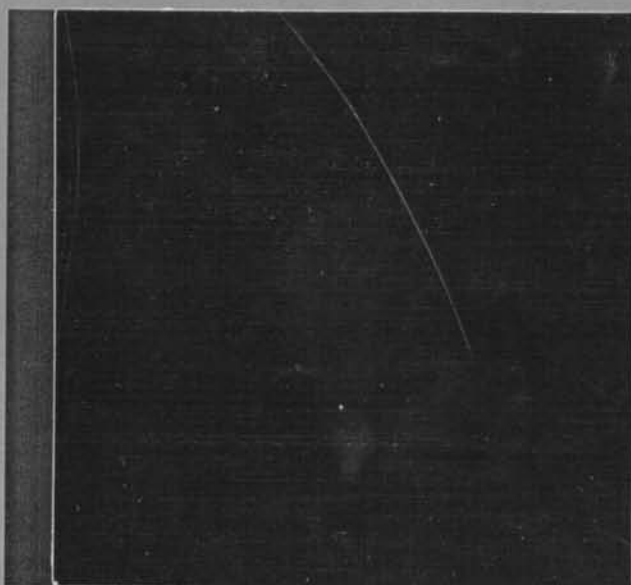




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**SOURCE TEST REPORT
MULTIPLE METALS EMISSIONS
TESTING**

**Sandia National Laboratories
P.O. Box 969, MS 9053
Livermore, CA 94551-0969**

**Prepared by:
Dames & Moore Emission Measurement Section
Submitted: April 14, 1997
Date of Test: March 10-14, 1997
Job No. 35746-001-131**

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- Appendix A: Field Data Sheets
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- Appendix C: Continuous Emission Data
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PREFACE

Firm Tested: Balboa Pacific Corporation
Address 11240 Bloomfield Ave
City: Santa Fe Springs, CA 90670
Contact: Dr. Shapoor Hamid (310) 929-1633
Source: BAL-PAC Pyrolysis Unit Exhaust

Date of Test: March 10, March 12, & March 14, 1997
Test Requested by: Sandia National Laboratories
Contact: David Hahn (510) 294-3337
Test Observed by: Sandia/Balboa Staff
Test Performed by: Dames & Moore
Team Members: Christopher J. Barth, Brian Satow, Jason Wirth
Test Objectives: Quantify and compare multiple metal emissions using reference EPA Method 29 and Sandia National Laboratory's multiple metal continuous emissions monitor.
Test Methods: EPA Method 29 "Determination of Multiple Metals Emissions from Stationary Sources"
CARB Method 100 "Instrumental Methods for Gaseous Emissions"

EXECUTIVE SUMMARY

Dames & Moore was retained by Sandia National Laboratories to conduct multiple metals emission testing on the BAL-PAC pyrolytic unit manufactured by Balboa Pacific Corporation. The emissions testing was conducted concurrently with Sandia's multiple metal Continuous Emissions Monitor (CEM) unit. The primary goal of the sampling project was to compare the metal emissions recorded by Sandia's CEM unit with the metals emissions determined by Dames & Moore using USEPA Reference Method 29 "Determination of Metals Emissions From Stationary Sources".

Testing was conducted during the week of March 10, 1997 and consisted of three test days utilizing one feed per day of the following: municipal waste, shredded tires, and a molybdenum-rich spent catalyst. Each day of testing entailed a 2-hr test run in which the CEM unit looked for six (6) specific target metals: Iron (Fe), Manganese (Mn), Molybdenum (Mo), Chromium (Cr), Nickel (Ni) and Cadmium (Cd). Table ES-1 summarizes the target metal results.

TABLE ES-1

Target Metal Sampling Summary

Multiple Metals Summary	Run-1		Run-2		Run-3	
Date Tested	3/10/97		3/12/97		3/14/97	
Feed	Municipal Waste		Shredded Tires		Spent Catalyst	
Metals Emissions, ug/m ³	RM	CEM	RM	CEM	RM	CEM
Iron, Fe	39.9	22.6	82.2	48.7	139.7	91.3
Manganese, Mn	3.3	3.1	4.9	9.3	4.5	7.0
Molybdenum, Mo	76.2	NA	20.2	NA	16.6	NA
Chromium, Cr	2.9	2.3	1.9	1.5	1.0	NA
Nickel, Ni	1.3	4.8	1.8	5.0	1.3	2.0
Cadmium, Cd	1.4	NA	1.6	NA	0.6	NA

RM = Metals results using Reference Method 29 "Multiple Metals Emissions from Stationary Sources"

CEM = Average metals results using Sandia's Multiple Metal Continuous Emissions Monitor

1.0 INTRODUCTION

Sandia National Laboratories is developing the technology of a multiple metals continuous emissions monitor (CEM). Dames & Moore was retained by Sandia National Laboratories (Sandia) to conduct multiple metal emissions testing, using a reference method, simultaneously with the multiple metals CEM. The purpose of the testing was to compare Sandia's CEM results with a reference method to fine tune this new CEM technology.

The emissions testing was performed on the exhaust stack of a pilot-model pyrolytic conversion unit used to treat hazardous and non-hazardous waste. The pyrolysis unit was manufactured and operated by Balboa Pacific Inc., Santa Fe Springs, CA.

In addition to the multiple metals testing, exhaust gas sampling was conducted and the results are included for Sandia's information. Constituents analyzed include oxides of nitrogen (NO_x), carbon monoxide (CO), total hydrocarbons (THC), oxygen (O_2), and carbon dioxide (CO_2). The following test methods were used:

- **EPA Method 29** "Determination of Multiple Metals Emissions from Stationary Sources"
- **CARB Method 100** "Instrumental Methods for Gaseous Emissions, NO_x , CO, THC, O_2 and CO_2 "

The testing was conducted at Balboa Pacific, 11240 Bloomfield Avenue, Santa Fe Springs, California 90670.

This document presents a detailed description of the system, a description of the operating parameters within which the system was operated during the course of the test, a description of the monitoring, sampling techniques and analytical procedures which were used, and the results of the testing.

2.0 EQUIPMENT AND PROCESS INFORMATION

2.1 Balboa Pacific Corporation Pyrolytic Conversion System

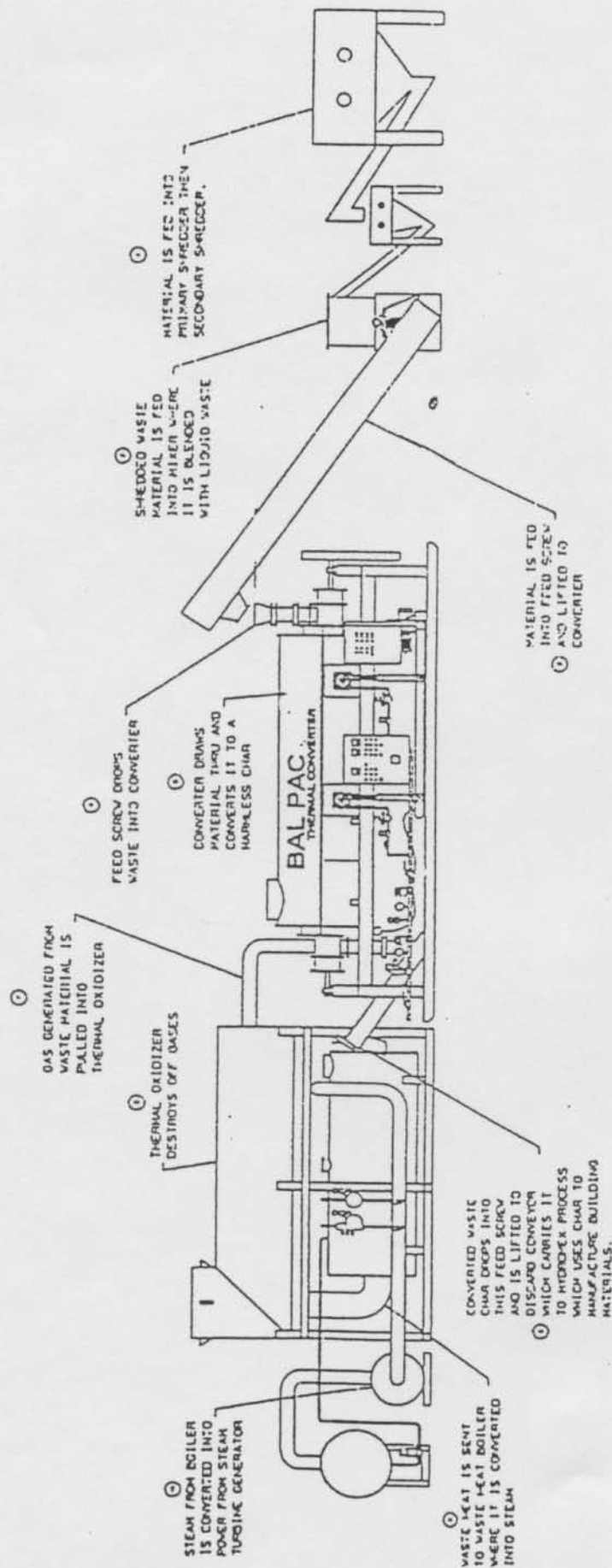
The BAL-PAC Pyrolytic Conversion System is based on patented technology. The basic principle of this technology is the destruction and stabilization of the waste stream using very high temperatures (1900-2000 °F) in an oxygen free environment. By this process the waste stream entering the unit is converted into combustible gases and a stabilized solid waste stream. The gases produced are further subjected to high temperatures (2250 °F) in a thermal oxidizer where the combustible gases are destroyed. There are four main subsystems to the BAL-PAC system described below. A diagram of the unit is shown in Figure 2-1.

- The Feed or Input System: The waste material is introduced into the system through a series of valves and gates that are synchronized to prevent unwanted oxygen or air from entering into the processing chamber.
- Pyrolytic Conversion Chamber (retort): A thermally insulated outer housing surrounding a retort or pyrolytic chamber containing a rotary screw that conveys the waste through the retort as pyrolysis occurs. The space between the outer housing and the internal retort chamber contains a heat chamber, through which natural gas at a maximum rate of 500 cubic feet per minute is routed for combustion, providing the heat source for pyrolysis.
- Thermal Oxidizer: The gases liberated by pyrolysis are drawn off by a "closed coupled thermal oxidizer" where they are ignited, converting them primarily to carbon dioxide and water. The temperature in the thermal oxidizer can reach 2500 degrees F. The retention time for the gases in the chamber is at least 2 seconds. The thermal oxidizer is also fired on natural gas.
- Output System: This system handles the solid by-product produced after the pyrolytic retort. It is characterized by air locks controlled by synchronized valves that expel inert residual pyrolysis matter for post-pyrolytic processing.

FIGURE 2-1

Diagram of BAL-PAC System

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DRAWING AS SHOWN IS A REPRESENTATION OF THE PROCESS FLOW AND IS NOT TO SCALE. FOR APPROXIMATE DIMENSIONS SEE NOTES BELOW.

WASTE HEAT BOILER, 87' x 25' L
THERMAL OXIDIZER, 87' x 6' x 20' L
BAL PAC BOILER, 810' x 6' x 40' L
STEAM TURBINE GENERATOR, 12' x 32' L x 10' H (HEIGHT x LENGTH x AREA, 100,000 sq ft)

2.2 Process Information

Mr. Jerry Hold of Balboa Pacific was responsible for the operation of the BAL-PAC during all testing and for the recording of specific process parameters during the testing. Table 4-1 "BAL-PAC Operating Parameters" tabulates the system parameters during the course of the testing.

TABLE 2-1

BAL-PAC Operating Conditions
During the Destruction of Several Waste Feeds

Feed Material	Feed Rate lb/hr	Retort Temperature		Oxidizer Temp. °F	Boiler Exh. Temp. °F	Stack Temp. °F
		Inlet °F	Outlet °F			
Waste/Sludge	21	1600	1589	1700	560	138
Shredded Tires	30	1500	1450	1700	520	138
Spent Catalyst	10	1700	1650	1700	560	136

Since the purpose of testing was to determine the accuracy of Sandia's multiple metal CEM, Balboa Pacific was operating the BAL-PAC unit at temperatures which were higher than what would be considered normal operating conditions for the purposes of generating metals emissions that could be detected by both Sandia's CEM and Dames & Moore's reference method testing.

3.0 SAMPLING METHODOLOGIES

3.1 Sampling Locations

The exhaust stack from the wet scrubber was 16 inches by 21 inches (18" equivalent diameter). Figure 3-1 shows the port locations on the exhaust stack. Three test ports were located on the 21 inch face, the ports being 4 inches in diameter and 1.5 inches in length. They were located 77" upstream (4.3 duct diameters) and 92" downstream (5.1 duct diameters) of the nearest flow disturbance, meeting EPA Method 1 requirements. Eight sampling points for each sample port were used for the Method 29 multiple metals testing as shown in Figure 3-2.

Sandia located their multiple metal CEM probe approximately 18 inches downstream of the three 4-inch diameter sampling ports.

FIGURE 3-1

Exhaust Stack Diagram

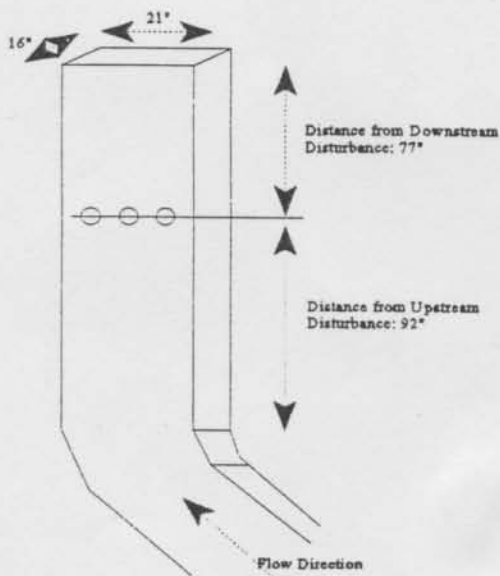
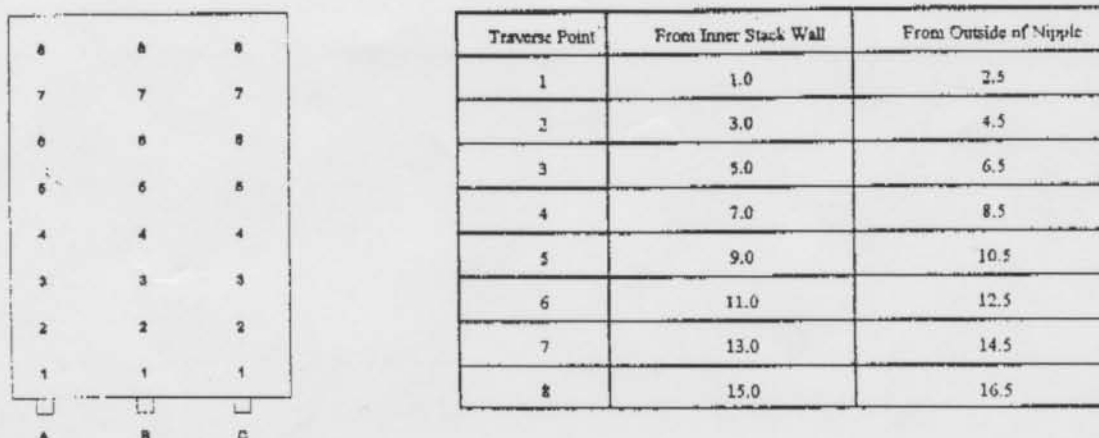


FIGURE 3-2

EPA Method 29 Traverse Point Locations



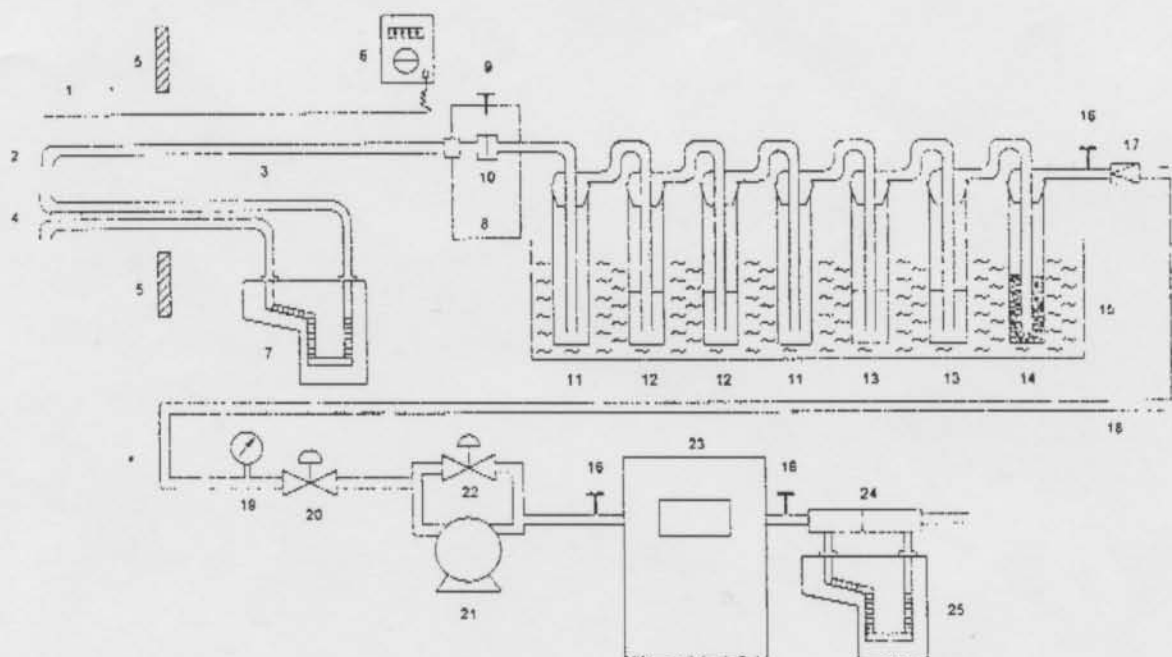
3.2 EPA Method 29: Determination of Metals Emissions from Stationary Sources

EPA Method 29 was used to collect samples for multiple metal determination. Multi-point isokinetic sampling was conducted at five minutes per traverse point for a total of 120 minutes. The sampling train consisted of a glass-lined, heat traced probe with a glass, button hook nozzle with an attached thermocouple and pitot tube assembly. Figure 3-3 shows the Method 29 sampling apparatus.

