# IXM MANUFACTURING PROCESSES POLYMERS STACK EMISSIONS TEST REPORT TEST DATES: 17-18 JANUARY 2019

# THE CHEMOURS COMPANY FAYETTEVILLE, NORTH CAROLINA

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#### 1. INTRODUCTION

#### 1.1 FACILITY AND BACKGROUND INFORMATION

The Chemours Fayetteville Works (Chemours) is located in Bladen County, North Carolina, approximately 10 miles south of the city of Fayetteville. The Chemours operating areas on the site include the Fluoromonomers, IXM and Polymers Processing Aid (PPA) manufacturing areas, Wastewater Treatment, and Powerhouse.

Chemours contracted Weston Solutions, Inc. (Weston) to perform HFPO Dimer Acid Fluoride, captured as HFPO Dimer Acid emission testing on the Polymers Stack. Testing was performed on 17-18 January 2019 and generally followed the "Emission Test Protocol" reviewed and approved by the North Carolina Department of Environmental Quality (NCDEQ). This report provides the results from the emission test program.

#### 1.2 TEST OBJECTIVES

The specific objectives for this test program were as follows:

- Measure the emissions concentrations and mass emissions rates of HFPO Dimer Acid Fluoride from the Polymers stack which is located in the IXM processes.
- Monitor and record process data in conjunction with the test program.
- Provide representative emissions data.

#### 1.3 TEST PROGRAM OVERVIEW

During the emissions test program, the concentrations and mass emissions rates of HFPO Dimer Acid Fluoride were measured on the Polymers stack.

Table 1-1 provides a summary of the test location and the parameters that were measured along with the sampling/analytical procedures that were followed.

Section 2 provides a summary of test results. A description of the processes is provided in Section 3. Section 4 provides a description of the test locations. The sampling and analytical procedures are provided in Section 5. Detailed test results and discussion are provided in Section 6.

Appendix C includes the summary reports for the laboratory analytical results. The full laboratory data packages are provided in electronic format and on CD with each hard copy.

Table 1-1
Sampling Plan for Polymers Stack

Sampling Point & Location		Poly	mers Stack						
Number of Tests:	3								
Parameters To Be Tested:	HFPO Dimer Acid Fluoride (HFPO-DAF)	Volumetric Flow Rate and Gas Velocity	Carbon Dioxide	Oxygen	Water Content				
Sampling or Monitoring Method	EPA M-0010	EPA M1, M2, M3A, and M4 in conjunction with M-0010 tests	EPA I	M3A	EPA M4 in conjunction with M-0010 tests				
Sample Extraction/ Analysis Method(s):	LC/MS/MS	NA <sup>6</sup>	N/	A	NA				
Sample Size	$> 1 \text{m}^3$	NA	NA	NA	NA				
Total Number of Samples Collected <sup>1</sup>	3	3	3	3	3				
Reagent Blanks (Solvents, Resins) <sup>1</sup>	1 set	0	0	0	0				
Field Blank Trains <sup>1</sup>	1 per source	0	0	0	0				
Proof Blanks <sup>1</sup>	1 per train	0	0	0	0				
Trip Blanks <sup>1,2</sup>	1 set	0	0	0					
Lab Blanks	1 per fraction <sup>3</sup>	0	0	0	0				
Laboratory or Batch Control Spike Samples (LCS)	1 per fraction <sup>3</sup>	0	0	0	0				
Laboratory or Batch Control Spike Sample Duplicate (LCSD)	1 per fraction <sup>3</sup>	0	0	0	0				
Media Blanks	1 set <sup>4</sup>	0	0	0	0				
Isotope Dilution Internal Standard Spikes	Each sample	0	0	0	0				
Total No. of Samples	75	3	3	3	3				

#### Key:

<sup>&</sup>lt;sup>1</sup> Sample collected in field.

<sup>&</sup>lt;sup>2</sup> Trip blanks include one XAD-2 resin module and one methanol sample per sample shipment.

<sup>&</sup>lt;sup>3</sup> Lab blank and LCS/LCSD includes one set per analytical fraction (front half, back half and condensate).

 $<sup>^{\</sup>rm 4}$  One set of media blank archived at laboratory at media preparation.

<sup>&</sup>lt;sup>5</sup> Actual number of samples collected in field.

<sup>&</sup>lt;sup>6</sup> Not applicable.

#### 2. SUMMARY OF TEST RESULTS

A total of three test runs were performed on the Polymers Stack. Table 2-1 provides a summary of the HFPO Dimer Acid emission test results. Detailed test results summaries are provided in Section 6.

It is important to note that emphasis is being placed on the characterization of the emissions based on the stack test results. Research conducted in developing the protocol for stack testing HFPO Dimer Acid Fluoride, HFPO Dimer Acid Ammonium Salt and HFPO Dimer Acid realized that the resulting testing, including collection of the air samples and extraction of the various fraction of the sampling train, would result in all three compounds being expressed as simply the HFPO Dimer Acid. However, it should be understood that the total HFPO Dimer Acid results provided on Table 2-1 and in this report include a percentage of each of the three compounds.

Table 2-1
Summary of HFPO Dimer Acid Test Results

Course	Dun No	Emission Rates					
Source	Run No.	lb/hr	g/sec				
	1	1.81E-04	2.28E-05				
Dalymana Staalr	2	1.51E-04	1.90E-05				
Polymers Stack	3	1.55E-04	1.95E-05				
	Average <sup>2</sup>	1.62E-04	2.04E-05				

#### 3. PROCESS DESCRIPTIONS

The IXM area is included in the scope of this test program.

#### 3.1 POLYMERS

The Polymers area consists of a polymerization process, finishing and recycle. There are two types of polymer produced, using products made in the Fluoromonomers and IXM Precursors areas: SR polymer and CR polymer. Both SR and CR polymerization processes take place in a solvent. The reaction is initiated and sustained by continuous addition of Dimer Peroxide initiator. There is a Recycle Still that takes solution and removes any impurities, allowing the solution to be used again. The finishing area takes the polymer produced during polymerization and transforms it into pellets.

#### 3.2 PROCESS OPERATIONS AND PARAMETERS

Source	Operation/Product	Batch or Continuous
Polymers Stack	SR Polymer	Continuous – Polymerization Batch – Recycle Still Batch – Line Four extrusion

During the test program, operations parameters were monitored by Chemours and are included in Appendix A.

4. DESCRIPTION OF TEST LOCATIONS

4.1 POLYMERS STACK

The Polymers stack is a 30-inch ID fiberglass stack located near the roof edge. Vent lines enter

the stack at various points and a significant straight run of vertical stack without flow

disturbances is not available. Two sample ports are installed in the stack 30 inches down from

the stack exit and 58 inches up from the last vent line entry point. Per EPA Method 1, 24 traverse

points, 12 per port, were used for sampling.

See Figure 4-1 for a schematic of the test port and traverse point locations.

Note: All measurements at the test location were confirmed prior to sampling.

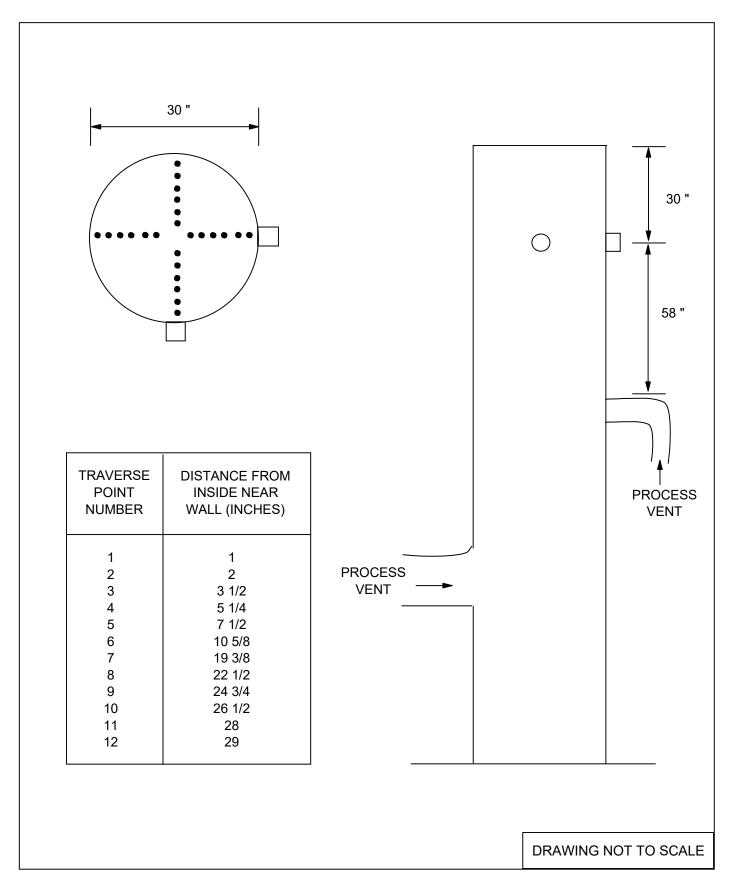


FIGURE 4-1
POLYMERS STACK TEST PORT
AND TRAVERSE POINT LOCATIONS

#### 5. SAMPLING AND ANALYTICAL METHODS

#### 5.1 STACK GAS SAMPLING PROCEDURES

The purpose of this section is to describe the stack gas emissions sampling trains and to provide details of the stack sampling and analytical procedures utilized during the emissions test program.

#### 5.1.1 Pre-Test Determinations

Preliminary test data were obtained at the test location. Stack geometry measurements were measured and recorded, and traverse point distances verified. A preliminary velocity traverse was performed utilizing a calibrated S-type pitot tube and an inclined manometer to determine velocity profiles. Flue gas temperatures were observed with a calibrated direct readout panel meter equipped with a chromel-alumel thermocouple. Preliminary water vapor content was estimated by wet bulb/dry bulb temperature measurements.

A check for the presence or absence of cyclonic flow was previously conducted at the test location. The cyclonic flow checks were negative ( $< 20^{\circ}$ ) verifying that the source was acceptable for testing.

Preliminary test data was used for nozzle sizing and sampling rate determinations for isokinetic sampling procedures.

Calibration of probe nozzles, pitot tubes, metering systems, and temperature measurement devices was performed as specified in Section 5 of EPA Method 5 test procedures.

#### 5.2 STACK PARAMETERS

#### 5.2.1 EPA Method 0010

The sampling train utilized to perform the HFPO Dimer Acid sampling was an EPA Method 0010 train (see Figure 5-1). The Method 0010 consisted of a borosilicate nozzle that attached directly to a heated borosilicate probe. In order to minimize possible thermal degradation of the HFPO Dimer Acid, the probe and particulate filter were heated above stack temperature to minimize water vapor condensation before the filter. The probe was connected directly to a heated borosilicate filter holder containing a solvent extracted glass fiber filter.

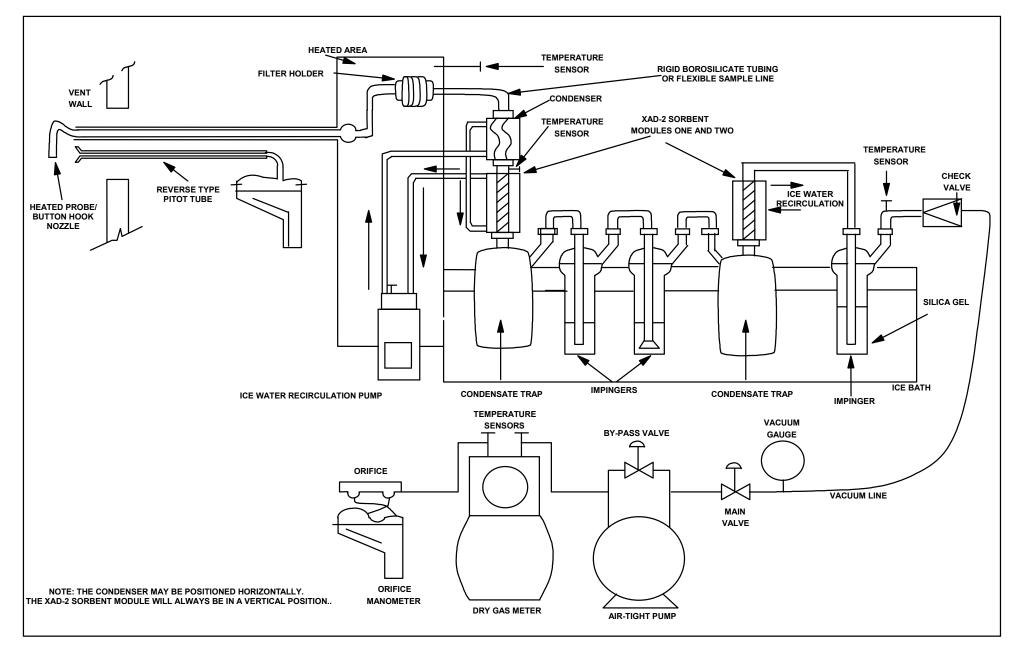


FIGURE 5-1
EPA METHOD 0010 SAMPLING TRAIN

A section of borosilicate glass or flexible polyethylene tubing connected the filter holder exit to a Grahm (spiral) type ice water-cooled condenser, an ice water-jacketed sorbent module containing approximately 40 grams of XAD-2 resin. The XAD-2 resin tube was equipped with an inlet temperature sensor. The XAD-2 resin trap was followed by a condensate knockout impinger and a series of two impingers that contained 100 milliliters of high purity distilled water. The train also included a second XAD-2 resin trap behind the impinger section to evaluate possible sampling train breakthrough. Each XAD-2 resin trap was connected to a 1-liter condensate knockout trap. The final impinger contained 300 grams of dry pre-weighed silica gel. All impingers and the condensate traps were maintained in an ice bath. Ice water was continuously circulated in the condenser and both XAD-2 modules to maintain method-required temperature. A control console with a leakless vacuum pump, a calibrated orifice, and dual inclined manometers was connected to the final impinger via an umbilical cord to complete the sample train.

HFPO Dimer Acid Fluoride (CAS No. 2062-98-8) that is present in the stack gas is expected to be captured in the sampling train along with HFPO Dimer Acid (CAS No. 13252-13-6). HFPO Dimer Acid Fluoride undergoes hydrolysis instantaneously in water in the sampling train and during the sample recovery step and will be converted to HFPO Dimer Acid such that the amount of HFPO Dimer Acid emissions represents a combination of both HFPO Dimer Acid Fluoride and HFPO Dimer Acid.

During sampling, gas stream velocities were measured by attaching a calibrated S-type pitot tube into the gas stream adjacent to the sampling nozzle. The velocity pressure differential was observed immediately after positioning the nozzle at each traverse point, and the sampling rate adjusted to maintain isokineticity  $\pm$  10. Flue gas temperature was monitored at each point with a calibrated panel meter and thermocouple. Isokinetic test data was recorded at each traverse point during all test periods, as appropriate. Leak checks were performed on the sampling apparatus according to reference method instructions, prior to and following each run, component change (if required), or during midpoint port changes.

