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## **REPORT OF NOX EMISSION TESTING ON THE #8 BOILER EXHAUST STACK AT THE PFIZER FACILITY LOCATED IN KALAMAZOO, MI**

**Prepared for:** 

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AIR QUALITY DIVISION

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#### 1.0 EXECUTIVE SUMMARY

On February 6, 2019, The Stack Test Group, Inc. performed nitrogen oxides (NOx) emission testing on the #8 boiler at the Pfizer facility located in Kalamazoo, Michigan. Testing was conducted on the exhaust stack of the #8 boiler serving this facility. Three one-hour tests were conducted on this stack. Testing was conducted while firing with natural gas. Presented below are the average results of these tests.

NOx Concentration:	93.1 Parts per Million
NOx Emission Rate:	0.126 Pounds per MMBTU
NOx Emission Limit:	0.20 lb/MMBTU

#### 2.0 INTRODUCTION

The Stack Test Group, Inc. conducted NOx emission testing on the exhaust stack of the #8 boiler (EUEBLR43-8-S1). Testing was performed at the Pharmacia & Upjohn Company, LLC, a subsidiary of Pfizer, Inc., located in Kalamazoo, MI on February 6, 2019. The Portage Steam Generating System is located at 7000 Portage Road, Kalamazoo, MI 49001. The facility generates 175 pounds per square inch gauge pressure steam at 377 degrees Fahrenheit utilizing six boilers. Three of the boilers are stoker coal boilers. Three use natural gas as the primary fuel.

Three NOx tests lasting one-hour in duration were conducted on the #8 boiler. The purpose of this testing was to determine the concentrations and emission rates of the NOx exhausting from the #8 boiler per the requirements of the renewable operating permit (ROP) for this source. The ROP number is MI-ROP-B361002008c. The ROP specifies the NOx testing be completed once during the five year term of this permit.

The exhaust stack for the #8 boiler shares a common stack with the #7 boiler. During this test series, the #7 boiler remained off.

Testing was conducted while Pfizer personnel operated the boiler at the maximum routine operating conditions. Testing was conducted while firing the boiler with natural gas. All pollutant emissions were calculated using the EPA Oxygen F-Factor found in U.S. EPA Reference Method 19. The F-Factor for the #8 boiler was determined to be 8,710 based on the published number in 40 CFR 60 Appendix A Method 19, for natural gas.

Testing was supervised by Mr. Gary A. Kohnke of the Stack Test Group, Inc. and coordinated by Mr. Jeff Robey, Pfizer Environmental of Pharmacia & Upjohn Company, LLC.

All testing followed the guidelines of U.S. EPA Reference Methods 3A and 7E. This report contains a summary of results for the above mentioned tests and all the supporting field, process, and computer generated data.

# 3.0 SAMPLING AND ANALYTICAL PROCEDURES

# 3.1 Oxygen (O<sub>2</sub>) & Nitrogen Oxides (NOx)

# 3.1.1 Sample Collection

Oxygen concentrations and nitrogen oxides (NOx) emissions were determined using U.S. EPA Methods 3A & 7E. A gas sample was drawn from the exhaust duct through a filter and transported to a Universal gas conditioner through a heated Teflon line set to  $250^{\circ}$ F. The gas conditioner removed moisture from the gas stream and pumped a dry gas sample through a Teflon line and manifold flow system to a Servomex Model 1440C paramagnetic oxygen analyzer and a Thermal Electron Corporation (TECO) Model 42C Chemiluminescent NO-NO<sub>2</sub> gas analyzer.

## 3.1.2 Sample Duration and Frequency

The samples were collected in triplicate with each test lasting sixty minutes in duration. A sample was drawn at least twice as long as the response time before the beginning of each test. The response time was approximately 40 seconds.

## 3.1.3 Calibration

At the beginning of the test series, the analyzers were calibrated and then checked for calibration error by introducing zero, mid-range and high-range calibration gases to the back of each analyzer. Following each test run, a system bias was performed by introducing a zero and mid-range calibration gases to the outlet of the probe. Calibration gases used were U.S. EPA Protocol 1 certified.

