FLUOROMONOMERS MANUFACTURING PROCESS DIVISION STACK EMISSIONS TEST REPORT TEST DATES: 16 AND 17 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. 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 Division Stack at the facility. Testing was performed on 16 and 17 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 Division stack which is located in the Fluoromonomers process area.
- Monitor and record process and emissions control 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 were measured at the Division 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 location. 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 package is provided in electronic format and on CD with each hard copy.

Table 1-1
Sampling Plan for Division Stack Testing

Sampling Point & Location		Divi	sion Stack						
Number of Tests:			3						
Parameters To Be Tested:	HFPO Dimer Acid (HFPO-DA)	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 M	13/3A	EPA M4 in conjunction with M-0010 tests				
Sample Extraction/ Analysis Method(s):	LC/MS/MS	NA ⁶	N/	4	NA				
Sample Size	$\geq 1.5 \text{m}^3$	NA	NA	NA	NA				
Total Number of Samples Collected ¹	3	3	3	3	3				
Reagent Blanks (Solvents, Resins)1	1 set	0	0	0	0				
Field Blank Trains ¹	0 per source	0	0	0	0				
Proof Blanks ¹	1 per train	0	0	0	0				
Trip Blanks ^{1,2}	1 set	0	0	0					
Lab Blanks	1 per fraction ³	0	0	0	0				
Laboratory or Batch Control Spike Samples (LCS)	1 per fraction ³	0	0	0	0				
Laboratory or Batch Control Spike Sample Duplicate (LCSD)	1 per fraction ³	0	0	0	0				
Media Blanks	1 set ⁴	0	0	0	0				
Isotope Dilution Internal Standard Spikes	Each sample	0	0	0	0				
Total No. of Samples	65	3	3	3	3				

Key:

¹ Sample collected in field.

² Trip blanks include one XAD-2 resin module and one methanol sample per sample shipment.

³ Lab blank and LCS/LCSD includes one set per analytical fraction (front half, back half and condensate).

⁴ One set of media blank archived at laboratory at media preparation.

⁵ Actual number of samples collected in field.

⁶ Not applicable.

2. SUMMARY OF TEST RESULTS

A total of three test runs were performed on the Division stack. Table 2-1 provides a summary of the HFPO Dimer Acid emissions 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

Source	Run No.	Emission Rates									
Source	Kuli 140.	lb/hr	g/sec								
Division Stack	1	4.70E-03	5.92E-04								
	2	4.90E-03	6.17E-04								
	3	4.45E-03	5.60E-04								
	Average	4.68E-03	5.90E-04								

3. PROCESS DESCRIPTIONS

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

3.1 FLUOROMONOMERS

These facilities produce a family of fluorocarbon compounds used to produce Chemours products such as Nafion®, Krytox®, and Viton®, as well as sales to outside customers.

Process emissions are vented to the Division waste gas scrubber system (which includes the secondary scrubber) and vents to the Carbon Bed and then onto the Division Stack. VE North building air also vents to the Carbon Bed and then onto the Division Stack.

3.2 PROCESS OPERATIONS AND PARAMETERS

The following table is a summary of the operation and products from the specific areas tested.

Source	Operation/Product	Batch or Continuous
VE North	PSEPVE	Condensation is continuous. Agitated Bed Reactor and Refining are batch.
HFPO Tower	HFPO	Continuous.

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

- Fluoromonomers Process
 - VEN Precurser Rate
 - VEN Condensation Rate
 - o VEN ABR Rate
 - o HFPO

4. DESCRIPTION OF TEST LOCATIONS

4.1 DIVISION STACK

Two 6-inch ID test ports were installed on the 36-inch ID fiberglass stack as shown below. The four vents that enter the top of the stack and the one vent ~11 feet below are catch pots which, under normal process operations, do not discharge to the stack. They are used to vent process gas to the stack in the event of a process upset and are not considered a flow contributor or a disturbance.

Per EPA Method 1, a total of 12 traverse points (six per axis) were used for M-0010 isokinetic sampling. Figure 4-1 provides a schematic of the test ports and traverse point locations.

Location	Distance from Fl	low Disturbance
Location	Downstream (B)	Upstream (A)
Division Stack	30 feet > 10 duct diameters	9 feet > 3 diameters

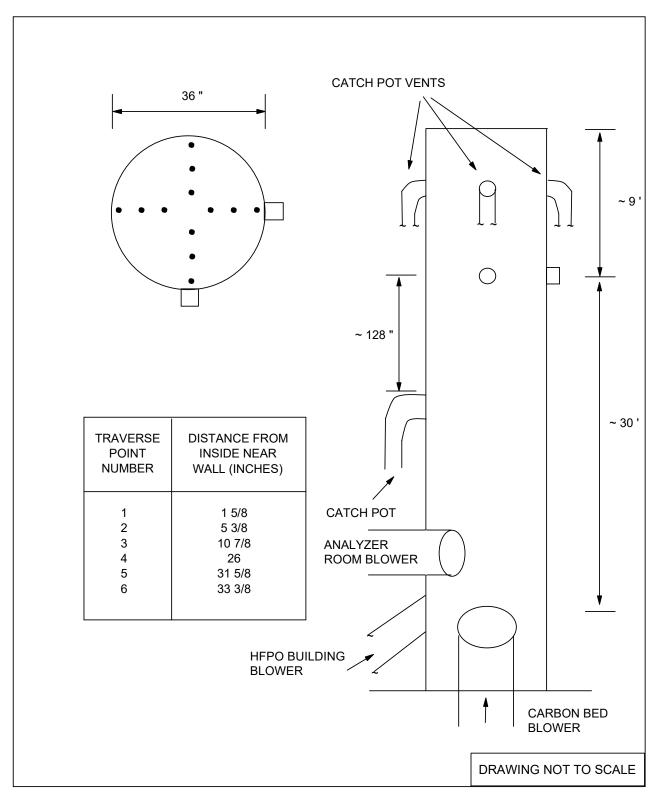


FIGURE 4-1
DIVISION STACK TEST PORT
AND TRAVERSE POINT LOCATIONS

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 each 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°) verifying that the test location 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.

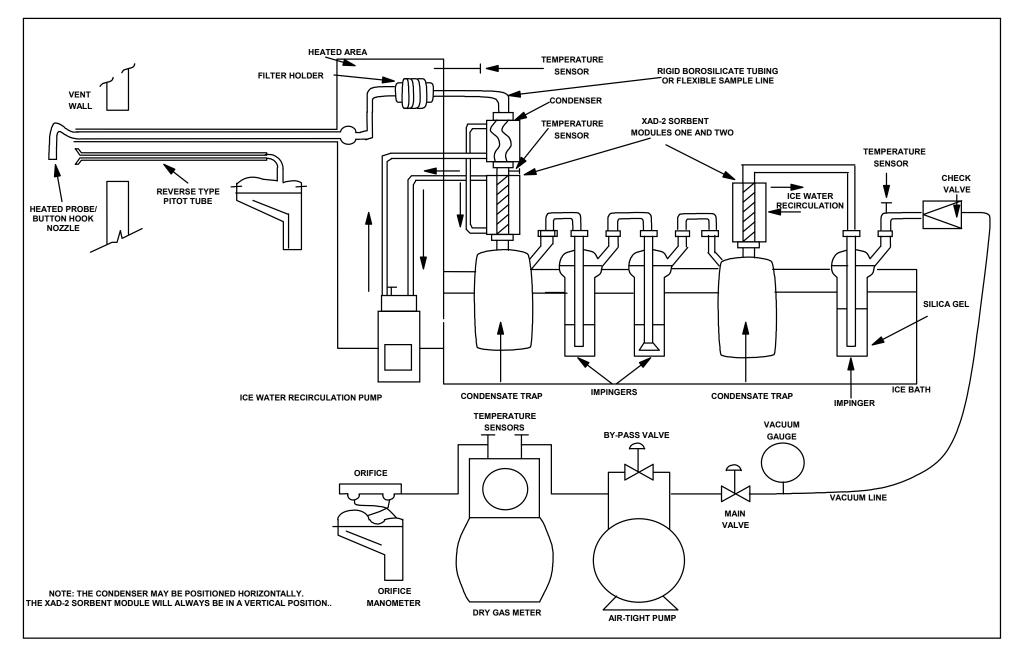


FIGURE 5-1
EPA METHOD 0010 SAMPLING TRAIN

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 underwent hydrolysis instantaneously in water in the sampling train and during the sample recovery step, and was converted to HFPO Dimer Acid such that the amount of HFPO Dimer Acid emissions represented 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 at $100\% \pm 10$. 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 remains. 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 impingers one and two, and the second condensate trap, were measured, the values recorded, and the sample was placed in the same container as Step 4 above, then 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.

See Figure 5-2 for a schematic of the Method 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 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 (CO₂) and oxygen (O₂) 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 maintained 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 insure 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.