Five 500-ml impingers were connected in series with leak-free ground-glass joints. The first impinger was used as a moisture drop-out impinger due to the high moisture content of the stack gas. The second and third impingers each contained 100 ml of a 5% HNO_3 /10% H_2O_2 absorbing solution. The fourth impinger was left empty and the fifth impinger contained a known amount of silica gel. The mercury absorbing impingers specified in Method 29 was left out as mercury was not requested. Also, the filter and glass filter holder were replaced with a filter bypass. All recovered fractions, probe rinse, moisture dropout, impinger contents, were combined for subsequent analysis. The collected sample was sent under chain-of-custody to West Coast Analytical, Santa Fe Springs, CA for predetermined target metal determination.

FIGURE 3-3

Multi Metals Sampling Apparatus



- | | | | |
|-----|--|-----|----------------|
| 1. | Temperature Sensor | 14. | Silica Gel |
| 2. | Glass Probe Tip | 15. | Ice Bath |
| 3. | Glass Probe Liner | 16. | Thermocouple |
| 4. | S-Type Pitot Tube | 17. | Check Valve |
| 5. | Stack Wall | 18. | Vacuum Line |
| 6. | Temperature Sensor Meter | 19. | Vacuum Gauge |
| 7. | Pitot Tube Inclined Manometer | 20. | Main Valve |
| 8. | Heated Area | 21. | Air Tight Pump |
| 9. | Thermometer | 22. | By-Pass Valve |
| 10. | Glass Filter Holder | 23. | Dry Gas Meter |
| 11. | Empty Impinger (optional) | 24. | Orifice |
| 12. | 5% HNO_3 /10% H_2O_2 | 25. | Manometer |
| 13. | 4% KMnO_4 /10% H_2SO_4 (eliminated) | | |

3.5 CARB Method 100: Instrumental Methods for Gaseous Emissions (NO_x , CO, THC, O_2 , CO_2)

Oxides of nitrogen (NO_x), carbon monoxide (CO), total hydrocarbons (THC), oxygen (O_2), and carbon dioxide (CO_2) gas concentrations in the flue gas were measured using an extractive Continuous Emissions Monitoring System (CEMS) in Dames & Moore's Mobile Laboratory. One 2-hour test was conducted simultaneous with the Method 29 testing.

The Dames & Moore Continuous Monitoring System is comprised of four major subsystems. They are: (1) the sample acquisition and conditioning system, (2) the calibration gas system, (3) the analyzers, and (4) the data recording system. A schematic of the Dames & Moore CEM system is presented in Figure 3-4.

The sample acquisition and conditioning system extracts a representative sample from the stack, removes moisture and particulate material from the sample, and transports the sample to the analyzers. The sample acquisition system consists of a 7 micron stainless steel filter and a 3/8 inch 316 stainless steel probe. The probe is insulated and heated between 250-275 °F to avoid condensation. From the probe, the sample gas is transported through a heat-traced Teflon sample line maintained at 240-260 °F from the probe to the Universal Analyzer Thermoelectric Gas Sample Cooler Model 3080 via a Teflon-lined diaphragm pump. The outlet temperature of the thermoelectric sample cooler is fully automatic and maintains the dew-point of sample gas below 37 °F.

Sample gas flow is controlled by a series of flow-meter, valves, and regulators upstream of the instrument manifold. Excess sample is vented through a back-pressure regulator, maintaining a constant pressure of 6-7 psig and flow of 2 scfh to each analyzer rotameter. Instrument response is permanently recorded using an online data acquisition system. All fittings and sample line which may contact the sample gas are constructed of stainless steel and Teflon.

Following system performance checks, preliminary calibration error checks were performed on each analyzer by introducing zero and high span gases and recording the response. A system bias check was then performed to determine the effect of the sample lines, pump, and sample cooler on the measurements. This was done by comparing the calibrations through a 3-way valve at the probe outlet, to calibrations done using calibration gases (zero and upscale concentrations) introduced directly to the instruments. EPA Protocol 1 calibration gases in current certification were used for

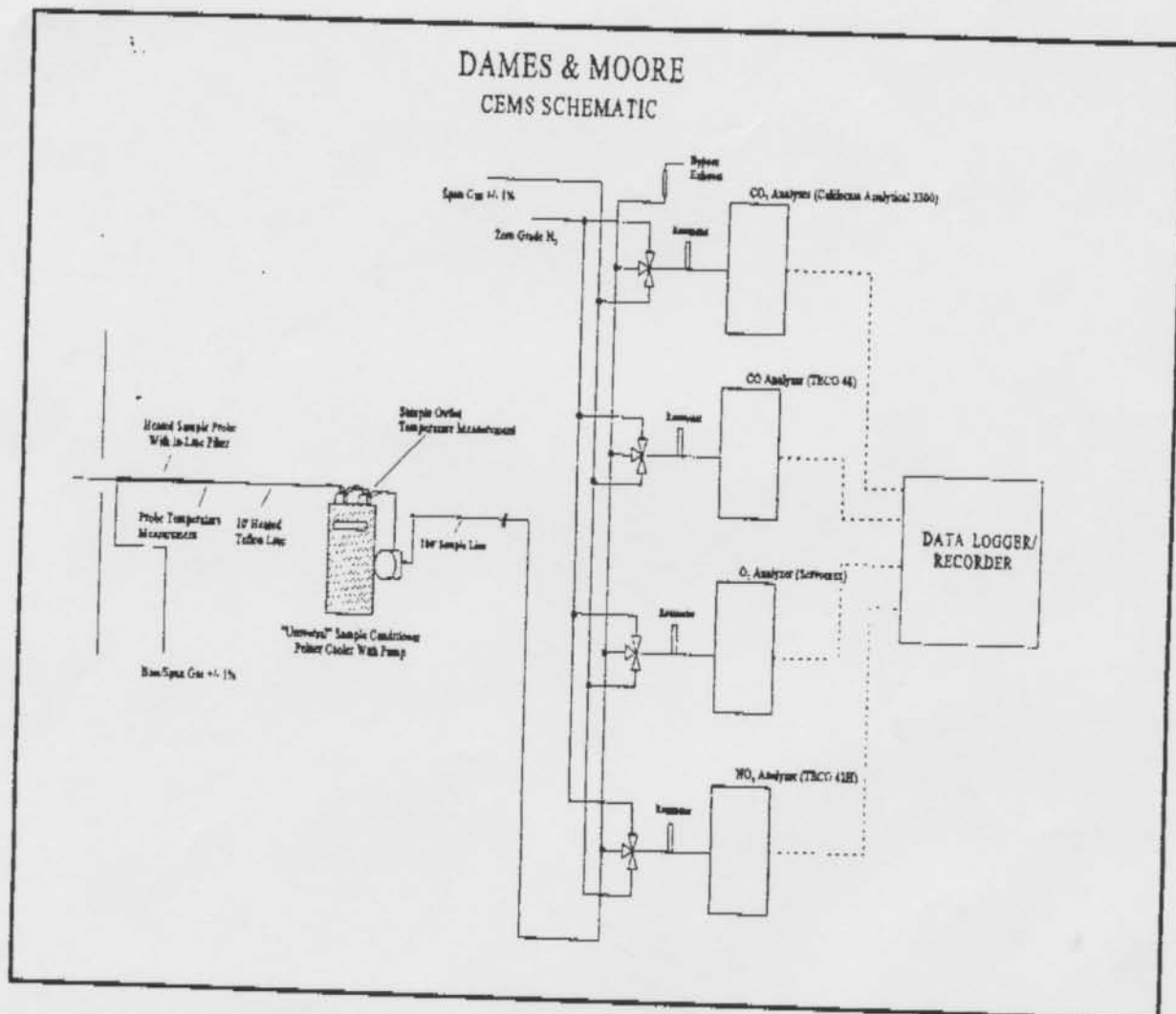
all calibration purposes. Calibration gas certifications are included in Appendix D. Post test calibration and bias checks were performed upon completion of the test.

Specifications for the continuous monitors used in Dames & Moore's Mobile Lab are presented in Table 3-1.

TABLE 3-1
Continuous Emissions Monitoring Instrumentation

Gas	Nitrogen Oxides (NO _x)	Carbon Monoxide (CO)	Oxygen O ₂	Carbon Dioxide CO ₂	Hydrocarbons THC
Instrument Manufacturer and Model Number	TECO 42H	TECO Model 48	Servomex Model 1400	California Analytical Model 3300	TECO 51
Detection Principle	Chemiluminescence Photo-Multiplier Tube	Gas filter Correlation Analyzer	Magneto-Dynamic Paramagnetic Analyzer	Nondisp. Infrared Absorption (NDIR)	Flame Ionization (FID)
Drift: Zero Span	50 ppb 24 hr <1 % full scale/24hr>	± 0.2 ppm ± 1 % full scale	<1 % full scale/24 hr <1 % full scale/24 hr	± 1 % full scale/ 24 hr ± 1 % full scale/24 hr	± 1 % full scale/ 24 hr
Output	0-10 V	0-10V	0-1V	0-1V	0-10 V
Ranges	0-10, 20, 50, 100, 200, 500, 1000, 2000, 5000 ppm	0-1, 2, 5, 10, 20, 50, 100, 200, 500, 1000 ppm	0-25 percent 0-10 percent	0-10 percent 0-20 percent	0 - 10,000 selectable
Response Time	30 sec	30 sec	30 sec	30 sec	5 sec

FIGURE 3-4
Dames & Moore CEMS Schematic



4.0 TEST RESULTS

All reference method (EPA Method 29) testing was conducted simultaneously with Sandia's multiple metal CEM testing. One 2-hour run was performed on each day of testing. At the end of each test day, Sandia supplied the Dames & Moore test crew with a printout of their CEM data collected during the 2-hour test run. This data as well as all Dames & Moore field data and calculations are included in the Appendices. On the third day of testing, March 4, the Sandia CEM Laser malfunctioned approximately 30-minutes into the run. Dames & Moore stopped sampling upon notification that the CEM was not operating. Approximately two hours later, Sandia informed Dames & Moore that the CEM would be inoperable for the remainder of the day. Dames & Moore proceeded with the Method 29 sampling in order to obtain a sufficient sample volume to achieve the project data quality objectives. The test results are summarized in Tables 4-1, 4-2 and 4-3.

TABLE 4-1

EPA Method 29 Metal Sampling and Stack Gas Data

Parameter	Units	Run-1	Run-2	Run-3
Test Date		03/10/97	03/12/97	03/04/97
Feed		Municipal Waste	Shredded Tires	Molybdenum Catalyst
Sampling Time	min	120.5	119	120
Sample Volume	ft ³	70.356	69.300	68.790
Static Pressure	in H ₂ O	-0.02	-.02	-.02
Isokinetics	%	98.3	100.5	100.5
Stack Temperature	°F	138	138	136
Moisture	%	18.4	17.3	17.2
Stack Gas Velocity	ft/sec	6.4	6.2	6.0
Stack Gas Flow Rate	ACFM	897	863	846
	DSCFM	648	633	622

TABLE 4-2

EPA Method 29 - Multiple Metal Emission Results

Test Date	Run-1 3/10/97		Run-2 3/12/97		Run-3 3/14/97	
Fuel	Municipal Waste		Shredded Tires		Spent Catalyst	
Emissions	mg/hr	ug/m ³	mg/hr	ug/m ³	mg/hr	ug/m ³
Aluminum (Al)	23.7	15.52	33.3	22.70	9.7	6.72
Antimony (Sb)	0.8	0.51	0.5	0.34	0.1	0.08
Arsenic (As)	<	1.1	<	1.1	<	1.1
Barium (Ba)	1.4	0.94	2.4	1.61	0.7	0.49
Beryllium (Be)	<	0.0	<	0.0	<	0.0
Cadmium (Cd)	2.1	1.38	2.4	1.64	0.8	0.57
Chromium (Cr)	4.4	2.90	2.8	1.94	1.5	1.02
Copper (Cu)	8.1	5.29	9.6	6.57	4.4	3.09
Lead (Pb)	16.0	10.52	5.4	3.66	2.2	1.51
Iron (Fe)	60.8	39.90	120.5	82.15	200.9	139.70
Manganese (Mn)	5.0	3.26	7.1	4.85	6.5	4.53
Molybdenum (Mo)	116.1	76.15	29.5	20.14	23.9	16.59
Nickel (Ni)	2.0	1.34	2.6	1.76	1.9	1.32
Phosphorus (P)	23.8	15.60	17.0	11.58	9.8	6.80
Selenium (Se)	<	0.5	<	0.5	<	0.5
Silver (Ag)	0.4	0.25	0.3	0.19	0.2	0.11
Thallium (Tl)	<	0.0	<	0.0	<	0.0
Zinc (Zn)	27.6	18.14	273.9	186.71	40.2	27.94

Note: Concentrations (ug/m³) based on stack conditions (i.e. stack temperature and moisture content)

< denotes non-detects, emissions based on 1/2 the detection limit

TABLE 4-3

Additional Gaseous Emissions Data

CONSTITUENT	UNITS	CONCENTRATION	
		03/10/97	03/12/97
Oxygen, O ₂	%	11.3	13.4
Carbon Dioxide, CO ₂	%	5.8	4.6
Carbon Monoxide, CO	ppmvd	< 1	< 1
Oxides of Nitrogen, NO _x as NO ₂	ppmvd	75.5	59.1
Total Hydrocarbons, THC as C ₃	ppmvd	< 1	< 1

Note: No gaseous emissions testing was performed on March 14.

5.0 QUALITY ASSURANCE AND QUALITY CONTROL PROCEDURES

5.1 Sampling Protocol

Dames & Moore is organized to facilitate sample management, analytical performance management, and data management. Personnel are assigned specific tasks to ensure implementation of the QA/QC program. The Senior Scientist in charge of air emission measurements reports directly to the Associate of the Air Quality Group and is the QA officer responsible for program effectiveness and compliance.

The sample custodian is responsible for the care, custody and control of all samples for analysis and for the accumulation and consolidation of the analytical results, including internal quality controls and approvals.

