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AP-42 Section 11.26
Reference 11
Report Sect. 4
Reference 4

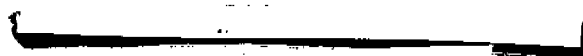
COMPLIANCE REPORT

PARTICULATE EMISSION COMPLIANCE PROGRAM - RETEST

for

ALPHA MILL

prepared for:



Purchase Order Number: E-20548

prepared by:

Air Quality Technical Services, Inc.

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September 28, 1990

FOREWORD

Air Quality Technical Services, Inc., an environmental consulting company specializing in the monitoring, measurement and evaluation of contaminants in source emissions and ambient air, was contracted by Cyprus-Windsor Minerals, Inc. to conduct a source emissions compliance program retest at the Columbia Mill facility which it owns and operates in

The following is a report of the results for the compliance program, the test methods and analytical procedures employed for determining the results, and all other pertinent data relevant to the tests conducted.

To the best of my knowledge, the data contained herein are correct and reliable.

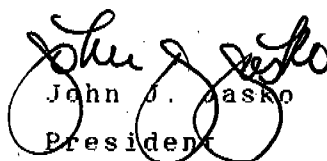

John J. Jasko
President

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1.0 INTRODUCTION

is a talc mineral processing facility. Raw minerals are reduced and classified to produce products which meet particle size range requirements of the end usage application for which they are intended.

On June 25, 26, 27, and 28, particulate emission testing was conducted on the Alpha Mill, Bagging Operation, Primary Crusher, and Screen & Belt Transfer to determine compliance with emission limits set forth in Condition 4 of an Air Pollution Control Permit (#AP-89-046) issued by the State of _____ Agency of Natural Resources. After completing a review of the reported results of the program, the Air Pollution Control Division (APCD) concluded that the Alpha Mill was operating in violation of its permit limits at the time of testing.

In a letter dated August 30, 1990, the APCD requested that I have the Alpha Mill source retested to demonstrate that repairs made to dust collector #1 after the initial compliance testing were effective in reducing emissions to within permitted limits.

On September 19 and 20, 1990, the Alpha Mill source was retested to determine compliance with the applicable particulate emission limits.

The major on-site representatives of the organizations participating in the program and their affiliations are as follows:

State of _____ Natural Resources
David Manning

Air Quality Technical Services, Inc.

Curtis Pujsto

Richard Stergas

2.0 SUMMARY OF RESULTS

2.1 ALPHA ROLLER MILL—Collector #1

The three test runs conducted on the Alpha Roller Mill resulted in particulate concentrations of 0.0007, 0.0011, and 0.0010 gr/dscf, and mass emission rates of 0.057, 0.091, and 0.081 lb/hr, respectively. The arithmetic means of the three test runs upon which compliance is judged were 0.0009 gr/dscf and 0.076 lb/hr. This represents a concentration that is 9.0 % of the limit of 0.01 gr/dscf and an emission rate that is 13.3 % of the limit of 0.57 lb/hr.

A summary of pertinent test results for the Alpha Roller Mill are found in Table 2-1.

TEST DATA SUMMARY SHEET

Alpha Roller Mill

Test/Run	1-1	1-2	1-3	Average
Date				
Test Time	1118-1352	1516-1745	0825-1053	

Plant Operating Data

Production Rate (TPH)				≈ 20
-----------------------	--	--	--	------

Test Measurements

Sample Volume (dscf)	67.154	69.295	68.829	
Isokinetics (%)	94.8	95.5	97.2	
Moisture (%)	2.5	3.2	3.4	3.0
Temperature (°F)	174.2	169.6	189.1	177.6
Velocity (fps)	54.4	55.4	56.8	55.5
Volumetric Flow (dscfm)	9491.2	9643.1	9484.1	9539.5
(acfm)	12565.3	12255.6	12034.4	12285.1
CO ₂ (%)	0.7	0.5	1.0	0.7
O ₂ (%)	20.7	20.5	20.0	20.4
N ₂ (%)	78.6	79.0	79.0	78.9

Particulate Emission Determinations

Concentration (gr/dscf)	0.0007	0.0011	0.0010	0.0009
Emission Rate (lb/hr)	0.057	0.091	0.081	0.076

Particulate Emission Standard

Emission Limit (gr/dscf)	0.01	0.01	0.01	0.01
Percent of Limit	7.0	11.0	10.0	9.0
Emission Limit (lb/hr)	0.57	0.57	0.57	0.57
Percent of Limit	10.0	16.0	14.2	13.3

Table 2-1

2.2 DATA REDUCTION

The results generated from the particulate emission compliance test program were computed by programmed methods. For comparative and illustrative purposes, Appendix A contains a data reduction example for the Alpha Mill test run 1-1 and a listing of calculation input parameters for each test run.

4.0 SOURCE DESCRIPTION

4.1 ALPHA MILL SYSTEM

The Alpha roller mill system consists of two sub-systems; a closed loop process system and an exhaust system. The closed loop process system consists of a furnace which supplies heated process makeup air, an induced draft fan which circulates the process air, a roller mill, a product recovery cyclone and two shaker screens. The exhaust system consists of a dust collector and an exhaust fan.

Crushed talc ore is fed into the roller mill from the raw product silo which has a storage capacity of approximately 800 tons. The talc ore is reduced by the interaction of a stationary circular race and rollers which rotate in a horizontal plane. The induced draft fan takes process air, combines it with a necessary volume of heated makeup air and forces the air stream up through the roller mill, entraining fine pulverized talc. The talc particles are removed from the air stream via the product recovery cyclone prior to returning to the process fan. Recovered product passes from the cyclone hopper into two bins, which, in turn feed the two shaker screens. Proper sized particles pass through the sieve and are conveyed pneumatically to storage. Oversized particles are screened off and pass back into the mill for further processing.

The exhaust system serves to prevent the buildup of humidity in the closed loop system by venting off a percentage of the recirculated air. Without this exchange, the moisture content of the conveying air stream would increase to saturation and would cause the talc product to agglomerate. That portion of the circulated air stream which is vented off passes through a fabric filter dust collector prior to being vented to the atmosphere. Recovered talc is discharged from the baghouse hopper through an air lock to the pneumatic conveying system and is carried to storage.

5.0 OPERATING AND PRODUCTION CAPACITY

The production rate for the source involved in the test program was subject to process variability between interconnected processes. Best efforts were made to test during maximum operating capacity of the source.

5.1 ALPHA ROLLER MILL SYSTEM

Alpha roller mill production rate was governed by the variable hardness of the raw ore entering the milling process and could not be controlled. A harder ore requires a longer milling time to achieve the desired grade of the final talc product and results in a lower production rate. The mill operates at 100% capacity as determined by this factor. The Alpha roller mill is used to produce one size grade of talc in the product 36 class.

Although the percentage of larger particles is varied, the percentage of fines present in the system for any given product size grade remains fairly constant. These fines, small enough to pass through the product recovery cyclone, are continuously transported through the circulating system. Fines are removed by the dust collector when air is drawn off from the closed loop process via the exhaust system. An estimate of the range of production rate for the Alpha Mill is 10 to 26 tons per hour. The estimated production rate during the test program was 20 tons per hour.

6.0 PARTICULATE EMISSION COMPLIANCE PROCEDURES

The procedures used in the particulate emission compliance program were conducted in accordance with the standard methods described in Appendix A of 40 CFR Part 60, STANDARDS OF PERFORMANCE FOR NEW STATIONARY SOURCES, as revised July 1, 1989.

6.1 SAMPLING LOCATIONS

The sampling locations were located according to the guidelines set forth in Method 1 - Sample and Velocity Traverses for Stationary Sources (40 CFR, Part 60, App. A, pp. 570 - 577).

6.1.1 Alpha Mill

The sample ports for the Alpha roller mill exhaust were located in the discharge duct approximately 18.4 feet downstream from the last disturbance and 4.6 feet upstream from a stack elbow. At the sampling location, the duct has an inside diameter of 26 inches. This positioned the sampling location 8.5 diameters downstream and 2.1 diameters upstream from the respective disturbances. A schematic of the sample port locations can be found in Figure 6-1. In accordance with Method 1, a minimum of 12 sample points were required for a particulate traverse. A cross-sectional view of the exhaust duct at the sampling location and the traverse point layout can be found in Figure 6-2.