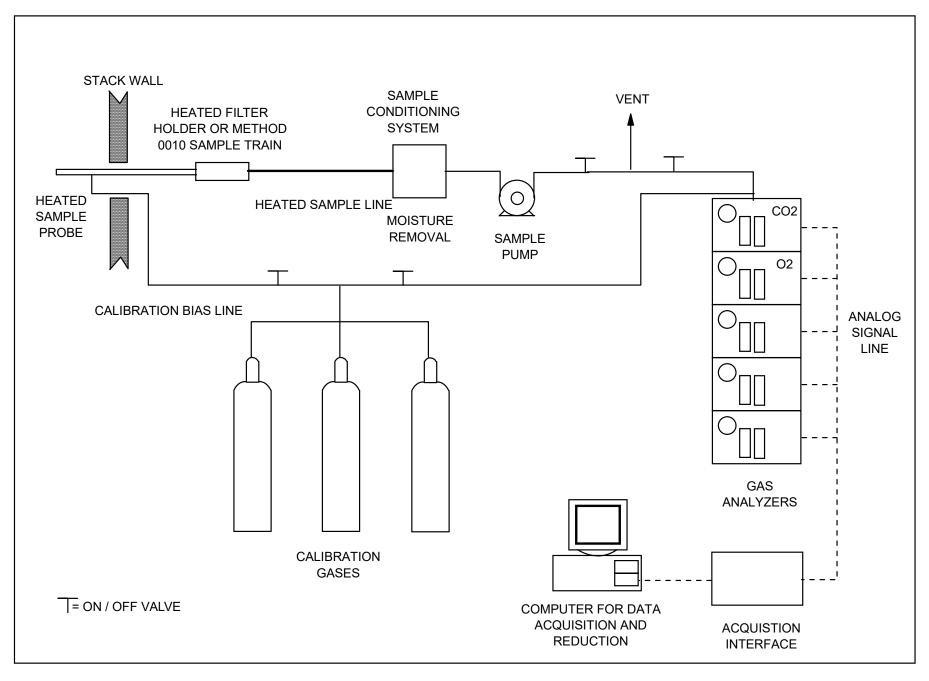


FIGURE 5-3
WESTON SAMPLING SYSTEM

6. DETAILED TEST RESULTS AND DISCUSSION

Each test was a minimum of 96 minutes in duration. A total of three test runs were performed at each location.

Table 6-1 provides detailed test data and test results for the Division stack.

The Method 3A sampling indicated that the O₂ and CO₂ concentrations were at ambient air levels (20.9% O₂, 0% CO₂), therefore, 20.9% O₂ and 0% CO₂ values were used in all calculations.

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

Test Data			
Run number	1	2	4
Location	Divison Stack	Divison Stack	Divison Stack
Date	1/16/19	1/16/19	1/17/19
Time period	0941-1140	1312-1513	0842-1035
SAMPLING DATA:			
Sampling duration, min.	96.0	96.0	96.0
Nozzle diameter, in.	0.160	0.160	0.160
Cross sectional nozzle area, sq.ft.	0.000140	0.000140	0.000140
Barometric pressure, in. Hg	30.15	30.06	30.18
Avg. orifice press. diff., in H ₂ O	1.14	1.13	1.15
Avg. dry gas meter temp., deg F	49.1	59.2	40.9
Avg. abs. dry gas meter temp., deg. R	509	519	501
Total liquid collected by train, ml	21.5	21.1	21.6
Std. vol. of H ₂ O vapor coll., cu.ft.	1.0	1.0	1.0
Dry gas meter calibration factor	1.0069	1.0069	1.0069
Sample vol. at meter cond., dcf	53.660	53.775	53.146
Sample vol. at std. cond., dscf (1)	56.604	55.451	57.033
Percent of isokinetic sampling	98.5	98.0	97.9
GAS STREAM COMPOSITION DATA:			
CO ₂ , % by volume, dry basis	0.0	0.0	0.0
O ₂ , % by volume, dry basis	20.9	20.9	20.9
N ₂ , % 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 ₂ 0 vapor in gas stream, prop. by vol.	0.018	0.018	0.018
Mole fraction of dry gas	0.982	0.982	0.982
Molecular wt. of wet gas, lb/lb mole	28.65	28.65	28.65
GAS STREAM VELOCITY AND VOLUMETRIC FLOW DATA:			
Static pressure, in. H ₂ O	-0.70	-0.70	-0.70
Absolute pressure, in. Hg	30.10	30.01	30.13
Avg. temperature, deg. F	44	46	37
Avg. absolute temperature, deg.R	504	506	497
Pitot tube coefficient	0.84	0.84	0.84
Total number of traverse points	12	12	12
Avg. gas stream velocity, ft./sec.	69.0	68.4	69.0
Stack/duct cross sectional area, sq.ft.	7.07	7.07	7.07
Avg. gas stream volumetric flow, wacf/min.	29268	29016	29252
Avg. gas stream volumetric flow, dscf/min.	30315	29840	30735

⁽¹⁾ 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 DIVISION STACK

TEST D	AIA	

120121111			
Run number	1	2	3
Location	Divison Stack	Divison Stack	Divison Stack
Date	1/16/19	1/16/19	1/17/19
Time period	0941-1140	1312-1513	0842-1035
LABORATORY REPORT DATA, ug.			
HFPO Dimer Acid	66.35	68.82	62.45
EMISSION RESULTS, ug/dscm.			
HFPO Dimer Acid	41.38	43.82	38.66
EMISSION RESULTS, lb/dscf.			
HFPO Dimer Acid	2.58E-09	2.74E-09	2.41E-09
EMISSION RESULTS, lb/hr.			
HFPO Dimer Acid	4.70E-03	4.90E-03	4.45E-03
EMISSION RESULTS, g/sec.			
HFPO Dimer Acid	5.92E-04	6.17E-04	5.60E-04

APPENDIX A PROCESS OPERATIONS DATA

Date 1/16/2019																																			
Time		800			900			10	.000		1100		.00		1200				130				14	00		1500				1600					
Stack Testing						RUN 1 941-1			1140				RUN 2 - 1312-1513							13															
HFPO																																			
VEN Product			PSEPVE																																
VEN Precursor																																			
VEN Condensation (HFPO)																																			
VEN ABR																																			
VEN Refining																																			
Stripper Column Vent																																			
Division WGS Recirculation Flow																14	400	00 kg	g/h																
Division WGS Inlet Flow	80	0 kg/l	kg/h 107 kg/h																																

Date 1/17/2019																
Time		80	00			900			10	00		1100				
Stack Testing					084	2-103										
HFPO																
VEN Product		PSEPVE														
VEN Precursor																
VEN Condensation (HFPO)																
VEN ABR																
VEN Refining		Ва	tch													
Stripper Column Vent																
Division WGS Recirculation Flow							1400	0 kg	/h							
Division WGS Inlet Flow				100 kg/h												

APPENDIX B RAW AND REDUCED TEST DATA

CHEMOURS - FAYETTEVILLE, NC INPUTS FOR HFPO DIMER ACID CALCULATIONS DIVISION STACK

Test Data			
Run number	1	2	3
Location	Divison Stack	Divison Stack	Divison Stack
Date	1/16/19	1/16/19	1/17/19
Time period	0941-1140	1312-1513	0842-1035
Operator	MW	MW	MW
Inputs For Calcs.			
Sq. rt. delta P	1.25716	1.24191	1.26546
Delta H	1.1436	1.1278	1.1545
Stack temp. (deg.F)	43.6	45.7	37.0
Meter temp. (deg.F)	49.1	59.2	40.9
Sample volume (act.)	53.660	53.775	53.146
Barometric press. (in.Hg)	30.15	30.06	30.18
Volume H ₂ O imp. (ml)	10.0	9.0	9.0
Weight change sil. gel (g)	11.5	12.1	12.6
% CO ₂	0.0	0.0	0.0
% O ₂	20.9	20.9	20.9
% N ₂	79.1	79.1	79.1
Area of stack (sq.ft.)	7.070	7.070	7.070
Sample time (min.)	96.0	96.0	96.0
Static pressure (in.H ₂ O)	-0.70	-0.70	-0.70
Nozzle dia. (in.)	0.160	0.160	0.160
Meter box cal.	1.0069	1.0069	1.0069
Cp of pitot tube	0.84	0.84	0.84
Traverse points	12	12	12