The analysts perform the analyses and initial data review. Each analyst must check and initial their work, making certain that it is complete, determining that any instrumentation has been properly calibrated, and ensuring that the analysis has been performed within the QA/QC limits. Completed work is placed in the job jacket and submitted for review.

The Senior Scientist evaluates the data submitted by the analyst by first assessing the validity of the analytical method chosen for the analysis. They then verifies that the data and documentation are complete, that each analysis has been performed within QA criteria specific to each method, check calculations, assembly and sign the data package and prepares the report.

5.1.2 Sample Management

5.1.2.1 Chain of Custody Definition

The highest quality analyses have no validity unless all aspects of sample collection, transport, receipt, analysis, and data compilation can be validated. The following procedures are employed to maintain "chain of custody" for samples received by Dames & Moore. Because of the nature of the data being collected, the possession of samples must be traceable from the time the samples are collected until final disposition. A sample is under custody if:

- It is in your actual possession, or

- It is in your view, after being in your possession, or
- It is locked in a secure area.

5.1.2.2 Sample Identification

Each sample shall have affixed a label containing the sample number and sample description to identify the contents of the container. Additionally, the sample number shall be marked on the outside of any special packaging container.

5.1.2.3 Sample Management: Sample Custody During Transport

To assure custody of samples during transport and shipping, each sample within a packing container is recorded on a chain of custody record. Each sample number is recorded along with the number of containers shipped. Chain of Custody record forms must always be signed and dated when samples change hands or are shipped to another location. Copies of the COC sheets are provided to sample delivery personnel if requested. The original custody sheet is then placed inside the package (protected from damage) and the package sealed. Sample containers, shipping boxes, coolers or other packages may be sealed by using custody seals. The seals must be placed so the container cannot be opened without breaking the seal.

5.2 Equipment Calibration and Maintenance

The Senior Field Technician and Field Technician are in charge of routine maintenance and calibrations of all source testing equipment. Most recent calibration information can be found in the Appendix.

5.2.1 Equipment Maintenance

All major pieces of equipment have maintenance logs where all maintenance activities are recorded and documented. Table 5-1 shows routine maintenance that is performed on source testing equipment.

TEST EQUIPMENT MAINTENANCE SCHEDULE

<u>Equipment</u>	<u>Acceptance Limits</u>	<u>Frequency of Service</u>	<u>Methods of Service</u>
Pump	1. Absence of leaks 2. Ability to draw manuf. Reg. Vacuum and flow	Every 500 hrs or operation or 6 months, whichever is less	1. Visual inspection
Flow Meter	1. Free mechanical movement 2. Absence of malfunction	Every 500 hrs. of operation or 5 mos whichever is less	1. Visual insp. 2. Clean 3. Calibrate
Sampling Instrument	1. Absence of malfunction 2. Proper response to aero, span gas	As recommended by manufacturer	As recommended by manufacturer
Mobile Van Sampling System	1. Absence of leaks	Depends on nature of use	1. Change filters 2. Change gas dryer 3. Leak check 4. Check for system contamination
Sampling Line	Sampling degradation after each test less than one percent of test series		Flush with solvent. Blow air through line until dry.

Table 5-1 Maintenance Schedule Information

5.2.2 Equipment Calibration

Current calibration information on equipment used during testing will be available for viewing during the source test. Examples of calibration current calibration information can be found in the appendix.

The S-Type pitot tubes are measured initially upon purchase and then semiannually. Visual measurements are taken prior to each use to insure accidental damage has not occurred. This check

is documented on the testing forms. Measurement is performed using a micrometer and compass.

Each temperature sensor is marked and identified. This is done by marking each thermocouple end connector with a number. This sensor is calibrated as a unit with the control box potentiometer and associated lead wire as an identified unit. Calibration are performed initially and annually at multiple three-points over the range of expected temperatures for that particular thermocouple. A non-multiple three-point check is performed bimonthly thereafter. As an alternative to the three baths, an Oyster Calibrator Thermometer may be used as a temperature reference source.

The field barometer is adjusted initially and semiannually to within 0.1 in Hg of the atmospheric pressure as reported by John Wayne Airport. There is no correction between John Wayne Airport and Dames & Moore.

The field dry gas test meter is calibrated before its initial use and semiannually thereafter. Its calibration is checked bimonthly. It is calibrated against a reference dry gas test meter.

The dry gas meter orifice is calibrated before its initial use and then annually. This calibration is performed during the calibration of the dry gas test meter. The unit is checked in the field after every series of tests using a field gas meter check procedure.

Probe nozzles are measured prior to use in the field using a micrometer.

Analytical balances are internally calibrated prior to use following the manufacturer's instructions. The balances are further checked using Class S-1 analytical weights prior to daily usage. Field top loading balances are also internally calibrated prior to use and checked with a field analytical weight prior to usage.

5.3 Instrument Calibration

The analytical range is selected so that the sample gas concentration for each run is between 10 and 95 percent of the range, for 95 percent of the test period. The run is considered invalid if the measured gas concentration exceeds the range during the test period. Data obtained below 10 percent of the range can be used only for qualitative purposes.

All continuous emission monitoring analyzers will be calibrated to meet the following specifications:

- Analyzer Calibration Error less than ± 2 percent of the range of the zero, mid-range, and high-range calibration gases.
- Sampling System Bias less than ± 5 percent of the range for the zero, and mid-or high range calibration gases.
- Zero Drift less than ± 3 percent of the range over the period of each run.
- Calibration Drift less than ± 3 percent of the range over the period of each run.
- Linearity less than ± 2.0 percent of the range for the pretest and post test values.

Bias calibration gases are introduced to the probe tip by flooding the probe with calibration gas during probe calibrations. Calibration and zero gases are introduced into the manifold through a valving system that shunts the sample flow to vent and allows calibration or zero gases into the sample manifold.

Calibration gases shall be certified to an analytical accuracy of ± 1 percent and be traceable to applicable NIST Standard Reference Materials (SRM's). Traceability shall include identification of applicable SRM and its cylinder number. Superblends may be used for simultaneous calibration of multiple analyzers. The average deviation of each component gas shall not exceed ± 1 percent of the tag value.

Operation of the continuous emission monitoring system is as follows:

- Cleaning of Sample Train as needed. Thoroughly flush the probe, heat-trace line, and sample conditioner with distilled water, followed by acetone. Dry with filtered dry air.

- Allow analyzers to warm up according to manufacturers instructions or until stable readings are obtained.
- Sampling System leak check.
- Calibrate instruments by Introducing zero and high range calibration gases directly to the instruments and make all necessary adjustments to calibrate the analyzer and the data recorder. Adjust system components to achieve individual analyzer sampling rates recommended by the instrument manufacturer.
- Conduct the analyzer calibration error check at the beginning and end of each test run by introducing calibration gases to the analyzers through the manifold system. The calibration error check should be considered invalid if the gas concentration displayed by the analyzer exceeds ± 2 percent of the range for any of the calibration gases. If an invalid calibration is exhibited, take corrective action and repeat the analyzer calibration error check until acceptable performance is achieved.
- Instrument Response Time is establish during semi-annual certification.
- Sampling System Bias Check is performed by flooding calibration gases into the sample probe. A zero gas and either the mid-range or high-range gas, whichever most closely approximates the effluent concentrations, is used. The sampling system bias check shall be considered invalid if the difference between the gas concentrations exhibited by the measurement system when a known concentration gas is introduced at the sampling probe tip and when the same gas is introduced directly to the analyzer, exceeds ± 5 percent of the analyzer range. If an invalid calibration is exhibited, take corrective action and repeat the sampling system bias check until acceptable performance is achieved.
- NO₂ to NO conversion efficiency conversion test is performed by introducing an NO₂ standard and measuring the NO_x concentration.

Switch the analyzer mode to NO and record the NO reading. The converter efficiency is the percent difference between the two readings.

- Zero and Calibration Drift Tests are performed immediately preceding and following each run. If either the zero or upscale calibration value exceeds the sampling system bias specification, then the run is considered invalid. Repeat both the analyzer calibration error check procedure and the sampling system bias check procedure before repeating the run.
- If both the zero and upscale calibration values are within the sampling system bias specification, then use the average of the initial and final bias check values to calculate the gas concentration for the run. If the zero or upscale calibration drift value exceeds the drift limits, based on the difference between the sampling system bias check responses immediately before and after the run, repeat both the analyzer calibration error check procedure and the sampling system bias check procedure before conducting additional runs.

5.4 Data Validation

The data presented on final reports are reviewed three times. First, the analyst reviews and certify that raw data complies to technical controls, documentation requirements, and standard group procedures. Second, the Senior Scientist reviews and certifies that data packages comply to specifications for sample holding conditions, chain of custody, data documentation, and the final report is free of transcription errors. Third, a QA review is performed by another senior personnel. This review thoroughly examines the entire completed data report. The report is signed off and sent out. All raw laboratory data and final reports are stored for 5 years.

APPENDIX A
FIELD DATA SHEETS

DAMES & MOORE
PRELIMINARY TRAVERSE

Company BALBOA PACIFIC
Source Exhibit - BALBOA
Date 3.10.97
Test Number Pre 1.4

Sampling Location BAL-12E FK 1451
 Static Pressure -0.025
 Barometric Pressure 29.89
 Engineer's Initials [Signature]

[illegible][illegible]

Equipment

Pressure Gauge
T/C Potentiometer
Pitot Tube
Pitot C.F.

1/4 MARCH
ATKINS
5-PT-CCS-3
0.84

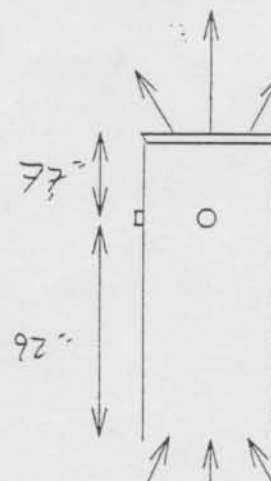
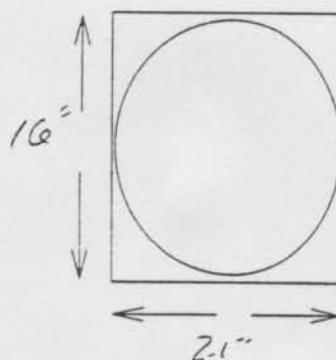
Leak Checks

Pre	
Post	

Nipple Dimensions

Length: 1.5"

Diameter: 3"



5.3.

$O_2 \sim 15\%$
 $CO_2 \sim 35\%$

DAMES & MOORE TRAVERSE SOURCE TEST DATA

Company: BALCO PACIFIC
Sampling Location: BAL PAC Exhaust
Test No: MULTIMETALS - RAW-1

Date: 3-10-97
Time Start: 1115
Time End: 1325

System Pre-Test Leak Check:
0.004 cfm @ 10 "Hg vac
(Pitot tube Leak Check: 0.0)

System Post-Test Leak Check
0.002 cfm @ 14 "Hg vac
(Pitot tube Leak Check: 0.0)

Time On	Source Point #	Gas Meter Reading (scf)	Stack				Probe Temp °F	Hot Box Temp °F	Dryer Temp °F	Meter Temp °F		Vacuum "Hg
			Velocity Head (H ₂ O)	Temp °F	Calculated dH	Set dH				in	out	
0		111.861	—	—	—	—	—	—	—	—	—	—
5	1	—	.015	141	1.64	1.6	250	250	61	92	92	1
10	2	118.59	.013	140	1.42	1.4	250	251	60	94	91	1
15	3	121.85	.013	141	1.4	1.4	250	250	47	97	92	1
20	4	—	.015	138	1.65	1.7	250	250	45	88	93	1
25	5	—	.018	139	1.98	2.0	250	251	46	99	94	1
30	6	132.85	.017	140	1.87	1.9	249	250	49	101	95	1
35	7	136.4	.015	137	1.66	1.7	250	250	49	101	96	1
40	8	140.015	.016	139	1.77	1.8	250	250	49	101	97	1
		(28.154)	(.01525)	(139)	—	(1.69)	—	—	(51)	—	(96)	(1)
45	1	143.35	.013	137	1.44	1.4	250	255	58	101	98	1
50	2	146.45	.012	138	1.33	1.3	249	251	50	101	98	1
55	3	149.86	.014	140	1.55	1.6	250	250	52	102	99	1
60	4	153.22	.013	138	1.44	1.4	251	251	50	102	99	1
65	5	156.39	.011	139	1.22	1.2	249	250	55	103	99	1
70	6	159.14	.008	138	0.88	0.88	250	248	56	102	99	1
75	7	161.8	.008	132	.88	0.9	251	250	57	102	100	1
80	8	164.890	.009	138	1.0	1.0	250	249	58	102	100	1
		(52.63)	(.0110)	(138)	—	(1.21)	—	—	(55)	—	(100)	(1)
85	1	—	.005	138	.55	.55	249	251	61	101	99	1
90	2	169.05	.007	136	.78	.78	250	253	67	101	99	1
95	3	171.61	.008	137	.88	0.89	250	247	67	101	99	1
100	4	—	.007	139	.78	.78	250	250	67	101	99	1
105	5	176.75	.007	132	.78	.78	249	251	57	101	99	1
110	6	179.47	.008	138	.88	.88	249	249	53	102	99	1
115	7	182.25	.008	139	.88	.88	250	250	54	102	99	1
120.5	8	185.339	.009	139	1.0	1.0	251	249	54	102	99	1
AVERAGES		73.478	.01121	138	—	1.24	—	—	55	—	100	1

PRE-TEST DATA

Nozzle Diameter: 0.626
Barometric Pressure: 29.99 "Hg
Static Pressure in Stack: -0.25 "H₂O
Pitot Factor: .84
Est. Moisture: 22% 70%, sat'd @
Est. MW: 88.26 140°F

CHAIN OF CUSTODY INFO

Impingers Loaded By: JA
Impingers Recovered By: JA
Filter Loaded By: JA
Filter No.: —
Filter Recovered By: JA
Probe Wash By: JA, JW

EQUIPMENT/CALIBRATION DATA

Pitot Tube No.: PF-003-1 (Cal: —)
Potentiometer No.: M5B-2 (Cal: 34.97)
Thermocouple No.: U4B-35-4C (Cal: 34.97)
Gas Meter No.: 2962389 (Cal: 34.97)
Meter Corr. Factor: 1.0083
Delta H @: 1.84263

Impinger #1
Impinger #2
Impinger #3
Impinger #4
Impinger #5
Impinger #6

Soln.	Final	Initial	Net
Dry	770.7	555.6	215.1
Dry	676.2	553.0	123.2
100% HNO ₃ /H ₂ O	699.0	668.3	30.7
"	782.5	778.4	4.1
dry	574.0	573.4	0.6
Silica	856.9	845.6	11.3g

323.7g

POST TEST INFO

Filter Appearance: NA
Impinger Appearance: Clear/whiskers
Silica Gel Spent (%): 30%

Total Wt Gain:

335.0g

Recorded by: Gregory Smith Date: 3-10-97
Checked by: JA Date: 4/8/97

Company: BALBIA PACIFIC
Sampling Location: BAL-BAL EXHIBIT
Test No. SC2 PM-1

System Pre-Test Leak Check:
0.00 ^{1/2 hr} 0.00 "Hg vac
 (Pitot tube Leak Check: _____)

[illegible]

Pilot Tube No.: 11A (Cal: —)
 Potentiometer No.: ATKIN'S (Cal: —)
 Thermocouple No.: 11A (Cal: —)
 Gas Meter No: 1231635 (Cal: 3652)
 Meter Corr. Factor: 4258353
 Delta H @: 11A

