	41.6		
		acfm:		131273	131273	125491	131445	131273	125491		
		%HOH:		24.53%	24.53%		25.15%	24.53%	24.53%		
		scfm:		86438	86438	82631	85845	86438	82631		
		Emission:		lb/hr							
	ppm	Mgas	pVM/RT*60	std	point	cyc					
NOx	48	46	7.1646	29.7	29.7	28.4	29.5	29.7	28.4		
CO	3.25	28	4.3610	1.2	1.2	1.2	1.2	1.2	1.2		
VOC	0.4	44	6.8531	0.2	0.2	0.2	0.2	0.2	0.2		
SO2	5.7	64	9.9681	4.9	4.9	4.7	4.9	4.9	4.7		
PART	0.020	gr/scf	0.0002	14.9	14.9	14.2	14.8	14.9	14.2		

COMPARISON CALCULATIONS

RUN: 3F 06/10/87 time 13:45 ref: 7A 6

COMBUSTION CALCULATIONS

	lb/mole	lb/lb	mole/lb fuel	oxygen required	mole/lb fuel	Btu/°F/#		
carbon	12	0.8690	0.0724	OCO	1 0.0724	OCO	0.21	Btu/lb
hydrogen	1	0.1047	0.1047	NO2	1 0.0002	HOH	0.45	Huv HOH 1015
nitrogen	14	0.0030	0.0002	OSO	1 0.0004	OO	0.23	
sulfur	32	0.0124	0.0004	-fuel O	-0.0003	NN	0.25	RT/p
oxygen	16	0.0109	0.0007	HOH	0.25 0.0262	rock	0.25	378.20
ash		0.0006				liq HOH	1	

Combustion Air	mole/lb	scf/lb	Comb Products	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
net OO	32	0.0989	cmb NN	0.3727	10.435	140.95	149	89	157
cmb NN	28	0.3727	OCO	44 0.0724	3.186	27.39	149	89	40
total		178.33	NO2	46 0.0002	0.010	0.08			
			OSO	64 0.0004	0.025	0.15			
			total dry			168.56			
			HOH	18 0.0524	0.942	19.80	149	89	982
			total products						1179

FUEL	Meter	Tank	
Expansion @ 60	95.5%	99.0%	18509 btu/lb
Fuel GPH	777	756	117 mmBtu/HR
lb/gal	8.105	-2.8%	1267 GPH @190 mmBtu/hr
lb/hr	6300		

SCRUBBER	gpm Inlet Twb	In hoh sat
	580 150	23.49%

ROCK	%HOH		mole/lb	lb/lb	scf/lb	tout	tin	btu/lb
Scale TPH	515	11.0% in						
Dry TPH	458							
Calc Bin TPH	478	4.2% out						
Meas Bin TPH	1022		Rock dried	145.42		179	79	3635
(Meas-Calc)/Meas	53.2%		Rock HOH	6.38		179	79	638
Evaporated TPH	36.5		Evap HOH	0.6443	11.60	243.7	149	79 12137

NOx CALCULATION	Fuel	thermal	ppm condition	lb/lb	scf/lb
	481		481 stoichometric	0.0099	0.0810
	England	300 225	525 @ 3% Excess Air	0.0111	0.0912
	AP42	217 132	349 stoichometric	0.0072	0.0588

Standard Flow at measured %HOH	wSCFM	mhoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	114534	24.53%							NOx	48	98	111	71	
Dry Flow					823.24				ppm	30.3	62.1	69.8	45.1	
Stk HOH			12.735	267.58					lb/hr					
Net HOH	0.0108	0.195	4.10	149	89	203	2.46	evap gpm						
Scrubber HOH	45.836			130	89	1879								
Excess Air		654.68							Stack gas composition					
EA OO	0.3629	11.613	137.25	149	89	160			%HOH	OO	OCO	Mdry	EA	
EA NN	1.3681	38.307	517.43	149	89	575			std	24.53%	16.67%	3.33%	29.20	3.671
						20406			p + rk	24.25%				
									inlet	23.49%				
									meas	24.53%	14.50%	4.50%	29.30	2.495
									ACE	0.00%	0.00%	0.00%	28.00	ERR