2/19/2019 9:32 AM 01162019 Division

Sample and Velocity Traverse Point Data Sheet - Method 1 Client CHCMOURS Operator Loaction/Plant Fagetteville N.C. Source W.O. Number Duct Type Circular Rectangular Duct Particulate Traverse Traverse Type Velocity Traverse □ CEM Traverse Distance from far wall to outside of port (In.) = C **Flow Disturbances** 1219 Port Depth (in.) = D Upstream - A (ft) Depth of Duct, diameter (in.) = C-D Downstream - B (ft) Area of Duct (ft²) Upstream - A (duct diameters) Downstream - B (duct diameters) Total Traverse Points Total Traverse Points per Port Diagram of Stack Port Diameter (in.) —(Flange-Threaded-Hole) 411 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 Distance from Outside of Inside Duct Traverse Point % of Duct Wall (in) /Port (in) 20 1,62 3 Duct Diameters Upstream from Flow Disturbance (Distance A) 5/er 5 54 3/E 6 Stack Diameter > 24 inches 40 7 MANA 8 Minimum Number of 30 Particulate Traverse Points 10 24 (ctroular) 25 (rectangular ducts.) 11 12 20 16 Traverse Points for Velocity CEM 3 Point(Long Measurment Line) Stratification Point Locations 12 0.167 2 0.50 10 (Disturbance =Bend, Expansion, Contraction, etc.) 0.833 Note: If stack dla < 12 inch use EPA Method 1A (Sample port upstream of pitot port) Note: If stack dia >24" then adjust traverse point to 1 inch from wall 0 If stack dia <24° then adjust traverse point to 0.5 inch from wall Duct Diameters Downstream from Flow Disturbance (Distance B) Traverse Point Location Percent of Stack -Circular Number of Traverse Points Traverse Point Location Percent of Stack -Rectangular Number of Traverse Points 1 1 2 1 6 11 | 12 146 6.7 4.4 3.2 1 2 6 1 21 25.0 | 16.7 | 12.5 | 10.0 | 8.3 | 7.1 | 6.3 | 5.6 | 5.0 | 4.5 | 4.2 1 82 2 85.4 - 36 14.5 10.5 6.7 75.0 | 30.0 | 37.5 | 30.0 | 26.0 | 21.4 | 18.8 | 16.7 | 15.0 | 13.6 | 12.5 3 | 4 | 2 | 5 | 121 29.6 19.4 14.6 1118 3 | 833 | 62.5 | 50.0 | 41.7 | 35.7 | 31.3 | 27.8 | 25.0 | 22.7 | 20.8 93.3 76.4 32.3 17.7 22.6 4 c 5 87.5 70.0 58.3 50.0 43.8 38.9 35.0 31.8 29.2 90.0 75.0 64.3 56.3 50.0 45.0 40.9 37.5 67.7 85.4 34.2 95.6 65.8 35.6 a 6 91.7 | 78.6 | 68.8 | 61.1 | 55.0 | 50.0 | 45.8 | a | 6 | t 7 89.5 77.4 64.4 92.9 | 81.3 | 72.2 | 65.0 | 59.1 | 54.2 8 | 9 | 85.A 75 В 93.8 | 83.3 | 75.0 | 68.2 | 62.5

91.8

97.4

10

111

82.3

88.2

1 93.3

9 1

101

111



[94.4 | 85.0 | 77.3 | 70.8

95.0 | 86.4 | 79.2

ISOKINETIC FIELD DATA SHEET EPA Method 0010 - HFPO Dimer Acid '2_ Client **Stack Conditions** Chemours Meter Box ID K Factor ,0069 V W.O.# 15418 Assumed Actual Meter Box Y Project ID Chemours % Moisture Meter Box Del H Final Mid-Point . VS Initial Mode/Source ID Division Impinger Vol (ml) Probe ID / Length Sample Train (ft3) 100.0 100.6 Silica gel (g) Samp, Loc. ID STK Probe Material Boro Leak Check @ (in Hg) 26 do 215 Run No.ID $0.1 \, \text{V}$ Pitot / Thermocouple ID 1 CO2, % by Vol no / no you / no Ves / no Pitot leak check good 2043 Test Method ID M0010 O2. % by Vol Pitot Coefficient (es) / no 0.84 Pitot Inspection good no / no **gea** / no Date ID 16JAN2019 Temperature (°F) Nozzie ID 7-160 Method 3 System good yes / no yes / no yes / no Meter Temp (°F) Source/Location Division Stack 40 Nozzie Measurements 0.160 0.160 0.160 Temp Check Pre-Test Set Post-Test Set Sample Date 116119 Static Press (in H₂O) -0.70J-07 Avg Nozzie Dia (in) Sy 0.160 / Meter Box Temp 36 Baro, Press (in Hg) Area of Stack (ft²) 1.07 Reference Temp 240 MULLIN CELEA J Ambient Temp (°F) Operator Pass / Fail Sample Time Pass/Fall (+/- 20) Case / Fall Total Traverse Pts 」っ Temp Change Response 7 yes / no Veg / no

TRAVERSE POINT NO.		CLOCK TIME (plant time)	VELOCITY PRESSURE Delta	ORIFICE PRESSURE	DRY GAS METER READING (ft ³)	STACK TEMP (°F)	DGM OUTLET TEMP (°F)	PROBE TEMP (°F)	FILTER BOX TEMP	IMPINGER EXIT TEMP	SAMPLE TRAIN VAC	XAD EXIT TEMP (F)		COMMENTS
100	0	0941	P (in H2O)	Delta H (in H2O)	278,115				(F)	(oF)	(in Hg)			
	4	011	14	(),957	280,13	40	35	228	755	66	3	(0/0		
(8		1,5	1.02	222,51	40	35-	258	255	66	3	66		
2	12		1.8	1,28	275,10	40	35	255	255	62	_3	67	4	0713
	16		1.8	1.28	286,96	40	35	755	255	62	4	62		4-New
3.	ಬ		1,9	1,35	239.51	40	40	255	255	60	7	50		KIFACTOR
3	24		1,9	1,35	292.10	44	45	255	255	60	\$-	60		
4	28		1.8	1.28	294,35	44	49	255	251	60	5	60		
<u> </u>	32	, , , , , , , , , , , , , , , , , , ,	1.8	1128	296.55	44	50	255	251	54	5	54		27.160
	36		446	1,14	299.00	44	51	255	251	<u> 54</u>	5-	57		
2	<u>40</u>		1.6	1.14	30115	44	51	254	टेडप	7	2	51		
<u> </u>		12 3 Cl	4.2	0.855	303.25	44	_ کے	254	252	51	_5_	_51		
<u> </u>	48	1029	1.2	0.855	305,278	40	51	283	ટઽઽ	ر ک	_5	Si		
	4	1052	1,5	109	305,400	50		2	25-					2 2 2 2 2
-	\$		1.5	1.09	309,60	48	54 54	255	255	25	3	55		E0,728
2			1.3	1.87	312.00	मुँ भ	37	255 251	355	7 P	3	5 4		K-FACTOR
2	5		1.8	1.31	314,44	44	37	251	551 521	48	5	48		
3	20		1 9	1,32	317.21	44	54	251	251	3/2/	3	44		
3	24		1. 9	1 38	र्वि ३ म	44	54	251	231	42	-3-	42		26.500
7	スを		1. 8	7:31	321.05	45	रॅप	251	251	40	-3-	40		4 6: 300
14	32		1/2 8	1.31	324168	45	54	250	250	40	5	40		
5	30		1.5	1.09	326, 14	45	54	251	251	40	- J	40		
S	40	328.18	7;5	1.09	326, Sh	445	5.5	251	251	40	4	40		
b	44		0.90	0.65	330,60	44	55		244	40	3	40		
6	43	1140	0.90	0.65	331,900	, 441	5.5	248	8 45	, 40	3	40		
			Avg Delta P	Avg Delta H	Total Volume	Avg Ts	Δy₩ [™] J	Min/Max	Min/Max	Max	Max Vac	Min/Max		
<u>\x\x\</u> ;			/, 59583 Avg Sgrt Delta P/	1.14363	53,660	43.5	1 7 (248	248/23	9 66		40 66		ľ

Avg Sqrt Delta P/ Avg Sqrt Del H Comments:

43.58

49.08 250

EPA Method 0010 from EPA SW-846

ISOKINET	TIC FIELD	DATA SHI	EET		EPA M	[ethod	0010 - H	HFPO I	oimer A	cid		Pageo	,
Client	Chemours		Stack Condi	tions	Meter Box ID			12			14.5		
W.O.#		<u> </u>	Assur	med Actual	Meter Box Y			0691	-		K Factor	0. ta	Dl
Project ID	CHEMOURS	% Moisture	#25	9	Meter Box Del	Н	7.8	8/2_	Leak Chec	ks	Initial	Mid-Poir	nt Final
Mode/Source ID		Impinger Vol (r	ml)	9.0	_ Probe ID / Len	gth	P695	5	Sample Train	n (ft ³)	1001	0.00	100.001
Samp. Loc. ID	STK	Silica gel (g)		12.1	Probe Material			oro	Leak Check	@ (in Hg)	015	17	e 7
Run No.ID	2	CO2, % by Vol		√	Pitot / Thermod	couple ID	P695	35.74	Pitot good		ve s∕ no	y(e)s / no	yes / no
Test Method ID	M0010	O2, % by Vol	20,2	7 · Y	Pitot Coefficier	nt		.84	Orsat good		ves / no	y Gaz / no	
Date ID	16JAN2019	Temperature (Nozzle ID		<u> </u>		Temp Che			est Set	Post-Test Set
Source/Location _	Division Stack	Meter Temp (°			Avg Nozzle Dia	, ,		1601	Meter Box To	•	<i>5</i> 0		51
Sample Date	1/16/19	Static Press (in	1 H ₂ O) _ - (5 , 7	01-0,7	Area of Stack	(ft²)		<u>37 V</u>	Reference T	<u>-</u>	<u> </u>		-51
Baro. Press (in Hg)	30.06	V	ND.	45555	Sample Time			76 J	Pass/Fail (+/			/ Fail	Pass / Fail
	WINKELIER	Ambient Temp	(°F) ••>>	42 70	_Total Traverse	Pts	1	<u>∠ √</u>	Temp Chang	ge Response '	X	/ no	∫es / no
13 / 2 2 2 2 1	2_	1,5 1,5 1,7 1,7	1.08 1.08 1.22 1.22	332,725 334,85 338,80 341,23	47 47 47 43 43	Wρ	56 55 55 56 56	259 259 259 258 258	PLYTER BOX TEMP	MPRIVO EXIL TEMP (15)	SAMPLE TRAIN VAC (R.Hg)	# 5-	COMMENTS
	А	1.8	1,29	346.44	44		56	255	521	7.	5	41	
	4	1: 2	1,29	348,30	45	<u> </u>		255		71		4/1	
	12	1.8	1.36		43		56		254	41	کِ		
	6		1.08	350,70			57	254	255	41	ج_	47	
	3	1,5		325.92	45		60	254	254	43	4	43	
		1, 5	1.08	355,10	45		60	284	254	4)3	4	43	
	9	1,2	0.864	357,15	45		60	254	253	43	3	43	
6 "	13 1400	1,2	0.864	+	45		60	254	253	4/3	്ച	43	
	1425			359,600					1				
	1 '	1, 4	60,1	361,25	45		60	254	233	43	3	43	26,4110
	3	(,)	1.00	364,11	45		60	254	253	L13	3	43	
	2	1.7	1,22	366,20	45		61	254	253	45	4)	45	··· t
2 ()		1,7	1,22	368.73	45		6	254	253	46	5	46	
3 2		1.9	7,41	371.01	47		61	255	255	46	5=	46€	0.145
3 2	1	1. a	1,41	373.58	47		61	255	255	46	-3-	46	K-19actors
4 2	i l	1.7	1.276	376.0	47		6	255	255	47	<u> </u>	4/7	7.1901FR
4 3.	2	1.7	1.26	378,21	47		62	234	253	47-	4	47-	
7VV-71-36T	v1.3	िंप	1:04	3 80.52	41	<u> </u>	62	254		47			
3 W.S. C.	<u> </u>	1,4	1,04	382.86	41				253		4	47	
6	44	1.0		384,64			102	255	255	47		47	
	2 8-15-20 W	1.0	0.748		48	-	62	255	2.5-1	47	3	47	
<u>v</u>	י טכלים סו	Avg Delta P	0.745) Painin	48/		62,	255	234	4.8	3	48	
	1513	Avg sun Delta P	Avg Delta H ルスフ名3	Total Volume	Avg Ts V	AVQ	Tm 7	Min/Max	Min/Max	Max Temp	Max Vac	Max Temp	4.
	l	" >2111	Avg Sqrt Del H	23118	17375	<u> </u>	9.2"	254/259	55) (25)	51		51	
	TOP-	1.24191	1.02300	Comments: •						EPA Method	0010 from EF	'A SW-846	į.

ISOKINE	ETIC FIELD	DATA SH	EET		EPA N	1ethod	0010 - 1	HFPO I	Dimer A	cid		Pageof	./
Client	Chemours		Stack Condit	tions	Meter Box ID		4	12			<u> </u>		$\overline{}$
W.O.#			Assur	med Actual	Meter Box Y		1:8	0691	_		K Factor	0,721	
Přoject ID	CHEMOURS	% Moisture	~1.9		Meter Box De	el H		812	Leak Ched	:ks	Initial	Mid-Poir	l nt Final
Mode/Source ID		Impinger Vol (ml)	9	Probe ID / Le		P695	1/5/	Sample Trai		0,001	0.00	
Samp. Loc. ID	STK	Silica gel (g)		126	Probe Materia	al		3oro	Leak Check		C)5	<u> </u>	<u>(</u> 0,001
Run No.ID	3	CO2, % by Vo	0.1	1	- Pitot / Thermo	ocouple ID	P695	695	Pitot good	G (g)	(yes / no	(ye) / no	
Test Method ID	M0010	O2, % by Vol	20.5	7. /	- Pitot Coefficie			0.84	Orsat good		yes / no	yes / no	
Date ID	16JAN2019	Temperature ((°F) タリし)	Nozzle ID			60.	Temp Che	ck	Pre-T	est Set	Post-Test Set
Source/Location	Division Stack	Meter Temp (F) 2 40		Avg Nozzle D	Dia (in)	0,	160 1	Meter Box T		36		48
Sample Date	1/17/19	Static Press (i	n H ₂ O) -0.7	01-0.7	Area of Stack	(ft²)	7.		Reference T			·	49
Baro. Press (in Hg					Sample Time	` '	a (<i>a</i>	– Pass/Fail (+			/ Fail	Pass / Fail
Operator M2	U WINKELE	2 √ Ambient Temp	o (°F) 🥕 💆	36	Total Travers		1	21		ge Response		/ no	yes / no
505 t	1	Laksania in a sakil	- 3.57%×9405 a.:	At the Control of the		ti sana		·		Je reoponse Take	<u> </u>	Ti and	
100 T	0 O842	Files pr			30 X	per Maria		elegedes .	And the Andrews		To the second	# 60 # 1 # V 1 # (30 0) # 1 # 1	san ting
	0892		A DO	387.510	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1			r series	- 1 m	100	70.00	1	** X 2.
- 12 		1,5	1,08	389,61	34	NA	33	255	249	60	1	160	
<u> </u>	ર	1,5	1,08	391.32	3-1	<u> </u>	34	755	251	63	4	63	
	12	107	1,22	393,52	35	1{	36	255-	258	63	5	63	
2	16	1,7	1,22	1396,51	35		36	255	258	63	5-	63	
ا م	20	1,8	1,29	398,30	35		37	259	258	63	5	63	26,48
3	2-1	1.2	1.29	400.86	300	1	3.5	25 9	258	63	5	63	20, 18
4	28	1.3	124	403.28	36	 	33	239	258	63	3		
4	32	1,3	1,29	405.28	377	 	3.5				5	63	
3	36	1.2	1.00	439.23	36	 		258	258	63		63	
Ś	40	1, 4				 	38	255	255	63	4	63	
	44	+ 1, 5	1,00	410,00	36		3.8	254	253	15	4	四	
<u> </u>		1.5	0,937	411.92	36		39	254	253	62	4	62	
<u> </u>	48 0930	1.3	0.937	413.990	37		39	252	523	62	3	62	
	0947			414,135	1								
A	4	1,6	1.02	416,27	79		39	255	255-	65-	Ч	05	
1	3	1,5	1,08	418,52	39		34	750-	255-	65-	-1	65-	
2	12	11.7	1,22	420.70	37		44	255	255-	63	\$	63	26.666
2.	16	1,7	1.72	422,91	37		पियो	255	25 =				20,000
7	20	1.9	127	425.26	77	 				63	چ	63	
	29	11 a	<u> </u>		3-	 	45	255	255	<u>C3</u>	~	63	
	28	1 / a	1,36	427.66			45	225	254	63	٤	63	
- 4 -			<u> 7,36</u>	430,07	37		46	255	258	62	5	62	
	32	1,9	1.36	432 51	3.5		46	258	258	62	3	62	
-100 -100 -100 -100 -100 -100 -100 -100	3 <u>0</u>	1,5	1,00	434,35	39	L. 1	47	825	528	61	5	61	
VVV-971/6	<u>N40</u>	1.5	1.08	426.70	40		143	258	288	62	2	62	
WASING	44	1.3	0.937	438,77	40	. /	42	258	258	63	-2	63	
* (2)	48 1035	7,3	0,937	440.801	40		42.	257	257	64	5	64	
	1,60233 V	Avg Sqrt Delta P	Avg Delta H		3°€.5°9	94	2.91	Min/Max.		Max Temp	Max Vac	Max Temp	 _
a hold			Avg Sqrt Del H	Comments	37.0			154 F	2MIL		0010 from EP	A C)A/ C40	
VO LAG		ļ	Avg Sqrt Del H	2	٧		•	•		LFA MEU100	OUTO ITOM EP	M 3VV-040	
r1		L	,,,,,	Ţ									
				7									

SAMPLE RECOVERY FIELD DATA

EPA Method 0010 - HFPO Dimer Acid

Client		Chem	ours	_	W.O. #					_
Location/Pla	ınt _	Fayette	rille, NC	Source	e & Location		Division	Stack		-
Run No.	1				Sample Date			Recovery Date		C/16/19
Sample I.D.	Chemours - I	Division - STK -	1 - M0010 -	•	Analyst	pour le	W	Fifter N	umber	<u> </u>
					Impinge					
<u></u>	1	2	3	4	5	6	7	Imp.Total	8	Total
Contents	Empty	HPLC H20	HPLC H20						Silica Gel	
Final	2	101	102	5				210	7115	
Initial	0	100	100	Q				200	300	
Gain	2	1	1	5				10	1h5	28.5
Impinger Cold	or	cloar			Labeled?		1	✓	V	
Silica Gel Co	_	ලංගේ	_		Sealed?		(- -
					·	1 <i>f</i>				1.1.
Run No.	_2_				Sample Date	1 /16/12	ļ	Recove	ry Date	1/16/18 NA
Sample I.D.	Chemours - [Division - STK -	2 - M0010 -		Analyst	i holu dun	(cu	Filter N	umber	<u>MA</u>
				•	Impinge	er (· · · · · · · · · · · · · · · · · · ·			
	1	2	3	4	5	6	7	Imp.Total	8	Total
Contents	Empty	HPLC H20	HPLC H20						Silica Gel	
Final	16	9-3	96	4					3171	
Initial	0	100	100	Ð				200	300	aw
Gain	16	~7	- ਪ	৭				29°	124	171
Impinger Col	or	سعيل	L		Labeled?			an 1	✓	2107
Silica Gel Co	ndition	Cean			Sealed?					-
						cha lse		_		, I la
Run No.					Sample Date	clurka ruul	P	Recove	y Date	1/17/18
Sample I.D.	Chemours - [Division - STK -	3 - M0010 -		Analyst	Pull	200	Filter N	umber	NA
					Impinge					
	11	2	3	4	5	6	7	Imp.Total	8	Total
Contents	Empty	HPLC H20	HPLC H20		-	, ,		ـ مـــــــــــــــــــــــــــــــــــ	Silica Gel	
Final	2	103	ior	~				24	312.6	
Initial	0	100	100	i)				200	300	
Gain	2	3	て	2				9	12.6	746
Impinger Cole	or	Cheer			Labeled?			/ /	V	_
Silica Gel Co	ndition —	Good			Sealed?					