### 5.2.2 EPA Method 0010 Sample Recovery

At the conclusion of each test, the sampling train was dismantled, the openings sealed, and the components transported to the field laboratory trailer for recovery.

A consistent procedure was employed for sample recovery:

- 1. The two XAD-2 covered (to minimize light degradation) sorbent modules (1 and 2) were sealed and labeled.
- 2. The glass fiber filter(s) were removed from the holder with tweezers and placed in a polyethylene container along with any loose particulate and filter fragments.
- 3. The particulate adhering to the internal surfaces of the nozzle, probe and front half of the filter holder were rinsed with a solution of methanol and ammonium hydroxide into a polyethylene container while brushing a minimum of three times until no visible particulate remained. Particulate adhering to the brush was rinsed with methanol/ammonium hydroxide into the same container. The container was sealed.
- 4. The volume of liquid collected in the first condensate trap was measured, the value recorded, and the contents poured into a polyethylene container.
- 5. All train components between the filter exit and the first condensate trap were rinsed with methanol/ammonium hydroxide. The solvent rinse was placed in a separate polyethylene container and sealed.
- 6. The volume of liquid in the impingers one, two, and second condensate trap were measured, the values recorded, and sample was placed in the same container as step 4 above and sealed.
- 7. The two impingers, condensate trap, and connectors were rinsed with methanol/ammonium hydroxide. The solvent sample was placed in a separate polyethylene container and sealed.
- 8. The silica gel in the final impinger was weighed and the weight gain value recorded.
- 9. Site (reagent) blank samples of the methanol/ammonium hydroxide, XAD resin, filter and distilled water were retained for analysis.

Each container was labeled to clearly identify its contents. The height of the fluid level was marked on the container of each liquid sample to provide a reference point for a leakage check during transport. All samples were maintained cool.

During each test campaign, an M-0010 blank train was setup near the test location, leak checked and recovered along with the respective sample train. Following sample recovery, all samples were transported to the TestAmerica Inc. for sample extraction and analysis.

See Figure 5-2 for a schematic of the M-0010 sample recovery process.

#### 5.2.3 EPA Method 0010 - Sample Analysis

Method 0010 sampling trains resulted in four separate analytical fractions for HFPO Dimer Acid analysis according to SW-846 Method 3542:

- Front-Half Composite—comprised of the Particulate Filter, and the probe, nozzle, and front-half of the filter holder solvent rinses,
- Back-Half Composite—comprised of the first XAD-2 resin material and the back-half of the filter holder with connecting glassware solvent rinses,
- Condensate Composite—comprised of the aqueous condensates and the contents of impingers one and two with solvent rinses,
- Breakthrough XAD-2 Resin Tube—comprised of the resin tube behind the series of impingers.

The second XAD-2 resin material was analyzed separately to evaluate any possible sampling train HFPO-DA breakthrough.

The front-half and back-half composites and the second XAD-2 resin material were placed in polypropylene wide-mouth bottles and tumbled with methanol containing 5% NH4OH for 18 hours. Portions of the extracts were processed analytically for the HFPO dimer acid by liquid chromatography and duel mass spectroscopy (HPLC/MS/MS). The Condensate composite was concentrated onto a solid phase extraction (SPE) cartridge followed by desorption from the cartridge using methanol. Portions of those extracts were also processed analytically by HPLC/MS/MS.

Samples were spiked with isotope dilution internal standard (IDA) at the commencement of their preparation to provide accurate assessments of the analytical recoveries. Final data was corrected for IDA standard recoveries.

TestAmerica Laboratories, Inc. (TestAmerica) developed detailed procedures for the sample extraction and analysis for HFPO Dimer Acid. These procedures were incorporated into the test protocol.

#### 5.3 GAS COMPOSITION

The Weston mobile laboratory equipped with instrumental analyzers was used to measure carbon dioxide (CO<sub>2</sub>) and oxygen (O<sub>2</sub>) concentrations. A diagram of the Weston sampling system is presented in Figure 5-3.

The sample was collected at the exhaust of the Method 0010 sampling system. At the end of the line, a tee permitted the introduction of calibration gas. The sample was drawn through a heated Teflon® sample line to the sample conditioner. The output from the sampling system was recorded electronically, and one-minute averages were recorded and displayed on a data logger.

Each analyzer was set up and calibrated internally by introduction of calibration gas standards directly to the analyzer from a calibration manifold. The calibration manifold is designed with an atmospheric vent to release excess calibration gas and maintains the calibration at ambient pressure. The direct calibration sequence consisted of alternate injections of zero and mid-range gases with appropriate adjustments until the desired responses were obtained. The high-range standards were then introduced in sequence without further adjustment.

The sample line integrity was verified by performing a bias test before and after each test period. The sampling system bias test consisted of introducing the zero gas and one up-range calibration standard in excess to the valve at the probe end when the system was sampling normally. The excess calibration gas flowed out through the probe to maintain ambient sampling system pressure. Calibration gas supply was regulated to maintain constant sampling rate and pressure. Instrument bias check response was compared to internal calibration responses to ensure sample line integrity and to calculate a bias correction factor after each run using the ratio of the measured concentration of the bias gas certified by the calibration gas supplier.

The oxygen and carbon dioxide content of each stack gas was measured according to EPA Method 3A procedures which incorporate the latest updates of EPA Method 7E. A Servomex Model 4900 analyzer (or equivalent) was used to measure oxygen content. A Servomex Model 4900 analyzer (or equivalent) was used to measure carbon dioxide content of the stack gas. Both analyzers were calibrated with EPA Protocol gases prior to the start of the test program and performance was verified by sample bias checks before and after each test run.

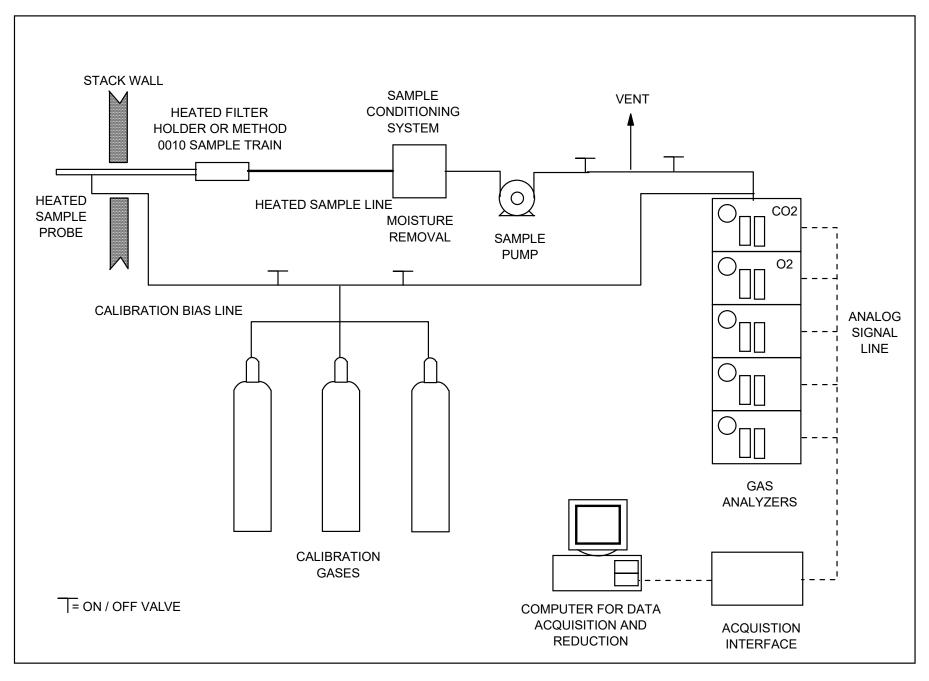


FIGURE 5-3
WESTON SAMPLING SYSTEM

### 6. DETAILED TEST RESULTS AND DISCUSSION

Preliminary testing and the associated analytical results required significant sample dilution to bring the HFPO Dimer Acid concentration within instrument calibration; therefore, sample times and sample volumes were reduced for the formal test program. This was approved by the North Carolina Department of Environmental Quality (NCDEQ).

Each test was a minimum of 96 minutes in duration. A total of three test runs were performed on the Polymers Stack.

Table 6-1 provides detailed test data and test results for the Polymers Stack.

The Method 3A sampling indicated that the O<sub>2</sub> and CO<sub>2</sub> concentrations were at ambient air levels (20.9% O<sub>2</sub>, 0% CO<sub>2</sub>), therefore, 20.9% O<sub>2</sub> and 0% CO<sub>2</sub> values were used in all calculations.

### TABLE 6-1 CHEMOURS - FAYETTEVILLE, NC SUMMARY OF HFPO DIMER ACID TEST DATA AND TEST RESULTS

Test Data			
Run number	1	2	3
Location	Polymers Stack	Polymers Stack	Polymers Stack
Date	1/17/2019	1/18/2019	1/18/2019
Time period	1443-1641	0835-1028	1111-1315
SAMPLING DATA:			
Sampling duration, min.	96.0	96.0	96.0
Nozzle diameter, in.	0.218	0.218	0.218
Cross sectional nozzle area, sq.ft.	0.000259	0.000259	0.000259
Barometric pressure, in. Hg	30.10	30.01	30.01
Avg. orifice press. diff., in H <sub>2</sub> O	1.31	1.43	1.50
Avg. dry gas meter temp., deg F	64.7	49.4	61.7
Avg. abs. dry gas meter temp., deg. R	525	509	522
Total liquid collected by train, ml	25.6	9.0	11.2
Std. vol. of H <sub>2</sub> O vapor coll., cu.ft.	1.2	0.4	0.5
Dry gas meter calibration factor	1.0069	1.0069	1.0069
Sample vol. at meter cond., dcf	57.360	58.671	60.644
Sample vol. at std. cond., dscf <sup>(1)</sup>	58.636	61.606	62.193
Percent of isokinetic sampling	97.2	95.4	94.9
GAS STREAM COMPOSITION DATA:			
CO <sub>2</sub> , % by volume, dry basis	0.0	0.0	0.0
O <sub>2</sub> , % by volume, dry basis	20.9	20.9	20.9
N <sub>2</sub> , % by volume, dry basis	79.1	79.1	79.1
Molecular wt. of dry gas, lb/lb mole	28.84	28.84	28.84
H <sub>2</sub> 0 vapor in gas stream, prop. by vol.	0.020	0.007	0.008
Mole fraction of dry gas	0.980	0.993	0.992
Molecular wt. of wet gas, lb/lb mole	28.62	28.76	28.74
GAS STREAM VELOCITY AND VOLUMETRIC I	FLOW DATA:		
Static pressure, in. H <sub>2</sub> O	-0.24	-0.24	-0.24
Absolute pressure, in. Hg	30.08	29.99	29.99
Avg. temperature, deg. F	62	56	62
Avg. absolute temperature, deg.R	522	516	522
Pitot tube coefficient	0.84	0.84	0.84
Total number of traverse points	24	24	24
Avg. gas stream velocity, ft./sec.	40.6	42.5	43.7
Stack/duct cross sectional area, sq.ft.	4.91	4.91	4.91
Avg. gas stream volumetric flow, wacf/min.	11957	12517	12872
Avg. gas stream volumetric flow, dscf/min. (1)	11910	12743	12928

 $<sup>^{(1)}</sup>$  Standard conditions = 68 deg. F. (20 deg. C.) and 29.92 in Hg (760 mm Hg)

# TABLE 6-1(cont.) CHEMOURS - FAYETTEVILLE, NC SUMMARY OF HFPO DIMER ACID TEST DATA AND TEST RESULTS

T	EST	DA	TA

Run number Location Date Time period	1 Polymers Stack 1/17/2019 1443-1641	2 Polymers Stack 1/18/2019 0835-1028	3 Polymers Stack 1/18/2019 1111-1315
LABORATORY REPORT DATA, ug. HFPO Dimer Acid	6.75	5.51	5.63
EMISSION RESULTS, ug/dscm. HFPO Dimer Acid	4.06	3.15	3.20
EMISSION RESULTS, lb/dscf. HFPO Dimer Acid	2.54E-10	1.97E-10	2.00E-10
EMISSION RESULTS, lb/hr. HFPO Dimer Acid	1.81E-04	1.51E-04	1.55E-04
EMISSION RESULTS, g/sec. HFPO Dimer Acid	2.28E-05	1.90E-05	1.95E-05

# APPENDIX A PROCESS OPERATIONS DATA

Date 1/17/2019

Time	1400 1500							16	1600			
Stack Testing						RUI	N 1 - 1	443-1	641			
Recycle Still				Star	ting u	p to d	istill S	R Solu	tion			
Polymerization				Stan	dard S	R Poly	ymer I	Produ	ction			
Line 4 Extrusion												
Line 3 Extrusion	Extrud CR 10											

Date 1/18/2019

Time		8	00			9	00			10	00			1100 1200						1300			
Stack Testing		RUN 2 - 0835-1028 RUN 3 - 1111-1315																					
Recycle Still									Di	stilling	SR Sc	lution	, takir	ng off	to was	ite							
Polymerization										Stan	dard S	R Poly	mer l	Produ	ction								
Line 4 Extrusion																							
Line 3 Extrusion																							

# APPENDIX B RAW AND REDUCED TEST DATA

# CHEMOURS - FAYETTEVILLE, NC INPUTS FOR HFPO DIMER ACID CALCULATIONS

Test Data			
Run number	1	2	3
Location	Polymers Stack	Polymers Stack	Polymers Stack
Date	1/17/2019	1/18/2019	1/18/2019
Time period	1443-1641	0835-1028	1111-1315
Operator	MW	MW	MW
Inputs For Calcs.			
Sq. rt. delta P	0.72582	0.76494	0.78166
Delta H	1.3125	1.4321	1.5013
Stack temp. (deg.F)	62.0	56.1	62.3
Meter temp. (deg.F)	64.7	49.4	61.7
Sample volume (act.)	57.360	58.671	60.644
Barometric press. (in.Hg)	30.10	30.01	30.01
Volume H <sub>2</sub> O imp. (ml)	6.0	-4.0	-3.0
Weight change sil. gel (g)	19.6	13.0	14.2
% CO <sub>2</sub>	0.0	0.0	0.0
% O <sub>2</sub>	20.9	20.9	20.9
% N <sub>2</sub>	79.1	79.1	79.1
Area of stack (sq.ft.)	4.910	4.910	4.910
Sample time (min.)	96.0	96.0	96.0
Static pressure (in.H <sub>2</sub> O)	-0.24	-0.24	-0.24
Nozzle dia. (in.)	0.218	0.218	0.218
Meter box cal.	1.0069	1.0069	1.0069
Cp of pitot tube	0.84	0.84	0.84
Traverse points	24	24	24

2/19/2019 3:02 PM 011719 polymer

### Sample and Velocity Traverse Point Data Sheet - Method 1

Clie Loaction/Pla Sour	ant	Chemours Fayethealle Polymers	<del></del>	Оря W.O. N	erator <u>SK</u> Date <u>3/2//18</u> Jumber
Duct Type		Circular		Rectangular Duct	indicate appropriate type
Traverse Type		Particulate Traverse		Velocity Traverse	☐ CEM Traverse

