

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

The file name refers to the reference number, the AP42 chapter and section. The file name "ref02_c01s02.pdf" would mean the reference is from AP42 chapter 1 section 2. The reference may be from a previous version of the section and no longer cited. The primary source should always be checked.

Background Report Reference

AP-42 Section Number: 9.10.1.1

Background Chapter: 4

Reference Number: 2

Title: Emission Compliance Tests, Char
Dust Collector #1 and #4 ACM Mill
Exhaust, Domino Sugar Corporation,
Chalmette Refinery, Arabi, LA

Emission Testing Service

Emission Testing Service

January 1997

**DOMINO SUGAR CORPORATION
CHALMETTE REFINERY
7417 NORTH PETERS STREET
ARABI, LOUISIANA 70032**

**EMISSION COMPLIANCE TESTS
CHAR DUST COLLECTOR #1 and
#4 ACM MILL EXHAUST
JANUARY 27-28, 1997**

Testing conducted by :

E T S

EMISSION TESTING SERVICES, INC.

P.O. BOX 15075

BATON ROUGE, LOUISIANA 70895

(504) 925-8405

File : 97010



Rec'd 4/23/97
JW

State of Louisiana
Department of Environmental Quality



M.J. "MIKE" FOSTER, JR.
GOVERNOR

J. DALE GIVENS
SECRETARY

Mr. Dallas Safreit
US EPA MD14
Research Triangle Park, North Carolina 27711

RE: Compliance Test, Domino Sugar, Louisiana

Dear Mr. Safreit:

The Louisiana Department of Environmental Quality received a request from you to forward a copy of the test report from Domino Sugar to include in your study. Enclosed in that copy. If you have any questions, please feel free to contact me at 504-765-0168.

Sincerely,

Jim Courville, PE
Engineering Section

JC/ea



recycled paper

OFFICE OF AIR QUALITY P.O. BOX 82135 BATON ROUGE, LOUISIANA 70884-2135

AN EQUAL OPPORTUNITY EMPLOYER



EMISSION TEST REPORT CERTIFICATION

for

DOMINO SUGAR CORPORATION
CHALMETTE REFINERY

EMISSION COMPLIANCE TESTS

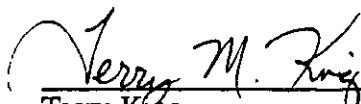
CHAR DUST COLLECTOR #1 and
#4 ACM MILL EXHAUST

JANUARY 27-28, 1997

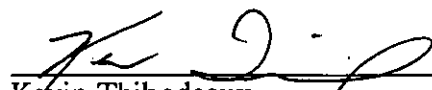
Testing conducted by :

EMISSION TESTING SERVICES, INC.

I certify that I have personally examined and am familiar with the information submitted herein, and based on my inquiries of those individuals immediately responsible for obtaining the information, I believe the submitted information is true, accurate, and complete.



Terry King
Domino Sugar Corporation
Plant Representative



Kevin Thibodeaux
Emission Testing Services, Inc.
Project Leader



Barry Gipson
Emission Testing Services, Inc.
Manager

CONTACT REPORT--MRI Project No. 4604-04

From: Brian Shrager, Environmental Engineering Department
Date of Contact: May 8, 1997
Contacted by: Telephone
Company/Agency: Domino Sugar Corporation
7417 North Peters Street
Arabi, LA 70032-1598
Telephone Number: (504) 271-5331

Person(s) Contacted/Title
Mr. Terry King

CONTACT SUMMARY: Mr. King was initially contacted in April 1997 to obtain process data for a test report that was supplied to EPA by the State of Louisiana Department of Environmental Quality. During the initial conversation with Mr. Tom Lapp of MRI, Mr. King stated that process data are not available for one of the sources, the confectioners sugar mill, that was tested. He also stated that he may be able to track down process data for the char dust collector emission test. Mr. King left a voice mail today stating that during the emission test, 15401.1 lb/hr of char went past the char dust collector. I called Mr. King back for clarification of the process and process rate, and he stated that the process rate represents the amount of char (that has come out of the kiln) transferred from a bucket elevator to a belt conveyor. The process that was tested is basically a conveyor transfer point, and does not include any kiln emissions. The belt conveyor conveys the char back to the filters that are used in decolorization.

DOMINO SUGAR CORPORATION
CHALMETTE REFINERY

EMISSION COMPLIANCE TESTS

CHAR DUST COLLECTOR #1 and
#4 ACM MILL EXHAUST

JANUARY 27-28, 1997

TABLE of CONTENTS

I.	Introduction	1.1
II.	Summary	
	Test Results Summary	2.1
	Quality Assurance/Quality Control	2.4
	Process Description	2.5
III.	Procedure	3.1
IV.	Analytical Technique	4.1
V.	Chain of Custody	5.1
VI.	Test Data and Calculations	6.0.1
	Char Dust Collector #1	
	EPA Method 1 (sample point location and traverse layout)	6.1.1
	EPA Methods 2-4 (stack gas flow rate, moisture, mole wt.)	
	Test Data & Calculations, Runs 1-3	6.1.2
	EPA Method 5 (particulate)	
	Data Summary & Calculations, Runs 1-3	6.1.11
	Laboratory Data, Runs 1-3	6.1.17
	#4 ACM Mill Exhaust	
	EPA Method 1 (sample point location and traverse layout)	6.2.1
	EPA Methods 2-4 (stack gas flow rate, moisture, mole wt.)	
	Test Data & Calculations, Runs 1-3	6.2.2
	EPA Method 5 (particulate)	
	Data Summary & Calculations, Runs 1-3	6.2.8
	Laboratory Data, Runs 1-3	6.2.14
VII.	Appendix	7.0.1
	Resumes of Test Personnel	7.1.1
	Test Equipment Calibrations	7.2.1

LIST of TABLES

I.	Emission Test Results Summary, Char Dust Collector #1	2.2
II.	Emission Test Results Summary, #4 ACM Mill Exhaust	2.3

LIST of FIGURES

I.	Emission Test Results Summary, Char Dust Collector #1	2.6
II.	Emission Test Results Summary, #4 ACM Mill Exhaust	2.7

I. INTRODUCTION

Emission Testing Services, Inc. (ETS) was contracted by Domino Sugar Corporation to conduct emission testing at Domino Sugar Corporation's Chalmette Refinery. Emissions from the Char Dust Collector #1, source ID 028-74; and #4 ACM Mill Exhaust, source ID 016-74 were tested to determine the rate of particulate emissions. Tests were conducted to demonstrate compliance with permit number 2500-00009-03 issued by the Louisiana Department of Environmental Quality (LDEQ).

Particulate emission rates were tested using EPA Method 5. EPA Methods 1 - 4 were used to determine the number and location of sample points, and the velocity, moisture, molecular weight, and volumetric flow rate of the stack gas.

The emission test team consisted of Kevin Thibodeaux, project leader, and a test crew consisting of Ryan Deaville and Jeff Savoie. Terry King represented Domino Sugar Corporation and coordinated testing with plant operations. The LDEQ was not represented on-site during the tests.

Emission tests on the Char Dust Collector #1 were conducted on January 27, 1997. Emission tests on the #4 ACM Mill Exhaust were conducted on January 28, 1997. Three, 1-hr test runs were conducted on each source.

II. SUMMARY

Emission tests were conducted on the Char Dust Collector #1 on January 27, 1997. Results of the tests indicate that particulate emissions are in compliance with permit limits. Particulate emissions averaged 2.000 lb/hr which is lower than the permit limit of 2.2 lb/hr. Test results are presented in Table I.

The #4 ACM Mill Exhaust was tested on January 28, 1997. Results of the tests indicate that particulate emissions are not in compliance with permit limits. Particulate emissions averaged 0.449 lb/hr which exceeds the permit limit of 0.04 lb/hr. A summary of test results is presented in Table II.

During tests on the #4 ACM Mill Exhaust, test personnel noted a residue buildup on the inside walls of the stack which reduced the stack ID. This did not have an affect on the particulate emission tests. A physical obstruction blocked one test port however, which resulted in a modification of the test method. Samples were collected from only one test port with samples collected from four traverse points while traversing into the stack and then from four points while traversing out of the stack.

EPA methods were used to determine all pollutant and stack gas data. All deviations from the test methods and unusual conditions have been noted above.

TABLE I EMISSION TEST RESULTS SUMMARY

DOMINO SUGAR CORPORATION CHALMETTE REFINERY

CHAR DUST COLLECTOR #1 (source ID 028-74)

<u>PARAMETER</u>	<u>Permit Limit Max./Avg.</u>	<u>RUN 1</u>	<u>RUN 2</u>	<u>RUN 3</u>	<u>AVERAGE</u>
DATE		1-27-97	1-27-97	1-27-97	
TIME		1102-1205	1241-1342	1415-1517	
PARTICULATE ⁽¹⁾					
grain/dscf		0.00533	0.02928	0.33309	0.12257
lb/hr	2.3/2.2	0.083	0.459	5.458	2.000
STACK GAS					
Oxygen, vol %		20.9	20.9	20.9	20.9
Carbon dioxide, vol %		0.0	0.0	0.0	0.0
Temperature, F		74	74	78	75
Moisture, vol %		2.24	2.23	1.78	2.08
Velocity, ft/sec		10.13	10.26	10.72	10.37
Flow rate, acfm		1869.6	1894.3	1979.0	1914.3
dscfm		1812.5	1827.1	1911.5	1850.4

- (1) Particulate samples were collected isokinetically to avoid bias due to particle size and density. The degree of which the actual sampling rate deviated from the ideal isokinetic sampling rate has been reported as a percentage of the ideal rate - 100% being ideal. The range of acceptable deviation is 90% to 110%. Sampling rates during the tests were; Run 1 - 89.43%; Run 2 - 98.92%; Run 3 - 86.57%.

TABLE II **EMISSION TEST RESULTS SUMMARY**

DOMINO SUGAR CORPORATION **CHALMETTE REFINERY**

#4 ACM MILL EXHAUST **(source ID 016-74)**

<u>PARAMETER</u>	<u>Permit Limit</u> <u>Max./Avg.</u>	<u>RUN 1</u>	<u>RUN 2</u>	<u>RUN 3</u>	<u>AVERAGE</u>
DATE		1-28-97	1-28-97	1-28-97	
TIME		0907-1007	1051-1151	1228-1328	
PARTICULATE ⁽¹⁾					
grain/dscf		0.08978	0.02165	0.01394	0.04179
lb/hr	0.04	0.949	0.242	0.156	0.449
STACK GAS					
Oxygen, vol %		20.9	20.9	20.9	20.9
Carbon dioxide, vol %		0.0	0.0	0.0	0.0
Temperature, F		121	122	118	120
Moisture, vol %		8.47	1.67	1.69	3.94
Velocity, ft/sec		41.14	40.37	40.23	40.58
Flow rate, acfm		1484.2	1456.5	1451.5	1464.1
dscfm		1233.5	1305.2	1307.9	1282.2

- (1) Particulate samples were collected isokinetically to avoid bias due to particle size and density. The degree of which the actual sampling rate deviated from the ideal isokinetic sampling rate has been reported as a percentage of the ideal rate - 100% being ideal. The range of acceptable deviation is 90% to 110%. Sampling rates during the tests were; Run 1 - 94.55%; Run 2 - 97.12%, Run 3 - 98.77%.

QUALITY ASSURANCE/ QUALITY CONTROL

Test results have been validated by following rigorous quality assurance (QA) and quality control (QC) programs. These programs ensure that the test data are representative of the source conditions by verifying the completeness, accuracy, and precision of the test data. QA/QC functions cover all aspects of the test including field testing, sampling, and analysis, sample transport and laboratory analysis, and data processing and reporting.

Emission testing at Domino Sugar Corporation included a series of QC functions. All QC functions were completed satisfactorily within the required limits.

Emission Testing Services's QA program includes participation in EPA's quarterly audits. Results of the audits, applicable to the test methods applied during testing at Domino Sugar Corporation, were within acceptable limits.

PROCESS DESCRIPTION

DOMINO SUGAR CORPORATION CHALMETTE REFINERY

Domino Sugar Corporation's Chalmette Refinery currently produces refined granulated sugar, confectionery sugar, blackstrap molasses, and several specialty products. Raw cane sugar, received by ship, barge, truck, and rail, is stored in sheds until processed. Raw sugar is refined through the processes of affination, carbonotation, press filtration, bone char decolorization, crystallization, and centrifugation. The drying operation of countercurrent flows of hot air and sugar creates dust which is recovered by cyclonic gravimetric separation and wet scrubbing.

Powdered sugars are produced by pulverizing granulated sugar along with cornstarch (or, in the case of a specialty product, malto-dextrin) in a rotary mill. The product is collected in a modified baghouse which also acts as an emissions control device.

Boilers No. 1 and 7 are natural gas fired boilers with maximum heat inputs of 78.2 MMBtu/hr and 151.3 MMBtu/hr respectively. The High Pressure Boilers, which exhaust to atmosphere through a common stack, are five identical natural gas fired boilers having a maximum heat input of 56.9 MMBtu/hr each. During normal operation, only four of the five boilers are fired.

The Char Kilns North are 18 identical kilns, each having a maximum heat input of 1.7 MMBtu/hr. During normal operation, only 14 of the 18 kilns are fired. The Char Kilns South are 6 identical kilns, each having a maximum heat input of 1.7 MMBtu/hr. During normal operation, only 4 of the 6 kilns are fired.

Following are schematics and brief process descriptions for the Char Dust Collector #1 (Figure I) and #4 ACM Mill (Figure II).

Domino Sugar Corporation Process Descriptions

Char Dust Collector #1

The char dust collectors are gravity collectors that are 80-95% efficient. They remove dust from bucket elevators utilized to transport char from the 1st floor of the filter to the 8th floor.

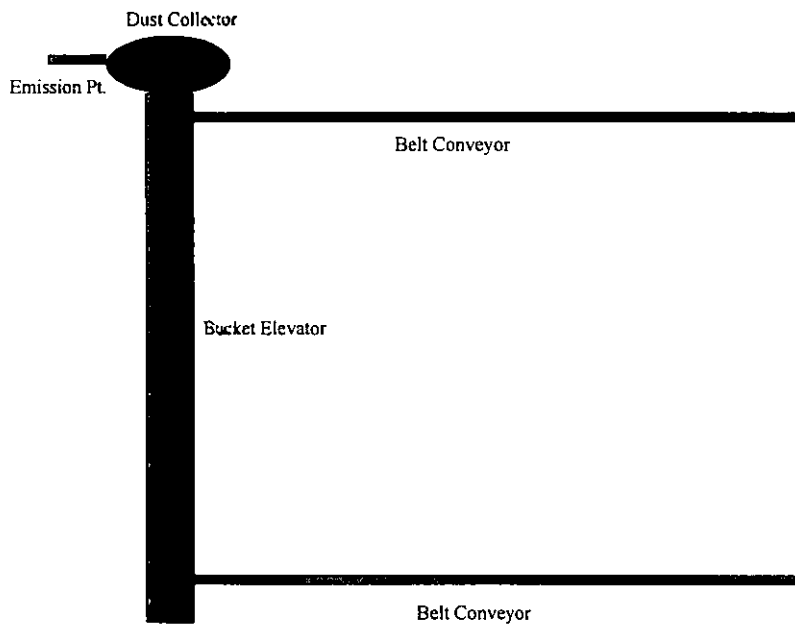


FIGURE 1

#4 ACM Mill

The ACM Mills are used to make and collect confectioner's, or powdered, sugar by employing a hammer-mill and fabric filters. The emissions are from the baghouse that collects the sugar. It is a product recovery system.