ALPHA MILL EXHAUST STACK PORT LOCATIONS

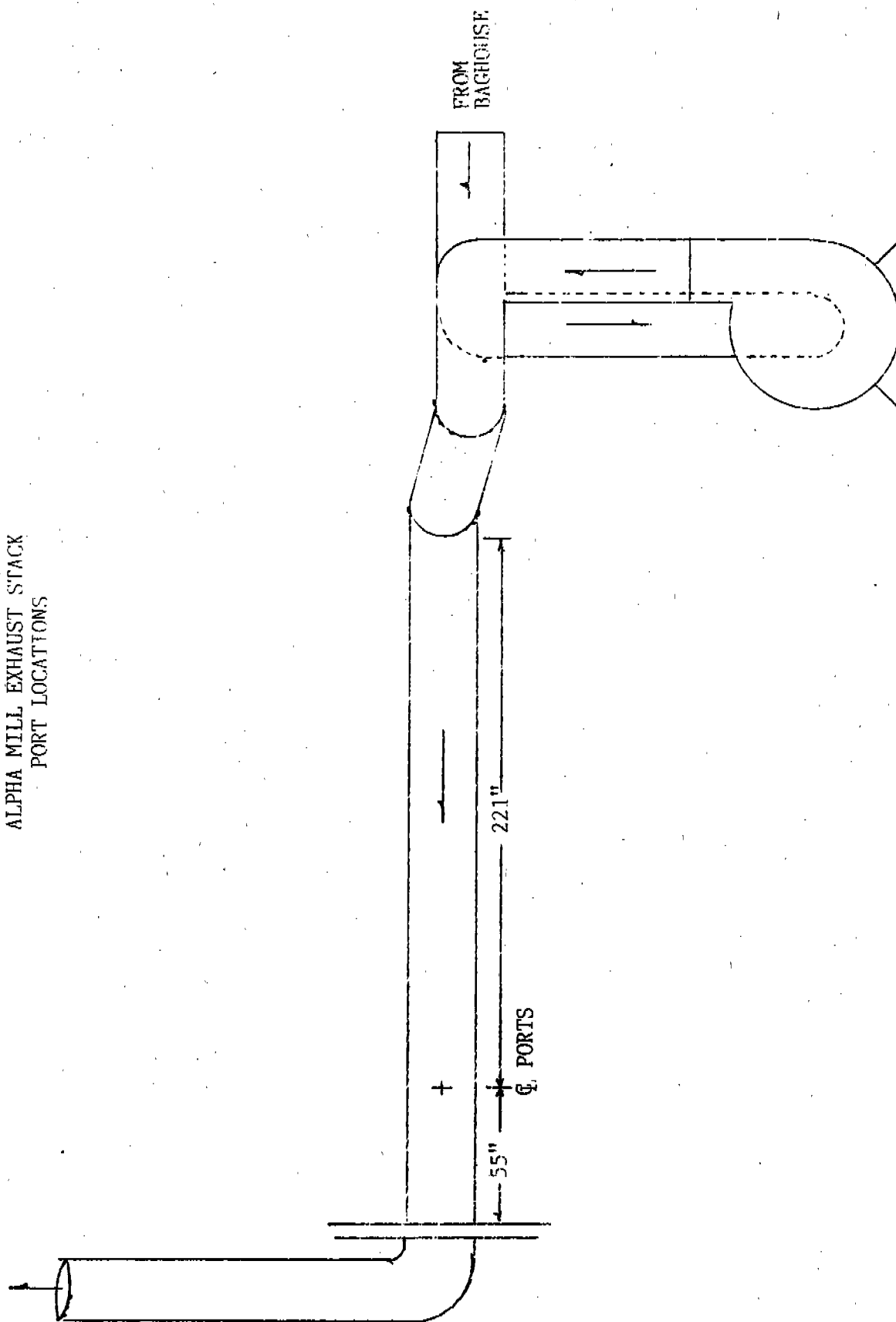
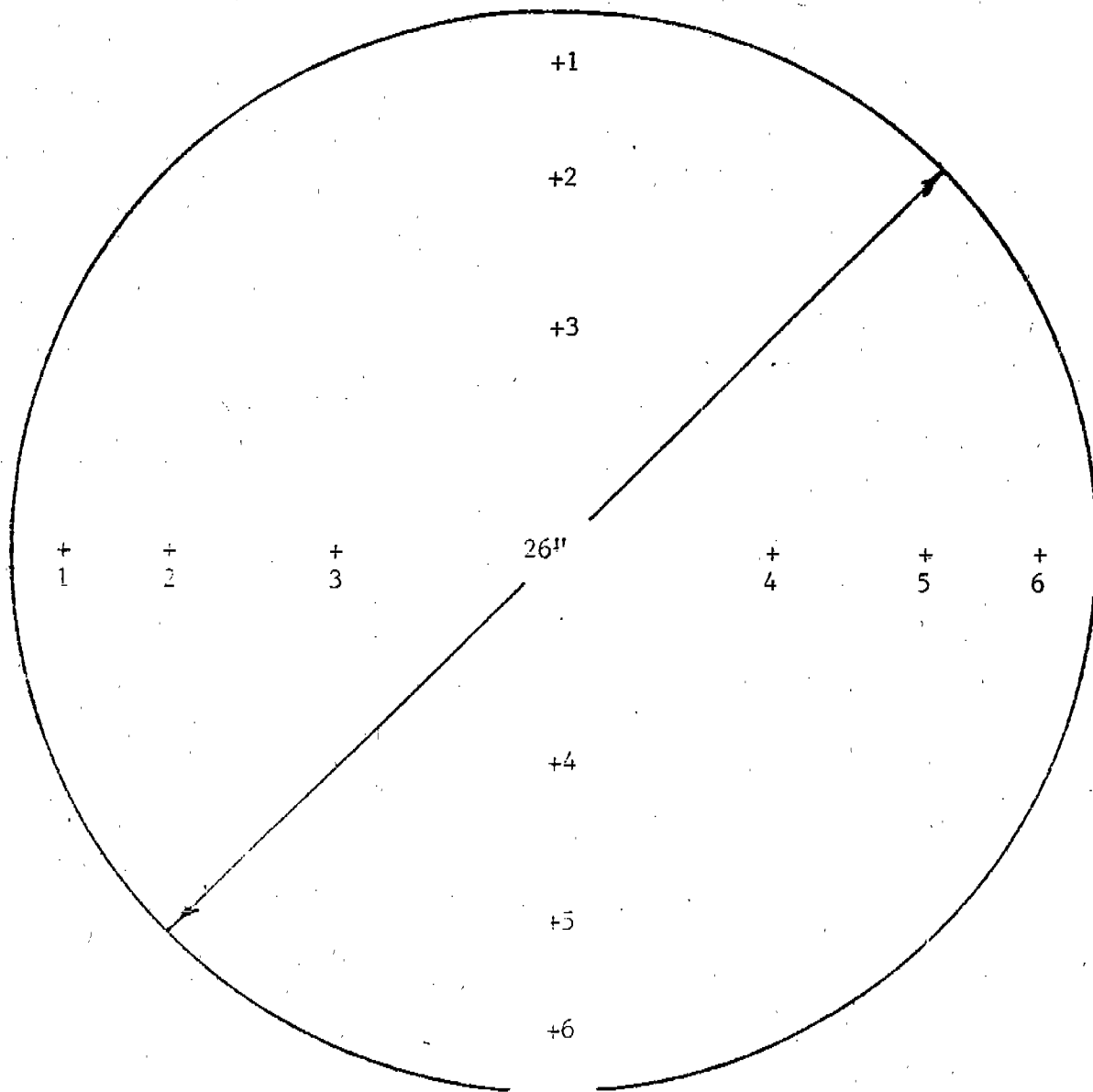


FIGURE 6-1

ALPHA MILL
TRAVERSE POINT LOCATION



<u>POINT</u>	<u>DISTANCE</u>
1.	1.1
2.	3.8
3.	7.7
4.	18.3
5.	22.2
6.	24.9

FIGURE 6-2

6.2 SAMPLE COLLECTION

Particulate sampling for the Alpha roller mill source was conducted in accordance with standard procedures described in EPA Method 17 - Determination of Particulate Emissions from Stationary Sources (in-stack filtration method) (40 CFR Part 60, App. A, pp. 851 - 868). The particulate sampling apparatus is depicted in Figure 6-3.

The sample train was operated so that an isokinetic rate $\pm 10\%$ was maintained over the course of each test run.

Copies of the field test data and summaries are found in Appendix B.