Distance from far wall to outside of port (in.) = C	48
Port Depth (in.) = D	14
Depth of Duct, diameter (in.) = C-D	30
Area of Duct (ft <sup>2</sup> )	4.91
Total Traverse Points	24
Total Traverse Points per Port	17
Port Diameter (in.) —(Flange-Threaded-Hole)	
Monorail Length	
Rectangular Ducts Only	
Width of Duct, rectangular duct only (in.)	
Total Ports (rectangular duct only)	
Equivalent Diameter = (2°L*W)/(L+W)	

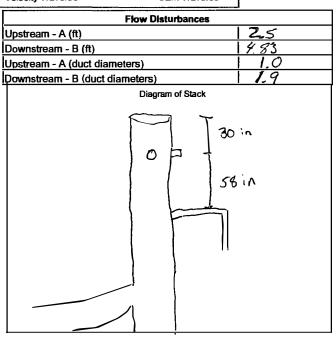
Traverse Point Locations											
		Distance from									
Traverse		Inside Duct	Distance from Outside of								
Point	% of Duct	Wall (in)	Port (in)								
1	21		19								
2	67	2	70								
3	119	3/2	21/12								
4	17.7	51/4	23 1/4								
5	25	7/2	251/2								
6	356	10 5/8	24 5/8								
7	64.4	193/8	37 3/8								
8	75	221/2	40 1/2								
9	82.3	2434	42 3/4								
10	S8 2	761/2	44								
11	933	28	46								
12	97.9	79	47								
CEM	3 Point(Long L	Annument Line) Str	dification Point Locations								
1	0.167										
2	0.50										
3	0.833										
N	ote: If stack	dia < 12 inch us	e EPA Method 1A								

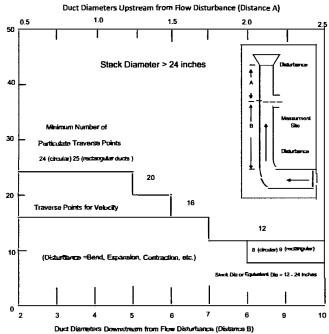
(Sample port upstream of pitot port)

Note: If stack dia >24" then adjust traverse point to 1 inch from wall

If stack dia <24" then adjust traverse point to 0.5 inch from wall

		$\Box$	Traverse Point Location Percent of Stack -Circular														
					_			N	umb	er of Tra	verse	Points					
		$\Box$	l	2	Π	3	4	Π	5	6	7	8	9	T	10	11	12
Т	1	1		14.6			6.7	Ι		4.4		3.2 [		ī	2.6		2.1
r	2	T		85.4			25	Γ		14.6		10.5		1	8.2	1	6.7
a	3			1	Γ	_	75	1		29.6		19.4		ī	14.6		11.8
V L	4	T			Г		93.3	Г		70.4		32.3		T	22.6		17.7
1, 0	15	Π			ī			Ī		85.4		67.7		ī	34.2		25
8 8	6	1		I				1		95.6	1	80.6		Ţ	65.8	l.	35.6
e t	7	Т		l .	Π			Ι				89.5		T	77.4	ļ	64.4
i	<u> 8</u>	Τ.						ı				96.8		T	85.4		75
0 1	T 9	П		l	ı			ı						1	91.8		82.3
1	10	ī			1			ı				1 1		1	97.4	l .	88.2
n	11	ī			Ī			1						ī			93.3
t	12	T		I	Π			1			l	1 1		T			97.9





		Traverse Point Location Percent of Stack -Rectangular											
		Number of Traverse Points											
	Г	1	2	3	4	l 5	6	7	8	9	10	111	12
т   1	Т		25.0	16.7	12.5	10.0	8.3	7.1	6.3	5.6	5.0	4.5	4.2
r   2	1	1	75.0	50.0	37.5	30.0	25.0	21.4	18.8	16.7	15.0	13.6	12.5
1 3				83.3	62.5	50.0	41.7	35.7	31.3	27.8	25.0	22.7	20.8
4					87.5	70.0	58.3	50.0	43.8	38.9	35.0	31.8	29.2
5	Т					90.0	75.0	64.3	56.3	50.0	45.0	40.9	37.5
a   6				l			91.7	78.6	68.8	61.1	55.0	50.0	45.8
1 7	T						}	92.9	81.3	72.2	65.0	59.1	54.2
. [8	Ŧ.			1			1		93.8	83.3	75.0	68.2	62.5
, , , 79	Т			I		l	l		1	94.4	85.0	77.3	70.8
i   10	) [			1		1				l	95.0	86.4	79.2
n	П						i			1	<b>I</b> !	95.5	87.5
1 12	! [	. 1					1. 3						95.8



#### ISOKINETIC FIELD DATA SHEET EPA Method 0010 - HFPO Dimer Acid **Stack Conditions** Meter Box ID Client Chemours K Factor 0069 15418.002.009 Assumed Actual Meter Box Y W.O.# 72,5 Initial Mid-Point Final Project ID Chemours % Moisture Meter Box Del H 0.001 P7DH Mode/Source ID Polymer Impinger Vol (ml) Probe ID / Length Sample Train (ft3) 0.00 D. DOI e15 **4** 5 Samp. Loc. ID STK Silica gel (g) Probe Material Boro Leak Check @ (in Hg) 47 P7041 Pitot / Thermocouple ID / no Run No.ID 1 CO2, % by Vol Pitot leak check good (Ves) / no 798 / no Test Method ID M0010 O2, % by Vol Pitot Coefficient 0.84 Pitot Inspection good 765 / no yes / no /03 / no G128 ves / no Date ID 17JAN2019 Temperature (°F) Nozzle ID Method 3 System good yes / no Ves / no Meter Temp (°F) 815,0 815,0 Pre-Test Set Post-Test Set Source/Location Polymer Stack Nozzle Measurements Temp Check Static Press (in H<sub>2</sub>O) Sample Date 17/19 24 -0,24 Avg Nozzle Dia (in) Meter Box Temp 60 53 -n60 Baro, Press (in Hg) Area of Stack (ft2) Reference Temp 260 A) LO CLIDAmblent Temp (°F) O Rads / Fail Operator Sample Time Pass/Fail (+/- 20) (Pase) / Fail Total Traverse Pts 24 Temp Change Response ? Cos / no / no

TRAVERSE POINT NO.	SAMPLE TIME (min)	CLOCK TIME (plant time)	VELOCITY PRESSURE Delta	PRESSURE	DRY GAS METER READING (ft³)	STACK TEMP (°F)	DGM OUTLET TEMP	PROBE TEMP (°F)	FILTER BOX TEMP	IMPINGER EXIT TEMP	SAMPLE TRAIN VAC	XAD EXIT TEMP (F)		COMMENTS
	0	1443	P (in H2O)	Delta H (in H2O)	441.130	remine (18)		HERRICAL FOR	(F)	(oF)	(in Hg)	(-110)		
17-1	ت ت	1 1-12	0.45	1.11	4143.41	63	65	258	242	ط7	3	56		
2	र्ष्ट		0,53	(2)	445.34	64	67	257	24.5	56	3	56		
3	12		0,55	1.35	443.40	64	66	257	24	56	3	36	****	28,324
4	16		0.48	1.18	450,40	64	66	257	258	50	7	50		
3	ZO		0,48	1.12	452.70	64	67	255-	257	44	3	44		
6	24		0,50	1.23	455.19	63	66	255	257	46	3	46		
7	58,		0,50	1,23	457 38	63	67	254	256	20	3	20		
8	32		0.51	1,25	459. 2	<u> </u>	68	८≤५	253	48	3	48		
9	36		84.0	1,18	462.01	62	67-	کحح	252	48	3	43		
10	40		Q. <i>S</i> Q	1,23	464,39	62	67	259	254	48	3	48		
14	44		0,59	1.46	466.81	63	6-)-	259	254	48	4	48		
12	48	1531	0.64	1,58	469.454		67-	259	254	48	34	43		
n   1	41	1543	A 2.	2 21	469.629		177	260	061=			6.		
1 15 31	1		0.35	0.86	471.82	91	64	259	255	24	3	54		
7 2	2		<u> </u>	0.86	475-22	6)	64	255-	255	54	3	الاك		
	12	*********	0.35	0.86	476.80	80	64	213	213		3	24		-6 631
1	10	*X1604	0,35	0.86		(0)	64	(03	207	2075	13	54		29.036
6	24	1689	0.65	1.61	480.20	-61	62	145	145	2 54 53	3	54 54		
1 3	28		0.69	1,70	415 56	6	52	135	122	53	3	<u> </u>		
q	32		0.69	* 1.70	428.70	61	62	135	145	32	3	32		
9	36		0,72	1: 75	491 15	Tol	52	125	145	\$2.	3	52		
10	40		0.65	1,100	494 01	61	62	135	145	52	3	52		
11	44		10.61	1.30	496.40	101	61	135	145	52	3	52	*****	
12	48	1641	52.0	1,22.	1498.66	61	161	133	145	52	3	50	****	
		0.53292	Avg Delta P	Avg Delta HV	Total Volume	/ 62.0	. Avg Tm . V	Min/Max	Min/Max JSO	Max グレ	Max Vac	Min/Max ろし		

Avg Sqrt Delta P Avg Sqrt Del H Comments

Block MBC ESSUES

EPA Method 0010 from EPA SW-846

/m

#### ISOKINETIC FIELD DATA SHEET EPA Method 0010 - HFPO Dimer Acid Page K Factor 2,5 Stack Conditions Meter Box ID Client Chemours ಯಠಿ 15418.002.009 Assumed Actual Meter Box Y W.O.# 18812 Initial Mid-Point Final Project ID Chemours % Moisture Meter Box Del H <u>7.013</u> 0.00 Mode/Source ID Polymer Impinger Vol (ml) Probe ID / Length POH Sample Train (ft3) 1001 13.0 W15 27 47 Silica gel (g) Probe Material Samp, Loc. ID STK Boro Leak Check (2) (in Hg) P704 **√** / no (96) / no ABO NO Pitot / Thermocouple ID Pitot leak check good Run No.ID 2 CO2, % by Vol 0. Test Method ID M0010 O2, % by Vol Pitot Coefficient 0.84 Pitot Inspection good Que / no yes / no yesy/ no (J 218 Method 3 System good 17JAN2019 Temperature (°F) > GD Nozzie ID ves / no yes / no Date ID ves / no Meter Temp (°F) = 61 Pre-Test Set Post-Test Set Polymer Stack Nozzie Measurements 0.28 0.218 Temp Check Source/Location Static Press (in H<sub>2</sub>O) -0.24 Meter Box Temp पठ Sample Date 7/18/19 -1),24 Avg Nozzle Dia (in) 0,218 55 30. h 4,91 Reference Temp 55 Baro, Press (in Hg) Area of Stack (ft<sup>2</sup>) ME, WINKELER (Pass / Fall Ambient Temp (°F) 96 Pass/Fail (+/- 20) Operator Sample Time Passo/ Fall Total Traverse Pts Temp Change Response 1 Yes / no y(3)/ no DRY GAS METER DGM OUTLET TEMP SAMPLE SAMPLE CLOCK TIME VELOCITY ORIFICE FILTER IMPINGER STACK PROBE XAD EXIT TRAVERSE TIME (min) PRESSURE Delta PRESSURE READING (ft<sup>3</sup>) (plant time) (Oil BOX TEMP **EXIT TEMP** TRAIN VAC COMMENTS POINT TEMP (oF) TEMP (F) NO. TEMP (°F) P (in H20) Delta H (in H2O (G)(oF) (in Hg) 0835 458.224 0 20 136 140 4 501.10 14 56 40 503.15 くつ 126 41 4 145 4 3 60,60 50 506,45 57 135 H 41 4 29.58 130 150 50826 56 13 o 0.60 4 1 25 36 0.03 410 ้งจ 100 41 0.13 56 46 100 100 41 30 50 100 100 417 Z2 60 <11b 41 45 4 4 $\mathbf{n}$ 518.70 100 37. ino 521 L 40 60 10 100 UD 100

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Avo Ts

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0.78467 | 1.45208 | 58.6 Avg Sqrt Delta PV | Avg Sqrt Del H | Comments: 0.76493 | 1.19504 |

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Avg Delta P \/

EPA Method 0010 from EPA SW-846

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#### EPA Method 0010 - HFPO Dimer Acid ISOKINETIC FIELD DATA SHEET **Stack Conditions** Chemours Meter Box ID Client K Factor 10069 Assumed Actual Meter Box Y W.O.# 15418.002.009 Mid-Point 22.2 Initial Final Meter Box Del H 8812 Project ID Chemours % Moisture 0.001 P702 Probe ID / Length Sample Train (ft3) 0,013 Polymer Impinger Vol (ml) Mode/Source ID e15 0.( ales Probe Material Boro Leak Check @ (in Hg) STK Silica gel (g) Samp, Loc, ID **P702** Vesy no Pitot leak check good yes) no West/ no CO2, % by Vol Pitot / Thermocouple ID Run No.ID yesy/ no (es) / no M0010 O2, % by Vol 2017 Pitot Coefficient 0.84 Pitot Inspection good yes y no Test Method ID GAIR Method 3 System good yes / no yes / no yes / no Temperature (°F) Nozzle ID Date ID 17JAN2019 クリマ Pre-Test Set 140 0.217 0.218 Temp Check Post-Test Set Meter Temp (°F) Nozzie Measurements 0.218 Polymer Stack Source/Location 63 55 Static Press (in H<sub>2</sub>O) Avg Nozzle Dia (in) 0.213 Meter Box Temp Sample Date 1/12/19 -0.24 1-0.24 62 4.91 Reference Temp Area of Stack (ft2) Baro, Press (in Hg) 20,01 Pass / Fall Pass / Fail Pass/Fail (+/- 20) NINKELER Amblent Temp (°F) Sample Time Operator Total Traverse Pts Temp Change Response (64) / no (yes) no SAMPLE DRY GAS METER **DGM OUTLET TEMP** IMPINGER SAMPLE CLOCK TIME VELOCITY ORIFICE FILTER XAD EXIT STACK PROBE TRAVERSE **EXIT TEMP** TRAIN VAC COMMENTS TIME (min) PRESSURE Delta PRESSURE READING (ft.) (oF) BOX TEMP (plant time) TEMP (F) TEMP (oF) TEMP (°F) POINT NO. (oF) (in Hg) Delta H (in H2O) P (in H2O) 558.112 0 1111

