

# **TRC Environmental Corporation**

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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/](http://www.epa.gov/ttn/chief/ap42/)

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## **EMISSIONS TESTING OF A PRESS REGENERATIVE THERMAL OXIDIZER**

*Presented to*

**Louisiana-Pacific Corporation**  
Houlton, Maine

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TRC Project No. 19624  
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## 1.0 INTRODUCTION

TRC Environmental Corporation (TRC) was retained by Louisiana-Pacific Corporation to conduct an emissions measurement program on the regenerative thermal oxidizer (RTO) at the Louisiana-Pacific facility in New Limerick, Maine. The purpose of this program was to measure emissions of formaldehyde, oxides of nitrogen (NO<sub>x</sub>), and carbon monoxide (CO) at the inlet and outlet of the RTO. Emissions measurements for NO<sub>x</sub> and CO were conducted in accordance with accepted USEPA test methodologies. Formaldehyde testing was conducted using the NCASI Acetylacetone Method for measuring formaldehyde in water.

The test program was conducted on December 14, 1995 and was supervised by Mr. Raul Baez of TRC. Ms. Sue Somers and Mr. Mark Stile, both of Louisiana-Pacific, provided the process and logistical support during the program.

Section 2.0 of this test report presents a summary and discussion of the test results. Section 3.0 describes the process and associated control equipment and the parameters that were monitored during testing. Section 4.0 details the test methods to be used, and Section 5.0 presents TRC's quality control plan for this program.

## 2.0 SUMMARY AND DISCUSSION OF RESULTS

The purpose of this program was to measure emissions of formaldehyde, oxides of nitrogen (NO<sub>x</sub>), and carbon monoxide (CO) at the inlet and outlet of the regenerative thermal oxidizer (RTO). Emissions measurements for NO<sub>x</sub> and CO were conducted in accordance with accepted USEPA test methodologies. Formaldehyde testing was conducted using the NCASI Acetylacetone Method for measuring formaldehyde in water. The following sections provide the results for each pollutant.

### 2.1 Regenerative Thermal Oxidizer Emissions

Emissions measurements were conducted at the inlet and outlet of the regenerative thermal oxidizer (RTO) to determine inlet and outlet concentrations of formaldehyde, NO<sub>x</sub> and CO. Triplicate 60-minute formaldehyde tests were conducted at each location in accordance with the NCASI Acetylacetone Method for measurement of formaldehyde in water. Triplicate 20-minute NO<sub>x</sub> and CO tests were conducted at each location in accordance with EPA Methods 7E and 10, respectively. In addition, TRC measured the volumetric flowrate, concentration of oxygen, and moisture content of the exhaust gases at each location in accordance with EPA Methods 1-4. The results of the inlet and outlet tests are presented in **Tables 2-1 and 2-2**, respectively. All associated field data are presented in **Appendix A**.

#### 2.1.1 Formaldehyde Emissions

The results of the formaldehyde testing indicate that the average inlet concentration was 6.6 parts per million (ppm) and that the average inlet loading rate was 2.25 lb/hr. The average outlet concentration was 0.3 ppm and the average outlet emission rate was 0.10 lb/hr. The average removal efficiency of formaldehyde was 95.5 %.

#### 2.1.2 NO<sub>x</sub> and CO Emissions

The results of the continuous emissions monitoring for NO<sub>x</sub> and CO at the inlet of the RTO indicate that the average NO<sub>x</sub> concentration was 1.6 ppm and the average inlet loading rate was 0.83 lb/hr. The average CO concentration was 9.8 ppm and the average inlet loading rate was 3.10 lb/hr.

Table 2-1

Summary of Formaldehyde, NOx, and CO Loading  
at the RTO Inlet

Louisiana-Pacific Corporation  
New Limerick, Maine  
December 14, 1995

Test Number	In-1	In-2	In-3	Average
Time	1035-1135	1200-1300	1338-1438	
Location	RTO -Inlet	RTO -Inlet	RTO -Inlet	
<u>Stack Conditions</u>				
Stack Temperature (°F)	102	104	104	103
CO <sub>2</sub> (%)	0.00	0.00	0.00	0.00
O <sub>2</sub> (%)	20.8	20.9	20.8	20.8
Moisture (%)	1.33	1.37	1.40	1.36
Volumetric Flowrate, Actual (ACFM)	78432	79065	79364	78954
Volumetric Flowrate, Standard (SCFM) <sup>a</sup>	73109	73420	73688	73406
Volumetric Flowrate, Dry Std. (DSCFM) <sup>b</sup>	72137	72415	72660	72404
<u>Formaldehyde Emissions</u>				
Concentration (ppm)	5.9	6.5	7.6	6.6
Mass Emission Rate (lb/hr) <sup>c</sup>	1.98	2.20	2.57	2.25
<u>NOx Emissions</u>				
Concentration (ppm)	1.8	2.1	0.9	1.6
Mass Emission Rate (lb/hr) <sup>c</sup>	0.93	1.09	0.47	0.83
<u>CO Emissions</u>				
Concentration (ppm)	16.3	7.1	6.1	9.8
Mass Emission Rate (lb/hr) <sup>c</sup>	5.13	2.24	1.93	3.10

a - ACFM Actual Cubic Feet per Minute.

b - DSCFM Dry Standard Cubic Feet per Minute at 68°F and 29.92 in. Hg.

c - lb/hr = conc(ppm) x MW x flow(DSCFM) x 15.58E-08

MW Formaldehyde = 30

MW NOx = 46

MW CO = 28

Table 2-2

Summary of Formaldehyde, NOx, and CO Emissions  
at the RTO Outlet

Louisiana-Pacific Corporation  
New Limerick, Maine  
December 14, 1995

Test Number	out-1	out-2	out-3	Average
Time	1035-1135	1200-1300	1338-1438	
Location	RTO-outlet	RTO-outlet	RTO-outlet	
<u>Stack Conditions</u>				
Stack Temperature (°F)	232	234	235	233
CO <sub>2</sub> (%)	0.00	0.00	0.00	0.00
O <sub>2</sub> (%)	20.3	20.4	20.4	20.4
Moisture (%)	2.11	2.04	2.42	2.19
Volumetric Flowrate, Actual (ACFM)	95318	94835	96425	95526
Volumetric Flowrate, Standard (SCFM) <sup>a</sup>	72896	72370	73477	72914
Volumetric Flowrate, Dry Std. (DSCFM) <sup>b</sup>	71357	70891	71700	71316
<u>Formaldehyde Emissions</u>				
Concentration (ppm)	0.3	0.3	0.3	0.3
Mass Emission Rate (lb/hr) <sup>c</sup>	0.10	0.10	0.11	0.10
<u>NOx Emissions</u>				
Concentration (ppm)	15.9	16.0	15.6	15.8
Mass Emission Rate (lb/hr) <sup>c</sup>	8.13	8.13	8.02	8.09
<u>CO Emissions</u>				
Concentration (ppm)	16.5	15.8	16.6	16.3
Mass Emission Rate (lb/hr) <sup>c</sup>	5.14	4.89	5.19	5.07

a - ACFM Actual Cubic Feet per Minute.

b - DSCFM Dry Standard Cubic Feet per Minute at 68°F and 29.92 in. Hg.

c - lb/hr = conc(ppm) x MW x flow(DSCFM) x 15.58E-08

MW Formaldehyde = 30

MW NOx = 46

MW CO = 28

The results of the continuous emissions monitoring for NO<sub>x</sub> and CO at the outlet of the RTO indicate that the average NO<sub>x</sub> concentration was 15.8 ppm and the average emission rate was 8.09 lb/hr. The average CO concentration was 16.3 ppm and the average inlet loading rate was 5.07 lb/hr.