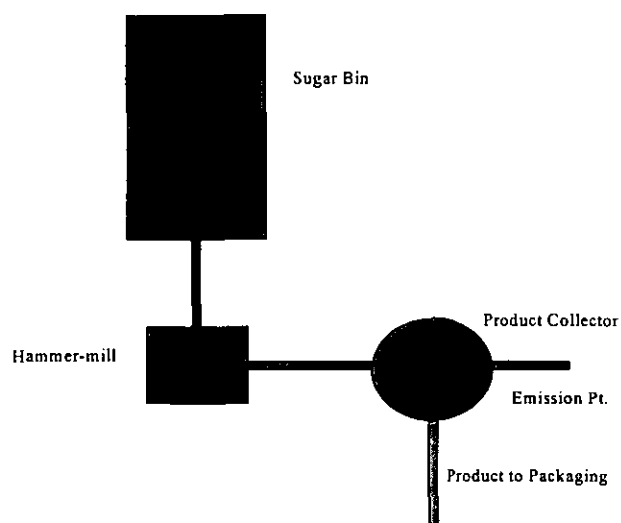


FIGURE 11

III. PROCEDURE

Emissions tests were conducted following EPA methods documented in Title 40 of the Code of Federal Regulations, Part 60, Appendix A. A brief description of each method used follows.

EPA METHOD 1:

SAMPLE and VELOCITY TRAVERSES for STATIONARY SOURCES

Method 1 is applicable to stacks having a diameter greater than 12 in. The method recommends that sampling points be located 8 diameters downstream and 2 diameters upstream from the nearest flow disturbances; however, any distances greater than the minimum 2 diameters downstream and 0.5 diameters upstream are acceptable. These distances help insure the collection of a representative sample in the absence of cyclonic flow. Cyclonic flow is detected by measuring the stack gas flow Yaw angle at each test point. The average yaw angle is required to be less than 20 degrees.

The Char Dust Collector # 1 has a stack diameter of 23.75 in. The sampling point was located 3.24 stack diameters downstream and 1.16 stack diameters upstream from the nearest flow disturbances. Based on the upstream and downstream position of the sampling plane, 24 sample points were selected and laid-out on 2 perpendicular traverses (diameters) of the stack - 12 points per traverse. Access to the sample points was obtained via 2 ports through the stack wall. Cyclonic flow was not detected.

EPA METHOD 1A:

SAMPLE and VELOCITY TRAVERSES for STATIONARY SOURCES with SMALL STACKS or DUCTS

Method 1A is applicable to stacks with a diameter greater than 4 in. and less than 12 in. The method recommends that sampling points be located 8 diameters downstream and 2 diameters upstream from the nearest flow disturbances; however, any distances greater than the minimum 2 diameters downstream and 0.5 diameters upstream are acceptable. These distances help insure the collection of a representative sample in the absence of cyclonic flow. Cyclonic flow is detected by measuring the stack gas flow Yaw angle at each test point. Average yaw angle is required to be less than 20 degrees.

The #4 ACM Mill Exhaust has a stack diameter of 10.5 in. The sampling point was located 8.19 stack diameters downstream and 3.81 stack diameters upstream from the nearest flow disturbances. Based on the upstream and downstream position of the sampling plane, 8 sample points were selected and laid-out on 2 perpendicular traverses (diameters) of the stack - 4 points per traverse. Access to the sample points was obtained via 1 ports through the stack wall. The other port was physically obstructed. Tests were conducted by sampling four points in and four points out. Cyclonic flow was not detected.

EPA METHOD 2:**DETERMINATION of STACK GAS VELOCITY and VOLUMETRIC FLOW RATE
(Type S pitot tube)**

The average gas velocity in the stack was determined from the gas molecular weight, moisture content, and the average velocity head measured using a type S pitot tube. Volumetric flow rate was determined from the velocity and stack cross-sectional area.

EPA METHOD 3:**GAS ANALYSIS FOR CARBON DIOXIDE, OXYGEN, EXCESS AIR, and DRY
MOLECULAR WEIGHT**

The dry molecular weight of the stack gas was determined using an Orsat analyzer which measured the concentration of oxygen, carbon monoxide, and carbon dioxide. The remaining gas components were assumed to be nitrogen. A gas sample was removed from the centroid of the stack using a stainless steel probe fitted with a particulate filter. The probe, sample lines, and Orsat were purged sufficiently to obtain a representative grab sample for analysis.

EPA METHOD 4:**DETERMINATION of MOISTURE CONTENT in STACK GASES**

A gas sample was extracted from the stack using a heated glass probe fitted with a particulate filter. Sample gas passed from the probe, through a series of four impingers immersed in an ice bath. The first two impingers contained known weights of water, the third was empty, and the fourth contained a known weight of silica gel. Any water vapor in the gas stream was condensed and trapped in the impingers.

EPA METHOD 5:**DETERMINATION of PARTICULATE EMISSIONS FROM STATIONARY
SOURCES**

Particulate samples were collected from the source isokinetically to insure that a representative sample of particulate was collected without bias due to particle size or density. During ideal isokinetic sampling conditions, the velocity of the gas entering the probe nozzle equals the velocity of the stack gas. The degree at which the actual sampling rate deviated from the ideal sampling rate was calculated as a percentage of the ideal isokinetic sampling rate with 100% being ideal. The range of acceptable sampling rates is 90 - 110% of the ideal isokinetic rate.

Sample gas was collected from the source using a heated probe and was transferred through a sample line to a heated glass fiber filter. The filter collected particulate and any other material which condensed at or above the filtration temperature. After filtration, the gas passed through an impinger train immersed in an ice bath. The train consisted of two impingers containing known weights of DI water, one empty impinger, and one impinger packed with a known weight of silica gel. Sample flow rate was established using a diaphragm pump and was controlled using a valve. A calibrated dry gas meter was used to determine the total gas sample volume.

After sampling, the filter was recovered from the filter holder. The front half of the filter holder and probe were washed and the washings were collected as the probe wash sample. Sample containers were labeled and sealed for shipment to the laboratory for analyses.

IV. ANALYTICAL TECHNIQUE

All sample analyses were conducted following EPA methodology. Presented below are brief descriptions of each method used.

EPA METHOD 5:

DETERMINATION of PARTICULATE EMISSIONS FROM STATIONARY SOURCES

The particulate filter was oven conditioned at 350 F and weighed before and after sampling. The filter weight gain represented the particulate catch on the filter. The probe wash sample was analyzed for total solids content by gravimetric analyses. An aliquot of the sample was placed into a tared flask and evaporated to dryness. The weight of the residue, corrected for sample volume and wash blank, represented the probe wash catch.

V. CHAIN of CUSTODY

For all samples transported to the laboratory for analyses, a Chain of Custody form was prepared during the sample recovery process. The sample ID, number of samples, liquid level when applicable, name of the sample recovery supervisor, name of the person responsible for sample recovery, and the signature of the sample trustee were recorded on the form.

During all subsequent handling, the custody form accompanied the samples. As personnel relinquished responsibility for the samples, the receiving sample trustee verified the number and integrity of the samples and signed the custody form.

Upon receipt at the analytical laboratory, the lab sample trustee verified the number, type, and integrity of the samples. These data were recorded on the custody form along with the date of receipt and lab sample trustee's signature.

At the time of analyses, the laboratory analyst recorded the analyses date and the laboratory supervisors name on the custody form. The analyst signed the custody form and attached it to the analytical results.

Chain of Custody forms for samples collected during testing are presented on the following pages along with the Analytical Request forms.

ANALYTICAL REQUEST FORM

Laboratory Receiving Samples Chemtex

Date Submitted/Submitted by: 1/30/97 / Ryan Dawitt

CLIENT Domino Sugar

Date of Collection 1/27/97

Data Number _____

Source Identification Dust Collector / ACM Vent

Parameters To Be Performed Total Solids

Sample Preservative (Media) Acetone Refrigerated ☒
 Not Refrigerated _____

Total Numbers of Samples 6

Total Numbers of Blanks 1

Other Spikes, Etc. _____

Date Analyzed 2/4/97

Analyst S.R. KOR

Report Date 2/5/97

Report Sample Data Number 37010168

Comments _____

EMISSION TESTING SERVICES, INC.
CHAIN OF CUSTODY
SAMPLE RECOVERY AND INTEGRITY DATA

CLIENT Domino Sugar SOURCE ID Dust Collector

FIELD DATA CHECKS

Field Sample Recovery Supervisor

Field Person Responsible for Recovered Samples

Kevin Philadeaux

Ryan Deerville

SAMPLE NUMBER	SAMPLE I.D.	DATE OF RECOVERY	LIQUID LEVEL MARKED	STORED IN LOCKED CONTAINER
3710171 Run-1	Probe wash w/acetone	1/27/97	✓	✓
3710172 Run-2	Probe wash w/acetone	1/27/97	✓	✓
3710173 Run-3	Probe wash w/acetone	1/27/97	✓	✓
3710174	Blank			

SIGNATURE OF FIELD SAMPLE TRUSTEE

Ryan Deerville

LABORATORY DATA CHECKS

SIGNATURE OF LAB SAMPLE TRUSTEE

[Signature]

DATE

1-31-97

DATE LAB REC'D SAMPLES

1/31/97

LAB SUPERVISOR SIGNATURE

AK Reddy

SAMPLE I.D.	SAMPLE I.D.	DATE OF ANALYSIS	LIQUID AT MARKED LEVEL	SAMPLE IDENTIFIED
1	Run 1 Probe wash	2/4/97	✓	✓
2	Run 2 Probe wash		✓	✓
3	Run 3 Probe wash		✓	✓
4	Blank		✓	✓

ANALYST SIGNATURE

URG

5.3

DATE

2/5/97

EMISSION TESTING SERVICES, INC.
CHAIN OF CUSTODY
SAMPLE RECOVERY AND INTEGRITY DATA

CLIENT Domino Sugar SOURCE ID ACM Vent

FIELD DATA CHECKS

Field Sample Recovery Supervisor

Field Person Responsible for Recovered Samples

Kevin Thibodeaux

Ryan Draville

SAMPLE NUMBER	SAMPLE I.D.	DATE OF RECOVERY	LIQUID LEVEL MARKED	STORED IN LOCKED CONTAINER
B7010168 Run - 1	Probe Wash w/Acetone	1/28/97	✓	✓
B7010169 Run - 2	Probe Wash w/Acetone	1/28/97	✓	✓
B7010170 Run - 3	Probe Wash w/Acetone	1/28/97	✓	✓

SIGNATURE OF FIELD SAMPLE TRUSTEE

Ryan Draville

LABORATORY DATA CHECKS

SIGNATURE OF LAB SAMPLE TRUSTEE

[Signature]

DATE

1-30-97 gordon

DATE LAB REC'D SAMPLES

1/30/97

LAB SUPERVISOR SIGNATURE

DR Reddy

SAMPLE I.D.	SAMPLE I.D.	DATE OF ANALYSIS	LIQUID AT MARKED LEVEL	SAMPLE IDENTIFIED
1	Run 1 Probe wash	2/4/97	✓	✓
2	Run 2 Probe wash	✓	✓	✓
3	Run 3 Probe wash	✓	✓	✓

ANALYST SIGNATURE
 FORMS CHN-CUST

[Signature]

5.4

DATE

2/5/97

**EMISSION TESTING SERVICES, INC.
CHAIN OF CUSTODY
SAMPLE RECOVERY AND INTEGRITY DATA**

CLIENT Dominio Sugar SOURCE ID Char Dust collector

FIELD DATA CHECKS

Field Sample Recovery Supervisor

Field Person Responsible for Recovered Samples

Kevin Thibodeaux

Ryan Deaville

SAMPLE NUMBER	SAMPLE I.D.	DATE OF RECOVERY	LIQUID LEVEL MARKED	STORED IN LOCKED CONTAINER
Run #1	Dust collector	1-27-97	N/A	✓
Run #2	" "	"	↓	↓
Run #3	" "	"	↓	↓

SIGNATURE OF FIELD SAMPLE TRUSTEE

[Signature]

LABORATORY DATA CHECKS

SIGNATURE OF LAB SAMPLE TRUSTEE

Fred Dowling

DATE 1-30-97

DATE LAB REC'D SAMPLES

1-30-97

LAB SUPERVISOR SIGNATURE

Fred Dowling

SAMPLE NUMBER	SAMPLE I.D.	DATE OF ANALYSIS	LIQUID AT MARKED LEVEL	SAMPLE IDENTIFIED
Run #1	3021 BCCollect	2-10-97		2-10-97 4p
2	3023	↓		↓
3	3022	↓		↓

ANALYST SIGNATURE
FORMS/CHN-CUST

Fred Dowling

DATE 2-10-97

EMISSION TESTING SERVICES, INC.
CHAIN OF CUSTODY
SAMPLE RECOVERY AND INTEGRITY DATA

CLIENT Domino Sugar SOURCE ID ACM vent

FIELD DATA CHECKS

Field Sample Recovery Supervisor

Field Person Responsible for Recovered Samples

Kevin Thibodeaux Ryan Deaville

SAMPLE NUMBER	SAMPLE I.D.	DATE OF RECOVERY	LIQUID LEVEL MARKED	STORED IN LOCKED CONTAINER
Run #1	ACM vent	1-28-97	N/A	✓
Run #2	" "	" "	↓	↓
Run #3	" "	" "	↓	↓

SIGNATURE OF FIELD SAMPLE TRUSTEE

[Signature]

LABORATORY DATA CHECKS

SIGNATURE OF LAB SAMPLE TRUSTEE

Fred Dowling

DATE

1-30-97

DATE LAB REC'D SAMPLES

1-30-97

LAB SUPERVISOR SIGNATURE

Fred Dowling

SAMPLE NUMBER	SAMPLE I.D.	DATE OF ANALYSIS	LIQUID AT MARKED LEVEL	SAMPLE IDENTIFIED
Run #	3024 ACM vent	2-10-97		124
1	3025 "	↓		↓
2	3026 "	↓		↓
3				

ANALYST SIGNATURE
 (FORMS)CHN-CUST

Fred Dowling

DATE

2-10-97

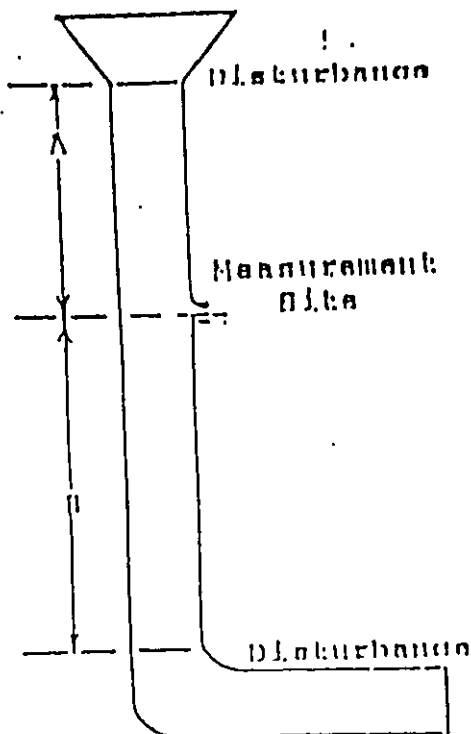
VI. TEST DATA and CALCULATIONS

All emission test field data, unit operations data, and laboratory data are presented in this section along with the associated data reduction calculations.