	U				1750,110	200			1000		0.000	The second second second	
14	4		0.55	1,35	560.62	59	58	100	100	50	4	SD	
2	S		0.55	1.35	563.21	59	58	100	100	70	4	20	
3	12		0.53	1,42	565, 49	59	58	100	100	43	7	<i>9</i> 8	29.887
4	16		0.63	1,54	568.00	59	58	180	/10	48	45	48	 <u>.</u>
اخ	25		0.63	1.54	570.60	59	58	100	100	47	ځ	47	
6	24		0.61	1.50	573,11	59	5-73	/00	101	ሃን	3	47	0.015
7	28		0.05	1,59	575.75	59	52	100	101	46	5	46	<b>12</b> (c)
2	32		0.68	1.67	578.49	59	52	100	101	46	S	46	
9	36		0,68	1,67	581,23	60	ર્ટ્ય	100	101	46	گ	46	586.430
10	40		0.63	1,54	523.76	GD	2.8	00	101	46	7	46	
1/	44		0.55	1.35	586114	60	<i>ই</i> হ	100	102	46	Ţ	46	
12	4%	1159	0.43	1,18	584,999	59	58	100	103	H6	4	46	 587,999 WF
2		1227			588.573								
15 1	Ч		0,58	1.42	591.08	61	53	100	1D1	56	4	بطك	
2	₹.		000	1.47	993,40	64	61	100	101	56	7	5 %	
3	72		0.65	1,59	596,21	64	63	(DD)	100	50	7	50	
Ч	16		0,68	1,67	598,86	65	65	100	100	50	Ż	50	4.4
3	20		0.70	1,72	601.55	67	65	100	185	50	5-	50	
6	24		0,65	1.59	6804,45	67	65	iDl	105	55	5	557	
7	28		0.06	662	606.94	67	67	101	101	66	5	66	
ર	3z		0.65	1,59	609,55	66	6-3	101	101	60	5	60	30.757
9	36		0,65	1,59	612,12	60	68	10)	i0	60	5-	60	
10	40		0.63	1,54	614 72	66	673	10)	10)	59	4	59	
	44		0.55	1, 1,35	617.13	lala	68	101	101	55	4	سي مي	
12	पर	1315	0.48	1)118/	1619.330	66	62/	101	101	35		-22	
			0.6/250	Avg Delta HV	GO, GHM	Avg Ts	Avo Tm	Min/Max	Min/Max	Max	Max Vac	Min/Max	
<b>LXX1</b>			0.61720	1120125	100.644	69.3	61.7 V	10/101	20 105	<u> </u>	5	46 00	1

(VVPQ)TABIVI

Avg Sqrt Delta P Avg Sqrt Del H Comments: EPA Method 0010 from EPA SW-846

## SAMPLE RECOVERY FIELD DATA

EPA Method 0010 - HFPO Dimer Acid

Client		Chemo	ours		W.O. #	1541	8.00	2,009,6		
Location/Pl	ant _	Fayettevi	lle, NC	Source	ce & Location			er Stack		_
Run No.	_1_				Sample Date			Recove	ery Date	clizhe
Sample I.D.	CHEMOURS	- Polymer - ST	K - 1 - M0010	-	Analyst	- Fore		Filter 1	Number	NA
					lmpinge					
	11	2	3	4	5	6	7	Imp.Total	8	Total
Contents	Empty	HPLC H20	HPLC H20						Silica Gel	
Final	l l	100	603	2				10C	719.6	
Initial	0	100	100	0				700	300	/
Gain	1	0	3	کم				6/	18.6	
Impinger Col	or	chean	_		Labeled?					
Silica Gel Co	ndition	000	<u> </u>		Sealed?					<b>-</b>
Run No.					Sample Date	1/18/19		Recove	ery Date	1/18/19
Sample I.D.	CHEMOURS	- Polymer - ST	K - 2 - M0010	-	Analyst	WF		Filter N	lumber	NA
				Impinge	r					
	11	2	3	4	5	6	7	Imp.Total	8	Total
Contents	Empty	HPLC H20	HPLC H20	·					Silica Gel	
Final	<u> </u>	96	100	0				196	3/30	
Initial	0	100	100	0				ræ	300	
Gain	ט	-4	0	0				1-4	13.0	<b>√</b>
Impinger Col	or	clear	_		Labeled?			<b>,</b>		_
Silica Gel Co	ndition	Blue			Sealed?					- -
Run No.	3_				Sample Date	1/18/19		Recove	ery Date	1/18/19
Sample I.D.	CHEMOURS	- Polymer - ST	K - 3 - M0010	-	Analyst	WF		Filter N	lumber	WA
					Impinge	r				
Cantonto	1	2	3	4	5	6	7	Imp.Total	8	Total
Contents	Empty	90	/02	4				197	Silica Gel	, /
Final Initial	5	100	100					700		1
Gain	WOSI	-10	2	4				-3	300 14.2	/
Impinger Col	nr	dea-			Labeled?	. /		1	·   · ·	š
	_	71	<del></del>		•					-
Silica Gel Co	ndition	Dluc			Sealed?					

## SAMPLE RECOVERY FIELD DATA

EPA Method 0010 - HFPO Dimer Acid

Client Location/Pla	ant —	Chemo Fayettevil		Sourc	W.O. # ce & Location		Polymer Stack					
		rayettevii	ie, NC		<u> </u>	11.1	- Polym			· · · · · · · · · · · · · · · · · · ·		
Run No.			BT		Sample Date	1/14/19	ï	Recove	ery Date	1/18/19		
Sample I.D.	CHEMOURS	6 - Polymer - ST	K M0010 -		Analyst	wr		Filter N	lumber	<u> </u>		
					Imping	1						
	1	2	3	4	5	6	7	Imp.Total	8	Total		
Contents	Empty	HPLC H20	HPLC H20						Silica Gel			
Final	0	/00	100	0					300			
Initial	0	100	100	0					300			
Gain	0	0	C	C					0			
Impinger Col	or _	clear_			Labeled?	/						
Silica Gel Co	ndition	Blue			Sealed?	<u></u>				•		
Run No.	2				Sample Date			Recove	ery Date			
Sample I.D.		5 - Polymer - ST	K - 2 - M0010 -		Analyst	····			lumber			
					Impinge							
	1	2	3	4	5	6	7	imp.Total	8	Total		
Contents	Empty	HPLC H20	HPLC H20						Silica Gel			
Final												
Initial		100	100						300			
Gain												
Impinger Col	or				Labeled?							
Silica Gel Co	ndition				Sealed?		··					
Run No.	3				Sample Date			Recove	ry Date			
Sample I.D.		i - Polymer - ST	K - 3 - M0010 -		Analyst				lumber			
•					Impinge	<del></del> er						
	11	2	3	4	5	6	7	Imp.Total	8	Total		
Contents	Empty	HPLC H20	HPLC H20					16	Silica Gel			
Final												
Initial	440000000000000000000000000000000000000	100	100						300			
Gain				was a same								
Impinger Col	or _				Labeled?					•		
Silica Gel Co	ndition				Sealed?							

Check COC for Sample IDs of Media Blanks



#### **METHODS AND ANALYZERS**

Client: Chemours Project Number: 15418.002.009

Operator: SR

Source: Polymers Date: 17 Jan 2019

ent Folders.A-F\Chemours Fayetteville\15418.002.009 Fayetteville Jan 2019 Carbon Bed Test\Data\Polymers\0117

Program Version: 2.1, built 19 May 2017 File Version: 2.03

Computer: WSWCAIRSERVICES Trailer: 27
Analog Input Device: Keithley KUSB-3108

#### **Channel 1**

Location: CHEMOURS

Analyte O<sub>2</sub>

Method EPA 3A, Using Bias Analyzer Make, Model & Serial No. Servomex 4900 Full-Scale Output, mv 10000 Analyzer Range, % 25.0 Span Concentration, % 21.0

#### **Channel 2**

Analyte CO<sub>2</sub>

Method
Analyzer Make, Model & Serial No.

Full-Scale Output, mv

Analyzer Pange %

EPA 3A, Using Bias
Servomex 4900
10000

Analyzer Range, % 20.0 Span Concentration, % 16.6



### **CALIBRATION DATA**

Number 1

Client: Chemours

Location: **CHEMOURS**Source: **Polymers** 

Project Number: 15418.002.009

Operator: **SR** 

Date: 17 Jan 2019

Start Time: 14:23

 $O_2$ 

Method: EPA 3A

Calibration Type: Linear Zero and High Span

Calibration Standards

%Cylinder ID12.0CC1805521.0SG9169108

**Calibration Results** 

**Zero** 2 mv **Span, 21.0 %** 8011 mv

**Curve Coefficients** 

Slope Intercept 381.4 2

 $CO_2$ 

Method: EPA 3A

Calibration Type: Linear Zero and High Span

Calibration Standards

**Cylinder ID**8.9 CC18055
16.6 SG9169108

Calibration Results

**Zero** -9 mv **Span, 16.6 %** 8288 mv

**Curve Coefficients** 

Slope Intercept 500.4 -9



### **CALIBRATION ERROR DATA**

Number 1

Client: Chemours

Project Number: 15418.002.009

Operator: SR

Location: CHEMOURS Source: Polymers

Calibration 1 Date: 17 Jan 2019

Start Time: 14:23

 $O_2$ 

Method: EPA 3A Span Conc. 21.0 %

**Slope** 381.4

Intercept 2.0

Standard %	Result %	Difference %	Error %	Status
Zero	0.0	0.0	0.0	Pass
12.0	12.1	0.1	0.5	Pass
21.0	21.0	0.0	0.0	Pass

 $CO_2$ 

Method: EPA 3A Span Conc. 16.6 %

**Slope** 500.4

Intercept -9.0

Standard %	Result %	Difference %	Error %	Status
Zero	0.0	0.0	0.0	Pass
8.9	8.6	-0.3	-1.8	Pass
16.6	16.6	0.0	0.0	Pass



# **BIAS**Number 1

Client: Chemours

Location: CHEMOURS

Source: Polymers

Project Number: 15418.002.009

Operator: **SR** 

Date: 17 Jan 2019

Calibration 1

Start Time: 14:26

 $O_2$ 

Method: EPA 3A Span Conc. 21.0 %

	Bias Results										
Standard	Cal.	Bias	Difference	Error							
Gas	%	%	%	%	Status						
Zero	0.0	0.0	0.0	0.0	Pass						
Span	12.1	12.0	-0.1	-0.5	Pass						

 $CO_2$ 

Method: EPA 3A Span Conc. 16.6 %

	Bias Results					
Standard	Cal.	Bias	Difference	Error		
Gas	%	%	%	%	Status	
Zero	0.0	0.1	0.1	0.6	Pass	
Span	8.6	8.4	-0.2	-1.2	Pass	



Number 1

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **17 Jan 2019** 

Tiı	me	O <sub>2</sub> %	CO <sub>2</sub> %
		Port A	
1.1	:43	20.9	0.0
	:44	20.9	0.0
	:45	20.8	0.0
	:46	20.8	0.0
	:47	20.9	0.0
	:48	20.9	0.0
	:49	20.9	0.0
	:50	20.9	0.0
	:51	20.9	0.0
	:52	21.0	0.0
	:53	21.0	0.0
	:54	21.0	0.0
	:55	21.0	0.0
	:56	21.0	0.0
	:57	21.0	0.0
	:58		0.0
		21.0	
	:59	21.0	0.0
	:00	21.0	0.0
	:01	21.0	0.0 0.0
	:02	21.0	
	:03	21.0	0.0
	:04	21.0	0.0
	:05	21.0	0.0
	:06	21.0	0.0
	:07	21.0	0.0
	:08	21.0	0.0
	:09	21.0	0.0
	:10	21.0	0.0
	5:11	21.0	0.0
	:12	21.0	0.0
	:13	21.0	0.0
	:14	21.0	0.0
	:15	21.0	0.0
	:16	21.0	0.0
	:17	21.0	0.0
	:18	21.0	0.0
	:19	21.0	0.0
	:20	21.0	0.0
15	:21	21.0	0.0



Number 1

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **17 Jan 2019** 

C	alibration	1	Date: <b>17 Jan 2019</b>
Time	<b>O</b> <sub>2</sub> %	CO <sub>2</sub> %	

	Time	O <sub>2</sub> %	CO <sub>2</sub> %	
		70	70	
	15:22	21.0	0.0	
	15:23	21.0	0.0	
	15:24	21.0	0.0	
	15:25	21.0	0.0	
	15:26	21.0	0.0	
	15:27	21.0	0.0	
	15:28	21.0	0.0	
	15:29	21.0	0.0	
	15:30	21.0	0.0	
	15:31	21.0	0.0	
		Port B		
	15:48	20.9	0.0	
	15:49	20.9	0.0	
	15:50	20.9	0.0	
	15:51	20.9	0.0	
	15:52	20.9	0.0	
	15:53	20.9	0.0	
	15:54	20.9	0.0	
	15:55	20.9	0.0	
	15:56	20.9	0.0	
	15:57	20.9	0.0	
	15:58 15:50	21.0	0.0	
	15:59 16:00	21.0 20.9	0.0 0.0	
	16:01	20.9	0.0	
	16:01	21.0	0.0	
	16:02	21.0	0.0	
	16:04	21.0	0.0	
ı			ibe blocked	
•	16:09	20.9	0.0	
	16:10	20.9	0.0	
	16:11	20.9	0.0	
	16:12	20.9	0.0	
	16:13	20.9	0.0	
	16:14	20.9	0.0	
	16:15	20.9	0.0	
	16:16	20.9	0.0	
	16:17	20.9	0.0	
	16:18	20.9	0.0	
	16:19	21.0	0.0	



Number 1

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **17 Jan 2019** 

		0-	
Tim	ne	<b>O</b> 2 %	CO <sub>2</sub>
16:2		21.0	0.0
16:2	21	21.0	0.0
16:2	22	21.0	0.0
16:2	23	21.0	0.0
16:2	24	21.0	0.0
16:2	25	21.0	0.0
16:2		20.9	0.0
16:2		21.0	0.0
16:2		20.9	0.0
16:2		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:3		20.9	0.0
16:		20.9	0.0
16:4		20.9	0.0
16:4		20.9	0.0
10.		End Run 1	0.0
Avç		21.0	0.0
Λ·ζ	y	21.0	0.0



### **RUN SUMMARY**

Number 1

Client: Chemours

Location: CHEMOURS

Project Number: **15418.002.009** 

Operator: SR

Source: Polymers Calibration 1 Date: 17 Jan 2019

O2CO2MethodEPA 3AEPA 3AConc. Units%%

Time: 14:42 to 16:41

**Run Averages** 

21.0 0.0

Pre-run Bias at 14:26

 Zero Bias
 0.0
 0.1

 Span Bias
 12.0
 8.4

 Span Gas
 12.0
 8.9

Post-run Bias at 17:03

 Zero Bias
 0.0
 0.1

 Span Bias
 12.0
 8.3

 Span Gas
 12.0
 8.9

Run averages corrected for the average of the pre-run and post-run bias

21.0 0.0



# **BIAS AND CALIBRATION DRIFT**

Number 2

Client: Chemours

Location: CHEMOURS

Source: Polymers

Project Number: 15418.002.009

Operator: **SR** 

Date: 17 Jan 2019

Calibration 1

 $O_2$ 

Start Time: 17:03

Method: EPA 3A Span Conc. 21.0 %

		Bias	Results		
Standard	Cal.	Bias	Difference	Error	
Gas	%	%	%	%	Status
Zero	0.0	0.0	0.0	0.0	Pass
Span	12.1	12.0	-0.1	-0.5	Pass
		Calibra	ation Drift		
Standard	Initial*	Final	Difference	Drift	
Gas	%	%	%	%	Status
Zero	0.0	0.0	0.0	0.0	Pass
Span	12.0	12.0	0.0	0.0	Pass
•	*Bias No. 1				