## 3.1.4 Data Reduction

The analyzer outputs were recorded on a data logger and laptop computer. These oneminute data logger readings were then average using an Excel spreadsheet. The raw data logger readings are included in Appendix E.

## 4.0 <u>TEST RESULTS</u>

Presented in this section are the results of this test series. Test results are reported in Table 4.1. Table 4.1 presents the NOx results from the #8 boiler exhaust stack. The NOx results in parts per million (ppm) and lb/MMBTU and the oxygen results in terms of percent.

Copies of the calculations used to determine these emission rates may be found in Appendix A. Copies of the calibration data are presented in Appendix B. Copies of the NOx and  $O_2$  data are presented in Appendix C. Copies of the calibration gas certification sheets are presented in Appendix D.

#### Table 4.1

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#### NOx Test Results Pfizer 02/06/19 #8 Boiler Exhaust Stack Firing Natural Gas

Test No: Start Time: Finish Time:	<u>T1</u> 08:05 AM 09:05 AM	<u>T2</u> 09:15 AM 10:15 AM	<u>T3</u> 10:22 AM 11:22 AM	<u>Avg.</u>
% Oxygen:	4.8	4.8	4.8	4.8
NOx Results: NOx, ppmv: NOx, lb/DSCF: LBS/MMBTU (F-Factor = 8,710):	94.0 1.12E-05 0.127	93.7 1.12E-05 0.126	91.6 1.09E-05 0.124	93.1 1.11E-05 0.126

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# APPENDIX A

# SAMPLE CALCULATIONS

# SAMPLE CALCULATIONS

The tables presenting the results are generated electronically from raw data. It may not be possible to exactly duplicate these results using a calculator. The reference method data, results and all calculations are carried to sixteen decimal places throughout. The final table is formatted to an appropriate number of significant figures.

1. Volume of water collected (wscf)

V<sub>wsid</sub>

 $= (0.04707) (V_{lc})$ 

Where:

Vic	total volume of liquid collected in impingers and silica gel (ml)
Vwstd	volume of water collected at standard conditions (ft <sup>3</sup> )
0.04707	conversion factor (ft <sup>3</sup> /ml)

2. Volume of gas metered, standard conditions (dscf)

 $=\frac{(17.64)(V_m)\left(P_{bar}+\frac{\Delta H}{13.6}\right)(Y_d)}{(460+T_m)}$ 

Pbar	barometric pressure (in. Hg)
Γm	average dry gas meter temperature (°F)
Vm	volume of gas sample through the dry gas meter at meter conditions $(ft^3)$
V <sub>mstd</sub>	volume of gas sample through the dry gas meter at standard conditions $(ft^3)$
$\mathbf{Y}_{\mathbf{d}}$	gas meter correction factor (dimensionless)
ΔH	average pressure drop across meter box orifice (in. H <sub>2</sub> O)
17.64	conversion factor (°R/in. Hg)
13.6	conversion factor (in. H <sub>2</sub> O/in. Hg)
460	°F to °R conversion constant

# **SAMPLE CALCULATIONS (CONTINUED)**

3. Volume of gas metered, standard conditions (dscm)

 $=\frac{\left(V_{mstd(ft)}\right)}{35.35}$ 

Where:

V <sub>mstd(ft)</sub>	volume of gas sample through the dry gas meter at standard conditions $(ft^3)$
V <sub>mstd(m)</sub>	volume of gas sample through the dry gas meter at standard conditions $(m^3)$
35.35	conversion factor ( $ft^3$ to $m^3$ )
13.6	conversion factor (in. H <sub>2</sub> O/in. Hg)

4. Sample gas pressure (in. Hg)

$$P_s = P_{bar} + \left(\frac{P_g}{13.6}\right)$$

Where:

$\mathbf{P}_{\mathbf{bar}}$	barometric pressure (in. Hg)
Pg	sample gas static pressure (in. H <sub>2</sub> O)
Ps	absolute sample gas pressure (in. Hg)
13.6	conversion factor (in. H <sub>2</sub> O/in. Hg)