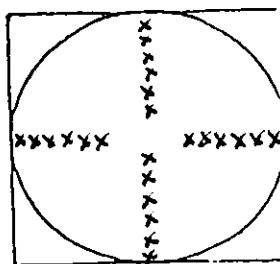
Data from the field data forms were manually input into a computer for data processing and calculations. Integrity of data input is maintained by using a double entry system. The computer performed all calculations and produced the calculations and results summary pages. Input data accuracy were further verified by comparison of raw field data to the computer data printed on the summary pages. Computer program integrity and calculations verification are validated by processing a known set of data. Further controls on the computer program are conducted by comparison of manual spot calculations of the raw data input to computer calculations results.

ETS

EMISSION TESTING SERVICES, INC. Traverse Point Layout



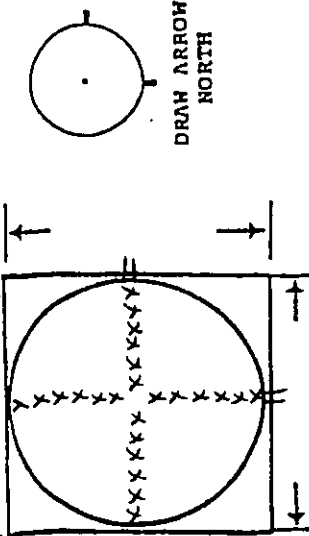
Client Domino Sugar
 Date 1/27/97
 Unit Tested CHAR Dust collector #1
 Location Arabi, NO. LA.
 Upstream Distance (A) 27.5"
 Downstream Distance (B) 77"
 Inside Diameter 23.75"
 Coupling Length 7.75"
 Coupling Type 3" male



Cross Section Schematic
of Stack

Traverse Point	Distance from Diameter	Stack Diameter	Coupling Length	Traverse Points
1	<u>0.021</u>	<u>23.75"</u> <u>6.1.1</u> <u>23.75"</u>	<u>7.75"</u>	<u>8.25</u>
2	<u>0.067</u>			<u>9.34</u>
3	<u>0.118</u>			<u>10.55</u>
4	<u>0.177</u>			<u>11.95</u>
5	<u>0.250</u>			<u>13.69</u>
6	<u>0.356</u>			<u>16.20</u>
7	<u>0.644</u>			<u>23.04</u>
8	<u>0.750</u>			<u>25.56</u>
9	<u>0.843</u>			<u>27.30</u>
10	<u>0.882</u>			<u>28.70</u>
11	<u>0.933</u>			<u>29.91</u>
12	<u>0.979</u>			<u>31.00</u>

CROSS SECTIONAL SCHEMATIC OF STACK
TRAVERSE POINT LAYOUT



ETS FIELD DATA, METHOD(S)

1-5

PLANT Dominion Sugar SAMPLE BOX NO. 85 STACK DIAMETER, in. 23.75
 DATE 1-27-97 METER BOX NO. 10 PITOT TUBE NO. P-3-1
 LOCATION Arabic, LA AMBIENT TEMPERATURE 69 PITOT FACTOR 0.834
 OPERATOR J. Saville BAROMETRIC PRESSURE 30.03 METER Δ H₀ 1.7089
 STACK NO. C.D. Dust Collector #1 FILTER TAG # 3021 D.G.M.G. FACTOR 1.0231
 RUN NO. 1 NOZZLE DIAMETER, in. 0.4095 K FACTOR 23.7774
0.42 T= 12554.5004

TRAVERSE POINT NUMBER	SAMPLING TIME (9), min.	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _s), °F	VELOCITY HEAD (ΔP _v) (1/ΔP _s)	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) (in. H ₂ O) ACTUAL DESIRED	GAS SAMPLE VOLUME (V _m), lb	GAS SAMPLE TEMPERATURE AT DRY GAS METER		SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM in. Hg gauge
1	11:02.5	-0.005	73	.056	1.200	749.200	87	89	249	50	3.5
2	11:05		75	.040	0.938	750.735	88	90	248	50	3.0
3	11:07.5		74	.035	0.822		88	90	250	49	3.0
4	11:10		77	.015	0.350	753.380	91	91	251	50	1.5
5	11:12.5		74	.020	0.470	754.280	90	90	252	51	1.5
6	11:15		74	.020	0.470	755.225	91	90	252	51	2.0
7	11:17.5		74	.025	0.587	756.175	92	91	252	52	2.0
8	11:20		75	.035	0.821	757.240	93	91	253	52	3.0
9	11:22.5		73	.035	0.824	758.485	94	90	253	52	3.0
10	11:25		74	.040	0.940	759.600	93	89	253	52	3.0
11	11:27.5		73	.040	0.940	761.120	95	90	253	53	3.0
12	11:30		73	.040	0.940	762.475	95	91	254	53	3.0
13	11:32.5					763.839					
13	11:35		75	.025	0.586	763.840	91	88	252	54	2.5
14	11:37.5		74	.020	0.470	764.940	89	86	253	54	2.0
15	11:40		76	.010	0.235	765.895	90	90	252	54	1.5
AVERAGE	11:00:00	-0.005	74		0.180	29.330		90			

COMMENTS:

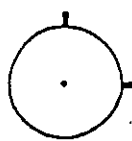
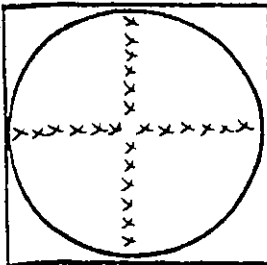
ORSAT MEASUREMENT

	CO ₂	O ₂	CO	N ₂
1	0.0	20.9		
2				
3				

VOLUME OF LIQUID WATER COLLECTED	IMPINGER WEIGHTS	
FINAL	545.3	576.3
INITIAL	541.0	573.9
WEIGHT COLLECTED	4.3	2.4

PRE-LEAK CHECK GOOD @ 0.008 CFM @ 12" H₂O
 POST-LEAK CHECK GOOD @ 0.005 CFM @ 10" H₂O

CROSS SECTIONAL SCHEMATIC OF STACK
TRAVERSE POINT LAYOUT



DRAW ARROW
NORTH

ETS FIELD DATA, METHOD(S)

1-5

PLANT Domino Sugar SAMPLE BOX NO. 85 STACK DIAMETER, in. 23.25
 DATE 1-27-97 METER BOX NO. 10 PITOT TUBE NO. P-3-1
 LOCATION Archie, LA AMBIENT TEMPERATURE 69 PITOT FACTOR 0.834
 OPERATOR J. S. Saville BAROMETRIC PRESSURE 30.03 METER Δ H. 1.7089
 STACK NO. 51 C. Dist Collector #1 FILTER TARE wt mg # 3021 D.G.M.C. FACTOR 1.0231
 RUN NO. 81 page 2 NOZZLE DIAMETER, in. 0.4099 K FACTOR 23.7774
OF 2 T → 12554.5004

TRAVERSE POINT NUMBER	SAMPLING TIME (B), min	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _s), °F	VELOCITY HEAD (ΔP _s) (1/ΔP _s)	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) in. H ₂ O ACTUAL DESIRED	GAS SAMPLE VOLUME (V _m), lb	GAS SAMPLE TEMPERATURE AT DRY GAS METER INLET (T _{inlet}), °F OUTLET (T _{outlet}), °F	SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM in. Hg gauge
16	11:42.5		74	.010	.235	766.975	88 85	254	55	1.5
17	11:45		73	.025	.588	767.255	88 86	254	56	2.5
18	11:47.5		75	.040	.938	768.315	91 86	254	56	3.0
19	11:50		76	.040	.936	769.635	92 89	254	56	3.0
20	11:52.5		73	.050	1.20	771.040	93 87	254	56	3.5
21	11:55		75	.060	1.40	772.525	93 87	253	57	4.0
22	11:57.5		74	.050	1.15	774.195	92 87	253	57	3.5
23	12:00		75	.045	1.05	775.475	92 87	253	57	3.5
24	12:02.5		75	.045	1.05	777.100	92 89	253	57	3.0
40	12:05					778.530				
AVERAGE	60 min	70.005	74	0.180	.7975	29.330			90	

COMMENTS:

PRE-LEAK CHECK GOOD @ 0.008 cfm @ 12" Hg
 POST-LEAK CHECK GOOD @ 0.005 cfm @ 10" Hg

ORIFICE MEASUREMENT

	CO ₂	O ₂	CO	N ₂
1	0.0	20.9		
2				
3				

VOLUME OF LIQUID WATER COLLECTED	IMPINGER WEIGHTS	
FINAL	545.3	576.3
INITIAL	541.0	573.9
WEIGHT COLLECTED		

DETERMINATION of STACK GAS MOISTURE, MOLECULAR WEIGHT, VELOCITY, and VOLUMETRIC FLOW RATE

A	- Cross-sectional area of the stack, ft ²	Pbar	- Barometric pressure, in. Hg	Vi	- Water vapor condensed, impinger 1, g
Acf	- Stack gas flow rate, actual, acfm	Pa	- Stack gas static pressure, in. H ₂ O	Vii	- Water vapor condensed, impinger 2, g
Bws	- Stack gas moisture fraction, %/100	Ps	- Stack gas absolute pressure, in. Hg	Viii	- Water vapor condensed, impinger 3, g
Cp	- Pitot tube coefficient	Pstd	- Standard absolute pressure, 29.92 in. Hg	Viv	- Water vapor condensed, impinger 4, g
dH	- Meter orifice pressure differential, in. H ₂ O	Qsd	- Stack gas volumetric flow rate, dry acfm	Vm	- Gas meter sample volume, dry ft ³
dP	- Stack gas velocity head, in. H ₂ O	R	- Ideal gas constant, 21.85	Vmstd	- Gas meter sample volume, dry std. ft ³
F1	- Conversion factor, 13.6 in. H ₂ O/ in. Hg	Tm	- Meter temperature, R	Vs	- Stack gas velocity, ft/sec
Kp	- Pitot tube constant, 85.49	Ts	- Stack gas temperature, R	Vw	- Total water vapor condensed, ft ³
MCO ₂	- Molecular weight of CO ₂ , 44.01 lb/ lb mole	Tstd	- Standard temperature, 528 R	Y	- Dry gas meter calibration factor
Md	- Stack gas molecular weight, dry lb/lb mole			%CO ₂	- Stack gas CO concentration, %/100
MH ₂ O	- Molecular weight of water, 18.02 lb/lb mole			%N ₂	- Stack gas N ₂ concentration, %/100
MN ₂	- Molecular weight of N ₂ , 28.01 lb/lb mole			O ₂	- Stack gas O ₂ concentration, %/100
MO ₂	- Molecular weight of O ₂ , 32.00 lb/lb mole				
Ms	- Stack gas molecular weight, wet lb/lb mole				

$$Vw = \left\{ \frac{(Vi + Vii + Viii + Viv) R Tstd}{(453.6 \text{ g/lb}) Pstd MH_2O} \right\} = \left\{ \frac{(4.3 + 2.4 + .4 + 7.0)(21.85)(528)}{(453.6)(29.92)(18.02)} \right\} = .66514$$

$$Vmstd = Vm Y \left\{ \frac{(Pb + dH/F1) Tstd}{Pstd Tm} \right\} = (29.329)(1.0231) \left\{ \frac{(30.030 + .7975/13.6)(528)}{(29.92)(550)} \right\} = 28.97078$$

$$Bws = \left\{ \frac{Vw}{Vw + Vmstd} \right\} = \left\{ \frac{(.66514)}{(.66514 + 28.97078)} \right\} = .0224$$

$$Md = (MCO_2)(\%CO_2) + (MO_2)(\%O_2) + (MN_2)(\%N_2) \\ = (44.01)(.000) + (32.00)(.209) + (28.01)(.791) = 28.844$$

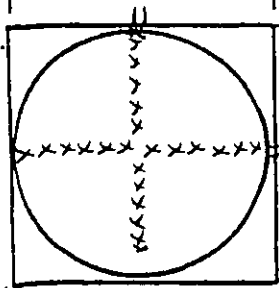
$$Ms = Md(1 - Bws) + (MH_2O Bws) = (28.844)(1 - .0224) + (18.02)(.0224) = 28.601$$

$$Vs = Kp Cp (dP)^{1/2} \left\{ \frac{Ts}{Ps Ms} \right\}^{1/4} = (85.49)(.8340)(.1801) \left\{ \frac{(534)}{(30.030)(28.601)} \right\}^{1/4} = 10.1282$$

$$Acf = (60 \text{ sec/min})(Vs A) \cdot (60)(10.1282)(3.0765) = 1869.6$$

$$Qsd = (3600 \text{ sec/hr})(1 - Bws)(Vs A) \left\{ \frac{Tstd Ps}{Ts Pstd} \right\} \\ = (3600)(1 - .0224)(10.1282)(3.0765) \left\{ \frac{(528)(30.030)}{(534)(29.92)} \right\} = 108752.8 \text{ dscf/hr} \\ 1812.5 \text{ dscf/min}$$

CROSS SECTIONAL SCHEMATIC OF STACK TRAVERSE POINT LAYOUT



ETS FIELD DATA, METHOD(S)

1-5

PLANT Dominion Sugar SAMPLE BOX NO. EE STACK DIAMETER, in. 23.75
 DATE 1-27-97 METER BOX NO. 10 PITOT TUBE NO. P-3-1
 LOCATION Arabic, LA AMBIENT TEMPERATURE 1/4 PITOT FACTOR 0.834
 OPERATOR J. SAVOIC BAROMETRIC PRESSURE 30.01 METER Δ H. 1.7089
 STACK NO. C. Dust Collector #1 FILTER 3023 D.G.M.C. FACTOR 1.0231
 RUN NO. 2 NOZZLE DIAMETER, in. 0.4099 K FACTOR 28.0942