Recorded by: [Signature] Date: 3-10-97
Checked by: [Signature] Date: 4-8-97

File	Time	Species	Line	Freq.	($\mu\text{g}/\text{acm}$)
Macintosh HD:L:	3/10/97 9:43	He	388.86	100.00	94.8
Macintosh HD:L:	3/10/97 9:44	He	388.86	100.00	93.6
Macintosh HD:L:	3/10/97 10:29	Fe	259.90	2.20	16.3
Macintosh HD:L:	3/10/97 11:19	Fe	259.90	1.17	7.5
Macintosh HD:L:	3/10/97 11:21	Fe	238.20	2.00	24.5
Macintosh HD:L:	3/10/97 11:25	Fe	259.90	2.00	7.0
Macintosh HD:L:	3/10/97 11:32	Mn	257.60	0.10	0.2
Macintosh HD:L:	3/10/97 11:38	Mn	257.60	1.50	3.9
Macintosh HD:L:	3/10/97 11:42	Mn	257.60	0.90	1.9
Macintosh HD:L:	3/10/97 11:47	Fe	238.20	2.80	21.7
Macintosh HD:L:	3/10/97 11:52	Fe	238.20	2.00	30.1
Macintosh HD:L:	3/10/97 11:58	Fe	238.20	3.70	33.8
Macintosh HD:L:	3/10/97 12:03	Fe	238.20	2.10	20.3
Macintosh HD:L:	3/10/97 12:09	Ni	243.80	1.42	9.5
Macintosh HD:L:	3/10/97 12:14	Ni	243.80	1.50	-5.4
Macintosh HD:L:	3/10/97 12:19	Mn	257.60	1.10	2.9
Macintosh HD:L:	3/10/97 12:23	Mn	257.60	2.00	4.1
Macintosh HD:L:	3/10/97 12:27	Fe	238.20	1.86	7.6
Macintosh HD:L:	3/10/97 12:32	Cr	283.56	1.88	1.6
Macintosh HD:L:	3/10/97 12:37	Cr	283.56	1.82	2.1
Macintosh HD:L:	3/10/97 12:41	Fe	238.20	2.62	15.5
Macintosh HD:L:	3/10/97 12:45	Fe	238.20	3.88	29.7
Macintosh HD:L:	3/10/97 12:49	Mn	257.60	1.80	3.7
Macintosh HD:L:	3/10/97 12:53	Mn	257.60	2.80	6.6
Macintosh HD:L:	3/10/97 12:57	Fe	238.20	3.25	39.1
Macintosh HD:L:	3/10/97 13:03	Cr	283.56	1.88	1.9
Macintosh HD:L:	3/10/97 13:07	Cr	283.56	2.70	3.1
Macintosh HD:L:	3/10/97 13:11	Cr	283.56	1.20	2.6
Macintosh HD:L:	3/10/97 13:17	Fe	238.20	1.71	30.2
Macintosh HD:L:	3/10/97 13:20	Fe	238.20	1.57	33.5
Macintosh HD:L:	3/10/97 13:24	Mn	257.60	0.40	1.6

22. OK 3/10/97 1:33pm

DAMES & MOORE TRAVERSE SOURCE TEST DATA

Company: BALBOA
Sampling Location: BALBOA BRIDGE
Test No.: 2 - Multi Metals (Times)

Date: 3.12.97
Time Start: 0930
Time End: 1710

System Pre-Test Leak Check:
0.002 cfm @ 14" Hg vac
(Pitot tube Leak Check: 0.0)

System Post-Test Leak Check:
0.000 cfm @ 14" Hg vac
(Pitot tube Leak Check: 0.0)

Time On	Source Point #	Gas Meter Reading (scf)	Stack				Probe Temp °F	Hot Box Temp °F	Dryer Temp °F	Meter Temp °F		Vacuum °Hg
			Velocity Head (H ₂ O)	Temp °F	Calculated dH	Set dH				in	out	
0		185.639	—	—	—	—	—	—	—	—	—	—
5	1	189.10	.016	138	1.72	1.7	247	251	55	72	69	1
10	2	192.55	.016	132	1.7	1.7	251	251	52	72	70	1
15	3	196.30	.018	138	1.91	1.9	250	251	50	78	71	1
20	4	199.70	.014	135	1.49	1.5	249	249	51	79	72	1
25	5	203.40	.017	139	1.80	1.8	250	250	53	80	73	1
30	6	207.02	.017	140	1.8	1.8	252	250	54	81	74	1
35	7	210.20	.013	136	1.3	1.3	251	249	54	82	79	1
40	8	213.615	.015	139	1.516	1.516	250	249	56	82	75	1
		(27.976)	(.016)	(138)	(1.7)	(1.7)	—	—	(53)	—	(70)	(1)
45	1	216.21	.008	134	.88	.88	250	250	59	81	72	1
50	2	219.17	.010	136	1.2	1.2	248	249	57	82	78	1
55	3	222.20	.011	140	1.22	1.2	252	250	57	83	78	1
60	4	224.92	.008	137	.88	.88	253	252	57	83	78	1
65	5	227.10	.008	139	.88	.9	248	250	56	84	79	1
70	6	230.23	.008	137	.88	.9	250	250	55	85	80	1
75	7	233.03	.009	140	1.01	1.0	248	250	56	85	80	1
80	8	235.844	0.080	143	1.124	1.0	250	250	56	85	81	1
			(0.009)	(138)	—	(1.01)	—	—	(57)	—	(81)	(1)
85	1	238.49	0.008	137	0.899	0.90	251	251	59	85	82	1
90	2	241.2	0.008	137	0.899	0.90	249	252	58	85	83	1
95	3	243.65	.008	138	.9	.9	250	250	58	85	83	1
100	4	246.70	.008	140	.9	.9	251	250	57	86	83	1
105	5	249.35	.007	135	.85	.85	251	249	52	86	84	1
110	6	251.7	.005	139	.56	.56	251	250	58	86	84	1
115	7	253.8	.005	140	.56	.56	247	250	59	87	84	1
119	8	255.604	.005	140	.56	.56	250	250	59	87	84	1
AVERAGES		69.965	.0106	138	—	1.112	—	—	58	—	81	1

PRE-TEST DATA
Nozzle Diameter: 0.626
Barometric Pressure: 30.03 *Hg
Static Pressure in Stack: -0.02 *H2O
Pitot Factor: 84
Est. Moisture: 18
Est. MW: 26

CHAIN OF CUSTODY INFO

Impingers Loaded By: JRW
Impingers Recovered By: JRW
Filter Loaded By: NA
Filter No.: —
Filter Recovered By: JRW
Probe Wash By: JRW

EQUIPMENT/CALIBRATION DATA

Pitot Tube No.: PT-003-1 (Cal: —)
Potentiometer No.: 458-2 (Cal: 3.492)
Thermocouple No.: 0.73-25-43 (Cal: 3.492)
Gas Meter No.: 2962389 (Cal: 3.492)
Meter Corr. Factor: 1.0083
Delta H @: 1.84263

	Soln.	Final	Initial	Net
Impinger #1	DRY	749.3	570.7	178.6
Impinger #2	DRY	534.0	534.0	0.0
Impinger #3	5% H ₂ O	749.3	691.2	58.1
Impinger #4	"	749.3	718.5	30.8
Impinger #5	DRY	621.2	625.1	-3.9
Impinger #6	SILICA	871.2	856.7	14.5

Total Wt Gain: 302.3

POST TEST INFO

Filter Appearance: Lot
Impinger Appearance: —
Silica Gel Spent (%): —

Recorded by: [Signature] Date: 3.12.97
Checked by: [Signature] Date: 4/18/97

Y = 112.40

Note: Some Readings are at 11:00 am on 3/12/97

Company: BALBOA
Sampling Location: BAL-DAC Ed
Test No. 2-502 (TIRES)

Date: 3.12.97
Time Start: 0932
Time End: 1122

System Post-Test Leak Check
0.02 in @ — *Hg vac
(Pitot tube Leak Check: —)

[illegible]

CHAIN OF CUSTODY INFO

Impingers Loaded By JRW
Impingers Recovered By JRW
Filter Loaded By N/A
Filter No. 1
Filter Recovered By JRW
Probe Wash By JRW

Pitot Tube No.: NA (Cal: —)
 Potentiometer No.: ↓ (Cal: —)
 Thermocouple No.: ↓ (Cal: —)
 Gas Meter No.: CM25B35F (Cal: 2.6.92)
 Meter Corr. Factor: 1.03163
 Delta H @: NA

POST TEST INFO

Filter Appearance light
Impinger Appearance clear/calc. - no partic.
Silica Gel Spent (%)

Recorded by: C. B. 274 Date: 3.17.97
Checked by: 9000 Date: 4.18.97

file	time	species	line	freq	($\mu\text{g}/\text{acm}$)
Macintosh HD:L:	3/12/97 8:53	He	388.86	100	102.1
Macintosh HD:L:	3/12/97 8:54	He	388.86	95	97.1
Macintosh HD:L:	3/12/97 9:26	Fe	238.2	5.4	25.8
Macintosh HD:L:	3/12/97 9:35	Fe	238.2	3.2	19.5
Macintosh HD:L:	3/12/97 9:42	Fe	238.2	5.87	46.9
Macintosh HD:L:	3/12/97 9:47	Mn	257.6	1.6	6.2
Macintosh HD:L:	3/12/97 9:52	Mn	257.6	1.8	8.9
Macintosh HD:L:	3/12/97 9:56	Fe	238.2	6.27	65.7
Macintosh HD:L:	3/12/97 10:00	Fe	238.2	4.4	30.4
Macintosh HD:L:	3/12/97 10:05	Mn	257.6	1.4	4.5
Macintosh HD:L:	3/12/97 10:09	Mn	257.6	2.1	6.4
Macintosh HD:L:	3/12/97 10:15	Mn	257.6	2.7	8.6
Macintosh HD:L:	3/12/97 10:21	Fe	259.9	5	21.9
Macintosh HD:L:	3/12/97 10:25	Ni	243.8	8.62	9.9
Macintosh HD:L:	3/12/97 10:30	Ni	243.8	13.8	-13.2
Macintosh HD:L:	3/12/97 10:35	Fe	238.2	5.62	66.3
Macintosh HD:L:	3/12/97 10:39	Fe	238.2	6.38	51.8
Macintosh HD:L:	3/12/97 10:43	Fe	238.2	4	68.8
Macintosh HD:L:	3/12/97 10:48	Mn	257.6	2	6.4
Macintosh HD:L:	3/12/97 10:52	Mn	257.6	3.2	11.8
Macintosh HD:L:	3/12/97 10:58	Mn	257.6	3.1	8.1
Macintosh HD:L:	3/12/97 11:02	Fe	238.2	6.87	76.4
Macintosh HD:L:	3/12/97 11:05	Fe	238.2	5.38	46.7
Macintosh HD:L:	3/12/97 11:11	Cr	283.56	3.3	1.6
Macintosh HD:L:	3/12/97 11:15	Cr	283.56	2.5	2.1
Macintosh HD:L:	3/12/97 11:20	Cr	283.56	2.42	0.8
Macintosh HD:L:	3/12/97 11:26	Mn	257.6	4.2	16.2
Macintosh HD:L:	3/12/97 11:29	Mn	257.6	4.4	15.6
Macintosh HD:L:	3/12/97 11:33	Fe	238.2	5.6	49
Macintosh HD:L:	3/12/97 11:36	Fe	238.2	6.71	64.1

22 AL 3/12/97 12pm

DAMES & MOORE TRAVERSE SOURCE TEST DATA

Company: BALBOA
Sampling Location: BAL-PAC Exh.
Test No. RVL-3 - C

Date: 3-14-97
Time Start: 1000
Time End: 1020

System Pre-Test Leak Check:
0.01 cfm @ 10 "Hg vac
(Pitot tube Leak Check: 0.0)

System Post-Test Leak Check
cfm @ "Hg vac
(Pitot tube Leak Check: 0.0)

Time On	Source Point #	Gas Meter Reading (scf)	Stack				Probe Temp °F	Hot Box Temp °F	Dryer Temp °F	Meter Temp °F		Vacuum °Hg
			Velocity Head (H ₂ O)	Temp °F	Calculated dH	Set dH				In	out	
0		358.53	—	—	—	—	—	—	—	—	—	—
5	1	347.0	.015	133	1.06	1.7	290	250	43	73	70	1
10	2	—	.012	130	1.32	1.3	292	250	51	70	71	1
15	3	348.33	.012	130	1.32	1.2	249	250	41	70	71	1
20	4	351.85	.010	132	1.22	1.8	249	250	41	70	72	1
25	5	355.5	.020	140	2.72	2.2	240	249	41	71	73	1
30	6	359.4	.012	135	1.40	1.0	237	251	41	72	74	1
35	7	365.49	.020	138	2.22	2.2	290	248	44	74	76	1
40	8	362.612	.019	135	2.12	2.1	251	251	44	75	77	1
		358.24	.024	132	—	1.81	—	—	42	—	76	1
45	9	374.42	.008	130	0.90	0.9	249	252	52	70	80	1
50	10	372.9	.008	135	2.3	0.8	249	252	43	72	85	1
55	11	376.1	.01	137	1.1	1.1	251	257	42	88	85	1
60	12	—	.008	133	0.9	0.9	252	247	42	90	85	1
65	13	380.0	.008	135	0.9	0.9	250	245	43	90	85	1
70	14	384.2	.008	135	—	—	249	240	45	91	85	1
75	15	387.0	.012	135	.80	.80	240	253	45	91	85	1
80	16	389.248	.018	135	9	9	249	246	42	91	86	1
		383.0	.004	136	—	(9.0)	—	—	45	88	88	1
85	17	393.1	.005	133	.52	.52	249	249	49	88	80	1
90	18	397.0	.005	135	2.4	1.4	247	245	40	80	80	1
95	19	399.2	.005	135	1.66	1.66	249	245	42	81	82	1
100	20	—	.005	135	4.2	5.2	247	244	48	81	82	1
105	21	—	.006	133	6.5	6.5	240	240	49	80	82	1
110	22	407.0	.006	135	6.0	6.0	251	242	40	80	82	1
115	23	409.5	.002	135	—	—	247	244	43	80	82	1
120	24	412.42	.002	135	2.4	2.4	247	240	40	80	82	1
AVERAGES		387.33	.010	130	—	1.4	—	—	44	—	84	1

PRE-TEST DATA

Nozzle Diameter: 0.25
Barometric Pressure: 29.8 "Hg
Static Pressure in Stack: -0.014 "H₂O
Pitot Factor: 0.9
Est. Moisture: 10
Est. MW: 20

CHAIN OF CUSTODY INFO

Impingers Loaded By [Signature]
Impingers Recovered By [Signature]
Filter Loaded By 1.6
Filter No. 1
Filter Recovered By [Signature]
Probe Wash By [Signature]

EQUIPMENT/CALIBRATION DATA

Pitot Tube No.: 20-003 (Cal: —)
Potentiometer No.: 4-3-2 (Cal: 5.0)
Thermocouple No.: UM-2-0.1 (Cal: 5.0)
Gas Meter No.: 1.000-3.5 (Cal: 0.0)
Meter Corr. Factor: 20.0-3.5
Delta H @ 1.000-3.5

Impinger #1
Impinger #2
Impinger #3
Impinger #4
Impinger #5
Impinger #6

Soln.	Final	Initial	Net
—	845.5	672.4	173.1
—	874.2	662.3	211.9
—	431.9	630.4	198.5
—	754.1	338.9	415.2
—	—	—	—
—	—	—	—

Total Wt Gain: 1413.7

POST TEST INFO

Filter Appearance 1.0
Impinger Appearance 0.0
Silica Gel Spent (%) 30%

Recorded by: [Signature] Date: 3/14/97
Checked by: [Signature] Date: 3/14/97

file	time	species	line	freq	($\mu\text{g/acm}$)
Macintosh HD:L:	3/14/97 9:11	He	388.9	100.00	99.3
Macintosh HD:L:	3/14/97 9:13	He	388.9	98.50	97.5
Macintosh HD:L:	3/14/97 10:05	Fe	238.2	7.10	97.6
Macintosh HD:L:	3/14/97 10:08	Fe	238.2	8.43	84.9
Macintosh HD:L:	3/14/97 10:12	Mn	257.6	1.80	7.1
Macintosh HD:L:	3/14/97 10:16	Mn	257.6	1.70	5.5
Macintosh HD:L:	3/14/97 10:21	Ni	243.8	8.20	-4.3
Macintosh HD:L:	3/14/97 10:26	Ni	243.8	8.30	4.0
Macintosh HD:L:	3/14/97 10:31	Mn	257.6	1.70	7.6
Macintosh HD:L:	3/14/97 10:36	Mn	257.6	1.90	7.8