Cyclonic Flow @ saturation %HOH	wSCFM	shoh	mole/lb	lb/lb	scf/lb	tout	tin	btu/lb	NOx emission model	meas	fuel	eng	ap42	
	109490	24.53%							NOx	48	103	116	75	
Dry Flow					786.98				ppm	28.9	62.1	69.8	45.1	
Stk HOH			12.174	255.80					lb/hr					
Net HOH	-0.0203	-0.366	-7.68	149	89	-381	-4.61	evap gpm						
Scrubber HOH	46.397			130	89	1902								
Excess Air		618.42							Stack gas composition					
EA OO	0.3428	10.970	129.65	149	89	151			%HOH	OO	OCO	Mdry	EA	
EA NN	1.2923	36.186	488.77	149	89	543			sat	24.53%				
						19804			cyc	24.53%	16.47%	3.48%	29.22	3.468
									p + rk	25.08%				

Individual Velocity Traverse Calculations

No: X 3R
 Date: 06/10
 Time: 13:48
 Pbar: 30.14
 Pst: -0.4
 Ps: 30.11 = Pbar+Pst/13.6
 Cp: 0.834
 Fv: 12.99 = 85.49*Cp*sqrt(1/Ps)
 Dia: 96 Area: 50.27
 %OO: 14.5%
 %OCO: 4.5%
 Mdry: 29.30
 H: 10.47%
 O: 1.09%
 C: 86.90% ZEA: 2.495 based on fuel analysis and %OO

Usages
 Rock: 508 TPH
 0.11 HOH
 452 Dry
 Oil: 795 GPH
 8.105 lb/gal
 18509 Btu/lb
 119 mmBtu/hr

Calculation equation summary

fps: std = Fv*sqrt(dp*(Ts ave+460)/Ms)
 point = Fv*sqrt(dp*(Ts pt+460)/(Mdry*(1-%HOH pt)+18*%HOH pt))
 cyc = fps pt*cos(alpha)
 sat %HOH = eqns from Lange's Handbook of Chemistry, pg 1436

		Assumed Twb:		151.4			150				
		sat %HOH:		26.04%			25.15%	23.56%			
Ms = Mdry*(1-%HOH)+18*%HOH		Ms:		26.36			26.46	26.64			
Avg:	17.0	151.4	26.10%			std	std	cyc			
point	dp	alpha	t	%HOH	std	point	cyc	inlet	meas	meas	
SE-1	0.40	16	154	27.74%	39.6	39.8	38.3	39.5	39.4	37.8	
2	0.45	16	155	28.43%	42.0	42.3	40.7	41.9	41.8	40.1	
3	0.45	18	155	28.43%	42.0	42.3	40.2	41.9	41.8	39.7	
4	0.45	18	155	28.43%	42.0	42.3	40.2	41.9	41.8	39.7	
5	0.45	18	155	28.43%	42.0	42.3	40.2	41.9	41.8	39.7	
6	0.45	16	149	24.53%	42.0	41.8	40.1	41.9	41.8	40.1	
SW-1	0.45	16	149	24.53%	42.0	41.8	40.1	41.9	41.8	40.1	
2	0.45	14	149	24.53%	42.0	41.8	40.5	41.9	41.8	40.5	
3	0.50	18	149	24.53%	44.3	44.0	41.9	44.2	44.0	41.9	
4	0.50	18	149	24.53%	44.3	44.0	41.9	44.2	44.0	41.9	
5	0.50	18	149	24.53%	44.3	44.0	41.9	44.2	44.0	41.9	
6	0.50	18	149	24.53%	44.3	44.0	41.9	44.2	44.0	41.9	
		avg									
		fps:		42.5	42.5	40.7	42.5	42.3	40.4		
		acfm:		128289	128293	122641	128044	127612	121991		
		%HOH:		26.04%	26.10%		25.15%	23.56%	23.56%		
		scfm:		82455	82395	78765	83293	84772	81038		
		Emission:		lb/hr							
	ppm	Mgas	pVM/RT*60	std	point	cyc					
NOx	48	46	7.1646	28.4	28.3	27.1	28.6	29.2	27.9		
CO	3.25	28	4.3610	1.2	1.2	1.1	1.2	1.2	1.1		
VOC	0.4	44	6.8531	0.2	0.2	0.2	0.2	0.2	0.2		
SO2	5.7	64	9.9681	4.7	4.7	4.5	4.7	4.8	4.6		
PART	0.020	gr/scf	0.0002	14.2	14.2	13.5	14.3	14.6	13.9		