CO<sub>2</sub>

Method: EPA 3A Span Conc. 16.6 %

		Bias	Results			
Standard	Cal.	Bias	Difference	Error		
Gas	%	%	%	%	Status	
Zero	0.0	0.1	0.1	0.6	Pass	
Span	8.6	8.3	-0.3	-1.8	Pass	
Calibration Drift						
Standard	Initial*	Final	Difference	Drift		
Gas	%	%	%	%	Status	
Zero	0.1	0.1	0.0	0.0	Pass	
Span	8.4	8.3	-0.1	-0.6	Pass	
•	*Bias No. 1					



#### **METHODS AND ANALYZERS**

Project Number: 15418.002.009 Client: Chemours

Operator: **SR** 

Date: 18 Jan 2019 Source: Polymers

ent Folders.A-F\Chemours Fayetteville\15418.002.009 Fayetteville Jan 2019 Carbon Bed Test\Data\Polymers\0118

Program Version: 2.1, built 19 May 2017 File Version: 2.03

Computer: WSWCAIRSERVICES Trailer: 27 Analog Input Device: Keithley KUSB-3108

16.6

#### Channel 1

Location: CHEMOURS

Analyte  $O_2$ 

EPA 3A, Using Bias Method Servomex 4900 Analyzer Make, Model & Serial No. Full-Scale Output, mv 10000 Analyzer Range, % 25.0 Span Concentration, % 21.0

#### Channel 2

Analyte  $CO_2$ 

EPA 3A, Using Bias Method Analyzer Make, Model & Serial No. Servomex 4900 Full-Scale Output, mv 10000 Analyzer Range, % 20.0 Span Concentration, %

## **CALIBRATION DATA**

Number 1

Client: Chemours

Location: **CHEMOURS**Source: **Polymers** 

Project Number: 15418.002.009

Operator: **SR** 

Date: 18 Jan 2019

Start Time: 07:19

 $O_2$ 

Method: EPA 3A

Calibration Type: Linear Zero and High Span

Calibration Standards

% Cylinder ID 12.0 CC18055 21.0 SG9169108

**Calibration Results** 

**Zero** 13 mv **Span, 21.0 %** 8012 mv

**Curve Coefficients** 

Slope Intercept 380.9 13

 $CO_2$ 

Method: EPA 3A

Calibration Type: Linear Zero and High Span

Calibration Standards

 %
 Cylinder ID

 8.9
 CC18055

 16.6
 SG9169108

**Calibration Results** 

**Zero** -10 mv **Span, 16.6 %** 8293 mv

**Curve Coefficients** 

Slope Intercept 500.8 -10



# **CALIBRATION ERROR DATA**

Number 1

Client: Chemours

Project Number: 15418.002.009

Location: CHEMOURS

Calibration 1

Operator: **SR** 

Source: **Polymers** Calibra

Date: 18 Jan 2019

Start Time: 07:19

 $O_2$ 

Method: EPA 3A Span Conc. 21.0 %

**Slope** 380.9

Intercept 13.0

Standard %	Result %	Difference %	Error %	Status
Zero	0.0	0.0	0.0	Pass
12.0	12.1	0.1	0.5	Pass
21.0	21.0	0.0	0.0	Pass

 $CO_2$ 

Method: EPA 3A Span Conc. 16.6 %

**Slope** 500.8

Intercept -10.0

Standard %	Result %	Difference %	Error %	Status
Zero	0.0	0.0	0.0	Pass
8.9	8.6	-0.3	-1.8	Pass
16.6	16.6	0.0	0.0	Pass



# **BIAS**Number 1

Client: Chemours

Location: CHEMOURS

Source: Polymers

Project Number: 15418.002.009

Operator: **SR** 

Date: 18 Jan 2019

Calibration 1

Start Time: 07:23

 $O_2$ 

Method: EPA 3A Span Conc. 21.0 %

Bias Results							
Standard	Cal.	Bias	Difference	Error			
Gas	%	%	%	%	Status		
Zero	0.0	0.0	0.0	0.0	Pass		
Span	12.1	12.0	-0.1	-0.5	Pass		

 $CO_2$ 

Method: EPA 3A Span Conc. 16.6 %

		Bias	Results		
Standard	Cal.	Bias	Difference	Error	
Gas	%	%	%	%	Status
Zero	0.0	0.1	0.1	0.6	Pass
Span	8.6	8.4	-0.2	-1.2	Pass



Number 2

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **18 Jan 2019** 

Time	$O_2$	$CO_2$					
Time	%	%					
Port B							
08:35		0.1					
08:36		0.1					
08:37		0.1					
08:38		0.1					
08:39		0.1					
08:40		0.1					
08:41		0.1					
08:42		0.1					
08:43		0.1					
08:44		0.1					
08:45		0.1					
08:46		0.1					
08:47		0.1					
08:48		0.1					
08:49		0.1					
08:50		0.1					
08:51		0.1					
08:52		0.1					
08:53		0.1					
08:54		0.1					
08:55		0.1					
08:56		0.1					
08:57		0.1					
08:58		0.1					
08:59		0.1					
09:00		0.1					
09:01		0.1					
09:02		0.1					
09:03		0.1					
09:04		0.1					
09:05		0.1					
09:06		0.1					
09:07		0.1					
09:08		0.1					
09:09		0.1					
09:10		0.1					
09:11		0.1					
09:12		0.1					
09:12		0.1					
09.10	20.0	0.1					



Number 2

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **18 Jan 2019** 

Time	<b>O</b> <sub>2</sub> %	CO <sub>2</sub> %	
09:14	20.9	0.1	
09:15			
09:16			
09:17		0.1	
09:18	20.9	0.1	
09:19	20.9	0.1	
09:20	20.9	0.1	
09:21	20.9	0.1	
09:22	20.9	0.1	
09:23	20.9	0.1	
	Port B		
09:40	20.9	0.1	
09:41	20.9	0.1	
09:42	20.9	0.1	
09:43	20.9	0.1	
09:44	20.9	0.1	
09:45	20.9	0.1	
09:46	20.9	0.1	
09:47	20.9	0.1	
09:48	20.9	0.1	
09:49	20.9	0.1	
09:50	20.9	0.1	
09:51	20.9	0.1	
09:52	20.9	0.1	
09:53	20.9	0.1	
09:54	20.9	0.1	
09:55	20.9	0.1	
09:58	20.9	0.1	
	20.9	0.1	
10:07	21.0	0.1	
10:08	21.0	0.1	
	09:14 09:15 09:16 09:17 09:18 09:19 09:20 09:21 09:22 09:23  09:40 09:41 09:42 09:43 09:44 09:45 09:45 09:46 09:47 09:48 09:49 09:50 09:51 09:52 09:53 09:54 09:55 09:56 09:57 09:58 09:57 09:58 09:59 10:00 10:01 10:02 10:03 10:04 10:05 10:06	09:14	09:14 20.9 0.1 09:15 20.9 0.1 09:16 20.9 0.1 09:17 20.9 0.1 09:18 20.9 0.1 09:19 20.9 0.1 09:20 20.9 0.1 09:21 20.9 0.1 09:22 20.9 0.1 09:23 20.9 0.1 09:44 20.9 0.1 09:42 20.9 0.1 09:45 20.9 0.1 09:45 20.9 0.1 09:46 20.9 0.1 09:47 20.9 0.1 09:48 20.9 0.1 09:48 20.9 0.1 09:49 20.9 0.1 09:50 20.9 0.1 09:50 20.9 0.1 09:51 20.9 0.1 09:52 20.9 0.1 09:53 20.9 0.1 09:55 20.9 0.1 09:55 20.9 0.1 09:56 20.9 0.1 09:57 20.9 0.1 09:58 20.9 0.1 09:59 20.9 0.1 09:59 20.9 0.1 09:59 20.9 0.1 09:59 20.9 0.1 09:59 20.9 0.1 09:50 20.9 0.1 09:50 20.9 0.1 09:51 20.9 0.1 09:52 20.9 0.1 09:53 20.9 0.1 09:54 20.9 0.1 09:55 20.9 0.1 09:55 20.9 0.1 09:55 20.9 0.1 09:56 20.9 0.1 09:57 20.9 0.1 09:59 20.9 0.1 09:59 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1 10:00 20.9 0.1



Number 2

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **18 Jan 2019** 

Tim	ne	<b>O</b> <sub>2</sub> %	CO <sub>2</sub> %	
10:0	 09	21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:3		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:3		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10:		21.0	0.1	
10		nd Run 2	J	
Avg		20.9	0.1	
A	9-9	_0.0	J	



### **RUN SUMMARY**

Number 2

Client: Chemours

Location: CHEMOURS
Source: Polymers

Project Number: **15418.002.009** 

Operator: SR

Calibration 1 Date: 18 Jan 2019

Method EPA 3A EPA 3A Conc. Units % %

Time: 08:34 to 10:28

**Run Averages** 

20.9 0.1

Pre-run Bias at 07:23

 Zero Bias
 0.0
 0.1

 Span Bias
 12.0
 8.4

 Span Gas
 12.0
 8.9

Post-run Bias at 11:08

 Zero Bias
 0.1
 0.0

 Span Bias
 12.0
 8.4

 Span Gas
 12.0
 8.9

Run averages corrected for the average of the pre-run and post-run bias

21.0 0.1



# **BIAS AND CALIBRATION DRIFT**

Number 2

Client: Chemours

Location: CHEMOURS

Source: Polymers

Project Number: 15418.002.009

Operator: **SR** 

Date: 18 Jan 2019

Calibration 1

Start Time: 11:08

 $O_2$ 

Method: EPA 3A Span Conc. 21.0 %

		Bias	Results		
Standard Gas	Cal. %	Bias %	Difference %	Error %	Status
Zero	0.0	0.1	0.1	0.5	Pass
Span	12.1	12.0	-0.1	-0.5	Pass
		Calibra	ation Drift		
Standard	Initial*	Final	Difference	Drift	
Gas	%	%	%	%	Status
Zero	0.0	0.1	0.1	0.5	Pass
Span	12.0 *Bias No. 1	12.0	0.0	0.0	Pass

 $CO_2$ 

Method: EPA 3A Span Conc. 16.6 %

	Bias Results						
Standard	Cal.	Bias	Difference	Error			
Gas	%	%	%	%	Status		
Zero	0.0	0.0	0.0	0.0	Pass		
Span	8.6	8.4	-0.2	-1.2	Pass		
	Calibration Drift						
Standard	Initial*	Final	Difference	Drift			
Gas	%	%	%	%	Status		
Zero	0.1	0.0	-0.1	-0.6	Pass		
Span	8.4	8.4	0.0	0.0	Pass		
•	*Bias No. 1						



Number 3

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **18 Jan 2019** 

Tim	10	$O_2$	$CO_2$	
11111	i <del>C</del>	%	%	
		Port A		
11:1		20.8	0.2	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:1		20.9	0.1	
11:2		20.9	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:2		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:3		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11:4		21.0	0.1	
11		_ 1.0	J. 1	



Number 3

Calibration 1

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **18 Jan 2019** 

 	, and a dion	•	
Time	<b>O</b> <sub>2</sub> %	CO <sub>2</sub> %	
11:50	21.0	0.1	
11:51	21.0	0.1	
11:52	21.0	0.1	
11:53	21.0	0.1	
11:54	21.0	0.1	
11:55	21.0	0.1	
11:56	21.0	0.1	
11:57	21.0	0.1	
11:58	21.0	0.1	
11:59	21.0	0.1	
	Port B		
12:27	20.9	0.1	
12:28	20.9	0.1	
12:29	20.9	0.1	
12:30	20.9	0.1	
12:31	20.9	0.1	
12:32	20.9	0.1	
12:33	20.9	0.1	
12:34	20.9	0.1	
12:35	21.0	0.1	
12:36	21.0	0.1	
12:37	21.0	0.1	
12:38	21.0	0.1	
12:39	21.0	0.1	
12:40	21.0	0.1	
12:41	21.0	0.1	
12:42	21.0	0.1	
12:43	21.0	0.1	
12:44	21.0	0.1	
12:45	21.0	0.1	
12:46	21.0	0.1	
12:47	21.0	0.1	
12:48	21.0	0.1	
12:49	21.0	0.1	
12:50	21.0	0.1	
12:51	21.0	0.1	
12:52	21.0	0.1	
12:53	21.0	0.1	
12:54	21.0	0.1	
12:55	21.0	0.1	



Number 3

Client: Chemours

Location: CHEMOURS Source: Polymers

Project Number: **15418.002.009**Operator: **SR**Date: **18 Jan 2019** 

Time	<b>O</b> <sub>2</sub> %	CO <sub>2</sub> %	
12:56	21.0	0.1	
12:57	21.0	0.1	
12:58	21.0	0.1	
12:59	21.0	0.1	
13:00	21.0	0.1	
13:01	21.0	0.1	
13:02	21.0	0.1	
13:03	21.0	0.1	
13:04	21.0	0.1	
13:05	21.0	0.1	
13:06	21.0	0.1	
13:07	21.0	0.1	
13:08	21.0	0.1	
13:09	21.0	0.1	
13:10	21.0	0.1	
13:11	21.0	0.1	
13:12	21.0	0.1	
13:13	21.0	0.1	
13:14	21.0	0.1	
13:15	21.0	0.1	
	End Run 3		
Avgs	21.0	0.1	
-			



### **RUN SUMMARY**

Number 3

Client: **Chemours** 

Location: CHEMOURS
Source: Polymers

Project Number: 15418.002.009

Operator: **SR** 

Calibration 1 Date: 18 Jan 2019

O2CO2MethodEPA 3AEPA 3AConc. Units%

Time: 11:10 to 13:15

## **Run Averages**

21.0 0.1

#### Pre-run Bias at 11:08

Zero Bias	0.1	0.0
Span Bias	12.0	8.4
Span Gas	12.0	8.9

## Post-run Bias at 13:34

Zero Bias	0.0	0.0
Span Bias	11.9	8.3
Span Gas	12.0	8.9

Run averages corrected for the average of the pre-run and post-run bias

21.1 0.1



# **BIAS AND CALIBRATION DRIFT**

Number 3

Client: Chemours

Location: CHEMOURS

Source: Polymers

Project Number: 15418.002.009

Operator: **SR** 

Date: 18 Jan 2019

Calibration 1

 $O_2$ 

Start Time: 13:34

Method: EPA 3A Span Conc. 21.0 %

Bias Results						
Standard	Cal.	Bias	Difference	Error		
Gas	%	%	%	%	Status	
Zero	0.0	0.0	0.0	0.0	Pass	
Span	12.1	11.9	-0.2	-1.0	Pass	
		Calibra	ation Drift			
Standard	Initial*	Final	Difference	Drift		
Gas	%	%	%	%	Status	
Zero	0.1	0.0	-0.1	-0.5	Pass	
Span	12.0	11.9	-0.1	-0.5	Pass	
•	*Bias No. 2					

 $CO_2$ 

Method: EPA 3A Span Conc. 16.6 %

Bias Results						
Standard	Cal.	Bias	Difference	Error		
Gas	%	%	%	%	Status	
Zero	0.0	0.0	0.0	0.0	Pass	
Span	8.6	8.3	-0.3	-1.8	Pass	
		Calibra	ation Drift			
Standard	Initial*	Final	Difference	Drift		
Gas	%	%	%	%	Status	
Zero	0.0	0.0	0.0	0.0	Pass	
Span	8.4	8.3	-0.1	-0.6	Pass	
•	*Bias No. 2					



# APPENDIX C LABORATORY ANALYTICAL REPORT

Note: The analytical report is included on the attached CD.