J. OLL 3-14-97
 10m

DAMES & MOORE
CEMS FIELD DATA

Page 1 of 2

Client: Balboa Pacific
Location: Santa Fe Springs
Source: INCINERATOR

Date: 3-10-97
Engineers: BJS

			ANALYZERS				
			NOx	CO	O2	CO2	THC
PRE-TEST INFORMATION							
INSTR	TECO 48-52479-291			0-100PPM			
	TECO 42H-52914-294		0-100PPM				
	CAI 8300 N3L24807					0-10%	
	SERVO-MEX 1400 014208/655				0-25%		
	TECO 5HT 55166-302						0-25PPM
GASSES	SA9772	10-20-98			20.95	9.02	
	SA 7849	3/99					20.1 PPM
	AAL7946	4-11-98		89.8			
	SA 7550	11-21-98	90.5				
CALIBRATION INFORMATION							
	Span		90.5	89.8	20.95	9.02	20.1
	High Mid						
	Low Mid						
TEST DATA							
LOCAL CAL "O"	LOG 0		.2	.4	.01	.01	0
	SPAN "		90.4	89.4	20.91	9.07	20.0
BIAS "O"	LOG 0		.3	.7	.02	.1	.5
	SPAN "		88.7	88.9	20.81	9.04	19.8
RUN 1 1114/1316	LOG 1						
BIAS "O"	LOG 3		.3	.4	.02	.4	.6
	SPAN "		85.5	87.8	20.7	8.7 9.5	20.7
LOCAL "O"	"		.2	.2	0	.4	
	SPAN		86.5	88.3	20.8	9.5	

CEMS FIELD DATA

Page 2 of 2

Client: Balboa Pacific
Location: Santa Fe Springs
Source: INCINERATOR

Date: 3-12-97

Engineers: BJS

[illegible]



12/20/2017

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

CHART NO. 89773AN

10 20 30 40 50 60 70 80 90 100

Mar. 10.97 11:13
NOx 72.5PPM
THC 0.5ppm
CO 0.3

CO2 6.4%
O2 10.5%

RUN 1 11:14
START

Mar. 10.97 09:54
NOx 38.8PPM
THC 20.1ppm
CO 1.2

CO2 3.7%
O2 14.8%

19.8 THC

18
THC

Mar. 10.97 13:17

NOx
THC
CO

65.9PPM
0.3ppm
0.2

CO2
O2

6.0%
11.4%

END RUN 1316

CHART NO. 891316

290cm

CHART NO. 891316

305cm

CHART NO. 891316

2

1

0

24

23

22

21

20

19

18

17

16

10

20

30

40

50

60

70

80

90

100

10

20

30

40

50

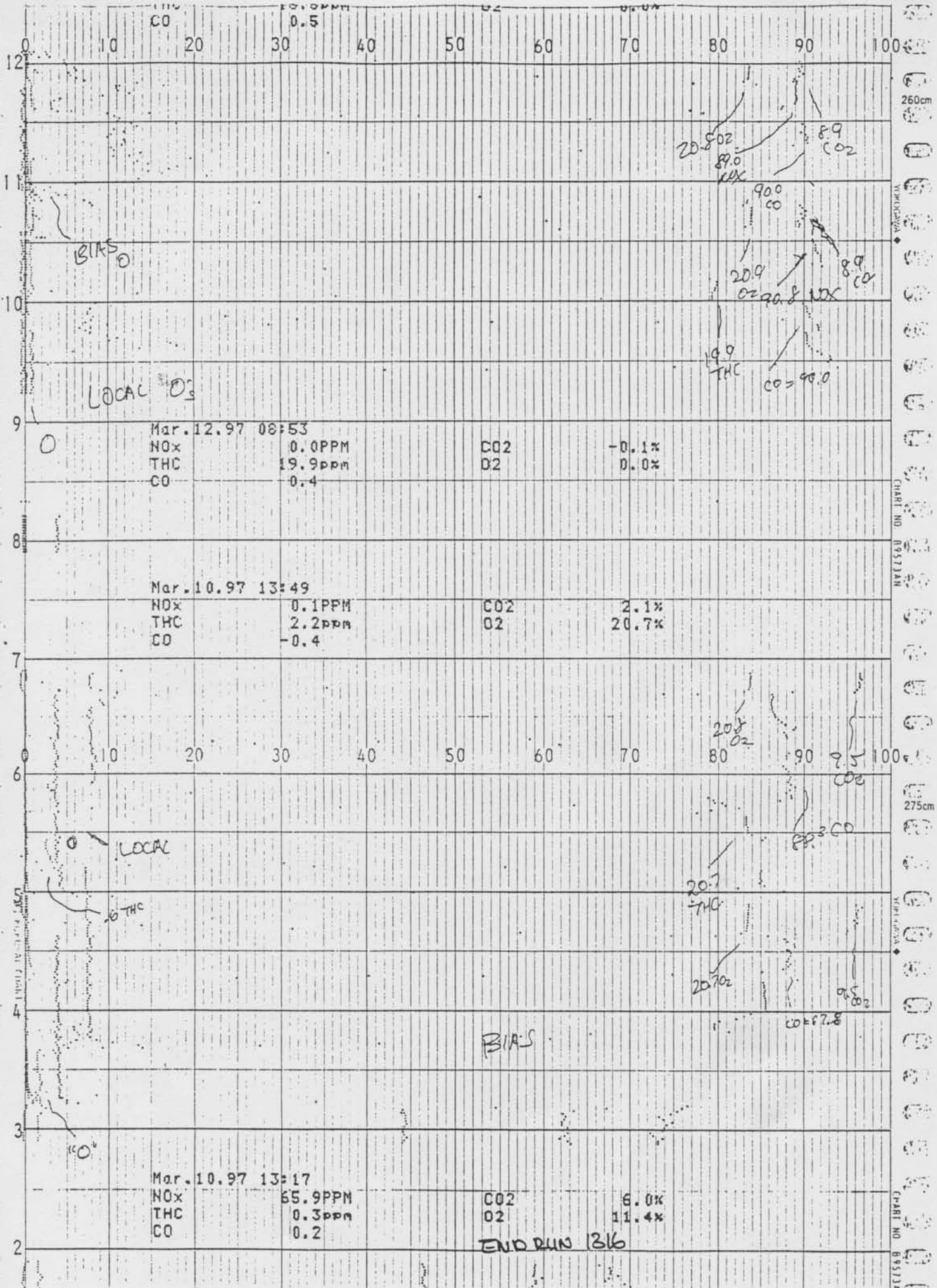
60

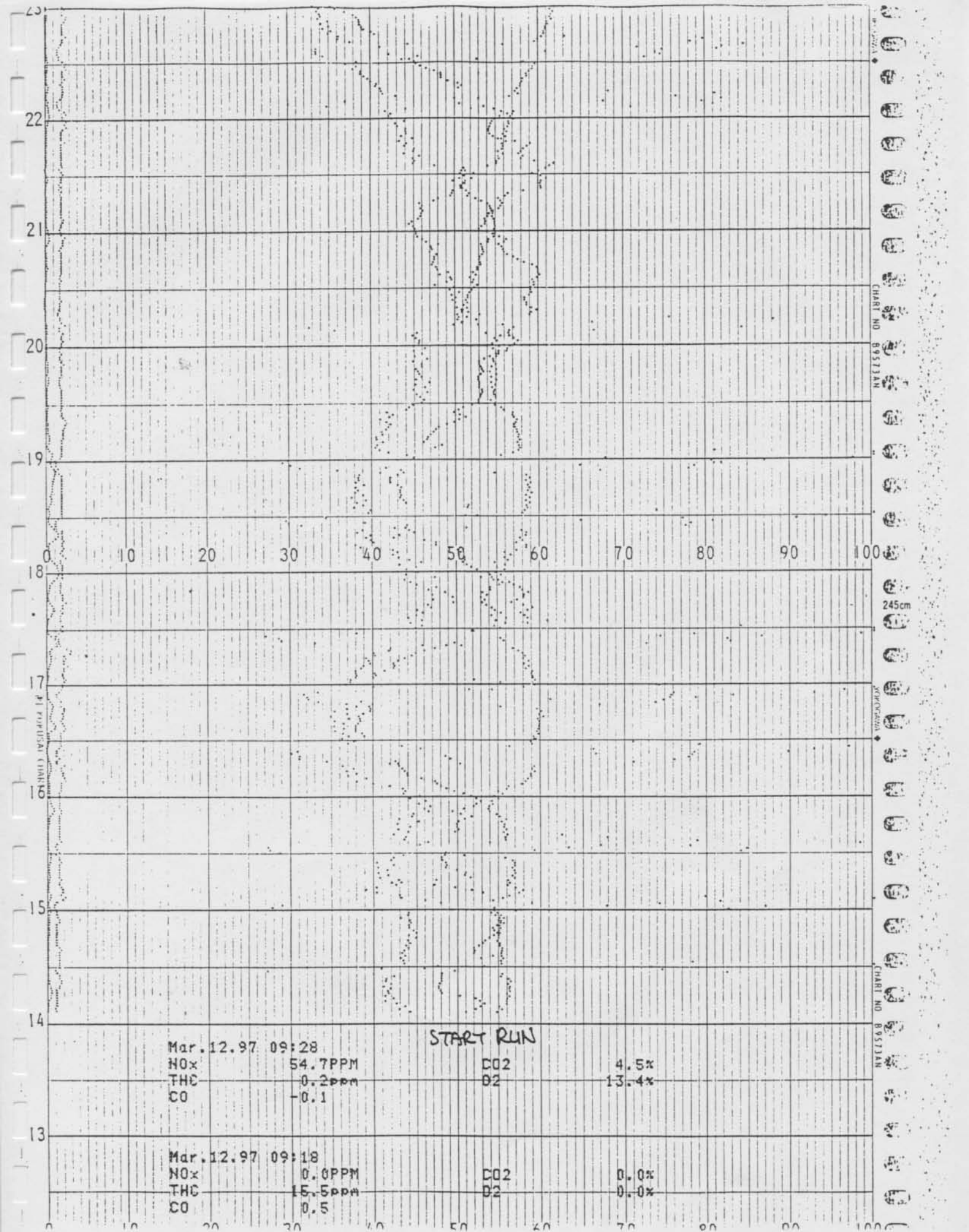
70

80

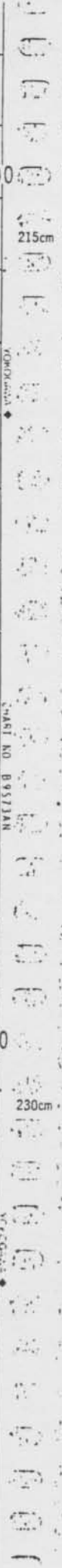
90

100





89.3
mk



11

Mar. 12.97 11:54
NOx 0.0PPM
THC 0.4ppm
CO -0.5

CO2 9.3%
O2 20.9%

10

9

8

7

6

5

4

3

2

Mar. 12.97 11:32
NOx 53.9PPM
THC 0.5ppm
CO -0.1

END RUN

CO2 4.3%
O2 14.1%

SMS
NOx
CO2

THC 20.3
O2 20.7
CO 27.0

89.3
NK

CO2 9.3
O2 20.9

CO2 11.9

200cm

215cm

YINCHUAN

CHARI NO. 85171

YINCHUAN

CHARI NO. 85171

200 MONI AT TIGHT

APPENDIX B
CALCULATIONS

DAMES & MOORE
EPA METHOD 29 - "DETERMINATION OF METAL EMISSIONS FROM STATIONARY SOURCES"

LOCATION: Balboa Pacific Corp. - Santa Fe Springs, CA
 SOURCE: BAL PAC Exhaust
 TRAIN: EPA Method 29; Multi-Metals

TEST DATE: 3/10/97 - 3/14/97
 SAMPLED BY: CB/TW
 CHECKED BY: AJH

Data Entry	SYMBOL	UNITS	DATA		
Test Number	--	--	R-1 - Munic. Waste	R-2 - Tires	R-3 - Moly. Cnt.
Time	--	--	1115-1225	0930-1200	1002-1430
Standard Temperature	t _{sd}	--	68	68	68
Round Stack Diameter	d _a	inches			
Rectangular Stack Length	L	inches	16.00	16.00	16.00
Width	W	inches	21.00	21.00	21.00
Nozzle Diameter	d _n	inches	0.626	0.626	0.626
Average Stack Temperature	t _s	deg F	138.0	138.0	136.0
Average Meter Temperature	t _m	deg F	99.0	81.0	84.0
Barometric Pressure	P _{bar}	in. Hg	29.99	30.03	29.99
Stack Static Pressure	P _g	in. H ₂ O	-0.025	-0.025	-0.025
Avg Delta H	dH	in. H ₂ O	1.240	1.160	1.140
Avg Velocity Head	dP	in. H ₂ O	0.011	0.011	0.010
Avg sqrt(dP) = sum(sqrt(dP))/N	sqrt(dP)	in. H ₂ O	0.104	0.101	0.098
Pitot Coefficient	C _p	none	0.84	0.84	0.84
Gas Sample Volume	V _m	cuft	73.478	69.965	69.932
Meter Calibration Factor	Y	none	1.0083	1.0083	1.0083
Total Sampling Time	min	minutes	120.5	119	120
Stack Gas Oxygen Content	O ₂	%	11.3	13.4	10.5
Stack Gas Carbon Dioxide Content	CO ₂	%	5.8	4.6	4.0
NOx Concentration	C _{nox}	ppmv	75.5	59.1	NA
CO Concentration	C _{co}	ppmv	-0.7	-0.3	NA
Net Weight Gain of Condenser Water (grams or milliliters)	g W _{wcg} or W _{wcm}	g/ml	323.7	292.7	287.0
Net Weight Gain of Silica Gel	W _{sg}	g	11.3	14.6	15.2
Calculated Data					
Nozzle Area, A _n = 3.14*(d _n /2) ²	A _n	sq. in.	0.3078	0.3078	0.3078
Stack Area, A _s = 3.1416*(d _a /2) ² /576 (round) = L*W/144 (Rectangular)	A _s	sq. ft.	2.333	2.333	2.333
Avg Stack Temperature, T _s = t _s + 460	T _s	deg R	598.0	598.0	596.0
Avg Meter Temperature, T _m = t _m + 460	T _m	deg R	559.0	541.0	544.0
Standard Temperature, T _{sd} = t _{sd} + 460	T _{sd}	deg R	528.0	528.0	528.0
Gas Sample Volume, V _m (std)=standard conditions, V _m (stack)=stack conditions V _m (std) = [T _{sd} /29.92]*Y*(V _m /T _m)*(P _{bar} +dH/13.6) V _m (stack) = (V _m (std) + V _{wc} (std)) * (T _s /T _{sd}) * (29.92/P _s)	V _m (std) V _m (stack)	cu.ft. cu.ft.	70.356 97.364	69.300 94.568	68.790 93.532
Water Collected, W _{wg} = [W _{wcg} + W _{sg} or (W _{wcm} *0.9982 g/ml) + W _{sg}]	W _{wc}	g	335.0	307.3	302.2
Volume of Water Vapor, V _{wc} (std) = W _{wc} *21.85*T _{sd} /(29.92*18.0*454)	V _{wc} (std)	cu. ft.	15.807	14.500	14.259
Moisture Fraction, B _{ws} = V _{wc} (std)/(V _m (std)+V _{wc} (std))	B _{ws}	none	0.1835	0.1730	0.1717
Dry Stack Gas Molecular Weight, M _d = (0.32*O ₂)+(0.44*CO ₂)+(0.28*(100-O ₂ -CO ₂))	M _d	g/g-mole	29.37318506	29.28	29.06
Wet Stack Gas Molecular Weight, M _w = M _d *(1-B _{ws}) + 18.0*(B _{ws})	M _w	g/g-mole	27.287	27.33	27.16
Absolute Stack Pressure, P _s = P _{bar} + P _g /13.6	P _s	in. Hg	29.99	30.03	29.99
Stack Gas Velocity, v _s = 85.49*C _p *sqrt(dP)*[sqrt(T _s /(P _s *M _w))] v _{sm} = 0.3048 * v _s	v _s v _{sm}	ft/sec m/sec	6.409 1.953	6.167 1.880	6.045 1.843
Actual Stack Gas Flow Rate, Q = 60*v _s *A _s	Q	acft/min	897	863	846
Dry Stack Gas Flow Rate (Dry, STP), Q _{sd} = 17.647*Q*(1-B _{ws})*(P _s /T _s) Q _{sdm} = Q _{sd} /35.32	Q _{sd} Q _{sdm}	dscft/min dscm/min	648 18.4	633 17.9	622 17.6
Isokinetic Rate, I = 100*A _s *V _m (std)/(min*(A _n /144)*Q _{sd})	I	%	98.31	100.48	100.54
NOx Emissions, as NO ₂ E _{nox} = C _{nox} * 46 / 385.3 * Q _{sd} * 60 * 1E-6 (@68 °F)	E _{nox}	lb/hr	0.35	0.27	NA
CO Emissions E _{co} = C _{co} * 28 / 385.3 * Q _{sd} * 60 * 1E-6 (@68 °F)	E _{co}	lb/hr	0.00	0.00	NA

