

Victorville #2
coal Precalciner

Note: This is a reference cited in AP 42, *Compilation of Air Pollutant Emission Factors, Volume I Stationary Point and Area Sources*. AP42 is located on the EPA web site at www.epa.gov/ttn/chief/ap42/

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AP-42 Section	<u>11.6</u>
Reference	<u>58</u>
Report Sect.	<u>4</u>
Reference	<u>63</u>

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EMISSION TESTS ON QUARRY PLANT #2 KILN

February 24, 1987

Prepared for

Southwestern Portland Cement
North End "E" Street
Victorville, California 92392

March 1987

Prepared by

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Report PS-87-1024/Project 5904-87

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*Ref 42 Precalciner Kiln
Down rated from
A to D by MAI but
I'm not sure why.
Report includes Calibration
data & good values for
CO & CO₂*

Pr
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5801 N
Bakersfield,

Report PS-87-1024/Project 5904-87

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SECTION 1

INTRODUCTION

At the request of Southwestern Portland Cement, Pape & Steiner Environmental Services undertook a series of emission tests on the effluent of #2 Kiln on February 4, 1987. Kiln #2 is in service at SwPC's quarry plant in Victorville, California. The purpose of these tests was measure the particulate emission rates and determine compliance with SBCAPCD Rules 404 and 405.

This test series was also meant to check the emission rate from the baghouse serving #2 Kiln after a preliminary report from a CARB Audit Test indicated noncompliance. The CARB audit tests were conducted November 11, 1986. A copy of CARB's final report was not available during this test series. The results of this test series shows compliance with San Bernadino County APCD Rules 404 and 405.

Messrs. Ralph Pape and Steve Carter conducted the source tests for Pape & Steiner Environmental Services on February 24, 1987. Triplicate tests for particulate were conducted using EPA Method 5. Triplicate tests for NO_x , SO_2 , CO , CO_2 and O_2 were conducted using CARB Method 1-100 continuous monitor procedures. Particulate data for the back half catch is reported for this test series. The back half catch was not requested

during the initial compliance demonstration testing in March 1985. Some problems were encountered during the analysis of the back half samples. Reactions between chlorides in the sample and the aluminum weighing dishes caused erroneously high weights. These analyses were repeated using glass and a much lower weight was found. The weights from the analysis in glass were used to calculate the total particulate emission rates.

Kiln #2 operations were increasing to full load during Test 1 and accounts for the lower average feed rate. Operations were fairly constant during Tests 2 and 3. Some changes to the fuel rates were being made to control temperature. These changes seemed only to affect the NO_x emission levels.

Section 2 of this report presents source test results. Sampling and analysis procedures are described in Sections 3 and 4. Section 5 presents the quality assurance procedures used during this program. Raw data sheets, continuous monitor strip charts, calculation worksheets and equipment calibration are included in the Appendix.

SECTION 2

RESULTS

Table 2-1 presents the results of particulate and continuous monitor source tests. The data is presented at SBCAPCD standard conditions (60°F, 29.92 inches of Mercury).

Kiln 2 operations were increasing to full load during Test 1 which accounts for the lower average feed rate. Operations were fairly constant during Tests 2 and 3. Some changes to the fuel rates were being made to control temperature. The changes seemed only to affect the NO_x emission levels.

The particulate mass is reported in two ways -- front half and front half plus back half. The front half catch (nozzle, probe, filter holder wash and filter) represents matter which is filterable at 250°F. EPA bases compliance on this value. The back half catch represents matter which is caught in the water impingers of the sample train. San Bernadino County bases compliance of the combined front and back half catches. Subsequent analysis of material found in the back half showed it was primarily ammonium chloride and sulfate. Since these materials are filterable, they should have been collected in the front half of the sample train. Finding them in the back half indicates they were formed there by stack gases

trapped in the impinger solutions. The back half catch should be disregarded from this source because the gaseous components form pseudo-particulate in the back half of the sample train.

NO_x , SO_2 and CO concentrations are presented on a dry volume basis. NO_x concentrations have also been standardized to 3-percent O_2 . Pound per hour emission rates were calculated from the concentrations and volumetric flowrate measurements.

$$KF \text{ LOI} = 34.55\%$$

$$\text{Clinker}_{TPH} = \frac{KF_{TPH} \times 0.89}{1.65}$$

TABLE 2-1. SUMMARY OF SOURCE EMISSION TEST DATA

Unit Tested: Kiln 2 Baghouse

Location: SWPC Quarry Plant

Test Number	1	2	3	Average
Date		February 24, 1987		
Test Condition (Tons Per-Hour Feed)	193	214	211	
Barometric Pressure (in. Hg)	26.08	26.04	26.00	26.04
Stack Pressure (in. Hg)	26.07	26.03	25.99	26.03
Stack Area (Ft ²)	139.63	139.63	139.63	139.63
Elapsed Sampling Time (min.)	72	72	72	72
Volume Gas Sampled (dscf)	48.4632	50.0944	49.1162	49.2246

GAS DATA

Average Gas Velocity (fps)	38.31	39.74	38.30	38.78
Average Gas Temperature (°F)	374	380	367	374
Gas Flowrate (dscfm)	165,108	169,820	165,749	166,893
Gas Analysis (Dry % Basis)				
Carbon Dioxide	20.2	20.1	20.6	20.3
Oxygen	8.9	8.7	8.8	8.8
Water	5.3	5.3	5.4	5.4

EMISSION CONCENTRATION

Filterable Particulate (gr/dscf)	0.0026	0.0022	0.0029	0.0026
Rule 404 -- Total Particulate (gr/dscf)	0.0039	0.0032	0.0042	0.0038
CO (ppm)	500	463	533	499
SO ₂ (ppm)	1.8	2.2	2.5	2.2
NO _x (ppm)	650	611	541	601
NO _x (ppm at 3% O ₂)	972	900	800	891

EMISSION RATE

Filterable Particulate (lb/hr)	3.61	3.20	4.08	3.63
Rule 405 -- Total Particulate (lb/hr)	5.46	4.72	6.00	5.39
CO (lb/hr)	365.53	347.77	391.39	368.23
SO ₂ (lb/hr)	3.06	3.74	4.20	3.67
NO _x (lb/hr)	781.04	754.53	652.04	729.20

Sunnyside Coal - @ 13050 BTU/lb = 26.1 MMBTU/Ton From Printout Last page of report	KF - T/hr	206	215.0	211.2	210.7
	Clinker - T/hr	111	116	114	113.7
	Coal - T/hr				
	" - pyroclastic	5.9	7.85	7.50	
	" - Kiln	7.60	7.70	7.50	
	" Total	13.5	15.55	15.0	14.7
	NO _x #/TCR	9.04	6.50	5.72	6.41
	NO _x #/MMBTU	2.22	1.86	1.67	1.91
	2-3				
	NO _x /TKF	3.79	3.51	3.09	3.46

SECTION 3

SAMPLING PROCEDURES

This section of the report describes the equipment and procedures used to conduct the particulate gaseous tests on this program.

3.1 PRELIMINARY MEASUREMENTS

Before conducting the stack tests a series of preliminary measurements were made to determine:

- The location of the sampling site and the number and location of the sampling points to be used (EPA Method 1)
- The velocity, temperature, and pressure of the gases in the stack (EPA Method 2)
- The composition of the stack gases (EPA Method 3)
- The moisture content of the stack gases (psychrometric chart)

Using the results of these preliminary measurements and the calibration constants for the sampling train, a series of calculations were made to determine the value of K, a constant, and N_d , ideal nozzle diameter, required to run an isokinetic test according to the equation:

$$\Delta H = \left\{ \frac{60^2 \pi^2 K_p^2 C_p^2 (1 - B_{wo})^2 P_s^{MW_d}}{576^2 K_o^2 MW_s P_m} \right\} (N_d^4) \left(\frac{T_m}{T_s} \right) (\Delta P)$$

where

$$K = \left\{ \frac{60^2 \pi^2 K_p^2 C_p^2 (1 - B_{wo})^2 P_s MW_d}{576^2 K_o^2 MW_s P_m} \right\}$$

An actual nozzle, whose diameter was as close as possible to the ideal nozzle diameter, was selected for the test. Isokinetic sampling rates for each sampling point in the stack were computed using the equation:

$$\Delta H = (K)(N_d^4) \left(\frac{T_m}{T_s} \right) (\Delta P)$$

Since K and N_d are known, and remain constant during a test, the only variables are the meter temperature, the stack temperature and the velocity pressure for each sampling point.