B. FUEL ANALYSES

PEMBROKE LABORATORY

PEMBROKE, FLORIDA

(813) 533-0969

Mailing address: 528 Gooch Rd., Ft. Meade, FL. 33841

IMCC - NORALYN
P. O. Box 867
Bartow, FL 33830

Date Received: 06-11-87

Date Reported: 06-23-87

Attn: J. N. Allen

P.O. #UCP7543

Lab Numbers	MC-3704	MC-3705	MC-3706	MC-3707
Sample I.D.	1A	2A	3A	4A
BTU/lb	18493.	18538.	18521.	18492.
Density	0.972	0.973	0.972	0.972
Carbon %	86.59	86.79	86.65	86.55
Hydrogen %	10.16	10.10	10.58	10.30
Nitrogen %	0.25	0.14	0.28	0.23
Oxygen %	1.72	1.68	1.23	1.62
Sulfur %	1.28	1.29	1.26	1.30
Ash %	0.054	0.042	0.021	0.023

Continued on page 2

PEMBROKE LABORATORY

PEMBROKE, FLORIDA

(813) 533-0969

Mailing address: 528 Gooch Rd., Ft. Meade, FL. 33841

INCO - NORALYN
P. O. Box 867
Bartow, FL 33830

Date Received: 06-11-87

Date Reported: 06-23-87

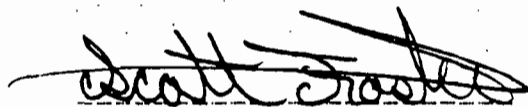
Attn: J. N. Allen

P.O. #UCF7543

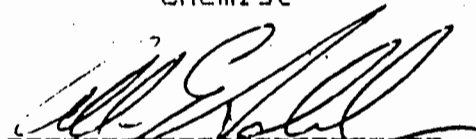
Lab Numbers	MC-3708	MC-3709	MC-3710
Sample I.D.	5A	6A	7A
BTU/lb	18497.	18545.	18509.
Density	0.971	0.969	0.973
Carbon %	86.79	86.67	86.90
Hydrogen %	10.54	10.52	10.47
Nitrogen %	0.15	0.23	0.30
Oxygen %	1.26	1.33	1.09
Sulfur %	1.26	1.25	1.24
Ash %	0.018	0.050	0.056

Thank you for this opportunity to serve you!

Respectfully submitted,



Scott Foster
Chemist



Allan E. Schreiber
Laboratory Director

BEST AVAILABLE COPY

P. E. LaMoreaux & Associates



Geochemistry Laboratory 4320 Old Highway 37

Lakeland, Florida 33803
Telephone 813/646-8526

REPORT OF ANALYSIS

TO: INTERNATIONAL MINERALS & CHEMICAL CORP
P.O. BOX 867
BARTOW, FL 33830
ATTN.: C.D. Turley
P.O. No.: UCP 6544

Project No.:
Sampled by: CLIENT
Date Collected: 04-03-87
Date Received: 04-07-87
Date Completed: 05-13-87
Lab No.: 04-07-IMC-56
Invoice No.: 5563

IDENTIFICATION: LONESOME DRYER FUEL OIL #2
OIL

BTU/Lb.	19,620
Density (lbs/gal)	8.12
% Carbon	86.84
% Hydrogen	10.33
% Nitrogen	0.40
% Oxygen	0.94
% Sulfur	1.84
% Ash	0.11

Results expressed in mg/l (ppm) unless otherwise designated.