DAMES & MOORE SOURCE TESTING
EPA Method 29 - Multi Metal Data Page

Company	Balboa Pacific	Test Date	3/10/97
Test Location	Santa Fe Springs, CA - BAL PAC	Test No.	Run-1, Municipal Waste
	start finish	Flow Rate, dscfm	648.37
Sampling Time	11:15 13:25	Sample Volume, dscf	70.36
		Sample Volume, stack	97.36

Metal	Molecular Weight	Detected Mass ug	Detection Limit ug	Detection Method	mg/hr	(in stack) ug/m ³	(in stack) ppbw
Aluminum (Al)	26.98	50.00	2.94	ICP-MS	23.67	15.52	13.84
Antimony (Sb)	121.75	1.40	0.10	ICP-MS	0.77	0.51	0.10
Arsenic (As)	74.92	NID	3.92	ICP-MS	1.08	1.42	0.46
Barium (Ba)	137.33	2.60	0.20	ICP-MS	1.44	0.94	0.17
Beryllium (Be)	9.01	NID	0.10	ICP-MS	0.03	0.04	0.09
Cadmium (Cd)	112.41	3.80	0.10	ICP-MS	2.10	1.38	0.29
Chromium (Cr)	52.00	8.00	1.96	ICP-MS	4.42	2.90	1.34
Copper (Cu)	63.55	15.00	0.39	ICP-MS	8.07	5.29	2.00
Lead (Pb)	207.20	29.00	0.10	ICP-MS	16.04	10.52	1.22
Iron (Fe)	55.85	110.00	29.40	ICP-MS	60.82	39.90	17.18
Manganese (Mn)	54.94	9.00	0.98	ICP-MS	4.98	3.26	1.43
Molybdenum (Mo)	95.94	210.00	0.10	ICP-MS	116.08	76.15	19.09
Nickel (Ni)	58.69	3.70	0.10	ICP-MS	2.05	1.34	0.55
Phosphorus (P)	30.97	67.00	29.40	ICP-MS	23.78	15.60	12.11
Selenium (Se)	78.96	ND	1.96	ICP-MS	0.54	0.71	0.22
Silver (Ag)	107.87	0.70	0.10	ICP-MS	0.39	0.25	0.06
Thallium (Tl)	204.38	ND	0.10	ICP-MS	0.03	0.04	0.00
Zinc (Zn)	65.39	50.00	1.96	ICP-MS	27.65	18.14	6.67

Mercury (Hg) M.W. = 200.59	Detected Mass ug	Detection Limit ug	Detection Method	mg/hr	ug/m ³	ppbw
Mercury Fraction # 1	NA	NA		0.00	0.00	0.0
Mercury Fraction # 2	NA	NA				
Mercury Fraction # 3	NA	NA				

* If the metal was not detected, 1/2 the detection limit was used for emission calculations.
* Emissions are blank corrected

Calculations

mg/hr = (Total ug) x (1 / Sample Vol) x (Flow Rate) x (60 min/hr) x (1 mg / 1000 ug)

mg/m³ = (Total ug) x (1 / Sample Vol (at stack conditions)) x (35.3 ft³ / 1 m³)

ppbw = (ug/m³) x (0.02405 m³/g-mol) x (1/MW) x (1g/1E6ug) x 1E9 ppbw

DAMES & MOORE SOURCE TESTING
EPA Method 29 - Multi Metal Data Page

Company Test Location	Balboa Pacific Santa Fe Springs, CA - BAL PAC		Test Date	3/12/97
	start	finish	Test No.	Run-2, Shredded Tires
Sampling Time	09:30	12:00	Flow Rate, dscfm	632.72
			Sample Volume, dscf	69.30
			Sample Volume, stack	94.57

Metal	Molecular Weight	Detected Mass ug	Detection Limit ug	Detection Method	mg/hr	(in stack) ug/m ³	(in stack) ppbw
Aluminum (Al)	26.98	68.00	2.94	ICP-MS	33.31	22.70	20.24
Antimony (Sb)	121.75	0.90	0.10	ICP-MS	0.49	0.34	0.07
Arsenic (As)	74.92	NID	3.92	ICP-MS	1.07	1.46	0.47
Barium (Ba)	137.33	4.30	0.20	ICP-MS	2.36	1.61	0.28
Beryllium (Be)	9.01	NID	0.10	ICP-MS	0.03	0.04	0.10
Cadmium (Cd)	112.41	4.40	0.10	ICP-MS	2.41	1.64	0.35
Chromium (Cr)	52.00	5.20	1.96	ICP-MS	2.85	1.94	0.90
Copper (Cu)	63.55	18.00	0.39	ICP-MS	9.64	6.57	2.49
Lead (Pb)	207.20	9.80	0.10	ICP-MS	5.37	3.66	0.42
Iron (Fe)	55.85	220.00	29.40	ICP-MS	120.52	82.15	35.38
Manganese (Mn)	54.94	13.00	0.98	ICP-MS	7.12	4.85	2.13
Molybdenum (Mo)	95.94	54.00	0.10	ICP-MS	29.55	20.14	5.05
Nickel (Ni)	58.69	4.70	0.10	ICP-MS	2.57	1.76	0.72
Phosphorus (P)	30.97	55.00	29.40	ICP-MS	16.98	11.58	8.99
Selenium (Se)	78.96	NID	1.96	ICP-MS	0.54	0.73	0.22
Silver (Ag)	107.87	0.50	0.10	ICP-MS	0.27	0.19	0.04
Thallium (Tl)	204.38	NID	0.10	ICP-MS	0.03	0.04	0.00
Zinc (Zn)	65.39	500.00	1.96	ICP-MS	273.91	186.71	68.67

Mercury (Hg) M.W. = 200.59	Detected Mass ug	Detection Limit ug	Detection Method	mg/hr	ug/m ³	ppbw
Mercury Fraction # 1	NA	NA		0.00	0.00	0.00
Mercury Fraction # 2	NA	NA				
Mercury Fraction # 3	NA	NA				

* If the metal was not detected, 1/2 the detection limit was used for emission calculations.

* Emissions are blank corrected

Calculations

mg/hr = (Total ug) x (1 / Sample Vol) x (Flow Rate) x (60 min/hr) x (1 mg / 1000 ug)

mg/m³ = (Total ug) x (1 / Sample Vol (at stack conditions)) x (35.3 ft³ / 1 m³)

ppbw = (ug/m³) x (0.02405 m³/g-mol) x (1/MW) x (1g/1E6ug) x 1E9 ppbw

DAMES & MOORE SOURCE TESTING

EPA Method 29 - Multi Metal Data Page

Company	Balboa Pacific	Test Date	3/14/97
Test Location	Santa Fe Springs, CA - BAL PAC	Test No.	Run-3, Catalyst
	start finish	Flow Rate, dscfm	622.45
Sampling Time	10:02 14:30	Sample Volume, dscf	68.79
		Sample Volume, stack	93.53

Metal	Molecular Weight	Detected Mass ug	Detection Limit ug	Detection Method	mg/hr	(in stack) ug/m ³	(in stack) ppbw
Aluminum (Al)	26.98	25.00	2.94	ICP-MS	9.66	6.72	5.99
Antimony (Sb)	121.75	0.20	0.10	ICP-MS	0.11	0.08	0.01
Arsenic (As)	74.92	ND	3.92	ICP-MS	1.06	1.48	0.48
Barium (Ba)	137.33	1.30	0.20	ICP-MS	0.71	0.49	0.09
Beryllium (Be)	9.01	ND	0.10	ICP-MS	0.03	0.04	0.10
Cadmium (Cd)	112.41	1.50	0.10	ICP-MS	0.81	0.57	0.12
Chromium (Cr)	52.00	2.70	1.96	ICP-MS	1.47	1.02	0.47
Copper (Cu)	63.55	8.60	0.39	ICP-MS	4.45	3.09	1.17
Lead (Pb)	207.20	4.00	0.10	ICP-MS	2.17	1.51	0.18
Iron (Fe)	55.85	370.00	29.40	ICP-MS	200.88	139.70	60.16
Manganese (Mn)	54.94	12.00	0.98	ICP-MS	6.51	4.53	1.98
Molybdenum (Mo)	95.94	44.00	0.10	ICP-MS	23.86	16.59	4.16
Nickel (Ni)	58.69	3.50	0.10	ICP-MS	1.90	1.32	0.54
Phosphorus (P)	30.97	42.00	29.40	ICP-MS	9.77	6.80	5.28
Selenium (Se)	78.96	ND	1.96	ICP-MS	0.53	0.74	0.23
Silver (Ag)	107.87	0.30	0.10	ICP-MS	0.16	0.11	0.03
Thallium (Tl)	204.38	ND	0.10	ICP-MS	0.03	0.04	0.00
Zinc (Zn)	65.39	74.00	1.96	ICP-MS	40.18	27.94	10.28

Mercury (Hg) M.W. = 200.59	Detected Mass ug	Detection Limit ug	Detection Method	mg/hr	ug/m ³	ppbw
Mercury Fraction # 1	NA	NA		0.00	0.00	0.00
Mercury Fraction # 2	NA	NA				
Mercury Fraction # 3	NA	NA				

* If the metal was not detected, 1/2 the detection limit was used for emission calculations.

* Emissions are blank corrected

Calculations

mg/hr = (Total ug) x (1 / Sample Vol) x (Flow Rate) x (60 min/hr) x (1 mg / 1000 ug)

mg/m³ = (Total ug) x (1 / Sample Vol (at stack conditions)) x (35.3 ft³ / 1 m³)

ppbw = (ug/m³) x (0.02405 m³/g-mol) x (1/MW) x (1g/1E6ug) x 1E9 ppbw

APPENDIX C
CONTINUOUS EMISSION DATA

DAMES & MOORE
CEMS GAS DATA FORM

LOCATION: Balboa Pacific Corp. - Santa Fe Springs, CA
SOURCE: BAL PAC Exhaust

TEST DATE: 3/12/97
Operator: CB

Test #	Gas	Full Scale	Cylinder Value	Analyzer Reading	Absolute Diff.	Diff. % Span	post	Analyzer Reading	Absolute Diff.	Diff. % Span
Zero	NOx	100	0	0.0	0.00	0.00%		0.0	0.00	0.00%
Mid Range			90.5	91	0.3	0.30%		90	0.20	0.20%
High Range	SA7550				0.00	0.00%			0.00	0.00%
Zero	CO	100	0	0.7	0.70	0.70%		0.0	0.00	0.00%
Mid Range			89.8	90	0.2	0.20%		88	2.00	2.00%
High Range	AAL7946				0.00	0.00%			0.00	0.00%
Zero	O2	25	0.01	0.01	0.00	0.00%		0.00	0.01	0.04%
Mid Range			20.95	20.90	0.05	0.20%		20.90	0.05	0.20%
High Range	SA9772				0.00	0.00%			0.00	0.00%
Zero	CO2	10	0	-0.10	0.10	0.40%		0.15	0.15	0.60%
Mid Range			9.02	8.90	0.12	0.48%		9.20	0.18	0.72%
High Range	SA9772				0.00	0.00%			0.00	0.00%
Pre Linearit NOx		NOx	#DIV/0!	CO	#DIV/0!	O2	84.0% CO2	#DIV/0!	#DIV/0!	
Post Linearit NOx		NOx	#DIV/0!	CO	#DIV/0!	O2	84.4% CO2	#DIV/0!	#DIV/0!	

DAMES & MOORE
CEMS BIAS DATA FORM

LOCATION: Balboa Pacific Corp. - Santa Fe Springs, CA
SOURCE: BAL PAC Exhaust

TEST DATE: 3/12/97
Operator: CB

Test #	Gas	Full Scale	Initial Cal Response	Initial Response	System Cal Bias	post	Final Response	System Cal Bias	Drift % of Range
Zero	NOx	100	0.20	0.2	0.00%		0.2	0.00%	0.00%
High Range			91	89	-1.80%		89.3	-1.50%	0.30%
Zero	CO	100	0.4	0.4	0.00%		0.2	-0.20%	-0.20%
High Range			90	90	0.00%		87	-3.00%	-3.00%
Zero	O2	25	0.02	0.02	0.00%		0.01	-0.04%	-0.04%
High Range			20.90	20.80	-0.40%		20.80	-0.40%	0.00%
Zero	CO2	10	0.01	-0.10	-0.44%		0.15	0.56%	1.00%
High Range			8.90	8.90	0.00%		9.19	1.16%	1.16%

Comments:		Uncorrected	Corrected
	NOx	58.25	59.06
	CO	0.03	-0.28
	O2	13.33	13.42
	CO2	4.65	4.62

BALBOA PACIFIC
 DAMES & MOORE CEMS DATA - TIRE BURN
 DATE: 3-12-1997
 TIME: 09:28:37.79

Time	NOX ppm	CO ppm	O2 %	CO2 %	THC ppm
9:28:38	54.13	-0.02	13.52	4.47	0.22
9:30:38	48.79	0.34	13.96	4.13	0.27
9:32:38	89.28	-0.09	13.58	4.44	0.34
9:34:38	54.48	0.07	13.70	4.42	0.27
9:36:38	71.55	-0.11	9.55	7.52	0.25
9:38:38	50.78	0.17	14.18	4.11	0.34
9:40:38	90.15	-0.13	12.55	5.34	0.40
9:42:38	49.63	0.50	13.50	4.51	0.39
9:44:38	47.15	0.04	14.51	3.87	0.44
9:46:38	80.29	-0.22	7.51	7.85	0.27
9:48:38	38.19	0.51	14.95	3.55	0.43
9:50:38	87.84	-0.14	7.90	7.63	0.32
9:52:38	42.16	0.24	14.56	3.85	0.60
9:54:38	86.35	0.13	7.42	8.41	0.29
9:56:38	57.10	0.29	13.20	4.79	0.54
9:58:38	53.81	1.27	13.68	4.43	0.44
10:00:38	47.30	0.08	14.42	3.88	0.47
10:02:38	97.06	0.19	13.88	4.36	0.44
10:04:38	42.29	0.59	14.63	3.83	0.44
10:06:38	100.07	0.56	14.02	4.20	0.41
10:08:38	49.14	0.07	14.21	4.17	0.55
10:10:38	53.01	-0.09	13.56	4.55	0.44
10:12:38	52.70	0.07	13.69	4.49	0.47
10:14:38	56.24	-0.02	11.78	5.09	0.48
10:16:38	58.19	-0.33	12.81	4.96	0.40
10:18:38	59.69	-0.15	13.07	4.84	0.44
10:20:38	55.84	0.17	13.46	4.57	0.53
10:22:38	54.76	-0.04	13.56	4.54	0.44
10:24:38	60.40	0.07	12.63	5.05	0.44
10:26:38	57.85	-0.08	13.78	4.47	0.44
10:28:38	54.47	0.21	14.13	4.26	0.59
10:30:38	59.12	0.12	14.24	4.14	0.51
10:32:38	44.53	0.11	13.39	4.13	0.45
10:34:38	39.45	-0.12	15.14	3.47	0.51
10:36:38	33.64	0.34	9.61	6.55	0.34
10:38:38	40.91	-0.21	14.66	3.77	0.52
10:40:38	41.31	0.06	14.53	3.90	0.51
10:42:38	62.31	-0.15	14.23	4.11	0.51
10:44:38	47.63	-0.05	14.36	4.06	0.54
10:46:38	94.68	-0.31	10.92	6.71	0.39
10:48:38	51.49	-0.15	14.25	4.17	0.49
10:50:38	55.19	-0.15	13.52	4.65	0.41
10:52:38	60.88	-0.15	13.23	4.87	0.45
10:54:38	58.92	-0.05	13.76	4.58	0.44
10:56:38	55.43	-0.17	14.14	4.29	0.44
10:58:38	49.45	0.03	10.44	5.86	0.35
11:00:38	47.72	-0.23	14.25	4.10	0.51
11:02:38	41.50	-0.05	14.71	3.79	0.51
11:04:38	71.68	-0.15	14.61	3.82	0.56
11:06:38	38.58	0.03	14.95	3.60	0.55
11:08:38	61.34	-0.21	14.90	3.62	0.56
11:10:38	37.39	0.05	15.19	3.45	0.53
11:12:38	57.74	-0.15	15.03	3.58	0.65
11:14:38	42.89	-0.05	14.41	4.01	0.55
11:16:38	92.30	0.08	14.21	4.16	0.51
11:18:38	44.90	-0.05	14.49	4.00	0.51
11:20:38	77.58	-0.24	14.23	4.17	0.56
11:22:38	52.14	0.16	14.14	4.30	0.60
11:24:38	80.48	-0.16	7.55	8.17	0.39
11:26:38	53.32	-0.24	13.89	4.45	0.51
11:28:38	57.61	-0.24	13.60	4.63	0.51