3.2 PREPARATION OF THE PARTICULATE SAMPLING TRAIN

All sampling train components were cleaned in the laboratory (soap and water, tap water rinse, distilled water rinse, and IPA rinse) to eliminate previous contamination. The sampling train components were sealed and transported to the sampling site in a mobile lab. The EPA Method 5 equipment used to measure particulates (filterable and total) consisted of:

- A calibrated 316 stainless steel nozzle for isokinetic sampling
- A heated Pyrex glass sampling probe (6 feet long) equipped with an S-type pitot tube and a thermocouple to measure stack velocity, pressure and temperature

- A heated Pyrex glass filter holder containing a weighed 100-mm Reeve Angel 934 AH glass fiber filter
- A Pyrex glass impinger train in an icebath (impingers 1 and 2 contain 100 ml distilled water; impinger 3 is dry; bubbler 4 contains a weighed amount of silica gel)
- An umbilical to connect the probe and sample box to the control module
- A control module containing a vacuum pump, a calibrated dry gas meter and a calibrated orifice meter to measure the pressure, temperature and flowrate throughout the train.

The sampling train was charged in the mobile lab using freshly prepared reagents. Each impinger and its contents was weighed to the nearest 0.1 gm on a calibrated electronic balance. Blanks of all filters and reagents were retained for subsequent analysis. The sampling point locations were marked on the probe using a high-temperature marker. The sampling train was completely assembled and lifted to the sampling site.

3.3 SAMPLING PROCEDURES FOR PARTICULATE SAMPLING TRAIN

Prior to a test, the sampling train was heated and leak-checked at 15 inches Mercury to insure leakage was less than 0.02 cfm. The S-type pitot tube was also leak-checked. The sampling train was installed on the unirail and the probe was inserted into the stack at the farthest point. An isokinetic sampling rate was calculated using an HP-41CV calculator for each sampling point on the traverse (6 points per traverse; 4 traverses at 90°). Each point was sampled for an equal period of time (3 minutes)

and all pertinent data was recorded on the data sheet for each point. The probe and sample box were maintained at 250°F throughout the traverse. The gases leaving the impinger train were maintained at <70°F. At the end of a traverse, the probe was withdrawn from the stack and the entire sampling train was transferred (in tact) to the next sampling port. Another traverse of the stack was completed and the sampling train was withdrawn for the final leak check. This leak check was performed at 15 inches Mercury or at the highest vacuum achieved during the test. The S-type pitot tube was also checked at this time. The sampling train was then purged with ambient air for 15 minutes using the highest ΔH measured during the test. After the train was purged, the probe, nozzle, filter holder, and impinger train was sealed with tin foil and lowered to the mobile lab for sample recovery.

3.4 SAMPLE RECOVERY PROCEDURES FOR PARTICULATE SAMPLING TRAIN

Sample recovery occurred in the mobile lab. The nozzle and probe were brushed and rinsed three times using ACS reagent grade acetone into a polyethylene sample bottle. The 100-mm filter was removed from the 4-inch filter holder and sealed in its petri dish. The front half of the 4-inch glass filter holder was brushed and rinsed with acetone. Each impinger was removed from the icebath, wiped dry and weighed to the nearest 0.1 gm. The contents of impingers 1, 2 and 3 were transferred to a polyethylene sample bottle. The back half of the 4-inch glass filter holder, the glass connectors, impingers 1, 2 and 3 were rinsed with distilled water and the rinsings were transferred to this same bottle. All sample bottles and

petri dishes were marked and labeled. A chain-of-custody log was completed and the field data sheet was also labeled with the sample ID numbers. The sampling train was then recharged in preparation for the next test.

3.5 SAMPLING PROCEDURES FOR CONTINUOUS MONITORING

The continuous monitors used in the Pape & Steiner Mobile Monitoring Lab are shown in Table 3-1. Figure 3-1 is a schematic of the continuous monitoring system. The procedures used to continuously monitor stack gases for NO_x , SO_2 , CO , CO_2 , and O_2 strictly follow CARB Method 1-100.

Sample was taken from the stack using a 316 stainless steel probe. A Balston filter holder and fiberglass filter (99.9999 percent efficiency retention of 0.6 micron particles) was connected to the outlet of the probe. Sample gas was transported through heated Teflon sample line (maintained at $>250^\circ\text{F}$) by a Teflon-lined diaphragm pump to a 316 stainless steel refrigeration type conditioner (Hankison Model E-4G-SS). The sample gas was passed through the conditioner two separate times under vacuum before entering the pump, then two additional times under pressure. The clean, dry sample gas (approximately 35°F) was then transported to the continuous analyzer system through an unheated Teflon line. A series of flowmeters, valves, and regulators maintain constant flow through the system at a constant pressure.

Calibrations of the continuous analyzers were performed using EPA Protocol 1 calibration gases (± 1 percent) for criteria pollutant analysis (NO_x/SO_2) and certified calibration gases (± 1 percent) for fixed gas analysis. Copies of the gas certifications are included in the Appendix of

TABLE 3-1. CONTINUOUS MONITORING LAB

NO_x CHEMILUMINESCENT ANALYZER -- THERMO ELECTRON MODEL 10

Response Time (0-90%)	1.5 sec -- NO mode 1.7 sec -- NO _x mode
Zero Drift	Negligible after 1/2 hour warmup
Linearity	+1% of full scale
Accuracy	Derived from the NO or NO ₂ calibration gas, +1% of full scale
Output	0.100 mV, 0-10 mV, 0-5 V, 0-10 V

O₂ ANALYZER, FUEL TYPE -- TELEDYNE MODEL 326

Response Time (0-90%)	60 seconds
Accuracy	+1% of scale at constant temperatures; +1% of scale of +5% of reading, whichever is greater, over the operating temperature range
Output	0-1 V

CO₂/CO INFRARED ANALYZER -- ANARAD MODEL AR-600

Response Time (0-90%)	5 seconds
Zero Drift	+1%
Span Drift	+1%
Linearity	1%
Resolution	Less than 1% of full scale
Output	0-1 V

SO₂ UV ANALYZER -- DUPONT MODEL 400

Response Time (0-90%)	Less than 1 second
Zero Drift	Less than 1% full scale in 24 hours
Linearity	+1% full scale
Accuracy	+2% full scale
Output	0-10 mV
Operating Ranges	0-100 ppm, 0-1000 ppm

STRIP CHART RECORDERS (3) -- LINEAR MODEL 486

Pen Response	20 inches/second
Span -- full scale	1 mV through 10 V
Zero Set	Electronically adjustable full scale with 1 full scale of zero suppression
Accuracy	Total limit of error $\pm 0.5\%$
Dead Band	
Linearity	
Repeatability	

SCOTSMAN TRAILER

Fully Insulated
Air conditioned
8-ft x 14-ft x 11-ft

1. Filter 0.6 μ , 99.9999 percent efficient
2. Duct
3. 316 stainless steel probe
4. 3/8-inch, heated (250°F) Teflon
5. Four-pass conditioner-dryer, 316 stainless steel internals
6. 3/8-inch, unheated Teflon
7. Teflon-lined sample pump
8. 3/8-inch unheated Teflon
9. Rotameter
10. 1/4-inch Teflon tubing
11. Calibration gas manifold
12. Calibration gas selector valve
13. Calibration gas cylinders
14. Backpressure regulator
15. Auxiliary analysis port