Respectfully Submitted

P. E. LaMoreaux & Associates, Inc.

State of Florida Certification #84183
State of Alabama Certification #40120

Chemist

METHODS: "Standard Methods for the Examination of Water and Wastewater", Latest Edition, APHA, AWWA, AND WPCF and/or other EPA approved methods which meet FDER or ADEM protocol, unless otherwise designated.



P. E. LaMoreaux & Associates

Geochemistry Laboratory 4320 Old Highway 37

Lakeland, Florida 33803
Telephone 813/646-8526

REPORT OF ANALYSIS

TO: INTERNATIONAL MINERALS & CHEMICAL CORP
P.O. BOX 867
BARTOW, FL 33830
ATTN.: C.D. Turley
P.O. No.: UCP 6544

Project No.:
Sampled by: CLIENT
Date Collected: 04-03-87
Date Received: 04-07-87
Date Completed: 05-13-87
Lab No.: 04-07-IMC-55
Invoice No.: 5563

IDENTIFICATION: LONESOME DRYER FUEL OIL #1
OIL

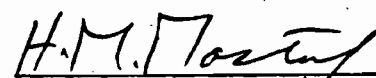
BTU/Lb.	19,360
Density (lbs/gal)	8.15
% Carbon	86.93
% Hydrogen	9.90
% Nitrogen	0.41
% Oxygen	1.25
% Sulfur	1.87
% Ash	0.13

Results expressed in mg/l (ppm) unless otherwise designated.

State of Florida Certification #84183
State of Alabama Certification #40120

Respectfully Submitted

P. E. LaMoreaux & Associates, Inc.


Chemist

METHODS: "Standard Methods for the Examination of Water and Wastewater", Latest Edition, APHA, AWWA, AND WPCF and/or other EPA approved methods which meet FDER or ADEM protocol, unless otherwise designated.

PEMBROKE LABORATORY

PEMBROKE, FLORIDA

(813) 533-0969

Mailing address: 528 Gooch Rd., Ft. Meade, FL. 33841

*4/3 Test Samples
CST*

IMCC - NORALYN
P. O. Box 867
Bartow, FL 33830

Attn: J. N. Allen

Date Received: 04-30-87

Date Reported: 05-29-87

P.O. #UCP7543

Lab Numbers

MC-3389

MC-3390

MC-3391

#3

#2

#1

Sample I.D.

Sample 9389

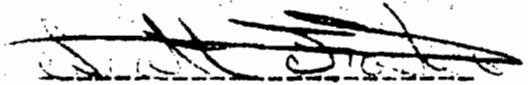
Dryer Oil

Dryer Oil

		MC-3389 #3 Sample 9389	MC-3390 #2 Dryer Oil	MC-3391 #1 Dryer Oil
PCB's	mg/l	15.6	-----	-----
BTU/lb.		-----	18246.	18346.
Density		-----	0.986	0.987
Carbon	%	-----	86.93	86.75
Hydrogen	%	-----	9.92	10.22
Nitrogen	%	-----	0.27	0.13
Oxygen	%	-----	1.34	1.49
Sulfur	%	-----	1.87	1.83
Ash	%	-----	0.055	0.048

Thank you for this opportunity to serve you!