### 2.1.3 Volumetric Flowrate

Volumetric flowrate measurements were conducted at the inlet and outlet of the RTO in accordance with EPA Method 2. The results indicated that the inlet flowrate was approximately 1000 DSCFM greater than the outlet flowrate. However, the outlet flowrate should have been greater than the inlet flowrate due to the addition of combustion air. This error can be partially explained by the accuracy of the EPA Method 2. EPA Method 2 has an accuracy of +/- 10%. It is possible that the inlet measurement was biased high and the outlet may have been biased low.

### 3.0 PROCESS DESCRIPTION

The Houlton, Maine facility operates one oriented strandboard line including two wafer dryers, a press, and two thermal oil heaters.

#### 3.1 Regenerative Thermal Oxidizer

The regenerative thermal oxidizer (RTO) is used to control emissions of volatile organic compounds and particulate matter from the press operation. The RTO is fired on natural gas. During testing the plant operated at approximately 90% of the daily production rate of 440 tons per day. Plant personnel monitored and recorded the following data:

Panels/Hour  
Panel Weight  
Press Temp.  
Tons/Hour

Copies of the recorded process data are presented in **Appendix C**.

#### 4.0 SAMPLING AND ANALYTICAL METHODS

Emissions measurements were conducted at the inlet and outlet of the RTO in accordance with the following methods.

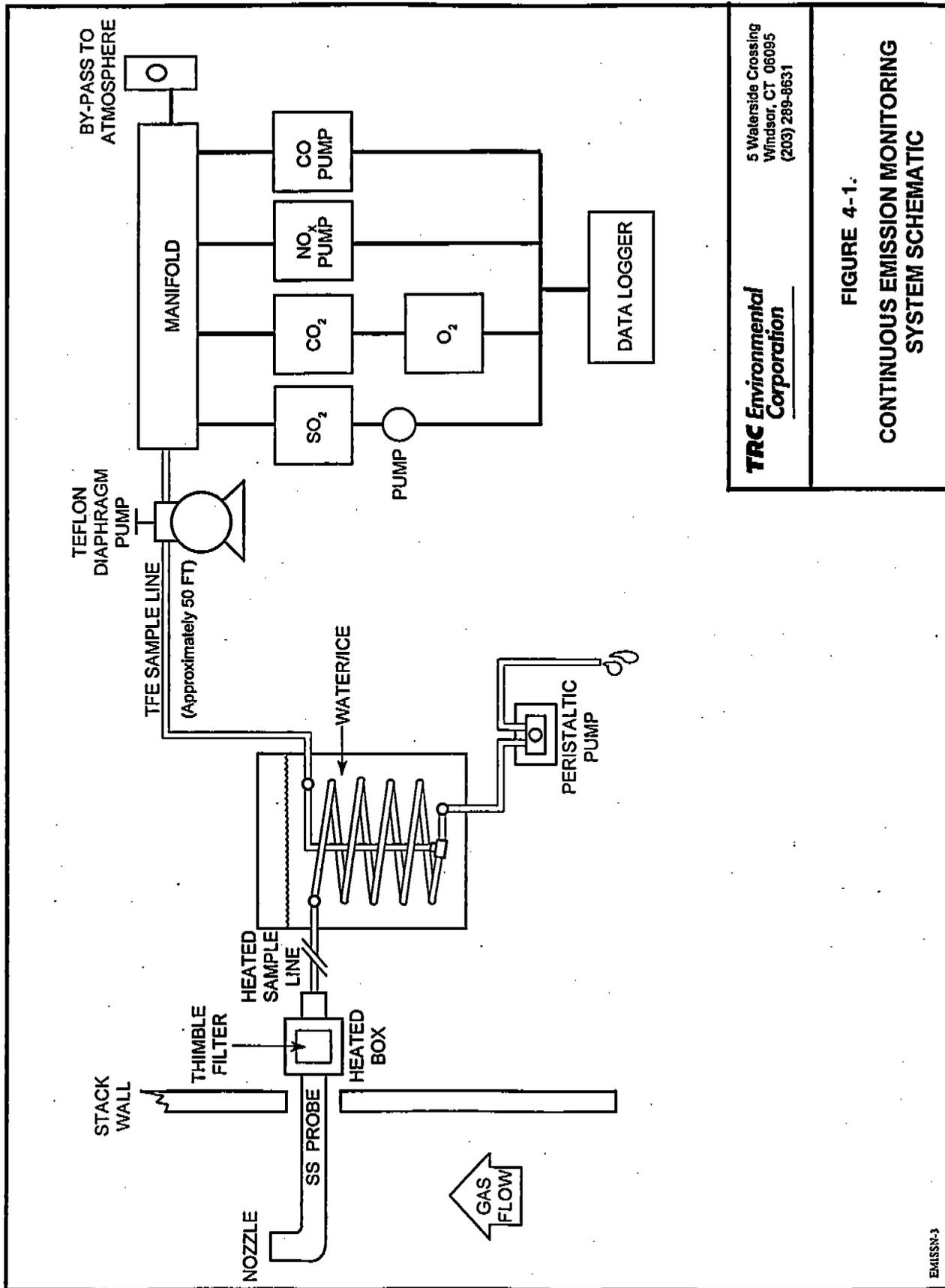
##### 4.1 NO<sub>x</sub> and CO Emissions Measurements - EPA Methods 7E and 10

Triplicate 20-minute tests were conducted at the inlet and outlet of the RTO to measure concentrations of NO<sub>x</sub> and CO. Testing was conducted in accordance with EPA Methods 7E and 10, respectively. Diluent concentrations of oxygen were measured in accordance with EPA Method 3A. A schematic of the measurement system is presented in **Figure 4-1**. A single CEMS was used to evaluate inlet and outlet concentrations. A three way valve was placed in line prior to the conditioning system, which allowed the operator to switch from the inlet to the outlet location and vice versa. Each sampling location had a dedicated sampling probe and heated sample line prior to the three way valve and conditioning system.

All CEM data was recorded by a Yokogawa Model 2300 strip chart/data logger which is capable of integrating 5-minute and 20-minute averages. The CEM system was housed in the TRC Mobile Environmental Laboratory (MEL) which was parked at the base of the RTO.

##### 4.1.1 Sample Conditioning System

An in-stack Alundum thimble filter with a stainless steel nozzle facing away from the stack gas flow was used to remove any PM from the sample gas stream. The thimble filter was mounted on the end of a stainless steel sampling probe. The sample was then drawn through 100 feet of heated (325°F ± 25°F) Teflon sample line through a condenser system to remove the moisture from the gas stream. The sample was drawn through the tubing by a leak-free Teflon double-diaphragm pump to a stainless steel sample manifold with an atmospheric bypass rotameter. The analyzers then drew their samples from the manifold.