Client: Chemours Company FC, LLC The Project/Site: Polymer Stack - M0010

TestAmerica Job ID: 140-14024-1

Client Sample ID: H-2501,2502 POLYMER STK R1 M0010 FH

Lab Sample ID: 140-14024-1

Date Collected: 01/17/19 00:00 Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit Prepared Analyzed Dil Fac **HFPO-DA** 0.151 0.0163 ug/Sample 01/28/19 10:24 02/04/19 10:49 1.67 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac

13C3 HFPO-DA 106 50 - 200

Client Sample ID: H-2503,2504,2506 POLYMER STK R1 M0010

Lab Sample ID: 140-14024-2

Lab Sample ID: 140-14024-4

01/28/19 10:24 02/04/19 10:49

Matrix: Air

BH

Date Collected: 01/17/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte RL **MDL** Unit Result Qualifier **Prepared** Analyzed Dil Fac 0.200 01/30/19 04:34 02/06/19 11:07 HFPO-DA 0.0400 ug/Sample 5.08 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac

13C3 HFPO-DA 42 X 50 - 200 01/30/19 04:34 02/06/19 11:07

Client Sample ID: H-2505 POLYMER STK R1 M0010 IMP 1,2&3 Lab Sample ID: 140-14024-3

CONDENSATE

Date Collected: 01/17/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - HFPO-DA Analyte Result Qualifier RL **MDL** Unit **Prepared** Analyzed

Dil Fac HFPO-DA 0.204 01/30/19 04:45 02/04/19 12:17  $\overline{\mathsf{ND}}$ 0.0104 ug/Sample Surrogate Qualifier Limits Prepared Analyzed Dil Fac %Recovery 13C3 HFPO-DA 50 - 200 01/30/19 04:45 02/04/19 12:17 96

Client Sample ID: H-2507 POLYMER STK R1 M0010

**BREAKTHROUGH XAD-2 RESIN TUBE** 

Date Collected: 01/17/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit **Prepared** Analyzed Dil Fac HFPO-DA ND 0.200 0.0400 ug/Sample 01/30/19 04:34 02/06/19 11:10 Surrogate Qualifier Limits

%Recovery Prepared Analyzed Dil Fac 13C3 HFPO-DA 88 50 - 200 01/30/19 04:34 02/06/19 11:10

Project/Site: Polymer Stack - M0010

Client: Chemours Company FC, LLC The TestAmerica Job ID: 140-14024-1

Client Sample ID: H-2508,2509 POLYMER STK R2 M0010 FH

Lab Sample ID: 140-14024-5

Date Collected: 01/18/19 00:00 Date Received: 01/20/19 10:00

Matrix: Air

Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit Prepared Analyzed **HFPO-DA** 0.126 0.0136 ug/Sample 01/28/19 10:24 02/04/19 10:52 0.975

Surrogate %Recovery Qualifier I imits Prepared Analyzed Dil Fac 13C3 HFPO-DA 101 50 - 200 01/28/19 10:24 02/04/19 10:52

Client Sample ID: H-2510,2511,2513 POLYMER STK R2 M0010

Lab Sample ID: 140-14024-6

BH

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit Prepared Analyzed Dil Fac HFPO-DA 0.200 0.0400 ug/Sample 01/30/19 04:34 02/06/19 11:13 4.53 %Recovery Qualifier I imits Analyzed Dil Fac Surrogate Prepared 48 X 13C3 HFPO-DA 50 - 200 01/30/19 04:34 02/06/19 11:13

Lab Sample ID: 140-14024-7 Client Sample ID: H-2512 POLYMER STK R2 M0010 IMP 1,2&3

**CONDENSATE** 

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - HFPO-DA

Analyte Result Qualifier RL MDL Unit **Prepared** Analyzed Dil Fac HFPO-DA ND 0 194 0.00989 ug/Sample 01/30/19 04:45 02/04/19 12:20 %Recovery Qualifier Surrogate Limits Dil Fac Prepared Analyzed 13C3 HFPO-DA 93 50 - 200 01/30/19 04:45 02/04/19 12:20

Client Sample ID: H-2514 POLYMER STK R2 M0010 Lab Sample ID: 140-14024-8

BREAKTHROUGH XAD-2 RESIN TUBE

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Result Qualifier **MDL** Unit Analyte RL Analyzed Prepared Dil Fac HFPO-DA ND 0.200 01/30/19 04:34 02/06/19 11:20 0.0400 ug/Sample

Qualifier %Recovery Surrogate Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 01/30/19 04:34 02/06/19 11:20 81 50 - 200

Client: Chemours Company FC, LLC The Project/Site: Polymer Stack - M0010

TestAmerica Job ID: 140-14024-1

Client Sample ID: H-2515,2516 POLYMER STK R3 M0010 FH

Lab Sample ID: 140-14024-9

Matrix: Air

Date Collected: 01/18/19 00:00

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit Prepared Analyzed **HFPO-DA** 0.151 0.0163 ug/Sample 01/28/19 10:24 02/04/19 10:55 1.07

Surrogate %Recovery Qualifier I imits Prepared Analyzed Dil Fac 13C3 HFPO-DA 103 50 - 200 01/28/19 10:24 02/04/19 10:55

Client Sample ID: H-2517,2518,2520 POLYMER STK R3 M0010

Lab Sample ID: 140-14024-10

BH

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit Prepared Analyzed Dil Fac HFPO-DA 0.250 0.0500 ug/Sample 01/30/19 04:34 02/06/19 11:23 4.42 %Recovery Qualifier I imits Analyzed Dil Fac Surrogate Prepared 13C3 HFPO-DA 51 50 - 200 01/30/19 04:34 02/06/19 11:23

Client Sample ID: H-2519 POLYMER STK R3 M0010 IMP 1,2&3

Lab Sample ID: 140-14024-11

Lab Sample ID: 140-14024-12

**CONDENSATE** 

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - HFPO-DA

Analyte Result Qualifier RL MDL Unit **Prepared** Analyzed Dil Fac **HFPO-DA** 0.0492 J 0 194 0.00989 ug/Sample 01/30/19 04:45 02/04/19 12:23 Surrogate %Recovery Qualifier Limits Dil Fac Prepared Analyzed 13C3 HFPO-DA 78 50 - 200 01/30/19 04:45 02/04/19 12:23

Client Sample ID: H-2521 POLYMER STK R3 M0010

BREAKTHROUGH XAD-2 RESIN TUBE

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

**MDL** Unit Analyte Result Qualifier RL Analyzed Prepared Dil Fac **HFPO-DA** 0.200 01/30/19 04:34 02/06/19 11:27 0.0939 .1 0.0400 ug/Sample Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 01/30/19 04:34 02/06/19 11:27 83 50 - 200

Client: Chemours Company FC, LLC The

Project/Site: Polymer Stack Field QC Samples

Lab Sample ID: 140-14026-1

TestAmerica Job ID: 140-14026-1

Client Sample ID: H-2522,2523 POLYMER STK QC M0010 FH

BT

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit D Prepared Analyzed Dil Fac HFPO-DA 0.159 0.0260 0.00281 ug/Sample 01/28/19 10:24 02/04/19 11:02

Surrogate Limits Prepared Dil Fac %Recovery Qualifier Analyzed 13C3 HFPO-DA 89 50 - 200 01/28/19 10:24 02/04/19 11:02

Client Sample ID: H-2524,2525,2527 POLYMER STK QC M0010

BH BT

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit Prepared Analyzed Dil Fac 0.200 01/30/19 04:34 02/06/19 11:30 **HFPO-DA** 0.412 0.0400 ug/Sample Limits Prepared Analyzed Dil Fac Surrogate %Recovery Qualifier

13C3 HFPO-DA 50 - 200 75

Client Sample ID: H-2526 POLYMER STK QC M0010 IMP 1,2&3

Lab Sample ID: 140-14026-3

Lab Sample ID: 140-14026-4

01/30/19 04:34 02/06/19 11:30

Lab Sample ID: 140-14026-2

**CONDENSATE BT** 

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - HFPO-DA

**Analyte** Result Qualifier RL **MDL** Unit **Prepared** Analyzed Dil Fac 0.00250 01/30/19 04:45 02/04/19 12:26 **HFPO-DA** 0.000392 J 0.000128 ug/Sample

Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 102 50 - 200 01/30/19 04:45 02/04/19 12:26

Client Sample ID: H-2528 POLYMER STK QC M0010

**BREAKTHROUGH XAD-2 RESIN TUBE BT** 

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL MDL Unit **Prepared** Analyzed Dil Fac HFPO-DA  $\overline{\mathsf{ND}}$ 0.200 0.0400 ug/Sample 01/30/19 04:34 02/06/19 11:33 %Recovery Surrogate Qualifier I imits Prepared Analyzed Dil Fac 13C3 HFPO-DA 50 - 200 01/30/19 04:34 02/06/19 11:33 79

Client: Chemours Company FC, LLC The

Project/Site: Polymer Stack Field QC Samples

TestAmerica Job ID: 140-14026-1

Client Sample ID: H-2529 POLYMER STK QC M0010 DI WATER

**RB** 

Date Collected: 01/18/19 00:00

Date Received: 01/20/19 10:00 Sample Container: Air Train

Lab Sample ID: 140-14026-5

Matrix: Air

Method: 8321A - HFPO-DA

Analyte Result Qualifier RL **MDL** Unit D Prepared Analyzed Dil Fac HFPO-DA ND 0.00250 0.000128 ug/Sample 01/30/19 04:45 02/04/19 12:30

Surrogate %Recovery Qualifier Limits 13C3 HFPO-DA 115 50 - 200

Prepared Analyzed Dil Fac 01/30/19 04:45 02/04/19 12:30

Analyzed

Lab Sample ID: 140-14026-7

Lab Sample ID: 140-14026-8

Dil Fac

Lab Sample ID: 140-14026-6

Client Sample ID: H-2530 POLYMER STK QC M0010 MEOH

WITH 5% NH4OH RB

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit D **Prepared** Analyzed Dil Fac HFPO-DA ND 0.0250 0.00500 ug/Sample 01/30/19 04:34 02/06/19 11:36

Surrogate %Recovery Qualifier Limits Prepared 13C3 HFPO-DA 95 50 - 200 01/30/19 04:34 02/06/19 11:36

Client Sample ID: H-2531 POLYMER STK QC M0010 XAD-2

**RESIN TUBE RB** 

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

Analyte Result Qualifier RL **MDL** Unit D **Prepared** Analyzed Dil Fac HFPO-DA  $\overline{\mathsf{ND}}$ 0.200 0.0400 ug/Sample 01/30/19 04:34 02/06/19 11:40 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 01/30/19 04:34 02/06/19 11:40 13C3 HFPO-DA 50 - 200 82

Client Sample ID: H-2532 POLYMER STK QC M0010 MEOH

WITH 5% NH4OH TB

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

**Analyte** Result Qualifier RL MDL Unit D **Prepared** Analyzed Dil Fac HFPO-DA  $\overline{\mathsf{ND}}$ 0.0250 0.00500 ug/Sample 01/30/19 04:34 02/06/19 11:43

Surrogate %Recovery Qualifier Limits Prepared Dil Fac Analyzed 13C3 HFPO-DA 50 - 200 01/30/19 04:34 02/06/19 11:43 96

Client: Chemours Company FC, LLC The TestAmerica Job ID: 140-14026-1

Project/Site: Polymer Stack Field QC Samples

Client Sample ID: H-2533 POLYMER STK QC M0010 XAD-2

**RESIN TUBE TB** 

Date Collected: 01/18/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

 Analyte
 Result
 Qualifier
 RL
 MDL
 Unit
 D
 Prepared
 Analyzed
 Dil Fac

 HFPO-DA
 ND
 0.200
 0.0400
 ug/Sample
 01/30/19 04:34
 02/06/19 11:46
 1

 Surrogate
 %Recovery
 Qualifier
 Limits

 13C3 HFPO-DA
 83
 50 - 200

Lab Sample ID: 140-14026-10

01/30/19 04:34 02/06/19 11:46

Analyzed

Dil Fac

Prepared

Lab Sample ID: 140-14026-9

Client Sample ID: H-2534 POLYMER STK QC M0010

COMBINED GLASSWARE RINSES (MEOH/5% NH4OH) PB

Date Collected: 01/17/19 00:00 Matrix: Air

Date Received: 01/20/19 10:00 Sample Container: Air Train

Method: 8321A - PFOA and PFOS

 Analyte
 Result
 Qualifier
 RL
 MDL
 Unit
 D
 Prepared
 Analyzed
 Dil Fac

 HFPO-DA
 0.00556
 J
 0.0250
 0.00500
 ug/Sample
 01/30/19 04:34
 02/06/19 11:49
 1

# APPENDIX D SAMPLE CALCULATIONS

# SAMPLE CALCULATIONS FOR HFPO DIMER ACID (METHOD 0010)

Client: Chemours
Test Number: Run 1
Test Location: Polymer Stack

Plant: Fayetteville, NC
Test Date: 1/17/2019
Test Period: 1443-1641

#### 1. HFPO Dimer Acid concentration, lbs/dscf.

$$C_1 = W \times 2.2046 \times 10^{-9}$$
 $Vm(std)$ 

$$C_1 = \frac{6.8 \times 2.2046 \times 10-9}{58.636}$$

$$= 2.54E-10$$

Where:

W = Weight of HFPO Dimer Acid collected in sample in ug.

C<sub>1</sub> = Polymer Stack HFPO Dimer Acid concentration, lbs/dscf.

 $2.2046 \times 10^{-9}$  = Conversion factor from ug to lbs.

#### 2. HFPO Dimer Acid concentration, ug/dscm.

 $C_2 = W / (Vm(std) \times 0.02832)$ 

 $C_2 = 6.8 / (58.636 \times 0.02832)$ 

= 4.06E+00

Where:

C<sub>2</sub> = HFPO Dimer Acid concentration, ug/dscm.

0.02832 = Conversion factor from cubic feet to cubic meters.

2/19/201911:29 AM O:\S\A\FMC\011719 polymer

#### 3. HFPO Dimer Acid mass emission rate, lbs/hr.

 $MR1 = C_1 \times Qs(std) \times 60 \text{ min/hr}$ 

 $MR1 = 2.54E-10 \times 11910 \times 60$ 

= 1.81E-04

Where:

MR1 = Polymer Stack HFPO Dimer Acid mass emission rate, lbs/hr.