T = 15002.3166

TRAVERSE POINT NUMBER	SAMPLING TIME (9), min.	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _s), °F	VELOCITY HEAD (ΔP), (1/ΔP)	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) in. H ₂ O		GAS SAMPLE VOLUME (V _m), ft ³	GAS SAMPLE TEMPERATURE AT DRY GAS METER		SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM in. Hg gauge
					ACTUAL DESIRED	IN. H ₂ O		INLET (T _{in}), °F	OUTLET (T _{out}), °F			
1	12:41	0.00	71	0.015	0.122	0.423	778.300	86	84	247	55	0.8
2	12:43.5		72	0.010	0.100	0.278	779.800	91	90	253	50	0.5
3	12:46		75	0.010	0.100	0.273	780.535	87	83	250	49	0.5
4	12:48.5		77	0.015	0.122	0.423	781.315	88	90	247	47	0.8
5	12:51		78	0.035	0.137	0.990	782.220	89	85	248	47	1.0
6	12:53.5		76	0.050	0.223	1.400	783.00	89	83	249	48	1.4
7	12:56		78	0.060	0.244	1.700	785.185	90	88	253	49	1.8
8	12:58.5		77	0.055	0.234	1.500	787.100	93	88	257	51	1.5
9	13:01		79	0.045	0.212	1.300	788.730	87	86	249	50	1.2
10	13:03.5		77	0.045	0.212	1.300	790.760	88	86	252	53	1.2
11	13:06		77	0.070	0.264	2.00	792.000	89	87	257	52	3.0
12	13:08.5		75	0.085	0.291	2.40	794.090	91	90	240	53	3.2
	13:10	off					796.178					
1	13:12.5	on	77	0.055	0.234	1.55	796.178	86	85	244	53	2.5
2	13:15		79	0.050	0.223	1.40	797.915	90	86	244	54	2.0
3	13:17.5		76	0.035	0.187	0.980	799.570	91	87	245	54	2.0
AVERAGE	10 min	-0.005	77		0.182	1.014	32.622		88.5			

COMMENTS:

ORFAT MEASUREMENT

	CO ₂	O ₂	CO	N ₂
1	0.0	20.9		
2				
3				

IMPINGER WEIGHTS

VOLUME OF LIQUID WATER COLLECTED	IMPINGER WEIGHTS
FINAL	574.8
INITIAL	571.8
WEIGHT COLLECTED	3.0
TOTAL WEIGHT	

PRE-LEAK CHECK GOOD @ 10" Hg. - 100 SCFH

POST-LEAK CHECK GOOD @ 0.001 CFM @ 10" Hg

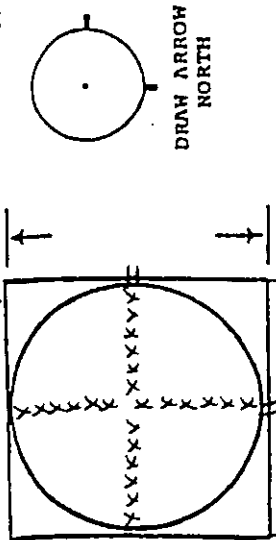
CROSS SECTIONAL SCHEMATIC OF STACK TRAVERSE POINT LAYOUT

1-5

STACK DIAMETER, IN. 23.75
 PITOT TUBE NO. P-3-1
 PITOT FACTOR 0.834
 METER Δ H. 1.7089
 D.G.M.C. FACTOR 1.0231
 K FACTOR 28.0742

PLANT Dominic Sugar DATE 1-27-97 LOCATION Arabic, LA OPERATOR J. Savio STACK NO. C. Dust Collector #1 RUN NO. 2 page 2 OF 2

SAMPLE BOX NO. EE METER BOX NO. 10 AMBIENT TEMPERATURE 71 BAROMETRIC PRESSURE 30.01 FILTER TARE w/mg # 3023 NOZZLE DIAMETER, in. 0.4099



TRAVERSE POINT NUMBER	SAMPLING TIME (hr, min)	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (°F)	VELOCITY HEAD (ΔP _s) (1/ΔP _t)	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (in. H ₂ O)	GAS SAMPLE VOLUME (Nm ³ , lb)	GAS SAMPLE TEMPERATURE AT DRY GAS METER INLET (T _{inlet}), °F OUTLET (T _{outlet}), °F	SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM in. Hg gauge
4	13:20		77	.030 .173	840 .838	801.000	89 86	245	54	2.0
5	13:22.5		75	.010 .100	280 .280	802.300	90 87	245	54	1.0
6	13:25		77	.010 .100	280 .279	803.050	90 87	247	54	1.0
7	13:27.5		78	.020 .141	520 .557	803.800	90 89	247	55	1.5
8	13:30		77	.030 .173	840 .838	804.800	90 90	248	55	2.0
9	13:32.5		77	.030 .173	840 .838	806.075	92 91	248	55	2.0
10	13:35		78	.030 .173	840 .836	807.375	92 91	249	56	2.0
11	13:37.5		78	.030 .173	840 .836	808.670	91 90	249	56	2.0
12	13:40		77	.040 .200	1,10 1,117	809.975	91 90	249	56	2.0
avg	13:42.5					811.422				
AVERAGE	13:42.5	7.005	77	.182	1,014	32.622	88.5			

COMMENTS:

OPSA? MEASUREMENT

	CO ₂	O ₂	CO	N ₂
1	0.0	20.3		
2				
3				
4				

IMPINGER
WET CENS

VOLUME OF LIQUID WATER COLLECTED	IMPINGER WEIGHTS	
FINAL	576.8	554.2
INITIAL	571.8	550.3
WEIGHT COLLECTED		

PRE-LEAK CHECK GOOD @ 0.005 cpm @ 10" Hg
POST-LEAK CHECK GOOD @ 0.001 cpm @ 10" Hg

DETERMINATION of STACK GAS MOISTURE, MOLECULAR WEIGHT, VELOCITY, and VOLUMETRIC FLOW RATE

A - Cross-sectional area of the stack, ft²
Acf - Stack gas flow rate, actual, acfm
Bws - Stack gas moisture fraction, %/100
Cp - Pitot tube coefficient
dH - Meter orifice pressure differential, in. H₂O
dP - Stack gas velocity head, in. H₂O
F1 - Conversion factor, 13.6 in. H₂O/ in. Hg
Kp - Pitot tube constant, 85.49
MCO₂ - Molecular weight of CO₂, 44.01 lb/ lb mole
Md - Stack gas molecular weight, dry lb/lb mole
MH₂O - Molecular weight of water, 18.02 lb/lb mole
MN₂ - Molecular weight of N₂, 28.01 lb/lb mole
MO₂ - Molecular weight of O₂, 32.00 lb/lb mole
Ms - Stack gas molecular weight, wet lb/lb mole

Pbar - Barometric pressure, in. Hg
Pa - Stack gas static pressure, in. H₂O
Ps - Stack gas absolute pressure, in. Hg
Pstd - Standard absolute pressure, 29.92 in. Hg
Qsd - Stack gas volumetric flow rate, dry scfm
R - Ideal gas constant, 21.85
Tm - Meter temperature, R
Ts - Stack gas temperature, R
Tstd - Standard temperature, 528 R

Vi - Water vapor condensed, impinger 1, g
Vii - Water vapor condensed, impinger 2, g
Viii - Water vapor condensed, impinger 3, g
Viv - Water vapor condensed, impinger 4, g
Vm - Gas meter sample volume, dry ft³
Vmstd - Gas meter sample volume, dry std. ft³
Vs - Stack gas velocity, ft/sec
Vw - Total water vapor condensed, ft³
Y - Dry gas meter calibration factor
%CO₂ - Stack gas CO concentration, %/100
%N₂ - Stack gas N₂ concentration, %/100
O₂ - Stack gas O₂ concentration, %/100

$$Vw = \left\{ \frac{(Vi + Vii + Viii + Viv) R Tstd}{(453.6 \text{ g/lb}) Pstd MH_2O} \right\} = \left\{ \frac{(5.0 + 3.9 + 1.4 + 5.3)(21.85)(528)}{(453.6)(29.92)(18.02)} \right\} = .73590$$

$$Vmstd = Vm Y \left\{ \frac{(Pb + dH/F1) Tstd}{Pstd Tm} \right\} = (32.622)(1.0231) \left\{ \frac{(30.010 + 1.0121/13.6)(528)}{(29.92)(549)} \right\} = 32.30350$$

$$Bws = \left\{ \frac{Vw}{Vw + Vmstd} \right\} = \left\{ \frac{(.73590)}{(.73590 + 32.30350)} \right\} = .0223$$

$$Md = (MCO_2)(\%CO_2) + (MO_2)(\%O_2) + (MN_2)(\%N_2) \\ = (44.01)(.000) + (32.00)(.209) + (28.01)(.791) = 28.844$$

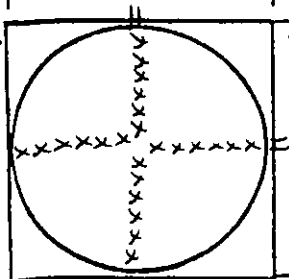
$$Ms = Md(1 - Bws) + (MH_2O Bws) = (28.844)(1 - .0223) + (18.02)(.0223) = 28.603$$

$$Vs = Kp Cp (dP)^{1/2} \left\{ \frac{Ts}{Ps Ms} \right\}^{1/2} = (85.49)(.8340)(.1820) \left\{ \frac{(537)}{(30.010)(28.603)} \right\}^{1/2} = 10.2625$$

$$Acf = (60 \text{ sec/min})(Vs A) = (60)(10.2625)(3.0765) = 1894.3$$

$$Qsd = (3600 \text{ sec/hr})(1 - Bws)(Vs A) \left\{ \frac{Tstd Ps}{Ts Pstd} \right\} \\ = (3600)(1 - .0223)(10.2625)(3.0765) \left\{ \frac{(528)(30.010)}{(537)(29.92)} \right\} = \begin{matrix} 109627.8 \text{ dscf/hr} \\ 1827.1 \text{ dscf/min} \end{matrix}$$

CROSS SECTIONAL SCHEMATIC OF STACK TRAVERSE POINT LAYOUT



DRAW ARROW NORTH

ETS FIELD DATA, METHOD(S)

PLANT Dominion Sugar SAMPLE BOX NO. 65
 DATE 1-27-97 METER BOX NO. 10
 LOCATION Archie, LA AMBIENT TEMPERATURE 72
 OPERATOR T. S. Sany BAROMETRIC PRESSURE 29.98
 STACK NO. C. Dist. Collector #1 FILTER TARE wt mg # 3022
 RUN NO. 3 page 1 of 2 NOZZLE DIAMETER, in. 0.4099

STACK DIAMETER, in. 23.75
 PITOT TUBE NO. P-3-1
 PITOT FACTOR 0.834
 METER Δ H₀ 1.7089
 D.G.M.C. FACTOR 1.0231
 K FACTOR 28.1546

T → 15119.0227

TRAVERSE POINT NUMBER	SAMPLING TIME (97) min.	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _{st}) °F	VELOCITY HEAD (DP _{st}) (1/2 DP _{st})	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) In. H ₂ O ACTUAL DESIRED	GAS SAMPLE VOLUME (V _{ml}) lb	GAS SAMPLE TEMPERATURE AT DRY GAS METER		SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM In. Hg gauge
1	14:15	-0.005	76	.020 .141	.565 .564	811.550	INLET (T _{met}) °F	OUTLET (T _{met}) °F	248	52	2.5
2	14:17.5		77	.020 .141	.565 .563	812.620	91	85	249	51	2.0
3	14:20		77	.015 .122	.420 .422	813.687	90	86	249	50	2.0
4	14:22.5		78	.010 .100	.280 .281	814.650	90	87	247	51	1.5
5	14:25		78	.010 .100	.280 .281	815.400	89	87	246	51	1.5
6	14:27.5		76	.010 .100	.280 .282	816.170	90	88	245	52	1.5
7	14:30		77	.055 .234	1.55 1.548	816.920	89	87	245	52	4.5
8	14:32.5		78	.060 .244	1.70 1.686	818.565	90	86	246	52	5.0
9	14:35		77	.075 .273	2.10 2.111	820.390	90	87	245	53	6.0
10	14:37.5		76	.070 .264	2.00 1.974	822.400	92	86	245	53	6.0
11	14:40		79	.070 .264	1.95 1.963	824.355	91	86	246	54	6.0
12	14:42.5		77	.075 .273	2.10 2.111	826.340	92	86	245	54	6.5
13	14:45					828.279					
13 cont	14:47.5		80	.055 .234	1.55 1.539	828.279	90	89	247	53	5.0
14	14:50		79	.055 .234	1.55 1.542	830.000	91	89	249	55	5.0
15	14:52.5		78	.035 .187	.985 .983	831.755	94	90	251	55	3.5
AVERAGE	10.0 min	-0.005	78	.1897	1.10	74.006	90				

COMMENTS:

ORSAT MEASUREMENT

VOLUME OF LIQUID WATER COLLECTED

IMPINGER WEIGHTS

	CO ₂	O ₂	CO	N ₂
1	0.0	20.8		
2				
3				

FINAL 548.0 578.9 471.0 702.5
 INITIAL 545.3 576.3 469.8 696.1

WEIGHT COLLECTED
 TOTAL WEIGHT

PRE-LEAK CHECK GOOD @ 0.000 CFM @ 11" H₂O
 POST-LEAK CHECK GOOD @ 0.000 CFM @ 10" H₂O

DETERMINATION of STACK GAS MOISTURE, MOLECULAR WEIGHT, VELOCITY, and VOLUMETRIC FLOW RATE

A - Cross-sectional area of the stack, ft²
 Acf - Stack gas flow rate, actual, acfm
 Bws - Stack gas moisture fraction, %/100
 Cp - Pitot tube coefficient
 dH - Meter orifice pressure differential, in. H₂O
 dP - Stack gas velocity head, in. H₂O
 F1 - Conversion factor, 13.6 in. H₂O/ in. Hg
 Kp - Pitot tube constant, 85.49
 MCO₂ - Molecular weight of CO₂, 44.01 lb/ lb mole
 Md - Stack gas molecular weight, dry lb/lb mole
 MH₂O - Molecular weight of water, 18.02 lb/lb mole
 MN₂ - Molecular weight of N₂, 28.01 lb/lb mole
 MO₂ - Molecular weight of O₂, 32.00 lb/lb mole
 Ms - Stack gas molecular weight, wet lb/lb mole

Pbar - Barometric pressure, in. Hg
 Pa - Stack gas static pressure, in. H₂O
 Ps - Stack gas absolute pressure, in. Hg
 Pstd - Standard absolute pressure, 29.92 in. Hg
 Qsd - Stack gas volumetric flow rate, dry acfm
 R - Ideal gas constant, 21.85
 Tm - Meter temperature, R
 Ts - Stack gas temperature, R
 Tstd - Standard temperature, 528 R