METHOD 17 SAMPLE TRAIN

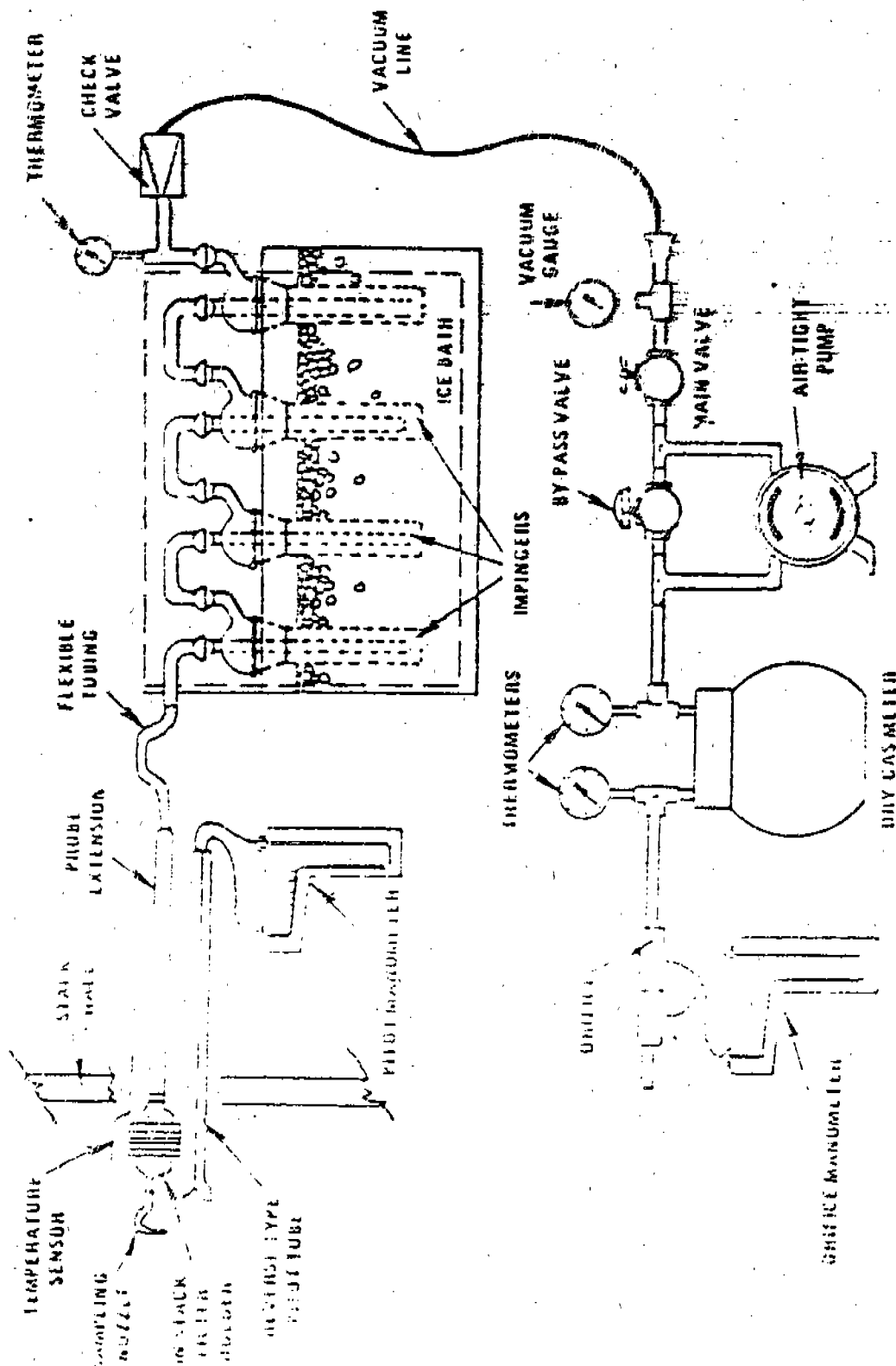


FIGURE 6-3

6.2.1 Collection System

The typical sample train consisted of:

- a) a nozzle properly sized to maintain isokinetic sampling;
- b) a stainless steel lined probe;
- c) an encased tared filter;
- d) a moisture condensing unit made up of four 500 ml impingers immersed in an ice water bath -
 - 1 modified Greenburg-Smith type containing 100 ml of water,
 - 1 Greenburg-Smith type containing 100 ml of water,
 - 1 modified Greenburg-Smith type empty,
 - 1 modified Greenburg-Smith type containing 200 grams of silica gel;
- e) an umbilical;
- f) a control console with:
 - a main on/off valve,
 - by-pass valve for flow adjustment,
 - leakless pump,
 - dry gas meter for measuring sample volume,
 - calibrated orifice equipped with pressure taps connected to an inclined manometer.

6.2.2 Sample Recovery

At the completion of each test run, the collected samples were recovered in the following manner:

Filter - The filter was removed from the filter holder, placed in it's original container, sealed and labeled for identification.

Front half rinse - The nozzle, probe, and front half of the filter holder were internally brushed and rinsed with acetone to remove any particulate matter which may have been deposited during the test run. The rinse collected in a glass jar, sealed and labeled for identification.

Impinger catch - The volume of water in the first three impingers was measured and the amount recorded.

Silica gel - The silica gel was removed from the fourth impinger, placed in it's original container, sealed and labeled for identification.

6.2.3 Quality Control/Quality Assurance

6.2.3.1 Equipment Calibration

The dry gas meter and orifice undergo semi-annual calibration according to procedures outlined in Method 5, Section 5 - Calibration (5.3 Metering System, §5.3.1 Calibration Prior to Use).

After the completion of the test program, the dry gas meter calibration was rechecked for accuracy according to procedures outlined in Method 5, Section 5 - Calibration (5.3 Metering System, §5.3.2 Calibration After Use).

Prior to use in the compliance test program the probe nozzle was calibrated according to procedures outlined in Method 5, Section 5 - Calibration (5.1 Probe Nozzle)

Copies of the pretest and post-test calibration sheets are found in Appendix C.

6.2.3.2 Equipment Leak Check

The sample train was leak checked according to procedures outlined in Method 5, Section 4 - Procedure (4.1 Sampling, §4.1.4 Leak-Check Procedures).

Before the start of each test run the inlet of the nozzle was plugged and a vacuum of 15" Hg was drawn and held. The metering dial was timed for a period of one minute and any movement during that period was noted. At the end of each test run the same procedure was followed using the

highest vacuum attained during the run. In each instance the maximum acceptable leakage rate was 0.02 cfm.

6.2.3.3 Chain of Custody

After recovery the samples collected during each test run were sealed in appropriate containers, labeled for identification, logged on a chain of custody record sheet, and placed in a secure container. The secure container remained in control of the project director in the field and until such time that it is returned to the lab. Any change of custody was recorded on the chain of custody record.

6.3 GAS VELOCITY/VOLUMETRIC FLOW DETERMINATIONS

Measurement of the stack gas velocity and volumetric flow was conducted in accordance with the standard procedures described in Method 2 - Determination of Stack Gas Velocity and Volumetric Flow Rate (40 CFR Part 60, App A, pp. 580 - 598).

These determinations were made simultaneous with the particulate sampling and were used to adjust the isokinetic sample rate.

6.3.1 Measurement Apparatus

The apparatus for measuring velocity head differential pressure and temperature profiles consisted of:

- a) a Type S (Stausscheibe) pitot tube with an assigned design coefficient of 0.84 connected to an inclined manometer;
- b) a thermocouple connected to a digital pyrometer.

6.3.2 Quality Control/Quality Assurance

6.3.2.1 Equipment Calibration

Prior to use in the test program the pitot tube assemblies were checked for conformity with the design specifications listed in Method 2, Section 4 - Calibration (4.1 Type S Pitot Tube, §4.1.1 Type S Pitot Tube Assemblies).

The thermocouple probe used in the test program undergoes annual calibration according to procedures outlined in the Quality Assurance Handbook, Section 3.1 - Method 2 (§3.1.2 Calibration of Apparatus).

6.3.2.2 Equipment Leak Check

The pitot tubes were leak checked according to procedures outlined in Method 2 (§3 Procedure).

The pitot tubes were subjected to leak checks prior to and after each sample run. The impact opening of the pitot was blown through until a minimum velocity pressure of 3" H₂O registered on the inclined manometer. The impact opening was then closed off and the pressure reading observed. The reading was required to remain stable for a period of 15 seconds. The same procedure was used to check the static pressure side of the pitot by applying suction to the static opening.

6.4 GAS COMPOSITION DETERMINATION

Stack gas composition was determined in accordance with the standard procedures described in Method 3 - Gas Analysis for Carbon Dioxide, Oxygen, Excess Air, and Dry Molecular Weight (40 CFR Part 60, App. A, pp. 606 - 613).

6.4.1 Fyrite Analysis

Several single-point grab samples were taken during the Alpha mill test runs and analyzed for CO_2 and O_2 by Fyrite analyzer, the balance considered N_2 . The results were used to determine the dry molecular weight of the effluent stream.

6.5 ANALYTICAL PROCEDURES

Sample analysis was conducted in accordance with the standard procedures described in Method 5 - Determination of Particulate Emissions From Stationary Sources.

Copies of the sample log sheets, filter tare weights, sample weight sheets, and lab analysis summary sheets are found in Appendix D.

6.5.1 Particulate Samples - Filters

Prior to use in the test program each glass fiber filter was:

- a) marked with an identifying number;
- b) desiccated for a minimum of 24 hours;
- c) weighed;
- d) re-desiccated for a minimum of 6 hours;
- e) reweighed to establish a final constant weight (<0.5 mg difference); and,
- f) sealed in a plastic petri dish container that was labeled with the identifying number and tare weight of the contained filter.

After sample collection each glass fiber filter used in the program was:

- a) desiccated for a minimum of 24 hours;
- b) weighed;
- c) re-desiccated for a minimum of 6 hours;
- d) reweighed to establish a final constant weight (<0.5 mg difference).

6.5.2 Particulate Samples - Front Half Rinse

The front half rinse of the sampling train from each test run, along with a blank sample of the acetone used in the recovery process, was:

- a) transferred from the sample jar to a tare weighed beaker that was marked with an identifying number;
- b) evaporated to dryness;
- c) desiccated for a minimum of 24 hours;
- d) weighed;
- e) re-desiccated for a minimum of 6 hours;
- d) reweighed to establish a final constant weight (<0.5 mg difference).

6.5.3 Moisture Content - Silica Gel

The silica gel quantities used in the test program were;

- a) initially tare weighed to 200 grams;
- b) re-weighed after sample collection to establish the net volume of moisture absorbed.

6.5.4 Quality Control/Quality Assurance

The filter media used for sample collection will be retained for a period of four months after analysis. A blank tare weighed filter from the group used in the test program will also be retained.