Respectfully submitted,



Scott Foster
Chemist



Allan E. Schreiber
Laboratory Director



P. E. LaMoreaux & Associates

Geochemistry Laboratory 4320 Old Highway 37

Lakeland, Florida 33803
Telephone 813/648-8526

REPORT OF ANALYSIS

TO: INTERNATIONAL MINERALS & CHEMICAL CORP
P.O. BOX 667
BRANTON, FL 33830
ATTN.: David Turley
P.O. No.: UCP 6543

Project No.:
Sampled by: CLIENT
Date Collected: 03-12-87
Date Received: 03-16-87
Date Completed: 04-09-87
Lab No.: 03-16-IMC-146
Invoice No.: 9446

IDENTIFICATION: LONESOME DRYER #1
#6 OIL

STU/LB.	19,107
Density (lbs/gal)	8.14
% Carbon	86.91
% Hydrogen	10.42
% Nitrogen	0.19
% Oxygen	1.07
% Sulfur	1.96
Asph	0.05

Results expressed in mg/l (ppm) unless otherwise designated.

Respectfully Submitted

P. E. LaMoreaux & Associates, Inc.

State of Florida Certification #84183
State of Alabama Certification #40120

H.M. Mast
Chemist

METHODS: "Standard Methods for the Examination of Water and Wastewater", Latest Edition, APHA, AWWA, AND WPCF and/or other EPA approved methods which meet FDER or ADEM protocol, unless otherwise designated.

BEST AVAILABLE COPY

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REPORT OF ANALYSIS

TO: INTERNATIONAL MINERALS & CHEMICAL CORP
P.O. BOX 667
DARTON, FL 33830
ATTN.: David Turley
P.O. No.: UCP 7544

Project No.:
Sampled by: CLIENT
Date Collected: 02-26-87
Date Received: 03-02-87
Date Completed: 03-20-87
Lab No.: 03-02-IMC-03
Invoice No.: 5391

IDENTIFICATION: FT. LONESOME WEST DRYER
OIL

SCRUBBER FUEL OIL

BTU/Lb.
Density (lbs/gal)
% Ash
% Sulfur
% Oxygen
% Nitrogen
% Hydrogen
% Carbon

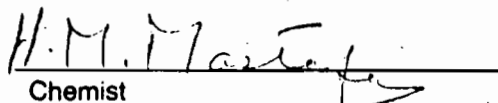
17,853
8.17
2.24
1.53
1.16
0.77
9.59
86.7

Results expressed in mg/l (ppm) unless otherwise designated.

State of Florida Certification #84183
State of Alabama Certification #40120

Respectfully Submitted

P. E. LaMoreaux & Associates, Inc.


Chemist

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C. OPERATING LOGS

Contents: Measurement Sheets

TESTING PARAMETER LOG for TEST: DATE 6/8/87 DRYER NO 2
 MARK STRIP CHARTS in CONTROL ROOM (time and for runs)

BY *George Jensen*

TIME	EVENT LOG	TIME	NOx DIVs	RUN	TIME	START RUN	SCF	TIT VOL
:		:		1 START	:	1		
:		:		1 MID	:	2		
:		:		1 END	:	3		
:		:		2 START	:	4		
:		:		2 MID	:	5		
:		:		2 END	:	6		
:		:		3 START	TIME	SAMPLE	TIME	SAMPLE
:		:		3 MID	13:55	OIL 1	14:35	U/F 1
:		:		3 END	:	OIL	:	U/F
:		TIME	CO DIVs	RUN	:	OIL	:	U/F
:		:		1 START	7:45	FEED 1	14:37	IN 1
:		:		1 MID	14:00	FEED 2	:	IN
:		:		1 END	:	FEED	:	IN
:		:		2 START	TIME	OO 1	OO 2	OO 3
:		:		2 MID	13:25	13	13	13
:		:		2 END	13:50	13	13	13
:		:		3 START	14:09	13	13	13
:		:		3 MID	14:30	13	13	13
:		:		3 END	:			
:		TIME	VOC DIVs	RUN	:			
:		:		1 START	:			
:		:		1 MID	TIME	OCO 1	OCO 2	OCO 3
:		:		1 END	13:25	4	4	4
:		:		2 START	13:50	4	4	4
:		:		2 MID	14:09	4	4	4
:		:		2 END	14:30	4	4	4
:		:		3 START	:			
:		:		3 MID	:			
:		:		3 END	:			
:		ROCK TEMPERATURE OUT			TIME	T AMB	TIME	T ROCK IN
:		14:05	145.4	194	14:06	87.8	14:07	84.2
:		:			:		:	
:		:			:		:	
:		:			:		:	