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FIGURE 4-1.  
 CONTINUOUS EMISSION MONITORING  
 SYSTEM SCHEMATIC

#### 4.1.2 NO<sub>x</sub> Analyzer

A Thermo-Electron Corporation Model 10A Chemiluminescent NO/NO<sub>x</sub> analyzer was used to determine NO<sub>x</sub> concentrations. The chemiluminescent reaction of NO and O<sub>3</sub> (ozone) provides the basis for the analytical method (NO + O<sub>3</sub> → NO<sub>2</sub> + O<sub>2</sub> + light). A photomultiplier-electrometer-amplifier produces a current proportional to the NO concentration. The output of the amplifier provides a signal for direct readout on a meter indicator, or for outputs to a recorder or computer.

#### 4.1.3 CO Analyzer

A TECO Model 48 nondispersive infrared gas analyzer was used to measure CO concentrations. The analyzer contains an infrared detector that uses the signal nondispersive beam technique with alternate modulations of the sample and reference cells. Radiation absorbed by CO in the sample cell results in a capacitance change in the detector which is proportional to the CO concentration.

#### 4.1.4 O<sub>2</sub> Analyzer

A Horiba Model PMA-200 O<sub>2</sub> analyzer was used to determine the concentration of O<sub>2</sub> in the stack gas. This instrument uses the paramagnetic principle, whereby the magnetic susceptibility of the gas volume is measured by the force acting on a nonmagnetic test body suspended in a magnetic field. The force is converted to an output current proportional to the O<sub>2</sub> concentration.

#### 4.1.5 Data Acquisition and Handling

All CEM data was monitored by a Yokagawa Model 2400 automatic data logger. Emissions data was "viewed" by the data logger every six seconds, averaged, and printed at 5-minute intervals and 20-minute intervals.

#### 4.1.6 CEM Calibrations

Calibrations (zero, low, mid, and high of span) of each analyzer were performed using EPA Protocol 1 gases at the beginning and end of each test period. Calibration gases were introduced to the system through a three-way valve at the back of the sample probe. The gas concentrations used were approximately 30%, 60%, and 90% of each analyzer's range. Each analyzer was multi-point calibrated at the beginning of the test program to establish linearity of each instrument.

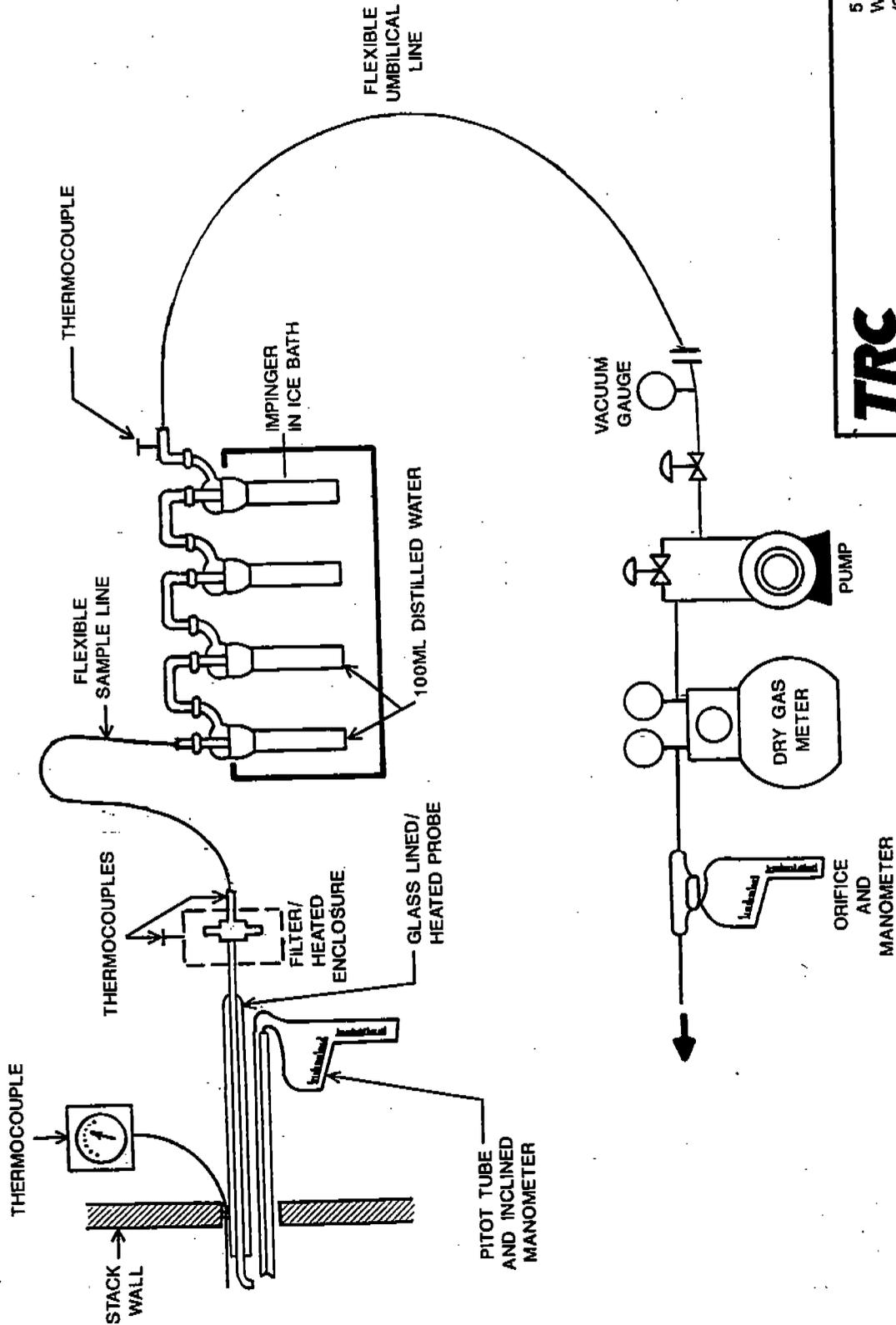
#### 4.2 Formaldehyde Measurements - NCASI Method

Triplicate 60-minute tests were conducted at the inlet and outlet of the RTO to measure concentrations of formaldehyde. Testing was conducted in accordance with the NCASI Acetylacetone Method (Determination of Formaldehyde in Water), with inlet and outlet testing conducted simultaneously. The sampling system describe in the NCASI method was modified by TRC. TRC's modification was to use a standard EPA Method 5 train with standard impingers instead of mini-impingers. This modification allowed for more accurate measurement of sample volume and the use of heated probes and filters.

##### 4.2.1 Sample Collection

Formaldehyde sampling was accomplished by use of an EPA Method 5 train. The sample train is shown schematically in **Figure 4-2** and consisted of a nozzle, probe, filter, four impingers, a vacuum pump, dry-gas meter, and an orifice flow meter.