APPENDIX A

PARTICULATE CALCULATION LEGEND

A_s	=	Area of stack, ft^2
B_{wo}	=	Stack gas proportion of water vapor, by volume
C_p	=	Pitot tube coefficient
C_s	=	Particulate matter concentration, $gr/dscf$
D_n	=	Diameter of nozzle, inches
ΔH	=	Pressure differential across orifice, inches H_2O
M_p	=	Weight of particulate matter in grams
MW_d	=	Molecular weight of dry stack gas, $lb/lb-mole$
MW_s	=	Molecular weight of stack gas, $lb/lb-mole$
P_o	=	Barometric pressure, inches Hg
P_s	=	Pressure of stack, inches Hg
P_{st}	=	Static pressure of stack, inches H_2O
P_{std}	=	Standard pressure, 29.92 inches Hg
ΔP	=	Stack differential pressure, inches H_2O
Q_{sd}	=	Stack flow, $dscfm$ $Q_{sa} = \text{Stack flow, acfm}$
T_m	=	Temperature of gas meter, $^{\circ}F$
T_s	=	Temperature offstack, $^{\circ}F$
T_{std}	=	Standard temperature, $68^{\circ}F$
Vl_c	=	Volume of liquid condensate, ml
V_m	=	Volume of gas meter, ft^3
$V_{m_{std}}$	=	Volume of gas meter at standard conditions, $dscf$
V_s	=	Velocity of stack, ft/min
$V_{w_{std}}$	=	Volume of water at standard conditions, scf
γ	=	Gas meter correction factor
t	=	Test time, minutes
0.04707	=	Conversion factor, ml to ft^3
0.00857	=	Conversion factor, grains to pounds and minutes to hours
17.65	=	Conversion factor, standard temperature and standard pressure
17.33	=	Percent isokenetic constant from factoring
85.49	=	Pitot tube constant, K_p
0.264	=	Ratio of O_2 to N_2 in air, v/v
15.432	=	Conversion factor, $grams/ft^3$ to $grains/ft^3$

PARTICULATE CALCULATIONS

Client: _____

Unit: Alpha Mill

Facility: I

Test: 1-1

1. Volume of dry gas sampled at standard conditions, 68°F, 29.92" Hg (scf)

$$V_{m_{std}} = 17.65 V_m Y \frac{P_b + \frac{\Delta H}{13.6}}{T_m + 460}$$

$$= 17.65 (73.814) (0.997) \frac{29.03 + \frac{0.838}{13.6}}{102.7 + 460} = \underline{67.154}$$

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

$$V_{w_{std}} = 0.04707 V_{l_c}$$

$$= 0.04707 (37.2) = \underline{1.751}$$

3. Stack gas proportion of water by volume

$$B_{w_o} = \frac{V_{w_{std}}}{V_{w_{std}} + V_{m_{std}}}$$

$$= \frac{1.751}{1.751 + 67.154} = \underline{0.025}$$

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

$$MW_d = 28.94 (1 - B_{w_o}) + 18 B_{w_o}$$

$$= 28.94 (1 - 0.025) + 18 (0.025) = \underline{28.67}$$

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

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

$$= 28.94 (1 - 0.025) + 18 (0.025) = \underline{28.67}$$

6. Pressure of the stack, in. Hg

$$P_s = P_b - \frac{\Delta H}{13.6}$$

$$= 29.03 - \frac{0.60}{13.6} = \underline{29.07}$$

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

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

$$= 85.49 (0.84) (0.868) \sqrt{\frac{174.2 + 460}{(29.07) (28.67)}} = 54.4$$

8. Stack gas volume at standard conditions (scfm)

$$Q_s = 60 (1 - \beta_{wo}) V_{s,avg} A_s \left(\frac{528}{T_{s,avg} + 460} \right) \left(\frac{P_s}{29.92} \right)$$

$$= 60 (1 - 0.025) (54.4) (3.687) \left(\frac{528}{174.2 + 460} \right) \left(\frac{29.07}{29.92} \right) = 9491.2$$

acfm = 12034.4

9. Test percent isokinetic

$$XI = \frac{17.33 (T_s - 460) \left[0.04707 (V_{lc}) + V_{m, std} \right]}{\theta V_s P_s D_n^2}$$

$$= \frac{17.33 (174.2 + 460) \left[0.04707 (37.2) + 67.154 \right]}{(144) (54.4) (29.07) (0.1873)^2} = 94.8$$

10. Particulate matter concentration, gr/scf

$$C_s = 15.432 \left(\frac{W_s}{V_{s, std}} \right)$$

$$= 15.432 \left(\frac{0.0030}{67.154} \right) = 0.0007$$

11. Dilution rate of particulate matter, 10/hr

$$DR = 0.00357 (Q_s) C_s$$

$$= 0.00357 (9491.2) (0.0007) = 0.057$$

CALCULATION INPUT PARAMETERS

Test/run	1-1	1-2	1-3
Dry gas meter volume (cu ft)	73.814	75.458	76.117
Meter correction factor	0.997	0.997	0.997
Barometric press. (in Hg)	29.93	29.95	29.70
Avg orifice press. diff. (in H ₂ O)	0.833	0.893	0.932
Avg water temperature (°F)	102.5	96.0	99.3
Volume H ₂ O condensed (cc) - syringe	23.0	38.0	39.0
" " " " " " " " gel		11.1	12.1
" " " " " " " " total	27	19.1	5.2
CO ₂ (%)		0.5	0
O ₂ (%)	20	20.5	20.3
N ₂ (%)		79.3	79.0
CO (%)		0.0	0.0
Static press. (in H ₂ O)	0.40	0.64	0.60
PV of correction factor	0.997	0.997	0.997
Avg. static temp. of air (°F)	102.5	96.0	99.3
Avg. stack temperature (°F)	171.5	89.5	189
Area of stack (sq ft)	3.627	3.627	3.627
Test duration (min)	143	144	144
Diameter of nozzle (in)	0.1875	0.1880	0.1875
Mass of particulate catch (g) wash	0.0021	0.0020	0.0020
" " " " filter	0.0009	0.0031	0.0025
" " " " total	0.0030	0.0031	0.0045

APPENDIX B

PARTICULATE TEST FIELD DATA SUMMARY

Client: CWM, Inc.
Facility:
Date:
Unit: ALPHA MILL
Test Run: 1-1

Vm1 =	67.154	Vm =	73.814
VW1 =	1.751	Avg. Delta P =	0.755
QWO =	0.025	Avg. Delta H =	0.838
WV =	28.94	Avg. Ts =	174.2
WWS =	28.67	Avg. Tm =	102.7
WS =	29.07	Avg. SQRT Delta P =	0.868
TS =	54.4		
QSCM =	9491.2		
QSCM =	12034.4		
SI =	94.8		
	0.0007		
TA =	0.057		

NOTES:



READ AND RECORD ALL DATA EVERY

[illegible]



AIR QUALITY TECHNICAL SERVICES

QUALITY ASSURANCE FIELD SHEET
PARTICULATE SAMPLECLIENT: _____ LOCATION: exhaustSOURCE: Alpha MillTEST/RUN: 1-1 DATE: _____

FILTER DATA

Sample Number: B 332Type: Glass Fiber Number 47-076 Tare wt. 1.0244Holder Number: 1 Recovery Site: lab (noted) field Date: 19 Sept 90Physical Description: white low-loading

IMPINGER DATA

Recovery Site: lab / field Date: 9/19/90Physical Description: slightly cloudy white#1 Final 123 ml Initial 100 ml Net: 23 ml#2 Final 102 ml Initial 100 ml Net: 2 ml#3 Final 0 ml Initial 0 ml Net: - mlSI Gel Final 212.2 g Initial 200 g Net: 12.2 gTotal: 37.2 ml

WASH DATA

Sample Number: B 333Solvent: Acetone Recovery Site: lab (noted) field Date: 19 Sept 90Rinse Components: nozzle / hose / front of holderNet Volume: 216 mlPhysical Description: relatively clear

PHYSICAL DATA

Meter Volume: Final: 376.393 Initial: 302.569 Net: 73.814Barometric Pressure: 29.03 inches / mm HgTemperature: Dry Bulb: 70°F Wet Bulb: _____ Rel Humid: _____

Client:
Facility:
Date:
Unit: ALPHA MILL
Test Run: 1-2

NOTES :

Alpha

AIR QUALITY CONSULTING SERVICES

PARTICULATE TEST FIELD DATA SHEET

Client: Alpha
 Facility: Alpha
 Date: 1-2
 Unit: Alpha
 Test Location: Alpha
 Load/Conditions: 1-2
 Test Number: 26
 Stack Dimensions: 26" ID
 Operator: AS

Barometric Pressure: 28.95
 Static Pressure: +0.65
 Pitot Coefficient: 0.84
 Pitot Reading: 29.15A
 Meter Box Number: 1084
 Meter Box: 1084
 Nozzle Size & No. of Orifices: 1/4" 10
 Leak Rate: 0.002 CFM @ 11 in Hg
no. 6, 0.02 cm @ 14 in Hg

READ AND RECORD ALL DATA EVERY 12 MINUTES Pitot

TRAVERSE POINT NUMBER	CLOCK TIME (24-hr CLOCK)	GAS METER READING (V.M.)	VELOCITY (AP) (ft/min)	ORIFICE PRESSURE		STACK (ft)	TEMPERATURE		SAMPLE BOX (ft)	PROBE (ft)	IMPINGER (ft)	PUMP VACUUM in Hg	VAP
				DESIRED	ACTUAL		INLET	OUTLET					
A1	0	15:16	0.72	0.82	0.82	160	92	73			51	4	8.49
2	12	15:28	0.85	0.97	0.97	165	98	76			45	4	8.22
3	24	15:40	0.84	0.96	0.96	167	104	82			45	4	9.17
4	36	15:52	0.83	0.95	0.95	169	106	86			45	4	9.11
5	48	16:04	0.79	0.90	0.90	170	104	88			45	4	8.89
6	60	16:16	0.85	0.98	0.97	171	107	90			45	4	9.22
END	72	16:28											
B1	72	16:33	0.79	0.90	0.90	170	107	90			45	3.5	8.89
2	84	16:45	0.79	0.90	0.90	172	108	91			50	4	8.89
3	96	16:57	0.74	0.84	0.84	173	109	91			48	4	8.60
4	108	17:09	0.80	0.91	0.91	173	109	92			50	4	8.94
5	120	17:21	0.70	0.80	0.80	173	108	91			50	4	8.37
6	132	17:33	0.70	0.80	0.80	172	108	91			50	4	8.37
END	144	17:45											

