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Segundo Generating Stating Unit 4

Carnot

Carnot

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Section 1.4
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**UREA PERMIT COMPLIANCE TESTING AT
EL SEGUNDO GENERATING STATION UNIT 4**

Prepared For:

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For Submittal To:

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SEPTEMBER 1993

REVIEW AND CERTIFICATION

All work, calculations, and other activities and tasks performed and documented in this report were carried out under my direction and supervision.

Hung M. Ha

Date 9/20/93

FOR Edward J. Filadelfia
Senior Engineer

I have reviewed, technically and editorially, details, calculations, results, conclusions and other appropriate written material contained herein, and hereby certify that the presented material is authentic and accurate.

Hung M. Ha

Date 9/20/93

FOR Edward J. Filadelfia
Senior Engineer

TABLE OF CONTENTS

<u>SECTION</u>	<u>PAGE</u>
1.0 INTRODUCTION	1
2.0 UNIT DESCRIPTION	3
3.0 TEST DESCRIPTION	4
3.1 TEST CONDITIONS	4
3.2 SAMPLE LOCATIONS	4
3.3 TEST PROCEDURES	4
3.3.1 Gaseous Emissions	4
3.3.2 Reactive Organic Gases (ROG) as Hydrocarbons	9
3.3.3 Ammonia	9
3.3.4 Moisture	9
4.0 RESULTS	10
 <u>APPENDICES</u>	
A MEASUREMENT PROCEDURES	A-1
B QUALITY ASSURANCE	B-1
B.1 Quality Assurance Program Summary	B-2
B.2 ARB Certification	B-7
B.3 Equipment Calibration	B-8
C DATA SHEETS	C-1
C.1 Sample Locations	C-2
C.2 Unit Data	C-3
C.3 CEM Data	C-4
C.4 Ammonia	C-5
C.5 Non-Methane Hydrocarbon	C-6
D CALCULATIONS	D-1
D.1 General Emissions Calculations	D-2
D.2 Specific Emissions Calculations	D-3
E LABORATORY REPORTS	E-1
F INSTRUMENT STRIP CHARTS	F-1
G CHAIN OF CUSTODY	G-1

SECTION 1.0

INTRODUCTION

Carnot was contracted to conduct a series of emissions tests on Unit 4 at the El Segundo Generating Station for Southern California Edison (SCE). Carnot conducted testing to comply with South Coast Air Quality Management District (SCAQMD) Permit To Construct for the urea injection system.

A test program was conducted to quantify the effect of urea injection on various stack emissions as stated in Conditions 11-B and 11-C(b) of the Permit to Construct. Emissions tests were performed on El Segundo Unit 4 for the following:

- NO_x, CO and O₂
- ROG as non-methane Hydrocarbons
- NH₃ (urea on only)
- Moisture

Table 1-1 summarizes the emissions test results.

Per verbal discussion with Mr. Darren Stroud of the SCAQMD, it was agreed that particulate emissions would be measured on one unit of each unit class because of the low potential for urea having an impact on particulate emissions. Therefore, particulates were not measured on this unit. The unit that was tested as representative of the 320/335 MW class of units was Alamitos No. 4. The results are contained in Carnot Report No. SCE1A-10840/R106B956.T and titled Urea Permit Compliance Testing at Alamitos Generating Station Unit 4.

The 256 MW and 150 MW load emissions testing was conducted by Matt McCune, Wayne Madden and Marc Rodabaugh of Carnot on May 24-25, 1993. Mike Escarcega of SCE coordinated the testing. Full load emission testing was conducted by Edward J. Filadelfia, Jim Mulligan and Marc Rodabaugh of Carnot on August 12, 1993. Larry Milleson of SCE coordinated the testing.

TABLE 1-1
EMISSIONS TEST RESULTS SUMMARY
EL SEGUNDO UNIT 4
MAY 1993

	Net Load 331 MW		Net Load 256 MW		Net Load 150 MW	
	Urea Off	Urea On	Urea Off	Urea On	Urea Off	Urea On
NO _x , ppm	73.5	65.8	67.0	55.3	43.9	39.3
NO _x , ppm @ 3% O ₂	83.5 16%	75.0	74.4 17%	61.8	53.0 11%	47.3
CO, ppm	6.5	4.7	23.4	18.5	0.0	0.0
CO, ppm @ 3% O ₂	7.4 28%	5.3	26.0 70%	20.7	0.0	0.0
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0	ND < 1.0	ND < 1.0	ND < 1.0
TNMHC as CH ₄ , ppm	8.6	8.6	20.8	21.4	17.4	17.6
Urea flow, gpm	0.0	2.2	0.0	1.3	0.0	0.7
NH ₃ slip, ppm	--	6.2	--	10.6	--	4.0
ppm @ 3% O ₂	--	7.1	--	11.9	--	4.8

Note: ND indicates that this species was not detected.

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SECTION 2.0

UNIT DESCRIPTION

El Segundo Generating Station Unit 4 is one of six similar Combustion Engineering inverted tangentially-fired boilers in the SCE system with a rated load of 335 net MW. The boiler is fired from the top and consists of three tiers of burners with eight burners in each tier. Each burner contains one oil gun and two gas injectors. The unit is divided into two furnaces by a division wall, with burners in each of the four corners to provide the "tangential" firing.

The boiler is equipped with burner tilts, which can be varied from the horizontal to 10 degrees up and 30 degrees down. All three burners in a furnace corner are controlled by one mechanism, with all the burners in a furnace set at the same position. Combustion air is supplied by two forced draft fans through primary and secondary air ducts. The primary air ducts supply air only to the immediate vicinity of the burners. The secondary air is introduced above and below each burner. Air flow to individual burners is controlled by dampers. Existing controls regulate the position of all the primary and secondary dampers separately. The unit has two flue gas recirculation (FGR) fans that mix flue gas with the secondary air supply. The furnace is operated under positive pressure.

SECTION 3.0

TEST DESCRIPTION

3.1 TEST CONDITIONS

The testing was conducted at two conditions and at three loads. Condition 1 was with urea off and Condition 2 was with optimum urea injection. Duplicate tests were performed at the approximate net loads: 256 and 150 MW. Loads below 150 MW were not tested as urea is first used at 150 MW. Due to a load limit (on the unit) imposed by operations, tests at full load were not performed until August 12, 1993. All testing was performed with the boiler fired on natural gas fuel. The test matrix and unit operating data are outlined in Tables 3-1 and 3-2. Unit operation was established by SCE operators, and unit data were collected by SCE personnel.

3.2 SAMPLE LOCATIONS

Measurements were made using two of four sample ports accessible from the newly constructed stack sampling platform. A stack configuration drawing is presented in Figure 3-1. The stack diameter at the cross-sectional plane of the sample ports is 21 feet, 1 inch. This sample location is 62 feet, 2 inches (2.9 diameters) from the nearest upstream flow disturbance and 117 feet, 10 inches (5.6 diameters) from the stack exit.

3.3 TEST PROCEDURES

The test procedures used for the program are presented in Table 3-3. Descriptions of standard procedures are included in Appendix A. Additional information and modifications to standard procedures are presented in the following section.

3.3.1 Gaseous Emissions

Gaseous species (NO_x , CO and O_2) were measured using Carnot's Continuous Emissions Monitor (CEM) described in Appendix A. This system meets SCAQMD, CARB and EPA requirements for gaseous species. Two 30-minute gaseous emission test runs were performed for each test condition and load.

TABLE 3-1
TEST MATRIX
SCE EL SEGUNDO UNIT 4
MAY AND AUGUST 1993

Test #	Date	Time	Type of Test	Net Load MW
4-1-CEM	5/24/93	1000/1045	NO _x , CO & O ₂	150
4-1-NH ₃	5/24/93	1000/1030	Ammonia	150
4-2-CEM	5/24/93	1115/1145	NO _x , CO, & O ₂	150
4-2-HC	5/24/93	1115/1145	Hydrocarbon	150
4-3-CEM	5/24/93	1200/1230	NO _x , CO, & O ₂	150
4-3-HC	5/24/93	1200/1230	Hydrocarbon	150
4-4-CEM	5/24/93	1345/1415	NO _x , CO & O ₂	150
4-4-HC	5/24/93	1345/1415	Hydrocarbon	150
4-4-NH ₃	5/24/93	1345/1415	Ammonia	150
4-5-CEM	5/24/93	1435/1505	NO _x , CO & O ₂	150
4-5-HC	5/24/93	1435/1505	Hydrocarbon	150
4-5-NH ₃	5/24/93	1435/1505	Ammonia	150
4-6-CEM	5/25/93	1015/1053	NO _x , CO & O ₂	256
4-6-HC	5/25/93	1015/1105	Hydrocarbon	256
4-6-NH ₃	5/25/93	1015/1045	Ammonia	256
4-7-CEM	5/25/93	1115/1145	NO _x , CO & O ₂	256
4-7-HC	5/25/93	1115/1145	Hydrocarbon	256
4-7-NH ₃	5/25/93	1115/1145	Ammonia	256
4-8-CEM	5/25/93	1210/1240	NO _x , CO & O ₂	256
4-8-HC	5/25/93	1210/1230	Hydrocarbon	256
4-9-CEM	5/25/93	1250/1320	NO _x , CO & O ₂	256
4-9-HC	5/25/93	1250/1310	Hydrocarbon	256
4-10-CEM	8/12/93	1045/1125	NO _x , CO, O ₂	331
4-10-HC	8/12/93	1050/1110	Hydrocarbon	331
4-10-NH ₃	8/12/93	1045/1125	Ammonia	331
4-11-CEM	8/12/93	1130/1200	NO _x , CO, O ₂	330
4-11-HC	8/12/93	1135/1155	Hydrocarbon	330
4-11-NH ₃	8/12/93	1130/1200	Ammonia	330
4-12-CEM	8/12/93	1220/1250	NO _x , CO, O ₂	330
4-12-HC	8/12/93	1230/1250	Hydrocarbon	330
4-13-CEM	8/12/93	1300/1330	NO _x , CO, O ₂	332
4-13-NHC	8/12/93	1300/1320	Hydrocarbon	332

TABLE 3-2
UNIT OPERATING CONDITIONS
EL SEGUNDO UNIT 4
MAY AND AUGUST 1993

Test No.	Date	Net Load, MW	Urea Status	Total Urea Flow gpm
4-2	5/24/93	150	off	0.0
4-3	5/24/93	150	off	0.0
4-4	5/24/93	150	on	0.7
4-5	5/24/93	150	on	0.7
4-6	5/24/93	245	on	1.34
4-7	5/24/93	246	on	1.34
4-8	5/24/93	246	off	0.0
4-9	5/24/93	246	off	0.0
4-10	8/12/93	331	on	2.24
4-11	8/12/93	330	on	2.24
4-12	8/12/93	330	off	0.0
4-13	8/12/93	332	off	0.0

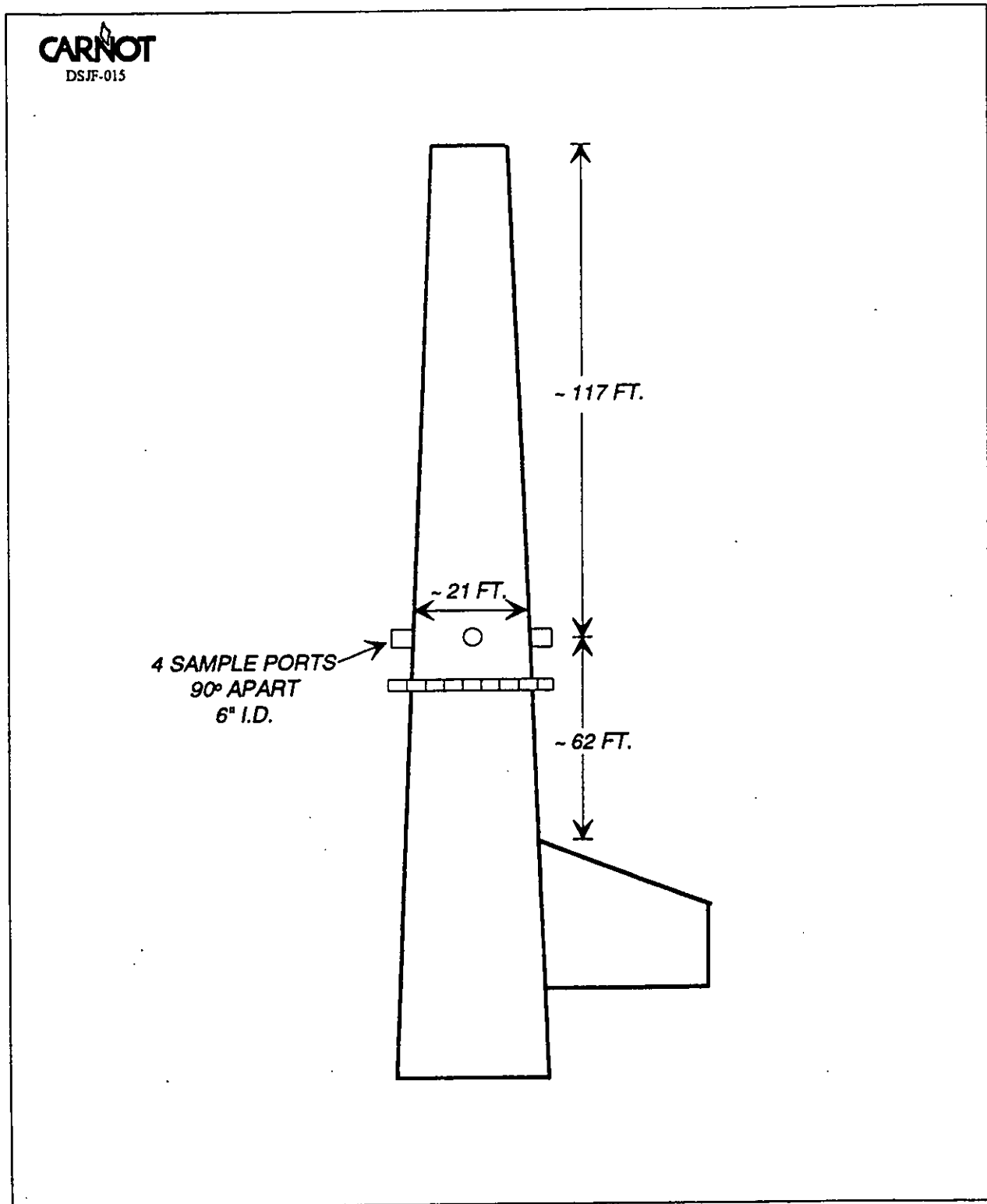


Figure 3-1. El Segundo Unit 4 Sample Location

TABLE 3-3
EMISSIONS TEST PROCEDURES - SCE/EL SEGUNDO UNIT 4
MAY 1993

Parameter	Number of Replicates	Duration of Test (Minutes)	Measurement Principle	Reference Method	Method Detection Limit
NO _x	2	30	Chemiluminescent	SCAQMD 100.1	Below 2% of instrument full scale
CO	2	30	NDIR/Gas filter correlation	SCAQMD 100.1	Below 2% of instrument full scale
O ₂	2	30	Electrochemical cell	SCAQMD 100.1 EPA 3E	Below 2% of instrument full scale
ROG as Hydrocarbons	2	20	Gas chromatography/FID	SCAQMD 25.2	0.1 ppm
Ammonia	2	30	Colorimetry spectrophotometry	SCAQMD Draft 207.1	0.1 ppm

3.3.2 Reactive Organic Gases (ROG) as Hydrocarbons

Hydrocarbon samples were collected in new tedlar bags and analyzed according to SCAQMD Method 25.2. The samples were analyzed within 72 hours of collection by Atmosphere Assessments in Chatsworth, California.

3.3.3 Ammonia

Ammonia samples were collected non-isokinetically, with NH_3 being absorbed in two impingers containing 0.1 N H_2SO_4 . The ammonia samples were analyzed on-site by SCAQMD NH_3 Draft Method.

3.3.4 Moisture

Moisture was determined by SCAQMD Method 4 and in conjunction with NH_3 sampling.

SECTION 4.0

RESULTS

Emissions test results are presented in Tables 4-1 through 4-6. Measurement procedures are detailed in Appendix A. Quality assurance and equipment calibrations are in Appendix B. All raw data sheets from Carnot's testing and from unit operation are included in Appendix C. Appendix D contains general and specific emissions calculations. Appendix E contains laboratory reports. Carnot's instrument strip charts are in Appendix F. Chain of Custody information is included in Appendix G.

TABLE 4-1
EMISSIONS TEST RESULTS AT 331 MW
UREA ON
EL SEGUNDO UNIT 4
AUGUST 12, 1993

Parameter	Test No. 4-10	Test No. 4-11	Test Average
Time	1045/1115	1130/1200	
NO _x , ppm	64.8	66.7	65.8
ppm @ 3% O ₂	73.9	76.0	75.0
CO, ppm	4.8	4.5	4.7
ppm @ 3% O ₂	5.5	5.1	5.3
O ₂ %	5.2	5.2	5.2
H ₂ O %	18.2	18.0	18.1
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0
Total Non-Methane Hydrocarbons as methane, ppm	10.3	7.7	8.6
NH ₃ , ppm	5.5	6.9	6.2
NH ₃ @ 3% O ₂ ⁽²⁾	6.3	7.9	7.1

Note: ND indicates that this species was not detected.

TABLE 4-2
EMISSIONS TEST RESULTS AT 331 MW
UREA OFF
EL SEGUNDO UNIT 4
AUGUST 12, 1993

Parameter	Test No. 4-12	Test No. 4-13	Test Average
Time	1220/1250	1300/1330	
NO _x , ppm	73.1	73.8	73.5
ppm @ 3% O ₂	83.9	83.1	83.5
CO, ppm	5.9	7.1	6.5
ppm @ 3% O ₂	6.8	8.0	7.4
O ₂ %	5.3	5.0	5.2
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0
Total Non-Methane Hydrocarbons as methane, ppm	10.0	7.2	8.6

Note: ND indicates that this species was not detected.

TABLE 4-3
EMISSIONS TEST RESULTS AT 256 MW
UREA ON
EL SEGUNDO UNIT 4
MAY 25, 1993

Parameter	Test No. 4-6	Test No. 4-7	Test Average
Time	1015/1045	1115/1145	
NO _x , ppm	55.4	55.1	55.3
ppm @ 3% O ₂	62.0	61.6	61.8
CO, ppm	17.5	19.5	18.5
ppm @ 3% O ₂	19.6	21.8	20.7
O ₂ %	4.9	4.9	4.9
H ₂ O %	18.3	17.9	18.1
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0
Total Non-Methane Hydrocarbons as methane, ppm	19.6	23.1	21.4
NH ₃ , ppm	9.7	11.5	10.6
NH ₃ @ 3% O ₂	10.8	12.9	11.9

Note: ND indicates that this species was not detected.

TABLE 4-4
EMISSIONS TEST RESULTS AT 256 MW
UREA OFF
EL SEGUNDO UNIT 4
MAY 25, 1993

Parameter	Test No. 4-8	Test No. 4-9	Test Average
Time	1115/1145	1210/1240	
NO _x , ppm	66.6	67.3	67.0
ppm @ 3% O ₂	74.0	74.8	74.4
CO, ppm	21.1	25.6	23.4
ppm @ 3% O ₂	23.5	28.5	26.0
O ₂ %	4.8	4.8	4.8
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0
Total Non-Methane Hydrocarbons as methane, ppm	21.2	20.4	20.8

Note: ND indicates that this species was not detected.

TABLE 4-5
EMISSIONS TEST RESULTS AT 150 MW
UREA ON
EL SEGUNDO UNIT 4
MAY 24, 1993

Parameter	Test No. 4-4	Test No. 4-5	Test Average
Time	1345/1415	1435/1505	
NO _x , ppm	38.6	40.0	39.3
ppm @ 3% O ₂	46.4	48.1	47.3
CO, ppm	0.0	0.0	0.0
ppm @ 3% O ₂	0.0	0.0	0.0
O ₂ %	6.0	6.0	6.0
H ₂ O%	18.1	16.1	17.1
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0
Total Non-Methane Hydrocarbons as methane, ppm	17.1	18.0	17.6
NH ₃ , ppm	2.8	5.2	4.0
NH ₃ , @ 3% O ₂	3.4	6.2	4.8

Note: ND indicates that this species was not detected.