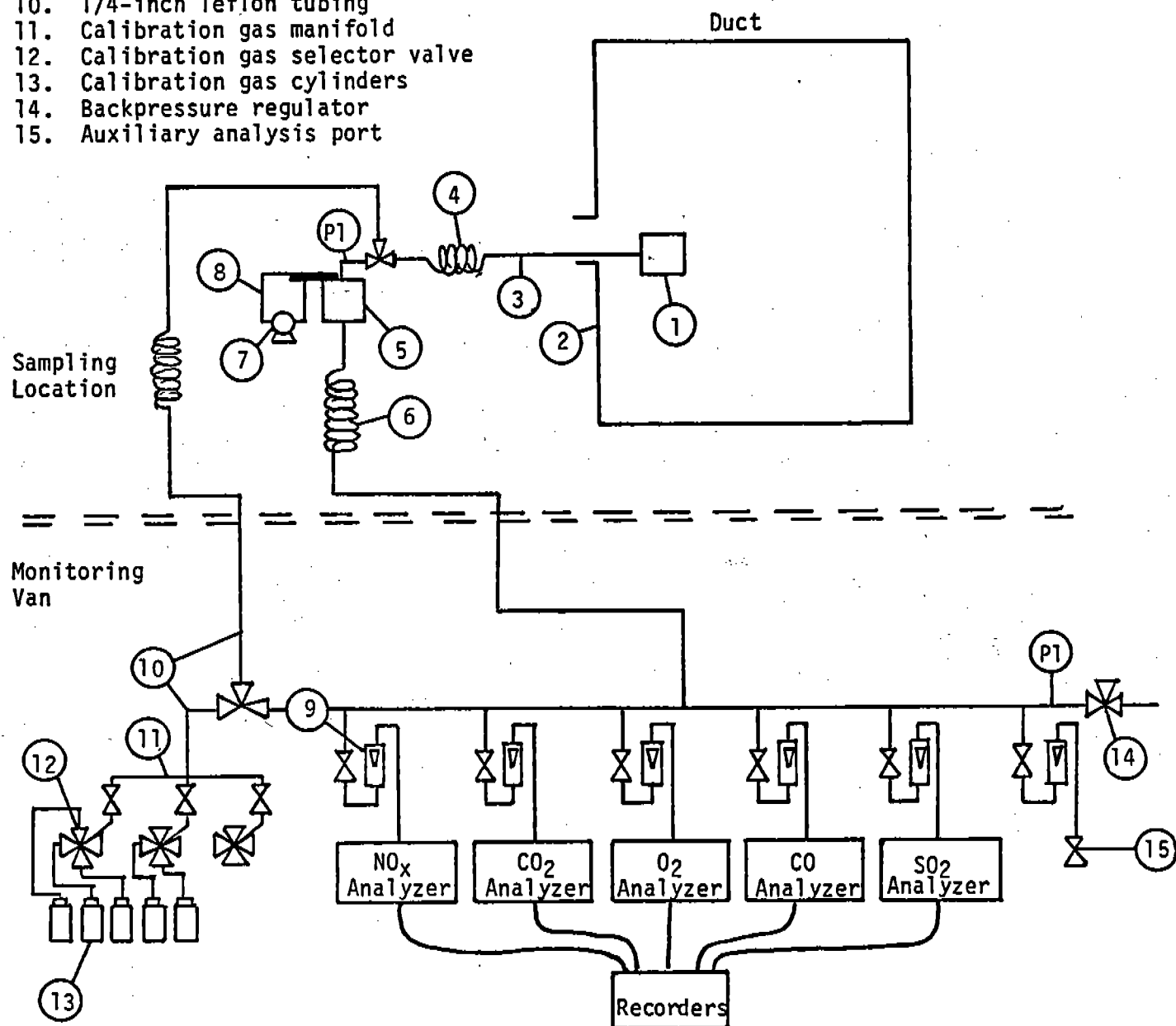


Figure 3-1. Schematic of Continuous Monitoring System.

this report. All pertinent data (date, time, test locations, analyzer range, cal gas value) were recorded on both the field data sheets and continuous analyzer strip charts in the field.

At the start of the test, a leak-check was performed. The sample probe was removed from the stack and the end was sealed with a Swagelok cap. A leak-check is successful only if pressure at the analyzer system and flow through the rotameters to the individual analyzers all drop to zero. A mandatory leak-check was performed at the completion of each test.

An initial calibration was performed at the start of the test period by introducing zero and span gases for each analyzer and making the necessary adjustments. Calibration gas values were recorded on the continuous monitor strip charts and field data sheets. A calibration check was completed at the end of the test and adjustments (if necessary) to the analyzers were made in preparation for another test.

Test data was collected by recording 10-minute averages from the strip chart recordings onto the field data sheets. Data collected over the 2-hour test period was averaged and reported. A fuel analysis was used to calculate the F-factor, dscf/MMBtu corrected to zero percent O_2 (standard conditions 60°F, 29.92 inches Mercury), as described in 40 CFR 60.45. The NO_x and the F-factor were used to calculate the emission factor in lb/MMBtu.

SECTION 4

ANALYTICAL PROCEDURES

This section of the report describes the procedures used to analyze the samples collected during the test program. All analyses were performed in the Pape & Steiner climate-controlled laboratory in Bakersfield.

4.1 ANALYSIS OF PARTICULATE SAMPLES

4.1.1 Nozzle, Probe, Filter Holder Wash

The volume of the acetone washings were measured and the washings were transferred to clean, tared, aluminum weighing dishes. The dishes were placed on a hot plate in a fume hood and gently heated to dryness. The dishes with the dry residue were desiccated and weighed repeatedly at 6-hour intervals until a constant weight was achieved (to the nearest 0.01 mg with a tolerance of <0.1 mg between weighings). The ACS reagent grade acetone blank was treated in the same manner.

4.1.2 Filter

The 100 mm filter was removed from its petri dish and transferred to an oven where it was heated for 2 hours at 105°C. The filter was then desiccated and weighed repeatedly at 6-hour intervals until a constant weight was achieved (to nearest 0.01 mg with a tolerance of <0.1 mg between weighings). An unused, tared blank filter was treated in the same manner.

221°F

4.1.3 Condensible Particulate

Acid mist filter

why?

The 47-mm glass fiber filter was leached with distilled water and the leach was added to the contents and washings from impinger 1. The volume was measured and an aliquot was transferred to a clean, tared evaporating dish. The dish was placed on a hot plate in a fume hood and gently heated to dryness. The dish with the dry residue was desiccated and weighed repeatedly at 6-hour intervals until a constant weight was achieved (to nearest 0.01 mg with a tolerance of <0.1 mg between weighings).

Is for Method 5-2 - is back filter between
IPA impingers & H_2O_2 impingers.

Does not belong in this report.

DGM per Ralph Pape 3/16/87

SECTION 5

QUALITY ASSURANCE

5.1 PARTICULATE SAMPLING EQUIPMENT

A detailed record of repair and maintenance to each sampling train is kept. Preventative maintenance to each system is performed periodically to avoid complete component breakdown during a field test.

A detailed record of sampling system calibrations is also kept. Calibration data for the sampling nozzles, pitot tubes, dry gas meters and orifice meters are available for review. Results of the EPA Quality Assurance Branch biannual audits of the dry gas meter and orifice meter combinations are also logged and verify our in-house calibration data. The calibration data for the equipment used on this program can be found at the end of this section.

5.2 LAB ANALYSIS

All field samples are assigned a label and an ID number. This ID is also affixed to a chain-of-custody log and to the field data sheet to eliminate any chance of sample mixup.

Prior to analysis, all glassware is thoroughly cleaned (soap and water, tap water rinse, distilled water rinse, IPA rinse) to eliminate any contamination. The evaporating dishes used to evaporate the washings are

treated the same as a sample (dried in an oven, desiccated and weighed repeatedly at 6-hour intervals until a constant weight is achieved). The glassware used to measure volumes and make transfers and dilutions are all NBS Class A to insure accurate measurements. All weighings are carried out on a Mettler Model H54AR analytical balance supported by a marble table in a separate room from the main analytical laboratory. The balance is calibrated regularly against an NBS Class S-1 weight.

All reagents used in the field and in the laboratory are ACS reagent grade and blanks of these reagents are evaluated for every set of tests. Blanks are taken in the field from the squeeze bottles and not the original container. Records are kept on these blanks to insure consistent quality of the reagents. Prior to use, the IPA is also analyzed to insure no peroxides are present which could lead to high SO_3 and low SO_2 values.

5.3 CONTINUOUS MONITORS

The NO_x and SO_2 analyzers are calibrated before and after each test using an EPA Protocol 1 gas (± 1 percent) traceable to NBS. The CO , CO_2 , and O_2 analyzers are calibrated before and after each test using a certified gas mixture (± 1 percent). Copies of these certifications are included in the Appendix of this report.

A system check was performed during calibration checks. This was done by introducing span gas at the inlet to the sampling handling system (after the heated line) and measuring the response. The purpose of this test is to check the sampling handling portion of the continuous monitor system for sample loss.

The CO interferences in NDIR-type analyzers described in EPA Method 10, caused by CO₂, have been eliminated by utilizing an electronic cross-talk feature in the dual CO/CO₂ analyzer. The crosstalk is setup by injecting a known amount of CO-free CO₂ and balancing out the CO interference response. Once this has been done, the CO values will automatically be corrected for the amount of CO₂ being analyzed at any particular instant.

A multipoint calibration linearity check of the continuous analyzers was performed on December 10, 1986. These results were well within CARB limitations of +2 percent of full scale. Table 5-1 lists the results of this check.