□	11:30:38	58.43	-0.15	13.80	4.53	0.56
		NOX ppm	CO ppm	O2 %	CO2 %	THC ppm
Average		58.25	0.03	13.33	4.65	0.46

APPENDIX D
CALIBRATION DATA

S-Type Pitot e Semi-Annual Calibration Sheet

Calibration Date 11/15/96 Calibrated By JRW

Pitot ID	Dt (in)	L (in)	dN (in)	dT (in)	dW (in)	P		Alpha		Beta		z (in)	w (in)
						A (in)	B (in)	A (d)	B (d)	A (d)	B (d)		
PT-003-1	0.374	95	0.827	0.885	6.750	0.456	0.456	6	1	0	0	0.000	0.010
PT-003-2	0.374	39	0.915	0.956	7.000	0.410	0.410	1	1	0	4	0.050	0.000
PT-005-1	0.253	58.5	0.800	0.903	7.000	0.308	0.308	1	0	1	3	0.040	0.010
PT-005-2	0.370	58.5	0.800	1.000	7.000	0.390	0.390	4	4	4	3	0.040	0.000
PT-008-1	0.375	95	0.827	0.930	7.250	0.456	0.456	3	2	2	2	0.000	0.025
PT-005-3	0.374	60	NA	0.775	60.00	0.472	0.472	3	2	1	0	0.030	0.015
PT-006-1	0.373	72	NA	NA	72.00	0.440	0.440	2	1	1	1	0.030	0.010

S-Type Criteria

Pitot ID	Assembly				Face Opening Planes			
	Dt	dN	dT	dW	P	Alpha	Beta	z
	<3/8	>3/4	>3/4	>3	<1.5Dt	<10	<5	<1/8
PT-003-1	PASS	PASS	PASS	PASS	PASS	PASS	PASS	PASS
PT-003-2	PASS	PASS	PASS	PASS	PASS	PASS	PASS	PASS
PT-005-1	PASS	PASS	PASS	PASS	PASS	PASS	PASS	PASS
PT-005-2	PASS	PASS	PASS	PASS	PASS	PASS	PASS	PASS
PT-008-1	PASS	PASS	PASS	PASS	PASS	PASS	PASS	PASS
PT-005-3	PASS	NA	PASS	PASS	PASS	PASS	PASS	PASS
PT-006-1	PASS	NA	NA	PASS	PASS	PASS	PASS	PASS

Dt - External Tubing Diameter
 L - Usable Length
 P - Distance from Long. Axis to Opening Plane
 dN - Distance from Pitot to Nozzle
 dT - Distance from Tubes to Thermocouple
 dW - Distance from End to First Obstruction
 Alpha - Opening Plane angle from Trans. Axis
 Beta - Opening Plane angle from Long. Axis
 z - Tube Ends Difference
 w - Tube Skew Separation

S-Type Pitot Tube - Semi-Annual Calibration Sheet

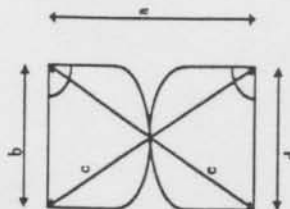
Pitot ID	Calibration Date				Calibrated By		JRW	
	L	P (A)	P (B)	dN	dT	dW	z	w
PT-003-1	95	0.456	0.456	0.827	0.885	6.75	0	0.01
PT-003-2	39	0.41	0.41	0.915	0.956	7	0.05	0
PT-005-1	58.5	0.308	0.308	0.8	0.903	7	0.04	0.01
PT-005-2	58.5	0.39	0.39	0.8	1	7	0.04	0
PT-008-1	95	0.456	0.456	0.827	0.93	7.25	0	0.025
PT-005-3	60	0.453	0.441	NA	0.775	60	0	0

PITOT MEASUREMENTS

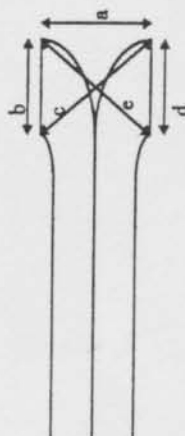
- L - Usable Length
- P - Distance from Long. Axis to Opening Plane
- dN - Distance from Pitot to Nozzle
- dT - Distance from Tubes to Thermocouple
- dW - Distance from End to First Obstruction
- z - Tube Ends Difference
- w - Tube Skew Separation

ANGLE CALCULATION MEASUREMENTS

Transverse Axis



Longitudinal Axis



Pitot ID	a	b	c	d	e
PT-003-1	0.832	0.374	0.876	0.374	0.905
PT-003-2	0.918	0.374	0.988	0.378	0.988
PT-005-1	0.615	0.253	0.66	0.252	0.663
PT-005-2	0.902	0.37	0.95	0.37	0.95
PT-008-1	0.927	0.375	0.979	0.375	0.99
PT-005-3	0.954	0.379	1.005	0.375	1.022

Pitot ID	a	b	c	d	e
PT-003-1	0.821	0.682	1.069	0.766	1.121
PT-003-2	0.915	0.615	1.099	0.617	1.067
PT-005-1	0.615	0.248	0.658	0.256	0.655
PT-005-2	0.883	0.643	1.049	0.62	1.048
PT-008-1	0.927	0.693	1.138	0.75	1.173
PT-005-3	0.954	0.611	1.118	0.562	1.117

Stack Temperature Sensor Calibration Data Form

Calibration Date 3/4/97
 Thermometer Model MicroComputer 7000P
 Umbilical Cord # UMB-003
 Method 5 Box ID M5B-1

Calibrator JRW
 Reference Extech #43141K
 Serial #T793272

Thermocouple Line #	Reference Thermometer Temp. (°C) ^a	Thermocouple Thermometer Temp. (°C)	Temperature Difference (%) ^b	Criteria ≤1.5%
1	0	0	0.00	PASS
1	100	100	0.00	PASS
1	260	260	0.00	PASS
2	0	0	0.00	PASS
2	100	100	0.00	PASS
2	260	262	0.38	PASS
3	0	0	0.00	PASS
3	100	100	0.00	PASS
3	260	262	0.38	PASS
4	0	0	0.00	PASS
4	100	98	0.54	PASS
4	260	260	0.00	PASS
5	0	0	0.00	PASS
5	100	100	0.00	PASS
5	260	260	0.00	PASS
6	0	0	0.00	PASS
6	100	100	0.00	PASS
6	260	260	0.00	PASS
7	0	0	0.00	PASS
7	100	100	0.00	PASS
7	260	262	0.38	PASS

^a Every 30° C for each reference point.

^b
$$\frac{(\text{ref temp } ^\circ\text{C} + 273) - (\text{test thermo temp } ^\circ\text{C} + 273)}{(\text{ref temp } ^\circ\text{C} + 273)} \times 100$$

BI-MONTHLY DRY GAS METER CALIBRATION

Meter ID	2962389
Box ID	M5B-2
Date	3/4/97
Cal. By	JRW

Ambient Temperature
Corrected Barometric Pressure

70 F
30.000 inHg

DRY GAS METER					REFERENCE METER # 2656207										
Flow	Time (min)	Volume (acf)	P-in (inH2O)	P-out (inH2O)	T-in (F)	T-out (F)	Volume (acf)	P-in (inH2O)	P-out (inH2O)	T-in (F)	T-out (F)	Y	Ave Y	Delta H @	Avg Delta H @
0.81 inH2O	0.00	897.00	0.81	0.73	65	64	590.98	0.07	0.00	71	71				
	10.46	902.20	0.81	0.73	68	65	596.24	0.07	0.00	72	72	1.01495		1.81949	
	20.48	907.20	0.81	0.73	70	66	601.26	0.07	0.00	72	72	1.00979	1.01065	1.83045	1.82511
0.49 scfm	30.05	912.00	0.81	0.73	71	67	606.06	0.07	0.00	73	73	1.00720		1.82540	
	2.00 inH2O	0.00	921.80	2.00	1.96	76	68	615.88	0.04	0.00	74	74			
0.76 scfm	5.40	926.00	2.00	1.96	77	69	620.09	0.04	0.00	74	74	1.00805		1.86189	
	10.13	929.70	2.00	1.96	78	70	623.77	0.04	0.00	74	74	1.00210	1.00590	1.86613	1.86015
	15.25	933.70	2.00	1.96	78	70	627.77	0.04	0.00	75	75	1.00755		1.85243	

* Standard Conditions are 29.92 inHg and 60 F

Ave Y of Reference Meter
Semi-Annual Double Ave Y
Bi-Monthly Double Ave Y
Average Delta H @

0.9981
1.0075
1.0083
1.842632

Validation Tests

[1-Y] < 0.05
[Y max - Y min] < = 0.01
Ave Y < 2 % Double Ave Y
Bi-Monthly < 2 % Semi-Annual
[H @ avg - H max, min] < = 0.15

PASS
PASS
PASS
PASS
PASS



213-585-2154
FAX: 213-585-0582

LIQUID CARBONIC

CYLINDER GAS PRODUCTS

5700 SOUTH ALAMEDA STREET • LOS ANGELES, CA 90058

CERTIFICATE OF ANALYSIS / EPA PROTOCOL GAS

CUSTOMER DAMES & MOORE

P.O NUMBER AIR 0049

REFERENCE STANDARD

COMPONENT	NIST SRM NO.	CYLINDER NO.	CONCENTRATION
PROPANE	GM1S vs. 1667b	SA 5183	49.7 ppm

ANALYZER READINGS

R=REFERENCE STANDARD

Z=ZERO GAS

C=GAS CANDIDATE

1. COMPONENT	PROPANE	GM1S	ANALYZER MAKE-MODEL-S/N	HORIBA, FIA-510, 851135122
ANALYTICAL PRINCIPLE		Flame Ionization Detector		LAST CALIBRATION DATE 02/01/96
FIRST ANALYSIS DATE		02/29/96		SECOND ANALYSIS DATE
Z 0	R 212049	C 85805	CONC. 20.1 ppm	Z R C CONC.
R 212017	Z 0	C 85650	CONC. 20.1 ppm	R Z C CONC.
Z 0	C 85557	R 211744	CONC. 20.1 ppm	Z C R CONC.
U/M mV		MEAN TEST ASSAY	20.1 ppm	U/M mV MEAN TEST ASSAY

Values not valid below 150 psig

THIS CYLINDER NO.	SA 7847	CERTIFIED CONCENTRATION	
HAS BEEN CERTIFIED ACCORDING TO SECTION	EPA-600/R93/224	PROPANE	20.1 ppm
OF TRACEABILITY PROTOCOL NO.	Rev. 9/93	NITROGEN	BALANCE
PROCEDURE	G1		
CERTIFIED ACCURACY	± 1 % NIST TRACEABLE		
CYLINDER PRESSURE	2000 PSIG		
CERTIFICATION DATE	02/29/96		
EXPIRATION DATE	02/29/99	TERM	36 MONTHS

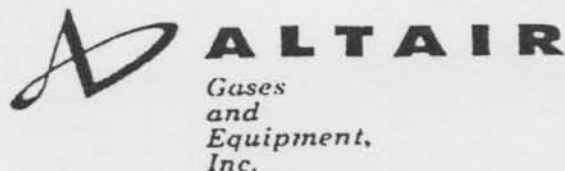
ANALYZED BY

MARIA E. ROCHON

CERTIFIED BY

KWAN T. YOUNG

2678 Bishop Drive, Suite 200
Caller Service 5161
San Ramon, California 94583-5161
Telephone: (510) 277-2100
Facsimile: (510) 866-8618



CERTIFICATE OF ANALYSIS / EPA PROTOCOL GAS

CUSTOMER AIR QUALITY ENGINEERING

P.O NUMBER 545492-00

REFERENCE STANDARD

COMPONENT		NIST SRM NO.	CYLINDER NO.	CONCENTRATION
CARBON DIOXIDE	GMIS	vs 1675b	SA 4193	9.02 %
OXYGEN	GMIS	vs 2659a	SA 19971	20.89 %

ANALYZER READINGS

R=REFERENCE STANDARD

Z=ZERO GAS

C=GAS CANDIDATE

1. COMPONENT	CARBON DIOXIDE	GMIS	ANALYZER MAKE-MODEL-S/N	Siemens Ultramat 5E	S/N A12-730
ANALYTICAL PRINCIPLE		NDIR		LAST CALIBRATION DATE	07/15/96
FIRST ANALYSIS DATE		08/14/96		SECOND ANALYSIS DATE	
Z 0.00	R 9.02	C 9.02	CONC. 9.02 %	Z	R C CONC.
R 9.02	Z 0.00	C 9.02	CONC. 9.02 %	R	Z C CONC.
Z 0.00	C 9.02	R 9.02	CONC. 9.02 %	Z	C R CONC.
U/M %		MEAN TEST ASSAY	9.02 %	U/M %	MEAN TEST ASSAY
2. COMPONENT	OXYGEN	GMIS	ANALYZER MAKE-MODEL-S/N	Siemens Oxymat 5c	S/N A12-839
ANALYTICAL PRINCIPLE		Paramagnetic		LAST CALIBRATION DATE	07/15/96
FIRST ANALYSIS DATE		08/14/96		SECOND ANALYSIS DATE	
Z 0.00	R 20.90	C 20.96	CONC. 20.95 %	Z	R C CONC.
R 20.90	Z 0.00	C 20.96	CONC. 20.95 %	R	Z C CONC.
Z 0.00	C 20.96	R 20.90	CONC. 20.95 %	Z	C R CONC.
U/M %		MEAN TEST ASSAY	20.95 %	U/M %	MEAN TEST ASSAY

Values not valid below 150 psig

THIS CYLINDER NO.	SA 9772	CERTIFIED CONCENTRATION	
HAS BEEN CERTIFIED ACCORDING TO SECTION	EPA-600/R93/224	CARBON DIOXIDE	9.02 %
OF TRACEABILITY PROTOCOL NO.	Rev. 9/93	OXYGEN	20.95 %
PROCEDURE	G1	NITROGEN	BALANCE
CERTIFIED ACCURACY	± 1 % NIST TRACEABLE		
CYLINDER PRESSURE	2000 PSIG		
CERTIFICATION DATE	08/14/96		
EXPIRATION DATE	08/14/99	TERM	36 MONTHS

ANALYZED BY

CERTIFIED BY

DOUG GRANT

IMPORTANT: The information contained herein has been prepared at your request by qualified experts at the Los Angeles Laboratory of Fluid Dynamics for Altair Gases & Equipment. While we believe that the information is accurate within the limits of the analytical methods employed and is complete to the extent of the specific analyses performed, we make no warranty of representation as to the suitability of the use of



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
DAMES & MOORE
6 HUTTON CENTER DRIVE SUITE 700
SANTA ANA, CA 92707

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

Purchase Order TONY
Project # 37304.004

ANALYTICAL INFORMATION

This certification was performed according to EPA Traceability Protocol For Assay and Certification of Gaseous Calibration Standards; Procedure G1; September 1993.