TABLE 4-6
EMISSIONS TEST RESULTS AT 150 MW
UREA OFF
EL SEGUNDO UNIT 4
MAY 24, 1993

Parameter	Test No. 4-2	Test No. 4-3	Test Average
Time	1115/1145	1200/1230	
NO _x , ppm	44.1	43.7	43.9
ppm @ 3% O ₂	53.0	52.6	53.0
CO, ppm	0.0	0.0	0.0
ppm @ 3% O ₂	0.0	0.0	0.0
O ₂ %	6.0	6.0	6.1
Methane, ppm	ND < 1.0	ND < 1.0	ND < 1.0
Total Non-Methane Hydrocarbons as methane, ppm	15.8	19.0	17.4

Note: ND indicates that this species was not detected.

APPENDIX A
MEASUREMENT PROCEDURES

Continuous Emissions Monitoring System

Oxygen (O_2) by Continuous Analyzer

NO/ NO_x by Continuous Analyzer

Carbon Monoxide (CO) by NDIR/Gas Filter Correlation

Hydrocarbons by SCAQMD Micro TCA Method

Ammonia

Continuous Emissions Monitoring System

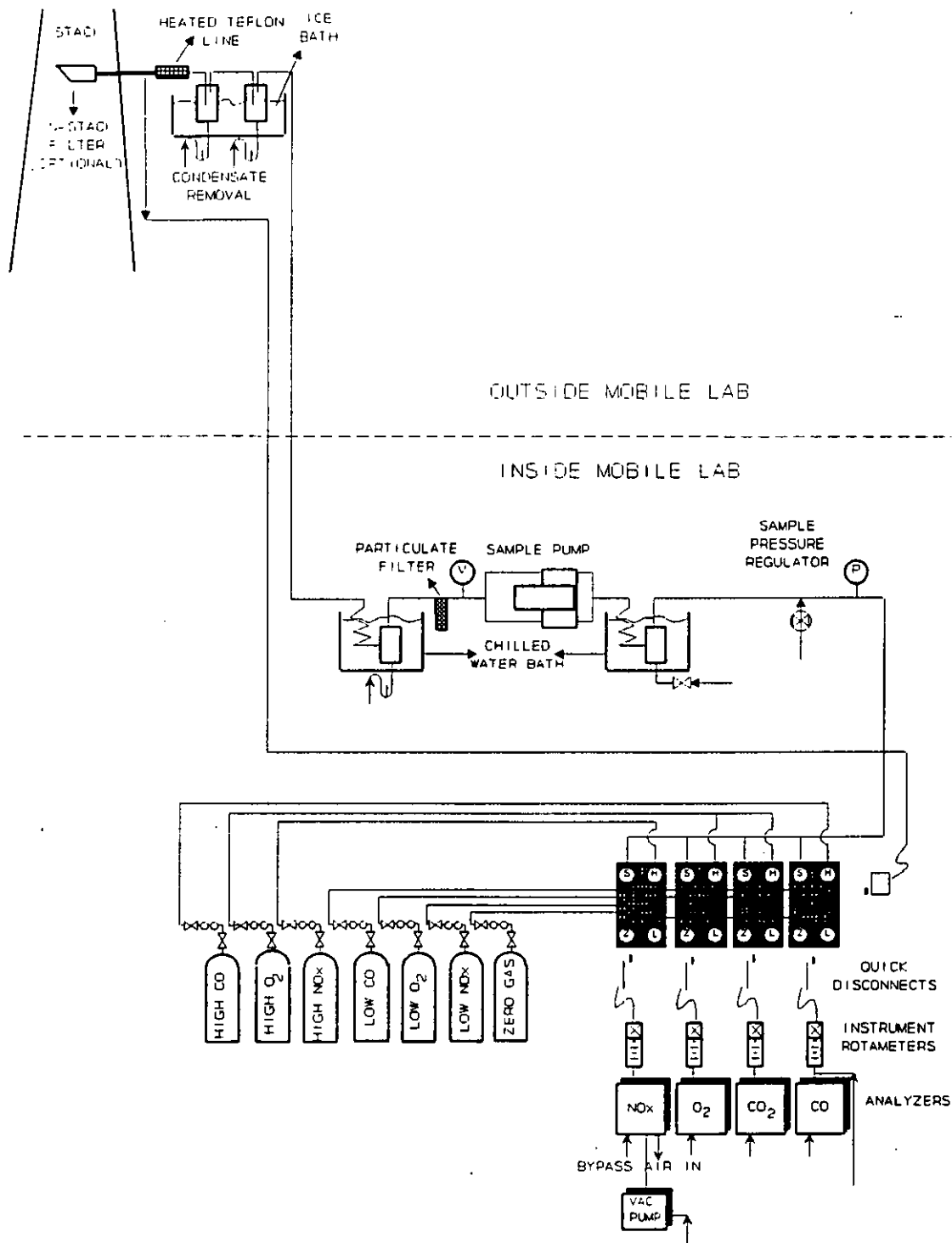
O₂, CO, and NO_x are measured using an extractive continuous emissions monitoring (CEM) package, shown in the following figure. This package is comprised of three basic subsystems. They are: (1) the sample acquisition and conditioning system, (2) the calibration gas system, and (3) the analyzers themselves. This section presents a description of the sampling and calibration systems. Descriptions of the analyzers used in this program and the corresponding reference test methods follow. Information regarding quality assurance information on the system, including calibration routines and system performance data follows.

The sample acquisition and conditioning system contains components to extract a representative sample from the stack or flue, transport the sample to the analyzers, and remove moisture and particulate material from the sample. In addition to performing the tasks above, the system must preserve the measured species and deliver the sample for analysis intact. The sample acquisition system extracts the sample through a stainless steel probe. The probe is insulated or heated as necessary to avoid condensation. If the particulate loading in the stack is high, a sintered stainless steel filter is used on the end of the probe.

Where water soluble NO₂ is to be measured, the sample is drawn from the probe through a heated teflon sample line into an on-stack cooled (approximately 35-40°F) water removal trap. The trap consists of stainless steel flasks in a bath of ice and water. This design removes the water vapor by condensation. The contact between the sample and liquid water is minimized and the soluble NO₂ is conserved. This system meets the requirements of EPA Method 20. The sample is then drawn through a teflon transport line, particulate filter, secondary water removal and into the sample pump. The pump is a dual head, diaphragm pump. All sample-wetted components of the pump are stainless steel or teflon. The pressurized sample leaving the pump flows through a third condensate trap in a refrigerated water bath ($\approx 38^\circ\text{F}$) for final moisture removal. A drain line and valve are provided to constantly expel any condensed moisture from the dryer at this point. After the dryer, the sample is directed into a distribution manifold. Excess sample is vented through a back-pressure regulator, maintaining a constant pressure of 5-6 psig to the analyzer rotameters.

The calibration system is comprised of two parts: the analyzer calibration, and the system bias check (dynamic calibration). The analyzer calibration equipment includes pressurized cylinders of certified span gas. The gases used are, as a minimum, certified to 1% by the manufacturer. Where necessary to comply with reference method requirements EPA Protocol 1 gases are used. The cylinders are equipped with pressure regulators which supply the calibration gas to the analyzers at the same pressure and flow rate as the sample. The selection of zero, span, or sample gas directed to each analyzer is accomplished by operation of the sample/calibration selector fittings.

The system bias check is accomplished by transporting the same gases used to zero and span the analyzers to the sample system as close as practical to the probe inlet. This is done either by attaching the calibration gas supply line to the probe top with flexible tubing or by actuation of a solenoid valve located at the sample conditioner inlet (probe exit). The span gas is exposed to the same elements as the sample and the system response is documented. The analyzer indications for the system calibration check must agree within 5% of the analyzer calibration. Values are adjusted and changes/repairs are made to the system to compensate for any difference in analyzer readings. Specific information on the analytical equipment and test methods used is provided in the following pages.



Schematic of CEM System

Method:	Oxygen (O₂) by Continuous Analyzer
Applicable Reference Methods:	EPA 3A, EPA 20, ARB 100, BA ST-14, SCAQMD 100.1
Principle:	A sample is continuously drawn from the flue gas stream, conditioned, and conveyed to the instrument for direct readout of O ₂ concentration.
Analyzer:	Teledyne Model 326A
Measurement Principle:	Electrochemical cell
Ranges:	0-5, 0-10, 0-25 % O ₂
Accuracy:	1 % of full scale
Output:	0-100 mV, linear
Interferences:	Halogens and halogenated compounds will cause a positive interference. Acid gases will consume the fuel cell and cause a slow calibration drift.
Response Time:	90% < 7 seconds
Sampling Procedure:	A representative flue gas sample is collected and conditioned using the CEM system described previously. If Method 20 is used, that method's specific procedures for selecting sample points are used. Otherwise, stratification checks are performed at the start of a test program to select single or multiple-point sample locations.
Analytical Procedure:	An electrochemical cell is used to measure O ₂ concentration. Oxygen in the flue gas diffuses through a Teflon membrane and is reduced on the surface of the cathode. A corresponding oxidation occurs at the anode internally, and an electric current is produced that is proportional to the concentration of oxygen. This current is measured and conditioned by the instrument's electronic circuitry to give an output in percent O ₂ by volume.
Special Calibration Procedure:	The measurement cells used with the O ₂ instrument have to be replaced on a regular basis. After extended use, the cell tend to produce a nonlinear response. Therefore, a three-point calibration is performed at the start of each test day to check for linearity. If the response is not linear ($\pm$ 2% of scale), the cell is replaced.

Method:	NO/NO _x by Continuous Analyzer
Applicable Reference Methods:	EPA 7E, EPA 20; ARB 100, BA ST-13A, SCAQMD 100.1
Principle:	A sample is continuously drawn from the flue gas stream, conditioned, and conveyed to the instrument for direct readout of NO or NO _x .
Analyzer:	Teco Model No. 10AR
Measurement Principle:	Chemiluminescence
Accuracy:	1% of full scale
Ranges:	0-2.5, 0-10, 0-25, 0-100, 0-250, 0-1000, 0-2500, 0-10,000 ppm
Output:	0-10 mV
Inferences:	Compounds containing nitrogen (other than ammonia) may cause interference.
Response Time:	90%, 1.5 seconds (NO mode) and 1.7 seconds (NO _x mode)
Sampling Procedure:	A representative flue gas sample is collected and conditioned using the CEM system described previously. If EPA Method 20 is used, that method's specific procedures for selecting sample points are used.
Analytical Procedure:	The oxides of nitrogen monitoring instrument is a chemiluminescent nitric oxide analyzer. The operational basis of the instrument is the chemiluminescent reaction of NO and ozone (O₃) to form NO₂ in an excited state. Light emission results when excited NO₂ molecules revert to their ground state. The resulting chemiluminescence is monitored through an optical filter by a high sensitivity photomultiplier tube, the output of which is electronically processed so it is linearly proportional to the NO concentration. The output of the instrument is in ppmV. When NO₂ is expected to be present in the flue gas, a supercooled water dropout flask will be placed in the sample line to avoid loss of NO₂. Since NO₂ is highly soluble in water, "freezing out" the water will allow the NO₂ to reach the analyzers for analysis. The analyzer measures NO only. In the NO_x mode, the gas is passed through a moly converter which converts NO₂ to NO and a total NO_x measurement is obtained. NO₂ is determined as the difference between NO and NO_x. Use of a moly converter instead of a stainless steel converter eliminates NH₃ interference; NH₃ is converted to NO with a stainless converter, but not with a moly converter.

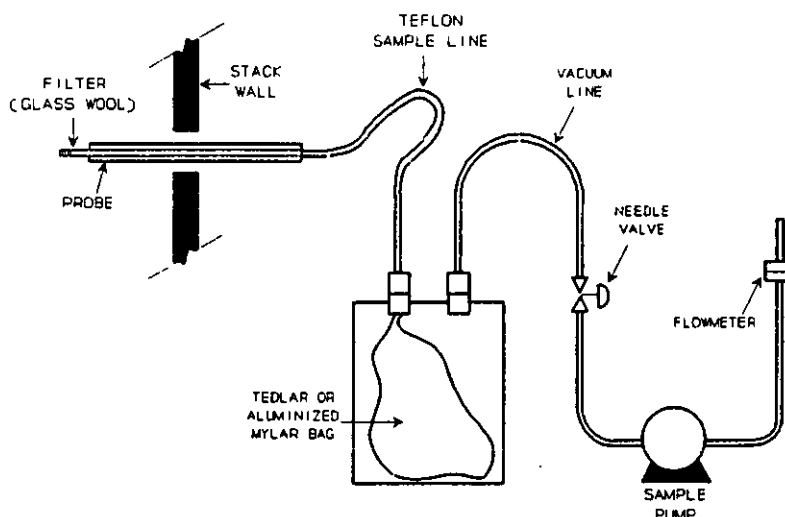
Method:	Carbon Monoxide (CO) by NDIR/Gas Filter Correlation
Applicable Reference Methods:	EPA 10; ARB 1-100; BA ST-6, SCAQMD 100.1
Principle:	A sample is continuously drawn from the flue gas stream, conditioned, and conveyed to the instrument for direct readout of CO concentration.
Analyzer:	TECO, Model 48
Measurement Principle:	NDIR/Gas Filter Correlation
Precision:	0.1% ppm
Ranges:	0-5, 0-10, 0-25, 0-50, 0-100, 0-250, 0-500, 0-1000, 0-5000 ppm
Output:	0-100 mV
Interferences:	Negligible interference from water and CO ₂
Rise/Fall Times (0-95%)	1 minute @ 1 lpm flow, 30 second integration time
Sampling Procedure:	A representative flue gas sample is collected and conditioned using the CEM system described previously. Sample point selection has been described previously.
Analytical Procedure:	Radiation from an infrared source is chopped and then passed through a gas filter which alternates between CO and N ₂ due to rotation of a filter wheel. The radiation then passes through a narrow band-pass filter and a multiple optical pass sample cell where absorption by the sample gas occurs. The IR radiation exits the sample cell and falls on a solid state IR detector.

Method: **Hydrocarbons by SCAQMD Micro Total Carbon Analysis (TCA/FID) Method**

Reference: SCAQMD Method 25.2 (draft Source Test Manual)

Principle: A Tedlar bag is filled with flue gas at a constant rate. The bag contents are analyzed by gas chromatography/flame ionization detection for CO, CO₂, methane and total gaseous non-methane hydrocarbons.

Sampling Procedure: Samples are collected using a lung-type sampling system shown in the attached figure. In this system, a bag is placed in a sealed container and the container is evacuated. Flue gas enters the bag as it expands to fill the container. Sampling rate is monitored by a rotameter on the container exhaust. This system allows sample collection without exposing the sample to pumps, flowmeters, oils, etc.



**Sample Train for Determination of Hydrocarbons
by SCAQMD Micro TCA Method**

Analytical Procedure:

In the analytical phase, gaseous carbon compounds from the tank are fractionated on a chromatographic column, eluting in the order: carbon monoxide, methane, carbon dioxide, non-methane hydrocarbons. All resulting vapors are passed through oxidizers where they are combusted to carbon dioxide, then converted to methane and measured by flame ionization detection. CO₂, CO, CH₄, and total gaseous non-methane hydrocarbons are reported separately.

This method provides lower detection limits than the standard TCA procedure, which uses GC/NDIR. Since this method does not use a steel condensate trap, it eliminates positive interferences that have been associated with carbon dioxide dissolved in water frozen in the condensate trap and released during analysis.

The TCA/FID method is appropriate for combustion sources and ambient sampling where low concentrations of hydrocarbons are expected.

Carnot subcontracts TCA analysis to qualified local laboratories experienced in the analytical procedures.

Method: Ammonia

Principle: Ammonia emissions are determined by collecting a dry, metered volume of flue gas containing ammonia vapor in dilute sulfuric acid solution. The ammonia concentration of the sample is then determined by ion selective electrode or colorimetric spectrophotometry.

Sampling Procedure: The ammonia sampling probe/train arrangement used during the test program is shown schematically in the following figure.

A sample is extracted through a glass probe fitted with a quartz wool plug or filter used to remove particulate matter. The plug is located and/or heated to avoid condensation of moisture at the plug. The sample is then passed through four impingers. The first two impingers are filled with 100 ml of 0.1 N sulfuric acid (H_2SO_4), the third is empty, and the fourth is filled with silica gel.

After each sample run, the quartz wool plug is removed and placed in a sample bottle. If a filter is used, it is generally saved but not analyzed. The impingers, glass probe, and sampling train glassware are washed with dilute H_2SO_4 into a separate sample bottle. The sample bottles containing the impinger wash solution and the quartz wool plug or filter are returned to the laboratory for analysis.

Analytical Procedure: Ammonia in source test samples is analyzed by ion selective electrode or colorimetrically. Source test samples are generally at low levels and titrations for ammonia are not applicable. The following sections discuss the two applicable analytical techniques.

Ion Selective Electrode

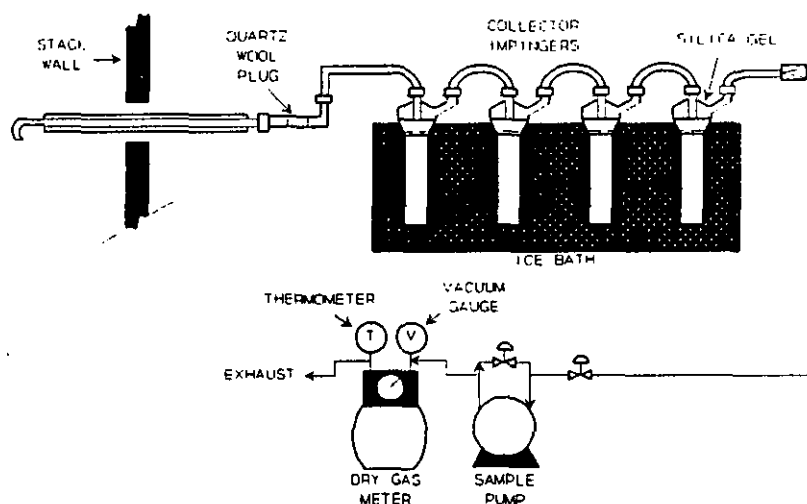
The ammonia samples are analyzed using an Orion Research Model 940 Microprocessor Ionanalyzer equipped with an ammonia ion selective electrode. The ammonia electrode is a glass electrode and reference electrode mounted behind a gas permeable membrane that separates the sample solution from the internal filling solution. After the addition of sodium hydroxide, ammonia in the sample diffuses across the electrode membrane. The measured pH change in the internal electrode solution is proportional to the concentration of ammonia in the sample solution.

The ammonia concentrations of the sample solutions are measured in units of ppm NH_3 by weight with the specific ion analyzer. By using the NH_3 concentration in ppm by weight, measuring the liquid volume of the sample solution, and obtaining the volume of the flue gas sample, the stack flue gas NH_3 concentration in ppm by volume is calculated using the equation:

$$NH_3, \text{ ppmV} = 0.049 \frac{(\text{ppm wt } NH_3, \text{ soln}) (\text{liq. vol., ml})}{\text{DSCF (at } 60^\circ\text{F) of sampled flue gas}}$$

**Colorimetric
Spectrophotometry:**

In the analytical phase, the resulting ammonium sulfate is mixed with Nessler reagent, forming a yellow-brown compound with a maximum absorbance in the 440 nm range. The ammonia concentration in the sample is determined colorimetrically by comparison to a standard calibration curve. Care will be taken to dilute the sample to the optimum observed range. If turbidity exists upon addition of Nessler reagent, the option of distillation or ion selective electrode will be used. Distillation will remove any aromatic and aliphatic amines along with other organic and inorganic compounds which can cause interference with the analysis.



Ammonia sampling probe/train arrangement

APPENDIX B
QUALITY ASSURANCE

Appendix B.1
Quality Assurance Program Summary

QUALITY ASSURANCE PROGRAM SUMMARY AND ARB CERTIFICATION

Carnot ensures the quality and validity of its emission measurement and reporting procedures through a rigorous quality assurance (QA) program. The program is developed and administered by an internal QA Officer and encompasses seven major areas:

1. Development and use of an internal QA manual.
2. QA reviews of reports, laboratory work, and field testing.
3. Equipment calibration and maintenance.
4. Chain of custody.
5. Training.
6. Knowledge of current test methods.
7. Agency certification.

Each of these areas is discussed individually below.

Quality Assurance Manual. Carnot has prepared a QA Manual according to EPA guidelines. The manual serves to document and formalize all of Carnot's QA efforts. The manual is constantly updated, and each member of the Source Test Division is required to read and understand its contents. The manual includes details on the other six QA areas discussed below.

QA Reviews. Carnot's review procedure includes review of each source test report by the QA Officer, and spot check reviews of laboratory and field work.