Vi - Water vapor condensed, impinger 1, g
 Vii - Water vapor condensed, impinger 2, g
 Viii - Water vapor condensed, impinger 3, g
 Viv - Water vapor condensed, impinger 4, g
 Vm - Gas meter sample volume, dry ft³
 Vmstd - Gas meter sample volume, dry std. ft³
 Vs - Stack gas velocity, ft/sec
 Vw - Total water vapor condensed, ft³
 Y - Dry gas meter calibration factor
 %CO₂ - Stack gas CO concentration, %/100
 %N₂ - Stack gas N₂ concentration, %/100
 O₂ - Stack gas O₂ concentration, %/100

$$Vw = \left\{ \frac{(Vi + Vii + Viii + Viv) R Tstd}{(453.6 \text{ g/lb}) Pstd MH_2O} \right\} = \left\{ \frac{(2.7 + 2.6 + 1.2 + 6.4)(21.85)(528)}{(453.6)(29.92)(18.02)} \right\} = .60854$$

$$Vmstd = Vm Y \left\{ \frac{(Pb + dH/F1) Tstd}{Pstd Tm} \right\} = (34.006)(1.0231) \left\{ \frac{(29.980 + 1.1000/13.6)(528)}{(29.92)(550)} \right\} = 33.54321$$

$$Bws = \left\{ \frac{Vw}{Vw + Vmstd} \right\} = \left\{ \frac{(.60854)}{(.60854 + 33.54321)} \right\} = .0178$$

$$Md = (MCO_2)(\%CO_2) + (MO_2)(\%O_2) + (MN_2)(\%N_2) \\ = (44.01)(.000) + (32.00)(.209) + (28.01)(.791) = 28.844$$

$$Ms = Md(1 - Bws) + (MH_2O Bws) = (28.844)(1 - .0178) + (18.02)(.0178) = 28.651$$

$$Vs = Kp Cp (dP)^{1/2} \left\{ \frac{Ts}{Ps Ms} \right\}^{1/2} = (85.49)(.8340)(.1900) \left\{ \frac{(538)}{(29.980)(28.651)} \right\}^{1/2} = 10.7211$$

$$Acf = (60 \text{ sec/min})(Vs A) = (60)(10.7211)(3.0765) = 1979.0$$

$$Qsd = (3600 \text{ sec/hr})(1 - Bws)(Vs A) \left\{ \frac{Tstd Ps}{Ts Pstd} \right\} \\ = (3600)(1 - .0178)(10.7211)(3.0765) \left\{ \frac{(528)(29.980)}{(538)(29.92)} \right\} = 114688.6 \text{ dscf/hr} \\ 1911.5 \text{ dscf/min}$$

DETERMINATION OF PARTICULATE EMISSIONS

page 1 of 2

dH - Average pressure differential, meter orifice, in-H₂O
Pbar - Barometric pressure at the sampling site, in Hg
Pstd - Standard absolute pressure, 29.92 in Hg
Tm - Average dry gas meter temperature, deg R
Tstd - Standard absolute temperature, 528 deg R
Vm - Volume of gas sample, actual dry ft³
Vmstd - Volume of gas sample, dry, Std. cond., DSCF
Y - Dry gas meter calibration factor

$$Vmstd = Vm Y \frac{Tstd}{Tm} \frac{Pbar + dH/13.6}{Pstd}$$

$$Vmstd = (29.329) (1.0231) \left(\frac{528}{550.} \right) \left(\frac{30.03 + (.7975/13.6)}{29.92} \right)$$

$$Vmstd = 28.97078$$

An - Cross-sectional area of nozzle, ft²
Ps - Absolute stack gas pressure, in Hg
Qsd - Dry volumetric stack gas flow rate, dscf/hr
St - Total sampling time, min
Ts - Absolute stack gas temperature, deg R
Vs - Average stack gas velocity, ft/sec

----- ISOKINETIC SAMPLING RATE -----

$$I = \frac{Ts Vmstd Pstd 100}{Tstd Vs An St Ps (60) (1-Bws)}$$

$$I = \frac{(534.) (28.97078) (29.92) (100)}{528 (10.1282) (.00091640) (60.) (30.030) 60 (1 - .0224)}$$

$$I = 89.43$$

DETERMINATION OF PARTICULATE EMISSIONS

page 2 of 2

Ep - Particulate emission rate, lb/hr
 Css - Concentration of particulate, grain/DSCF
 Ca -Particulate concentration, blank DI water, mg/l
 Mf -Particulate catch on filter, mg
 Mn -Total particulate collected, mg
 Mp -particulate in probe wash, mg Vp -Volume, probe wash,ml

---- Solids, probe wash ----

$$Mp = (Mp) - (Ca Vp) = (7.71) - (.000) (35.0) = 7.710$$

---- Solids, filter ----

$$Mf = 2.3$$

---- Total Solids collected ----

$$Mn = Mp + Mf$$

$$= (7.710) + (2.3) = 10.010$$

----- CONCENTRATION and EMISSION RATE -----

$$Css = (Mn/Vmstd)(0.01543 \text{ grain/mg})$$

$$= (10.010 / 28.97078) (0.01543)$$

$$= .00533$$

$$Ep = (Mn) (2.2046 \times 10^{-6} \text{ lb/mg}) (Qsd) / Vmstd$$

$$= (10.010) (2.2046 \times 10^{-6}) (108752.8) / (28.97078)$$

$$= .083$$

DETERMINATION OF PARTICULATE EMISSIONS

page 1 of 2

dH - Average pressure differential, meter orifice, in-H₂O
Pbar - Barometric pressure at the sampling site, in Hg
Pstd - Standard absolute pressure, 29.92 in Hg
Tm - Average dry gas meter temperature, deg R
Tstd - Standard absolute temperature, 528 deg R
Vm - Volume of gas sample, actual dry ft³
Vmstd - Volume of gas sample, dry, Std. cond., DSCF
Y - Dry gas meter calibration factor

$$Vmstd = Vm Y \frac{Tstd}{Tm} \frac{Pbar + dH/13.6}{Pstd}$$

$$Vmstd = (32.622) (1.0231) \left(\frac{528}{549.} \right) \left(\frac{30.01 + (1.0121/13.6)}{29.92} \right)$$

$$Vmstd = 32.30350$$

An - Cross-sectional area of nozzle, ft²
Ps - Absolute stack gas pressure, in Hg
Qsd - Dry volumetric stack gas flow rate, dscf/hr
St - Total sampling time, min
Ts - Absolute stack gas temperature, deg R
Vs - Average stack gas velocity, ft/sec

----- ISOKINETIC SAMPLING RATE -----

$$I = \frac{Ts Vmstd Pstd 100}{Tstd Vs An St Ps (60) (1 - Bws)}$$

$$I = \frac{(537.) (32.30350) (29.92) (100)}{528 (10.2625) (.00091640) (60.) (30.010) 60 (1 - .0223)}$$

$$I = 98.92$$

DETERMINATION OF PARTICULATE EMISSIONS

page 2 of 2

Ep - Particulate emission rate, lb/hr
 C_{ss} - Concentration of particulate, grain/DSCF
 Ca -Particulate concentration, blank DI water, mg/l
 M_f -Particulate catch on filter, mg
 M_n -Total particulate collected, mg
 M_p -particulate in probe wash, mg V_p -Volume, probe wash,ml

----- Solids, probe wash -----

$$M_p = (M_p) - (C_a V_p) = (39.60) - (.000)(40.0) = 39.600$$

----- Solids, filter -----

$$M_f = 21.7$$

----- Total Solids collected -----

$$\begin{aligned} M_n &= M_p + M_f \\ &= (39.600) + (21.7) = 61.300 \end{aligned}$$

----- CONCENTRATION and EMISSION RATE -----

$$\begin{aligned} C_{ss} &= (M_n / V_{mstd})(0.01543 \text{ grain/mg}) \\ &= (61.300 / 32.30350)(0.01543) \\ &= .02928 \end{aligned}$$

$$\begin{aligned} E_p &= (M_n)(2.2046 \times 10^{-6} \text{ lb/mg})(Q_{sd}) / V_{mstd} \\ &= (61.300)(2.2046 \times 10^{-6})(109627.8) / (32.30350) \\ &= .459 \end{aligned}$$

DETERMINATION OF PARTICULATE EMISSIONS

page 1 of 2

dH - Average pressure differential, meter orifice, in-H2O
Pbar - Barometric pressure at the sampling site, in Hg
Pstd - Standard absolute pressure, 29.92 in Hg
Tm - Average dry gas meter temperature, deg R
Tstd - Standard absolute temperature, 528 deg R
Vm - Volume of gas sample, actual dry ft3
Vmstd - Volume of gas sample, dry, Std. cond., DSCF
Y - Dry gas meter calibration factor

$$Vmstd = Vm Y \frac{Tstd}{Tm} \frac{Pbar + dH/13.6}{Pstd}$$

$$Vmstd = (34.006) (1.0231) \left(\frac{528}{550.} \right) \left(\frac{29.98 + (1.1000/13.6)}{29.92} \right)$$

$$Vmstd = 33.54321$$

An - Cross-sectional area of nozzle, ft2
Ps - Absolute stack gas pressure, in Hg
Qsd - Dry volumetric stack gas flow rate, dscf/hr
St - Total sampling time, min
Ts - Absolute stack gas temperature, deg R
Vs - Average stack gas velocity, ft/sec

----- ISOKINETIC SAMPLING RATE -----

$$I = \frac{Ts Vmstd Pstd 100}{Tstd Vs An St Ps (60) (1-Bws)}$$

$$I = \frac{(538.) (33.54321) (29.92) (100)}{528 (12.1625) (.00091640) (60.) (29.980) 60 (1 - .0178)}$$

$$I = 86.57$$

DETERMINATION OF PARTICULATE EMISSIONS

page 2 of 2

Ep - Particulate emission rate, lb/hr
 Css - Concentration of particulate, grain/DSCF
 Ca -Particulate concentration, blank DI water, mg/l
 Mf -Particulate catch on filter, mg
 Mn -Total particulate collected, mg
 Mp -particulate in probe wash, mg Vp -Volume, probe wash,ml

---- Solids, probe wash ----

$$Mp = (Mp) - (Ca Vp) = (298.80) - (.000) (90.0) = 298.800$$

---- Solids, filter ----

$$Mf = 425.3$$

---- Total Solids collected ----

$$\begin{aligned} Mn &= Mp + Mf \\ &= (298.800) + (425.3) = 724.100 \end{aligned}$$

----- CONCENTRATION and EMISSION RATE -----

$$\begin{aligned} Css &= (Mn/Vmstd)(0.01543 \text{ grain/mg}) \\ &= (724.100 / 33.54321) (0.01543) \\ &= .33309 \end{aligned}$$

$$\begin{aligned} Ep &= (Mn) (2.2046 \times 10^{-6} \text{ lb/mg}) (Qsd) / Vmstd \\ &= (724.100) (2.2046 \times 10^{-6}) (114688.6) / (33.54321) \\ &= 5.458 \end{aligned}$$

During sampling, particulate was collected in the probe wash and on the filter. The probe wash sample was transported to Chemtex's laboratory in Baton Rouge, Louisiana for analysis of total solids. The filter sample was transported to ETS' laboratory in Baton Rouge, Louisiana for analysis of total mass gain. Results of these analysis are presented on the following pages.

CHEMTEX

Environmental & Industrial Hygiene Services

Client: **ETS Inc.**

10461 Mammoth Dr.

Baton Rouge, LA 70814

16550 Highland Road, Baton Rouge, Louisiana 70810

(504) 752-5100 FAX (504) 753-1782

Report Date: 2/6/97

Sample Source: Impingers

Date Collected: 1/27&28/97

Collected By: KT/RD

Date Received: 1/31/97

CHEMTEX FILE #: B7020168

RESULTS OF ANALYSIS

CLIENT: DOMINO SUGAR

Source Identification: ACM Vent & C Dust Collector

<u>CHEMTEX #</u>	<u>Sample Identification</u>	<u>Parameter</u>	<u>Sample Volume ml</u>	<u>Concentration mg/l</u>	<u>Results mg/sample</u>
B7010168	Run-1 Probe Wash w/Acetone (1/28/97)	Total Solids	165	644	106.26
B7010169	Run-2 Probe Wash w/Acetone (1/28/97)	Total Solids	165	228	37.62
B7010170	Run-3 Probe Wash w/Acetone (1/28/97)	Total Solids	125	198	24.75
B7010171	Run-1 Probe Wash w/Acetone (1/27/97)	Total Solids	35	220	7.71
B7010172	Run-2 Probe Wash w/Acetone (1/27/97)	Total Solids	40	990	39.6
B7010173	Run-3 Probe Wash w/Acetone (1/27/97)	Total Solids	90	3,320	298.8
B7010174	Blank	Total Solids	---	< 2	----

ACM
Vent

CHAB
Dust
Collector

QC-Check Sample

True Value

Obtained Value

Total Solids (mg/l)

50

48/46*

* Duplicate Analysis

Parameter

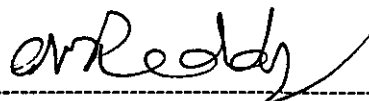
Method Reference

Date Analyzed/Analysts

Total Solids

EPA Method 160.3

2/4/97 KR



Dr. C. N. Reddy, Ph.D, CIH, ASP
Director

pap/CNR

The analytical results, opinions or interpretations contained in this report are based upon information and material supplied by the client for whose exclusive and confidential use this report has been made. The analytical results, opinions or interpretations expressed represent the best judgment of CHEMTEX. CHEMTEX, however, makes no warranty or representation, express or implied, of any type, and expressly disclaims same. This report shall not be reproduced, in whole or in part, without the written approval of CHEMTEX.

1

AIHA Accredited

Organic, Metal, Wet Chemical, Biological and Petro-
Chemical Analyses for Multi-media Environmental
and Industrial Hygiene Samples.

Facilities are also available at:

3082 25th Street, Port Arthur, Texas 77642

6.1.18

Christi, Texas 78408
Ir, Louisiana 70663

(409) 983-4575 FAX (409) 982-1522
(512) 299-9900 FAX (512) 299-1155
(318) 625-8898 FAX (318) 625-5582

ETS

EMISSION TESTING SERVICES, INC.

