

N1685

Filterable Particulate Matter 40 CFR Part 63 Subpart UUUUU LEE Demonstration

EUBOILER01 and 02 T.E.S. Filer City Station

T.E.S. Filer City Station 700 Mee Street Filer City, Michigan 48634 Test Dates: October 20-21, 2015

> December 17, 2015 Work Order No. 4101301

> > **Revision 0**

Test Performed by the Consumers Energy Company Regulatory Compliance Testing Section – Air Emissions Testing Body Engineering Services Department Written by G.A. Koteskey, Engineering Technical Analyst II

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1.0 INTRODUCTION

Consumers Energy Company (CECo) Regulatory Compliance Testing Section (RCTS) performed the Filterable Particulate Matter (FPM) Low Emitting Electric Generating Unit (LEE) demonstration per Subpart UUUUU, 40 CFR Part 63 (commonly referred to as the Mercury and Air Toxics Standard [MATS] Rule) at the stack exhausts associated with emissions units EUBOILER01 (Unit 1) and EUBOILER02 (Unit 2) in operation at the Tondu Energy Systems (TES) Filer City Station, located in Filer City, Michigan.

The FPM test was performed to demonstrate qualification as a LEE for FPM. This was the first test performed of the quarterly testing regimen. The FPM LEE demonstration requires quarterly sampling at each unit over a period of three calendar years. The results of each quarterly test must be less than or equal to 50 percent of the applicable FPM standard listed in Table 2 of the MATS Rule (see Table 1.1 below), equating to 0.015 lb/mmBtu for each of Units 1 and 2. A test protocol was submitted to the Michigan Department of Environmental Quality (MDEQ) on September 9th, 2015 and subsequently approved by Mr. Jeremy Howe, MDEQ Environmental Quality Analyst, in his letter dated October 2nd, 2015.

EGU Subcategory	Pollutant Being Sampled	Emission Limit
Existing Unit, Coal-fired not low rank virgin coal	Filterable Particulate Matter	0.030 lb/mmBtu

Table 1.1 - UUUUU, 40 CFR Part 63 (MATS Rule) Emission Limit

1.1 Summary of Test Program

The test program was conducted in accordance with applicable MATS Rule requirements and followed the sampling, calibration and quality assurance procedures specified in U.S. EPA CFR Part 60, Appendix A, Reference Methods (RM) 1, 2, 3A, 4, 5 (MATS Modified) and 19. In addition, to compare filterable particulate matter (FPM) emission rates to permit limits, equations contained in MDEQ Air Pollution Control Rules, Part 10, § R336.2011, Reference Test Method 5B were utilized to determine the amount of excess air and correct the particulate matter concentration to 50% excess air (Attachment 1).



1.2 Key Personnel

RCTS representatives Brian Glendening and Gregg Koteskey conducted the testing on October 20 and 21, 2015. Mr. Richard Brown, TES Environmental Health & Safety Coordinator, coordinated the test program with plant personnel. Mr. Jeremy Howe of the MDEQ observed portions of the testing.

Responsible Party	Address	Contact	
Test Facility	TES Filer City Station 700 Mee Street Manistee, Michigan 49634	Mr. Richard Brown 231-723-6573 Environmental Health & Safety richard.brown@cmsenergy.com	
Test Representative	Consumers Energy Company RCTS - AETB	Mr. Brian Glendening, QSTI Senior Technical Analyst II 616-738-3234 brian.glendening@cmsenergy.com	
& Qualified Individuals	17000 Croswell Street West Olive, Michigan 49460	Mr. Gregg Koteskey, QI Technical Analyst 616-738-3712 gregg.koteskey@cmsenergy.com	
Regulatory Agency Representative	Michigan Department of Environmental Quality 120 W. Chapin Street Cadillac, Michigan 49601	Mr. Jeremy Howe Environmental Quality Analyst 231-876-4416 howej1@michigan.gov	

Table 1.2 – Key Personnel Contact Information

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2.0 SOURCE DESCRIPTION

2.1 Process Description

TES Filer City Station operates a cogeneration power plant with a rated output of 60-megawatts (MW) net and 50,000 pounds of process steam per hour. At full load, each of Units 1 and 2 are capable of producing approximately 320,000 pounds per hour of steam, and this steam is fed to a common steam turbine and electrical generator. The electricity and process steam are sold under contract to public and/or private companies.