The most important review is the one that takes place before a test program begins. The QA Officer works closely with Source Test Division personnel to prepare and review test protocols. Test protocol review includes selection of appropriate test procedures, evaluation of any interferences or other restrictions that might preclude use of standard test procedures, and evaluation and/or development of alternate procedures.

Equipment Calibration and Maintenance. The equipment used to conduct the emissions measurements is maintained according to the manufacturer's instructions to ensure proper operation. In addition to the maintenance program, calibrations are carried out on each measurement device according to the schedule outlined by the California Air Resources Board (CARB). The schedule for maintenance and calibrations are given in Tables B-1 and B-2. Quality control checks are also conducted in the field for each test program. The following is a partial list of checks made as part of each CEM system test series.

- Sample acquisition and conditioning system leak check.
- 2-point analyzer calibrations (all analyzers)
- 3-point analyzer calibrations (analyzers with potential for linearity errors).
- Complete system calibration check ("dynamic calibration" through entire sample system).

- Periodic analyzer calibration checks (once per hour) are conducted at the start and end of each test run. Any change between pre- and post-test readings are recorded.
- All calibrations are conducted using gases certified by the manufacturer to be + 1% of label value (NBS traceable).

Calibration and CEM performance data are fully documented, and are included in each source test report.

Chain of Custody. Carnot maintains full chain of custody documentation on all samples and data sheets. In addition to normal documentation of changes between field sample custodians, laboratory personnel, and field test personnel, Carnot documents every individual who handles any test component in the field (e.g., probe wash, impinger loading and recovery, filter loading and recovery, etc.).

Samples are stored in a locked area to which only Source Test Division personnel have access. Neither other Carnot employees nor cleaning crews have keys to this area.

Data sheets are copied immediately upon return from the field, and this first generation copy is placed in locked storage. Any notes made on original sheets are initialed and dated.

Training. Personnel training is essential to ensure quality testing. Carnot has formal and informal training programs which include:

1. Attendance at EPA-sponsored training courses.
2. Enrollment in EPA correspondence courses.
3. A requirement for all technicians to read and understand Carnot's QA Manual.
4. In-house training and QA meetings on a regular basis.
5. Maintenance of training records.

Knowledge of Current Test Methods. With the constant updating of standard test methods and the wide variety of emerging test methods, it is essential that any qualified source tester keep abreast of new developments. Carnot subscribes to services which provide updates on EPA and CARB reference methods, and on EPA, CARB and SCAQMD rules and regulations. Additionally, source test personnel regularly attend and present papers at testing and emission-related seminars and conferences. Carnot personnel maintain membership in the Air Pollution Control Association, the Source Evaluation Society, and the ASME Environmental Control Division.

AGENCY CERTIFICATION

Carnot is certified by the CARB as an independent source test contractor for gaseous and particulate measurements. Carnot also participates in EPA QA audit programs for Methods 5, 6 and 7.

TABLE B-1
SAMPLING INSTRUMENTS AND EQUIPMENT CALIBRATION SCHEDULE
As Specified by the CARB

Instrument Type	Frequency of Calibration	Standard of Comparison or Method of Calibration	Acceptance Limits
Orifice Meter (large)	12 months	Calibrated dry test meter	$\pm 2\%$ of volume measured
Dry Gas Meter	12 months or when repaired	Calibrated dry test meter	$\pm 2\%$ of volume measured
S-Type Pitot (for use with EPA-type sampling train)	6 months	EPA Method 2	Cp constant (+5%) over working range; difference between average Cp for each leg must be less than 2%
Vacuum Gauges Pressure Gauges	6 months	Manometer	$\pm 3\%$
Field Barometer	6 months	Mercury barometer	$\pm 0.2''$ Hg
Temperature Measurement	6 months	NBS mercury thermometer or NBS calibrated platinum RTD	$\pm 4^{\circ}\text{F}$ for $< 400^{\circ}\text{F}$ $\pm 1.5\%$ for $> 400^{\circ}\text{F}$
Temperature Readout Devices	6 months	Precision potentiometer	$\pm 2\%$ full scale reading
Analytical Balance	12 months (check prior to each use)	Should be performed by manufacturer or qualified laboratory	± 0.3 mg of stated weight
Probe Nozzles	12 Months	Nozzle diameter check micrometer	Range $< \pm 0.10$ mm for three measurements
Continuous Analyzers	Depends upon use, frequency and performance	As specified by manufacturers operating manuals, EPA NBS gases and/or reference methods	Satisfy all limits specified in operating specifications

TABLE B-2
EQUIPMENT MAINTENANCE SCHEDULE
Based on Manufacturer's Specifications and Carnot Experience

Equipment	Performance Requirement	Maintenance Interval	Corrective Action
Pumps	1. Absence of leaks 2. Ability to draw manufacturer required vacuum and flow	Every 500 hours of operation or 6 months, whichever is less	1. Visual inspection 2. Clean 3. Replace worn parts 4. Leak check.
Flow Measuring Device	1. Free mechanical movement 2. Absence of malfunction	Every 500 hours of operation or 6 months, whichever is less After each test, if used in H ₂ S sampling or other corrosive atmospheres	1. Visual inspection 2. Clean 3. Calibrate
Sampling Instruments	1. Absence of malfunction 2. Proper response to zero, span gas	As required by the manufacturer	As recommended by manufacturer
Integrated Sampling Tanks	Absence of leaks	Depends on nature of use	1. Steam clean 2. Leak check
Mobile Van Sampling Systems	Absence of leaks	Depends on nature of use	1. Change filters 2. Change gas dryer 3. Leak check 4. Check for system contamination
Sampling Lines	Sample degradation less than 2%	After each test or test series	Blow filtered air through line until dry

Appendix B.2
ARB Certification

State of California
AIR RESOURCES BOARD

Executive Order G-836

Approval to Carnot
To Conduct Testing as an Independent Contractor

WHEREAS, the Air Resources Board ("Board"), pursuant to Section 41512 of the California Health and Safety Code, has established the procedures contained in Section 91200-91220, Title 17, California Code of Regulations, to allow the use of independent testers for compliance tests required by the Board; and

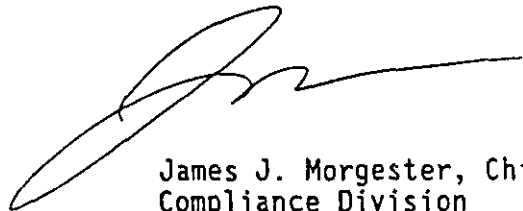
WHEREAS, pursuant to Sections 91200-91220, Title 17, California Code of Regulations, the Executive Officer has determined that Carnot meets the requirements of the Board for conducting Board Test Methods 1, 2, 3, 4, 5, 6, 8, 10, and 100 (NOx, O2) when the following conditions are met:

1. Carnot conducts Board Test Method 100 for O2 using a Teledyne 326 analyzer with either a A5 or a B1 sensor, or a paramagnetic analyzer.

NOW, THEREFORE, BE IT ORDERED that Carnot is granted an approval, from the date of execution of this order, until June 30, 1994 to conduct the tests listed above, subject to compliance with Section 91200-91220, Title 17, California Code of Regulations.

BE IT FURTHER ORDERED that during the approved period the Executive Officer or his or her authorized representative may field audit one or more tests conducted pursuant to this order for each type of testing listed above.

Executed this 14th day of July 1993, at Sacramento, California.



James J. Morgester, Chief
Compliance Division

State of California
Air Resources Board
Approved Independent Contractor

Carnot

This is to certify that the company listed above has been approved by the Air Resources Board to conduct compliance testing pursuant to Section 91207, Title 17, California Code of Regulations, until June 30, 1994, for those test methods listed below:

ARB Source Test Methods:
1, 2, 3, 4, 5, 6, 8, 10, 100(NOx, O2)


James J. Morgester, Chief
Compliance Division


Laura McKinney, Manager
Investigation and Certification Section

Appendix B.3
Equipment Calibration

CARNOT SPAN GAS RECORD

CLIENT/LOCATION: SCE - EL SEGUNDO #4

DATE: 5-24-93

BY: M. C. C.

GAS	SPAN CYLINDER		AUX. SPAN CYLINDER	
	CYLINDER NO.	CONCENTRATION	CYLINDER NO.	CONCENTRATION
ZERO		N ₂		
NOx	ALM 7741	51.96	ALM 33901	93.49
O ₂	ALM 27279	5.032	AAL 3966	8.013
CO	AAL 7957	55.07	ALM 12534	85.12
CO ₂	ALM 27279	14.97	AAL 3966	22.74
SO ₂				

CARNOT INSTRUMENT LINEARITY

	ANALYZER				
	O ₂	CO ₂	CO	NOx	SO ₂
ANALYZER RANGE	10%	25%	100 ppm	100 ppm	
SET TO HIGH STD (80-90% OF RANGE)	8.0	22.75	85.1	93.5	
ACTUAL VALUE OF LOW STD	5.032	14.97	55.07	51.96	
AS-FOUND LOW STD (50-60% OF RANGE)	5.1	15.1	53.4	50.7	
DIFFERENCE IN % OF FULL SCALE	+0.7%	+0.1%	-1.7%	-1.3%	

% ERROR CALCULATION:

$$\frac{(\text{AS FOUND} - \text{ACTUAL VALUE OF SPAN})}{\text{RANGE}} \times 100$$

ALLOWABLE DEVIATION IS 2% OF FULL SCALE (2 SQUARES ON STRIP CHART).

CARNOT SPAN GAS RECORD

CLIENT/LOCATION: SCE / EL SEGUNDO 4

DATE: 5-25-93

BY: M'CUNE

GAS	SPAN CYLINDER		AUX. SPAN CYLINDER	
	CYLINDER NO.	CONCENTRATION	CYLINDER NO.	CONCENTRATION
ZERO		N ₂		
NOx	ALN 7741	51.96	ALN 33901	93.49
O ₂	ALN 27279	5.032	AAL 3966	8.013
CO	AAL 7957	55.07	ALN 12534	85.12
CO ₂	ALN 27279	14.97	AAL 3966	22.74
SO ₂				

CARNOT INSTRUMENT LINEARITY

	ANALYZER				
	O ₂	CO ₂	CO	NOx	SO ₂
ANALYZER RANGE	10%	25%	100 ppm	100 ppm	
SET TO HIGH STD (80-90% OF RANGE)	8.0	22.75	85.7	93.5	
ACTUAL VALUE OF LOW STD	5.032	14.97	55.07	51.96	
AS-FOUND LOW STD (50-60% OF RANGE)	5.09	15.0	54.0	50.2	
DIFFERENCE IN % OF FULL SCALE	+0.6%	0.0%	-1.1%	-1.8%	

% ERROR CALCULATION:

$$\frac{(\text{AS FOUND} - \text{ACTUAL VALUE OF SPAN})}{\text{RANGE}} \times 100$$

ALLOWABLE DEVIATION IS 2% OF FULL SCALE (2 SQUARES ON STRIP CHART).

CARNOT CEM PERFORMANCE DATA

CLIENT/LOCATION: SCE - EL SEGUNDO UNIT 4 DATE: 5-24-93
BY: McCune

SYSTEM CONFIGURATION				
ANALYZERS IN SERVICE				
ANALYZERS:	O ₂	CO ₂	CO	- NOx
MODEL:	TELEDYNE	HORIBA P12-2000	TECO 48	TECO 10AR
SERIAL NO.:	46124	912017		30333-237
PROBE		SAMPLE CONDITIONER		
LENGTH: 10'		CONDENSER-VACUUM SIDE (CHECK FLOW): ☒		
LINER MATERIAL: S.S.		CONDENSER-PRESSURE SIDE (CHECK FLOW): ☒		
HEATED PROBE (Y/N): N		CONDENSER TEMPERATURE: 5°C		
HEATED LINE (Y/N): Y		FILTER CONDITION (COND. OR DATE LAST CHANGED):		
SAMPLE LINE		SYSTEM LEAK CHECK		
LENGTH: 100'		PRE-TEST (cfh): 0.0 @ 24" Hg		
LINER MATERIAL: TEFLON		POST-TEST (cfh): 0.0 @ 24" Hg		
SYSTEM BIAS LINE: 100' / POLY FLOW		LEAK RATE (%) = $\frac{\text{POST-TEST (cfh)}}{\text{SYSTEM FLOW RATE (cfm)} \times 60} \times 100 = \underline{\hspace{2cm}}\%$		
ON-STACK CONDITIONER		NOx CONVERSION EFFICIENCY		
IN SERVICE (Y/N): ☒		HIGH CAL NOx <u>PERFORMED IN SHOP</u>		
KNOCK-OUT CONDITION (CHECK FLOW):		HIGH CAL NO (AS FOUND) <u>1</u>		
COOLANT: ICE		LOW CAL NOx <u> </u>		
		LOW CAL NO (AS FOUND) <u>1</u>		
OPERATING CONDITIONS				
SAMPLE PRESSURE: 5 psi		SYSTEM RESPONSE TIME CHECK		
SAMPLE VACUUM: 11" Hg		UPSCALE: <u>45</u> sec.		
NOx VACUUM: 29.8" Hg		DOWNSCALE: <u>40</u> sec.		

FAX 714-887-0549
PHONE: 714-887-2571

2600 CAJON BLVD., SAN BERNARDINO, CA 92405

CARNOT

ATTN: RICK MADRIGAL
15991 RED HILL AVE

CA 92680

Shipped From: Scott SAN BERNARDINO, CA
 Date Shipped 12/10/92
 Our Project No: 22565 / 23160
 Your P.O. No: 9113 ITEM # 8
 Page 1 of 1
 Expiration Date: 5/94

CERTIFICATE OF ANALYSIS -- EPA PROTOCOL GASES

Certified Per Traceability Protocol No. 1 Procedure No. G1

Cylinder No. ALM033901

Cylinder Pressure 2000 PSIG

Certified Accuracy +1% % NBS Traceable

REFERENCE STD

**CERTIFIED
CONC**

SRM/CRM NO.

CONC.

MAKE/MODEL/SERIAL NO.

GAS ANALYZER

LAST CAL. DATE	ANALYTICAL PRINCIPLE

10/29/92 CHEMI-LUMINESCEN.
10AR 14853-150
THERMO-ELECTRON

BALANCE GAS NITROGEN

ANALYZER READINGS: Z = Zero Gas T = Test Gas R = Reference Gas

Component NITRIC OXIDEFirst Analysis Date 11/19/92 Units mv

Z	0.00	R	95.31	T	90.70
---	------	---	-------	---	-------

R	95.21	
Z	0.00	
T	90.98	

Z 0.00 T 91.15 B 95.26

Mean Test Assay 92.13 ppm

Second Analysis Date 11/30/92 Units mv

Z 0.00 R 95.57 T 90 89

R 95.61
 Z -0.02
 T 90 95

Z	0.00
T	91.02
B	95.63
	90.93

Mean Test Assoc 01 81 nrm

Small Test Assay 91.8 2 PM

Chronology: Date

Criminology: Date _____
Assay _____

Assay

Chronology: Date

Assay

Analyst G. KING

Analyst G. KING

Approved By:

A. F. Lange
A. F. LANGE, Ph.D.

A. F. LANGE, Ph.D. MANAGER, OA



Scott Specialty Gases, Inc.

2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX: (909) 887-0549

CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS

Customer: Carnot
15991 Red Hill Ave.
Suite 110
Tustin, CA 92680-0740

Assay Laboratory
Scott Specialty Gases
2600 Cajon Boulevard
San Bernardino, CA 92411

Purchase Order 8957
Scott Project # 24740.001

ANALYTICAL INFORMATION

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number ALM 007741
Cylinder Pressure 1900psig

Certification Date 03-22-93
Previous Certification Dates NONE

Expiration Date 09-22-94

ANALYZED CYLINDER

Components
Nitric Oxide

Certified Concentration
50.80 ppm

Analytical Uncertainty*
± 1 % NIST Traceable

Balance Gas: Nitrogen

NOX

51.96 ppm

*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes.

REFERENCE STANDARD

Type CRM1683
Expiration Date 01-94

Cylinder Number
ALM10462

Concentration
47.53ppm Nitric Oxide

INSTRUMENTATION

Instrument/Model/Serial #
TECO / 10AR / 14853-150

Last Date Calibrated
01-20-93

Analytical Principle
Chemi-Luminescent

ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

Components

Nitric Oxide

First Triad Analysis

Date: 3/15/93 Response Units: mv
Z1= 0.00 R1= 47.58 T1= 50.92
R2= 47.76 Z2= 0.04 T2= 50.93
Z3= 0.06 T3= 50.97 R3= 47.71
Avg. Conc. of Cust Cyl 50.78 ppm

Second Triad Analysis

Date: 03-22-93 Response Units: mv
Z1= 0.00 R1= 47.53 T1= 50.85
R2= 47.55 Z2= 0.00 T2= 50.81
Z3= 0.00 T3= 50.83 R3= 47.53
Avg. Conc. of Cust Cyl 50.82ppm

Calibration Curve

Concentration= Ax^2+Bx+C
A = -0.00022788
B = 1.011308772
C = 0.213935299

Date: Response Units: mv
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl

Date: Response Units: mv
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl

Concentration=

Date: Response Units:
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl

Date: Response Units:
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl

Concentration=

Special Notes


Analyst George King



Scott Specialty Gases, Inc.

2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX: (909) 887-0549

CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS

Customer
CARNOT
15991 RED HILL AVE
SUITE 110
TUSTIN, CA 92680
ATTN: RICK MADRIGAL

Assay Laboratory
Scott Specialty Gases
2600 Cajon Boulevard
San Bernardino, CA 92411

Purchase Order 9764 REL#5
Project # 25145.001

ANALYTICAL INFORMATION

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number ALM012534
Cylinder Pressure 2000 psig

Certification Date 04-02-93
Previous Certification Dates NONE

Expiration Date 10-02-94
** 04-02-96

ANALYZED CYLINDER

Components
CARBON MONOXIDE

Certified Concentration
85.12 ppm

Analytical Uncertainty*
± 1 % NIST Traceable

Balance Gas: Nitrogen

*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes.

REFERENCE STANDARD

Type CRM1679
Expiration Date 10-94

Cylinder Number
ALM10524

Concentration
96.67 ppm

INSTRUMENTATION

Instrument/Model/Serial #
Horiba / OPE-135D / 56565502

Last Date Calibrated
02-10-92

Analytical Principle
NDIR

ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

Components

Carbon Monoxide

First Triad Analysis

Date: 03-26-92 Response Units: mv
Z1= 00.00 R1= 40.20 T1= 35.50
R2= 40.20 Z2= 00.01 T2= 35.50
Z3= 00.01 T3= 35.50 R3= 40.20
Avg. Conc. of Cust Cyl. 84.99 ppm

Second Triad Analysis

Date: 04-02-92 Response Units: mv
Z1= 00.00 R1= 40.20 T1= 35.60
R2= 40.20 Z2= 00.09 T2= 35.60
Z3= 00.07 T3= 35.60 R3= 40.20
Avg. Conc. of Cust Cyl. 85.24 ppm

Calibration Curve

Concentration= Ax²+Bx+C
A=0.00005011
B=0.50595305
C=-0.0100275

Date: Response Units: mv
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl.

Date: Response Units: mv
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl.

Concentration=

Date: Response Units:
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl.

Date: Response Units:
Z1= R1= T1=
R2= Z2= T2=
Z3= T3= R3=
Avg. Conc. of Cust Cyl.

Concentration=

Special Notes : * Acid Rain Certification Period
** Non-Acid Rain Certification Period

Analyst Maynard Johnson



Scott Specialty Gases, Inc.