Check COC for Sample IDs of Media Blanks



METHODS AND ANALYZERS

Client: Chemours Project Number: 15418.002.009

Operator: CW

Source: Division Stack Date: 16 Jan 2019

ent Folders.A-F\Chemours Fayetteville\15418.002.009 Fayetteville Jan 2019 Carbon Bed Test\Data\Division\011619

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 O₂

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 CO₂

Method **EPA 3A**, Using Bias Analyzer Make, Model & Serial No. **Servomex 4900**

Full-Scale Output, mv
Analyzer Range, %
Span Concentration, %
10000
20.0
16.6



CALIBRATION DATA

Number 1

Client: Chemours

Location: CHEMOURS
Source: Division Stack

Project Number: **15418.002.009**

Operator: CW

Date: 16 Jan 2019

Start Time: 07:51

 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 5 mv **Span, 21.0 %** 7991 mv

Curve Coefficients

Slope Intercept 380.3 5

 CO_2

Method: EPA 3A

Calibration Type: Linear Zero and High Span

Calibration Standards

Cylinder ID8.9 CC18055
16.6 SG9169108

Calibration Results

Zero 5 mv **Span, 16.6 %** 8383 mv

Curve Coefficients

Slope Intercept 505.3 5



CALIBRATION ERROR DATA

Number 1

Client: Chemours

Project Number: 15418.002.009

Operator: **CW**

Location: CHEMOURS

Source: **Division Stack**

Calibration 1 Date: 16 Jan 2019

Start Time: 07:51

 O_2

Method: EPA 3A Span Conc. 21.0 %

Slope 380.3

Intercept 5.0

Standard	Result	Difference	Error	
%	%	%	%	Status
Zero	0.0	0.0	0.0	Pass
12.0	12.0	0.0	0.0	Pass
21.0	21.0	0.0	0.0	Pass

 CO_2

Method: EPA 3A Span Conc. 16.6 %

Slope 505.3

Intercept 5.0

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



BIASNumber 1

Client: Chemours

Location: CHEMOURS

Source: Division Stack

Project Number: 15418.002.009

Operator: **CW**

Date: 16 Jan 2019

Calibration 1
Start Time: 07:58

 O_2

Method: EPA 3A Span Conc. 21.0 %

Bias Results									
Standard Gas	Cal. %	Bias %	Difference %	Error %	Status				
Zero	0.0	0.0	0.0	0.0	Pass				
Span	12.0	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	8.5	-0.1	-0.6	Pass					



Number 1

Client: Chemours

Location: CHEMOURS

Calibration 1 Source: Division Stack

Time	O ₂	CO ₂	
Time	%	%	
	Port A		
09:41		0.0	
09:42		0.0	
09:43		0.1	
09:44		0.1	
09:45		0.1	
09:46		0.1	
09:47		0.1	
09:48		0.1	
09:49		0.1	
09:50		0.1	
09:51		0.1	
09:52		0.1	
09:53		0.1	
09:54		0.1	
09:55		0.1	
09:56		0.1	
09:57		0.1	
09:58		0.1	
09:59		0.1	
10:00		0.1	
10:01		0.1	
10:02		0.1	
10:03		0.1	
10:04		0.1	
10:05		0.1	
10:06		0.1	
10:07		0.1	
10:08		0.1	
10:09		0.1	
10:10	21.0	0.1	
10:11		0.1	
10:12		0.1	
10:13	21.0	0.1	
10:14		0.1	
10:15		0.1	
10:16		0.1	
10:17		0.1	
10:18		0.1	
10:19		0.1	



Number 1

Calibration 1

Client: Chemours

Location: CHEMOURS

Source: Division Stack

Time	O ₂ %	CO ₂ %	
10:20	21.0	0.1	
10:21	21.0	0.1	
10:22	21.0	0.1	
10:23	21.0	0.1	
10:24	21.0	0.1	
10:25	20.9	0.1	
10:26	21.0	0.1	
10:27	21.0	0.1	
10:28	21.0	0.1	
10:29	21.0	0.1	
	Port B		
10:52	20.9	0.0	
10:53	20.9	0.0	
10:54	20.9	0.1	
10:55	20.9	0.1	
10:56	20.9	0.1	
10:57	20.9	0.1	
10:58	20.9	0.1	
10:59	20.9	0.1	
11:00	20.9	0.1	
11:01	20.9	0.1	
11:02	20.9	0.1	
11:03	20.9	0.1	
11:04	20.9	0.1	
11:05	20.9	0.1	
11:06	20.9	0.1	
11:07	20.9	0.1	
11:08	20.9	0.1	
11:09	21.0	0.1	
11:10	21.0	0.1	
11:11	21.0	0.1	
11:12	21.0	0.1	
11:13	21.0	0.1	
11:14	21.0	0.1	
11:15	21.0	0.1	
11:16	21.0	0.1	
11:17	21.0	0.1	
11:18	21.0	0.1	
11:19	21.0	0.1	
11:20	21.0	0.1	



Number 1

Client: Chemours

Location: CHEMOURS Source: Division Stack

Calibration 1

Time	O ₂ %	CO ₂ %	
11:21	21.0	0.1	
11:22	21.0	0.1	
11:23	21.0	0.1	
11:24	21.0	0.1	
11:25	21.0	0.1	
11:26	21.0	0.1	
11:27	21.0	0.1	
11:28	21.0	0.1	
11:29	21.0	0.1	
11:30	21.0	0.1	
11:31	21.0	0.1	
11:32	21.0	0.1	
11:33	21.0	0.1	
11:34	21.0	0.1	
11:35	21.0	0.1	
11:36	21.0	0.1	
11:37	21.0	0.1	
11:38	21.0	0.1	
11:39	21.0	0.1	
11:40	21.0	0.1	
	End Run 1		
Avgs	21.0	0.1	



RUN SUMMARY

Number 1

Client: **Chemours**

Location: CHEMOURS

Source: **Division Stack**

Project Number: 15418.002.009

Operator: **CW**

Date: **16 Jan 2019**

Calibration 1

O₂ EPA 3A

%

Method

Conc. Units

A

CO₂ EPA 3A %

Time: 09:40 to 11:40

Run Averages

21.0 0.1

Pre-run Bias at 07:58

 Zero Bias
 0.0
 0.0

 Span Bias
 12.0
 8.5

 Span Gas
 12.0
 8.9

Post-run Bias at 12:04

 Zero Bias
 0.0
 0.1

 Span Bias
 12.0
 8.5

 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: **Division Stack**

Project Number: 15418.002.009

Operator: CW

Date: 16 Jan 2019

Calibration 1
Start Time: 12:04

 O_2

Method: EPA 3A Span Conc. 21.0 %

		Bias	Results		
Standard Gas	Cal. %	Bias %	Difference %	Error %	Status
Zero	0.0	0.0	0.0	0.0	Pass
Span	12.0	12.0	0.0	0.0	Pass
Standard	Initial*	Final	ation Drift Difference	Drift	
Gas	%	% 0.0	%	%	Status
Zero Span	0.0 12.0 *Bias No. 1	0.0 12.0	0.0 0.0	0.0 0.0	Pass 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	8.5	-0.1	-0.6	Pass
		Calibra	ation Drift		
Standard	Initial*	Final	Difference	Drift	
Gas	%	%	%	%	Status
Zero	0.0	0.1	0.1	0.6	Pass
Span	8.5	8.5	0.0	0.0	Pass
•	*Bias No. 1				



Number 2

Client: Chemours

Location: CHEMOURS Source: Division Stack

Calibration 1

Tim	е	O ₂	CO ₂	
		%	%	
		Port A		
13:1		20.8	0.0	
13:1		20.8	0.0	
13:1		20.9	0.0	
13:1		20.9	0.0	
13:1		20.9	0.1	
13:1		20.9	0.1	
13:1		20.9	0.1	
13:1		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:2		20.9	0.1	
13:3		20.9	0.1	
13:3		20.9	0.1	
13:3	32	20.9	0.1	
13:3	33	20.9	0.1	
13:3		20.9	0.1	
13:3	35	20.9	0.1	
13:3	36	20.9	0.1	
13:3		20.9	0.1	
13:3	38	20.9	0.1	
13:3	39	20.9	0.1	
13:4		20.9	0.1	
13:4	11	20.9	0.1	
13:4	12	20.9	0.1	
13:4	13	20.9	0.1	
13:4	14	20.9	0.1	
13:4	1 5	20.9	0.1	
13:4	16	20.9	0.1	
13:4	17	20.9	0.1	
13:4	18	20.9	0.1	
13:4	19	20.9	0.1	
13:5	50	20.9	0.1	