TABLE 5-1. MULTIPOINT CALIBRATION LINEARITY CHECKS
(TRAILER 2 - December 10, 1986)

Gas	Cal Gas (ppm)	Final Reading (ppm)	Difference (ppm)	% Difference (CARB allowable ±2% Full Scale)
NO _x	0	0	0	0
	40.0	40.0	0	0
	90.3	92	1.7	1.7
	203.6	205	1.4	0.6
SO ₂	0	0	0	0
	7*	5	-2	-2.0
	51.53	51	-0.53	-0.5
CO	0	0	0	0
	24.96	24	-0.96	0.2
	201	202	1.0	0.2
CO ₂	0	0	0	0
	8.04	8.0	-0.04	-0.2
	14.1	14.1	0	0
O ₂	0	0	0	0
	4.03	4.0	-0.03	-0.3
	20.9	20.8	-0.1	-0.4

Method 20 NO_x Checks:

	NO _x Response	NO ₂ Response
201 ppm CO	0	
51.53 ppm SO ₂	0	
18.5 ppm NO ₂ Converter Efficiency		18

*EPA Method 5/8

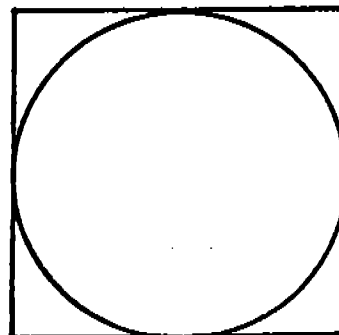
BPR14

APPENDIX

SAMPLING POINT LOCATION DATA SHEET

#1853/.3217

Plant SWPC
 Date 2/27/87
 Test Location #2 Km Outlet
 Upstream Dist./Dia. 92"/.6
 Downstream Dist./Dia. 416"/2.6
 No. of Sampling Points 24
 Stack Dimension 160"
 Coupling Length 5.75
 _____-inch MPT/FPT/Flange



Sample Point	Dist	Sample Point	Dist	Sample Point	Dist	Sample Point	Dist
1	3.9	9.2					
2	16.2	16.5					
3	24.4	24.6					
4	33.9	34.1					
5	45.5	45.8					
6	62.4	62.7					

SAMPLING POINT RELOCATION:

K FACTOR/NOZZLE CALCULATION

$$K = \frac{782.687(C_p)^2(1 - B_{wo})^2 P_s MW_d}{K_o^2 MW_s P_m}$$

B
 $\frac{\Delta P}{T_s}$
.30 380
.30 "
.38 "
.44 "
.41 "

XEQ KC

CP .84
 B_{wo} .045
 $P_s H_2O$ -.18
 P_b 26.08

O_2 9.0
 CO_2 17.0
 K_o 0.8331

MW_d 31.080
 $K(R10)$ 735.602

$$N_d = \left(\frac{H T_s}{K T_m \Delta P} \right)^{.25}$$

XEQ ND

ΔP 0.38
 T_s 379
 T_m 75

Ideal Nozzle @ 1.1 ΔH = 0.280
 Ideal Nozzle @ 1.8 ΔH = 0.317

H
 $\frac{\Delta P}{T_s}$
.27 376
.36 379
.36 "
.45 "
.48 "
.50 "

XEQ KND

(1) Actual Nozzle I.D. .3217
 $K(N_d)^4(R09)$ = 7.879

(2) Actual Nozzle I.D. _____
 $K(N_d)^4(R09)$ = _____

XEQ DH

Nozzle 1

Nozzle 2

	High	Low
T_m in	<u>75</u>	<u>75</u>
T_m out	<u>75</u>	<u>75</u>
T_s	<u>380</u>	<u>376</u>
ΔP	<u>.51</u>	<u>.27</u>
ΔH	<u>2.56</u>	<u>1.36</u>

High	Low
_____	_____
_____	_____
_____	_____
_____	_____
_____	_____

Av
 $\sqrt{\Delta P} = 0.6165$
 $\Delta P = 0.38$
 $T_s = 379$ CLΣ

Date 2/24/87

Test Location 12 km

Run Number 1

Stack Diameter | 60"

Operator Σ_{WP}

Filter No. 814

Barometric Pressure 26.08

Static in. wg. - 1.8

Probe Type/Length (2-6')

Pitot Coefficient .84

Meter Box No. / 8699 / 9483

Nozzle No./Size 853/3217

Impinger Volumes/Weights				Gas Composition			
Contents	Final	Initial	Net	Time	CO ₂	O ₂	CO
DI	581.1	542.4	38.7		20.2	8.9	
DI	536.6	520.8	7.8				
Blank	154.3	453.6	0.7				
				Leak Rate	cfm		"Hg
				Initial		0.009	16.0
S.G. silica gel	689.6	682.2	11.4	Final			
		Total	98.6				

[illegible]

Impinger Volumes/Weights				Gas Composition			
Contents	Final	Initial	Net	Time	CO ₂	O ₂	CO
S.G.				Leak Rate		cfm	"Hg
				Initial			
				Final			
Total						0.013	8.0

Sample Point	Time	ΔP in wg	ΔH in wg	Gas Meter Volume Ft ³	Temperature °F					Pump Vacuum in. Hg	$\sqrt{\Delta P}$	Comments	
					Stack	Probe	Oven	Imp.	Gas Meter				
									In	Out			
C-1	0	.24	1.18	690.746	372	258	245	16	62	58	3.5	0.490	
2	3	.21	1.04	692.828	372	259	247	19	71	60	3.0	0.458	
3	6	.18	0.90	694.878	371	262	258	23	74	61	3.0	0.424	
4	9	.18	0.90	696.801	371	256	251	23	75	61	3.0	0.424	
5	12	.14	0.70	698.720	372	254	257	25	75	61	2.5	0.374	
6	15	.10	0.50	700.458	370	255	254	26	75	62	2.0	0.316	
A-1	18	.24	1.19	702.961	371	256	254	23	63	59	3.5	0.490	1047 standard
2	21	.24	1.19	704.111	376	253	252	18	73	60	3.5	0.490	
3	24	.22	1.09	706.258	381	251	256	18	75	61	3.5	0.469	
4	27	.20	0.99	708.357	381	250	253	20	77	61	3.0	0.447	
5	30	.22	1.09	710.381	381	248	257	20	77	63	3.5	0.469	
6	33	.24	1.19	712.470	380	242	247	23	77	63	3.5	0.490	
	36			714.666									
			1.40	55.957	373.79					64.52		0.5240	

506.70

Date 2/24/87
 Test Location #2 Kiln
 Run Number 2
 Stack Diameter 160"
 Operator SWC
 Filter No. 815
 Barometric Pressure 26.04
 Static in. wg. -.18
 Probe Type/Length 6-16'
 Pitot Coefficient .84
 Meter Box No. 18609/9383
 Nozzle No./Size 853/3217

Impinger Volumes/Weights				Gas Composition			
Contents	Final	Initial	Net	Time	CO ₂	O ₂	CO
H ₂ O	585.6	58.1	37.5		20.15	8.75	
H ₂ O	555.3	545.9	9.4				
Blank	426.4	425.2	1.2				
				Leak Rate	cfm		"Hg
				Initial	0.014		10.0
S.G.	684.2	671.7	12.5	Final			
		Total	60.6				

Sample Point	Time	ΔP in wg	ΔH in wg	Gas Meter Volume Ft ³	Temperature °F				Stack	Probe	Oven		Imp.	Gas Meter		Pump Vacuum in. Hg	$\sqrt{\Delta P}$	Comments
														In	Out			
P-1	0	.24	1.18	715.172	377	266	246	31	63	59						4.0	0.490	pilot OK
2	3	.24	1.18	717.4-	385	261	248	33	72	60						4.0	0.490	
3	6	.21	1.03	719.811	385	261	252	33	73	61						3.5	0.458	
4	9	.21	1.03	721.911	384	256	247	35	74	61						3.5	0.458	
5	12	.23	1.13	724.052	385	247	244	35	75	61						4.0	0.480	
6	15	.20	0.99	726.312	380	248	249	34	75	60						3.5	0.447	
C-1	18	.24	1.18	728.279	376	247	245	33	68	59						4.0	0.490	1156 - "c"
2	21	.21	1.04	730.509	378	251	252	37	73	60						3.5	0.458	
3	24	.18	0.89	732.442	382	261	255	40	75	60						3.5	0.424	
4	27	.18	0.89	734.371	382	261	255	42	76	60						3.5	0.424	
5	30	.15	0.74	736.274	382	257	259	43	76	61						3.0	0.387	
6	33	.12	0.60	738.051	378	263	254	43	76	61						2.5	0.346	
	36			739.672														

515

Nozzle No. / Size	853	3215
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[illegible]

21512

Nozzle No./Size 1853/3217

[illegible]

Date 2-24-87

Test Location #2 K2

Run Number 3

Stack Diameter 16.0Operator owe

Filter No.

Barometric Pressure

Static in. wg.