2600 CAJON BLVD., SAN BERNARDINO, CA 92411

CARNOT

ATTN: RICK MADRICAL
15991 RED HILL AVE
SUITE 110
TUSTIN

CA 92680

FAX 887-0549
PHONE: 714-887-2571

Shipped From: Scott

SAN BERNARDINO, CA

Date Shipped: 10/15/92

Our Project No: 21912

Your P.O. No: 8957

Page 1 of 1

Expiration Date: 4/94

CERTIFICATE OF ANALYSIS - EPA PROTOCOL GASES

Cylinder No. ALI.7957

Cylinder Pressure 1900 PSIG

Certified Accuracy $\pm 1\%$ % NIST Traceable

Certified Per Traceability Protocol No. 1

Procedure No. G1

REFERENCE STD

COMPONENTS	CERTIFIED CONC	SRM/CRM NO.	CYL. NO.	CONC.	MAKE/MODEL/SERIAL NO.
CARBON MONOXIDE	55.07 PPM	CRM1679	ALM016524	86.62 PPM	HORIBA AIA-24

4564403

GAS ANALYZER

LAST CAL DATE 6/30/92

ANALYTICAL PRINCIPLE INFRARED

NITROGEN

BALANCE GAS ANALYZER READINGS: Z = Zero Gas T = Test Gas R = Reference Gas

Component	CARBON MONOXIDE	Units	mv
First Analysis Date 10/7/92			
Z 00.00	R 39.4	T 22.9	
R 39.4	Z 00.01	T 22.9	
Z 00.02	T 22.9	R 39.4	
Mean Test Assay	55.07 PPM		
Second Analysis Date 10/14/92			
Z 00.00	R 39.4	T 22.9	
R 39.4	Z 00.01	T 22.9	
Z -00.03	T 22.9	R 39.4	
Mean Test Assay	55.07 PPM		

Component	Units	mv
Date		
Z	R	T
R	Z	T
Z	T	R
Mean Test Assay		
Date		
Z	R	T
R	Z	T
Z	T	R
Mean Test Assay		

Chronology: Date _____
Assay _____

Analyst M. JOHNSON

Approved By:

C. Lange

A.F. LANGE, Ph.D., MANAGER, QA



Scott Specialty Gases, Inc.

Shipped
From:

2600 CAJON BLVD.
SAN BERNARDINO
Phone: 714-887-2571

CA 92411

Fax: 714-887-0549

CERTIFICATE OF ANALYSIS

CARNOT
ATTN: RICK MADRIGAL
15991 RED HILL AVE
SUITE 110
TUSTIN

CA 92680

PROJECT #: 02-22304
PO#: 8957
ITEM #: 0202C3013601AL
DATE: 10/29/92

CYLINDER #: AAL3966
FILL PRESSURE: 1950 PSIG

ANALYTICAL ACCURACY: +/-1%

COMPONENT

CARBON DIOXIDE
OXYGEN
NITROGEN

REQUESTED GAS

CONC	MOLES
22.5	PCT
8.	PCT
	BAL

ANALYSIS

(MOLES)
22.74 PCT
8.013 PCT
BAL

CERTIFIED

CARNOT 11/5 BIN#1
CERTIFIED TO HAVE BEEN BLENDED
AND VERIFIED TO BE CORRECT BY
ANALYSIS.

GRAVIMETRIC MASTER GAS -
AGAINST NIST CERTIFIED WEIGHTS
INDEPENDENT LABORATORY

ANALYST:

JOSEPH DE LA TORRE

APPROVED BY:

DR. ARMAND LANGE, PH.D.

PLUMSTEADVILLE, PENNSYLVANIA / TROY, MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA
SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO
BATON ROUGE, LOUISIANA



Scott Specialty Gases, Inc.

9113

opped
F:

2600 CAJON BLVD.
SAN BERNARDINO
Phone: 714-887-2571

CA 92411

Fax: 714-887-0549

CERTIFICATE OF ANALYSIS

CARNOT
ATTN: RICK MADRIGAL
15991 RED HILL AVE
SUITE 110
DUSTIN

CA 92680

PROJECT #: 02-21678
FOR: 8880 ITEM # 1
ITEM #: 020203000701AL
DATE: 9/24/92

CYLINDER #: ALM027279
FILL PRESSURE: 2000 PSIG

ANALYTICAL ACCURACY: +/-1%

COMPONENT
CARBON DIOXIDE
OXYGEN
NITROGEN

REQUESTED GAS	
CONC. MOLES	
15.	PCT
5.	PCT
	BAL

ANALYSIS	
(MOLES)	
14.97	PCT
5.032	PCT
	BAL

CERTIFIED

CARNOT 9/28 BIN#9
CERTIFIED TO HAVE BEEN BLENDED
AND VERIFIED TO BE CORRECT BY
ANALYSIS.

GRAVIMETRIC MASTER GAS -
AGAINST HIST CERTIFIED WEIGHTS
INDEPENDENT LABORATORY

ANALYST:

JOSEPH DE LA TORRE

APPROVED BY:

DR ARMAND LANGE PH D

PLUMSTEADVILLE, PENNSYLVANIA / TROY, MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA
SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO

POST TEST DRY GAS METER CALIBRATION CHECK

CALIBRATED BY WM

DATE 5/27/93

FIELD GAS METER I.D. ES-31

INITIAL Yd 1.006

INITIAL H@ 2.110

DATA ENTRY BY

BAROMETRIC PRESSURE WM 29.897

TEST METETER ES-13

TEST METER Y (Yt) 0.9980

TEST METER LAST CAL. 4/05/93

FIELD METER					TEST METER			RESULTS	
VOLUME cu.ft.	TEMP. IN	TEMP. OUT	DELTA "H2O	TIME min.	VOLUME cu.ft.	TEMP. Avg.	DELTA "H2O	Q cfm	Y
5.185	80.0	75.0	1.00	10	5.268	73.5	1.5	0.519	1.025
5.237	82.0	75.0	1.00	10	5.268	71.5	1.5	0.524	1.020
5.257	82.0	75.0	1.00	10	5.256	71.5	1.5	0.526	1.014

AVERAGES 0.523 1.020

PASS - INDIVIDUAL Yd VALUES ACCEPTABLE

PASS - INDIVIDUAL DELTA H@ VALUES ACCEPTABLE

PASS - POST TEST Yd WITHIN LIMITS

ES-31

YD = 1.0064

$\Delta H @ = 2.11$

$\Delta H @ 1.00 = 0.510$

POST-TEST DRY GAS METER CALIBRATION DATA

CALIBRATED BY WM
DATE 5-27-93
FIELD METER ID ES-31
BAROMETRIC PRESS 29.897

TEST METER ID ES-13
TEST METER Y (Y) .9980
TEST METER LAST CAL 4/5/93

TEST PROGRAM PRECEDING CALIBRATION CHECK EL SECONDO / SCE
AVERAGE ΔH FROM TEST RUNS 2.4
MAXIMUM VACUUM FROM TEST RUNS 15"
LEAK CHECK VAC 15" INIT. WM

FIELD METER							TEST METER						
TEMPERATURE							TEMPERATURE						
ΔH	FINAL VOLUME	INITIAL VOLUME	VOLUME (V _w)	INLET (I)	OUTLET (O)	VACUUM	TIME (t)	FINAL VOLUME	INITIAL VOLUME	VOLUME (V _w)	INLET	OUTLET	METER PRESS.
1.0	113. 253	108. 068	5.185	80	75	5"	1:44 / 1:54	5.268	0000	5.268	72	75	1.50
1.0	118. 490	113. 253	5.237	82	75	5"	1:55:25 / 2:05:05	5.268	0000	5.268	70	73	1.50
1.0	123. 747	118. 490	118. 5.257	82	75	5"	2:06:10 / 2:16:10	5.256	0000	5.256	70	73	1.50

APPENDIX C
DATA SHEETS

Appendix C.1
Sample Locations

CARNOT

SAMPLING POINT LOCATION DATA - EPA METHOD 1

PLANT: SCE / El Segundo / UNIT 4

DATA BY: KRM/R

DATE: 11/20/92 - 11/22/92

TEST LOCATION: STACK

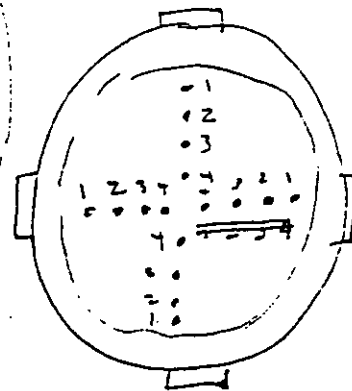
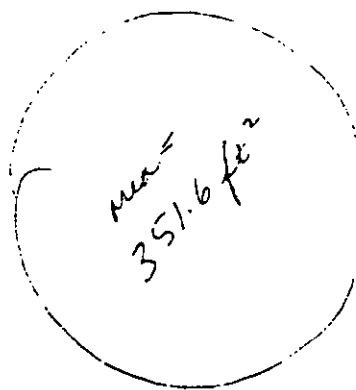
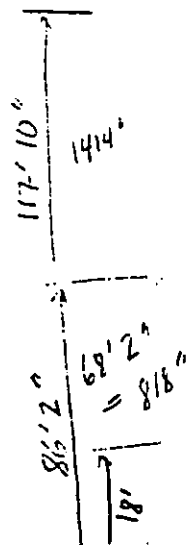


Diagram of Sampling Location

UPSTREAM DIST/DIA: 1414 / 5.57

DOWNSTREAM DIST/DIA: 818 / 3.22

COUPLING LENGTH: 10"

NO. OF SAMPLING PTS.: 16

STACK DIMENSION: 253.9" 21' 2"

STACK AREA, FT²: 351.6 ft²

SAMPLE POINT	% OF DIAMETER	IN. FROM NEAR WALL	IN. FROM NOZZLE*
1	3.2	8.125	26.125
2	10.5	26.667	44.667
3	17.4	49.257	67.257
4	32.3	82.010	109.01

*INCHES FROM WALL PLUS
COUPLING LENGTH

CARNOT

Appendix C.2
Unit Data

CARNOT

1

UNIT DATA

Client: SCELocation: EL SEGUNDOUnit: EL SEG. #4Data by: M. ESCARCEGA

TEST #	4-2	4-3	4-4	4-5
DATE	5/24/93	5/24/93	5/24/93	5/24/93
TIME	1115	1200	1345	1435
LOAD MW	151	150	150	-150
AUX. LOAD	5.6	5.8	5.7	5.4
UREA ON/OFF	OFF	OFF	ON	ON
UREA LOOP /MODE	—	—	3/3	3/3
UREA FLOW GPM	—	—	0.70	.68
H ₂ O FLOW /PRESS. GPM	—	—	4.6/64.9	4.4/64.0
O ₂ % NORTH	2.0	2.2	2.2	2.2
O ₂ % SOUTH	2.2	2.2	2.3	2.3
STM. PRES PSI	2201	2200	2200	2200
STM FLOW KLB/HR	942	945	939	938
FDWTR FLOW KLB/HR	1064	1063	1050	1050
FUEL FLOW 10 ⁶ SCF	1425	1421	1415	1432
FUEL FLOW %	40.8	41.0	40.6	40.2
AIR FLOW %	41.0	40.9	41.0	40.3
SH STM TEMP °F	1049	1050	1049	1050
RH STM TEMP °F	998	1000	998	1001
FD FAN % NORTH	—	—	—	—
FD FAN % SOUTH	—	—	—	—
FD FAN AMPS NORTH	135	134	134	137
FD FAN AMPS SOUTH	119	119	118	119

CARNOT

2

	4-1	4-2	4-3	
O ₂ /CO BIAS SET POINT	0	0	0	0
BLR MASTER	42.02	41.49	41.17	41.34
FGR AMPS NORTH	55	55	55	55
FGR AMPS SOUTH	54	54	54	54
FGR OUTLET DAMPER NORTH	—	—	—	—
FGR OUTLET DAMPER SOUTH	—	—	—	—
FGR INLET DAMPER NORTH	—	—	—	—
FGR INLET DAMPER SOUTH	—	—	—	—
BNR TILTS NORTH 1-4	-19/-32/-19/-22	-19/-20/-22	-19/-20/-22	-16/-30/-17/-19
BNR TILTS SOUTH 1-4	-24/-26/-25/-26	-25/-27/-25/-26	-23/-26/-24/-24	-18/-19/-18/-18
WP/FURN dP N/S	3.9/4.3	4.0/4.3	4.2/4.3	4.1/4.3
BOOS	NONE	NONE	NONE	NONE
APH OUT °F NORTH	216	216	218	219
APH OUT °F SOUTH	186	186	187	187

NOTES:

CARNOT

1

UNIT DATA

Client: SCELocation: EL SEGUNDOUnit: EL SEG. #4Data by: M. ESCOBAR

TEST #	4-6	4-7	4-8	4-9
DATE	5/23/93	5/25/93	5/25/93	5/25/93
TIME	1217	1115	1210	1250
LOAD MW	245	242	246	- 246
AUX. LOAD	5.0	5.0	5.0	5.1
UREA ON/OFF	ON	ON	OFF	OFF
UREA LOOP / MODE	1 / 2	1 / 2	—	—
UREA FLOW GPM	1.34	1.34	2	2
H ₂ O FLOW / PRES. GPM / PSI	6.32 / 55.0	5.94 / 55.0	—	—
O ₂ % NORTH	1.7	1.8	1.7	1.7
O ₂ % SOUTH	1.6	1.6	1.6	1.6
STM. PRES PSI	2249	2350	2247	2350
STM FLOW KLB/HR	1520	1515	1527	1517
FDWTR FLOW KLB/HR	1685	1680	1690	1695
FUEL FLOW 10 ⁶ SCF	1270	1225	1235	1215
FUEL FLOW %	63.9	63.5	64.1	64.8
AIR FLOW %	64.9	65.1	63.7	64.7
SH STM TEMP °F	1050	1049	1050	1050
RH STM TEMP °F	1000	1000	1000	999
FD FAN % NORTH	—	—	—	—
FD FAN % SOUTH	—	—	—	—
FD FAN AMPS NORTH	169	171	173	174
FD FAN AMPS SOUTH	168	167	168	169

CARNOT

2

	4-1	4-2	4-3	4-4
O ₂ /CO BIAS SET POINT	$\ominus$	$\ominus$	$\ominus$	$\ominus$
BLR MASTER	65.0	63.9	64.6	64.37
FGR AMPS NORTH	52	52	52	52
FGR AMPS SOUTH	51	51	51	51
FGR OUTLET DAMPER NORTH	—	—	—	—
FGR OUTLET DAMPER SOUTH	—	—	—	—
FGR INLET DAMPER NORTH	—	—	—	—
FGR INLET DAMPER SOUTH	—	—	—	—
BNR TILTS NORTH	-6/-19/-7/-9	-6/-19/-7/-9	-5/-19/-6/-9	-6/-21/-7/-10
BNR TILTS SOUTH	-10/-11/-9/-10	-8/-7/-9/-9	-8/-10/-8/-10	-9/-10/-9/-10
WP/FURN dP N/S	5.9/5.7	5.7/5.6	5.0/5.6	5.0/5.6
BOOS	NONE	NONE	NONE	NONE
APH OUT °F NORTH	242	241	240	241
APH OUT °F SOUTH	210	210	209	209

NOTES:

CARNOT

1

UNIT DATA

Client: SCELocation: EL SEGUNDOUnit: EL SEG #4Data by: L. Milleron

TEST #	4-10	4-11	4-12	4-13
DATE	8-12-93	8-12-93	8-12-93	8-12-93
TIME	10:45	11:30	12:20	13:00
LOAD MW	331	330	330	332
AUX. LOAD	8.7	8.7	8.7	8.7
UREA ON/OFF	ON	ON	OFF	OFF
UREA LOOP / Mode	1/1	1/1	—	—
UREA FLOW GPM	2.24	2.24	0	0
H ₂ O FLOW GPM	4.38	4.42	—	—
O ₂ % NORTH	1.7	1.6	1.7	1.7
O ₂ % SOUTH	2.1	1.9	1.9	2.0
STM. PRES PSI	2400	2398	2400	2400
STM FLOW KLB/HR	2259	2258	2258	2258
FDWTR FLOW KLB/HR	2446	2442	2436	2482
FUEL FLOW 10 ⁶ SCF	3124	3150	3150	3152
FUEL FLOW %	87.2	87.2	87.1	88.1
AIR FLOW %	91	91	91	91
SH STM TEMP °F	1050	1050	1050	1050
RH STM TEMP °F	1002	1003	1003	1003
FD FAN % NORTH	—	—	—	—
FD FAN % SOUTH	—	—	—	—
FD FAN AMPS NORTH	271	279	264	265
FD FAN AMPS SOUTH	270	277	269	262

SCREENS 164H

CARNOT

2

	4-10	4-11	4-12	4-13
O ₂ /CO BIAS SET POINT	0	0	0	0
BLR MASTER	86.25	86.98	78.23	88.66
FGR AMPS NORTH	59	58	59	61
FGR AMPS SOUTH	55	54	56	56
FGR OUTLET DAMPER NORTH	-	-	-	-
FGR OUTLET DAMPER SOUTH	-	-	-	-
FGR INLET DAMPER NORTH	-	-	-	-
FGR INLET DAMPER SOUTH	-	-	-	-
BNR TILTS NORTH	10/3/11/9	10/3/11/8	10/3/11/9	10/3/11/8
BNR TILTS SOUTH	5/6/11/4	4/11/11/9	5/6/12/5	6/8/11/6
WP/FURN dP	7.7	7.8	7.6	7.6
BOOS	2N 45	2N 45	2N 45	2N 45
APH OUT °F NORTH	270	270	268	268
APH OUT °F SOUTH	231	232	230	230

NOTES:

Appendix C.3
CEM Data

CARNOT

PAGE 1 OF 1

REFERENCE METHOD GASEOUS MEASUREMENTS

IDENT/LOCATION: SCE / EL SEGUNDO CACONDITION: ~150 MWDATE: 5-24-93OPERATOR: McLureTEST NUMBER: 4-1-CRMTEST LOCATION: STALL

TEST NO.	TIME.		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO ____%____, DRY			
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION				5.07	14.97	55.07	51.96						
			AS FOUND ANAL. SPAN	5.03	15.0	55.0	52.0						
			SYSTEM ZERO	0.0	0.0	0.0	0.9						
			SYSTEM SPAN	5.08	14.91	55.0	49.9						
	BEGIN	1000	S-4	6.0	8.5	0	40.3						
	END	1002											
	BEGIN	1002	S-3	6.0	8.5	0	40.3						
	END	1004											
	BEGIN	1004	S-2	6.0	8.5	0	40.5						
	END	1006											
	BEGIN	1006	S-1 *	7.2	7.9	0	38.0						
	END	1008											
	BEGIN	1012	W-4	5.9	8.6	0	41.1						
	END	1014											
	BEGIN	1014	W-3	6.0	8.6	0	41.0						
	END	1018											
	BEGIN	1018	W-2	5.9	8.6	0	41.0						
	END	1020											
	BEGIN	1020	W-1	5.9	8.6	0	41.0						
	END	1022											
	BEGIN	1022	N-4	6.0	8.6	0	41.3						
	END	1024											
	BEGIN	1024	N-3	6.0	8.6	0	41.5						
	END	1026											
	BEGIN	1030	N-2	6.0	8.6	0	41.5						
	END	1032											
	BEGIN	1032	E-4	6.0	8.6	0	41.0						
	END	1034											
	BEGIN	1034	E-3	6.0	8.6	1.0	41.0						
	END	1036											
	BEGIN	1043	E-2	6.0	8.6	0	41.0						
	END	1045											
			SYSTEM ZERO	0.0	0.0	0.0	1.4						
			SYSTEM SPAN	5.08	14.8	55.0	50.0						
RAW AVERAGE				5.98	8.58	0.2	41.0						
CORRECTED AVERAGE				5.92	8.64	0.2	42.5						
COMMENTS:													

CARNOT

CARNOT REFERENCE METHOD GASEOUS MEASUREMENTS

PAGE 1 OF

CLIENT/LOCATION: SEE EL SEGUNDO 4 2

CONDITION: ~150 MW

DATE: 5-24-93

OPERATOR: MCCUNE

TEST NUMBER: 4-2-CEN

TEST LOCATION: STACK

TEST NO.	TIME		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO % DRY			
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION				5.032	14.97	55.07	51.90						
			AS FOUND ANAL. SPAN										
			SYSTEM ZERO	0.0	0.0	0.0	0.0						
			SYSTEM SPAN	5.03	14.8	55.0	50.0						
	BEGIN	1115	S-4 ↓	6.1	8.5	0	42.5						
	END	1120											
	BEGIN	1120		6.1	8.5	0	42.5						
	END	1125											
	BEGIN	1125		6.1	8.5	0	42.5						
	END	1130											
	BEGIN	1130		6.1	8.5	0	42.5						
	END	1135											
	BEGIN	1135		6.1	8.5	0	42.3						
	END	1140											
	BEGIN	1140		6.1	8.5	0	42.1						
	END	1145											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
			SYSTEM ZERO	0.0	0.0	0.0	0.0						
			SYSTEM SPAN	5.13	14.8	54.9	50.0						
RAW AVERAGE				6.1	8.5	0	42.4						
CORRECTED AVERAGE				6.0	8.6	0	44.1						
COMMENTS:													