Units 1 and 2 are capable of firing mixtures of coal (bituminous and subbituminous), wood and wood waste, construction/demolition (C/D) material, petroleum (pet) coke and tire-derived-fuel (TDF) and are classified as "coal-fired unit not low rank virgin coal" in Item 1 of Table 2 Subpart UUUUU. Starting in 2016, Units 1 and 2 will have the capability to fire natural gas as a clean startup fuel under MATS, as well as at other times for flame stabilization and other purposes. Each unit has a nominal heat input rating of approximately 384 mmBtu.

2.2 Control Device Description

The exhaust gas from each boiler is vented to an individual baghouse for PM control and a spray dryer absorber (SDA) flue gas desulfurization (FGD) system for sulfur dioxide (SO₂) and acid gas control. The abated exhaust is discharged through separate circular stacks which are approximately 250 feet in height.



3.0 SUMMARY OF TEST RESULTS

During the test program, Units 1 and 2 burned a mixture of coal, petroleum coke, tire-derived fuel, and wood. The fuel blend firing rate data for each of the runs is included in Attachment 4. Fuel analysis reports are included in Attachment 3. Please note that the fuel analysis for the mixed coal pile was a composite sample consisting of the mixed coal and pet coke. Testing was conducted at as close to full load as possible with Units 1 and Unit 2 operating at a combined average load of 65.4 MW gross over three test runs. The Unit 1 average steam flow 305 klbs/hr (95% of full load), while the Unit 2 average steam flow was 299 klbs/hr (93% of full load).

3.1 Objectives

The objective of this test was to demonstrate qualification as a LEE for FPM. This was the first test performed of the three-year duration, quarterly testing regimen. The results of each quarterly test must be less than or equal to 50% the particulate matter emission limit of 0.030 lb/mmBtu.

3.2 Test Results and Discussion

As shown in Table 3.1 below, the results of each individual run, as well as the average of the three runs for each unit were below the 40 CFR Part 63 Subpart UUUUU limit of 0.030 lb/mmBtu for Units 1 and 2. Both units demonstrated eligibility for Low Emitting EGU qualification as emission rates were below 0.015 lb/mmBtu (i.e., 50% of the FPM limit). This test program was the first of a series of 12 filterable particulate matter tests that will take place over a period of approximately three years to qualify Units 1 and 2 for LEE status.

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Summary of Filterable PM Emission Test Results									
Source	PM Run Concentratior (gr/dscf)	PM Concentration	PM Emission	PM Concentration (lb/1,000 lbs Gas Flow [*])	PM Emission Rate (lb/mmBtu)				
		(lb/hr)	Result	Result	LEE Qualifi- cation				
	Filterable Particulate Matter								
UNIT 1	1	0.0004	0.29	0.0006	0.0008	-			
	2	0.0003	0.24	0.0005	0.0007	-			
	3	0.0004	0.29	0.0006	0.0008	-			
	Average	0.0004	0.28	0.0006	0.0007	0.015			
	Filterable Particulate Matter								
UNIT 2	1	0.0012	0.90	0.0019	0.0023	-			
	2	0.0005	0.41	0.0009	0.0011	_			
	3	0.0003	0.22	0.0005	0.0006	-			
	Average	0.0007	0.51	0.0011	0.0013	0.015			

Table 3.1 - TES Filer City

* Emissions in pounds of particulate per 1000 pounds gas flow corrected to 50 % excess air.



4.0 SAMPLING AND ANALYTICAL PROCEDURES

PM test runs were performed on the Unit 1 Stack October 20, 2015 and the Unit 2 stack on October 21, 2015. During the testing, each boiler was operating under routine operating conditions as close to full load as possible. Operating data collected at 1-minute intervals during the test runs included CO_2 concentrations, fuel feed rates, steam flow and pressure, stack opacity readings and CEMS derived heat input. It should be noted that the run start and stop times for the CEMS data were adjusted by the response time of the respective CEMS (i.e., 3 minutes for Unit 1 and 4 minutes for Unit 2). In addition, combined Units 1 and 2 gross and net electrical output and unit specific SDA slurry flow rates were logged manually; all process data is presented in Attachment 4. Although the test protocol acceptance letter requested that natural gas fuel flow rate be recorded, installation of the natural gas-fired burners has not yet been completed and no such data was therefore recorded.