5. Actual vapor pressure (in. Hg)<sup>1</sup>

 $= P_s$ 

 $P_{v}$ 

Where:

<u>vi</u> v.	
Pv	vapor pressure, actual (in. Hg)
Ps	absolute sample gas pressure (in. Hg)

6. Moisture content (%)  $B_{wo} = \frac{V_{wsid}}{V_{msid} + V_{wsid}}$ 

Bwo	proportion of water vapor in the gas stream by volume (%)
V <sub>mstd</sub>	volume of gas sample through the dry gas meter at standard conditions (ft <sup>3</sup> )
Vwstd	volume of water collected at standard conditions (ft <sup>3</sup> )

<sup>&</sup>lt;sup>1</sup> For effluent gas temperatures over 212°F,  $P_v$  is assumed to be equal to  $P_s$ .

# SAMPLE CALCULATIONS (CONTINUED)

7. Saturated moisture content (%)

 $=\frac{\left(P_{\nu}\right)}{\left(P_{\nu}\right)}$ 

Where:

B <sub>ws</sub>	proportion of water vapor in the gas stream by volume at saturated conditions (%)
Ps	absolute sample gas pressure (in. Hg)
Pv	vapor pressure, actual (in. Hg)

Whichever moisture value is smaller is used for  $B_{wo}$  in the following calculations.

8. Molecular weight of dry gas stream (lb/lb·mole)

$$M_{d} = M_{CO_{2}} \frac{(CO_{2})}{(100)} + M_{O_{2}} \frac{(O_{2})}{(100)} + M_{CO+N_{2}} \frac{(CO+N_{2})}{(100)}$$

Where:

Mđ	dry molecular weight of sample gas (lb/lb·mole)
Mco <sub>2</sub>	molecular weight of carbon dioxide (lb/lb mole)
Mo2	molecular weight of oxygen (lb/lb·mole)
Mco+N <sub>2</sub>	molecular weight of carbon monoxide and nitrogen (lb/lb·mole)
$CO_2$	proportion of carbon dioxide in the gas stream by volume (%)
O <sub>2</sub>	proportion of oxygen in the gas stream by volume (%)
CO+N <sub>2</sub>	proportion of carbon monoxide and nitrogen in the gas stream by volume (%)
100	conversion factor (%)

9. Molecular weight of sample gas (lb/lb·mole)

 $= (M_d)(1 - B_{wo}) + (M_{H_2O})(B_{wo})$ 

$\mathbf{B}_{wo}$	proportion of water vapor in the gas stream by volume
Md	dry molecular weight of sample gas (lb/lb mole)
М <sub>Н2</sub> о	molecular weight of water (lb/lb·mole)
Ms	molecular weight of sample gas, wet basis (lb/lb·mole)

# SAMPLE CALCULATIONS (CONTINUED)

# 10. Velocity of sample gas (ft/sec)

$$V_{s} = \left(K_{p}\right)\left(C_{P}\right)\left(\overline{\sqrt{\Delta P}}\right)\left(\sqrt{\frac{\left(\overline{T_{s}}+460\right)}{\left(M_{s}\right)\left(P_{s}\right)}}\right)$$

.

Where:

Kp	velocity pressure coefficient (dimensionless)
Cp	pitot tube constant
Ms	molecular weight of sample gas, wet basis (lb/lb·mole)
Ps	absolute sample gas pressure (in. Hg)
Ts	average sample gas temperature (°F)
Vs	sample gas velocity (ft/sec)
$\sqrt{\Delta P}$	average square roots of velocity heads of sample gas (in. H <sub>2</sub> O)
460	°F to °R conversion constant

11. Total flow of sample gas (acfm)

 $= (60) (A_s) (V_s)$ Q,

Where:

As	cross sectional area of sampling location (ft <sup>2</sup> )
Qa	volumetric flow rate at actual conditions (acfm)
Vs	sample gas velocity (ft/sec)
60	conversion factor (sec/min)