CARNOT REFERENCE METHOD GASEOUS MEASUREMENTS

PAGE 1 OF

CLIENT/LOCATION: SCE/EL Segundo 4

CONDITION: ~150 MW

DATE: 5-24-93

OPERATOR: McLure

TEST NUMBER: 4-3-CEN

TEST LOCATION: STACK

TEST NO.	TIME		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO % ____ DRY			
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION				5.032	14.97	53.57	51.96						
			AS FOUND ANAL. SPAN										
			SYSTEM ZERO	0.0	0.0	0.0	0.0						
			SYSTEM SPAN	5.03	14.8	55.0	50.0						
	BEGIN	1200	S-4										
	END	1205		6.1	8.5	0	42.2						
	BEGIN	1205											
	END	1210		6.1	8.5	0	42.2						
	BEGIN	1210											
	END	1215		6.1	8.5	0	42.0						
	BEGIN	1215											
	END	1220		6.1	8.5	0	41.7						
	BEGIN	1220											
	END	1225		6.1	8.5	0	41.9						
	BEGIN	1225											
	END	1230		6.1	8.5	0	42.1						
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
			SYSTEM ZERO	0.0	0.0	0.0	-0.1						
			SYSTEM SPAN	5.18	14.8	54.5	50.0						
RAW AVERAGE				6.1	8.5	0.0	42.0						
CORRECTED AVERAGE				6.0	8.6	0.0	43.7						
COMMENTS:													

CARNOT

CARNOT

REFERENCE METHOD GASEOUS MEASUREMENTS

PAGE 1 OF 1

CLIENT/LOCATION: SCE / EL SEGUNDO 4

CONDITION: ~150 MW w/ UREA

DATE: 5-24-93

OPERATOR: McLure

TEST NUMBER: 4-4-CM

TEST LOCATION: Stack

TEST NO.	TIME		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO % DRY			
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION				5.022	14.97	55.07	51.96						
			AS FOUND ANAL. SPAN										
			SYSTEM ZERO	0.0	0.0	0.0	-0.1						
			SYSTEM SPAN	5.18	14.8	54.5	50.0						
	BEGIN	1345	S-4										
	END	1350		6.2	8.5	0	36.8						
	BEGIN	1350											
	END	1355		6.2	8.5	0	37.1						
	BEGIN	1355											
	END	1400		6.2	8.5	0	37.1						
	BEGIN	1400											
	END	1405		6.1	8.5	0	37.0						
	BEGIN	1405											
	END	1410		6.1	8.5	0	37.0						
	BEGIN	1410	V										
	END	1415		6.1	8.5	0	37.0						
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
	BEGIN	----											
	END	----											
			SYSTEM ZERO	0.0	0.0	0.0	-0.5						
			SYSTEM SPAN	5.12	14.8	55.0	49.8						
RAW AVERAGE				6.15	8.5	0	37.0						
CORRECTED AVERAGE				6.0	8.6	0	38.6						
COMMENTS:													

REFERENCE METHOD GASEOUS MEASUREMENTS

CLIENT/LOCATION: SCE / EL SEGUNDO 4CONDITION: 2150 kWDATE: 5-24-93

OPERATOR: _____

TEST NUMBER: 4-5-221

TEST LOCATION: _____

TEST NO.	TIME	SAMPLE POINT/ CONDITION	DRY, UNCORRECTED							CORRECTED TO ____%____, DRY		
			O ₂	CO ₂	CO	NO _x	NO	NO ₂	SO ₂	CO	NO _x	SO ₂
SPAN GAS CONCENTRATION			5.032	14.97	55.07	51.96						
		AS FOUND ANAL. SPAN										
		SYSTEM ZERO	0.0	0.0	0.0	-0.5						
		SYSTEM SPAN	5.12	14.9	55.0	49.8						
	BEGIN <u>1435</u> END <u>1440</u>	S-4	6.15	8.5	0	37.8						
	BEGIN <u>1440</u> END <u>1445</u>		6.15	8.5	0	38.0						
	BEGIN <u>1445</u> END <u>1450</u>		6.15	8.5	0	38.5						
	BEGIN <u>1450</u> END <u>1455</u>		6.15	8.5	0	38.5						
	BEGIN <u>1455</u> END <u>1500</u>		6.15	8.5	0	38.2						
	BEGIN <u>1500</u> END <u>1505</u>	↓	6.15	8.5	0	38.2						
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
		SYSTEM ZERO	0.0	0.0	0.0	-0.5						
		SYSTEM SPAN	5.12	14.9	55.0	49.8						
RAW AVERAGE			6.15	8.5	0	38.2						
CORRECTED AVERAGE			6.0	8.6	0	40.0						
COMMENTS:												

REFERENCE METHOD GASEOUS MEASUREMENTS

CLIENT/LOCATION: SCE / EL SEGUNDO 4CONDITION: ~ 240 MW w/ UREADATE: 5-25-93OPERATOR: McCuneTEST NUMBER: 4-6-LEMTEST LOCATION: STACK

TEST NO.	TIME		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO ____%____, DRY			
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION				5.032	14.97	55.07	51.96						
			AS FOUND ANAL. SPAN	5.03	15.0	55.0	52.0						
			SYSTEM ZERO	0.0	0.0	0.2	0.2						
			SYSTEM SPAN	5.12	14.9	55.1	50.4						
	BEGIN END	10:15 10:17	S-4	5.0	9.1	14	53.5						
	BEGIN END	10:17 10:19	S-3	5.0	9.1	14	53.5						
	BEGIN END	10:19 10:21	S-2	5.0	9.1	19	53.0						
	BEGIN END	10:21 10:23	S-1	5.0	9.1	17	52.5						
	BEGIN END	10:25 10:27	E-4	5.05	9.1	17	53.8						
	BEGIN END	10:27 10:29	E-3	5.0	9.1	18	53.8						
	BEGIN END	10:29 10:31	E-2	5.0	9.1	17	52.8						
	BEGIN END	10:31 10:33	E-1	5.05	9.1	14	53.3						
	BEGIN END	10:35 10:37	N-4	4.95	9.1	18	53.1						
	BEGIN END	10:37 10:39	N-3	5.0	9.1	20	53.2						
	BEGIN END	10:39 10:41	N-2	5.0	9.1	22	53.0						
	BEGIN END	10:41 10:43	N-1	5.0	9.1	18	54.0						
	BEGIN END	10:45 10:47	W-4	4.95	9.1	18	53.0						
	BEGIN END	10:47 10:49	W-3	5.0	9.1	17	53.9						
	BEGIN END	10:49 10:51	W-2	4.95	9.1	18	54.5						
	BEGIN END	10:51 10:53	W-1	4.95	9.1	22	52.8						
			SYSTEM ZERO	0.0	0.0	0.2	0.0						
			SYSTEM SPAN	5.1	14.8	55.2	49.5						
RAW AVERAGE				5.0	9.1	17.7	53.4						
CORRECTED AVERAGE				4.9	9.2	17.5	55.4						
COMMENTS:													

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PAGE 1 OF 1

REFERENCE METHOD GASEOUS MEASUREMENTS

CLIENT LOCATION: SCC/EL SEGUNDO 4CONDITION: 240 MW w/ OPENDATE: 5-25-13OPERATOR: McCuneTEST NUMBER: 4-7-BEMTEST LOCATION: STACK

TEST NO.	TIME	SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO % DRY			
			O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION			5.032	14.97	55.07	51.96						
		AS FOUND ANAL. SPAN										
		SYSTEM ZERO	0.0	0.0	0.2	0.0						
		SYSTEM SPAN	5.1	14.8	55.2	49.5						
	BEGIN 1115 END 1120	W-4	4.95	9.1	24	52.5						
	BEGIN 1120 END 1125		4.95	9.1	23	52.0						
	BEGIN 1125 END 1130		4.95	9.1	19	52.5						
	BEGIN 1130 END 1135		5.0	9.1	17	52.2						
	BEGIN 1135 END 1140		5.0	9.1	17	52.5						
	BEGIN 1140 END 1145	◇	4.95	9.1	18	53.0						
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
	BEGIN END											
		SYSTEM ZERO	0.0	0.0	0.2	0.0						
		SYSTEM SPAN	5.1	14.8	55.0	49.4						
RAW AVERAGE			4.97	9.1	19.7	52.5						
CORRECTED AVERAGE			4.98	9.2	19.5	55.1						
COMMENTS:												

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REFERENCE METHOD GASEOUS MEASUREMENTS

CLIENT LOCATION: SCE / EL SEGUNDO

CONDITION: 240 MW

DATE: 5-25-93

OPERATOR: McLure

TEST NUMBER: 4-B-CEN

TEST LOCATION: Stack

TEST NO.	TIME		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO % DRY				
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂	
SPAN GAS CONCENTRATION				5.032	14.97	55.07	519.6							
			AS FOUND ANAL. SPAN											
			SYSTEM ZERO	0.0	0.0	0.2	0.0							
			SYSTEM SPAN	5.1	14.8	55.0	49.4							
	BEGIN	1210	W-4 / ↓	4.9	9.1	22	63.5							
	END	1215		4.9	9.1	21	63.5							
	BEGIN	1220		4.95	9.1	19	63.8							
	END	1225		4.9	9.1	22	63.5							
	BEGIN	1230		4.9	9.1	22	64.0							
	END	1235		4.9	9.1	22	63.2							
	BEGIN	1235												
	END	1240												
	BEGIN	---												
	END	---												
	BEGIN	---												
	END	---												
	BEGIN	---												
	END	---												
	BEGIN	---												
	END	---												
			SYSTEM ZERO	0.0	0.0	0.2	0.0							
			SYSTEM SPAN	5.1	14.8	55.3	49.8							
RAW AVERAGE				4.9	9.1	21.3	63.6							
CORRECTED AVERAGE				4.8	9.2	21.1	66.6							
COMMENTS:														

CARNOT REFERENCE METHOD GASEOUS MEASUREMENTS

PAGE 1 OF 1

IDENTIFICATION: SCE / EL SEGUNDO 4

CONDITION: 250 MW

DATE: 5-25-93

OPERATOR: McCune

TEST NUMBER: 4-9-C2M

TEST LOCATION: STACK

TEST NO.	TIME		SAMPLE POINT/ CONDITION	DRY, UNCORRECTED						CORRECTED TO ____%____, DRY			
				O ₂	CO ₂	CO	NOx	NO	NO ₂	SO ₂	CO	NOx	SO ₂
SPAN GAS CONCENTRATION				5.032	14.97	55.07	51.96						
			AS FOUND ANAL. SPAN										
			SYSTEM ZERO	0.0	0.0	0.2	0.0						
			SYSTEM SPAN	5.1	14.9	55.3	49.9						
	BEGIN	1250	W-4	4.9	9.2	23	64.2	-					
	END	1255											
	BEGIN	1255											
	END	1300											
	BEGIN	1300											
	END	1305											
	BEGIN	1305	↓	4.9	9.2	25	64.0						
	END	1310											
	BEGIN	1310											
	END	1315											
	BEGIN	1315											
	END	1320											
	BEGIN	---											
	END	---											
	BEGIN	---											
	END	---											
	BEGIN	---											
	END	---											
	BEGIN	---											
	END	---											
			SYSTEM ZERO	0.0	0.0	0.2	0.0						
			SYSTEM SPAN	5.1	14.8	55.1	49.9						
RAW AVERAGE				4.9	9.2	25.3	64.5						
CORRECTED AVERAGE				4.9	9.3	25.6	67.3						
COMMENTS:													

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Reference Method Gaseous Measurements

Client: SCE EL SEGUNDO		Condition: Urea On			
Unit: #4		Load: 320 MW			
Location: STACK					
		O2%	CO2%	CO ppm	NOx ppm
Analyzer Range:	10.00		100.0	100.0	
Span Value:	4.97		55.6	50.0	
Test #:	4-10-CEM	Time: 1045/1115		Date:	8/10/93
	O2%	CO2%	CO ppm	NOx ppm	
System Zero	0.10		0.0	-0.2	
System Span	5.20		55.0	48.5	
Start	Stop				
1045	1050		4.5	63.0	
1050	1055		5.0	64.2	
1055	1100		5.0	64.5	
1100	1105		4.0	62.5	
1105	1110		5.0	62.0	
1110	1115		5.2	64.0	
Raw Average	5.39		4.7	63.2	
As founds	4.92		55.0	51.0	
System Zero	0.08		0.0	-0.1	
System Span	5.10		54.9	49.0	
Corrected Average	5.21		4.8	64.8	
System Bias Check					
Zero Pre-test	0.01		0.0	0.0	< 5% Pass
Zero Post-test	0.01		0.0	0.0	< 5% Pass
Span Pre-test	0.02		0.0	0.0	< 5% Pass
Span Post-test	0.01		0.0	0.0	<5% Pass

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Reference Method Gaseous Measurements

Client: SCE EL SEGUNDO		Condition: Urea On			
Unit: #4		Load: 320 MW			
Location: STACK					
		O2%	CO2%	CO ppm	NOx ppm
Analyzer Range:	10.00		100.0	100.0	
Span Value:	4.97		55.6	50.0	
Test #:	4-11-CEM	Time:			Date: 8/10/93
		O2%	CO2%	CO ppm	NOx ppm
System Zero		0.08		0.00	-0.10
System Span		5.10		54.90	49.00
Start	Stop				
1130	1135	5.35		5.0	62.0
1135	1140	5.40		4.9	64.2
1140	1145	5.40		5.0	65.0
1145	1150	5.42		3.5	68.2
1150	1155	5.40		3.8	67.4
1155	1200	5.38		7.1	61.1
Raw Average		5.39		4.4	65.4
As founds		4.95		55.0	51.0
System Zero		0.10		0.0	0.0
System Span		5.15		55.0	49.0
Corrected Average		5.24		4.5	66.7
System Bias Check					
Zero Pre-test		0.01		0.0	0.0
Zero Post-test		0.01		0.0	0.0
Span Pre-test		0.01		0.0	0.0
Span Post-test		0.02		0.0	0.0
					< 5% Pass
					< 5% Pass
					< 5% Pass
					< 5% Pass

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Reference Method Gaseous Measurements

Client: SCE EL SEGUNDO		Condition: Urea Off					
Unit: #4		Load: 320 MW					
Location: STACK							
		O2%	CO2%	CO ppm	NOx ppm		
Analyzer Range:		10.00		100.0	100.0		
Span Value:		4.97		55.6	50.0		
Test #:		4-13-CEM		Time: 1300/1330		Date: 8/10/93	
		O2%	CO2%	CO ppm	NOx ppm		
System Zero		0.10		0.0	0.1		
System Span		5.10		55.0	50.0		
Start	Stop						
1300	1305	5.10		7.5	73.5		
1305	1310	5.10		6.0	74.3		
1310	1315	5.10		7.4	74.0		
1315	1320	5.10		7.5	73.5		
1320	1325	5.10		6.6	73.6		
1325	1330	5.12		7.5	74.0		
Raw Average		5.10		7.0	73.8		
As founds		4.95		55.0	51.2		
System Zero		0.10		0.0	0.0		
System Span		5.10		55.0	50.0		
Corrected Average		4.97		7.1	73.8		
System Bias Check							
Zero Pre-test		0.01		0.0	0.0	< 5%	Pass
Zero Post-test		0.01		0.0	0.0	< 5%	Pass
Span Pre-test		0.01		0.0	0.0	< 5%	Pass
Span Post-test		0.01		0.0	0.0	<5%	Pass

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Reference Method Gaseous Measurements

Client: SCE EL SEGUNDO		Condition: Urea Off			
Unit: #4		Load: 320 MW			
Location: STACK					
		O2%	CO2%	CO ppm	NOx ppm
Analyzer Range:	10.00			100.0	100.0
Span Value:	4.97			55.6	50.0
Test #:	4-12-CEM	Time: 1220/1250			Date: 8/10/93
	O2%	CO2%	CO ppm	NOx ppm	
System Zero	0.10		0.0	0.0	
System Span	5.15		55.0	49.0	
Start	Stop				
1220	1225	5.40	5.0	72.0	
1225	1230	5.45	5.8	73.5	
1230	1235	5.45	6.0	72.2	
1235	1240	5.42	6.1	72.1	
1240	1245	5.40	6.5	72.0	
1245	1250	5.32	6.0	73.5	
Raw Average	5.42		5.9	72.4	
As founds	4.92		55.0	51.2	
System Zero	0.10		0.0	0.1	
System Span	5.10		55.0	50.0	
Corrected Average	5.27		5.9	73.1	
System Bias Check					
Zero Pre-test	0.01		0.0	0.0	< 5% Pass
Zero Post-test	0.01		0.0	0.0	< 5% Pass
Span Pre-test	0.02		0.0	0.0	< 5% Pass
Span Post-test	0.01		0.0	0.0	< 5% Pass

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Appendix C.4
Ammonia

CARNOT

SAMPLE TRAIN TEST DATA

CLIENT SCE/EL Spindle

UNIT 4

TEST NO. 4-1-14

METHOD

ROUND PAGE 1 OF 1

PROJECT # 10852

DATE 5/24/93

SAMPLE LOCATION Stack

TEST CONDITION

AMB. TEMP., °F

METER VOL (START/END)

OPERATOR/ASSISTANT EFC

PRE-TEST DATA:

Barometric Press., in. Hg.

Assumed Stack Press.

Assumed Moisture

Assumed Molecular Wt

Assumed ΔP Assumed ΔH

Stack Diameter, in.

Sample Time: Total

per point

Total of Traverso Points

EQUIPMENT INFO:

Meter No.

Meter, Yd.

CFM @ $\Delta H = 1.0$

Pilot ID, Cp

O₂/CO₂ Method

Tollon Connecting

Line (Y/N)

Probe: Matl

Length

Nozzle: Matl

Diam.

No.

Filter: Matl

IMP. Matl WL(End) WL(Start) WL(g)

#1 0.1458 603.4 582.0

#2 " 598.7 597.3

#3 " 546.1 545.6

#4 50 686.4 682.4

#5

Total

POST TEST INFO:

Filter Appearance

Impinger Appearance

Silica Gel Spent (Y/N)

SAMPLE TRAIN LEAK CHECK:

CEM Vac. Pilot Inlet

Pre-Test 0.06 15" = Wks

Post-Test

PRE-TEST CALIBRATION CHECK:

Meter Meter

Imp ΔH Reading In Out

Init.