#### 4. HFPO Dimer Acid mass emission rate, g/sec.

 $MR2 = PMR1 \times 453.59 / 3600$ 

 $MR2 = 1.81E-04 \times 453.59 / 3600$ 

= 2.28E-05

Where:

MR2 = Polymer Stack HFPO Dimer Acid mass emission rate, g/sec.

454 = Conversion factor from pounds to grams.

### = Conversion factor from hours to seconds.

2/19/201911:29 AM O:\S\A\FMC\011719 polymer

# EXAMPLE CALCULATIONS FOR VOLUMETRIC FLOW AND MOISTURE AND ISOKINETICS

 Client: Chemours
 Facility: Fayetteville, NC

 Test Number: Run 1
 Test Date: 1/17/19

 Test Location: Polymer Stack
 Test Period: 1443-1641

#### 1. Volume of dry gas sampled at standard conditions (68 deg F, 29.92 in. Hg), dscf.

#### 2. Volume of water vapor in the gas sample corrected to standard conditions, scf.

Vw(std) =	(0.04707  x Vwc) + (0.04715  x Wwsg)
Vw(std) =	$(0.04707 \times 6.0) + (0.04715 \times 19.6) = 1.21$
Where:	
Vw(std) =	Volume of water vapor in the gas sample corrected to standard conditions, scf.
Vwc =	Volume of liquid condensed in impingers, ml.
Wwsg =	Weight of water vapor collected in silica gel, g.
0.04707 =	Factor which includes the density of water
	(0.002201 lb/ml), the molecular weight of water
	(18.0 lb/lb-mole), the ideal gas constant
	21.85 (in. Hg) (ft <sup>3</sup> )/lb-mole)(deg R); absolute
	temperature at standard conditions (528 deg R), absolute
	pressure at standard conditions (29.92 in. Hg), ft <sup>3</sup> /ml.
0.04715 =	Factor which includes the molecular weight of water
	(18.0 lb/lb-mole), the ideal gas constant
	21.85 (in. Hg) (ft <sup>3</sup> )/lb-mole)(deg R); absolute
	temperature at standard conditions (528 deg R), absolute
	pressure at standard conditions (29.92 in. Hg), and
	$453.6 \text{ g/lb, ft}^3/\text{g.}$

2/19/201911:17 AM 011719 polymer

#### 3. Moisture content

$$bws = \begin{array}{c} Vw(std) \\ ------ \\ Vw(std) + Vm(std) \\ \\ bws = \\ ----- = 0.020 \\ 1.21 + 58.636 \\ \end{array}$$

Where:

bws = Proportion of water vapor, by volume, in the gas stream, dimensionless.

#### 4. Mole fraction of dry gas.

$$Md = 1 - bws$$

$$Md = 1 - 0.020 = 0.980$$

Where:

Md = Mole fraction of dry gas, dimensionless.

#### 5. Dry molecular weight of gas stream, lb/lb-mole.

$$MWd = (0.440 \times \% CO_2) + (0.320 \times \% O_2) + (0.280 \times (\% N_2 + \% CO))$$

$$MWd = (0.440 \times 0.0) + (0.320 \times 20.9) + (0.280 \times (79.1 + 0.00))$$

$$MWd = 28.84$$

Where:

MWd = Dry molecular weight, lb/lb-mole.

% CO2 = Percent carbon dioxide by volume, dry basis.

 $\% O_2 =$  Percent oxygen by volume, dry basis.

% N<sub>2</sub> = Percent nitrogen by volume, dry basis.

% CO = Percent carbon monoxide by volume, dry basis. 0.440 = Molecular weight of carbon dioxide, divided by 100.

0.320 = Molecular weight of carbon dioxide, divided by 100.

0.280 = Molecular weight of nitrogen or carbon monoxide,

divided by 100.

#### 6. Actual molecular weight of gas stream (wet basis), lb/lb-mole.

$$MWs = (MWd x Md) + (18 x (1 - Md))$$

$$MWs = (28.84 \times 0.980) + (18(1 - 0.980)) = 28.62$$

Where:

MWs = Molecular weight of wet gas, lb/lb-mole. 18 = Molecular weight of water, lb/lb-mole.

#### 7. Average velocity of gas stream at actual conditions, ft/sec.

Vs = 
$$85.49 \times 0.84 \times 0.72582 \times (------)^1/2 = 40.6$$

Where:

Vs = Average gas stream velocity, ft/sec.

(lb/lb-mole)(in. Hg)<sup>1/2</sup>

85.49 = Pitot tube constant, ft/sec x -----

(deg R)(in H<sub>2</sub>O)

Cp = Pitot tube coefficient, dimensionless.

 $Ts = \qquad \quad Absolute \ gas \ stream \ temperature, \ deg \ R = Ts, \ deg \ F + 460.$ 

P(static

delt p = Velocity head of stack, in. H<sub>2</sub>O.

#### 8. Average gas stream volumetric flow rate at actual conditions, wacf/min.

$$Qs(act) = 60 x Vs x As$$

$$Qs(act) = 60 \times 40.6 \times 4.91 = 11957$$

Where:

Qs(act) = Volumetric flow rate of wet stack gas at actual

conditions, wacf/min.

As = Cross-sectional area of stack,  $ft^2$ .

60 = Conversion factor from seconds to minutes.

#### 9. Average gas stream dry volumetric flow rate at standard conditions, dscf/min.

$$Qs(std) = \begin{array}{c} P_S \\ 17.64 \text{ x Md x} & ---- \text{ x Qs(act)} \\ T_S \end{array}$$

$$Qs(std) = 11910$$

Where:

Qs(std) = Volumetric flow rate of dry stack gas at standard conditions, dscf/min.

#### ${\bf 10.}\ Is okinetic\ variation\ calculated\ from\ intermediate\ values,\ percent.$

17.327 x Ts x Vm(std) I = Vs x O x Ps x Md x (Dn)<sup>2</sup> 17.327 x 522 x 58.636 ----= 97.2 I = 40.6 x 96 x 30.08 x 0.980 x (0.218)^2 Where: I = Percent of isokinetic sampling. 0 = Total sampling time, minutes. Dn = Diameter of nozzle, inches. 17.327 = Factor which includes standard temperature (528 deg R), standard pressure (29.92 in. Hg), the formula for calculating area of circle  $D^{2/4}$ , conversion of square feet to square inches (144), conversion of seconds to minutes (60), and conversion to percent (100), (in. Hg)(in2)(min) (deg R)(ft<sup>2</sup>)(sec)

2/19/201911:17 AM 011719 polymer

# APPENDIX E EQUIPMENT CALIBRATION RECORDS

# Sample and Velocity Traverse Point Data Sheet - Method 1

Client Chemours  Loaction/Plant Fayetteville  Source Polymers			<del></del>	Operator <u>SK</u> Date <u>3/2//18</u> W.O. Number			
Duct Type		Circular		Rectangular Duct	indicate appropriate type		
Traverse Type		Particulate Traverse		Velocity Traverse	☐ CEM Traverse		

Distance from far wall to outside of port (in.) = C	48
Port Depth (in.) = D	14
Depth of Duct, diameter (in.) = C-D	30
Area of Duct (ft <sup>2</sup> )	4.91
Total Traverse Points	24
Total Traverse Points per Port	17
Port Diameter (in.) —(Flange-Threaded-Hole)	
Monorail Length	
Rectangular Ducts Only	
Width of Duct, rectangular duct only (in.)	
Total Ports (rectangular duct only)	
Equivalent Diameter = (2°L*W)/(L+W)	

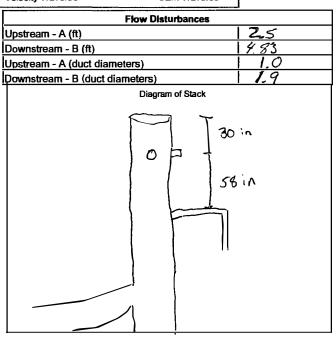
Traverse Point Locations										
		Distance from								
Traverse		Inside Duct	Distance from Outside of							
Point	% of Duct	Wall (in)	Port (in)							
1	21		19							
2	67	2	70							
3	119	3/2	21/12							
4	17.7	51/4	23 1/4							
5	25	7/2	251/2							
6	356	10 5/8	24 5/8							
7	64.4	193/8	37 3/8							
8	75	221/2	40 1/2							
9	82.3	2434	42 3/4							
10	S8 Z	761/2	44							
11	933	28	46							
12	97.9	79	47							
CEM	3 Point(Long L	Annument Line) Str	dification Point Locations							
1	0.167									
2	0.50									
3	0.833									
N	ote: If stack	dia < 12 inch us	e EPA Method 1A							

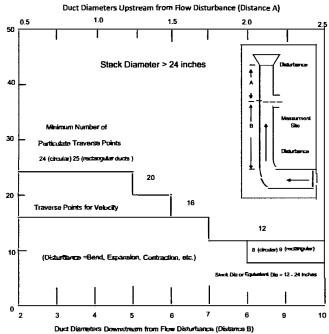
(Sample port upstream of pitot port)

Note: If stack dia >24" then adjust traverse point to 1 inch from wall

If stack dia <24" then adjust traverse point to 0.5 inch from wall

		Traverse Point Location Percent of Stack -Circular															
			Number of Traverse Points														
		$\Box$	l	2	Π	3	4	Π	5	6	7	8	9	T	10	11	12
Т	1	1		14.6			6.7	Ι		4.4		3.2 [		ī	2.6		2.1
r	2	T		85.4			25	Γ		14.6		10.5		1	8.2	1	6.7
a	3			1	Γ	_	75	1		29.6		19.4		ī	14.6		11.8
V L	4	T			Г		93.3	Г		70.4		32.3		T	22.6		17.7
1, 0	15	Π			ī			Ī		85.4		67.7		ī	34.2		25
8 8	6	1		I				1		95.6	1	80.6		Ţ	65.8	l.	35.6
e t	7	Т		l .	Π			Ι				89.5		T	77.4	ļ	64.4
i	<u> 8</u>	Τ.						ı				96.8		T	85.4		75
0 1	T 9	П		l	ı			ı						1	91.8		82.3
1	10	ī			1			ı				1 1		1	97.4	l .	88.2
n	11	ī			Ī			1						ī			93.3
t	12	T		I	Π			1			1	1 1		T			97.9

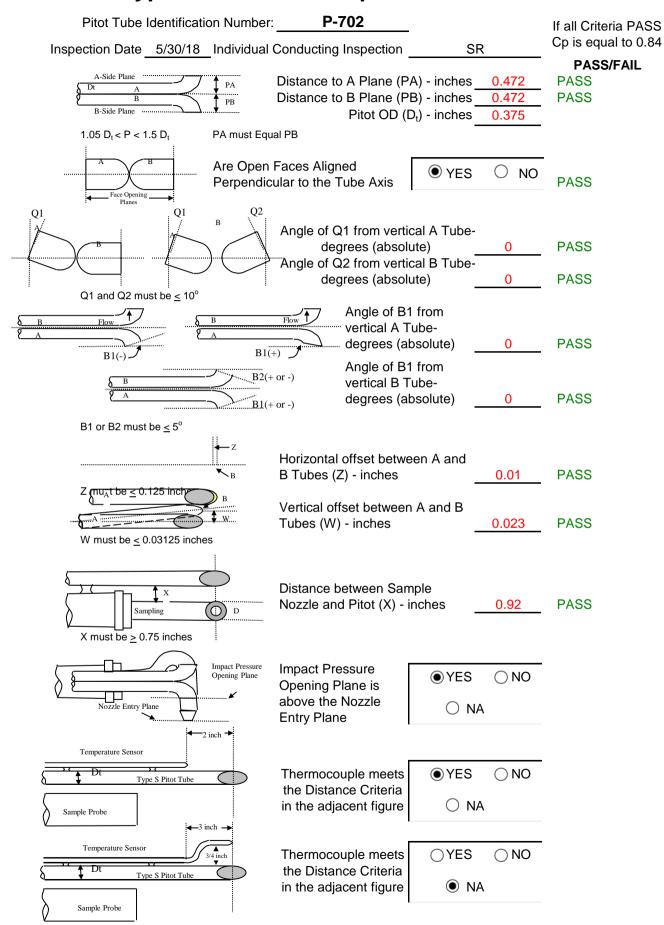




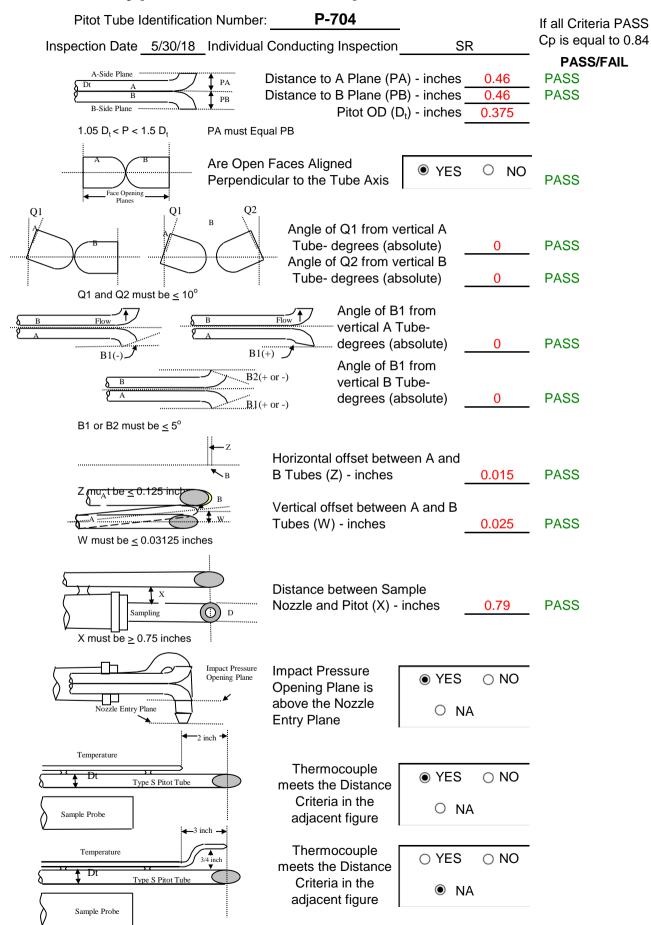
		Traverse Point Location Percent of Stack -Rectangular											
		Number of Traverse Points											
	Г	1	2	3	4	l 5	6	7	8	9	10	111	12
т   1	Т		25.0	16.7	12.5	10.0	8.3	7.1	6.3	5.6	5.0	4.5	4.2
r   2	1	1	75.0	50.0	37.5	30.0	25.0	21.4	18.8	16.7	15.0	13.6	12.5
1 3				83.3	62.5	50.0	41.7	35.7	31.3	27.8	25.0	22.7	20.8
4					87.5	70.0	58.3	50.0	43.8	38.9	35.0	31.8	29.2
5	Т					90.0	75.0	643	56.3	50.0	45.0	40.9	37.5
a   6				l			91.7	78.6	68.8	61.1	55.0	50.0	45.8
1 7	T						}	92.9	81.3	72.2	65.0	59.1	54.2
. [8	Ŧ.			1					93.8	83.3	75.0	68.2	62.5
, , , 79	Т			I		l	l		1	94.4	85.0	77.3	70.8
i   10	) [			1		1				l	95.0	86.4	79.2
n	П						i			1	<b>I</b> !	95.5	87.5
1 12	! [	. 1					1						95.8



# **Type S Pitot Tube Inspection Data Form**



# Type S Pitot Tube Inspection Data Form





# **CERTIFICATE OF ANALYSIS**

## **Grade of Product: EPA Protocol**

Part Number: E03NI79E15A00E4 Reference Number: 82-401288926-1

Cylinder Number: CC18055 Cylinder Volume: 150.5 CF Laboratory: 124 - Riverton (SAP) - NJ Cylinder Pressure: 2015 PSIG

PGVP Number: B52018 Valve Outlet: 590

Gas Code: CO2,O2,BALN Certification Date: Sep 04, 2018

Expiration Date: Sep 04, 2026

Certification performed in accordance with "EPA Traceability Protocol for Assay and Certification of Gaseous Calibration Standards (May 2012)" document EPA 600/R-12/531, using the assay procedures listed. Analytical Methodology does not require correction for analytical interference. This cylinder has a total analytical uncertainty as stated below with a confidence level of 95%. There are no significant impurities which affect the use of this calibration mixture. All concentrations are on a volume/volume basis unless otherwise noted.