FILTER WEIGHTS

CLIENT Domino Sugar
UNIT ID C Dust Collector TEST DATE 1-27-97

Filter Number	Sample ID	Initial Weight		Final Weight		Weight Gained
3021	Run #1	3.8417	3.8418	3.8439	3.8441	2.3
3023	#2	3.8631	3.8634	3.8655	3.8651	21.7
3022	#3	3.8056	3.8059	4.2316	4.2312	425.3

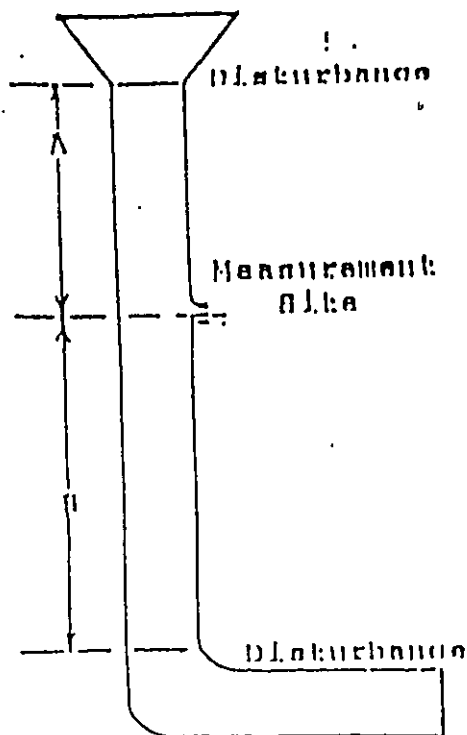
CLIENT Domino Sugar
UNIT ID ACM Vent TEST DATE 1-28-97

Filter Number	Sample ID	Initial Weight		Final Weight		Weight Gained
3024	Run 1	3.8850	3.8847	3.9703	3.9701	85.4
3025	2	3.8811	3.8813	3.8440	3.8439	12.6
3026	3	3.8272	3.8269	3.8350	3.8351	8.2

ANALYST Fred Dowling ANALYSIS DATE 2-10-97

ETS

EMISSION TESTING SERVICES, INC. Traverse Point Layout



Client Domino Sugar

Date 1-28-97

Unit Tested ACM Vent

Location Arabic, LA

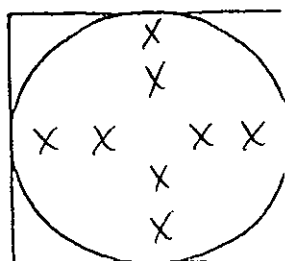
Upstream Distance (A) 40"

Downstream Distance (B) 86"

Inside Diameter 10.5 → due to build-up of materials inside pipe (Actual ~12-13")

Coupling Length 8.5"

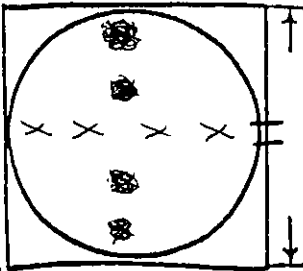
Coupling Type 3" Male



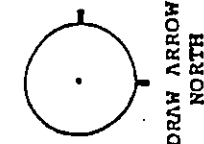
Cross Section Schematic
of Stack

Traverse Point	Distance to of Diameter	Stack Diameter	Coupling Length	Traverse Points
1	<u>.067</u>	<u>10.5</u>	<u>8.5</u>	<u>9.20</u>
2	<u>.250</u>			<u>11.13</u>
3	<u>.750</u>			<u>16.38</u>
4	<u>.933</u>			<u>18.30</u>
5				
6				
7				
8				
9				
10				
11				

CROSS SECTIONAL SCHEMATIC OF STACK
TRAVERSE POINT LAYOUT



ETS FIELD DATA, METHOD(S)



PLANT Domino Sugar
DATE 1-28-87
LOCATION Arabis, LA
OPERATOR J. Smoot
STACK NO. ACM Vent
RUN NO. 1

STACK DIAMETER, in. 10.5
PITOT TUBE NO. 0-3-1
PITOT FACTOR 0.834
METER Δ H, 1.7089
D.G.M.C. FACTOR 1.0231
K FACTOR 2.0878

T → 1202.6096

TRAVERSE POINT NUMBER	SAMPLING TIME (9), min.	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _s), °F	VELOCITY HEAD (ΔP _v) (1/ΔF)	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) in. H ₂ O ACTUAL DESIRED	GAS SAMPLE VOLUME (V _m), ft ³	GAS SAMPLE TEMPERATURE AT DRY GAS METER INLET (T _{inlet}), °F OUTLET (T _{outlet}), °F		SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM in. Hg gauge
1	09:07	7.315	118	0.520	1.10	1.081	879.200	88	87	259	2.0
2	09:14.5		122	0.530	1.10	1.095	883.340	89	88	257	2.0
3	09:22		122	0.435	0.90	0.898	888.030	89	85	258	2.0
4	09:29.5		121	0.420	0.870	0.869	892.000	87	84	259	1.5
5	09:37		123	0.490	1.05	1.052	895.925	87	84	258	1.5
6	09:44.5		123	0.510	1.05	1.052	900.035	88	84	258	2.0
7	09:52		121	0.420	0.870	0.869	904.300	88	84	260	1.5
8	09:59.5		121	0.510	1.05	1.055	908.230	88	84	259	2.0
	10:07						912.475				
AVERAGE	09:07	7.315	121	0.515	1.0925		33.275			86.5	

ORSAT MEASUREMENT

	CO ₂	O ₂	CO	N ₂
1	0.0	20.9		
2				
3				
4				

VOLUME OF LIQUID WATER COLLECTED

WEIGHT COLLECTED	IMPINGER WEIGHTS
FINAL 578.1	555.9
INITIAL 574.7	554.3
WEIGHT COLLECTED	
TOTAL WEIGHT	

COMMENTS:

PRE-LEAK CHECK GOOD @ 0.000 CFM @ 11" Hg
POST-LEAK CHECK GOOD @ 0.005 CFM @ 10" Hg
Could transcribe only one port (IN & OUT)

DETERMINATION of STACK GAS MOISTURE, MOLECULAR WEIGHT, VELOCITY, and VOLUMETRIC FLOW RATE

A - Cross-sectional area of the stack, ft²
 Acf - Stack gas flow rate, actual, acfm
 Bws - Stack gas moisture fraction, %/100
 Cp - Pitot tube coefficient
 dH - Meter orifice pressure differential, in. H₂O
 dP - Stack gas velocity head, in. H₂O
 F1 - Conversion factor, 13.6 in. H₂O/ in. Hg
 Kp - Pitot tube constant, 85.49
 MCO₂ - Molecular weight of CO₂, 44.01 lb/ lb mole
 Md - Stack gas molecular weight, dry lb/lb mole
 MH₂O - Molecular weight of water, 18.02 lb/lb mole
 MN₂ - Molecular weight of N₂, 28.01 lb/lb mole
 MO₂ - Molecular weight of O₂, 32.00 lb/lb mole
 Ms - Stack gas molecular weight, wet lb/lb mole

Pbar - Barometric pressure, in. Hg
 Pa - Stack gas static pressure, in. H₂O
 Ps - Stack gas absolute pressure, in. Hg
 Pstd - Standard absolute pressure, 29.92 in. Hg
 Qsd - Stack gas volumetric flow rate, dry acfm
 R - Ideal gas constant, 21.85
 Tm - Meter temperature, R
 Ts - Stack gas temperature, R
 Tstd - Standard temperature, 528 R

Vi - Water vapor condensed, impinger 1, g
 Vii - Water vapor condensed, impinger 2, g
 Viii - Water vapor condensed, impinger 3, g
 Viv - Water vapor condensed, impinger 4, g
 Vm - Gas meter sample volume, dry ft³
 Vmstd - Gas meter sample volume, dry std. ft³
 Vs - Stack gas velocity, ft/sec
 Vw - Total water vapor condensed, ft³
 Y - Dry gas meter calibration factor
 %CO₂ - Stack gas CO concentration, %/100
 %N₂ - Stack gas N₂ concentration, %/100
 O₂ - Stack gas O₂ concentration, %/100

$$V_w = \left\{ \frac{(V_i + V_{ii} + V_{iii} + V_{iv}) R T_{std}}{(453.6 \text{ g/lb}) P_{std} \text{ MH}_2\text{O}} \right\} = \left\{ \frac{(3.4 + 3.6 + 2.1 + 55.5)(21.85)(528)}{(453.6)(29.92)(18.02)} \right\} = 3.04739$$

$$V_{mstd} = V_m Y \left\{ \frac{(P_b + dH/F_1) T_{std}}{P_{std} T_m} \right\} = (33.275)(1.0231) \left\{ \frac{(29.890 + .9925/13.6)(528)}{(29.92)(547)} \right\} = 32.93842$$

$$B_{ws} = \left\{ \frac{V_w}{V_w + V_{mstd}} \right\} = \left\{ \frac{(3.04739)}{(3.04739 + 32.93842)} \right\} = .0847$$

$$M_d = (M_{CO_2})(\%CO_2) + (M_{O_2})(\%O_2) + (M_{N_2})(\%N_2) \\ = (44.01)(.000) + (32.00)(.209) + (28.01)(.791) = 28.844$$

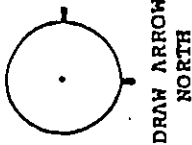
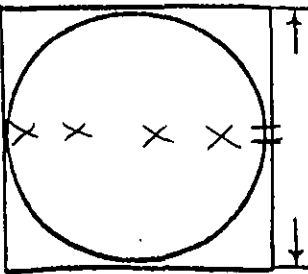
$$M_s = M_d(1 - B_{ws}) + (MH_2O B_{ws}) = (28.844)(1 - .0847) + (18.02)(.0847) = 27.927$$

$$V_s = K_p C_p (dP)^{1/2} \left\{ \frac{T_s}{P_s M_s} \right\}^{1/2} = (85.49)(.8340)(.6916) \left\{ \frac{(581)}{(29.913)(27.927)} \right\}^{1/2} = 41.1379$$

$$A_{cf} = (60 \text{ sec/min})(V_s A) = (60)(41.1379)(.6013) = 1484.2$$

$$Q_{sd} = (3600 \text{ sec/hr})(1 - B_{ws})(V_s A) \left\{ \frac{T_{std} P_s}{T_s P_{std}} \right\} \\ = (3600)(1 - .0847)(41.1379)(.6013) \left\{ \frac{(528)(29.913)}{(581)(29.92)} \right\} = \begin{matrix} 74011.7 \text{ dscf/hr} \\ 1233.5 \text{ dscf/min} \end{matrix}$$

CROSS SECTIONAL SCHEMATIC OF STACK
TRAVERSE POINT LAYOUT



ETS FIELD DATA, METHOD(S)

1-5

PLANT Dominion Sugar SAMPLE BOX NO. 85 STACK DIAMETER, in. 10.5
 DATE 1-28-97 METER BOX NO. 10 PITOT TUBE NO. 1-3-1
 LOCATION Arden, LA AMBIENT TEMPERATURE 79 PITOT FACTOR 0.834
 OPERATOR S. Savore BAROMETRIC PRESSURE 30.03 METER Δ H₀ 1.7089
 STACK NO. ACM Vent FILTER TARE wt mg #3025 D.G.M.C. FACTOR 1.0231
 RUN NO. 2 NOZZLE DIAMETER, in. .2278 K FACTOR 2.4925
 T → 1448.1499

TRAVERSE POINT NUMBER	SAMPLING TIME (9), min.	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _s), °F	VELOCITY HEAD (ΔP ₁) (1/ΔP ₂)	PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) in. H ₂ O ACTUAL DESIRED	GAS SAMPLE VOLUME (V _m), ft ³	GAS SAMPLE TEMPERATURE AT DRY GAS METER		SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM in. Hg gauge
1	10:51	4.265	122	.425	1.05	916.600	86	84	254	50	2.5
2	10:58.5		122	.430	1.05	920.895	90	86	255	49	2.5
3	11:06		124	.525	1.30	925.175	91	86	258	51	3.0
4	11:13.5		122	.540	1.35	929.955	93	87	259	52	3.0
5	11:21		121	.450	1.10	934.800	92	86	257	54	3.0
6	11:28.5		123	.535	1.35	939.255	92	86	258	55	3.0
7	11:36		121	.455	1.15	943.700	92	87	257	56	2.5
8	11:43.5		120	.445	1.10	948.280	93	86	259	61	2.5
9	11:51					952.713					
AVERAGE	60 min	4.265	122		1.088	36.113				89	

VOLUME OF LIQUID WATER COLLECTED		IMPINGER WEIGHTS	
FINAL	549.3	581.3	472.3
INITIAL	548.0	579.0	471.0
WEIGHT COLLECTED			
TOTAL WEIGHT			

ORSAT MEASUREMENT

	CO ₂	O ₂	CO	N ₂
1	0.0	20.9		
2				
3				
4				

COMMENTS:

PRE-LEAK CHECK GOOD @ 0.010 CFM @ 11" Hg
 POST-LEAK CHECK GOOD @ 0.008 CFM @ 11" Hg
 Could traverse only one port (Ex 5 Out)

DETERMINATION of STACK GAS MOISTURE, MOLECULAR WEIGHT, VELOCITY, and VOLUMETRIC FLOW RATE

A - Cross-sectional area of the stack, ft²
 Acf - Stack gas flow rate, actual, acfm
 Bws - Stack gas moisture fraction, %/100
 Cp - Pitot tube coefficient
 dH - Meter orifice pressure differential, in. H₂O
 dP - Stack gas velocity head, in. H₂O
 F1 - Conversion factor, 13.6 in. H₂O/in. Hg
 Kp - Pitot tube constant, 85.49
 MCO₂ - Molecular weight of CO₂, 44.01 lb/lb mole
 Md - Stack gas molecular weight, dry lb/lb mole
 MH₂O - Molecular weight of water, 18.02 lb/lb mole
 MN₂ - Molecular weight of N₂, 28.01 lb/lb mole
 MO₂ - Molecular weight of O₂, 32.00 lb/lb mole
 Ms - Stack gas molecular weight, wet lb/lb mole

Pbar - Barometric pressure, in. Hg
 Pa - Stack gas static pressure, in. H₂O
 Ps - Stack gas absolute pressure, in. Hg
 Pstd - Standard absolute pressure, 29.92 in. Hg
 Qsd - Stack gas volumetric flow rate, dry acfm
 R - Ideal gas constant, 21.85
 Tm - Meter temperature, R
 Ts - Stack gas temperature, R
 Tstd - Standard temperature, 528 R