OPERATING PARAMETER LOG for TEST: DATE 6/8/87 DRYER NO 2

BY Arny Jones

TIME	PRESSURE PSI	TIME	FLOW GPM	TIME	U/F pH	TIME	DELTA P in HOH	TIME	U/F TEMP
14:18	87	14:16	500	14:21	4.8	13:50	25	14:20	148
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	

TIME	INLET TEMP	TIME	OIL METER	TIME	OIL LEVEL	TIME	SCALE TONS	TIME	FLAME PATTERN
13:51	188	11:51	487272	Tank being filled		13:23	2500	:	
:		13:25	487431	:		13:49	2713	:	
:		13:45	487464	:		14:15	2935	:	
:		14:14	487511	:		14:47	3210	:	
:		14:50	487568	:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	

TIME	INLET WET BULB	TIME	LOCATION	MOISTURE DETERMINATION WTS			TIME	BIN LEVEL			
				TARE	INITIAL	FINAL		1	2	3	4
13:56	152	7:35	IN	2.5	52.5	46.7	13:25			44	
:		9:25	OUT	2.5	52.5	50.7	14:43			35	
:		10:18	IN	2.5	52.5	47	:				
:		:	OUT				:				
:		11:25	IN	2.5	52.5	46.6	:				
:		:	OUT				:				
:		14:00	IN	2.5	52.5	46.5	:				
:		14:05	OUT	2.5	52.5	50.4	:				
:		:	IN				:				
:		:	OUT				:				

OPERATING PARAMETER LOG for TEST: DATE 6/9/87 DRYER NO 2 BY CEK

TIME	PRESSURE PSI	TIME	FLOW GPM	TIME	U/F pH	TIME	DELTA P in HOH	TIME	U/F TEMP
14:57	112	14:20	570	15:04	4.9	14:50	24.5	15:04	136
16:09	112	16:00	570	16:08	5	16:08	25.75	16:08	136
18:37	112	17:27	570	17:27	5	18:30	25	18:40	134
:	:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:	:
:	:	:	:	:	:	:	:	:	:

TIME	INLET TEMP	TIME	OIL METER	TIME	OIL LEVEL	TIME	SCALE TONS	TIME	FLAME PATTERN
14:52	189	11:54	487917	13:55	22'7 5/8	14:09	2779	:	
16:05	188	13:53	488128	14:08	22'7 1/4	15:01	3203	:	
18:30	182	14:06	488147	14:59	22'5 5/8	15:25	3402	:	
:	:	14:57	488226	15:23	22'4 13/16	16:16	3824	:	
:	:	15:21	488262	16:54	22'1 3/4	17:09	4267	:	
:	:	16:18	488351	18:25	21'11 13/16	18:23	4629	:	
:	:	16:52	488405	19:18	21'10 1/2	19:15	5066	:	
:	:	18:25	488501	:	:	:	:	:	
:	:	19:17	488582	:	:	:	:	:	
:	:	:	:	:	:	:	:	:	