A stainless steel nozzle was attached to a glass-lined stainless steel probe which was heated to prevent condensation. Teflon mat filter papers supported in 4½-inch glass filter holders were used to remove particulate matter from the gas stream. The filter assembly was enclosed in a heated box to maintain temperatures at  $248^{\circ}\text{F} \pm 25^{\circ}\text{F}$ . A thermocouple located inside the back half of the filter holder monitored the gas stream temperature and verified that the temperature was kept at  $248^{\circ}\text{F} \pm 25^{\circ}\text{F}$ .



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Figure 4-2  
 NCASI Formaldehyde Train

An ice bath containing four impingers was attached to the back end of the filter. The first and second impingers contained 100 milliliters (ml) each of HPLC grade deionized water. The third impinger was empty, and the fourth impinger contained silica gel to remove any remaining moisture. Flexible tubing, a vacuum gauge, a needle valve, a leakless vacuum pump, a bypass valve, a dry-gas meter calibration orifice, and an inclined manometer completed the sampling train. The stack velocity pressure and temperature were monitored by an S-type pitot and a thermocouple connected to a potentiometer. A check valve was not used in the TRC sampling train.

Sampling flow was adjusted by means of the bypass valve. Before and after each particulate test run, the sampling train was leak checked to meet the 0.02 cfm limit. All pertinent test data were recorded on the appropriate field data sheets.

#### 4.2.2 Sample Recovery

Two sample containers were used, as follows:

- Container No. 1: The impinger contents were measured volumetrically to the nearest milliliter and deposited into a sample jar. Each impinger was rinsed three times with HPLC grade deionized water. The water rinses were also deposited into the sample jar.
- Container No. 2: Silica gel from the fourth impinger was transferred to its original container and weighed to the nearest 0.5 milligram (mg).

#### 4.2.3 Sample Analysis

The samples were transported to TRC's laboratory, where the following analyses were performed:

- Container No. 1: The impinger contents were analyzed in accordance with the NCASI Acetylacetone Method.

Container No. 2: Silica gel was weighed to the nearest 0.5 mg. The weight of the moisture entrapped in the silica gel, along with the impingers, was used to calculate the moisture content of the stack gas.

#### 4.3 Flowrate and Moisture Measurements

Flowrate measurements were conducted according to EPA Method 2. An S-type pitot and an inclined manometer were used to measure the velocity pressure head, and a thermocouple was used to monitor gas stream temperature.

#### 4.4 Sampling Locations

Sampling points for isokinetic and/or flowrate measurements at each location were determined in accordance with EPA Method 1.

##### 4.4.1 Regenerative Thermal Oxidizer Inlet and Outlet

Inlet sampling was conducted from two ports in the 54-inch diameter stack. The ports were located 2.8 diameters downstream from any flow disturbances and 5.4 diameters upstream from any flow disturbances. Twenty four traverse points were sampled with each flowrate measurement. The inlet sampling point locations are presented in **Figure 4-3**.

Outlet sampling was conducted from two ports in the 76-inch diameter stack. The ports were located 10.4 diameters downstream from any flow disturbances and 6.9 diameters upstream from the stack exhaust. Twelve traverse points were sampled with each flowrate measurement. The outlet sampling point locations are presented in **Figure 4-4**.

Figure 4-3

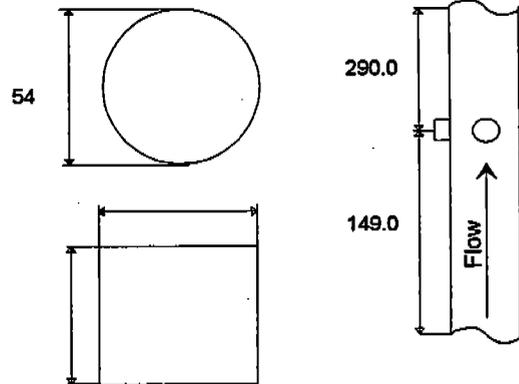
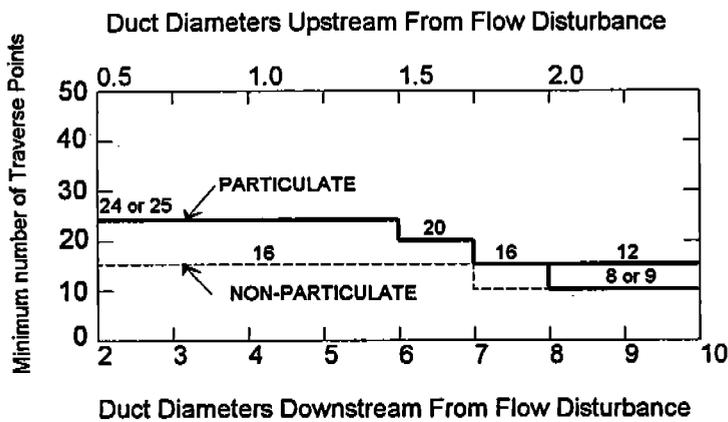
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EPA Method 1 Data Sheet

Firm Louisiana-Pacific  
 Location RTO Inlet  
 Diameters Upstream 5.4  
 Diameters Downstream 2.8  
 Nipple Size (in.) 6

Total Traverse Points Required 24  
 Number of Ports 2  
 Points Per Port 12  
 Traverse (Horizontal or Vertical) Horizontal/Vertical

Minimum Number of Traverse Points For Particulate and Non-Particulate Traverses



Cross-Sectional Layout For Rectangular Stacks	
Total Traverse Points	Matrix
9	3x3
12	4x3
16	4x4
20	5x4
25	5x5

Location of Points on a Circular Stack

Point Number On a Diameter	(Percent of Stack Diameter from Inside Wall to Traverse Point)			
	No. of Traverse Points on a Diameter			
	4	8	10	12
1	6.7	3.2	2.6	2.1
2	25.0	10.5	8.2	6.7
3	75.0	19.4	14.6	11.8
4	93.3	32.3	22.6	17.7
5		67.7	34.2	25.0
6		80.6	65.8	35.6
7		89.5	77.4	64.4
8		96.8	85.4	75.0
9			91.8	82.3
10			97.4	88.2
11				93.3
12				97.9

Traverse Point Location

Point Number	Distance From Wall	Total Distance
1	1.1	7.1
2	3.6	9.6
3	6.4	12.4
4	9.6	15.6
5	13.5	19.5
6	19.2	25.2
7	34.8	40.8
8	40.5	46.5
9	44.4	50.4
10	47.6	53.6
11	50.4	56.4
12	52.9	58.9

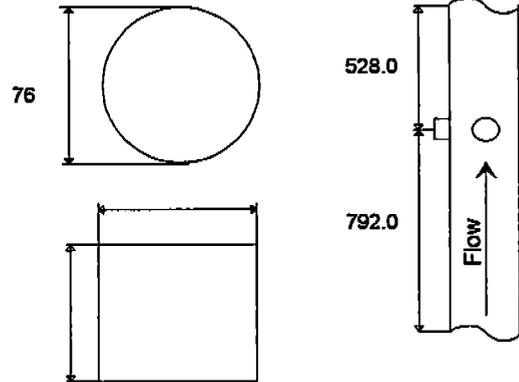
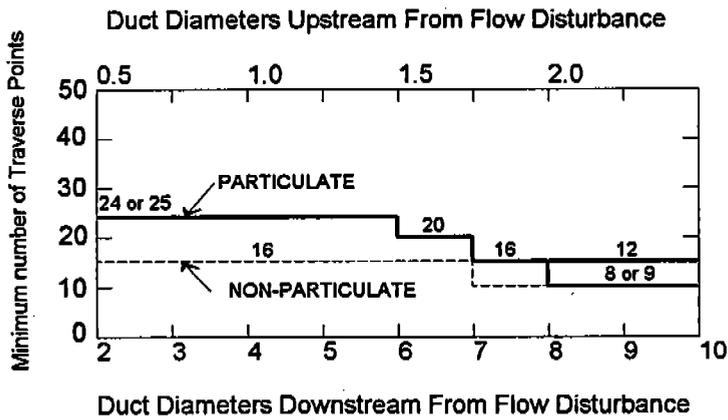
Figure 4-4