Number 2

Client: Chemours

Location: CHEMOURS Source: Division Stack

Calibration 1

Jource. Division Stack		alibration	1	Date. 16 Jan 2019
	Time	O ₂ %	CO ₂ %	
	13:51	20.9	0.1	
	13:52	20.9	0.1	
	13:53	20.9	0.1	
	13:54	20.9	0.1	
	13:55	20.9	0.1	
	13:56	20.9	0.1	
	13:57	20.9	0.1	
	13:58	20.9	0.1	
	13:59	20.9	0.1	
	14:00	20.9	0.1	
		Port B		
	14:25	20.8	0.0	
	14:26	20.8	0.0	
	14:27	20.9	0.1	
	14:28	20.9	0.1	
	14:29	20.9	0.1	
	14:30	20.9	0.1	
	14:31	20.9	0.1	
	14:32	20.9	0.1	
	14:33	20.9	0.1	
	14:34	20.9	0.1	
	14:35	20.9	0.1	
	14:36	20.9	0.1	
	14:37	20.9	0.1	
	14:38	20.9	0.1	
	14:39	20.9	0.1	
	14:40	20.9	0.1	
	14:41	20.9	0.1	
	14:42	20.9	0.1	
	14:43	20.9	0.1	
	14:44	20.9	0.1	
	14:45	20.9	0.1	
	14:46	20.9	0.1	
	14:47	20.9	0.1	
	14:48	20.9	0.1	
	14:49	20.9	0.1	
	14:50	20.9	0.1	
	14:51	20.9	0.1	
	14:52	20.9	0.1	
	14:53	20.9	0.1	



Number 2

Client: Chemours

Location: CHEMOURS Source: Division Stack

Project Number: **15418.002.009**Operator: **CW**Date: **16 Jan 2019** Calibration 1

14:54 20.9 0.1	
14:55 20.9 0.1 14:56 20.9 0.1	
14:57 20.9 0.1	
14:58 20.9 0.1	
14:59 20.9 0.1	
15:00 20.9 0.1	
15:01 20.9 0.1	
15:02 20.9 0.1	
15:03 20.9 0.1	
15:04 20.9 0.1	
15:05 20.9 0.1	
15:06 20.9 0.1	
15:07 20.9 0.1	
15:08 20.9 0.1	
15:09 20.9 0.1	
15:10 20.9 0.1	
15:11 20.9 0.1	
15:12 20.9 0.1	
15:13 20.9 0.1	
End Run 2	
Avgs 20.9 0.1	



RUN SUMMARY

Number 2

Client: Chemours

Location: CHEMOURS

Source: Division Stack

Project Number: **15418.002.009**

Operator: CW

Date: 16 Jan 2019

Calibration 1

O₂ EPA 3A

Method

Conc. Units

CO₂ EPA 3A %

Time: 13:11 to 15:13

%

Run Averages

20.9 0.1

Pre-run Bias at 12:04

 Zero Bias
 0.0
 0.1

 Span Bias
 12.0
 8.5

 Span Gas
 12.0
 8.9

Post-run Bias at 15:16

 Zero Bias
 0.0
 0.0

 Span Bias
 12.0
 8.5

 Span Gas
 12.0
 8.9

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

20.9 0.0



BIAS AND CALIBRATION DRIFT

Number 3

Client: Chemours

Location: CHEMOURS

Source: **Division Stack**

Project Number: 15418.002.009

Operator: **CW**

Date: 16 Jan 2019

Start Time: 15:16

Calibration 1

 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	12.0	12.0	0.0	0.0	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. 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	8.5	-0.1	-0.6	Pass		
	Calibration Drift						
Standard	Initial*	Final	Difference	Drift			
Gas	%	%	%	%	Status		
Zero	0.1	0.0	-0.1	-0.6	Pass		
Span	8.5	8.5	0.0	0.0	Pass		
•	*Bias No. 2						



METHODS AND ANALYZERS

Client: Chemours Project Number: 15418.002.009

Operator: CW

Source: Division Stack Date: 17 Jan 2019

ent Folders.A-F\Chemours Fayetteville\15418.002.009 Fayetteville Jan 2019 Carbon Bed Test\Data\Division\011719

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 O₂

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 CO₂

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

Span Concentration, % 16.6



CALIBRATION DATA

Number 1

Client: Chemours

Location: CHEMOURS
Source: Division Stack

Project Number: 15418.002.009

Operator: CW

Date: 17 Jan 2019

Start Time: 07:32

 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 8 mv **Span, 21.0 %** 8020 mv

Curve Coefficients

Slope Intercept 8

 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 1 mv **Span, 16.6 %** 8293 mv

Curve Coefficients

Slope Intercept 500.1 1



CALIBRATION ERROR DATA

Number 1

Client: Chemours

Project Number: 15418.002.009

Location: CHEMOURS

Operator: **CW**

Source: Division Stack

Date: 17 Jan 2019

Calibration 1

Start Time: 07:32

 O_2

Method: EPA 3A Span Conc. 21.0 %

Slope 381.5

Intercept 8.0

Standard %	Result %	Difference %	Error %	Status
Zero	0.0	0.0	0.0	Pass
12.0	12.0	0.0	0.0	Pass
21.0	21.0	0.0	0.0	Pass

 CO_2

Method: EPA 3A Span Conc. 16.6 %

Slope 500.1

Intercept 1.0

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



BIASNumber 1

Client: Chemours

Location: CHEMOURS

Source: **Division Stack**

Project Number: 15418.002.009

Operator: **CW**

Date: 17 Jan 2019

Calibration 1

Start Time: 07:36

 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	12.0	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	8.5	-0.1	-0.6	Pass				



Number 3

Client: Chemours

Location: CHEMOURS Source: Division Stack

Project Number: **15418.002.009**Operator: **CW**Date: **17 Jan 2019** Calibration 1

Time	O ₂ %	CO ₂ %	
	Port A		
08:42	20.9	0.0	
08:43	20.9	0.0	
08:44	20.9	0.0	
08:45	20.9	0.0	
08:46	20.9	0.1	
08:47	20.9	0.1	
08:48	20.9	0.1	
08:49	20.9	0.1	
08:50	20.9	0.1	
08:51	20.9	0.1	
08:52	20.9	0.1	
08:53	20.9	0.1	
08:54	20.9	0.1	
08:55	21.0	0.1	
08:56	20.9	0.1	
08:57	21.0	0.1	
08:58	21.0	0.1	
08:59	21.0	0.1	
09:00	21.0	0.1	
09:01	21.0	0.1	
09:02	20.9	0.1	
09:03	20.9	0.1	
09:04	21.0	0.1	
09:05	20.9	0.1	
09:06	20.9	0.1	
09:07	21.0	0.1	
09:08	20.9	0.1	
09:09	21.0	0.1	
09:10	21.0	0.1	
09:11	21.0	0.1	
09:12	21.0	0.1	
09:13	21.0	0.1	
09:14	21.0	0.1	
09:15	21.0	0.1	
09:16	21.0	0.1	
09:17	21.0	0.1	
09:18	21.0	0.1	
09:19	21.0	0.1	
09:20	21.0	0.1	



Number 3

Calibration 1

Client: Chemours

Location: CHEMOURS

Source: Division Stack

 		•	
Time	O ₂ %	CO ₂ %	
09:21	21.0	0.1	
09:22	21.0	0.1	
09:23	21.0	0.1	
09:24	21.0	0.1	
09:25	21.0	0.1	
09:26	21.0	0.1	
09:27	21.0	0.1	
09:28	21.0	0.1	
09:29	21.0	0.1	
09:30	21.0	0.1	
	Port B		
09:47	20.8	0.0	
09:48	20.8	0.0	
09:49	20.8	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	
09:56	20.9	0.1	
09:57	20.9	0.1	
09:58	21.0	0.1	
09:59	21.0	0.1	
10:00	20.9	0.1	
10:01	21.0	0.1	
10:02	21.0	0.1	
10:03	21.0	0.1	
10:04	21.0	0.1	
10:05	21.0	0.1	
10:06	21.0	0.1	
10:07	21.0	0.1	
10:08	21.0	0.1	
10:09	21.0	0.1	
10:10	21.0	0.1	
10:11	21.0	0.1	
10:12	21.0	0.1	
10:13	21.0	0.1	
10:14	21.0	0.1	
10:15	21.0	0.1	



Number 3

Client: Chemours

Location: CHEMOURS Source: Division Stack

Calibration 1

Time	O ₂ %	CO ₂ %	
10:16	21.0	0.1	
10:17	21.0	0.1	
10:17	21.0	0.1	
10:19	21.0	0.1	
10:20	21.0	0.1	
10:21	21.0	0.1	
10:22	21.0	0.1	
10:23	21.0	0.1	
10:24	21.0	0.1	
10:25	21.0	0.1	
10:26	21.0	0.1	
10:27	21.0	0.1	
10:28	21.0	0.1	
10:29	21.0	0.1	
10:30	21.0	0.1	
10:31	21.0	0.1	
10:32	21.0	0.1	
10:33	21.0	0.1	
10:34	21.0	0.1	
10:35	21.0	0.1	
	End Run 3	0.1	
Avgs	21.0	0.1	