4.1 Sampling Location

The number and location of traverse points for determining exhaust gas velocity/volumetric air-flow and particulate concentrations were determined in accordance with U.S. EPA Reference Method 1, Sample and Velocity Traverses for Stationary Sources. The area of the stack was determined and the cross-section divided into a number of equal areas based on existing air flow disturbances. The test location for Units 1 and 2 is on the stack at an elevation approximately 100 feet above stack grade. Each stack is 76 inches in diameter with two 6-inch internal diameter ports apiece that extend 20 inches from the stack interior wall. At this sample location, USEPA Reference Method 1 required a minimum of 12 traverse points for isokinetic particulate sampling. A schematic depicting the Unit 1 and 2 stacks and test port locations is shown in Figures 1-3.

4.2 Velocity and Temperature

The exhaust gas velocity and temperature were determined using U.S. EPA Reference Method 2, *Determination of Stack Gas Temperature and Velocity (Type S Pitot Tube)*. The exhaust gas pressure differential (delta P) was measured at each traverse point during PM testing using an "S Type" Pitot tube connected to a manometer. Exhaust gas temperatures were also measured in conjunction with delta P determinations using a "Type K" thermocouple and a temperature indicator.

Attachment 3 of this report includes cyclonic flow test data as verification of the absence of cyclonic flow at the Units 1 and 2 stack test locations. Method 1, § 11.4.2 indicates *if the average* (null angle) *is greater than 20°, the overall flow condition in the stack is unacceptable, and alternative methodology...must be used.* The average null yaw angle measured at the Unit 1 exhaust on August 20, 2012 was observed to be 3° and the average null yaw angle measured at the Unit 2 exhaust on August 20, 2012 was observed to be 8°, thus meeting the less than 20° requirement. There have been

no ductwork and/or stack configuration changes, so the preceding null angle information is considered to be valid and additional cyclonic flow verification was not performed prior to the PM test.

4.3 Molecular Weight

The exhaust gas composition was determined using U.S. EPA Reference Method 3A, Determination of Oxygen and Carbon Dioxide Concentrations in Emissions from Stationary Sources (Instrumental Analyzer Procedure). Integrated bag samples were obtained by sampling at each traverse point for purposes of determining flue gas molecular weight. The bag samples were analyzed for oxygen using a paramagnetic analyzer. The reference monitor used was calibrated with certified gas standards at three levels and operated following the guidelines of Method 3A.

4.4 Moisture

The exhaust gas moisture content was determined using U.S. EPA Reference Method 4, *Determination of Moisture in Stack Gases* in conjunction with the Method 5 sample apparatus. Exhaust gas was drawn through a series of: one modified Greenburg-Smith (GS) impinger filled with 100 mL of water, one standard GS impinger filled with 100 mL of water, an empty modified Greenburg-Smith (GS) impinger, and one modified GS impinger containing approximately 300 g of silica gel. The impingers were immersed in an ice bath to ensure condensation of exhaust gas moisture and the amount of water vapor collected was determined gravimetrically to calculate exhaust gas percent moisture.

4.5 Filterable Particulate Matter

Filterable PM was collected utilizing 40 CFR Part 60, U.S. EPA Method 5, *Determination of Particulate Matter Emissions from Stationary Sources* with the necessary modifications specified in the MATS Rule for qualifying for low emitting EGU (LEE) status. Specifically, the Method 5 front half temperature was maintained at 320 °F, ± 25 °F, throughout the duration of each test run. A minimum of 2 dry standard cubic meters (dscm) or 70.629 dry standard cubic feet (dscf) of sample volume was drawn through a stainless steel nozzle, a heated stainless steel probe, and a heated glass filter holder containing an 83 millimeter (mm) quartz glass fiber filter followed by a Teflon frit filter support. After each run, filterable PM collected in the nozzle and probe was brushed and rinsed into an appropriately labeled sample bottle using acetone and a Teflon brush. After recovering the quartz FPM filter into a Petri dish labeled "Container #1, Filter", the front half filter holder was recovered with acetone rinses and combined with the probe and nozzle rinse in the sample bottle labeled "Container #2, Probe and Nozzle Rinse". At the laboratory, Method 5 gravimetric analytical procedures were followed to analyze the filters and rinses. All filters and rinses were weighed



multiple times (to ensure a constant weight) in a weighing room maintained at less than 50% relative humidity.