12.	Total	flow	of	sample	gas	(dscfm)
						-

$$Q_{std}$$

$$=\frac{(Q_{s})(P_{s})(17.64)(1-B_{wo})}{(\overline{T_{s}}+460)}$$

Bwo	proportion of water vapor in the gas stream by volume
Ps	absolute sample gas pressure (in. Hg)
Qa	volumetric flow rate at actual conditions (acfm)
Qstd	volumetric flow rate at standard conditions, dry basis (dscfm)
Ts	average sample gas temperature (°F)
17.64	conversion factor (°R/in. Hg)
460	°F to °R conversion constant

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# SAMPLE CALCULATIONS (CONTINUED)

# 13. NOx concentration (lb/dscf)

E <sub>ibidscf</sub>

 $=\frac{(ppm)(MW)}{(385.3\times10^6)}$ 

Where:

E <sub>lb/dsef</sub> Conm	emission rate measured concentration in the gas stream (ppmdv)
MW	molecular weight of NOx (46)
385.3	conversion factor

# 14. NOx emission (lb/hr)

$$E_{lb/hr} = (lb/dscf)(60)(dscfm)$$

Where:

E <sub>lb/hr</sub>	emission rate
Elb/dscf	concentration
Edsof	concentration
60min/hr	conversion factor

# 15. NOx emission (lb/MMBTU)

E<sub>Ib / MMBTU</sub>

= (lb/dscf)(F - Factor)(20.9/(20.9 - 02))

E16/MMBTU	emission rate
Elb/dscf	concentration

# APPENDIX B

# CALIBRATION DATA

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# Pfizer Inc. Project #18-3114A Kalamazoo, Mi 2/6/2019 Boller #8

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Analyte	Oxygen		Dry	
Lablel Collection	Makua	Permonto	Calibration	
High	24.01	24	0.04%	
Mid	12.18	12.2	0.08%	
Low (Zero)	0	0,1	0.42%	
Тура	Run Averaga	Pre Zero	Post Zero	Up
Initial System Bias Check			0.1	
Run 1	4.9	0.1	0.2	
Run 2	4.9	0.2	0,1	
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\*Corrected Run Average and Upscale System Bias cell formulas must be adjusted if the high calibration gas is used for post calibration upscale checks.

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Type	Run Averaga	Pre Zero	Post Zero	Pre Upscale	Post Upscale	"Corrected Run Average	Zero System Blas	*Upscale System Blas	Zero Drift	Upscale Drift
Initial System Blas Check			0.1		12.2		0.00%	0.00%		
Run 1	4.9	0.1	0.2	12.2	12.3	4.8	0.42%	0.42%	0.42%	0.42%
Run 2	4.9	0.2	0.1	12.3	12.1	4.8	0.00%	0.42%	0.42%	0.83%
Run 3	4.8	0.1	0.1	12.1	12,1	4,8	0.00%	0.42%	0.00%	0.00%

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Pfizer Inc. Project #18-3114A Kalamazoo, Ml 2/6/2019 Boiler #8

			Calibration
Initial Calibration	Value	Response	Error
High	97.5	97.8	0.31%
Mid	50.25	50.3	0.05%
cw (Zero)	0	0.3	0.31%

\*Corrected Run Average and Upscale System Blas cell formulas must be adjusted if the high calibration gas is used for post calibration upscale checks.

Туре	Run Average	Pre Zero	Post Zero	Pre Upscale	Post Upscale	Corrected Run Average	Zoro System Blas	"Upscale System Blas	Zero Drift	Upecale Drift	lb/decf
Initial System Blas Check			0.4		50.6		0.10%	0.31%			
Run 1	94.0	0.4	0.2	50.6	50.2	0.40	0.10%	0.10%	0.21%	0.41%	1.12E-05
Run 2	94.1	0.2	0.2	50.2	50.9	93.7	0.10%	0.62%	0.00%	0.72%	1.12E-05
Run 3	91.9	0.2	0.3	50.9	50.2	91.6	0,00%	0.10%	0.10%	0.72%	1.09E-05