Final

 $\Delta H =$ x ΔP

SAMPLE POINT	TIME	METER CONDITIONS				TEMPERATURES, °F					O ₂	VAC.	STATIC PRESS. inwg	CHAIN OF CUSTODY INFORMATION
		ΔP	ΔH	METER READING	STACK	PROBE	METER		OVEN	IMP. OUT				
							IN	OUT						
	10:00		2.4	653.58	200	251	71	66	—	RED	—	12.5	—	Impingers Loaded <i>ES</i>
	10:10				199	265			—		—		—	Impingers Recovered <i>ML</i>
	10:20								—		—		—	Filter Loaded
	10:30			660.965					—		—		—	Filter Recovered
									—		—		—	Probe Wash
									—		—		—	TEST SUMMARY
													—	Calculated by:
														Checked by:
														Sample Vol., c.f. <i>6.494</i>
														Stack Press., inwg <i>2.4</i>
														ΔH, inwg
														ΔP, inwg
														Motor Temp., °F <i>66.5</i>
														Stack Temp., °F
														Water Collected, g
														O ₂ /CO ₂
														Comments:

491

Appendix C.5
Non-Methane Hydrocarbon

**CARNOT
TEDLAR BAG SAMPLE DATA**

CLIENT/PROJECT: SCE / E / 50961000

DATE: 8/12/93 UNIT: 4 BY: S. Mulligan

	BAG ID <u>4-10-THC</u>	BAG ID <u>4-11-THC</u>	BAG ID _____
TEST NO.	<u>4-10-THC</u>	<u>4-11-THC</u>	
SAMPLE RATE	<u>.5 c/m</u>	<u>.5 c/m</u>	
START TIME	<u>1050</u>	<u>1135</u>	
STOP TIME	<u>1110</u>	<u>1155</u>	
SAMPLE LOCATION	<u>STACH</u>	<u>STACH</u>	

DATE: 8/12/93 UNIT: 4 BY: S. Mulligan

	BAG ID _____	BAG ID _____	BAG ID _____
TEST NO.	<u>4-12-THC</u>	<u>4-13-THC</u>	
SAMPLE RATE	<u>.5 c/m</u>	<u>.5 c/m</u>	
START TIME	<u>1230</u>	<u>1300</u>	
STOP TIME	<u>1250</u>	<u>1320</u>	
SAMPLE LOCATION	<u>STACH</u>	<u>STACH</u>	

**CARNOT
TEDLAR BAG SAMPLE DATA**

CLIENT/PROJECT: _____

DATE: 5-24-93 UNIT: 4 BY: Wm

	BAG ID <u>1</u>	BAG ID <u>2</u>	BAG ID <u>3</u>

BAG ID	BAG ID	BAG ID	BAG ID
3	2	1	

DATE: 5-24-93 UNIT: 4 BY: LSH

CLIENT/PROJECT:

CARNOT
TEDLAR BAG SAMPLE DATA



APPENDIX D
CALCULATIONS

Appendix D.1
General Emissions Calculations

EMISSION CALCULATIONS

1. Sample Volume and Isokinetics

- a. Sample gas volume, dscf

$$V_{m\ scd} = 0.03342 V_m \left(P_{bar} + \frac{H}{13.6} \right) \left(\frac{T_{ref}}{T_m} \right) (Y)$$

- b. Water vapor volume, scf

$$V_{w\ scd} = 0.0472 V_k \left(\frac{T_{ref}}{528^\circ R} \right)$$

- c. Moisture content, nondimensional

$$B_{wo} = \frac{V_{w\ scd}}{V_{m\ scd} + V_{w\ scd}}$$

- d. Stack gas molecular weight, lb/lb mole

$$MW_{dry} = 0.44 (\%CO_2) + 0.32 (\%O_2) + 0.28 (\%N_2)$$

$$MW_{wet} = MW_{dry} (1 - B_{wo}) + 18 (B_{wo})$$

- e. Absolute stack pressure, iwg

$$P_s = P_{bar} + \frac{P_{sg}}{13.6}$$

- f. Stack velocity, ft/sec

$$V_s = 2.90 C_p \sqrt{\Delta PT_s} \sqrt{\left(\frac{29.92}{P_s} \right) \left(\frac{28.95}{MW_{wet}} \right)}$$

- g. Actual stack flow rate, wacfm

$$Q = (V_s)(A_s)(60)$$

- h. Standard stack gas flow rate, dscfm

$$Q_{sd} = Q (1 - B_{wo}) \left(\frac{T_{ref}}{T_s} \right) \left(\frac{P_s}{29.92} \right)$$

- i. Percent isokinetic

$$I = \left(\frac{17.32 (T_s) (V_{m\ scd})}{(1 - B_{wo}) (\Theta) (V_s) (P_s) (D_n^2)} \right) \left(\frac{528^\circ R}{T_{ref}} \right)$$

2. Particulate Emissions

- a. Grain loading, gr/dscf

$$C = 0.01543 \left(\frac{M_n}{V_{m \text{ std}}} \right)$$

- b. Grain loading at 12% CO
- ₂
- , gr/dscf

$$C_{12\% \text{ CO}_2} = C \left(\frac{12}{\% \text{ CO}_2} \right)$$

- c. Mass emissions, lb/hr

$$M = C(Q_{\text{std}}) \left(\frac{60 \text{ min/hr}}{7000 \text{ gr/lb}} \right)$$

3. Gaseous Emissions, lb/hr

$$M = (\text{ppm})(10^{-6}) \left(\frac{MW_i \text{ lb/lb mole}}{SV} \right) (Q_{\text{std}}) (60 \text{ min/hr})$$

where,

SV = specific molar volume of an ideal gas:

$$SV = 385.3 \text{ ft}^3/\text{lb mole} \quad \text{for } T_{\text{ref}} = 528^\circ \text{R}$$

$$SV = 379.5 \text{ ft}^3/\text{lb mole} \quad \text{for } T_{\text{ref}} = 520^\circ \text{R}$$

4. Emissions Rates, lb/10⁶ Btu

- a. Fuel factor at 68 °F, dscf/10
- ⁶
- Btu at 0% O
- ₂

$$F_{68} = \frac{10^6 [3.64(\%H) + 1.53(\%C) + 0.14(\%N) + 0.57(\%S) - 0.46(\%O_{2, \text{fuel}})]}{HHV, \text{ Btu/lb}}$$

- b. Fuel factor at 60 °F

$$F_{60} = F_{68} \left(\frac{520^\circ \text{R}}{528^\circ \text{R}} \right)$$

- c. Gaseous Emissions factor

$$\left(\frac{\text{lb}}{10^6 \text{ Btu}} \right)_i = (\text{ppm})_i (10^{-6}) \left(\frac{MW_i \text{ lb}}{\text{lb mole}} \right) \left(\frac{1}{SV} \right) (F) \left(\frac{20.9}{20.9 - \%O_2} \right)$$

- d. Particulate emission factor

$$\left(\frac{\text{lb}}{10^6 \text{ Btu}} \right) = C \left(\frac{1 \text{ lb}}{7000 \text{ gr}} \right) (F) \left(\frac{20.9}{20.9 - \%O_2} \right)$$

Nomenclature:

A_s	=	stack area, ft^2
B_{wo}	=	flue gas moisture content
$C_{12\% \text{ CO}_2}$	=	particulate grain loading, gr/dscf corrected to 12% CO_2
C	=	particulate grain loading, gr/dscf
C_p	=	pitot calibration factor, dimensionless
D_n	=	nozzle diameter, in.
F	=	fuel F factor, $\text{dscf}/10^6 \text{ Btu}$ at 0% O_2
H	=	orifice pressure differential, iwg
I	=	% isokinetics
M_s	=	mass of collected particulate, mg
M_i	=	mass emissions of species i , lb/hr
MW	=	molecular weight of flue gas
MW_i	=	molecular weight of species i :
		$\text{NO}_x : 46$
		$\text{CO} : 28$
		$\text{SO}_x : 64$
		$\text{HC} : 16$
θ	=	sample time, min.
ΔP	=	average velocity head, $\text{iwg} = (\sqrt{\Delta P})^2$
P_{bar}	=	barometric pressure, in.Hg
P_s	=	stack absolute pressure, in.Hg
P_{sg}	=	stack static pressure, iwg

Nomenclature (Continued):

Q	=	wet stack gas flow rate at actual conditions, wacfm
Q_{sd}	=	dry stack gas flow rate at standard conditions, dscfm
SV	=	specific molar volume of an ideal gas at standard conditions, ft ³ /lb mole
T_m	=	meter temperature, °R
T_{ref}	=	reference temperature, °R
T_s	=	stack temperature, °R
V_s	=	stack velocity, ft/sec
V_{lc}	=	volume of liquid collected in impingers, ml
V_m	=	dry meter volume uncorrected, dcf
$V_{m\ sd}$	=	dry meter volume at standard conditions, dscf
$V_{w\ sd}$	=	volume of water vapor at standard conditions, scf
Y	=	meter calibration coefficient

Appendix D.2
Specific Emissions Calculations

CARNOT SOURCE TEST DATA SUMMARY

AMMONIA

Client/Location.....	SCE/EL SEG 4	Reference Temp (F).....			60
Unit.....	4	Fuel.....			NAT GAS
Sample Location.....	STACK	Data By.....			MAR
		Date of Data Entry.....			24-May-93
Test No.....	4-4-NH3	4-5-NH3	4-6-NH3	4-7-NH3	
Date.....	24-May-93	24-May-93	25-May-93	25-May-93	
Test Method.....	SCAQMD	SCAQMD	SCAQMD	SCAQMD	
Sample Train.....	ES-31	ES-31	ES-31	ES-31	
Pitot Factor	0.841	0.841	0.841	0.841	
Meter Cal Factor.....	1.02	1.02	1.02	1.02	
Sample Time (Min).....	30	30	30	30	
Bar Press (in Hg).....	29.95	29.95	29.86	29.86	
Start/Stop Time.....	1345/1415	1453/1505	1015/1045	1115/1145	
Stack Press (iwg).....	0.02	0.02	0.02	0.02	
Stack Temp (F).....	203.3	204.7	221.7	223.7	
Stack O2 (%).....	6.00	6.00	4.90	4.90	
Stack CO2 (%).....	8.60	8.60	9.10	9.10	
Meter Vol (acf).....	23.074	23.204	23.270	21.905	
Meter Temp (F).....	70.7	70.0	71.3	70.2	
Meter Press (iwg).....	2.40	2.40	2.33	2.13	
Liquid Vol (ml).....	110.3	96.8	112.0	103.0	
Std Sample Vol (SCF).....	23.218	23.380	23.314	21.983	
Std Sample Vol (Nm^3).....	0.613	0.617	0.615	0.580	
Moisture Fraction.....	0.181	0.161	0.183	0.179	
Ammonia Conc (ug/ml)	3.14	5.62	11.90	9.97	
Sample Vol (ml)	414	432	379	509	
Ammonia Conc (ppmv)	2.80	5.19	9.66	11.54	
Ammonia Conc @ 3% O2	3.36	6.24	10.81	12.91	

CARNOT

8/13/93

Client: SCE		BP in Hg: 29.81		Blank: 0.00	
Date: 8/12/93		Tref R: 520			
Location: Elsegundo 4		Meter No ES-47			
Unit: Natural Gas		Yd: 1.004			

Test #	Vm (adcf)	Tm (F)	Hm (in. H2O)	Tot Vol (ml)	Solution NH3 (ppm)	Vmstd (SCF)	O2 (%)	NH3 (ppm)	NH3 @ 3% O2
4-10-NH3	23.660	65.9	2.50	525	5.05	23.549	5.21	5.5	6.3
4-11-NH3	23.690	69.3	2.50	460	7.20	23.427	5.24	6.9	7.9

15991 Red Hill Ave., Suite 110
Tustin, California 92680-7388
(714) 259-9520
FAX (714) 259-0372

191

APPENDIX E
LABORATORY REPORTS

1:54 PM 7/1/93

AMMONIA DETERMINATION USING NESSLER'S REAGENT

CLIENT: SCE/EL SEG
 LOCATION: UNIT 4
 PROJECT: 10 852

ANALYST: MAR
 CALIBRATION DATE: 5/24/93
 ANALYSIS DATE: 5/24/93
 CONC. OF STD. (ug/ml)= 10.031

CALIBRATION:

Vol. std added (ml)	Expected conc. (ug/ml)	Absorb. (OD)	ABS-BLK	Measured conc. (ug/ml)	%diff	R.F. (mg/OD)
0	0.00	0.300	0.000	---	---	
4	0.80	0.348	0.048	0.75	-6.15%	0.836
6	1.20	0.397	0.097	1.22	1.74%	0.620
10	2.01	0.482	0.182	2.04	1.81%	0.551
12	2.41	0.523	0.223	2.44	1.23%	0.540
14	2.81	0.564	0.264	2.83	0.82%	0.532
16	3.21	0.597	0.297	3.15	-1.89%	0.540
20	4.01	0.670	0.370	3.85	-4.00%	0.542
slope=	0.1039				Avg R.F.=	0.603
y-intercept=	-0.0303					
r2=	0.9979					
					% of std	
RICCA STD	2.42		0.233	2.53	104.68%	

ANALYSIS:

Test no.	Absorbance	Dil factor	Conc (ug/ml)	TV
4-1-NH3 (ABORT)	0.017	1.1	0.51	366
4-4-NH3 DUP	0.126	2.0	3.01	414
4-4-NH3	0.263	1.1	3.14	414
4-5-NH3	0.262	2.0	5.62	432
4-6-NH3	0.217	5.0	11.90	379
4-7-NH3	0.177	5.0	9.97	509
				% diff
Ricca Std Dup	0.244	1.0	2.64	4.18%
Ricca Std Dup	0.249	1.0	2.69	6.08%

SCE El Segundo

8/12/93

std= 9.485

ml added	conc.	abs	abs-blk	conc fnd	% diff
0	0.00	0.041	0.000		
6	1.14	0.119	0.078	1.10	-3.10%
8	1.52	0.171	0.130	1.55	2.31%
10	1.90	0.211	0.170	1.90	0.08%
12	2.28	0.258	0.217	2.30	1.26%
14	2.66	0.299	0.258	2.66	0.14%
16	3.04	0.339	0.298	3.01	-0.98%
18	3.41	0.382	0.341	3.38	-1.09%
20	3.79	0.434	0.393	3.83	0.87%

slope= 0.116
y-int= -0.050
r2= 0.999

RICCA STD 2.44 0.288 0.247 2.56 5.10%

sample #	abs	dil ftr	conc	TV	ug/mL
4-A-NH3	0.095	2.50	3.12	513	0.006091
4-A-NH3	0.238	1.25	3.11	513	0.006059
4-10-NH3	0.248	2.00	5.15	525	0.009802
4-10-NH3	0.411	1.25	4.98	525	0.009482
4-11-NH3	0.369	2.00	7.24	460	0.015737
4-11-NH3	0.447	1.67	7.16	460	0.015558



AtmAA, Inc.

21354 Nordhoff St., Suite 113, Chatsworth, CA 91311 (818) 718-6070 • FAX (818) 718-9779

environmental consultants
laboratory services

June 2, 1993

LTR/201/93

Jody Hone
CARNOT
15991 Red Hill Ave. Suite 110
Tustin, CA 92680

RECEIVED

JUN 03 1993

CARNOT

re: 10852

Dear Jody:

Please find enclosed the laboratory analysis report, quality assurance summary, and the original chain of custody form for four Tedlar bag samples received on May 25, 1993.

The samples were analyzed for methane and total gaseous non-methane organics.

Sincerely,

AtmAA, Inc.

Michael L. Porter
Laboratory Director

Encl.
MLP/krp



AtmAA Inc.

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LABORATORY ANALYSIS REPORT

Methane & Total Gaseous Non-Methane Organics Analysis in Tedlar Bag Samples

Report Date: June 2, 1993
Client: CARNOT
P.O. No.: 9912
Project No.: 10852
Date Received: May 25, 1993
Date Analyzed: May 26, 1993

ANALYSIS DESCRIPTION

Methane and total gaseous non-methane organics were measured by flame ionization detection/total combustion analysis portion (FID/TCA).

AtmAA Lab No.:	91453-38	91453-39	91453-40	91453-41
Sample I.D.:	4-2-HC	4-3-HC	4-4-HC	4-5-HC
Components	(Concentration in ppmv)			
Methane	<1	<1	<1	<1
TGNMO	15.8	19.0	17.1	18.0

TGNMO is total gaseous non-methane organics measured and reported as ppm methane.


Michael L. Porter
Laboratory Director

QUALITY ASSURANCE SUMMARY
(Repeat Analysis)

Project No.: 10852
Date Received: May 25, 1993
Date Analyzed: May 26, 1993

Components	Sample ID	Repeat	Analysis	Mean	% Diff.
		Run #1	Run #2	Conc.	From Mean
(Concentration in ppmv)					
Methane	4-3-HC	< 1	< 1	---	---
TGNMO	4-3-HC	19.2	18.8	19.0	1.0

A set of 4 Tedlar bag samples laboratory numbers, 91453-(38-41) was analyzed for methane and TGNMO. Agreement between repeat analyses is a measure of precision and is shown above in the column "% Difference from Mean." Repeat analyses are an important part of AtmAA's quality assurance program. The average % Difference from Mean for 1 repeat measurement from the sample set of 4 Tedlar bag samples is 1.0%.



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LABORATORY ANALYSIS REPORT

Methane & Total Gaseous Non-Methane Organics Analysis in Tedlar Bag Samples

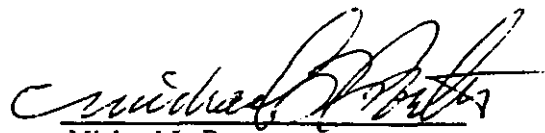
Report Date: June 2, 1993
Client: CARNOT
P.O. No.: 9916
Project No.: 10852
Date Received: May 26, 1993
Date Analyzed: May 26, 1993

ANALYSIS DESCRIPTION

Methane and total gaseous non-methane organics were measured by flame ionization detection/total combustion analysis portion (FID/TCA).

AtmAA Lab No.:	91463-1	91463-2	91463-3	91463-4
Sample I.D.:	4-6-NH3	4-7-NH3	4-8-HC	4-9-HC
Components	(Concentration in ppmv)			
Methane	<1	<1	<1	<1
TGNMO	19.6	23.1	21.2	20.4

TGNMO is total gaseous non-methane organics measured and reported as ppm methane.


Michael L. Porter
Laboratory Director

QUALITY ASSURANCE SUMMARY
(Repeat Analysis)

Project No.: 10852
Date Received: May 26, 1993
Date Analyzed: May 26, 1993

Components	Sample ID	Repeat	Analysis	Mean	% Diff.
		Run #1	Run #2	Conc.	From Mean
(Concentration in ppmv)					
Methane	4-6-HH3	<1	<1	---	---
	4-9-HC	<1	<1	---	---
TGNMO	4-6-HH3	19.1	20.1	19.6	2.6
	4-9-HC	21.7	19.2	20.4	6.1

A set of 4 Tedlar bag samples laboratory numbers, 91463-(1-4) was analyzed for methane and TGNMO. Agreement between repeat analyses is a measure of precision and is shown above in the column "% Difference from Mean." Repeat analyses are an important part of AtmAA's quality assurance program. The average % Difference from Mean for 2 repeat measurements from the sample set of 4 Tedlar bag samples is 4.3%.



AtmAA Inc.

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RECEIVED

AUG 16 1993

August 16, 1993

LTR/305/93

Ed Filadelfia
CARNOT
15991 Red Hill Ave. Suite 110
Tustin, CA 92680

re: 10-852

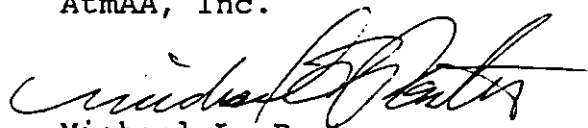
Dear Ed:

Please find enclosed the laboratory analysis report, quality assurance summary, and the original chain of custody form for four Tedlar bag samples received on August 13, 1993.

The samples were analyzed for methane and total gaseous non-methane organics.

Sincerely,

AtmAA, Inc.


Michael L. Porter
Laboratory Director

Encl.
MLP/krp



AtmAA Inc.

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LABORATORY ANALYSIS REPORT

Methane & Total Gaseous Non-Methane Organics Analysis in Tedlar Bag Samples

Report Date: August 16, 1993
Client: CARNOT
P.O. No.: 1175
Project No.: 10-852
Date Received: August 13, 1993
Date Analyzed: August 13, 1993

ANALYSIS DESCRIPTION

Methane and total gaseous non-methane organics were measured by flame ionization detection/total combustion analysis portion (FID/TCA).

AtmAA Lab No.:	92253-11	92253-12	92253-13	92253-14
Sample I.D.:	4-10-HC	4-11-HC	4-12-HC	4-13-HC
Components	(Concentration in ppmv)			
Methane	<1	<1	<1	<1
TGNMO	10.3	7.73	10.0	7.15

TGNMO is total gaseous non-methane organics measured and reported as ppm methane.