Probe Type/Length

Pitot Coefficient

Meter Box No. / 8

Nozzle No./Size

Impinger Volumes/Weights				Gas Composition			
Contents	Final	Initial	Net	Time	CO ₂	O ₂	CO
S.G.				Leak Rate		cfm	"Hg
				Initial			
				Final			
Total						0.200	11

Sample Point	1926 Time	ΔP in wg	ΔH in wg	Gas Meter Volume Ft ³	Temperature °F						Pump Vacuum in. Hg	$\sqrt{\Delta P}$	Comments
					Stack	Probe	Oven	Imp.	Gas Meter				
									In	Out			
C-1	0	.18	0.89	807.876	367	258	255	30	59	57	2.5	0.424	
2	3	.18	0.90	809.846	368	256	253	29	69	59	2.5	0.424	
3	6	.18	0.90	811.831	368	248	258	30	72	59	2.5	0.424	
4	9	.15	0.75	813.727	367	241	256	30	73	59	2.0	0.387	
5	12	.12	0.61	815.498	361	244	257	30	72	59	2.0	0.346	
6	15	.11	0.55	817.148	360	245	259	30	72	58	1.5	0.332	
A-1	16	.21	1.04	818.682	360	240	252	29	59	56	3.0	0.458	
2	21	.21	1.05	820.711	362	249	259	26	70	57	3.0	0.458	
3	24	.18	0.91	822.891	358	251	257	26	72	57	3.0	0.424	
4	27	.20	1.01	824.727	358	255	251	27	72	58	3.5	0.447	
5	30	.20	1.01	826.809	358	256	254	27	73	58	3.5	0.447	
10	33	.18	0.92	828.818	354	253	256	28	73	58	3.0	0.424	
	36			830.573									
	T/A		1.45	56.8160	366.88				63.96				0.5250

273.957

CONTINUOUS MONITOR DATA SHEET

Plant SWPC Kiln 2
 Date 2-24-87 Run No. 1,2
 Test Location Baghouse
 Operator JP
 Fuel Type Coal Trailer No. 2

APCD Witness/Number
 Generator Type Kiln 2
 Burner Type
 O₂ Controller Type
 Gas Cylinder Nos. NOx 13472, SO₂ 13435, O₂ 13460 584

Time	Sample Point	Fuel Flow	Dry Uncorrected					3% O ₂ Dry		Hydrocarbons			Comments
			O ₂ %	CO ₂ %	CO ppm	SO ₂ ppm	NO ppm	NO _x ppm	ppm	NO _x ppm	THC	Non CH ₄	
0930	Initial Coal		8.06	15.03	40.1	51.9		807.8					
			0			50.5		600					
			20.7										
0955			9.6	19.2	37	2		640					Extend response
1005			9.0	20.0	490	41		640					Air
1015			8.8	20.5	550	2		678					Test 1
1025			8.7	20.7	650	2		660					TPH Feed 206
1035			8.7	20.6	550	2		640					f Coal
1045			8.8	20.3	450	2		630					
1055	Coal Check		-0.1	0	-2	0		-5					Zero
			8.3	20.1	374	53		804					Spun
			14.75										Reset after lock
1205			8.8	20.0	400	2		640					Test 2
1215			8.8	19.8	490	2		600					
1225			8.7	20.3	475	2		640					
1235			8.8	20.3	475	2		625					
1245			8.7	20.3	500	2		605					
1255			8.7	20.2	475	3		615					
1315	Coal Check		0	-0.1	0	0		-5					Zero
			7.9	14.7	400	52		930					Spun
			8.93	20.2	500	1.83		650.3					
			8.75	20.5	425	2.17		610.8					

CONTINUOUS MONITOR DATA SHEET

Plant	SWPC		
Date	2-24-87	Run No.	3
Test Location	Kiln 2 Baghouse		
Operator	RP		
Fuel Type	Coal	Trailer No.	2
APCD Witness/Number			
Generator Type			
Burner Type			
O ₂ Controller Type			
Gas Cylinder Nos.			

[illegible]

SAMPLE HANDLING/LOG-IN

Date 2-24-87 Test Location Suppc Kila 2

	Sample Type	Volume	Comments
1.	17794 <u>5</u> <u>FHA</u> <u>1</u> Meth Sample Test	90	
2.	17795 <u>5</u> <u>MF</u> <u>1</u> Meth Sample Test	N/A	
3.	17796 <u>5</u> <u>BHW</u> <u>1</u> Meth Sample Test	390	
4.	17797 <u>5</u> <u>FHA</u> <u>2</u> Meth Sample Test	79	
5.	17798 <u>5</u> <u>MF</u> <u>2</u> Meth Sample Test	N/A	
6.	17799 <u>5</u> <u>BHW</u> <u>2</u> Meth Sample Test	318	
7.	17800 <u>5</u> <u>FHA</u> <u>3</u> Meth Sample Test	168	
8.	17801 <u>5</u> <u>MF</u> <u>3</u> Meth Sample Test	N/A	
9.	17802 <u>5</u> <u>BHW</u> <u>3</u> Meth Sample Test	405	
10.	17803 <u>5</u> <u>FHA</u> <u>Blank</u> Meth Sample Test	100	
11.	17805 <u>5</u> <u>BHW</u> <u>Blank</u> Meth Sample Test	N/A	
12.	17804 <u>5</u> <u>MF</u> <u>Blank</u> Meth Sample Test	100	

CHAIN OF CUSTODY



JP 3/2

ANALYTICAL REPORT

SAMPLE TYPE Front Half Wash
 SAMPLE COMPONENT Aceton
 REQUESTED BY SWPC
 ANALYTICAL METHOD Gravimetric

DATE 3/4/87
 ANALYST JP

Sample ID No.	Test No. #	Sample Volume mls	Sample Aliquot	Titer mls or (Absorbance)	Analytical Result (total sample) gms	
					Uncorrected	Blank corrected
	1	90			0.00801	0.00750 ✓
	2	79			0.00583	0.00538 ✓
	3	168			0.00874	0.00778 ✓
	BL	100			0.00057	

JP

ANALYTICAL REPORT

SAMPLE TYPE Filterable Particulate
 SAMPLE COMPONENT 100mm Filter
 REQUESTED BY SWCP
 ANALYTICAL METHOD Gravimetric

DATE 3/4/87
 ANALYST JP

Sample ID No. <i>Filter #</i>	Test No. <i>#</i>	Sample Volume	Sample Aliquot	Titer mls or (Absorbance)	Analytical Result (total sample) <i>gms</i>	
					Uncorrected	Blank corrected
10-814	1				0.00050	0.00051
10-815	2				0.00174	0.00175
10-816	3				0.00136	0.00137
10-817	Bl				-0.00001	

ANALYTICAL REPORT

SAMPLE TYPE Condensable Particulates
 SAMPLE COMPONENT _____
 REQUESTED BY SWPC
 ANALYTICAL METHOD Gravimetric

DATE 3/4/87
 ANALYST JP

Sample ID No.	Test No. #	Sample Volume mls	Sample Aliquot mls	Titer mls or (Absorbance)	Analytical Result (total sample) gmo.	
					Uncorrected	Blank corrected
Analysis in Tin	1	390	100		0.01928	0.07504 ✓
	2	318	100		0.01931	0.06128 ✓
	3	405	100		0.00984	0.03969 ✓
	Bl	100	100		0.00004	
Analysis in Glass	1	390				.0041
	2	318				.0034
	3	405				.0043
						<u>.0118</u>

ISOKINETIC CALCULATIONS

Standard temperature, S_t 60 (XEQ 60 or 68); 29.92 inches Hg

Test 1 Date 2-24-87 Location SWPC Kiln 2 Baghouse

T_m <u>64.52</u>	V_m <u>55.957</u>	O_2 <u>8.9</u>	Stack ID " <u>160</u>
T_s <u>373.79</u>	γ <u>.9983</u>	CO_2 <u>20.2</u>	Stack Area Ft^2 <u>139.63</u>
$\sqrt{\Delta P}$ <u>.5240</u>	P_b <u>26.08</u>	Static <u>-.18</u>	Test Time <u>72</u>
ΔH <u>1.40</u>	V_{1c} <u>58.6</u>	C_p <u>.84</u>	N_d <u>.3217</u>

1. Volume of dry gas sampled at standard conditions (dscf)

(R-17)

$$V_{m, std} = \left(\frac{S_t + 460}{29.92} \right) (\gamma) V_m \left(\frac{P_b + \frac{\Delta H}{13.6}}{T_m + 460} \right) = 48.4632$$

2. Stack gas moisture condensed at standard conditions (scf)

(R-18)

$$(V_{m, std} + V_{w, std}; R-04) \quad V_{w, std} = 8.9148 \times 10^{-5} (S_t + 460) (V_{1c}) = 2.7165$$