Do Not Use This Cylinder below 100 psig, i.e. 0.7 megapascals.

			ANALYTICA	L RESULTS						
Compon	ent	Requested Actual Protocol Total Relative Assay Concentration Method Uncertainty Dates								
CARBON	DIOXIDE	9.000 %	8.864 %	G1	+/- 0.7% NIST Traceab	le 09/04/2018				
OXYGEN		12.00 %	12.00 %	G1	+/- 0.4% NIST Traceab	le 09/04/2018				
NITROGE	N	Balance			-					
CALIBRATION STANDARDS										
Type	Lot ID	Cylinder No	Concentration		Uncertainty	<b>Expiration Date</b>				
NTRM	13060629	CC413730	13.359 % CARBON [	DIOXIDE/NITROGEN	+/- 0.6%	May 09, 2019				
			ANALYTICAL	EQUIPMENT	1					
Instrume	ent/Make/Mod	el	Analytical Princi	•	Last Multipoint Calibration					
Horiba VIA	A 510-CO2-19G	YCXEG	NDIR		Aug 09, 2018					
Horiba MF	PA 510-O2-7TW	MJ041	Paramagnetic		Aug 09, 2018					

**Triad Data Available Upon Request** 





# **CERTIFICATE OF ANALYSIS**

## **Grade of Product: EPA Protocol**

Part Number: E03NI62E15A0224 Reference Number: 82-401044874-1

Cylinder Number: SG9169108 Cylinder Volume: 157.2 CF Laboratory: 124 - Riverton (SAP) - NJ Cylinder Pressure: 2015 PSIG

PGVP Number: B52017 Valve Outlet: 590

Gas Code: CO2,O2,BALN Certification Date: Nov 18, 2017

Expiration Date: Nov 18, 2025

Certification performed in accordance with "EPA Traceability Protocol for Assay and Certification of Gaseous Calibration Standards (May 2012)" document EPA 600/R-12/531, using the assay procedures listed. Analytical Methodology does not require correction for analytical interference. This cylinder has a total analytical uncertainty as stated below with a confidence level of 95%. There are no significant impurities which affect the use of this calibration mixture. All concentrations are on a volume/volume basis unless otherwise noted.

Do Not Use This Cylinder below 100 psig, i.e. 0.7 megapascals.

			ANALYTICA	L RESULTS		
Compon	ent	Requested Concentration	Actual Concentration	Protocol Method	Total Relative Uncertainty	Assay Dates
CARBON	RBON DIOXIDE 17.00 % 16.58 % G1 +/- 0.7% NIST Traceable				11/18/2017	
OXYGEN 21.00 %		21.00 %	G1	+/- 0.5% NIST Traceable	11/18/2017	
NITROGE	OGEN Balance -					
			CALIBRATION	STANDARDS	8	
Type	Lot ID	Cylinder No	Concentration		Uncertainty	<b>Expiration Date</b>
NTRM	12061336	CC360792	11.002 % CARBON [	DIOXIDE/NITROGEN	+/- 0.6%	Jan 11, 2018
NTRM	09061415	CC273526	22.53 % OXYGEN/NI	TROGEN	+/- 0.4%	Mar 08, 2019

ANALYTICAL EQUIPMENT
Instrument/Make/Model Analytical Principle Last Multipoint Calibration

Horiba VIA 510-CO2-19GYCXEG NDIR Oct 30, 2017
Horiba MPA 510-O2-7TWMJ041 Paramagnetic Oct 27, 2017

**Triad Data Available Upon Request** 



#### INTERFERENCE CHECK

Date: 12/4/14-12/5/14 Analyzer Type: Servomex - O<sub>2</sub> Model No: 4900 Serial No: 49000-652921 Calibration Span: 21.09 % Pollutant: 21.09% O<sub>2</sub> - CC418692

: Check 2014O2-Servomex 4900

	ANALYZEF	RESPONSE	.,				
INTERFERENT GAS	INTERFERENT GAS RESPONSE (%)	INTERFERENT GAS RESPONSE, WITH BACKGROUND POLLUTANT (%)	% OF CALIBRATION SPAN <sup>(a)</sup>				
CO <sub>2</sub> (30.17% CC199689)	0.00	-0.01	0.00				
NO (445 ppm CC346681)	0.00	0.02	0.11				
NO <sub>2</sub> (23.78 ppm CC500749)	NA	NA	NA				
N <sub>2</sub> O (90.4 ppm CC352661)	0.00	0.05	0.24				
CO (461.5 ppm XC006064B)	0.00	0.02	0.00				
SO <sub>2</sub> (451.2 ppm CC409079)	0.00	0.05	0.23				
CH <sub>4</sub> (453.1 ppm SG901795)	NA	NA	NA				
H <sub>2</sub> (552 ppm ALM048043)	0.00	0.09	0.44				
HCl (45.1 ppm CC17830)	0.00	0.03	0.14				
NH <sub>3</sub> (9.69 ppm CC58181)	0.00	0.01	0.03				
	TOTAL INTERFERENCE RESPONSE						
	METHOD SPECIFICATION		< 2.5%				

<sup>(</sup>a) The larger of the absolute values obtained for the interferent tested with and without the pollutant present was used in summing the interferences.

Chad Walker

2/19/2019

#### INTERFERENCE CHECK

Date: 12/4/14-12/5/14 Analyzer Type: Servomex - CO<sub>2</sub> Model No: 4900 Serial No: 49000-652921 <u>Calibration Span: 16.65%</u> <u>Pollutant: 16.65% CO<sub>2</sub> - CC418692</u>

	ANALYZER RESPONSE										
INTERFERENT GAS	INTERFERENT GAS RESPONSE (%)	INTERFERENT GAS RESPONSE, WITH BACKGROUND POLLUTANT (%)	% OF CALIBRATION SPAN <sup>(a)</sup>								
CO <sub>2</sub> (30.17% CC199689)	NA ·	NA	NA								
NO (445 ppm CC346681)	0.00	0.02	0.10								
NO <sub>2</sub> (23.78 ppm CC500749)	0.00	0.00	0.02								
N <sub>2</sub> O (90.4 ppm CC352661)	0.00	0.01	0.04								
CO (461.5 ppm XC006064B)	0.00	0.01	0.00								
SO <sub>2</sub> (451.2 ppm CC409079)	0.00	0.11	0.64								
CH <sub>4</sub> (453.1 ppm SG901795)	0.00	0.07	0.44								
H <sub>2</sub> (552 ppm ALM048043)	0.00	0.04	0.22								
HCl (45.1 ppm CC17830)	0.10	0.06	0.60								
NH <sub>3</sub> (9.69 ppm CC58181)	0.00	0.02	0.14								
	2.19										
	METHOD SPECIFICATION		< 2.5%								

<sup>(</sup>a) The larger of the absolute values obtained for the interferent tested with and without the pollutant present was used in summing the interferences.

Chad Walker

: Check 2014CO2-Servomex 4900 2/19/2019

# Long Cal and Temperature Cal Datasheet for Standard Dry Gas Meter Console

Calibrator MDW

Meter Box Number 12

Ambient Temp

Thermocouple Simulator

Date 10-Sep-18

**Wet Test Meter Number** P-2952 Temp Reference Source

(Accuracy +/- 1°F)

**Dry Gas Meter Number** 14244707

Baro Press, in	29.96
Hg (Pb)	29.90

Setting	Gas	Volume	Tempe	ratures				
Orifice	Wet Test	Dry gas Meter	Wet Test	Dry Gas				
Manometer	Meter	Dry guo motor	Meter	Meter		Calibration	Calibration Results	
in H <sub>2</sub> 0	ft <sup>3</sup>	ft <sup>3</sup>	°F	Outlet, °F	Time, min	Υ	ΔН	
(∆H)	(Vw)	(Vd)	(Tw)	(Td <sub>o</sub> )	(O)	'	ΔП	
		885.853		75.00				
0.5	5.0	890.822	73.0	76.00	12.60	1.0097	1.7823	
	4.969		75.50					
		892.810		76.00		1.0071	1.8559	
1.0 5.0	5.0	897.795	73.0	77.00	9.1			
		4.985		76.50				
		898.799		77.00		1.0036	1.9381	
1.5	10.0	908.810	73.0	78.00				
		10.011		77.50				
		915.870		78.00				
2.0	10.0	925.830	73.0	79.00	13.1	1.0094	1.9158	
		9.960		78.50				
		926.870		79.00				
3.0	10.0	936.870	73.0	80.00	10.70	1.0048	1.9137	
		10.000		79.50				
						1.0069	1.8812	

Vw - Gas Volume passing through the wet test meter

Vd - Gas Volume passing through the dry gas meter

Tw - Temp of gas in the wet test meter

Tdi - Temp of the inlet gas of the dry gas meter

Tdo - Temp of the outlet gas of the dry gas meter

Td - Average temp of the gas in the dry gas meter

0 - Time of calibration run

Pb - Barometric Pressure

ΔH - Pressure differential across

orifice

Y - Ratio of accuracy of wet test

meter to dry gas meter

$$Y = \frac{Vw * Pb * (td + 460)}{Vd * \left[Pb + \frac{(\Delta H)}{13.6}\right] * (tw + 460)}$$

$$\Delta H = \left[\frac{0.0317 * \Delta H}{Pb * (td + 460)}\right] * \left[\frac{\left(tw + 460\right) * O}{Vw}\right]^{2}$$

Reference Temperature	ire	Temperature l	Reading from I	ndividual Therr	nocouple Input	1	Average Temperature	Temp Difference <sup>2</sup>
0℃			Chann	el Number			Reading	(%)
	1	2	3	4	5	6		` ′
32	32	32	32	32	32	32	32.0	0.0%
212	212	212	212	212	212	212	212.0	0.0%
932	932	932	932	932	932	932	932.0	0.0%
1832	1834	1834	1834	1834	1834	1834	1834.0	-0.1%

1 - Channel Temps must agree with +/- 5°F or 3°C

2 - Acceptable Temperature Difference less than 1.5 %

(Reference Temp(°F)+460) – (Test Temp(°F)+460) Temp Diff = Reference Temp(°F)+460



#### **Y Factor Calibration Check Calculation**

#### POLYMER STACK METER BOX NO. 12 1/17/2019-1/18/2019

	Kun i	Kun 2	Kuli 3
MWd = Dry molecular weight source gas, lb/lb-mole.			
0.32 = Molecular weight of oxygen, divided by 100.			
0.44 = Molecular weight of carbon dioxide, divided by 100.			
0.28 = Molecular weight of nitrogen or carbon monoxide, divided by 100.			
% CO <sub>2</sub> = Percent carbon dioxide by volume, dry basis.	0.0	0.0	0.0
% O <sub>2</sub> = Percent oxygen by volume, dry basis.	20.9	20.9	20.9

 $MWd = (0.32 * O_2) + (0.44 * CO_2) + (0.28 * (100 - (CO_2 + O_2)))$ 

MWd = (0.32 \* 20.9) + (0.44 \* 0) + (0.28 \* (100 - (0 + 20.9)))

MWd = (6.69) + (0.00) + (22.15)

MWd =

Tma = Source Temperature, absolute(°R)

Tm = Average dry gas meter temperature, deg F.

64.7 49.4 61.7

Tma = Ts + 460

Tma = 64.67 + 460

**Tma** = 524.67 509.42 521.67

28.84

28.84

28.84

Ps = Absolute meter pressure, inches Hg.			
13.60 = Specific gravity of mercury.			
delta H = Avg pressure drop across the orifice meter during sampling, in H2O	1.31	1.43	1.50
Pb = Barometric Pressure, in Hg.	30.10	30.01	30.01

Pm = Pb + (delta H / 13.6)

Pm = 30.1 + (1.3125 / 13.6)

Pm = 30.20 30.12 30.12

Yqa = dry gas meter calibration check value, dimensionless.			
0.03 = (29.92/528)(0.75)2 (in. Hg/°/R) cfm2.			
29.00 = dry molecular weight of air, lb/lb-mole.			
Vm = Volume of gas sample measured by the dry gas meter at meter conditions, dcf.	57.360	58.671	60.644
Y = Dry gas meter calibration factor (based on full calibration)	1.0069	1.0069	1.0069
Delta H@ = Dry Gas meter orifice calibration coefficient, in. H2O.	1.8812	1.8812	1.8812
avg SQRT Delta $H = Avg \ SQRT \ press.$ drop across the orifice meter during sampling , in. $H_2O$	1.1390	1.1950	1.2238
O = Total sampling time, minutes.	96	96	96

Yqa = (O / Vm) \* SQRT (0.0319 \* Tma \* 29) / (Delta H@ \* Pm \* MWd) \* avg SQRT Delta H

Yqa = (96.00 / 57.36) \* SQRT (0.0319 \* 524.67 \* 29) / (1.88 \* 30.20 \* 28.84) \* 1.14

Yqa = 1.674 \* SQRT 485.369 / 1,638.238 \* 1.14

**Yqa** = 1.0376 1.0501 1.0501

Diff = Absolute difference between Yqa and Y 3.05 4.29 4.29

Diff = (( Y - Yqa ) / Y ) \* 100

Diff = (( 1.0069 - 1.038 ) / 1.0069 ) \* 100

Average Diff = 3.88

Allowable = 5.0

2/19/201911:47 AM 011719 polymer

# APPENDIX F LIST OF PROJECT PARTICIPANTS

The following WESTON employees participated in this project.

Paul Meeter	Senior Project Manager
Wes Fritz	Team Member
Matt Winkeler	Team Member
Steve Rathfon	Team Member
Kyle Schweitzer	Team Member