Michael L. Porter
Laboratory Director

QUALITY ASSURANCE SUMMARY
(Repeat Analysis)

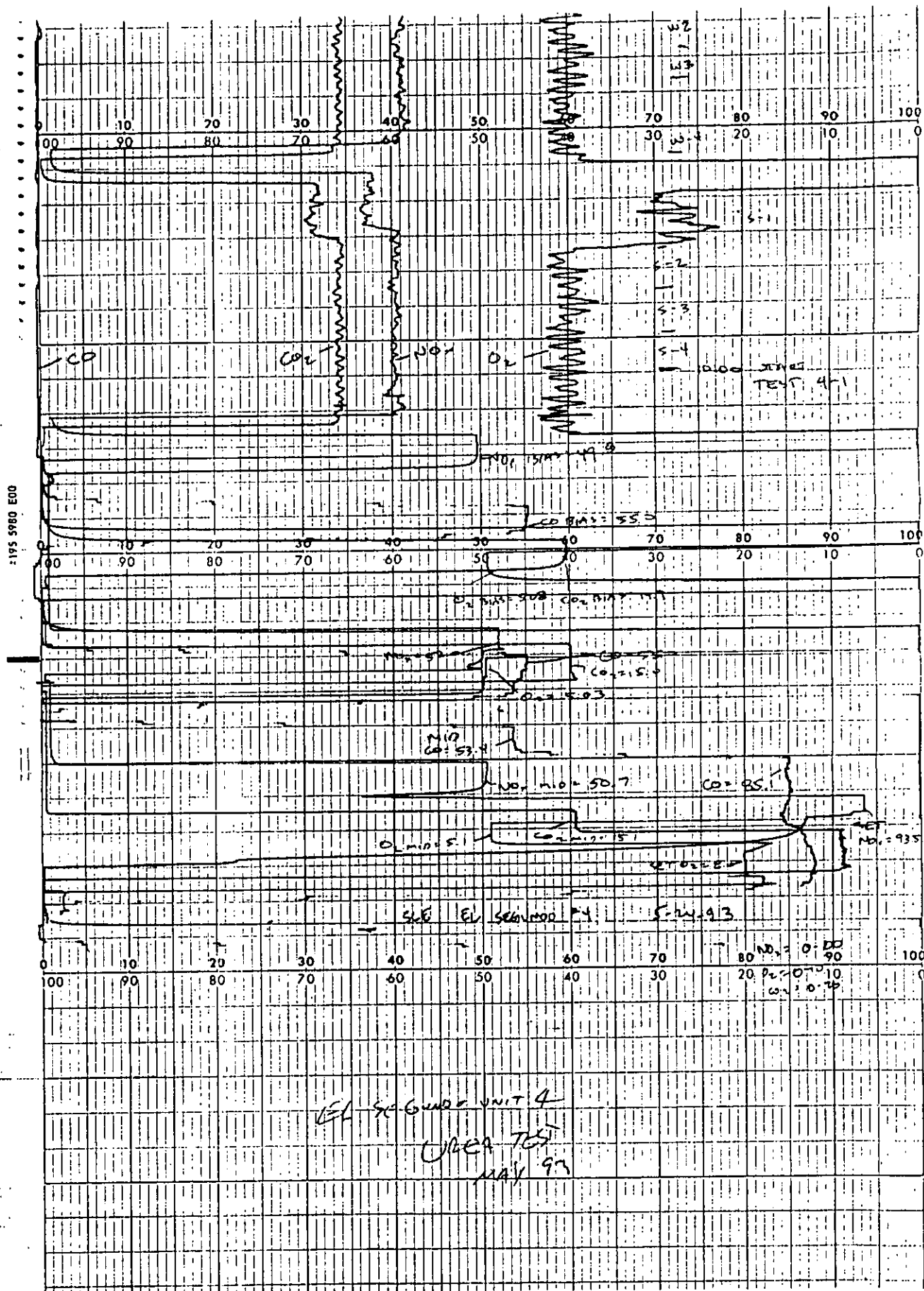
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Date Received: August 13, 1993
Date Analyzed: August 13, 1993

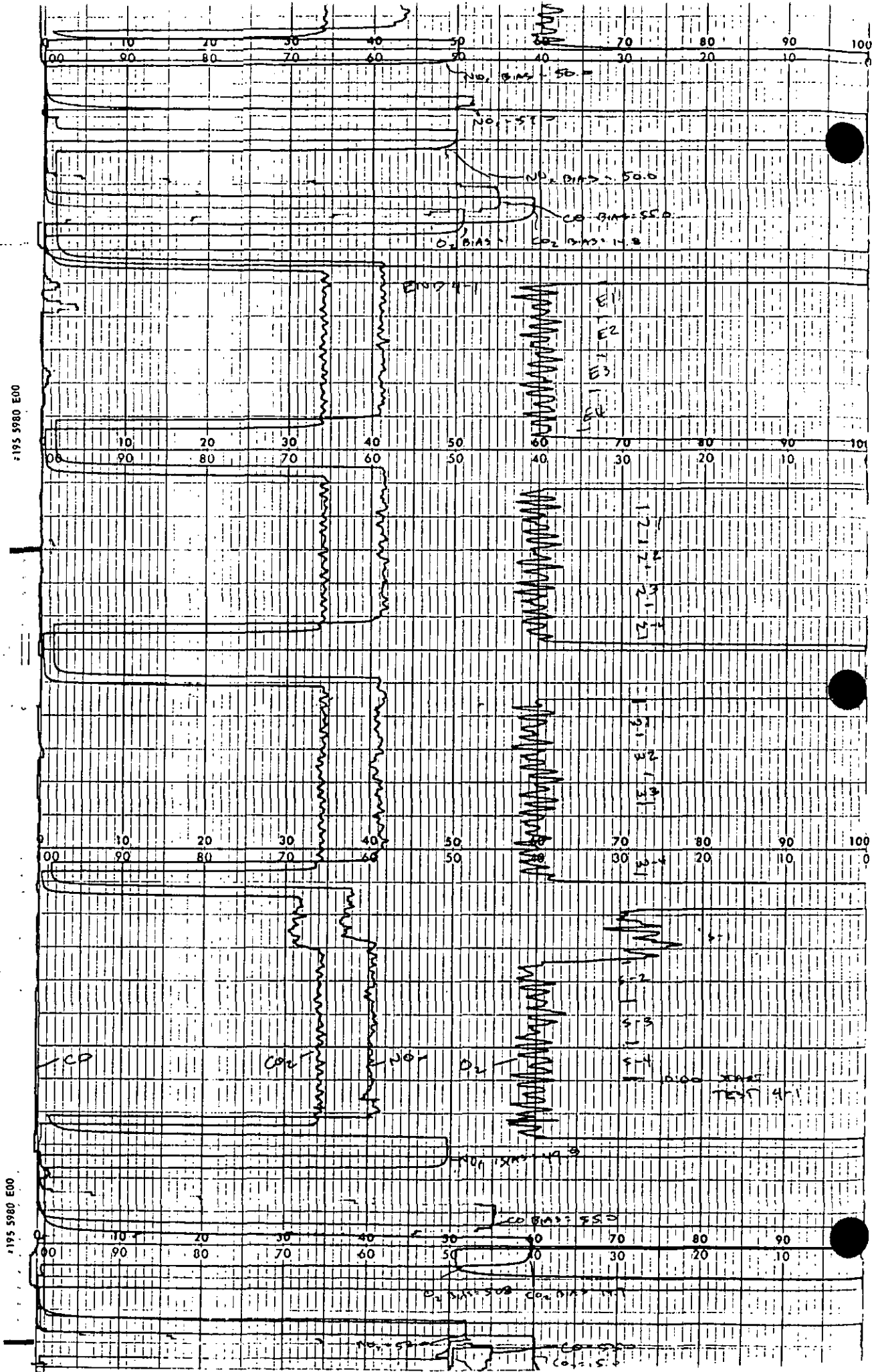
Components	Sample ID	Repeat	Analysis	Mean	% Diff.
		Run #1	Run #2	Conc.	From Mean
(Concentration in ppmv)					
Methane	4-11-HC	<1	<1	---	---
TGNMO	4-11-HC	7.36	8.10	7.73	4.8

A set of 4 Tedlar bag samples laboratory numbers, 92253-(11-14) was analyzed for methane and TGNMO. Agreement between repeat analyses is a measure of precision and is shown above in the column "% Difference from Mean." Repeat analyses are an important part of AtmAA's quality assurance program. The average % Difference from Mean for 1 repeat measurement from the sample set of 4 Tedlar bag samples is 4.8%.



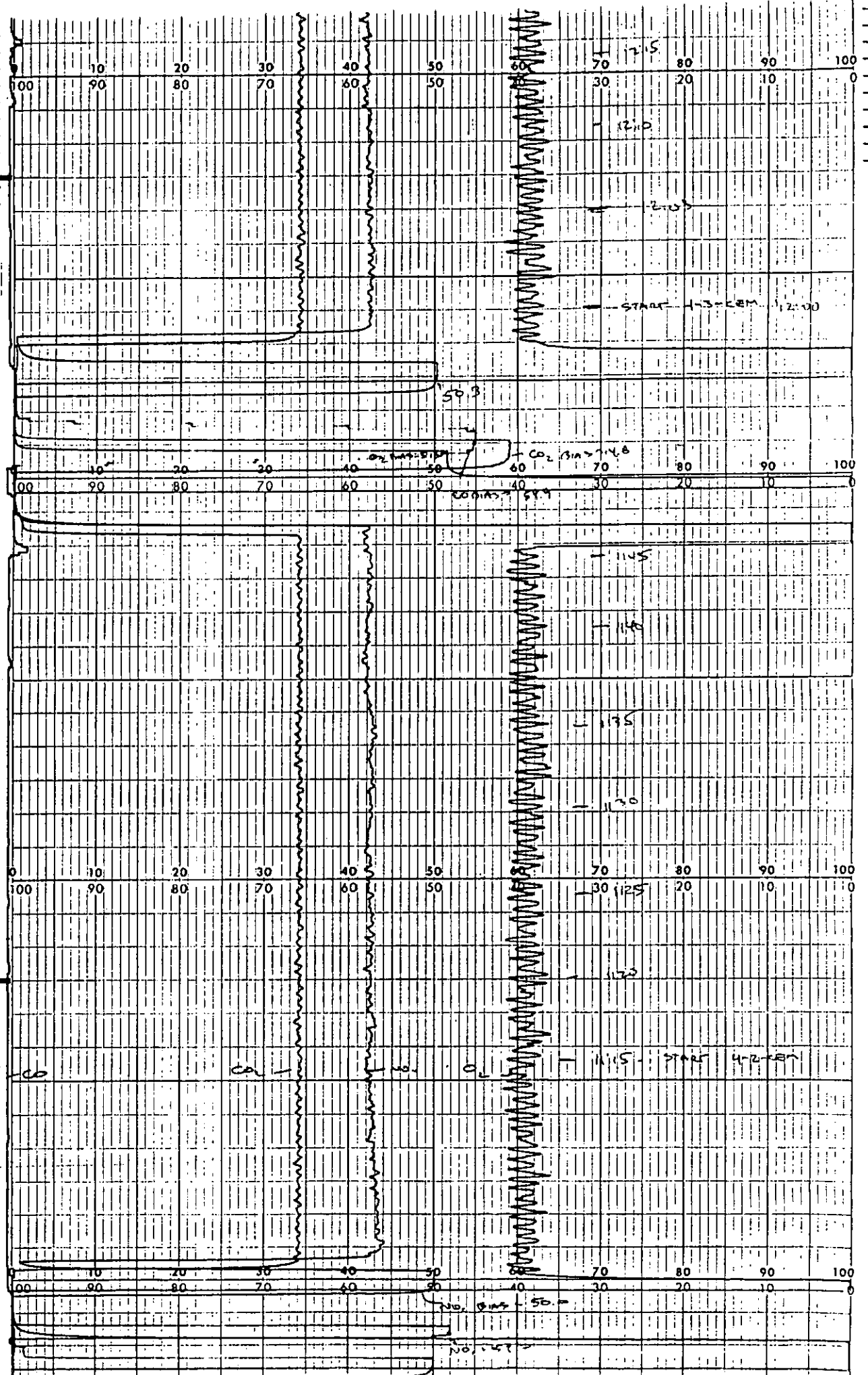
APPENDIX F
INSTRUMENT STRIP CHARTS

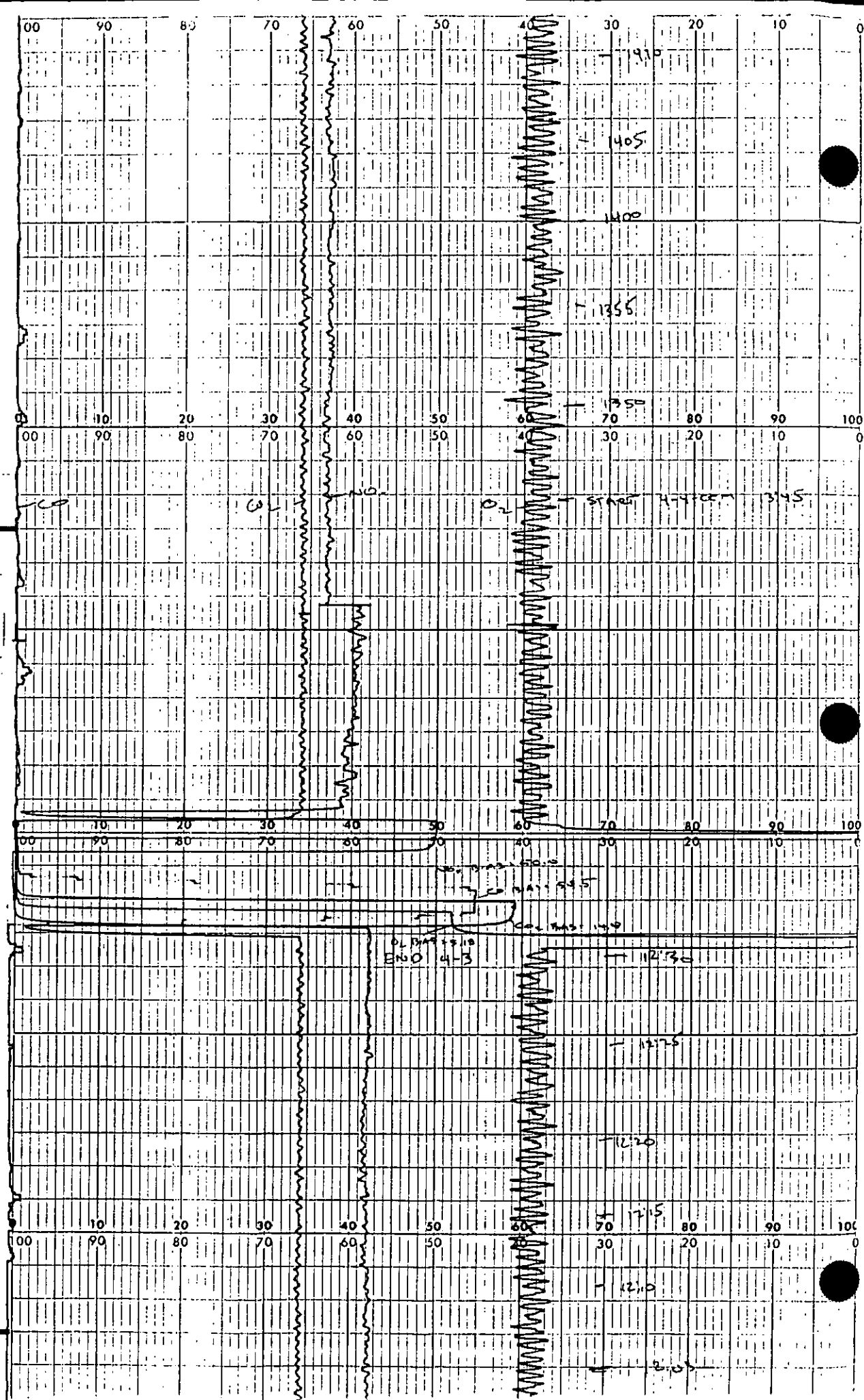
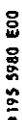




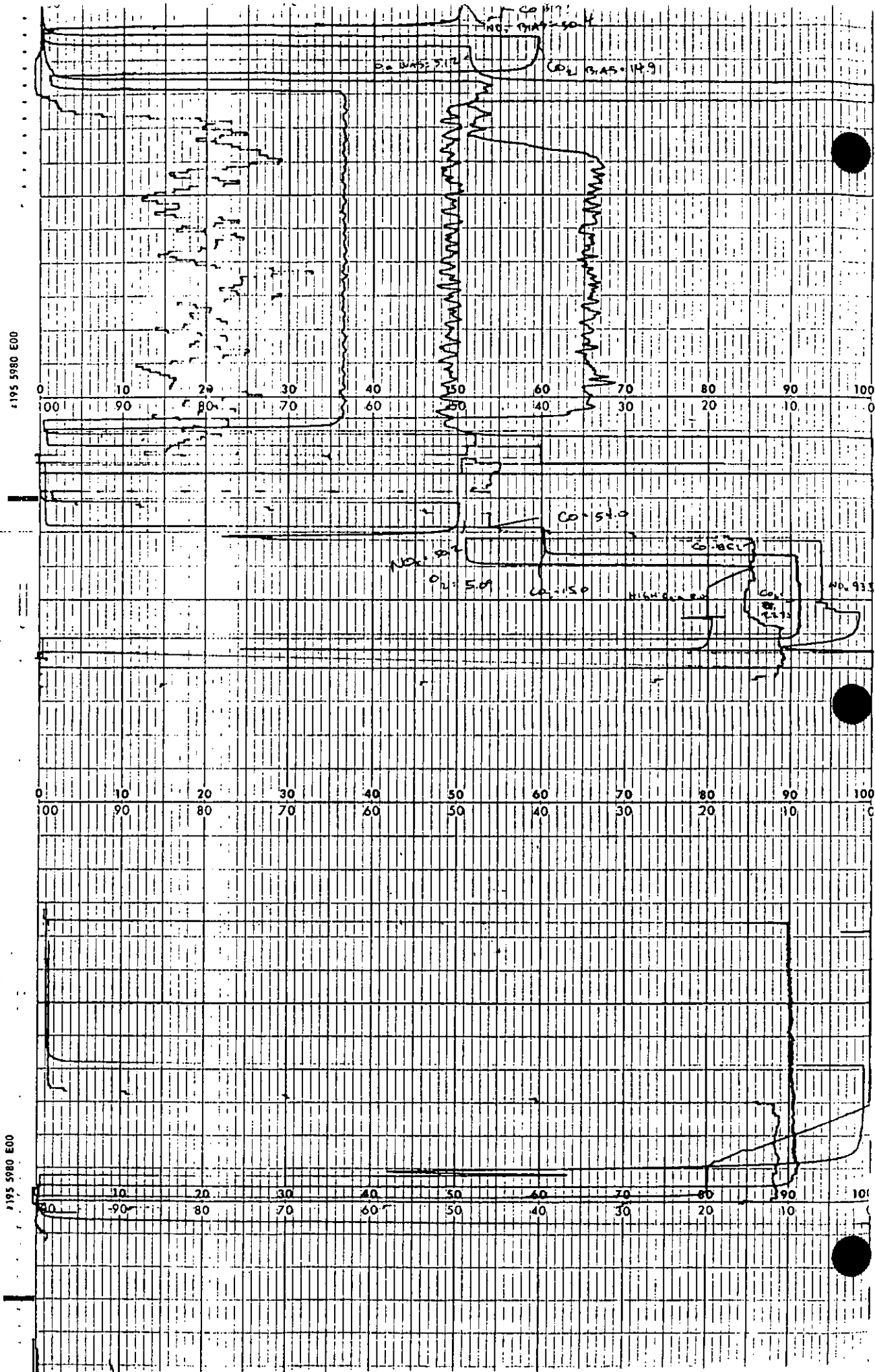
195 5980 E

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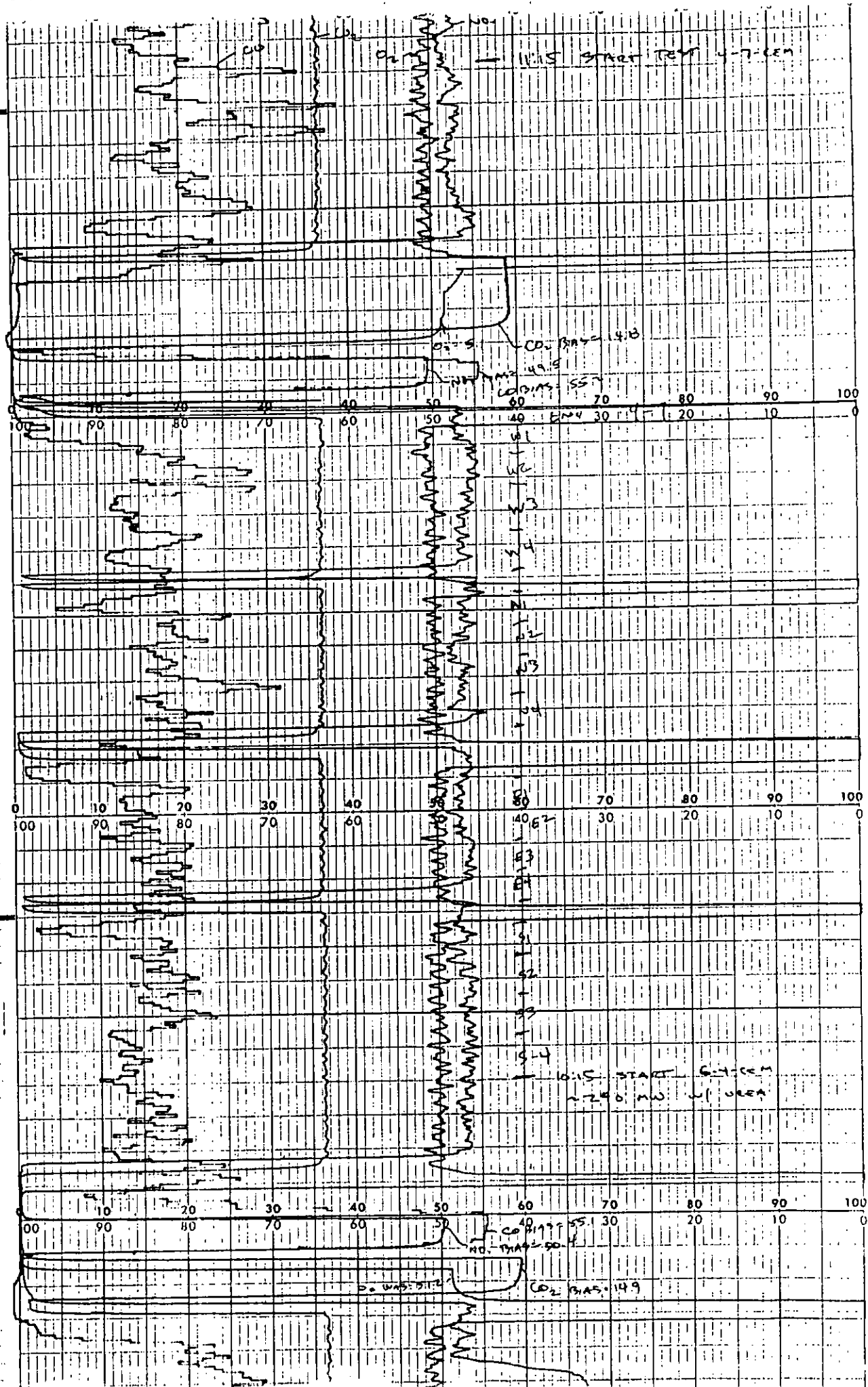