3. Stack gas proportion of water vapor, by volume

(R-19)

$$B_{wo} = \frac{V_{w, std}}{V_{w, std} + V_{m, std}} = .0531$$

4. Stack gas dry molecular weight (lb/lb mole)

(R-21)

$$MW_d = 0.44(\%CO_2) + 0.32(\%O_2) + 0.28(\%N_2 + \%CO) = 31.59$$

5. Stack gas molecular weight (lb/lb mole)

(R-22)

$$MW_s = MW_d (1 - B_{wo}) + 18(B_{wo}) = 30.87$$

6. Pressure stack, in. Hg

(R-23)

$$P_s = P_b + \left(\frac{P_{static}}{13.6} \right) = 26.07$$

7. Stack gas velocity, at stack conditions (ft/sec)

(R-24)

$$V_s = 85.49(C_p)(\sqrt{\Delta P})_{avg} \sqrt{\frac{T_s + 460}{P_s(MW_s)}} = 38.31$$

8. Stack gas volume at standard conditions (dscfm)

(R-26)

(ACFM; R-25)

320,910.27

$$Q_s = 60(1 - B_{wo}) V_s A_s \left(\frac{S_t + 460}{T_s + 460} \right) \left(\frac{P_s}{29.92} \right) = 165,108.21$$

9. Test percent isokinetic

(R-02)

$$\%I = \frac{9142.88(T_s + 460)(V_{w, std} + V_{m, std})}{(S_t + 460) \theta V_s P_s (D_n)^2} = 100.92$$

ISOKINETIC CALCULATIONS

Standard temperature, S_t 60 (XEQ 60 or 68); 29.92 inches Hg

Test 2 Date 2/24/87 Location SWPC Kiln 2

T_m <u>67.44</u>	V_m <u>58.235</u>	O_2 <u>8.75</u>	Stack ID " <u>160</u>
T_s <u>379.79</u>	δ <u>.9983</u>	CO_2 <u>20.15</u>	Stack Area Ft^2 <u>139.63</u>
$\sqrt{\Delta P}$ <u>.5412</u>	P_b <u>26.04</u>	Static <u>-.18</u>	Test Time <u>72</u>
ΔH <u>1.50</u>	V_{1c} <u>60.6</u>	C_p <u>.87</u>	N_d <u>.3217</u>

1. Volume of dry gas sampled at standard conditions (dscf)

(R-17)

$$V_{m,std} = \left(\frac{S_t + 460}{29.92} \right) (\delta) V_m \left(\frac{P_b + \frac{\Delta H}{13.6}}{T_m + 460} \right) = 50.0944$$

2. Stack gas moisture condensed at standard conditions (scf)

(R-18)

$$(V_{m,std} + V_{w,std}; R-04) \quad V_{w,std} = 8.9148 \times 10^{-5} (S_t + 460) (V_{1c}) = 2.8092$$

3. Stack gas proportion of water vapor, by volume

(R-19)

$$B_{wo} = \frac{V_{w,std}}{V_{w,std} + V_{m,std}} = .0531$$

4. Stack gas dry molecular weight (lb/lb mole)

(R-21)

$$MW_d = 0.44(\%CO_2) + 0.32(\%O_2) + 0.28(\%N_2 + \%CO) = 31.57$$

5. Stack gas molecular weight (lb/lb mole)

(R-22)

$$MW_s = MW_d (1 - B_{wo}) + 18(B_{wo}) = 30.85$$

6. Pressure stack, in. Hg

(R-23)

$$P_s = P_b + \left(\frac{P_{static}}{13.6} \right) = 26.03$$

7. Stack gas velocity, at stack conditions (ft/sec)

(R-24)

$$V_s = 85.49(C_p)(\sqrt{\Delta P})_{avg} \sqrt{\frac{T_s + 460}{P_s(MW_s)}} = 39.74$$

8. Stack gas volume at standard conditions (dscfm)

(R-26)

(ACFM; R-25)

332963.08

$$Q_s = 60(1 - B_{wo}) V_s A_s \left(\frac{S_t + 460}{T_s + 460} \right) \left(\frac{P_s}{29.92} \right) = 169.820.31$$

9. Test percent isokinetic

(R-02)

$$\%I = \frac{9142.88(T_s + 460)(V_{w,std} + V_{m,std})}{(S_t + 460) \theta V_s P_s (D_n)^2} = 101.43$$

ISOKINETIC CALCULATIONS

Standard temperature, S_t 60 (XEQ 60 or 68); 29.92 inches Hg

Test 3 Date 2/24/87 Location SWPC Rtn 2

T_m <u>63.96</u>	V_m <u>56.816</u>	O_2 <u>8.80</u>	Stack ID " <u>160</u>
T_s <u>346.88</u>	γ <u>.9983</u>	CO_2 <u>20.58</u>	Stack Area Ft^2 <u>139.63</u>
$\sqrt{\Delta P}$ <u>.5256</u>	P_b <u>26.00</u>	Static <u>-1.8</u>	Test Time <u>72</u>
ΔH <u>1.45</u>	V_{1c} <u>60.7</u>	C_p <u>.84</u>	N_d <u>.3217</u>

1. Volume of dry gas sampled at standard conditions (dscf)

(R-17)

$$V_{m,std} = \left(\frac{S_t + 460}{29.92} \right) (\gamma) V_m \left(\frac{P_b + \frac{\Delta H}{13.6}}{T_m + 460} \right) = 49.1162$$

2. Stack gas moisture condensed at standard conditions (scf)

(R-18)

$$(V_{m,std} + V_{w,std}; R-04) \quad V_{w,std} = 8.9148 \times 10^{-5} (S_t + 460) (V_{1c}) = 2.8139$$

3. Stack gas proportion of water vapor, by volume

(R-19)

$$B_{wo} = \frac{V_{w,std}}{V_{w,std} + V_{m,std}} = .0542$$

4. Stack gas dry molecular weight (lb/lb mole)

(R-21)

$$MW_d = 0.44(\%CO_2) + 0.32(\%O_2) + 0.28(\%N_2 + \%CO) = 31.64$$

5. Stack gas molecular weight (lb/lb mole)

(R-22)

$$MW_s = MW_d (1 - B_{wo}) + 18(B_{wo}) = 30.91$$

6. Pressure stack, in. Hg

(R-23)

$$P_s = P_b + \left(\frac{P_{static}}{13.6} \right) = 25.99$$

7. Stack gas velocity, at stack conditions (ft/sec)

(R-24)

$$V_s = 85.49(C_p)(\sqrt{\Delta P})_{avg} \sqrt{\frac{T_s + 460}{P_s(MW_s)}} = 38.30$$

8. Stack gas volume at standard conditions (dscfm)

(R-26)

(ACFM; R-25)

$$Q_s = 60(1 - B_{wo}) V_s A_s \left(\frac{S_t + 460}{T_s + 460} \right) \left(\frac{P_s}{29.92} \right) = 165,749.47$$

9. Test percent isokinetic

(R-02)

$$\%I = \frac{9142.88(T_s + 460)(V_{w,std} + V_{m,std})}{(S_t + 460) \theta V_s P_s (D_n)^2} = 101.89$$

EMISSION RATE CALCULATIONS

1. Fuel factor at 68°F, 29.92 in. Hg (dscf/mmBtu)

$$F \text{ Factor} = \frac{10^6 [3.64(\%H) + 1.53(\%C) + 0.57(\%S) + 0.14(\%N) - 0.46(\%O_2)]}{Btu/lb}$$

2. Fuel factor at 60°F, 29.92 in. Hg (dscf/mmBtu)

$$F \text{ Factor}_{600} = F \text{ Factor}_{680} \left(\frac{60 + 460}{68 + 460} \right)$$

3. Emission concentration wet at standard conditions* (gr/scf)

$$gr/scf = 15.432 \left(\frac{Comp (g)}{V_{m,std} + V_{w,std}} \right)$$

4. Emission concentration dry at standard conditions* (gr/dscf)

$$gr/dscf = 15.432 \left(\frac{Comp (g)}{V_{m,std}} \right)$$

5. Emission concentration corrected to 12% CO₂

$$C @ 12\% CO_2 = C_{measured} \left(\frac{12}{\%CO_2 \text{ measured}} \right)$$

6. Emission Rate lb/hr from gr/dscf

$$lb/hr = 0.00857(gr/dscf)(Q_s)$$

7. Gaseous concentration in ppm

$$ppm = \frac{1.60864(S_t + 460)*(mg \text{ comp})}{MW(V_{m,std})}$$

8. Emission rate lb/hr from ppm

$$lb/hr = \frac{8.223 \times 10^{-5}(Q_s)(MW)(ppm)}{(S_t + 460)*}$$

9. Emission concentration corrected to 3% O₂

$$ppm @ 3\% O_2 = ppm \text{ measured} \left(\frac{20.9 - 3}{20.9 - \%O_2 \text{ measured}} \right)$$