TYPE

TIME	CO ₂	O ₂	CO
A	0.5	20.5	
B	0.5	20.5	

IMPINGER VOLUMES	IMPINGER	FINAL WT.
100	138	
100	100	
0	0	

WATER DATA
 2005 21.1
 47-077 101W



AIR QUALITY TECHNICAL SERVICES
QUALITY ASSURANCE FIELD SHEET
PARTICULATE SAMPLE

CLIENT: _____ LOCATION: exhaust

SOURCE: _____

TEST/RUN: 1-2 DATE: _____

FILTER DATA

Sample Number: B 335

Type: Glass Fiber Number 47-077 Tare wt. 1.0164

Holder Number: 2 Recovery Site: lab / field Date: 19 Sept 90

Physical Description: white low-load

IMPINGER DATA

Recovery Site: lab / field Date: 9/19/90

Physical Description: _____

#1 Final 138 ml Initial 100 ml Net: 38 ml

#2 Final 100 ml Initial 100 ml Net: 0 ml

#3 Final 0 ml Initial 0 ml Net: 0 ml

SI Gel Final 211.1 g Initial 200 g Net: 11.1 g

Total: 49.1 ml

WASH DATA

Sample Number: B 336

Solvent: Acetone Recovery Site: lab / field Date: 19 Sept 90

Rinse Components: nozzle / probe / front of holder

Net Volume: 11 ml

Physical Description: relatively clear

PHYSICAL DATA

Meter Volume: Final: 451.985 Initial: 376.527 Net: 75.458

Barometric Pressure: 28.95 inches mm Hg

Temperature: Dry Bulb: 270°F Wet Bulb: _____ Rel Humid: _____

PARTICULATE TEST FIELD DATA SUMMARY

Client:
 Facility:
 Date:
 Unit: ALPHA MILL
 Test Run: 1-3

VMstd =	68.829	VM =	76.117
VWstd =	2.41	Avg. Delta P	0.793
Bwo =	0.034	Avg. Delta S	0.903
MWd =	26.96	Avg. Is	189.1
MWs =	28.59	Avg. Im	99.8
Ps =	28.74	Avg. SQRT Delta P	0.89
Vs =	56.8		
Qscfm =	9484.1		
Qacfm =	12565.3		
%I =	97.2		
Cs =	0.001		
ER =	0.081		

NOTES:



AIR QUALITY TECHNICAL SERVICES
QUALITY ASSURANCE FIELD SHEET
PARTICULATE SAMPLE

CLIENT: _____ LOCATION: exhaust

SOURCE: _____

TEST/RUN: 1-3 DATE: _____

FILTER DATA Sample Number: B 338

Type: Glass Fiber Number 47-078 Tare wt. 1.0147

Holder Number: L Recovery Site: lab field Date: 20 Sept 90

Physical Description: low-loading of white / off-white

IMPINGER DATA

Recovery Site: lab field Date: 9/20/90

Physical Description: slightly cloudy

#1 Final 139 ml Initial 100 ml Net: 39 ml

#2 Final 100 ml Initial 100 ml Net: 0 ml

#3 Final 0 ml Initial 0 ml Net: 0 ml

Sol Gel Final 212.2 g Initial 200 g Net: 12.2 g

Total: _____

WASH DATA

Sample Number: B 339

Solvent: acetone Recovery Site: lab / field Date: 20 Sept

Rinse Components: nozzle probe / front of holder

Net Volume: 17 ml

Physical Description: relatively clean

PHYSICAL DATA

Meter Volume: Final: 529.072 Initial: 452.955 Net: 76.117

Barometric Pressure: 28.70 inches mm Hg

Temperature: Dry Bulb: 75°F Wet Bulb: _____ Rel Humid: _____

APPENDIX C

METER BOX CALIBRATION DATA AND CALCULATION FORM

(English units)

Date

Meter box number 1084-239

Barometric pressure, $P_b = 29.67$ in. Hg

Calibrated by PA

Orifice manometer setting (ΔH), in. H ₂ O	Gas volume		Temperatures				Time (θ), min	Y_i	ΔH_{t_i} in. H ₂ O
	Wet test meter (V_w), ft ³	Dry gas meter (V_d), ft ³	Wet test meter (t_w), °F	Inlet (t_{d_i}), °F	Outlet (t_{d_o}), °F	Avg ^a (t_d), °F			
0.5	5	5.122	70.2 70.2	88.41 75	73.75 78	83.5	13.1	0.999	1.897
1.0	5	5.192	70.1 70.1	77.101 74	80.81 82	79.3	4.4	0.998	1.927
1.5	10	10.465	70.1 70.1	79.102 76	83.55 85.7	83.8	15.7	0.995	2.007
2.0	10	10.504	70.1 70.1	79.102 78	87.88 90	87	3.7	0.995	2.023
3.0	10	10.517	70.1 70.1	79.102 78	89.90 90	89.5	1.3	0.997	2.059
4.0	10	10.536	70.1 70.1	79.102 78	92.92 93	92.4	7.8	1.000	2.045
Avg								0.997	1.995

ΔH , in. H ₂ O	$\frac{\Delta H}{13.6}$	$Y_i = \frac{V_w P_b (t_d + 460)}{V_d (P_b + \frac{\Delta H}{13.6}) (t_w + 460)}$	$\Delta H_{t_i} = \frac{1.0317 \Delta H}{Y_i} \left[\frac{t_w + 460}{t_d + 460} \right]$
0.5	0.155	$\frac{5(29.67)(513.5)}{5.122(29.67 + \frac{0.155}{13.6})(530.1)}$	$\frac{1.0317(0.155)}{0.999} \left[\frac{530.1}{513.5} \right]$
1.0	0.155	$\frac{5(29.67)(513.5)}{5.192(29.67 + \frac{0.155}{13.6})(530.1)}$	$\frac{1.0317(0.155)}{0.998} \left[\frac{530.1}{513.5} \right]$
1.5	0.110	$\frac{10(29.67)(553.8)}{10.465(29.67 + \frac{0.110}{13.6})(530.1)}$	$\frac{1.0317(0.110)}{0.995} \left[\frac{530.1}{553.8} \right]$
2.0	0.147	$\frac{10(29.67)(557)}{10.504(29.67 + \frac{0.147}{13.6})(530.1)}$	$\frac{1.0317(0.147)}{0.995} \left[\frac{530.1}{557} \right]$
3.0	0.221	$\frac{10(29.67)(558.5)}{10.517(29.67 + \frac{0.221}{13.6})(530.1)}$	$\frac{1.0317(0.221)}{0.997} \left[\frac{530.1}{558.5} \right]$
4.0	0.234	$\frac{10(29.67)(564)}{10.536(29.67 + \frac{0.234}{13.6})(530.1)}$	$\frac{1.0317(0.234)}{1.000} \left[\frac{530.1}{564} \right]$

^a If there is only one thermometer on the dry gas meter, record the temperature under t_d .

POSTTEST DRY GAS METER CALIBRATION DATA FORM (English units)

Test numbers 17943 Date 9/9/90 Meter box number 1084-239 Plant

Barometric pressure, $P_b = 29.69$ in. Hg Dry gas meter number

Pretest $Y = 0.777$

Orifice manometer setting, (ΔH), in. H ₂ O	Gas volume		Temperature		Time (θ), min	Vacuum setting, in. Hg	Y_i	V_i
	Wet test meter (V_w), ft ³	Dry gas meter (V_d), ft ³	Wet test meter (t_w), °F	Dry gas meter Inlet Outlet Average (t_{d_i}), (t_{d_o}), (t_d), °F				
-1.0	10	10.364	67.5	86.104 86.5 76.82	20.3		0.997	$V_w P_b (t_d + 460)$ $V_d (P_b + \frac{\Delta H}{13.6}) (t_w + 460)$ $\frac{10(29.69)(86.5 + 460)}{10.364(29.61 + \frac{1.0}{13.6})(67.5 + 460)}$
-1.0	10	10.499	67.9	101.104 94.7 83.87	20.4		0.998	$\frac{10(29.69)(94.7 + 460)}{10.499(29.61 + \frac{1.0}{13.6})(67.9 + 460)}$
-1.0	10	10.564	68.0	103.104 96.8 88.90	20.7		0.996	$\frac{10(29.69)(96.8 + 460)}{10.564(29.61 + \frac{1.0}{13.6})(68 + 460)}$
							$Y = 0.997$	

POST CAL CORRECTION 9/10/90

^a If there is only one thermometer on the dry gas meter, record the temperature under t_d .

V_w = Gas volume passing through the wet test meter, ft³.

V_d = Gas volume passing through the dry gas meter, ft³.

t_w = Temperature of the gas in the wet test meter, °F.

t_{d_i} = Temperature of the inlet gas of the dry gas meter, °F.

t_{d_o} = Temperature of the outlet gas of the dry gas meter, °F.

t_d = Average temperature of the gas in the dry gas meter, obtained by the average of t_{d_i} and t_{d_o} , °F.