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EPA Method 1 Data Sheet

Firm	<u>Louisiana-Pacific</u>	Total Traverse Points Required	<u>12</u>
Location	<u>RTO Outlet</u>	Number of Ports	<u>2</u>
Diameters Upstream	<u>6.9</u>	Points Per Port	<u>6</u>
Diameters Downstream	<u>10.4</u>	Traverse (Horizontal or Vertical)	<u>Horizontal</u>
Nipple Size (in.)	<u>6</u>		

Minimum Number of Traverse Points For Particulate and Non-Particulate Traverses



Cross-Sectional Layout For Rectangular Stacks	
Total Traverse Points	Matrix
9	3x3
12	4x3
16	4x4
20	5x4
25	5x5

Location of Points on a Circular Stack

Point Number On a Diameter	(Percent of Stack Diameter from Inside Wall to Traverse Point)			
	No. of Traverse Points on a Diameter			
	4	8	10	12
1	6.7	3.2	2.6	2.1
2	25.0	10.5	8.2	6.7
3	75.0	19.4	14.6	11.8
4	93.3	32.3	22.6	17.7
5		67.7	34.2	25.0
6		80.6	65.8	35.6
7		89.5	77.4	64.4
8		96.8	85.4	75.0
9			91.8	82.3
10			97.4	88.2
11				93.3
12				97.9

Traverse Point Location

Point Number	Distance From Wall	Total Distance
1	3.3	9.3
2	11.1	17.1
3	22.5	28.5
4	53.5	59.5
5	64.9	70.9
6	72.7	78.7
7		
8		
9		
10		
11		
12		

## 5.0 QUALITY ASSURANCE

The project manager was responsible for implementating the TRC quality assurance (QA) program as applied to this project. The program was designed to ensure that emission measurement work was performed by qualified people using proper equipment following written procedures in order to provide accurate, defensible data.

At the beginning of the test day, a meeting was held to orient TRC and Louisiana-Pacific personnel to the activities scheduled for that day and to determine if any special considerations were appropriate for the day's work.

### 5.1 Sampling Quality Assurance

The TRC quality assurance program for source measurements was designed so that the work was done by competent, trained individuals experienced on the specific methodologies being used; using properly calibrated equipment; and using approved procedures for sample handling and documentation.

TRC's measurement devices, pitot tubes, dry-gas meters, thermocouples, and portable gas analyzers were uniquely identified and calibrated with documented procedures and acceptance criteria before and after each field effort. Records of all calibration data are maintained in TRC files.

Data were recorded on standard forms. Bound field notebooks were used to record observations and miscellaneous elements affecting data, calculations, or evaluation.

Specific details of TRC's QA program for stationary air pollution sources may be found in *Quality Assurance Handbook for Air Pollution Measurement Systems*, Volume III (EPA-600/4-77-027b).

In the mobile laboratory, analysts recorded calibration and analysis data in notebooks and summarized specific data on prepared data sheets. Notebooks, data sheets, and calculations were reviewed by the project manager.

## 5.2 Analytical Quality Control

Compressed gases used as fuels and carriers were purchased at specific purities, according to application. Compressed gases used as calibration standards are always National Institute for Standards and Technology (NIST) traceable, either directly or indirectly.

In the TRC and subcontract laboratories, all QC samples including field blank samples, reagent and filter blanks, and any audit samples were analyzed with the actual test samples. Each laboratory maintains a continuous QC program to monitor instrument response and analyst proficiency and to ensure the precision and accuracy of all analytical results. These programs were developed in consultation with EPA, NIOSH, and various state departments of health.

TRC used some or all of the following quality control procedures during the field and laboratory sampling program:

*Field Blanks*—For each set of samples taken, a field blank was also be collected. Field blanks included filter materials and solutions used for sample collection and recovery.

*Replicates*—When the analytical procedure permitted, a prescribed number of the analyses were duplicated to indicate the precision of the method used.

Appendix A

Field Data

ISOKINETIC SAMPLING DATA

PLANT NAME, CITY, STATE		SAMPLING LOCATION				Method No.		RUN NUMBER		TEST DATE		Notes		
L.P. Harbor ME		inlet				IN 1		IN 1		12-14-85				
OPERATORS	SIB RB	DGM No.	DGM CAL	Static P (In. H2O)	Ambient T. (°F)	Filter Type & Filter Number(s)	Stack (D. (In.))	Piton Tubs Cp	Probe Length and Liner Type	Nozzle Number	Diameter	K Factor	Notes	
														DGM No.
PITOT LEAK CHECK	OK	8823	1.88	1.00	2	C-7	1.88	100	240	235	25	NA	33	30
POINT NUMBER	1	10:35	11.00	14.700	18.300	23.900	25.600	29.200	32.800	36.300	39.800	43.300	46.900	50.300
CLOCK TIME (24-hour)	11:35	0	11.00	14.700	18.300	23.900	25.600	29.200	32.800	36.300	39.800	43.300	46.900	50.300
STOP	11:35	60	53.663	53.663	53.663	53.663	53.663	53.663	53.663	53.663	53.663	53.663	53.663	53.663
Total Run Time		DGM Volume		Ave. SQR AP		Average ΔH		Stack Temp		Ave. DGM Temp		40		
60		42.663		42.663		42.663		42.663		42.663		42.663		