Vi - Water vapor condensed, impinger 1, g
 Vii - Water vapor condensed, impinger 2, g
 Viii - Water vapor condensed, impinger 3, g
 Viv - Water vapor condensed, impinger 4, g
 Vm - Gas meter sample volume, dry ft³
 Vmstd - Gas meter sample volume, dry std. ft³
 Vs - Stack gas velocity, ft/sec
 Vw - Total water vapor condensed, ft³
 Y - Dry gas meter calibration factor
 %CO₂ - Stack gas CO concentration, %/100
 %N₂ - Stack gas N₂ concentration, %/100
 O₂ - Stack gas O₂ concentration, %/100

$$Vw = \left\{ \frac{(Vi + Vii + Viii + Viv) R Tstd}{(453.6 \text{ g/lb}) Pstd MH_2O} \right\} = \left\{ \frac{(1.3 + 2.3 + 1.3 + 8.0)(21.85)(528)}{(453.6)(29.92)(18.02)} \right\} = .60853$$

$$Vmstd = Vm Y \left\{ \frac{(Pb + dH/F1) Tstd}{Pstd Tm} \right\} = (36.113)(1.0231) \left\{ \frac{(30.030 + 1.1812/13.6)(528)}{(29.92)(549)} \right\} = 35.79628$$

$$Bws = \left\{ \frac{Vw}{Vw + Vmstd} \right\} = \left\{ \frac{(.60853)}{(.60853 + 35.79628)} \right\} = .0167$$

$$Md = (MCO_2)(\%CO_2) + (MO_2)(\%O_2) + (MN_2)(\%N_2) \\ = (44.01)(.000) + (32.00)(.209) + (28.01)(.791) = 28.844$$

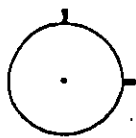
$$Ms = Md(1 - Bws) + (MH_2O Bws) = (28.844)(1 - .0167) + (18.02)(.0167) = 28.663$$

$$Vs = Kp Cp (dP)^{1/2} \left\{ \frac{Ts}{Ps Ms} \right\}^{1/2} = (85.49)(.8340)(.6889) \left\{ \frac{(582)}{(30.049)(28.663)} \right\}^{1/2} = 40.3696$$

$$Acf = (60 \text{ sec/min})(Vs A) = (60)(40.3696)(.6013) = 1456.5$$

$$Qsd = (3600 \text{ sec/hr})(1 - Bws)(Vs A) \left\{ \frac{Tstd Ps}{Ts Pstd} \right\} \\ = (3600)(1 - .0167)(40.3696)(.6013) \left\{ \frac{(528)(30.049)}{(582)(29.92)} \right\} = \begin{matrix} 78310.7 \text{ dscf/hr} \\ 1305.2 \text{ dscf/min} \end{matrix}$$

CROSS SECTIONAL SCHEMATIC OF STACK
TRAVERSE POINT LAYOUT



**DRAW ARROW
NORTH**

PLANT Domino Sugar
DATE 1-28-97
LOCATION Archie, LA
OPERATOR J. Sanoie
STACK NO. ACM Vent
RUN NO. 3

SAMPLE BOX NO. EE
METER BOX NO. 10
AMBIENT TEMPERATURE 69
BAROMETRIC PRESSURE 30.02
FILTER TARE wt mg # 3026
NOZZLE DIAMETER in. 0.2278

STACK DIAMETER, IN. 10.5
 PITOT TUBE NO. P-3-1
 PITOT FACTOR 0.834
 METER Δ H. 1.7089
 D.G.M.C. FACTOR 1.0231
 K FACTOR 2.4908

$T \rightarrow 1449.6812$

TRAVERSE POINT NUMBER	SAMPLING TIME (B), min.	STATIC PRESSURE (in. H ₂ O)	STACK TEMP (T _s), °F	VELOCITY HEAD (ΔP _s) (1/ΔP _s)		PRESSURE DIFFERENTIAL ACROSS ORIFICE METER (ΔH) In. H ₂ O	GAS SAMPLE VOLUME (V _m), ft ³	GAS SAMPLE TEMPERATURE AT DRY GAS METER		SAMPLE BOX TEMPERATURE °F	TEMPERATURE OF GAS LEAVING CONDENSER OR LAST IMPINGER °F	PUMP VACUUM In. Hg gauge	
				ACTUAL	DESIR ^d			INLET (T _{in}), °F	OUTLET (T _{out}), °F				
1	12:28	+280	121	.410	.640	1.05	1.023	952.900	86	86	249	55	2.0
2	12:35.5		120	.545	.738	1.35	1.362	957.195	87	84	252	55	2.0
3	12:43		118	.465	.681	1.15	1.160	962.110	87	83	251	56	2.0
4	12:50.5		119	.545	.738	1.35	1.364	966.875	87	82	251	56	2.0
5	12:58		117	.475	.689	1.20	1.193	971.445	86	82	250	57	2.0
6	13:05.5		116	.465	.681	1.15	1.170	976.050	86	82	253	58	2.0
7	13:13		118	.465	.681	1.15	1.160	980.530	86	83	254	58	2.0
8	13:20.5		118	.430	.655	1.10	1.078	985.000	88	83	255	59	2.0
460	13:28							989.468					
AVERAGE	12:28	+280	118		.688	1.19		36.568					

COMMENTS:

ORSAT MEASUREMENT				
	CO ₂	O ₂	CO	N ₂
1	0.0	20.9		
2				
3				

VOLUME OF LIQUID WATER COLLECTED	IMPINGER WEIGHTS	
.FINAL	579.7	633.5
INITIAL	578.1	625.7
WEIGHT COLLECTED		

PRE-LEAK CHECK GOOD @ 0.005 cfm @ 10" Hg
POST-LEAK CHECK GOOD @ 0.005 cfm @ 10" Hg
Could traverse only one part (In + Out)

DETERMINATION of STACK GAS MOISTURE, MOLECULAR WEIGHT, VELOCITY, and VOLUMETRIC FLOW RATE

A	- Cross-sectional area of the stack, ft ²	Pbar	- Barometric pressure, in. Hg	Vi	- Water vapor condensed, impinger 1, g
Acf	- Stack gas flow rate, actual, acfm	Pa	- Stack gas static pressure, in. H ₂ O	Vii	- Water vapor condensed, impinger 2, g
Bws	- Stack gas moisture fraction, %/100	Ps	- Stack gas absolute pressure, in. Hg	Viii	- Water vapor condensed, impinger 3, g
Cp	- Pitot tube coefficient	Pstd	- Standard absolute pressure, 29.92 in. Hg	Viv	- Water vapor condensed, impinger 4, g
dH	- Meter orifice pressure differential, in. H ₂ O	Qsd	- Stack gas volumetric flow rate, dry scfm	Vm	- Gas meter sample volume, dry ft ³
dP	- Stack gas velocity head, in. H ₂ O	R	- Ideal gas constant, 21.85	Vmstd	- Gas meter sample volume, dry std. ft ³
F1	- Conversion factor, 13.6 in. H ₂ O/ in. Hg	Tm	- Meter temperature, R	Vs	- Stack gas velocity, ft/sec
Kp	- Pitot tube constant, 85.49	Ts	- Stack gas temperature, R	Vw	- Total water vapor condensed, ft ³
MCO ₂	- Molecular weight of CO ₂ , 44.01 lb/ lb mole	Tstd	- Standard temperature, 528 R	Y	- Dry gas meter calibration factor
Md	- Stack gas molecular weight, dry lb/lb mole			%CO ₂	- Stack gas CO concentration, %/100
MH ₂ O	- Molecular weight of water, 18.02 lb/lb mole			%N ₂	- Stack gas N ₂ concentration, %/100
MN ₂	- Molecular weight of N ₂ , 28.01 lb/lb mole			O ₂	- Stack gas O ₂ concentration, %/100
MO ₂	- Molecular weight of O ₂ , 32.00 lb/lb mole				
Ms	- Stack gas molecular weight, wet lb/lb mole				

$$Vw = \left\{ \frac{(Vi + Vii + Viii + Viv) R Tstd}{(453.6 \text{ g/lb}) Pstd MH_2O} \right\} = \left\{ \frac{(1.6 + 2.7 + 1.2 + 7.8)(21.85)(528)}{(453.6)(29.92)(18.02)} \right\} = .62740$$

$$Vmstd = Vm Y \left\{ \frac{(Pb + dH/F1) Tstd}{Pstd Tm} \right\} = (36.568)(1.0231) \left\{ \frac{(30.020 + 1.1875/13.6)(528)}{(29.92)(545)} \right\} = 36.48100$$

$$Bws = \left\{ \frac{Vw}{Vw + Vmstd} \right\} = \left\{ \frac{(.62740)}{(.62740 + 36.48100)} \right\} = .0169$$

$$Md = (MCO_2)(\%CO_2) + (MO_2)(\%O_2) + (MN_2)(\%N_2) \\ = (44.01)(.000) + (32.00)(.209) + (28.01)(.791) = 28.844$$

$$Ms = Md(1 - Bws) + (MH_2O Bws) = (28.844)(1 - .0169) + (18.02)(.0169) = 28.661$$

$$Vs = Kp Cp (dP)^{1/2} \left\{ \frac{Ts}{Ps Ms} \right\}^{1/2} = (85.49)(.8340)(.6884) \left\{ \frac{(578)}{(30.041)(28.661)} \right\}^{1/2} = 40.2299$$

$$Acf = (60 \text{ sec/min})(Vs A) = (60)(40.2299)(.6013) = 1451.5$$

$$Qsd = (3600 \text{ sec/hr})(1 - Bws)(Vs A) \left\{ \frac{Tstd Ps}{Ts Pstd} \right\} \\ = (3600)(1 - .0169)(40.2299)(.6013) \left\{ \frac{(528)(30.041)}{(578)(29.92)} \right\} = 78473.5 \text{ dscf/hr} \\ 1307.9 \text{ dscf/min}$$

DETERMINATION OF PARTICULATE EMISSIONS

page 1 of 2

dH - Average pressure differential, meter orifice, in-H₂O
Pbar - Barometric pressure at the sampling site, in Hg
Pstd - Standard absolute pressure, 29.92 in Hg
Tm - Average dry gas meter temperature, deg R
Tstd - Standard absolute temperature, 528 deg R
Vm - Volume of gas sample, actual dry ft³
Vmstd - Volume of gas sample, dry, Std. cond., DSCF
Y - Dry gas meter calibration factor

$$Vmstd = Vm Y \frac{Tstd}{Tm} \frac{Pbar + dH/13.6}{Pstd}$$

$$Vmstd = (33.275) (1.0231) \left(\frac{528}{547.} \right) \left(\frac{29.89 + (.9925/13.6)}{29.92} \right)$$

$$Vmstd = 32.93842$$

An - Cross-sectional area of nozzle, ft²
Ps - Absolute stack gas pressure, in Hg
Qsd - Dry volumetric stack gas flow rate, dscf/hr
St - Total sampling time, min
Ts - Absolute stack gas temperature, deg R
Vs - Average stack gas velocity, ft/sec

----- ISOKINETIC SAMPLING RATE -----

$$I = \frac{Ts Vmstd Pstd 100}{Tstd Vs An St Ps (60) (1-Bws)}$$

$$I = \frac{(581.) (32.93842) (29.92) (100)}{528 (41.1379) (.00028303) (60.) (29.913) 60 (1 - .0847)}$$

$$I = 94.55$$

DETERMINATION OF PARTICULATE EMISSIONS

page 2 of 2

Ep - Particulate emission rate, lb/hr
 Css - Concentration of particulate, grain/DSCF
 Ca -Particulate concentration, blank DI water, mg/l
 Mf -Particulate catch on filter, mg
 Mn -Total particulate collected, mg
 Mp -particulate in probe wash, mg Vp -Volume, probe wash,ml

----- Solids, probe wash -----

$$Mp = (Mp) - (Ca Vp) = (106.26) - (.000) (165.0) = 106.260$$

----- Solids, filter -----

$$Mf = 85.4$$

----- Total Solids collected -----

$$Mn = Mp + Mf$$

$$= (106.260) + (85.4) = 191.660$$

----- CONCENTRATION and EMISSION RATE -----

$$Css = (Mn / Vmstd) (0.01543 \text{ grain/mg})$$

$$= (191.660 / 32.93842) (0.01543)$$

$$= .08978$$

$$Ep = (Mn) (2.2046 \times 10^{-6} \text{ lb/mg}) (Qsd) / Vmstd$$

$$= (191.660) (2.2046 \times 10^{-6}) (74011.7) / (32.93842)$$

$$= .949$$

DETERMINATION OF PARTICULATE EMISSIONS

page 1 of 2

dH - Average pressure differential, meter orifice, in-H2O
Pbar - Barometric pressure at the sampling site, in Hg
Pstd - Standard absolute pressure, 29.92 in Hg
Tm - Average dry gas meter temperature, deg R
Tstd - Standard absolute temperature, 528 deg R
Vm - Volume of gas sample, actual dry ft3
Vmstd - Volume of gas sample, dry, Std. cond., DSCF
Y - Dry gas meter calibration factor

$$Vmstd = Vm Y \frac{Tstd}{Tm} \frac{Pbar + dH/13.6}{Pstd}$$

$$Vmstd = (36.113) (1.0231) \left(\frac{528}{549.} \right) \left(\frac{30.03 + (1.1812/13.6)}{29.92} \right)$$

$$Vmstd = 35.79629$$

An - Cross-sectional area of nozzle, ft2
Ps - Absolute stack gas pressure, in Hg
Qsd - Dry volumetric stack gas flow rate, dscf/hr
St - Total sampling time, min
Ts - Absolute stack gas temperature, deg R
Vs - Average stack gas velocity, ft/sec

----- ISOKINETIC SAMPLING RATE -----

$$I = \frac{Ts Vmstd Pstd 100}{Tstd Vs An St Ps (60) (1-Bws)}$$

$$I = \frac{(582.) (35.79629) (29.92) (100)}{528 (40.3696) (.00028303) (60.) (30.049) 60 (1 - .0167)}$$

$$I = 97.12$$

DETERMINATION OF PARTICULATE EMISSIONS

page 2 of 2

Ep - Particulate emission rate, lb/hr
C_{ss} - Concentration of particulate, grain/DSCF
Ca -Particulate concentration, blank DI water, mg/l
M_f -Particulate catch on filter, mg
M_n -Total particulate collected, mg
M_p -particulate in probe wash, mg V_p -Volume, probe wash,ml

---- Solids, probe wash ----

$$M_p = (M_p) - (C_a V_p) = (37.62) - (.000)(165.0) = 37.620$$

---- Solids, filter ----

$$M_f = 12.6$$

---- Total Solids collected ----

$$M_n = M_p + M_f$$

$$= (37.620) + (12.6) = 50.220$$

----- CONCENTRATION and EMISSION RATE -----

$$C_{ss} = (M_n / V_{mstd})(0.01543 \text{ grain/mg})$$

$$= (50.220 / 35.79629)(0.01543)$$

$$= .02165$$

$$E_p = (M_n)(2.2046 \times 10^{-6} \text{ lb/mg})(Q_{sd}) / V_{mstd}$$

$$= (50.220)(2.2046 \times 10^{-6})(78310.7) / (35.79629)$$

$$= .242$$

DETERMINATION OF PARTICULATE EMISSIONS

page 1 of 2

dH - Average pressure differential, meter orifice, in-H2O
Pbar - Barometric pressure at the sampling site, in Hg
Pstd - Standard absolute pressure, 29.92 in Hg
Tm - Average dry gas meter temperature, deg R
Tstd - Standard absolute temperature, 528 deg R
Vm - Volume of gas sample, actual dry ft3
Vmstd - Volume of gas sample, dry, Std. cond., DSCF
Y - Dry gas meter calibration factor

$$Vmstd = Vm Y \frac{Tstd}{Tm} \frac{Pbar + dH/13.6}{Pstd}$$

$$Vmstd = (36.568) (1.0231) \left(\frac{528}{545.} \right) \left(\frac{30.02 + (1.1875/13.6)}{29.92} \right)$$

$$Vmstd = 36.48100$$

An - Cross-sectional area of nozzle, ft2
Ps - Absolute stack gas pressure, in Hg
Qsd - Dry volumetric stack gas flow rate, dscf/hr
St - Total sampling time, min
Ts - Absolute stack gas temperature, deg R
Vs - Average stack gas velocity, ft/sec