EMISSION RATE CALCULATIONS (Concluded)

10. Emission factor, lb/mmBtu at 0% O₂

$$\text{lb/mmBtu} = F(1.4286 \times 10^{-4}) \left(\frac{20.9}{20.9 - \%O_2} \right) (\text{gr/dscf})$$

$$\text{lb/mmBtu} = F(\text{MW}) \left(\frac{1.3711 \times 10^{-6}}{S_t + 460^*} \right) \left(\frac{20.9}{20.9 - \%O_2} \right) (\text{ppm})$$

11. Heating value of fuel, mmBtu/bbl

$$\text{mmBtu/bbl} = 349.786 \times 10^{-6} (\text{Btu/lb})(\text{SpGr})$$

12. Emission Factor, lb/bbl

$$\text{lb/bbl} = (\text{lb/mmBtu})(\text{mmBtu/bbl})$$

13. Brake Specific Emission Rate = grams per brake horsepower hour

$$= (\text{lb/MMbtu})(453.592 \times 10^{-6})(\text{Btu/brake horsepower hour})$$

14. Maximum Mass Emission Rate (lb/hr)

$$= (\text{grams/BHP hr})(\text{maximum BHP})(1 \text{ lb}/453.592 \text{ g})$$

15. Kilograms per thousand Barrels of Feed (Kg/Kbbl)

$$= \frac{\text{lb/hr emission (10.8864)}}{\text{KBPD Feed}}$$

*Standard temperature to be defined

EMISSION RATE DATA

Standard Temperature, S_e 60 (60°F or 68°F); 29.92 inches Hg

LOCATION

SWPC Kln 2 baghouse

DATA	Test 1	Test 2	Test 3	Average
$V_{m, std}$	48.4632	50.0944	49.1162	49.2246
Bwo	.0531	.0531	.0542	.0535
Qs	165108.21	169820.31	165749.47	166892.66
O ₂	8.93	8.75	8.80	8.83
CO ₂	20.20	20.15	20.58	20.31
F-Factor				

LAB DATA

Front Half Wash	g	.00750 ^{.00801}	.00538 ^{.00713}	.00778 ^{.00915}	
Mass Filter	g	.00051	.00175	.00137	
Back Half Catch	g	.0041	.0034	.0043	
Front Half Sulfate, mg H ₂ SO ₄					
Back Half Sulfate, mg H ₂ SO ₄					
H ₂ O ₂ Catch	mg H ₂ SO ₄				

RESULTS

Filt. Particulate	gr/dscf	.0026	.0022	.0029	.0026
	gr/dscf @ 12% CO ₂	.0015	.0013	.0017	.0015
	gr/scf	.0024	.0021	.0027	.0024
	lb/hr	3.61	3.20	4.08	3.63
	lb/MMBtu				

Total Particulate	gr/dscf	.0039	.0032	.0042	.0038
	gr/dscf @ 12% CO ₂	.0023	.0019	.0025	.0022
	gr/scf	.0037	.0031	.0040	.0036
	lb/hr	5.46	4.72	6.00	5.39
	lb/MMBtu				

Total Sulfate	gr/dscf				
	gr/dscf @ 12% CO ₂				
	gr/scf				
	lb/hr				
	lb/MMBtu				
	lb/MMBtu as Sulfur				

SO ₃	ppm				
	ppm, wet				
	lb/hr				

SO ₂	ppm				
	ppm, wet				
	lb/hr				
	lb/MMBtu				
	lb/MMBtu, as Sulfur				

TEST SWPC
LOCATION Kiln 2

CM SO₂, NO_x, HC, CO EMISSION RATE DATA

Standard Temperature, S_t 60 (XEQ 60 or 68); 29.92 inches Hg

ENTER			Test 1	Test 2	Test 3	Average
R-22	O ₂	%	<u>8.93</u>	<u>8.75</u>	<u>8.80</u>	<u>8.83</u>
R-26	Q _s		<u>165108.21</u>	<u>169820.31</u>	<u>165749.47</u>	<u>166892.66</u>
R-15	NO _x	ppm	<u>650.3</u>	<u>610.8</u>	<u>540.8</u>	<u>600.63</u>
R-15	SO ₂	ppm	<u>1.83</u>	<u>2.17</u>	<u>2.50</u>	<u>2.17</u>
R-15	CO	ppm	<u>500.0</u>	<u>462.5</u>	<u>533.3</u>	<u>498.6</u>
R-15	HC	ppm				

XEQ FF

XEQ SO₂

F-FACTOR

RESULTS:

MW = 46

NO _x	lb/hr	<u>781.04</u>	<u>754.53</u>	<u>652.04</u>	<u>729.20</u>
NO _x	ppm @ 3% O ₂	<u>972.46</u>	<u>899.86</u>	<u>800.03</u>	<u>890.78</u>
NO _x	lb/MMBtu				
NO _x	lb/Bbl				

MW = 64.1

SO ₂	lb/hr	<u>3.06</u>	<u>3.74</u>	<u>4.20</u>	<u>3.67</u>
SO ₂	ppm @ 3% O ₂				
SO ₂	lb/MMBtu				
SO ₂	lb/Bbl				

MW = 28

CO	lb/hr	<u>365.53</u>	<u>347.77</u>	<u>391.39</u>	<u>369.23</u>
CO	ppm @ 3% O ₂				
CO	lb/MMBtu				
CO	lb/Bbl				

MW = 16

HC	lb/hr				
HC	ppm @ 3% O ₂				
HC	lb/MMBtu				
HC	lb/Bbl				

DRY GAS METER/ORIFICE METER CALIBRATION DATA

Date 2/6/87 Dry Gas Meter No. 56001
 Barometric Pressure, in. Hg 29.80 Standard Test Meter No. 69279
 Meter Box No. 310609 Operator CARTER

Standard Test Meter			Meter Box			Time (min)	γ	K_o
Pres. ΔH_s ("Wg)	Temp. T_s (°F)	Volume V_s (Ft ³)	Pres. ΔH ("Wg)	Temp. T_{di} T_{do} (°F)	Volume V_d (Ft ³)			
	75	119.299		108.92	952.152			
-2.3	72	97.637	0.5	101.90	929.459	46	0.9911	0.8554
	73.5	21.662		97.75	22.693			
	73	95.940		109.88	927.684			
-3.9	73	73.1663	1.0	102.82	904.556	34	0.9913	0.8339
	73	22.277		95.25	23.128			
	75	147.675		115.93	975.343			
-5.2	75	120.856	1.5	112.92	953.751	26	0.9980	0.8319
	75	20.819		103	21.592			
	75	167.883		118.94	1002.284			
-6.6	75	144.803	2.0	117.93	978.553	25	1.0063	0.8280
	75	23.080		105.5	23.731			
	75	189.609		119.94	24.483			
-9.1	75	169.401	3.0	117.94	3.835	18	1.0047	0.8163
	75	20.208		106.0	20.648			
			4.0					

Average 0.9983 0.8331
 Std.Dev. 0.0072 0.0142
 (0.7%) (1.7%)

$$\gamma = \frac{V_s \left(P_{\text{bar}} + \frac{\Delta H_s}{13.6} \right) \left(\frac{T_{di} + T_{do}}{2} + 460 \right)}{V_d \left(P_{\text{bar}} + \frac{\Delta H}{13.6} \right) (T_s + 460)}$$