TRC

ISOKINETIC SAMPLING DATA

PLANT NAME, CITY, STATE		SAMPLING LOCATION			METHOD NO.		RUN NUMBER		TEST DATE		Notes	
Kosimo Paper		Inlet			In # 2		12/14/95		Forum/bhp.k			
OPERATORS	Static P. (in H <sub>2</sub> O)	Ambient T. (°F)	Filter Type & Filter Number(s)	Stack (D. in.)	Plot Tube Co.	Probe Length and Liner Type	Nozzle Number	Diameter				
Genco/LEPZ	45	10		34	087	3/4 Teflon / MANTLE	A4	N/A				
PITOT LEAK CHECK	DGM GAL	Assumed Moist. %	Plot No.	Stack Tharm. No.	Leak Check (In.) (in Hg)	Leak Check (In.) (in Hg)	Leak Check (In.) (in Hg)	K-Factor				
OK	1.88	2	C-7		10	0.001	5	0.000				
CLOCK TIME (24-hour)	ELAPSED TEST TIME (min)	DRY-GAS METER READING (Cubic Feet)	AP Velocity (in H <sub>2</sub> O)	AH Office (in H <sub>2</sub> O)	Stack Temp. (°F)	Filter Oven Temp. (°F)	Impinger Temp. (°F)	AUX Temp. (°F)	DGM Temp. (°F)	DGM In.	DGM Out	Sample Treh Vacuum (in Hg)
12:00	0	53,800		1.88	100	230	36	NA	39	38	38	3
	5	57,400		1.88	100	230	35		39	38	38	3
	10	60,900		1.88	100	240	35		41	37	37	3
	15	64,400		1.88	100	240	36		42	37	37	3
	20	67,900		1.88	100	238	40		44	37	37	3
	25	71,400		1.88	100	237	40		44	37	37	3
	30	74,800		1.88	100	243	39		45	37	37	3
	35	77,200		1.88	100	247	39		46	38	38	3
	40	81,600		1.88	100	252	38		46	38	38	3
	45	85,100		1.88	100	254	37		46	38	38	3
	50	88,500		1.88	102	256	37		47	39	39	3
	55	91,900		1.88	100	257	37		47	39	39	3
STOP	13:00	95,312										
Total Run Time		DGM Volume	Ave. SQR AP	Average ΔH	Stack Temp							
60		41,512										A
										Ave. DGM Temp.		A









PLANT LP  
 DATE 12-14-95 9 30AM  
 LOCATION Press - Inlet stack  
 STACK DIMENSIONS 54.5" AREA \_\_\_\_\_ ft<sup>2</sup>  
 BAROMETRIC PRESSURE, P<sub>b</sub> = 30.04 in. Hg  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± - 4.5 in. H<sub>2</sub>O  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> 289  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = 10%  
 PITOT NO. \_\_\_\_\_ C<sub>p</sub> = 0.99  
 TESTER \_\_\_\_\_

SCHMATIC OF TRAVERSE POINT LOCATION  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>s</sub> (°F)	± θ	Pitots reversed FOR Negative Flow?	√ΔP · COS θ
A	1	1.5		90			
	2	1.7		100			
	3	1.9		100			
	4	2.0		100			
	5	2.7		105			
	6	2.2		105			
	7	2.3		105			
	8	2.4		105			
B	1	1.3		100			
	2	1.4		102			
	3	2.2		100			
	4	2.2		100			
	5	1.9		100			
	6	1.9		100			
	7	1.8		100			
	8	1.7		103			
Average			2.756	T <sub>s</sub> = 101°F T <sub>sr</sub> = 561°R	*		

\*Avg. of absolute values including zeroes

Absolute Gas Temperature, T<sub>sr</sub> = T<sub>s</sub> + 460° = 29.709  
 Absolute Gas Pressure, P<sub>s</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 = 30.01  
 Gas Velocity, V<sub>s</sub> = (85.49) C<sub>p</sub> (√ΔP · COS θ) avg =  $\frac{79.98}{1.5916}$  ft/sec = 59.97  
 Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>s</sub>) (60) (A) =  $\frac{16780}{5003}$  ACFM  
 Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528^{\circ}R}{T_{sr}}\right) \left(\frac{P_s}{29.92}\right) \left(\frac{100 - \% H_2O}{100}\right)$  = 30464 DSCFM

PLANT LP  
 DATE 12-14-95 1200  
 LOCATION Inlet  
 STACK DIMENSIONS 54.5 AREA \_\_\_\_\_ ft<sup>2</sup>  
 BAROMETRIC PRESSURE, P<sub>b</sub> = 30.04 in. Hg  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± -4.6 in. H<sub>2</sub>O  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> 28.9  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = \_\_\_\_\_  
 PITOT NO. \_\_\_\_\_ Cp = 0.64  
 TESTER \_\_\_\_\_

SCHMATIC OF TRAVERSE POINT LOCATION  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>s</sub> (°F)	± θ	Pitots reversed for Negative Flow?	√ΔP · COS θ
A	1	1.6		100			
	2	1.7		100			
	3	1.9		100			
	4	2.1		100			
	5	2.2		105			
	6	2.2		102			
	7	2.3		106			
	8	2.4		107			
B	1	1.4		100			
	2	1.4		100			
	3	2.2		102			
	4	2.2		103			
	5	1.9		104			
	6	1.9		105			
	7	1.8		107			
	8	1.7		104			
Average			1.385	T <sub>s</sub> = 103°F	*		
				T <sub>sr</sub> = 563°R			

\*Avg. of absolute values including zeroes

Absolute Gas Temperature, T<sub>sr</sub> = T<sub>s</sub> + 460°  
 Absolute Gas Pressure, P<sub>s</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 = 29.70  
 Gas Velocity, V<sub>s</sub> = (85.49) C<sub>p</sub> (√ΔP · COSθ) avg  $\sqrt{\frac{T_{sr} \text{ avg}}{P_s M_w}}$  = 80.57 ft/sec  
 Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>s</sub>) (60) (A) = 16502 ACFM  
 Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528^\circ R}{T_{sr}}\right) \left(\frac{P_s}{29.92}\right) \left(\frac{100 - \% H_2O}{100}\right)$  = 15053 DSCFM

PLANT LP  
 DATE 12/14/81 1300  
 LOCATION Inlet  
 STACK DIMENSIONS 54.5 AREA \_\_\_\_\_ ft<sup>2</sup>  
 BAROMETRIC PRESSURE, P<sub>b</sub> = 30.04 in. Hg  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± -4.7 in. H<sub>2</sub>O  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> 28.9  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = \_\_\_\_\_  
 PITOT NO. \_\_\_\_\_ Cp = 0.84  
 TESTER \_\_\_\_\_

SCHMATIC OF TRAVERSE POINT LOCATION  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>s</sub> (°F)	± θ	Pitots reversed for Negative Flow?	√ΔP · COS θ
A	1	1.6		100			
	2	1.7		102			
	3	1.8		102			
	4	1.7		103			
	5	2.2		102			
	6	2.2		105			
	7	2.3		105			
	8	2.4		106			
B	9	1.5		102			
	2	1.5		108			
	3	2.2		109			
	4	2.2		108			
	5	2.2		108			
	6	1.9		108			
	7	1.9		108			
	8	1.8		108			
Average				T <sub>s</sub> = °F *			
				T <sub>sr</sub> = °R		*Avg. of absolute values including zeroes	

Absolute Gas Temperature, T<sub>sr</sub> = T<sub>s</sub> + 460°

Absolute Gas Pressure, P<sub>s</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 =

Gas Velocity, V<sub>s</sub> = (85.49) C<sub>p</sub> (√ΔP<sup>3</sup> · COS θ) avg  $\sqrt{\frac{T_{sr} \text{ avg}}{P_s M_w}}$  = \_\_\_\_\_ ft/sec

Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>s</sub>) (60) (A) = \_\_\_\_\_ ACFM

Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528^{\circ}R}{T_{sr}}\right) \left(\frac{P_s}{29.92}\right) \left(\frac{100 - \% H_2O}{100}\right)$  = \_\_\_\_\_ DSCFM