ΔH = Pressure differential across orifice, in. H₂O.

Y_i = Ratio of accuracy of wet test meter to dry gas meter for each run.

Y = Average ratio of accuracy of wet test meter to dry gas meter for all three runs;
tolerance = pretest $Y \pm 0.05Y$

P_b = Barometric pressure, in. Hg.

θ = Time of calibration run, min.

**STACK TEMPERATURE SENSOR
CALIBRATION DATA FORM**

METER BOX
1084-239

Date: 2 Jan 89 Thermocouple number: DRY GAS INLET

Ambient temperature: 73 F Barometric pressure: 29.84

Calibrated by: C Purste Reference: mercury-in-glass ✓

other _____

Reference Point Number	Source ^a	Reference Thermometer Temperature		Thermocouple Potentiometer Temperature		Temperature Difference ^b %
		°C	°F	°C	°F	
1	ICE H ₂ O	0.0		32.0	F	0.0
2	MIS H ₂ O	17.0		62		0.0
3	32.0 H ₂ O	0.0		32		0.0

Where:

a = Type of calibration system used

b = Metric units

$$\frac{(\text{ref temp } ^\circ\text{C} + 273) - (\text{test thermom temp } ^\circ\text{C} + 273)}{\text{ref temp } ^\circ\text{C} + 273} \quad 100 \leq 1.5\%$$

b = English units

$$\frac{(\text{ref temp } ^\circ\text{F} + 460) - (\text{test thermom temp } ^\circ\text{F} + 460)}{\text{ref temp } ^\circ\text{F} + 460} \quad 100 \leq 1.5\%$$

**STACK TEMPERATURE SENSOR
CALIBRATION DATA FORM**

METER BOX
1084-237

Date: _____

Thermocouple number: OUTLET GAS METER

Ambient temperature: 73°F

Barometric pressure: 29.84

Calibrated by: C. PISTO

Reference: mercury-in-glass ✓

other _____

Reference Point Number	Source ^a	Reference Thermometer Temperature		Thermocouple Potentiometer Temperature		Temperature Difference ^b %
		°C	°F	°C	°F	
1	ICE H ₂ O	0.0	C	32.3	F	0.0
2	AIR IN ₂ S	76.0	C			0.0
3	BOIL H ₂ O	100.0	C			0.1

Where:

a = Type of calibration system used

b = Metric units

$$\frac{(\text{ref temp } ^\circ\text{C} + 273) - (\text{test thermom temp } ^\circ\text{C} + 273)}{\text{ref temp } ^\circ\text{C} + 273} \quad 100 \leq 1.5\%$$

b = English units

$$\frac{(\text{ref temp } ^\circ\text{F} + 460) - (\text{test thermom temp } ^\circ\text{F} + 460)}{\text{ref temp } ^\circ\text{F} + 460} \quad 100 \leq 1.5\%$$

ITOT # 29 15 DATE 11

OBSERVATIONS BY J.P.

- 1) All construction criteria for an isolated "S" type pitot are within given tolerances prescribed in Federal Register, Vol. 42, No. 160. Thursday, August 18, 1977.

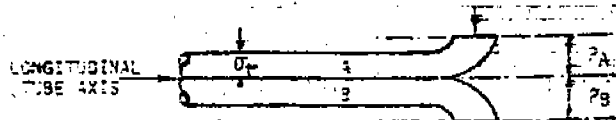
REQUIRED MEASUREMENTS

- a. External tubing diameter, D_t , $0.23 \text{ in.} < D_t < 0.38 \text{ in.}$
 b. Base to plane opening Distance, P_A and P_B , $0.40 \text{ in.} < P_A=P_B < 0.60 \text{ in.}$

$D_t = 0.375$

$P_A = 0.475$

$P_B = 0.450$



- 2) All assembly criteria to prevent aerodynamic interference for a sampling arrangement of an "S" type pitot, nozzle and thermocouple, are within given tolerances prescribed in Federal Register, Vol. 42, No. 160. Thursday, August 18, 1977.

REQUIRED MEASUREMENTS

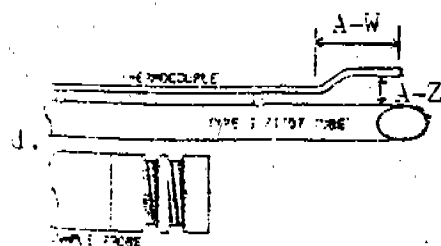
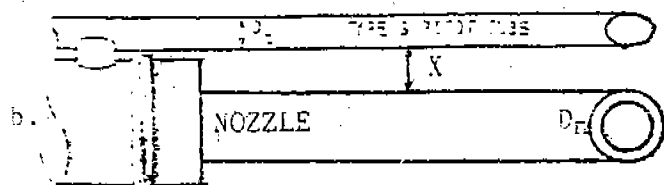
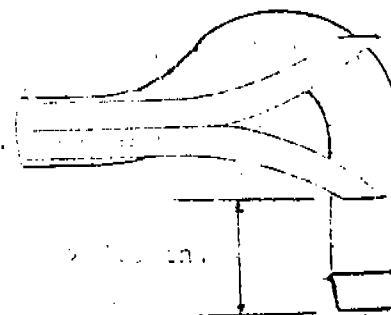
- a. External tubing diameter. See 1.1. above
 b. Pitot / nozzle separation, X , $X \geq 0.75 \text{ in.}$ for nozzle diam $D_n = 0.50 \text{ in.}$
 c. Plane of impact side of pitot in relation to plane of tube opening.
 d. Thermocouple placement, Z , $A-Z \geq 0.75 \text{ in.}$, $A-W \geq 3.0 \text{ in.}$, $B-Z \geq 1.0 \text{ in.}$
 e. Pitot / probe sheath distance, Y , $Y \geq 3.0 \text{ in.}$

$D_n = 0.1875$

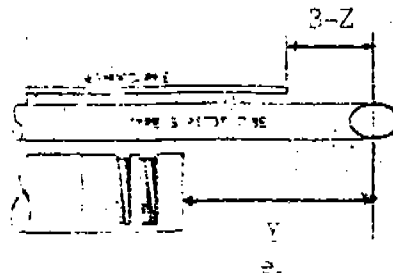
$X = 1.39$

$Z = 1.19 \text{ A-Z } A-W \geq 5.0$

$Y = 3.0$



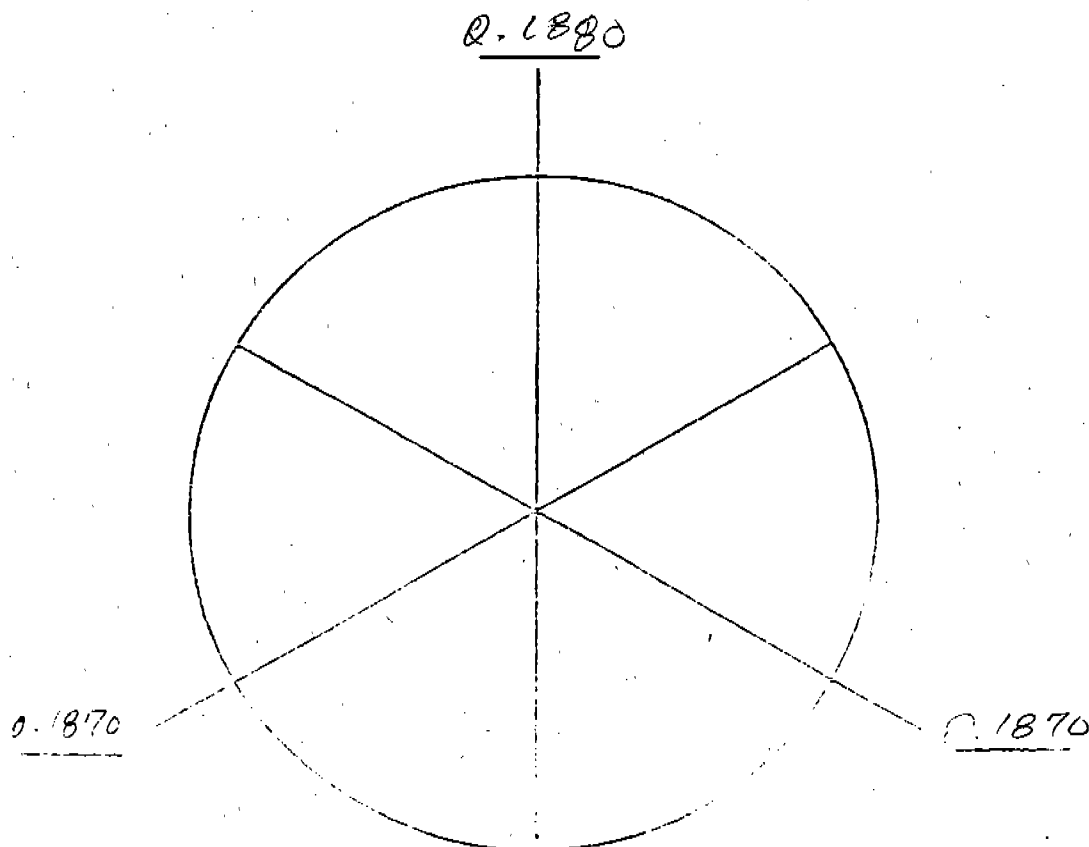
OR



NOZZLE CALIBRATION DATA FORM

Date: _____

Calibrated by: CFE



Nozzle Identification Number	Nozzle Diameter ^a			ΔD^b in (in.)	D_{avg}^c
	D_1 in (in.)	D_2 in (in.)	D_3 in (in.)		
A - 3	0.1880	0.1870	0.1870	0.001	0.1873