RUN SUMMARY

Number 3

Client: **Chemours**

Location: CHEMOURS

Source: Division Stack

Project Number: 15418.002.009

Operator: CW

Date: 17 Jan 2019

Calibration 1

 O_2

EPA 3A

%

Method

Conc. Units

CO₂ EPA 3A %

Time: 08:41 to 10:35

Run Averages

21.0 0.1

Pre-run Bias at 07:36

 Zero Bias
 0.0
 0.0

 Span Bias
 12.0
 8.5

 Span Gas
 12.0
 8.9

Post-run Bias at 10:43

 Zero Bias
 0.0
 0.1

 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 0.0



BIAS AND CALIBRATION DRIFT

Number 2

Client: Chemours

Location: CHEMOURS

Source: **Division Stack**

Project Number: 15418.002.009

Operator: **CW**

Date: 17 Jan 2019

Start Time: 10:43

Calibration 1

 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	12.0	12.0	0.0	0.0	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_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
		Calibra	ation Drift		
Standard	Initial*	Final	Difference	Drift	
Gas	%	%	%	%	Status
Zero	0.0	0.1	0.1	0.6	Pass
Span	8.5	8.4	-0.1	-0.6	Pass
•	*Bias No. 1				



APPENDIX C LABORATORY ANALYTICAL REPORT

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

Client Sample Results

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

TestAmerica Job ID: 140-14017-1

Client Sample ID: Q-1877,1878 DIV STACK R1 M0010 FH Lab Sample ID: 140-14017-1

Date Collected: 01/16/19 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** 0.126 0.0136 ug/Sample 01/23/19 14:18 02/01/19 10:50 12.0 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac

Client Sample ID: Q-1879,1880,1882 DIV STACK R1 M0010 BH

92

Lab Sample ID: 140-14017-2 Date Collected: 01/16/19 00:00 Matrix: Air

50 - 200

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

Method: 8321A - PFOA and PFOS

Analyte RL**MDL** Unit D Result Qualifier Prepared Analyzed Dil Fac HFPO-DA 54.1 0.500 0.100 ug/Sample 01/22/19 10:37 01/30/19 11:49 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 75 D 50 - 200 01/22/19 10:37 01/30/19 11:49

Lab Sample ID: 140-14017-3 Client Sample ID: Q-1881 DIV STACK R1 M0010 IMP 1,2&3

CONDENSATE

13C3 HFPO-DA

Date Collected: 01/16/19 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 0.210 01/30/19 04:42 02/04/19 11:11 **HFPO-DA** 0.0964 J H 0.0107 ug/Sample Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 88 50 - 200 01/30/19 04:42 02/04/19 11:11

Client Sample ID: Q-1883 DIV STACK R1 M0010 Lab Sample ID: 140-14017-4

BREAKTHROUGH XAD-2 RESIN TUBE

Date Collected: 01/16/19 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 01/22/19 10:37 01/30/19 11:53 **HFPO-DA** 0.150 J 0.0400 ug/Sample Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 01/22/19 10:37 01/30/19 11:53 13C3 HFPO-DA 77 50 - 200

01/23/19 14:18 02/01/19 10:50

Client Sample Results

Project/Site: Division Stack - M0010

Client: Chemours Company FC, LLC The TestAmerica Job ID: 140-14017-1

Client Sample ID: Q-1884,1885 DIV STACK R2 M0010 FH

Lab Sample ID: 140-14017-5 Matrix: Air

Date Collected: 01/16/19 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** 9.22 0.102 0.0110 ug/Sample 01/23/19 14:18 02/01/19 10:53

Surrogate %Recovery Qualifier I imits Prepared Analyzed Dil Fac 13C3 HFPO-DA 96 50 - 200 01/23/19 14:18 02/01/19 10:53

Client Sample ID: Q-1886,1887,1889 DIV STACK R2 M0010 BH

Lab Sample ID: 140-14017-6 Date Collected: 01/16/19 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 0.900 HFPO-DA 0.180 ug/Sample 01/22/19 10:37 01/30/19 11:56 59.6 Prepared Analyzed Dil Fac

Surrogate %Recovery Qualifier Limits 13C3 HFPO-DA 82 D 50 - 200

Lab Sample ID: 140-14017-7 Client Sample ID: Q-1888 DIV STACK R2 M0010 IMP 1,2&3

CONDENSATE

Date Collected: 01/16/19 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 Dil Fac ח Prepared Analyzed HFPO-DA 0.218 $\overline{\mathsf{ND}}$ $\overline{\mathsf{H}}$ 0.0111 ug/Sample 01/30/19 04:42 02/04/19 11:15

Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 01/30/19 04:42 02/04/19 11:15 87 50 - 200

Client Sample ID: Q-1890 DIV STACK R2 M0010

BREAKTHROUGH XAD-2 RESIN TUBE

Date Collected: 01/16/19 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 01/22/19 10:37 01/30/19 11:59 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac

13C3 HFPO-DA 76 50 - 200 01/22/19 10:37 01/30/19 11:59

01/22/19 10:37 01/30/19 11:56

Lab Sample ID: 140-14017-8

Client Sample Results

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

TestAmerica Job ID: 140-14017-1

Client Sample ID: Q-1891,1892 DIV STACK R3 M0010 FH

Lab Sample ID: 140-14017-9

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** 0.101 0.0109 ug/Sample 01/23/19 14:18 02/01/19 10:56 1.93

Surrogate %Recovery Qualifier I imits Prepared Analyzed Dil Fac 13C3 HFPO-DA 103 50 - 200 01/23/19 14:18 02/01/19 10:56

Client Sample ID: Q-1893,1894,1896 DIV STACK R3 M0010 BH

Lab Sample ID: 140-14017-10 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 0.800 **HFPO-DA** 0.160 ug/Sample 01/22/19 10:37 01/30/19 12:02 60.1 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 67 D 01/22/19 10:37 01/30/19 12:02 50 - 200

Lab Sample ID: 140-14017-11 Client Sample ID: Q-1895 DIV STACK R3 M0010 IMP 1,2&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 Dil Fac ח Prepared Analyzed 0.222 01/30/19 04:42 02/04/19 11:18 **HFPO-DA** 0.115 J 0.0113 ug/Sample Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 50 - 200 01/30/19 04:42 02/04/19 11:18 92

Client Sample ID: Q-1897 DIV STACK R3 M0010 Lab Sample ID: 140-14017-12

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** 0.200 0.0400 ug/Sample 01/22/19 10:37 01/30/19 12:06 0.306 Surrogate %Recovery Qualifier Limits Prepared Analyzed Dil Fac 13C3 HFPO-DA 85 50 - 200 01/22/19 10:37 01/30/19 12:06

APPENDIX D SAMPLE CALCULATIONS

SAMPLE CALCULATIONS FOR HFPO DIMER ACID (METHOD 0010)

Client: Chemours
Test Number: Run 3
Test Location: Divison Stack

Plant: Fayetteville, NC Test Date: 1/17/19 Test Period: 0842-1035

1. HFPO Dimer Acid concentration, lbs/dscf.

$$Conc1 = 2.41E-09$$

Where:

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

Conc1 = Division Stack HFPO Dimer Acid concentration, lbs/dscf.

 2.2046×10^{-9} = Conversion factor from ug to lbs.

${\bf 2.\ HFPO\ Dimer\ Acid\ concentration,\ ug/dscm.}$

Conc2 = $W / (Vm(std) \times 0.02832)$

Conc2 = $62.5 / (57.033 \times 0.02832)$

Conc2 = 3.87E+01

Where:

Conc2 = Division Stack HFPO Dimer Acid concentration, ug/dscm.

0.02832 = Conversion factor from cubic feet to cubic meters.

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

 $MR1_{(Outlet)}$ = Conc1 x Qs(std) x 60 min/hr

 $MR1_{(Outlet)} = 2.41E-09 \times 30735 \times 60$

 $MR1_{(Outlet)} = 4.45E-03$

Where:

MR1_(Outlet) = Division Stack HFPO Dimer Acid mass emission rate, lbs/hr.

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

 $MR2_{(Outlet)} = PMR1 \times 453.59 / 3600$

 $MR2_{(Outlet)} = 4.45E-03 \times 453.59 / 3600$

 $MR2_{(Outlet)} \ = \ 5.60E\text{-}04$

Where:

MR2_(Outlet) = Division Stack HFPO Dimer Acid mass emission rate, g/sec.

453.6 = Conversion factor from pounds to grams.

3600 = Conversion factor from hours to seconds.

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EXAMPLE CALCULATIONS FOR VOLUMETRIC FLOW AND MOISTURE AND ISOKINETICS

 Client: Chemours
 Facility: Fayetteville, NC

 Test Number: Run 3
 Test Date: 1/17/19

 Test Location: Division Stack
 Test Period: 842-1035

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) =	$(0.04707 \times 9.0) + (0.04715 \times 12.6) = 1.02$
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 ³)/lb-mole)(deg R); absolute
	temperature at standard conditions (528 deg R), absolute
	pressure at standard conditions (29.92 in. Hg), ft ³ /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 ³)/lb-mole)(deg R); absolute
	temperature at standard conditions (528 deg R), absolute
	pressure at standard conditions (29.92 in. Hg), and
	$453.6 \text{ g/lb, ft}^3/\text{g.}$

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3. Moisture content

$$bws = \begin{array}{c} Vw(std) \\ \hline Vw(std) + Vm(std) \\ \\ bws = \\ \hline 1.02 \\ \hline 1.02 + 57.033 \\ \\ \end{array} = 0.018$$

Where:

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

4. Mole fraction of dry gas.

$$Md = 1 - bws$$

$$Md = 1 - 0.018 = 0.982$$

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.