TIME	INLET WET BULB	TIME	LOCATION	MOISTURE DETERMINATION WTS			TIME	BIN LEVEL			
				TARE	INITIAL	FINAL		1	2	3	4
14:55	151	8:50	IN	2.5	52.5	46.9	14:00				54
16:05	150	8:50	OUT	2.5	52.5	50.6	15:33				44
18:35	148	11:00	IN	2.5	52.5	46.5	17:02				34
:	:	14:45	OUT	2.5	52.5	50.5	19:18				28
:	:	14:45	IN	2.5	52.5	46.5	:				
:	:	15:55	OUT	2.5	52.5	50.5	:				
:	:	15:55	IN	2.5	52.5	46.6	:				
:	:	17:35	OUT	2.5	52.5	50.4	:				
:	:	17:35	IN	2.5	52.5	48.1	:				
:	:	:	OUT	:	:	:	:				

TESTING PARAMETER LOG for TEST: DATE 6/9/87 DRYER NO 2
 MARK STRIP CHARTS in CONTROL ROOM (time and for runs)

BY CEK

TIME	EVENT LOG	TIME	NOx DIVs	RUN	TIME	START RUN	E-6 LB/SCF	%HOH	
13:53	Temp start down	:		1 START	14:13	1	0.96	0.30125	
		:		1 MID					
14:04	Start Traverse 1	:		1 END	14:53	2	1.23	0.2899	
14:15	Start Run 1	:		2 START	15:38	3	1.99	0.30735	
15:20	Start Traverse 2	:		2 MID	16:16	4	1.16	0.30399	
15:40	Start Run 2	:		2 END	17:18	5	1.1	0.27865	
17:20	Start Run 3	:		3 START	18:21	6	1.44	0.28509	
17:38	Flame Out	:		3 MID					
18:05	Dryer start up	:		3 END	14:25	OIL 2	15:04	U/F 2	
18:20	486 TPH	:			15:50	OIL 3	16:08	U/F 3	
		TIME	CO DIVs	RUN	17:30	OIL 4	18:40	U/F 4	
		:		1 START	14:25	FEED 3	15:08	IN 2	
		:		1 MID	15:44	FEED 4	16:15	IN 3	
		:		1 END	17:25	FEED 5	18:43	IN 4	
		:		2 START					
		:		2 MID	TIME	OO 1	OO 2	OO 3	
		:		2 END	14:10	13.5	13.5	13.6	
		:		3 START	14:53	13.8	13.8		
		:		3 MID	15:30	13.8	13.8	13.8	
		:		3 END	16:16	14	14		
		:			17:10	14	14	14	
		TIME	VOC DIVs	RUN	18:21	14.4	14.4		
		:		1 START	19:30	13	13	13	
		:		1 MID					
		:		1 END	TIME	OCO 1	OCO 2	OCO 3	
		:		2 START	14:10	4.5	4.6	4.5	
		:		2 MID	14:53	4.6	4.6		
		:		2 END	15:30	4.5	4.5	4.5	
		:		3 START	16:16	4.6	4.6		
		:		3 MID	17:10	4.5	4.6	4.5	
		:		3 END	18:21	3.6	3.6		
		:			19:30	4	4	4	
		ROCK TEMPERATURE OUT							
		TIME	BELT	METER	TIME	T AMB	TIME	T ROCK IN	
		14:32	132	179	15:12	90	14:25	83	
		15:52	145	184	16:15	90	15:44	82	
		17:28	133	175	18:46	78	17:25	81	
		:			:		:		

OPERATING PARAMETER LOG for TEST: DATE 6/10/87 DRYER NO 1 BY CEK

TIME	PRESSURE PSI	TIME	FLOW GPM	TIME	U/F pH	TIME	DELTA P in HOH	TIME	U/F TEMP
11:36	114	11:35	580	11:45	6.3	11:42	28.4	11:44	132
13:03	115	12:54	580	13:00	6.3	13:01	28.4	13:02	134
14:20	115	14:15	580	14:24	6.3	14:22	28.5	14:26	130
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	