----- ISOKINETIC SAMPLING RATE -----

$$I = \frac{Ts Vmstd Pstd 100}{Tstd Vs An St Ps (60) (1-Bws)}$$

$$I = \frac{(578.) (36.48100) (29.92) (100)}{528 (40.2299) (.00028303) (60.) (30.041) 60 (1 - .0169)}$$

$$I = 98.77$$

DETERMINATION OF PARTICULATE EMISSIONS

page 2 of 2

Ep - Particulate emission rate, lb/hr
 Css - Concentration of particulate, grain/DSCF
 Ca -Particulate concentration, blank DI water, mg/l
 Mf -Particulate catch on filter, mg
 Mn -Total particulate collected, mg
 Mp -particulate in probe wash, mg Vp -Volume, probe wash,ml

----- Solids, probe wash -----

$$Mp = (Mp) - (Ca Vp) = (24.75) - (.000) (125.0) = 24.750$$

----- Solids, filter -----

$$Mf = 8.2$$

----- Total Solids collected -----

$$Mn = Mp + Mf$$

$$= (24.750) + (8.2) = 32.950$$

----- CONCENTRATION and EMISSION RATE -----

$$Css = (Mn/Vmstd)(0.01543 \text{ grain/mg})$$

$$= (32.950 / 36.48100) (0.01543)$$

$$= .01394$$

$$Ep = (Mn) (2.2046 \times 10^{-6} \text{ lb/mg}) (Qsd) / Vmstd$$

$$= (32.950) (2.2046 \times 10^{-6}) (78473.5) / (36.48100)$$

$$= .156$$

During sampling, particulate was collected in the probe wash and on the filter. The probe wash sample was transported to Chemtex's laboratory in Baton Rouge, Louisiana for analysis of total solids. The filter sample was transported to ETS' laboratory in Baton Rouge, Louisiana for analysis of total mass gain. Results of these analysis are presented on the following pages.

CHEMTEX

16550 Highland Road, Baton Rouge, Louisiana 70810
(504) 752-5100 FAX (504) 753-1782

Environmental & Industrial Hygiene Services

Client: **ETS Inc.**

10461 Mammoth Dr.
Baton Rouge, LA 70814

Report Date: 2/6/97
Sample Source: Impingers
Date Collected: 1/27&28/97
Collected By: KT/RD
Date Received: 1/31/97
CHEMTEX FILE #: B7020168

RESULTS OF ANALYSIS

CLIENT: DOMINO SUGAR

Source Identification: ACM Vent & BC Dust Collector

<u>CHEMTEX #</u>	<u>Sample Identification</u>	<u>Parameter</u>	<u>Sample Volume ml</u>	<u>Concentration mg/l</u>	<u>Results mg/sample</u>
B7010168	Run-1 Probe Wash w/Acetone (1/28/97)	Total Solids	165	644	106.26
B7010169	Run-2 Probe Wash w/Acetone (1/28/97)	Total Solids	165	228	37.62
B7010170	Run-3 Probe Wash w/Acetone (1/28/97)	Total Solids	125	198	24.75
B7010171	Run-1 Probe Wash w/Acetone (1/27/97)	Total Solids	35	220	7.71
B7010172	Run-2 Probe Wash w/Acetone (1/27/97)	Total Solids	40	990	39.6
B7010173	Run-3 Probe Wash w/Acetone (1/27/97)	Total Solids	90	3,320	298.8
B7010174	Blank	Total Solids	---	< 2	----

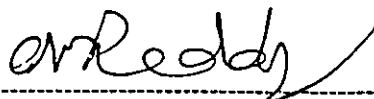
ACM
Vent

CHAR
Dust
Collector

<u>QC-Check Sample</u>	<u>True Value</u>	<u>Obtained Value</u>
Total Solids (mg/l)	50	48/46*

* Duplicate Analysis

<u>Parameter</u>	<u>Method Reference</u>	<u>Date Analyzed/Analysts</u>
Total Solids	EPA Method 160.3	2/4/97 KR



Dr. C. N. Reddy, Ph.D, CIH, ASP
Director

pap/CNR

The analytical results, opinions or interpretations contained in this report are based upon information and material supplied by the client, for whose exclusive and confidential use this report has been made. The analytical results, opinions or interpretations expressed represent the best judgment of CHEMTEX. CHEMTEX, however, makes no warranty or representation, express or implied, of any type, and expressly disclaims same. This report shall not be reproduced, in whole or in part, without the written approval of CHEMTEX.

1

AIHA Accredited

Organic, Metal, Wet Chemical, Biological and Petrol
Chemical Analyses for Multi-media Environmental
and Industrial Hygiene Samples.

Facilities are also available at:

3082 25th Street, Port Arthur, Texas 77642

6.2.15

Texas 78408
Louisiana 70663

(409) 983-4575 FAX (409) 982-1522
(512) 299-9900 FAX (512) 299-1155
(318) 625-8898 FAX (318) 625-5582

ETS

EMISSION TESTING SERVICES, INC.

FILTER WEIGHTS

CLIENT Domino Sugar
UNIT ID C Dust Collector TEST DATE 1-27-97

Filter Number	Sample ID	Initial Weight		Final Weight		Weight Gained
<u>3021</u>	<u>Run #1</u>	<u>3.8417</u>	<u>3.8418</u>	<u>3.8439</u>	<u>3.8441</u>	<u>2.3</u>
<u>3023</u>	<u>#2</u>	<u>3.8631</u>	<u>3.8634</u>	<u>3.8655</u>	<u>3.8651</u>	<u>21.7</u>
<u>3022</u>	<u>#3</u>	<u>3.8056</u>	<u>3.8059</u>	<u>4.2316</u>	<u>4.2312</u>	<u>425.3</u>

CLIENT Domino Sugar
UNIT ID ACM Vent TEST DATE 1-28-97

Filter Number	Sample ID	Initial Weight		Final Weight		Weight Gained
<u>3024</u>	<u>Run 1</u>	<u>3.8816</u>	<u>3.8847</u>	<u>3.9703</u>	<u>3.9701</u>	<u>85.4</u>
<u>3025</u>	<u>2</u>	<u>3.8311</u>	<u>3.8313</u>	<u>3.8440</u>	<u>3.8439</u>	<u>12.6</u>
<u>3026</u>	<u>3</u>	<u>3.8272</u>	<u>3.8269</u>	<u>3.8350</u>	<u>3.8351</u>	<u>8.2</u>

ANALYST Fred Dowling ANALYSIS DATE 2-18-97

VII. APPENDIX

The following appendices are presented as supporting documentation to the emission test report.

7.1.1 Resumes of Test Personnel

7.2.1 Test Equipment Calibrations

Kevin Thibodeaux
Sampling Specialist
Emission Testing Services, Incorporated
Baton Rouge, Louisiana

EDUCATION

U.S. EPA Round Robin sample integrity audits

Completed Emission Testing Services Air Sampling Techniques Seminar

Participates in on going Air-Pollution Training Institute Courses (APTI)

High School Graduate

CERTIFICATIONS

EPA Certified Opacity Reader (Method 9)
General Safety & Health Orientation Program
Respiratory Equipment Medical Certification
MSA Qualitative Fit Test Record

EXPERIENCE

1987 to Present - Testing Specialist for Emission Testing Services, Inc.
Performs EPA approved reference methods for compliance testing as well as other non-routine source evaluations. Responsibilities also include job preparation, equipment calibration and equipment maintenance.

Ryan Deaville
Sampling Technician Trainee
Emission Testing Services, Incorporated
Baton Rouge, Louisiana

EDUCATION

High School Graduate

CERTIFICATIONS

General Safety & Health Orientation Program
Respiratory Equipment Medical Certification
MSA Qualitative Fit Test Record

EXPERIENCE

06/96 to Present - Testing trainee technician for Emission Testing Services, Inc.
Supports in performing EPA approved reference methods for compliance testing
as well as other non-routine source evaluations. Responsibilities also include job
preparation, equipment calibration and equipment maintenance.

Jeff M. Savoie
Sampling Technician Trainee
Emission Testing Services, Incorporated
Baton Rouge, Louisiana

EDUCATION

Louisiana State University
B.S. -- Environmental Management Systems

CERTIFICATIONS

EPA Certified Opacity Reader (Method 9)
General Safety & Health Orientation Program
Respiratory Equipment Medical Certification
MSA Qualitative Fit Test Record

EXPERIENCE

10/95 to Present - Testing trainee technician for Emission Testing Services, Inc.
Supports in performing EPA approved reference methods for compliance testing
as well as other non-routine source evaluations. Responsibilities also include job
preparation, equipment calibration and equipment maintenance.

ETS

EMISSION TESTING SERVICE, INC.

METER BOX CALIBRATION DATA AND CALCULATION FORM (English Units)

Leak Check Good @: .0012 ft³
Barometric Pres., P_b: 30.28 in. Hg

Meter Box Number: 10
Date Calibrated: 1-16-97

Orifice Delta H Setting (ΔH) in. H ₂ O	Wet Test Meter Reading (V _w) ft ³	Dry Gas Meter Readings			Wet Test Meter Reading (T _w) °F	Dry Gas Meter			Run Time (Ø) Min.
		Initial (V _i) ft ³	Final (V _f) ft ³	Total (V _t) ft ³		Inlet (T _i) °F	Outlet (T _o) °F	Avg (T _a) °F	
0.5	5	673.691	678.885	5.194	72°	102	108	105.5	12.70
1.0	5	678.985	684.07	5.185	72°	108	106	105.5	8.71
1.5	10	684.167	694.511	10.344	72°	101	107	106.7	14.75
2.0	10	694.511	704.848	10.337	72°	102	106	105.25	13.05
3.0	10	704.845	715.174	10.329	72°	102	105	105.5	10.62

ΔH In. H ₂ O	ΔH 13.6 (F)	$Y = \frac{V_w \cdot P_b \cdot (T_a + 460)}{V_d \cdot (P_b + F_1) \cdot (T_w + 460)}$	$\Delta H@ = \frac{0.0317 \cdot \Delta H}{P_b \cdot (T_a + 460)} \cdot \left[\frac{(T_w + 460) \cdot \phi}{V_w} \right]^2$
0.5	.0368	1.022	1.6902
1.0	.0737	1.023	1.5900
1.5	.110	1.0266	1.7061
2.0	.147	1.0225	1.7854
3.0	.221	1.0216	1.7728
Average		1.0231	1.7089

Y = Ratio of reading of wet test meter to dry test meter; tolerance for individual values ± 0.02 from average.

ΔH@ = Orifice pressure differential that equates to 0.75 cfm of air @ 68 °F and 29.92 inches of mercury, in. H₂O; tolerance for individual values ± 0.20 from average.

Calibrated by: Craig & Jonathan

THERMOCOUPLE CALIBRATION FORM

CALIBRATED BY: Craig & Jonathan DATE: 1-16-99

METER BOX NUMBER: 10 BAROMETRIC PRESSURE: 30.53

TRIAL	THERMOMETER ICE BATH (T_f)	THERMOCOUPLE ICE BATH (TC_f)	THERMOMETER BOILING WATER (T_b)	THERMOCOUPLE BOILING WATER (TC_b)
1	<u>30°</u>	<u>30°</u>	<u>215</u>	<u>214</u>
2	<u>30°</u>	<u>31°</u>	<u>217</u>	<u>216</u>
3	<u>30°</u>	<u>31°</u>	<u>217</u>	<u>216</u>
AVG	<u>30°</u>	<u>92° 30.6</u>	<u>216.3</u>	<u>215.3</u>

CALCULATIONS

$$\% \text{ ERROR @ BOILING POINT} = \frac{\overline{T_b} - \overline{TC_b}}{\overline{T_b}} \times 100 = \underline{1.0\% \text{ error}}$$

$$\% \text{ ERROR @ FREEZING POINT} = \frac{\overline{T_f} - \overline{TC_f}}{\overline{T_f}} \times 100 = \underline{-1.0\% \text{ error}}$$

OKAY IF $-1.5\% < \% \text{ ERROR} < +1.5\%$

REFERENCE: METHOD 2 SECTION 4.3

PITOT TUBE CALIBRATION FORM

Pitot Tube Number: P-3-1

Calibrated By: SAVOIE

$C_{p(Std)}$: 0.99

Date Calibrated: 8-28-96

Point Number	Standard Pitot ΔP (in. H_2O)	"S" Type Pitot ΔP , in H_2O	
		Impact Side	Static Side
Low	<u>0.105</u>	<u>0.150</u>	<u>0.150</u>
Mid	<u>0.585</u>	<u>0.82</u>	<u>0.815</u>
High	<u>1.10</u>	<u>1.55</u>	<u>1.55</u>

CALCULATIONS

$$C_{p(s)} = C_{p(Std)} \times \sqrt{\frac{\Delta P_{(Std)}}{\Delta P_{(s)}}}$$

$$\text{Deviation} = C_{p(s)} - \bar{Cp} \quad (\text{Impact Side or Static Side})$$

$$\text{Avg. Dev.} = \frac{\sum_{i=1}^3 |C_{p(s)} - \bar{Cp}|}{3} \quad (\text{Impact Side or Static Side})$$

Impact Side

$C_{p(s)}$	Deviation
<u>0.828</u>	<u>0.005</u>
<u>0.836</u>	<u>0.003</u>
<u>0.834</u>	<u>0.001</u>
$\bar{Cp} = $ <u>0.833</u>	

Static Side

$C_{p(s)}$	Deviation
<u>0.828</u>	<u>0.006</u>
<u>0.839</u>	<u>0.005</u>
<u>0.834</u>	<u>0.000</u>
$\bar{Cp} = $ <u>0.834</u>	

$$| \bar{Cp} (\text{Static Side}) - \bar{Cp} (\text{Impact Side}) | = \underline{0.001} \quad (\text{Must be } \leq 0.01)$$

$$\text{Average Deviation (Impact Side)} = \underline{0.003} \quad (\text{Must be } \leq 0.01)$$

$$\text{Average Deviation (Static Side)} = \underline{0.004} \quad (\text{Must be } \leq 0.01)$$

$$C_p = \frac{\bar{Cp} (\text{Static Side}) + \bar{Cp} (\text{Impact Side})}{2} \quad (\text{Should be } 0.80 \leq C_p \leq 0.85) = \underline{0.834}$$