% N₂ = Percent nitrogen by volume, dry basis.

% CO = Percent carbon monoxide by volume, dry basis. 0.440 = Molecular weight of carbon dioxide, divided by 1

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

0.280 = Molecular weight of oxygen, divided by 100.

Molecular weight of nitrogen or carbon monoxide,

divided by 100.

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

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

$$MWs = (28.84 \times 0.982) + (18(1 - 0.982)) = 28.65$$

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 =
$$85.49 \times 0.84 \times 1.26546 \times (------)^1/2 = 69.0$$

30.13 x 28.65

Where:

Vs = Average gas stream velocity, ft/sec.

(lb/lb-mole)(in. Hg)^{1/2}

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

(deg R)(in H₂O)

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. H₂O.

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

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

$$Qs(act) = 60 \times 69.0 \times 7.07 = 29252$$

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.

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

$$Qs(std) = 30735$$

Where:

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

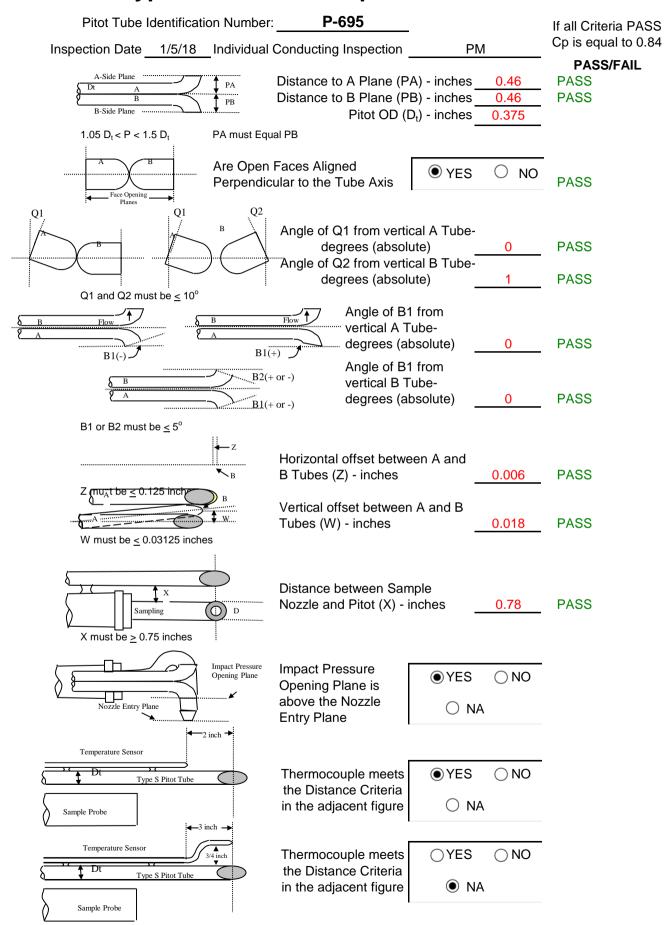
${\bf 10.}\ Is okinetic\ variation\ calculated\ from\ intermediate\ values,\ percent.$

17.327 x Ts x Vm(std) I = Vs x O x Ps x Md x (Dn)² 17.327 x 497 x 57.033 I = 69.0 x 96 x 30.13 x 0.982 x (0.160)^2 Where: I = Percent of isokinetic sampling. O = 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)(ft²)(sec)

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APPENDIX E EQUIPMENT CALIBRATION RECORDS

Type S Pitot Tube Inspection Data Form



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		Temperatures					
Orifice	Wet Test	Dry gas Meter	Wet Test	Dry Gas				
Manometer	Meter	Dry guo motor	Meter	Meter		Calibration	on Results	
in H ₂ 0	ft ³	ft ³	°F	Outlet, °F	Time, min	Υ	ΔН	
(∆H)	(Vw)	(Vd)	(Tw)	(Td _o)	(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				
1.0	5.0	897.795	73.0	77.00	9.1	1.0071	1.8559	
		4.985		76.50				
		898.799		77.00				
1.5	10.0	908.810	73.0	78.00	15.20	1.0036	1.9381	
		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	Select Temperature							
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

MODIFIED METHOD 0010 TEST TRAIN

DIVISION STACK

METER BOX NO. 12

1/16/2019 + 1/17/2019

	Kun I	Run 2	Run 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 ₂ = Percent carbon dioxide by volume, dry basis.	0.0	0.0	0.0
% O ₂ = Percent oxygen by volume, dry basis.	20.9	20.9	20.9
MWd = (0.32 * O2) + (0.44 * CO2) + (0.28 * (100 - (CO2 + O2)))			
MWd = (0.32 * 20.9) + (0.44 * 0) + (0.28 * (100 - (0 + 20.9)))			
MWd = (6.69) + (0.00) + (22.15)			
MWd =	28.84	28.84	28.84
Tma = Source Temperature, absolute(°R)			
Tm = Average dry gas meter temperature, deg F.	49.1	59.2	40.9
Tma = Ts + 460			
Tma = 49.08 + 460			
Tma =	509.08	519.21	500.92
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	1.14	1.13	1.15
Pb = Barometric Pressure, in Hg.	30.15	30.06	30.18
Pm = Pb + (delta H / 13.6)			
Pm = 30.15 + (1.143625 / 13.6)			
Pm =	30.23	30.14	30.26

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.	53.660	53.775	53.146
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	1.0641	1.0581	1.0722
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 / 53.66) * SQRT (0.0319 * 509.08 * 29) / (1.88 * 30.23 * 28.84) * 1.06 Yqa = 1.789 * SQRT 470.953 / 1,639.865 * 1.06 Yqa = 1.0202 1.0238

1.0291

Diff = Absolute difference between Yqa and Y 1.32 1.68 2.20

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

Diff = ((1.0069 - 1.020) / 1.0069) * 100

Average Diff = 1.73

Allowable = 5.0

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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 Concentration	Actual Concentration	Protocol Method	Total Relative Uncertainty	Assay 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					
Туре	Lot ID	Cylinder No	Concentration		Uncertainty	Expiration Date
NTRM	13060629	CC413730	13.359 % CARBON E	DIOXIDE/NITROGEN	+/- 0.6%	May 09, 2019
			ANALYTICAL	EQUIPMENT		
Instrume	ent/Make/Mod	el	Analytical Princi	•	Last Multipoint Cali	bration
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	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	N	Balance			-	
			CALIBRATION	STANDARDS	8	
Type	Lot ID	Cylinder No	Concentration		Uncertainty	Expiration Date
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 - O2 Model No: 4900 Serial No: 49000-652921 Calibration Span: 21.09 % Pollutant: 21.09% O₂ - CC418692

INTERFERENT GAS	INTERFERENT GAS RESPONSE (%)	INTERFERENT GAS RESPONSE, WITH BACKGROUND POLLUTANT (%)	% OF CALIBRATION SPAN ^(a)	
CO ₂ (30.17% CC199689)	0.00	-0.01	0.00	
NO (445 ppm CC346681)	0.00	0.02	0.11	
NO ₂ (23.78 ppm CC500749)	NA	NA	NA	
N ₂ O (90.4 ppm CC352661)	0.00	0.05	0.24	
CO (461.5 ppm XC006064B)	0.00	0.02	0.00	
SO ₂ (451.2 ppm CC409079)	0.00	0.05	0.23	
CH ₄ (453.1 ppm SG901795)	NA	NA	NA	
H ₂ (552 ppm ALM048043)	0.00	0.09	0.44	
HCl (45.1 ppm CC17830)	0.00	0.03	0.14	
NH ₃ (9.69 ppm CC58181)	0.00	0.01	0.03	
	1.20			
	< 2.5%			

⁽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 2014O2-Servomex 4900 2/15/2019

INTERFERENCE CHECK

Date: 12/4/14-12/5/14
Analyzer Type: Servomex - CO₂
Model No: 4900
Serial No: 49000-652921
Calibration Span: 16.65%
Pollutant: 16.65% CO₂ - CC418692

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

⁽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 2/15/2019

BALANCE CALIBRATION LOG

Balance ID:

Balance I	ט:			
Date	Initials	Calibration Weight	Measured Weight ⁽¹⁾	Maintenance and Adjustments
1/16/19	CSW	500.0	419.6	NA Chamsons
1/17/19	USW.	500-0	499.6	NA Camors
1/14/19	WF	500.0	499.7	NA Chemans
1/19/19	Nf	500.0	499-7	NA Chrinous

⁽¹⁾ Must be within \pm 0.5 grams of calibration weight

APPENDIX F LIST OF PROJECT PARTICIPANTS

The following WESTON employees participated in this project.

Paul Meeter	Senior Project Manager
Steve Rathfon	Team Member
Matt Winkeler	Team Member
Chad Walker	Team Member