PLANT LP  
 DATE 12/4/85 1200  
 LOCATION OUTLET  
 STACK DIMENSIONS 77 1/4 AREA \_\_\_\_\_ ft<sup>2</sup>  
 BAROMETRIC PRESSURE, P<sub>b</sub> = 30.02 in. Hg  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± -0.42 in. H<sub>2</sub>O  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> 25.9  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = 10  
 PITOT NO. \_\_\_\_\_ Cp = 0.99  
 TESTER \_\_\_\_\_

**SCHEMATIC OF TRAVERSE POINT LOCATION**  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>sr</sub> (°F)	± θ	Pitots reversed for Negative Flow?	√ΔP · COS θ
A	1	0.44		230			
	2	0.55		231			
	3	0.55		232			
	4	0.60		230			
	5	0.60		231			
	6	0.61		235			
	7	0.61		236			
	8	0.65		237			
B	1	0.42		230			
	2	0.43		231			
	3	0.48		232			
	4	0.57		234			
	5	0.59		235			
	6	0.61		235			
	7	0.67		235			
	8	0.66		235			
Average				T <sub>s</sub> = °F *			
				T <sub>sr</sub> = °R	*Avg. of absolute values including zeroes		

Absolute Gas Temperature, T<sub>sr</sub> = T<sub>s</sub> + 460°  
 Absolute Gas Pressure, P<sub>s</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 = 30.01  
 Gas Velocity, V<sub>s</sub> = (85.49) C<sub>p</sub> (√ΔP · COS θ) avg  $\sqrt{\frac{T_{sr} \text{ avg}}{P_s M_w}}$  = 48.25 ft/sec  
 Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>s</sub>) (60) (A) = 9.882 ACFM  
 Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528^\circ R}{T_{sr}}\right) \left(\frac{P_s}{29.92}\right) \left(\frac{100 - \% H_2O}{100}\right)$  = 7513 DSCFM

PLANT LF  
 DATE 12/14/95 1300  
 LOCATION OUTLET  
 STACK DIMENSIONS 77/14 AREA \_\_\_\_\_ ft<sup>2</sup>  
 BAROMETRIC PRESSURE, P<sub>b</sub> = 30.02 in. Hg  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± -0.42 in. H<sub>2</sub>O  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> 28.9  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = 10  
 PITOT NO. \_\_\_\_\_ Cp = 0.84  
 TESTER \_\_\_\_\_

SCHMATIC OF TRAVERSE POINT LOCATION  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>s</sub> (°F)	± θ	Pitots reversed for Negative Flow?	√ΔP · COS θ
A	1	0.47		230			
	2	0.54		231			
	3	0.54		231			
	4	0.59		231			
	5	0.60		231			
	6	0.61		232			
	7	0.62		233			
	8	0.65		233			
B	1	0.42		233			
	2	0.48		236			
	3	0.49		236			
	4	0.56		236			
	5	0.60		236			
	6	0.61		237			
	7	0.66		237			
	8	0.66		235			
Average				T <sub>s</sub> = °F *			
				T <sub>SR</sub> = °R			*Avg. of absolute values including zeroes

Absolute Gas Temperature, T<sub>SR</sub> = T<sub>s</sub> + 460°

Absolute Gas Pressure, P<sub>s</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 = \_\_\_\_\_

Gas Velocity, V<sub>s</sub> = (85.49) C<sub>p</sub> (√ΔP<sup>3</sup> · COS θ) avg  $\sqrt{\frac{T_{SR} \text{ avg}}{P_s M_w}}$  = \_\_\_\_\_ ft/sec

Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>s</sub>) (60) (A) = \_\_\_\_\_ ACFM

Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528^\circ R}{T_{SR}}\right) \left(\frac{P_s}{29.92}\right) \left(\frac{100 - \% H_2O}{100}\right)$  = \_\_\_\_\_ DSCFM

PLANT LP  
 DATE 12/14/85  
 LOCATION OUTLET AREA 1400 ft<sup>2</sup>  
 STACK DIMENSIONS \_\_\_\_\_ in. Hg  
 BAROMETRIC PRESSURE, P<sub>b</sub> = \_\_\_\_\_ in. H<sub>2</sub>O  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± \_\_\_\_\_  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> \_\_\_\_\_  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = \_\_\_\_\_  
 PITOT NO. \_\_\_\_\_ Cp = \_\_\_\_\_  
 TESTER \_\_\_\_\_

SCHMATIC OF TRAVERSE POINT LOCATION  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>s</sub> (°F)	± θ	Pitots reversed for Negative Flow?	√ΔP · COS θ
	1	0.45		232			
	2	0.47		230			
	3	0.66		230			
	4	0.64		233			
	5	0.67		230			
	6	0.67		233			
	7	0.67		237			
	8	0.7		230			
	9	0.60		232			
	2	0.52		232			
	3	0.63		240			
	4	0.69		241			
	5	0.66		241			
	6	0.62		241			
	7	0.59		241			
	8	0.60		241			
Average				T <sub>s</sub> = °F *	*Avg. of absolute values including zeroes		
				T <sub>sr</sub> = °R			

Absolute Gas Temperature, T<sub>sr</sub> = T<sub>s</sub> + 460°

Absolute Gas Pressure, P<sub>s</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 = \_\_\_\_\_

Gas Velocity, V<sub>s</sub> = (85.49) C<sub>p</sub> (√ΔP · COSθ) avg  $\sqrt{\frac{T_{sr} \text{ avg}}{P_s M_w}}$  = \_\_\_\_\_ ft/sec

Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>s</sub>) (60) (A) = \_\_\_\_\_ ACFM

Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528 \text{ OR}}{T_{sr}}\right) \left(\frac{P_s}{29.92}\right) \left(\frac{100 - \% \text{ H}_2\text{O}}{100}\right)$  = \_\_\_\_\_ DSCFM

Imp - weight

T-1 Inlet

Before 3171

after 3184  
13

Outlet

3187

3207 / 20

T-2 Inlet

Before 3126

after 3139  
13

Outlet

3185

3205 / 20

T-3 Inlet

Before 3109

after 3123  
14

Outlet

3113

3136 / 23

PLANT LP  
 DATE 12/14/95 1300  
 LOCATION 007115  
 STACK DIMENSIONS 7.714 AREA \_\_\_\_\_ ft<sup>2</sup>  
 BAROMETRIC PRESSURE, P<sub>b</sub> = 30.02 in. Hg  
 STACK STATIC PRESSURE, P<sub>g</sub> = ± -0.42 in. H<sub>2</sub>O  
 STACK GAS MOLECULAR WEIGHT (Wet), M<sub>w</sub> 28.9  
 STACK GAS MOISTURE CONTENT, % H<sub>2</sub>O = 10  
 PITOT NO. \_\_\_\_\_ Cp = 0.84  
 TESTER \_\_\_\_\_

SCHEMATIC OF TRAVERSE POINT LOCATION  
 + θ Clockwise  
 Cyclonic Flow Angle - θ Counterclockwise

PORT	POINT	ΔP Inch H <sub>2</sub> O	√ΔP	T <sub>SR</sub> (°F)	± θ	Pitots reversed for Negative Flow?	√ΔP · COS θ
A	1	0.47		230			
	2	0.54		231			
	3	0.54		231			
	4	0.59		231			
	5	0.60		231			
	6	0.61		232			
	7	0.62		233			
	8	0.65		233			
B	1	0.42		233			
	2	0.48		236			
	3	0.49		236			
	4	0.56		236			
	5	0.60		236			
	6	0.61		237			
	7	0.66		237			
	8	0.66		235			
Average				T <sub>S</sub> = °F *			
				T <sub>SR</sub> = °R			*Avg. of absolute values including zeroes