Where:

a = nozzle diameters b = maximum difference c = average diameter

STACK TEMPERATURE SENSOR
CALIBRATION DATA FORM

Date: _____

Thermocouple number: T2-3

Ambient temperature: 69°F

Barometric pressure: 29.80 in Hg

Calibrated by: R. STERBAS

Reference: mercury-in-glass ✓

other _____

Reference Point Number	Source ^a	Reference Thermometer Temperature		Thermocouple Potentiometer Temperature		Temperature Difference ^b %
		°C	°F	°C	°F	
1	HOT OIL	150		153		0.71
2	Boil H ₂ O	100		101		0.27
3	Amb H ₂ O	22		22		0.0
4	ICE H ₂ O	1		1		0.0

Where:

a = Type of calibration system used

b = Metric units

$$\frac{(\text{ref temp } ^\circ\text{C} + 273) - (\text{test thermom temp } ^\circ\text{C} + 273)}{\text{ref temp } ^\circ\text{C} + 273} \quad 100 \leq 1.5\%$$

b = English units

$$\frac{(\text{ref temp } ^\circ\text{F} + 460) - (\text{test thermom temp } ^\circ\text{F} + 460)}{\text{ref temp } ^\circ\text{F} + 460} \quad 100 \leq 1.5\%$$

PITOT # 29 15 A DATE

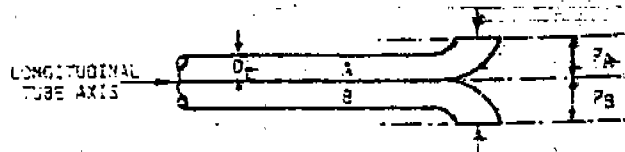
OBSERVATIONS BY C. J. [Signature]

- 1) All construction criteria for an isolated "S" type pitot are within given tolerances prescribed in Federal Register, Vol. 42, No. 160. Thursday, August 18, 1977.

REQUIRED MEASUREMENTS

- a. External tubing diameter, D_E , $0.23 \text{ in.} < D_E < 0.38 \text{ in.}$
 b. Base to plane opening Distance, P_A and P_B , $0.40 \text{ in.} < P_A=P_B < 0.60 \text{ in.}$

$D_E = 0.375$
 $P_A = 0.440$
 $P_B = 0.430$

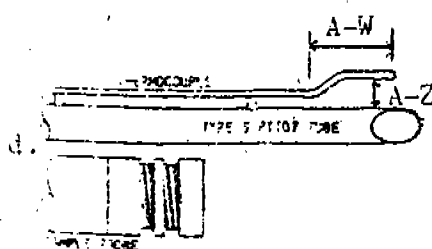
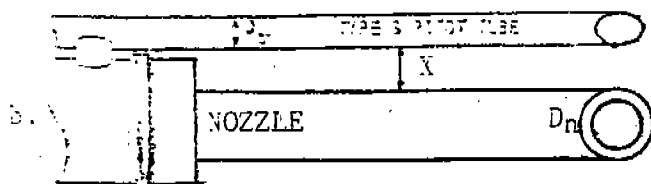
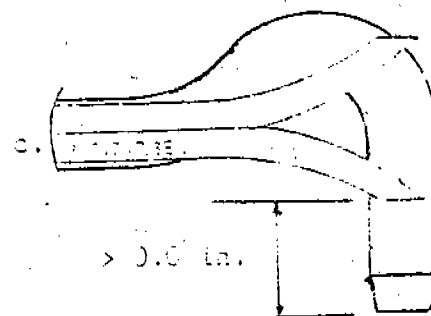


- 2) All assembly criteria to prevent aerodynamic interference for a sampling arrangement of an "S" type pitot, nozzle and thermocouple, are within given tolerances prescribed in Federal Register, Vol. 42, No. 160. Thursday, August 18, 1977.

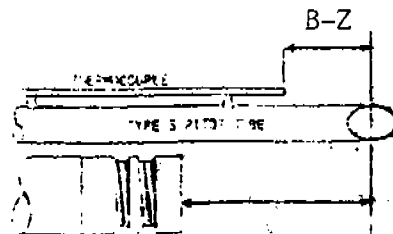
REQUIRED MEASUREMENTS

- a. External tubing diameter. See 1.a. above
 b. Pitot / nozzle separation, X , $X \geq 0.75 \text{ in.}$ for nozzle diameter, $D_n = 0.50 \text{ in.}$
 c. Plane of impact side of pitot in relation to plane of nozzle opening.
 d. Thermocouple placement, Z , $A-Z \geq 0.75 \text{ in.}$, $A-W \geq 3.0 \text{ in.}$, $B-Z \geq 2.0 \text{ in.}$
 e. Pitot / probe sheath distance, Y , $Y \geq 3.0 \text{ in.}$

$D_n = 0.1875$
 $X = 1.39$
 $Z = 1.36 \text{ A-Z, } 5.0$
 $Y = 3.0$



OR



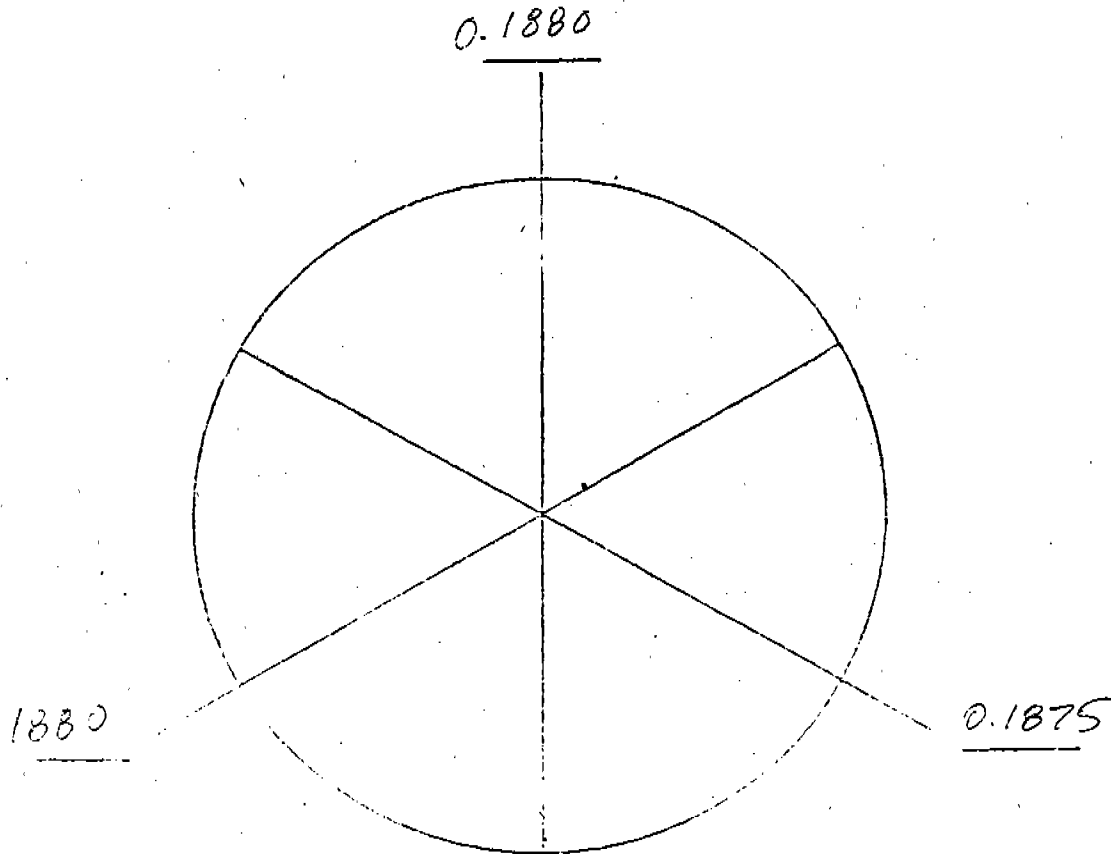
e.