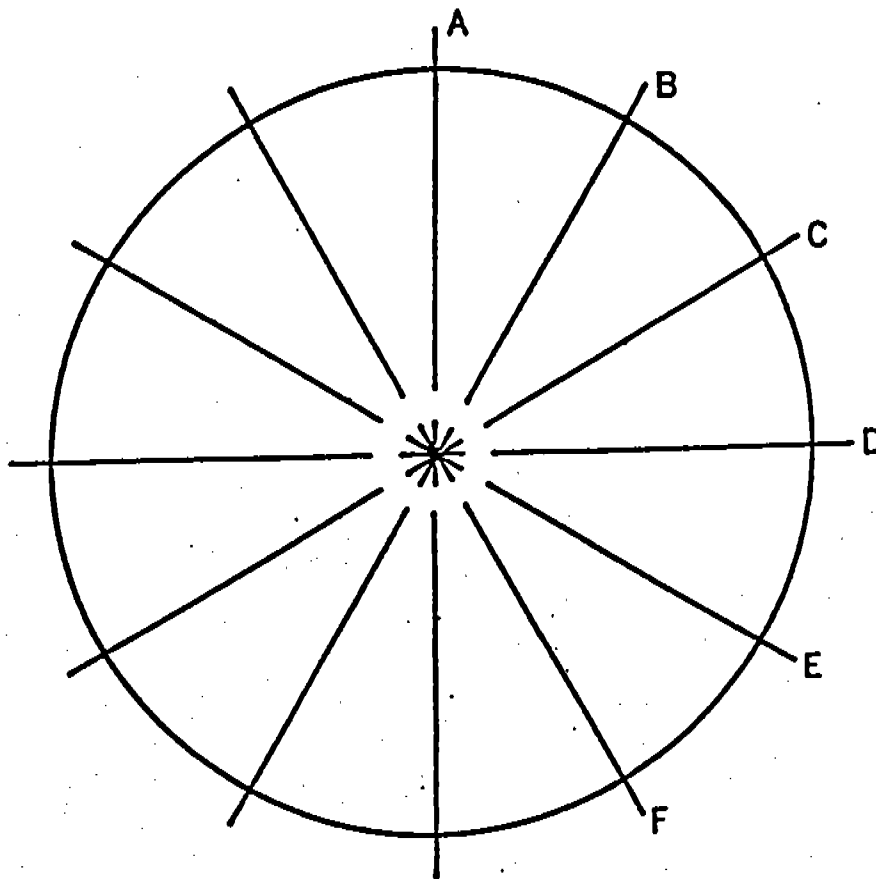
$$K_o = \frac{\left(\frac{V_s}{t} \right) \left(\frac{T_{do}}{T_s} \right) \left(\frac{P_{\text{bar}} + \frac{\Delta H_s}{13.6}}{P_{\text{bar}} + \frac{\Delta H}{13.6}} \right)}{\left(\frac{T_{do} \Delta H}{\left(P_{\text{bar}} + \frac{\Delta H}{13.6} \right) (M_m)} \right)^{0.5}}$$

where

ΔH_s = Standard meter pressure drop
 T_s = Standard meter temperature
 V_s = Standard test meter dry gas volume
 ΔH = Meter box orifice meter pressure drop
 T_{di} = Meter box inlet temperature
 T_{do} = Meter box outlet temperature
 V_d = Meter box dry gas volume

γ = Meter box correction factor
 K_o = Orifice meter calibration constant
 t = Calibration time in minutes
 M_m = Molecular weight of air (28.96 lb/lb-mole)
 P_{bar} = Barometric pressure

NOZZLE CALIBRATION DATA



DIAMETER (in.)

A .3205

B _____

C .3225

D .3218

E .3208

F _____

AVG. .3217

DATE 2-19-86

NOZZLE NO. 18153

OPERATOR JCO

2600 CAJON BLVD. SAN BERNARDINO, CA 92411 PHONE (714) 887-2571

PAPE & STEINER
5801 Norris Road
Bakersfield, CA 93308
Attn: Sue Powers

Date Shipped 1-16-87
Our Project No: 404936
Your P.O. No: SP 8864
Page 1 of 1

CERTIFICATE OF ANALYSIS - EPA PROTOCOL GASES*
(Concentrations are in mole % or ppm)

Cylinder Number AAL13472 Certified Accuracy ±1 % NBS Traceable Analysis Dates: First 01-06-87 Last 01-16-87

COMPONENTS	CERTIFIED CONC	EXPIRATION DATE	ANALYTICAL PRINCIPLE	PRIMARY STANDARD NBS/SRM's	REPLICATE CONCENTRATIONS	
					FIRST	SECOND
NITRIC OXIDE	807.8 ppm	07-16-87	CHEM.	16878	810.6 ppm	807.8 ppm
NOX	813.1 ppm				813.6 ppm	813.1 ppm
NITROGEN	BALANCE					

A-28

COMPONENTS	CERTIFIED CONC	EXPIRATION DATE	ANALYTICAL PRINCIPLE	PRIMARY STANDARD NBS/SRM's	REPLICATE CONCENTRATIONS	
					FIRST	SECOND

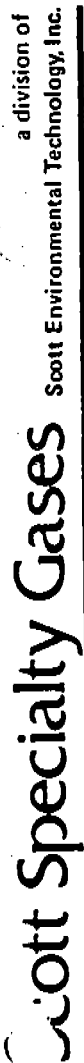
*We hereby certify the cylinder gas has been analyzed according to EPA Protocol No: 1

Analyst _____ Approved By *[Signature]*

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 / PLUMSTEADVILLE, PENNSYLVANIA



PHONE (714) 887-2571

Attn: Sue Powers

Page 3 of 7

(Concentrations are in mole % or ppm)

TROY, MICHIGAN / PLUMSTEADVILLE, PENNSYLVANIA



Scott Specialty Gases

a division of
Scott Environmental Technology, Inc.

PLUMSTEADVILLE, PA. 18949 PHONE: (215) 766-8861 TWX: 510-665-9344
SAN BERNARDINO, CA 714-887-2571

PAPE & STEINER
5801 Norris Road
Bakersfield, CA 93308
Attn: Linda

Date: 9-12-86
Our Project No.: 403673
Your P.O. No.: SP5203

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. AAL584 Analytical Accuracy ±1%
Component Concentration

CARBON MONOXIDE 400.1ppm

NBS TRACEABLE TO SRM 1680B

CARBON DIOXIDE 15.03%

OXYGEN 8.060%

NITROGEN BALANCE

Cyl. No. _____ Analytical Accuracy _____
Component Concentration

Cyl. No. _____ Analytical Accuracy _____
Component Concentration

Cyl. No. _____ Analytical Accuracy _____
Component Concentration

Analyst _____

Approved By D. J. Smith

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 REF A-30 ROTOCOL GASES
ACUBLEND® CALIBRATION & SPECIALTY GAS MIXTURES PURE GASES
ACCESSORY PRODUCTS CUSTOM ANALYTICAL SERVICES
TROY, MICHIGAN / SAN BERNARDINO, CALIFORNIA / HOUSTON, TEXAS

FINAL DATA RUN SWPC

QUARRY
PCT

KILN
#2

87 FEB 24 15:01

HOURLY AVERAGE

TIME	FIC-309	FIC-471	FIC-472	SIC-337	J1-313	SI-371	J1-314	T1-450
Hour	K2 FEED	K2 COAL	PYROCOAL	PREHEATR	PREHEATR	EXHT FAN	EXHT FAN	EXHT FAN
ENDING	TPH	TPH	TPH	ID FAN	ID FAN	EXHT FAN	EXHT FAN	EXHT FAN
15:01	205.0	7.400	7.200	747.0	1092.	693.0	674.7	2400.
14:01	217.5	7.599	7.799	783.0	1222.	711.0	709.3	2560.
13:01	215.0	7.699	7.799	801.0	1300.	711.0	726.6	2620.
12:01	212.5	7.699	7.900	809.9	1352.	720.0	743.9	2560.
11:01	212.5	7.599	8.100	792.0	1274.	702.0	709.3	2580.
10:01	172.5	7.599	5.900	711.0	961.9	693.0	657.3	2680.
09:01	167.5	7.000	6.099	711.0	936.0	693.0	640.1	2340.
08:01	177.5	6.300	6.199	702.0	909.9	693.0	640.1	2460.
07:01	192.5	7.500	6.600	747.0	1066.	702.0	674.7	2600.
06:01	190.0	7.599	6.499	737.9	1040.	693.0	657.3	2600.
05:01	192.5	7.599	6.700	747.0	1066.	702.0	674.7	2620.
04:01	180.0	7.599	6.199	693.0	883.9	675.0	622.8	2640.
03:01	175.0	7.599	6.000	684.0	892.0	684.0	622.8	2620.
02:01	200.0	7.599	7.100	747.0	1092.	702.0	674.7	2380.
01:01	217.5	7.599	7.900	783.0	1222.	702.0	692.0	2560.
00:01	217.5	7.599	8.100	809.9	1323.	711.0	709.3	2580.
23:01	215.0	7.500	8.100	809.9	1326.	720.0	726.6	2620.
22:01	215.0	7.500	8.100	809.9	1326.	720.0	726.6	2700.
21:01	210.0	7.500	8.000	809.9	1300.	720.0	726.6	2580.
20:01	217.5	7.400	8.100	809.9	1300.	711.0	709.3	2560.
19:01	217.5	7.400	8.000	809.9	1300.	711.0	709.3	2620.
18:01	217.5	7.400	8.000	809.9	1300.	711.0	709.3	2680.
17:01	217.5	7.400	8.000	809.9	1300.	711.0	692.0	2700.
16:01	215.0	7.400	7.900	783.0	1196.	702.0	692.0	2640.