TIME	INLET TEMP	TIME	OIL METER	TIME	OIL LEVEL	TIME	SCALE TONS	TIME	FLAME PATTERN
14:25	176	10:50	303206	10:47	20'4 3/8	10:51	4702	:	
:		11:42	303288	12:22	20'1 3/8	11:43	5128	:	
:		12:20	303347	13:39	19'11	12:19	5445	:	
:		12:59	303408	14:56	19'8 7/8	12:58	5777	:	
:		13:37	303465	:		13:36	6095	:	
:		14:21	303522	:		14:21	6481	:	
:		14:54	303567	:		14:54	6755	:	
:		:		:		:		:	
:		:		:		:		:	
:		:		:		:		:	

TIME	INLET WET BULB	TIME	LOCATION	MOISTURE DETERMINATION WTS			TIME	BIN LEVEL			
				TARE	INITIAL	FINAL		1	2	3	4
10:50	150	9:20	IN	2.5	52.5	47	10:50	29		23	
14:25	150	:	OUT				12:18	24	70	18	
:		11:30	IN	2.5	52.5	46.9	13:25			11	
:		11:30	OUT	2.5	52.5	50.8	13:36	20	70		
:		12:50	IN	2.5	52.5	46.9	14:55		46		
:		12:50	OUT	2.5	52.5	51	:				
:		14:15	IN	2.5	52.5	47	:				
:		14:15	OUT	2.5	52.5	50.4	:				
:		:	IN				:				
:		:	OUT				:				

TESTING PARAMETER LOG for TEST: DATE 6/10/87 DRYER NO 1 BY CEK
 MARK STRIP CHARTS in CONTROL ROOM (time and for runs)

TIME	EVENT LOG	TIME	NOx DIVs	RUN	TIME	START RUN	E-6 LB/SCF	%HOH
11:13	Start Run 1	:		1 START	11:30	1	1.51	0.29864
12:33	Start Run 1	:		1 MID	12:06	2	1.6	0.27762
13:36	Switch to Bin 2	:		1 END	12:40	3	1.82	0.29866
13:49	Start Run 1	:		2 START	13:16	4	1.71	0.28867
:		:		2 MID	13:50	5	0.8	0.25836
:		:		2 END	14:21	6	1.09	0.25635
:		:		3 START	TIME	SAMPLE	TIME	SAMPLE
:		:		3 MID	11:30	OIL 5	11:45	U/F 5
:		:		3 END	12:45	OIL 6	12:55	U/F 6
:		TIME	CO DIVs	RUN	14:04	OIL 7	16:22	U/F 7
:		:		1 START	11:17	FEED 6	11:49	IN 5
:		:		1 MID	12:40	FEED 7	12:55	IN 6
:		:		1 END	13:55	FEED 8	16:28	IN 7
:		:		2 START	TIME	OO 1	OO 2	OO 3
:		:		2 MID	11:00	14.5	14.5	14.5
:		:		2 END	12:06	14.6	14.6	
:		:		3 START	12:20	14.5	14.5	14.5
:		:		3 MID	12:40	14.6	14.6	
:		:		3 END	13:38	14.5	14.5	14.5
:		TIME	VOC DIVs	RUN	14:21	16.2	16.2	
:		:		1 START	TIME	OCO 1	OCO 2	OCO 3
:		:		1 MID	11:00	4.5	4.5	4.5
:		:		1 END	12:06	4.2		
:		:		2 START	12:20	4.5	4.5	4.5
:		:		2 MID	12:40	4.2	4.4	
:		:		2 END	13:38	4.5	4.5	4.5
:		:		3 START	14:21	3.4	3.4	
:		:		3 MID	:			
:		:		3 END	:			
:		TIME	ROCK TEMPERATURE	OUT	TIME	T AMB	TIME	T ROCK IN
:		TIME	BELT	METER	11:46	87	11:17	78
:		11:25	136	179	13:00	90	12:40	79
:		12:45	136	184	14:38	89	13:55	79
:		14:07	165	179	:		:	