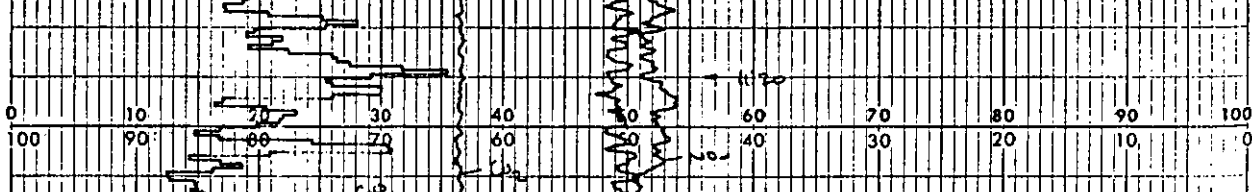
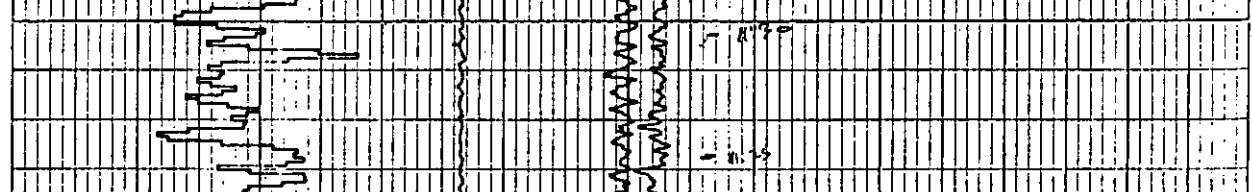
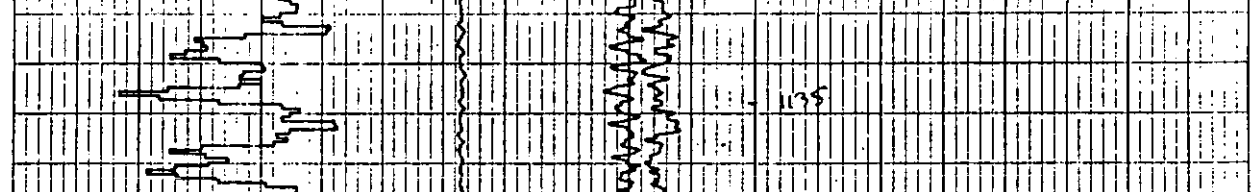
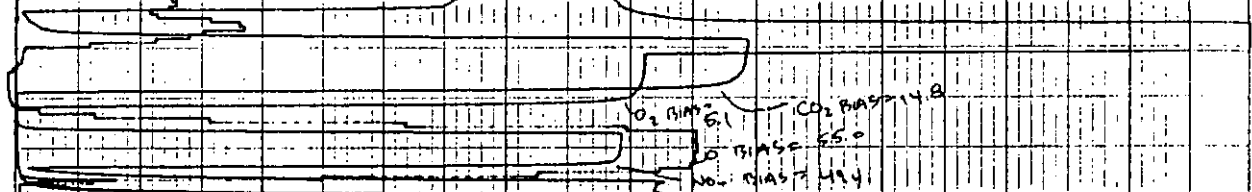
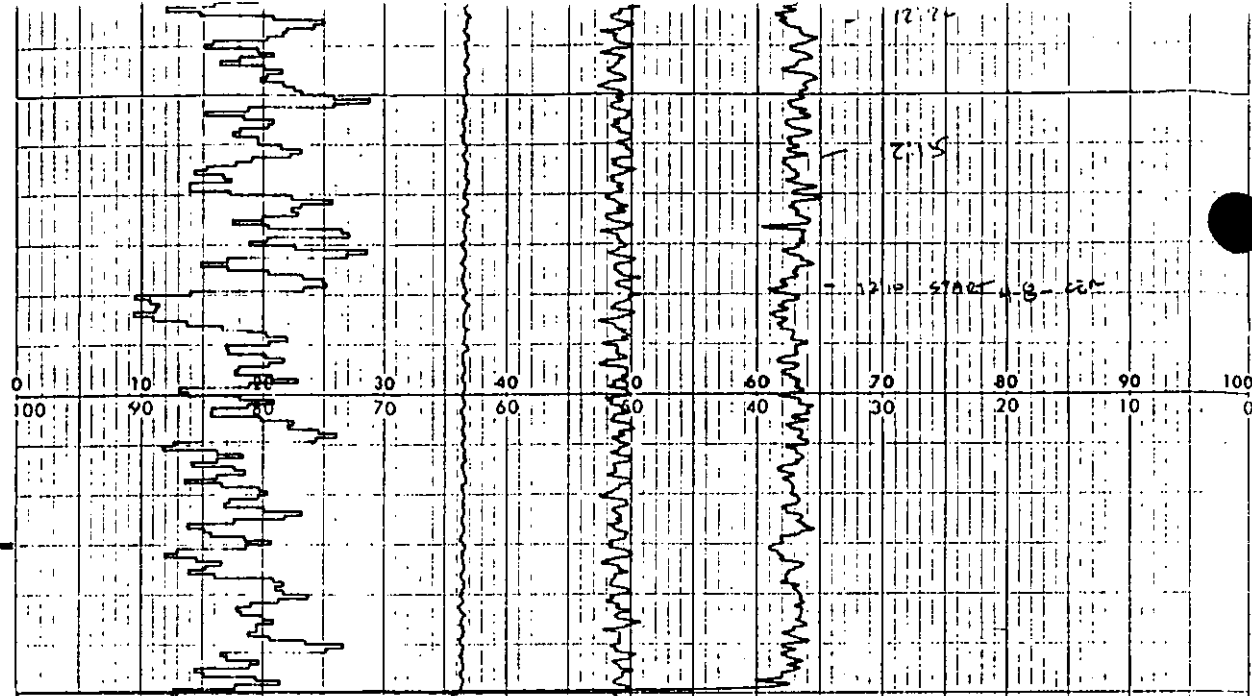
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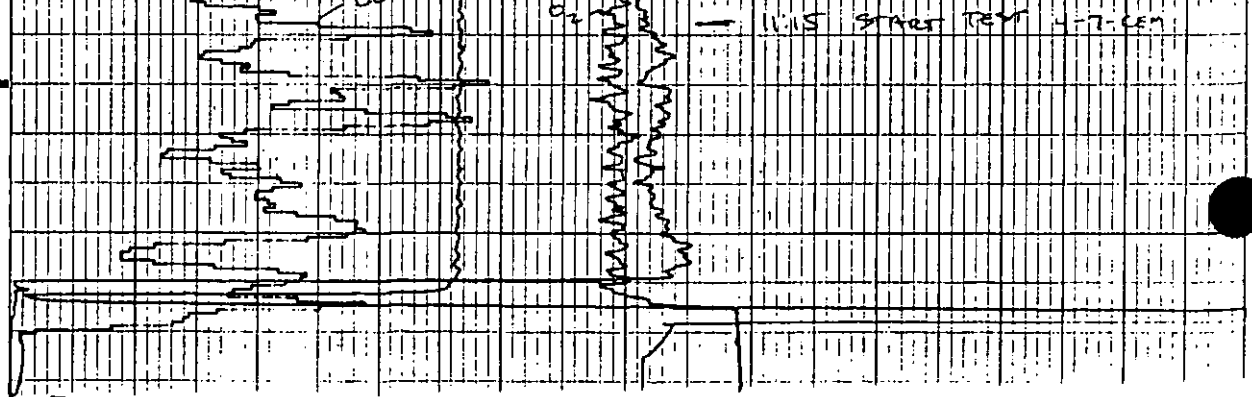
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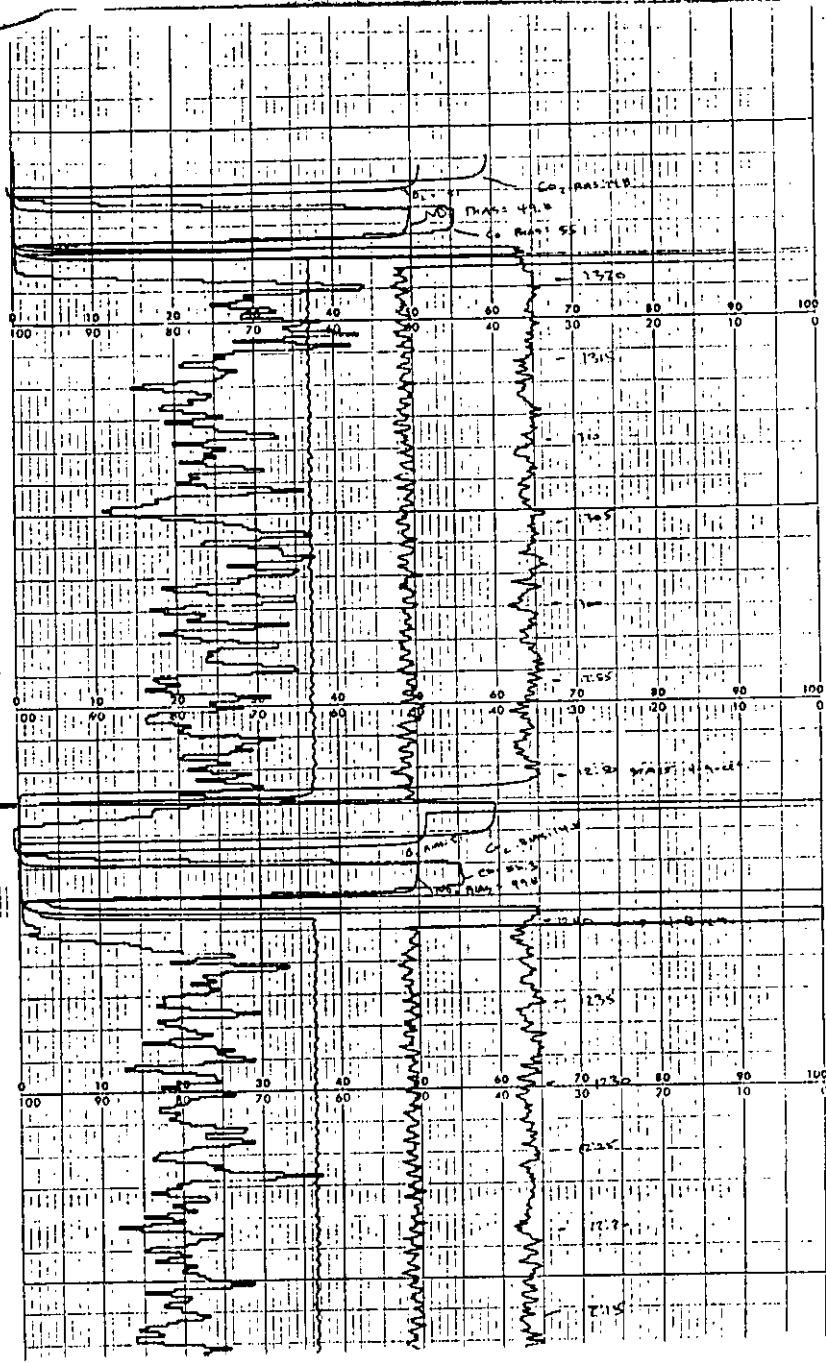
#195 \$980 E00



#195 \$980 E00



1175 5000 500



APPENDIX G
CHAIN OF CUSTODY

SCEJA-10852/R106C688.T

G-1

CANNOT

CHAIN OF CUSTODY

PROJECT # 10852 OUTSIDE LAB REQUIRED (Y/N) Y SAMPLE DATE 5/25/93
 CLIENT/LOCATION EL SEGUNDO PROJECT MANAGER EF
 SAMPLE LOCATION STALK METHOD(S) ZS2 TECHNICIAN WM
 COMPLIANCE TEST (Y/N) Y DATE DUE _____

DATE	TIME	TEST #	SAMPLE DESCRIPTION	CONTAINERS	SAMPLER	COMMENTS
5/25	10:15	4-6	TECLAC BAG		WM	
5/25	11:15	4-7	"	1	WM	
5/25	12:10	4-8	"	1	WM	
5/25	12:50	4-9	"	1	WM	
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					
/	:					

TRANSFER OF SAMPLES FROM FIELD SAMPLE CUSTODIAN (FSC):

RELEASED BY (FSC)	DATE AND TIME	RECEIVED BY	DATE AND TIME
W. R. Madden	5/25/93 4:00	W. R. Madden	5/25/93 4:00
	/ / :		/ / :
	/ / :		/ / :

ANALYSIS REQUIRED: _____

CERTIFICATION OF SAMPLE RECEIPT

PROJECT # 10852 OUTSIDE LAB REQUIRED (Y/N) Y SAMPLE DATE 5/24/93
 CLIENT/LOCATION SCE El Segundo 4 PROJECT MANAGER EF
 SAMPLE LOCATION Stack METHOD(S) SCARMDZS.2 TECHNICIAN WUM
 COMPLIANCE TEST (Y/N) Y DATE DUE 5/2

TEST #	COMPLETE DESCRIPTION
1453-38 4-2-HC	Tedlar bag sample 5/24/93 1115
-39 4-3-HC	" 1200
-40 4-4-HC	" 1345
-41 4-5-HC	" 1435

CHAIN OF CUSTODY PRIOR TO SHIPMENT:

RELEASED BY	DATE	TIME	RECEIVED BY	DATE	TIME
Jody R. Hone	5/25/93	:		/ /	:
	/ /	:		/ /	:
	/ /	:		/ /	:

OUTSIDE LAB INFORMATION

SAMPLES SHIPPED TO Atmospheric Assessment Assoc. CARNOT SHIPPER Jody Hone
21354 Nordhoff St. Ste 1B DATE 5/25/93
Chatsworth, CA 91311 CARRIER # Rite Way
 PHONE: 818-718-1070 AIR BILL # _____
 RECIPIENT Karl Pater
 COMPANY AmAA
 DATE 5/25/93

15991 Red Hill Ave., Suite 110, Tustin, California 92680-7388 (714) 259-9520 - FAX (714) 259-0372

CERTIFICATION OF SAMPLE RECEIPT

PROJECT # 10852 OUTSIDE LAB REQUIRED (Y/N) Y SAMPLE DATE 5/25/93
 PROJECT MANAGER EF
 SAMPLE LOCATION STACK METHOD(S) SCA QMD 25.2 TECHNICIAN Wm
 COMPLIANCE TEST (Y/N) Y DATE DUE 5-28-93

TEST #	COMPLETE DESCRIPTION
4-6-NH ₃	1 TEDLAR BAG 91463 - 1
4-7-NH ₃	1 TEDLAR BAG 2
4-8-HC	1 TEDLAR BAG 3
4-9-HC	1 TEDLAR BAG 4

CHAIN OF CUSTODY PRIOR TO SHIPMENT:

RELEASED BY	DATE	TIME	RECEIVED BY	DATE	TIME
Wg R. Mark	5/26/93	8:57	J. K. [Signature]	5/26/93	8:57

OUTSIDE LAB INFORMATION

SAMPLES SHIPPED TO MIKE PORTER
ATM AA, INC.
CHATS WORTH, CA
 PHONE: (818) 718-6070

CARNOT SHIPPER Wayne Madden
 DATE 5/26/93
 CARRIER # Right O' Way
 AIR BILL #

RECIPIENT [Signature]
 COMPANY ATM AA
 DATE 5/26/93

CERTIFICATION OF SAMPLE RECEIPT

PROJECT # 10852 OUTSIDE LAB REQUIRED (Y/N) Y SAMPLE DATE 5/25/93
 SAMPLE LOCATION STACK METHOD(S) SCA QMD 25.2 PROJECT MANAGER EF
 COMPLIANCE TEST (Y/N) Y TECHNICIAN Wm
 ANALYZE BY ANALYZE BY
 DATE DUE 5-28-93

TEST #	COMPLETE DESCRIPTION
4-6-NH3	1 TEDLAR BAG
4-7-NH3	1 TEDLAR BAG
4-8-HC	1 TEDLAR BAG
4-9-HC	1 TEDLAR BAG

CHAIN OF CUSTODY PRIOR TO SHIPMENT:

RELEASED BY	DATE	TIME	RECEIVED BY	DATE	TIME
Wg R. Mark	5/26/93	8:57	<i>[Signature]</i>	5/26/93	8:57
	/ /	:		/ /	:
	/ /	:		/ /	:

OUTSIDE LAB INFORMATION

SAMPLES SHIPPED TO MIKE PORTER CARNOT SHIPPER Wayne Madden
ATM AA, INC. DATE 5/26/93
CHATS WORTH, CA CARRIER # Right O'Way
 PHONE: (818) 718-6070 AIR BILL #
 RECIPIENT
 COMPANY
 DATE

15991 Red Hill Ave., Suite 110, Tustin, California 92680-7388 (714) 259-9520 • FAX (714) 259-0372

WORK AUTHORIZATION AND ANALYSIS INSTRUCTIONS FOR CONTRACT LABORATORIES

P.O. # 1175 AMOUNT \$ 340

CHAIN OF CUSTODY AND
SAMPLE IDENTIFICATION

☒ ATTACHED ☐ LISTED BELOW ☐ TO FOLLOW

PROJECT # 10 852

TYPE OF SAMPLE TEDLAR BAGS

OF SAMPLES IN TOTAL SHIPMENT 4

ANALYSIS INSTRUCTIONS:

ANALYZE THESE FOUR TEDLAR BAGS FOR
COMPLIANCE FOLLOWING SCAQMD DRAFT 25.2
FOR CH₄ + TCMHC

REFERENCE METHOD REQUIRED:

SCAQMD 25.2

COMMENTS:

DELIVER REPORT TO ED PHILADELPHIA

SIGNED W. R. M. M.

15991 Red Hill Ave., Suite 110, Tusin, California 92680-7388 (714) 259-9520 - FAX (714) 259-0372

CHAIN OF CUSTODY

PROJECT # 10852 OUTSIDE LAB REQUIRED (Y/N) Y SAMPLE DATE 2/1/11
 CLIENT/LOCATION _____ PROJECT MANAGER EF
 SAMPLE LOCATION STACK METHOD(S) SCAQMD 25.2 TECHNICIAN JM / MR
 COMPLIANCE TEST (Y/N) Y DATE DUE 4/1/11

DATE	TIME	TEST #	SAMPLE DESCRIPTION	CONTAINERS	SAMPLER	COMMENTS
2/1	:	1	✓	1	✓	
1/1	:	2	✓	1	✓	
1/1	:	40	✓	1	✓	
2/1	:	100	✓	1	✓	
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					
1	:					

TRANSFER OF SAMPLES FROM FIELD SAMPLE CUSTODIAN (FSC):

RELEASED BY (FSC)	DATE AND TIME	RECEIVED BY	DATE AND TIME
	1 1 :		1 1 :
	1 1 :		1 1 :
	1 1 :		1 1 :

ANALYSIS REQUIRED: ANALYZE FOUR (4) TENDR BARS FOR
COMPLIANCE RESULTS ACCORDING TO SCAQMD
25.2 (REPORT CH₄ + TMMHC IN PPM)