NOZZLE CALIBRATION DATA FORM

Date: _____

Calibrated by: _____

[Signature]



Nozzle Identification Number	Nozzle Diameter ^a			ΔD^b in (in.)	D_{avg}^c
	D_1 in (in.)	D_2 in (in.)	D_3 in (in.)		
125-3	0.1880	0.1875	0.1880	0.0005	0.1880

Where:

a = nozzle diameters b = maximum difference c = average diameter

**STACK TEMPERATURE SENSOR
CALIBRATION DATA FORM**

Date: _____

Thermocouple number: T2-2

Ambient temperature: 69°F

Barometric pressure: 29.80 inHg

Calibrated by: R. STERGAS

Reference: mercury-in-glass ✓

other _____

Reference Point Number	Source ^a	Reference Thermometer Temperature (°C) (°F)	Thermocouple Potentiometer Temperature (°C) (°F)	Temperature Difference ^b %
1	HOT OIL	167	170	0.68
2	Boil H ₂ O	100	100	0.0
3	AMB H ₂ O	22	22	0.0
4	Ice H ₂ O	1	1	0.0

Where:

a = Type of calibration system used

b = Metric units

$$\frac{(\text{ref temp } ^\circ\text{C} + 273) - (\text{test thermom temp } ^\circ\text{C} + 273)}{\text{ref temp } ^\circ\text{C} + 273} \quad 100 \leq 1.5\%$$

b = English units

$$\frac{(\text{ref temp } ^\circ\text{F} + 460) - (\text{test thermom temp } ^\circ\text{F} + 460)}{\text{ref temp } ^\circ\text{F} + 460} \quad 100 \leq 1.5\%$$

APPENDIX D

LAB ANALYSIS SHEET - METHOD 5

Test/Run: 1-1
Unit: Alpha Mill

Date:
Location: Exhaust

Sample Identification

ID Number	Description
B 332	Glass Fiber Filter - # 47-076
B 233	Front Half Acetone Wash - 20 ml
B 234	Silica Gel

Moisture Catch Data

Impingers	Silica Gel	Net
Gross Vol: 225 ml	Gross Weight: 212.2 g	Impingers: 25
Tare Vol: 200 ml	Tare Weight: 200.0 g	Silica Gel: 12.2
Net Vol: 25 ml	Net Weight: 12.2 g	Total: 37.2 ml

Acetone Background Data

Manufacturer: Fisher Scientific
Lot Number: 902245 Density: 0.7855 g/ml
Ca = Acetone blank residue concentration (mg/ml)
Ma = Mass of residue after evaporation (mg)
Va = Volume of acetone blank (ml)
pa = Density of acetone (mg/ml)

$$Ca = Ma / [(Va)(pa)] = (0.2 \text{ g}) / (20 \text{ ml} \times 0.7855 \text{ g/ml}) = 0.00013$$

Acetone Wash Data

Wa = Weight of calculated residue in acetone wash (g)
Vaw = Volume of acetone wash (ml)

$$Wa = (Ca)(Vaw)(pa) = (0.00013 \text{ g/ml}) \times (20 \text{ ml}) \times (0.7855 \text{ g/ml}) = 0.0002 \text{ g}$$

Exhaust Mill Data

3-Stage Harber: 15-1	Gross Weight: 62.0015 g
	Tare Weight: 62.0022 g
	Less Wa: 0.0002 g
Weight of particulate in wash:	0.0021 g

Filter Data

Filter Number: 47-076	Gross Weight: 1.0253 g
	Tare Weight: 1.0244 g
Weight of particulate on filter:	0.0009 g

Particulate Weight Summary

Weight of particulate in front half wash:	0.0021 g
Weight of particulate on filter:	0.0009 g
Total weight of particulate caught:	0.0030 g

Analyst: R Stergas

Date:

LAB ANALYSIS SHEET - METHOD 5

Test/Run: 1-2
Unit: Alpha Mill

Date:
Location: Exhaust

Sample Identification

ID Number	Description
B 335	Glass Fiber Filter - # 47-077
B 336	Front Half Acetone Wash - 22 ml
B 337	Silica Gel

Moisture Catch Data

Impingers	Silica Gel	Net
Gross Vol: 238 ml	Gross Weight: 211.1 g	Impingers: 38
Tare Vol: 200 ml	Tare Weight: 200.0 g	Silica Gel: 11.1
Net Vol: 38 ml	Net Weight: 11.1 g	Total: 49.1 ml

Acetone Background Data

Manufacturer: Fisher Scientific
Lot Number: 902245 Density: 0.7855 g/ml
Ca Acetone blank residue concentration (mg/ml)
Ma Mass of residue after evaporation (mg)
Va Volume of acetone blank (ml)
pa Density of acetone (mg/ml)

$$Ca = (Ma / (Va(pa))) = (0.0002 / ((20)(0.7855))) = 0.000013$$

Acetone Blank Data

Ca Calculated residue in acetone wash (g)
Va Volume of acetone wash (ml)

$$Ca = (0.000013)(22)(0.7855)/(1000) = 0.0002$$

Front Half Wash Data

Sample Number: 15-5
Gross Weight: 63.1032 g
Tare Weight: 63.1000 g
Less Wa: 0.0002 g
Weight of particulate in wash: 0.0020 g

Filter Data

Filter Number: 47-077
Gross Weight: 1.0185 g
Tare Weight: 1.0164 g
Weight of particulate on filter: 0.0031 g

Particulate Weight Summary

Weight of particulate in front half wash: 0.0020 g
Weight of particulate on filter: 0.0031 g
Total weight of particulate catch: 0.0051 g

Analyst: R. Stergas

Date:

LAB ANALYSIS SHEET - METHOD 5

Test/Run: 1-3
Unit: Alpha Mill

Date:
Location: Exhaust

Sample Identification

ID Number	Description
B 338	Glass Fiber Filter - # 47-078
B 339	Front Half Acetone Wash - 30 ml
B 340	Silica Gel

Moisture Catch Data

Impingers	Silica Gel	Net
Gross Vol: 239 ml	Gross Weight: 212.2 g	Impingers: 39
Tare Vol: 200 ml	Tare Weight: 200.0 g	Silica Gel: 12.2
Net Vol: 39 ml	Net Weight: 12.2 g	Total: 51.2 ml

Acetone Background Data

Manufacturer: Fisher Scientific
Lot Number: 902215 Density: 0.7855 g/ml
Ca = Acetone blank residue concentration (mg/mg)
Ma = Mass of residue after evaporation (mg)
Va = Volume of acetone blank (ml)
da = Density of acetone (mg/ml)
 $Ca = Ma / [Va(da)] = (0.2) / [(20)(.7855)] = 0.000013$

Acetone Blank Data

W = Weight of calculated residue in acetone wash (g)
Vw = Volume of acetone wash (ml)
 $W = [Ca(Vw)] = (0.000013)(30)(.7855)/(1000) = 0.0003$ g

Front Half Wash Data

Beaker Number: 156
Gross Weight: 65.0512 g
Tare Weight: 65.0510 g
Less Wt: 0.0003 g
Weight of particulate in wash: 0.0020 g

Filter Data

Filter Number: 47-078
Gross Weight: 1.0172 g
Tare Weight: 1.0147 g
Weight of particulate on filter: 0.0025 g

Particulate Weight Summary

Weight of particulate in front half wash:	0.0020	g
Weight of particulate on filter:	0.0025	g
Total weight of particulate catch:	0.0045	g

Analyst: R. Stergas

Date:

SAMPLE LOG SHEET

Client: _____

Facility:

Facility: _____
Project: Small particulate emission compliance - Re-test

[illegible]

FILTER TARE WEIGHT LOG SHEET

Client: ^

Location: _____

Project: 901037

Filter Type: Glass fiber w/ springs

Size: 47 mm

Filter Number	Date/Time				
	Weight				
47 - 007	6/20/90 1000	6/21/90 0800			
	1.0376 g	1.0378 g			
47 - 064	6/20/90 1000	6/21/90 0800			
	1.0620 g	1.0618 g			
47 - 065	6/20/90 1000	6/21/90 0800			
	1.0675 g	1.0676 g			
47 - 076	6/20/90 1000	6/21/90 0800			
	1.0243 g	1.0244 g			
47 - 077	6/20/90 1000	6/21/90 0800			
	1.0163 g	1.0164 g			
47 - 078	6/20/90 1000	6/21/90 0800			
	1.0150 g	1.0147 g			
47 - 079	6/20/90 1000	6/21/90 0800			
	1.0070 g	1.0070 g			
47 - 080	6/20/90 1000	6/21/90 0800			
	1.0015 g	1.0016 g			
47 - 081	6/20/90 1000	6/21/90 0800			
	1.0111 g	1.0111 g			
47 - 082	6/20/90 1000	6/21/90 0800			
	0.9972 g	0.9973 g			

$$y_1 = \frac{1}{2} \left(\frac{1}{2} + \frac{1}{2} \right) = \frac{1}{2}$$

Facility:

Project: X Mill Compl. Retest

Sample Type: G.F. Filter 47mm w/ Teflon Rings

[illegible]

1. $\sqrt{100} = 10$ 2. $\sqrt{16} = 4$ 3. $\sqrt{25} = 5$

Facility:

1

1. *Phragmites australis* (Cav.) Trin. ex Steud.

Facility: _____

Project: 901037

Sample Type: front wash

[illegible]

Source category: **TALC** Filename: **TALC_R13.WQ1** Date: 13/

Plant name: Location: Ref. No.: 13/

Process: Test date: Process rate basis: production

Source	Type of control	Pollutant	Run No.	Test Method	Isokinetic, %	Gas volume, DSCF	Volum. flow rate, DSCFM	Mass, g	Concen., gr/DSCF	Emission rate, lb/hr	Process rate, ton/hr	Emission factor		
												kg/Mg	lb/ton	Rat.
Roller mill	fabric filter	filterable PM	1	EPA 5	94.8	67.15	9,491	0.0030	0.00069	0.056	20.00	0.0014	0.0028	
		filterable PM	2		95.5	69.30	9,643	0.0051	0.0011	0.094	20.00	0.0023	0.0047	
		filterable PM	3		97.2	68.83	9,484	0.0045	0.0010	0.082	20.00	0.0021	0.0041	
									%		Average	0.0019	0.0039	B
	CO2		1	Fyrite	NA	NA	9,491	NA	0.7	456	20.00	11	23	
	CO2		2		NA	NA	9,643	NA	0.5	331	20.00	8.3	17	
	CO2		3		NA	NA	9,484	NA	1.0	651	20.00	16	33	
											Average	12	24	C

Basis for rating: Average process rates provided; run-by-run rates not specified.

Problems noted:

Other notes:

Test conducted after repairs made to APCD, which had been malfunctioning during previous test. (See Reference 12.)