5.0 QUALITY ASSURANCE PROCEDURES

Each U.S. EPA reference method performed contains specific language stating reliable results are obtained by persons equipped with a thorough knowledge of the techniques associated with each method. To that end, factors which could potentially cause sampling errors were minimized by implementing quality assurance (QA) programs into every applicable component of field testing possible. The following QA components were included in this test program.

While not directly required, each PM sample apparatus was leak-checked before each test run as well as immediately after. Extreme care was exercised to minimize effects of stray or ambient particulate at the sampling site, such as ensuring the sample ports are cleaned thoroughly, maintaining enough distance from duct walls and/or other sources of PM so that bias was not introduced artificially. Time, meter box temperature, sample rate, barometric pressure, source temperature and total sample volume was documented for each run. Isokinetic variation was verified to be within Method 5 requirements. Field recovery of the impingers and nozzle/filter particulate catch were carefully performed in an enclosed laboratory area, prior to analysis.

All manual test equipment was calibrated before the test program in accordance with appropriate U.S. EPA procedures. Pitot tubes and thermocouples used to measure the exhaust gas were calibrated following the handbook requirements outlined in Stationary Source-Specific Methods, Method 2, Type S Pitot Tube Inspection, and in ALT - 011, Alternative Method 2 Thermocouple Calibration Procedure Calibration Procedure. Dry test meters used for moisture determination were calibrated using ALT - 009 as described in Method 5, § 16.1, using the procedures in Method 5, § 10.3.2. All applicable equipment calibration documents are included in this report in Attachment 5.

All RM instruments measuring gaseous concentrations were calibrated and operated following applicable methodology based in part on specific quality assurance and quality control requirements contained in Method 7E. Although not required for MATS testing, U.S. EPA Protocol gas standards used by RCTS were purchased from an outside vendor participating in the U.S. EPA Protocol Gas Verification Program (PGVP) calibration gas audit program described 40 CFR Part 75 § 75.21(g). The standards are certified to have a total relative uncertainty of ± 1 percent according to the U.S. EPA Traceability Protocol for Assay & Certification of Gaseous Calibration Standards; EPA – 600/R-97/121; September, 1997 or EPA Traceability Protocol for Assay & Certification of Gaseous Calibration Standards; EPA – 600/R-12/531; May, 2012.



Before beginning the sampling, a three-point analyzer calibration error check was conducted on the RM analyzer by injecting zero, mid and high-level calibration gases directly into the instrument and measuring the response. The instrument response must be within $\pm 2.0\%$ of the respective analyzer span or within 0.5 ppmv absolute difference to be acceptable.

5.1 Field Test Issues

On October 20th, during the analyzer calibration error check prior to testing Unit 1, RCTS noted that the CO₂ analyzer was not passing linearity requirements, however the O₂ analyzer was operating appropriately. RCTS discussed this with MDEQ representative, Mr. Jeremy Howe, and requested permission to use the O₂ values from the RM analyzer and the CO₂ values from the site CEMS equipment. RCTS had performed a CEMS RATA at Filer City Units 1 and 2 August 17-20, 2015. During the RATA testing, the Units 1 and 2 CO₂ CEMS passed with relative accuracies of 1.61 and 5.87 percent, respectively. Mr. Howe approved this request. The CO₂ CEMS data for each sample run is included in Attachment 4. Note that as the plant CEMS sample on a wet basis, the per run moisture determinations were used to correct the wet plant CO₂ concentrations to a dry basis.



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FIGURE 1



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FIGURE 2



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FIGURE 3



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TES Filer City Unit 1 & 2 Particulate Regulatory Compli.

FIGURE 4

Method 5 Filterable Particulate Matter Sample Apparatus