Absolute Gas Temperature, T<sub>SR</sub> = T<sub>S</sub> + 460°  
 Absolute Gas Pressure, P<sub>S</sub> = P<sub>b</sub> + P<sub>g</sub>/13.6 = \_\_\_\_\_  
 Gas Velocity, V<sub>S</sub> = (85.49) C<sub>p</sub> (√ΔP · COS θ) avg  $\sqrt{\frac{T_{SR} \text{ avg}}{P_S M_w}}$  = \_\_\_\_\_ ft/sec

Actual Gas Flowrate, Q<sub>a</sub> = (V<sub>S</sub>) (60) (A) = \_\_\_\_\_ ACFM  
 Standard Gas Flowrate, Q<sub>std</sub> = Q<sub>a</sub>  $\left(\frac{528^{\circ}R}{T_{SR}}\right) \left(\frac{P_S}{29.92}\right) \left(\frac{100 - \% H_2O}{100}\right)$  = \_\_\_\_\_ DSCFM



Imp. weight

T-1 Inlet

Before 3171

after 3184  

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13

Outlet

3187

3207  

---

20

T-2 Inlet

Before 3126

after 3139  

---

13

Outlet

3185

3205  

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20

T-3 Inlet

Before 3109

after 3123  

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14

Outlet

3113

3136  

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23

**TRC Environmental Corporation**  
CEM Data Sheet

Firm: L Pacific  
 Location: Houlton, ME  
 Tester: S. Paterson  
 Test No.: Inlet-1  
 Sample Loc.: RTO Inlet Duct  
 Date: December 14, 1995  
 TIME: 1035-1055

Ambient Temp, deg. F = 10  
 MEL Temp, deg. F = 70  
 Bar. Pressure, in Hg = 30.28  
 Vacuum Gauge = 10  
 Pressure Gauge =  
 % Moisture =

Calibration Gases			Tank ID
Mid Cal	High Cal	Mid	High
9.1	20.8	ALM037884	ALM057653
142	451	ALM010642	AAL 20965
49.5	88.6	ALM43797	ALM43127

	Calibration Data				Emissions Summary	
	Initial Values		Final Values		Analyzer Units & Range	Corrected Effluent Gas Conc.
	(Rack) Analyzer Cal. Response	System Cal. Bias % of Span	System Cal. Response	System Cal. Bias % of Span		
O <sub>2</sub>	0.1	0.1	0.14	0.3	%	-
Upscale	20.9	20.8	20.78	-0.5	25	20.8
CO <sub>2</sub>				0.0	%	-
Upscale				0.0	20	ERR
CO	0.3	0.6	-5.5	-1.2	ppm	-
Upscale	451.8	448.3	437.3	-2.9	500	13.6
SO <sub>2</sub>				0.0	ppm	-
Upscale				0.0	1000	ERR
NOX	0.2	0.9	1.5	1.3	ppm	-
Upscale	89.2	89.2	90.8	1.6	100	3.0
THC				0	ppm	-
Upscale				0	100	ERR
		<b>LIMITS</b>				
		+/- 5%		+/- 3%		

Comments:  
 NOx (lb/hr) = K x ppm x MW x Flow  
 NOx (lb/hr) = 0.19  
 CO (lb/hr) = K x ppm x MW x Flow  
 CO (lb/hr) = 0.76  
 K = 2.59E-09  
 MW for NOx = 46.01 g/gmol  
 Flow (dscfm) = 15,028  
 MW for CO = 20.00 g/gmol

	ZERO Cal. Gas Analyzer Response	Analyzer Calib. Error	MID Cal. Gas Analyzer Response	Analyzer Calib. Error	HIGH Cal. Gas Analyzer Response	Analyzer Calib. Error
O <sub>2</sub>	0.1	0.2	9.1	0.2	20.9	0.4
CO <sub>2</sub>		0.0		0.0		0.0
CO	0.3	0.1	140.7	-0.3	451.8	0.2
SO <sub>2</sub>		0.0		0.0		0.0
NOX	0.2	0.2	47.6	-1.9	89.2	0.6
THC		0.0		0.0		0.0
<b>LIMIT</b>		+/- 2%		+/- 2%		+/- 2%

**TRC Environmental Corporation**  
CEM Data Sheet

Firm: L Pacific  
 Location: Houston, ME  
 Tester: S. Paterson  
 Test No.: Inlet-3  
 Sample Loc.: RTO Inlet Duct  
 Date: December 14, 1995  
 TIME: 13:38 - 13:58

Ambient Temp, deg. F = 10  
 MEL Temp, deg. F = 70  
 Bar. Pressure, in Hg = 30.28  
 Vacuum Gauge = 10  
 Pressure Gauge =  
 % Moisture =

Calibration Gases		
Mid Cal	High Cal	Tank ID
9.1	20.8	ALM037884 / ALM057653
142	451	ALM010642 AAL 20965
49.5	88.6	ALM43797 / ALM43127

	Calibration Data						Emissions Summary
	Initial Values			Final Values			
	(Rack) Analyzer Cal. Response	System Cal. Bias % of Span	System Cal. Response	System Cal. Bias % of Span	System Cal. Response	Drift % of Span	
O <sub>2</sub>	0.1	0.2	0.4	0.13	0.3	-0.1	
Upscale	20.9	20.7	-0.7	20.68	-0.9	-0.2	25
Zero			0.0		0.0	0.0	
Upscale			0.0		0.0	0.0	
Zero	0.3	1.4	0.2	-1.7	-0.4	-0.6	20
Upscale	450.7	448.3	-0.5	442.3	-1.7	-1.2	500
Zero			0.0		0.0	0.0	
Upscale			0.0		0.0	0.0	
Zero	0.2	2.2	2.0	2.2	2.0	0.0	1000
Upscale	89.2	91.9	2.7	91.8	2.6	-0.1	100
Zero			0		0	0	
Upscale			0		0	0	
LIMITS			+/- 5%			+/- 3%	

O<sub>2</sub>  
CO<sub>2</sub>  
CO  
SO<sub>2</sub>  
NOx  
THC

ZERO	MID	HIGH
Cal. Gas Analyzer Response	Cal. Gas Analyzer Response	Cal. Gas Analyzer Response
0.1	9.1	20.9
0.3	140.6	450.7
0.2	47.6	89.2
LIMIT		
+/- 2%		

O<sub>2</sub>  
CO<sub>2</sub>  
CO  
SO<sub>2</sub>  
NOx  
THC

Comments:  
 NOx (lb/hr) = K x ppm x MW x Flow  
 NOx (lb/hr) = 0.10  
 O (lb/hr) = K x ppm x MW x Flow  
 O (lb/hr) = 0.29  
 = 2.59E-09  
 MW for NOx = 46.01 g/mol  
 Flow (dscfm) = 15,143  
 MW for CO = 20.00 g/mol