Cylinder Number AAL7946
Cylinder Pressure+ 2000 PSIG

Certification Date 04/11/95

Exp. Date 04/11/98

ANALYZED CYLINDER

Components
(CARBON MONOXIDE)

Certified Concentration
89.80 PPM

Analytical Uncertainty*
±1% NIST TRACEABLE

(Nitrogen)

Balance Gas

-Do not use when cylinder pressure is below 150 psig.

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

REFERENCE STANDARD

Type/SRM Sample No.
CRM 1679

Expiration Date
03/96

Cylinder Number
ALM024790

Concentration
96.20 PPM CO IN N2

INSTRUMENTATION

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

Last Date Calibrated
03/23/95

Analytical Principle
NDIR

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

Components	First Triad Analysis	Second Triad Analysis	Calibration Curve
CARBON MONOXIDE	Date: 04/04/95 Response Units: mv Z1= 0.00 R1= 42.0 T1= 39.2 R2= 42.0 Z2= 0.00 T2= 39.2 Z3= 0.00 T3= 39.2 R3= 42.0 Avg. Conc. of Cust Cyl. 89.76 PPM	Date: 04/11/95 Response Units: mv Z1= 0.00 R1= 42.0 T1= 39.2 R2= 42.0 Z2= 0.00 T2= 39.3 Z3= 0.00 T3= 39.2 R3= 42.0 Avg. Conc. of Cust Cyl. 89.84 PPM	Concentration= Ax^2+Bx+C A=0.0005612 B=0.9397 C=-0.01058
	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: 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=

Special Notes:

Analyst: _____

CERTIFICATE OF ANALYSIS / EPA PROTOCOL GAS

CUSTOMER DAMES & MOORE

P.O NUMBER

REFERENCE STANDARD

COMPONENT	NIST SRM NO.	CYLINDER NO.	CONCENTRATION
NITRIC OXIDE	GM13	SA 10284	85.0 ppm

ANALYZER READINGS

R=REFERENCE STANDARD

Z=ZERO GAS

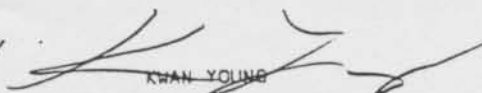
C=GAS CANDIDATE

I. COMPONENT	NITRIC OXIDE	GM13	ANALYZER MAKE-MODEL-S/N	Beckman 951A S/N 0101354	LAST CALIBRATION DATE	10/29/96
ANALYTICAL PRINCIPLE			Chemiluminescence		SECOND ANALYSIS DATE	11/21/96
FIRST ANALYSIS DATE			11/14/96			
Z 0	R 770	C 820	CONC. 90.5 ppm	Z 0	R 762	C 812
R 771	Z 0	C 820	CONC. 90.4 ppm	R 764	Z 1	C 813
Z 0	C 822	R 771	CONC. 90.6 ppm	Z 1	C 807	R 758
U/M	mv	MEAN TEST ASSAY 90.5 ppm		U/M	mv	MEAN TEST ASSAY 90.5 ppm

Values not valid below 150 psig
NOx values for reference only

THIS CYLINDER NO. SA 7550	EPA-600/R93/224	CERTIFIED CONCENTRATION
HAS BEEN CERTIFIED ACCORDING TO SECTION		NITRIC OXIDE 90.5 ppm
OF TRACEABILITY PROTOCOL NO. Rev. 9/93		NITROGEN BALANCE
PROCEDURE G1		NOx 91.1 ppm
CERTIFIED ACCURACY ± 1 % NIST TRACEABLE		
CYLINDER PRESSURE 2000 PSIG		
CERTIFICATION DATE 11/21/96		
EXPIRATION DATE 11/21/98	TERM 24 MONTHS	

ANALYZED BY



KWAN YOUNG

CERTIFIED BY



PHU-TIEN NGUYEN

APPENDIX E
LABORATORY DATA

March 27, 1997

DAMES & MOORE
6 Hutton Centre Drive
Suite 700
Santa Ana, CA 92707

Attn: Chris Barth

Job No: 34399

S

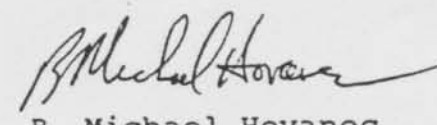
LABORATORY REPORT

Samples Received: Three (3) Impinger Samples
Date Received: 3-18-97
Purchase Order No: Proj. No: Sandia (Balboa Pacific)

The samples were analyzed as follows:

<u>Samples Analyzed</u>	<u>Analysis</u>	<u>Page</u>
Three (3) samples	Selected Metals by ICPMS	2 - 5

Page 1 of 5


Michael Shelton
Technical Director
B. Michael Hovanec
Senior Staff Chemist

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WEST COAST ANALYTICAL SERVICE, INC.

DAMES & MOORE
Attn: Chris BarthJob No: 34399
March 27, 1997

LABORATORY REPORT

Selected Metals
Quantitative Analysis Report
Inductively Coupled Plasma - Mass Spectrometry

Element	BALPAC- M29-R1 (ug/l)	Detection Limit (ug/l)	Impinger Vol (l)	BALPAC- M29-R1 (ug/sample)
Aluminum	51	3	0.98	50
Antimony	1.4	0.1	0.98	1.4
Arsenic	ND	4	0.98	ND<4
Barium	2.7	0.2	0.98	2.6
Beryllium	ND	0.1	0.98	ND<0.1
Cadmium	3.9	0.1	0.98	3.8
Chromium	8	2	0.98	8
Copper	14.6	0.4	0.98	15
Lead	29.5	0.1	0.98	29
Iron	111	30	0.98	110
Manganese	9	1	0.98	9
Molybdenum	214	0.1	0.98	210
Nickel	3.8	0.1	0.98	3.7
Phosphorus	68	30	0.98	67
Selenium	ND	2	0.98	ND<2
Silver	0.7	0.1	0.98	0.7
Thallium	ND	0.1	0.98	ND<0.1
Zinc	51	2	0.98	50

Date Analyzed: 3-20-97

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WEST COAST ANALYTICAL SERVICE, INC.

DAMES & MOORE
Attn: Chris BarthJob No: 34399
March 27, 1997

LABORATORY REPORT

Selected Metals
Quantitative Analysis Report
Inductively Coupled Plasma - Mass Spectrometry

Element	BALPAC- M29-R2 (ug/l)	Detection Limit (ug/l)	Impinger Vol (l)	BALPAC- M29-R2 (ug/sample)
Aluminum	70	3	0.97	68
Antimony	0.9	0.1	0.97	0.9
Arsenic	ND	4	0.97	ND<4
Barium	4.4	0.2	0.97	4.3
Beryllium	ND	0.1	0.97	ND<0.1
Cadmium	4.5	0.1	0.97	4.4
Chromium	5.4	2	0.97	5.2
Copper	18.3	0.4	0.97	18
Iron	231	30	0.97	220
Lead	10.1	0.1	0.97	9.8
Manganese	13.6	1	0.97	13
Molybdenum	56	0.1	0.97	54
Nickel	4.8	0.1	0.97	4.7
Phosphorus	57	30	0.97	55
Selenium	ND	2	0.97	ND<2
Silver	0.5	0.1	0.97	0.5
Thallium	ND	0.1	0.97	ND<0.1
Zinc	520	2	0.97	500

Date Analyzed: 3-20-97

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Page 3 of 5

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WEST COAST ANALYTICAL SERVICE, INC.

DAMES & MOORE
Attn: Chris BarthJob No: 34399
March 27, 1997

LABORATORY REPORT

Selected Metals
Quantitative Analysis Report
Inductively Coupled Plasma - Mass Spectrometry

Element	BALPAC- M29-R3 (ug/l)	Detection Limit (ug/l)	Impinger Vol (l)	BALPAC- M29-R3 (ug/sample)
Aluminum	37	3	0.67	25
Antimony	0.3	0.1	0.67	0.2
Arsenic	ND	4	0.67	ND<3
Barium	2	0.2	0.67	1.3
Beryllium	ND	0.1	0.67	ND<0.1
Cadmium	2.2	0.1	0.67	1.5
Chromium	4	2	0.67	2.7
Copper	12.8	0.4	0.67	8.6
Iron	550	30	0.67	370
Lead	6	0.1	0.67	4.0
Manganese	18	1	0.67	12
Molybdenum	65	0.1	0.67	44
Nickel	5.2	0.1	0.67	3.5
Phosphorus	63	30	0.67	42
Selenium	ND	2	0.67	ND<1
Silver	0.5	0.1	0.67	0.3
Thallium	ND	0.1	0.67	ND<0.1
Zinc	110	2	0.67	74

Date Analyzed: 3-20-97

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WCAS

WEST COAST ANALYTICAL SERVICE, INC.

DAMES & MOORE
Attn: Chris Barth

Job No: 34399
March 27, 1997

LABORATORY REPORT

Quality Control Summary

Sample: BALPAC-M29-R1 (Quality Control)

Matrix: Impinger

Parts Per Billion (ug/l)

	Sample	Duplicate	Avg	RPD %	Spike Conc.	MS	% Recovery
	-----	-----	-----	-----	-----	-----	-----
Aluminum	51	49	50	4.0	100	146	96
Antimony	1.37	1.48	1.43		100	86	85
Arsenic	ND	ND	ND		100	86	86
Barium	2.73	2.96	2.85		100	98	95
Beryllium	ND	ND	ND		100	96	96
Cadmium	3.9	4.2	4.05	7.4	100	93	89
Chromium	8.1	8.1	8.1		100	108	100
Copper	14.6	14.3	14.5	2.1	100	114	100
Iron	111	110	110.5		10000	10400	103
Lead	29.5	30.3	29.9	2.7	100	123	93
Manganese	8.6	8.7	8.65		100	108	98
Molybdenum	214	226	220	5.5	100	310	
Nickel	3.8	3.8	3.8		100	106	102
Selenium	ND	ND	ND		1000	780	78
Silver	0.68	0.73	0.71		100	96	96
Phosphorus	68	66	67		10000	8900	88
Thallium	ND	ND	ND		100	99	99
Zinc	51	50	50.5	2.0	100	136	86

Date Analyzed: 3-20-97

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Page 5 of 5

WCAS

Abbreviations Summary

General Reporting Abbreviations:

- B Blank - Indicates that the compound was found in both the sample and the blank. The sample value is reported without blank subtraction. If the sample value is less than 10X the blank value times the sample dilution factor, the compound may be present as a laboratory contaminant.
- D Indicates that the sample was diluted, and consequently the surrogates were too dilute to accurately measure.
- DL Detection Limit - Is the minimum value which we believe can be detected in the sample with a high degree of confidence, taking into account dilution factors and interferences. The reported detection limits are equal to or greater than Method Detection Limits (MDL) to allow for day to day and instrument to instrument variations in sensitivity.
- J Indicates that the value is an estimate.
- ND Not Detected - Indicates that the compound was not found in the sample at or above the detection limit.
- ppm Parts per million (billion) in liquids is usually equivalent to mg/l (ug/l), or in solids to mg/kg (ug/kg). In the gas phase it is equivalent to ul/l (ul/m³).
- ppb
- TR Trace - Indicates that the compound was observed at a value less than our normal reported Detection Limit (DL), but we feel its presence may be important to you. These values are subject to large errors and low degrees of confidence.
- | | | | | | | | |
|----|----------|----|-----------|----|------------|---|-------|
| kg | kilogram | mg | milligram | l | liter | m | meter |
| g | gram | ug | microgram | ul | microliter | | |

QC Abbreviations:

- Control & Warning Limits QC Limits are determined from historical data. The test value must be within the Control Limits for the test to be considered valid. Based on historical data, the confidence intervals are 95% for warning limits and 99% for control limits.
- % Error Percent Error - This is a measure of accuracy based on the analysis of a Laboratory Control Standard (LCS). An LCS is a reference sample of known value such as an NIST Standard Reference Material (SRM). The % Error is expressed in percent as the difference between the known value and the experimental value, divided by the known value. The LCS may simply be a solution based standard which confirms calibration (ICV or CCV - initial or continuing calibration verification), or it may be a reference sample taken through preparation and analysis.

CHAIN-OF-CUSTODY

PINK COPY - Project Manager

[illegible]



WEST COAST
ANALYTICAL
SERVICE, INC.
Analytical Chemists

April 9, 1997

DAMES & MOORE
6 Hutton Centre Drive
Suite 700
Santa Ana, CA 92707

Attn: Chris Barth

Job No: 34519

S

LABORATORY REPORT

Samples Received: One (1) Liquid Sample
Date Received: 4-1-97
Project No: 08040-131/QC-Balboa/Sandia

The sample was analyzed as follows:

<u>Samples Analyzed</u>	<u>Analysis</u>	<u>Page</u>
One (1) sample	Selected Metals by ICPMS	2 - 3

Page 1 of 3


Michael Shelton
Technical Director


D. J. Northington, Ph.D.
President

This report is to be reproduced in its entirety.

Client: DAMES & MOORE
Job No: 34519

Selected Metals
Quantitative Analysis Report
Inductively Coupled Plasma - Mass Spectrometry

Element	BAL-001 (ug/l)	Detection Limit (ug/l)	Impinger Vol (l)	BAL-001 (ug/sample)
Aluminum	35	3	0.207	7.2
Antimony	ND	0.1	0.207	ND<0.02
Arsenic	ND	4	0.207	ND<0.83
Barium	ND	2	0.207	ND<0.41
Beryllium	ND	0.1	0.207	ND<0.02
Cadmium	ND	0.1	0.207	ND<0.02
Chromium	ND	1	0.207	ND<0.21
Copper	2	0.2	0.207	0.41
Lead	ND	0.1	0.207	ND<0.02
Iron	ND	30	0.207	ND<6.2
Manganese	ND	0.2	0.207	ND<0.04
Molybdenum	0.3	0.1	0.207	0.06
Nickel	ND	0.7	0.207	ND<0.14
Phosphorus	115	30	0.207	24
Selenium	ND	2	0.207	ND<0.41
Silver	ND	0.1	0.207	ND<0.02
Tellurium	ND	0.1	0.207	ND<0.02
Thallium	ND	0.1	0.207	ND<0.02
Vanadium	ND	3	0.207	ND<0.62
Zinc	ND	4	0.207	ND<1.4

Date Analyzed: 4-3-97

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Page 2 of 3

WCAS

Client: DAMES & MOORE
Job No: 34519

Quality Control Summary

Sample: Batch QC
Matrix: Impinger

Parts Per Billion (ug/l)

	Sample	Duplicate	Avg	RPD %	Spike Conc.	MS	% Recovery
	-----	-----	-----	-----	-----	-----	-----
Aluminum	51	49	50	4.0	100	146	96
Antimony	1.37	1.48	1.43		100	86	85
Arsenic	ND	ND	ND		100	86	86
Barium	2.73	2.96	2.85		100	98	95
Beryllium	ND	ND	ND		100	96	96
Cadmium	3.9	4.2	4.05	7.4	100	93	89
Chromium	8.1	8.1	8.1		100	108	100
Copper	14.6	14.3	14.5	2.1	100	114	100
Iron	111	110	110.5		10000	10400	103
Lead	29.5	30.3	29.9	2.7	100	123	93
Manganese	8.6	8.7	8.65		100	108	98
Molybdenum	214	226	220	5.5	100	310	
Nickel	3.8	3.8	3.8		100	106	102
Selenium	ND	ND	ND		1000	780	78
Silver	0.68	0.73	0.71		100	96	96
Phosphorus	68	66	67		10000	8900	88
Thallium	ND	ND	ND		100	99	99
Vanadium	ND	ND	ND		100	128	128
Zinc	51	50	50.5	2.0	100	136	86

Date Analyzed: 3-20-97

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Page 3 of 3

WCAS

Abbreviations Summary

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- | | | | | | | | |
|----|----------|----|-----------|----|------------|---|-------|
| kg | kilogram | mg | milligram | l | liter | m | meter |
| g | gram | ug | microgram | ul | microliter | | |

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WHITE COPY - Original (Accompanies Samples)

6 HUTTON CENTRE DRIVE, SUITE 700
SANTA ANA, CALIFORNIA 92707
(714) 433-2000 FAX (714) 433-2364