Report

Emissions Test Federal-Mogul Corporation Test Date: March 5-6, 2014

> Federal-Mogul Corporation 510 E. Grove Street Greenville, Michigan 48838

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APR 1 8 2014 AIR QUALITY DIV. NTH Project No. 73-130459-01 April 16, 2014



NTH Consultants, Ltd. 608 S. Washington Avenue Lansing, MI 48933





1.0 INTRODUCTION

NTH Consultants, Ltd (NTH) has been retained by Federal-Mogul Corporation (Federal-Mogul) to perform emissions testing at the exhaust of a Torrit cartridge collector associated with EU-BABBITTLINE in Permit to Install (PTI) No. 176-07. The facility is located in Greenville, Michigan.

1.1 Purpose of Test

The purpose of this emissions program is to evaluate current emissions at EU-BABBITLINE. Emission testing at this process has been requested by Michigan Department of Environmental Quality (MDEQ).

1.2 Test Date Requirement

This test program was performed March 5-6, 2014.

1.3 Project Contact Information

Location	Address	Contact
Test Facility	Federal-Mogul Corporation 510 E. Grove Street Greenville, Michigan 48838	Ms. Shanda Jennings (616) 754-1240 Shanda.jennings@federalmogul.com
Testing Company Representative	NTH Consultants, Ltd. 1430 Monroe Avenue NW, Suite 180 Grand Rapids, Michigan 49505	Ms. Lori Myott (517) 702-2957 Imyott@nthconsultants.com
State Agency Representative	MDEQ - Air Quality Division Constitution Hall, 2nd Floor South 525 W. Allegan Lansing, Michigan 48909	Mr. Tom Gasloli (517) 335-4861 gaslolit@michigan.gov

The names and affiliations for personnel associated with the test program are presented below.

Mr. Chris Occhipinti, Mr. Tyler Hanna and Ms. Rhiana Dornbos of NTH conducted the testing. Ms. Shanda Jennings of Federal-Mogul provided assistance and coordinated plant operating conditions. Mr. Tom Gasloli and Mr. Eric Grinstern of the MDEQ observed the testing.

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1.4 Summary of Results

A summary of results is presented in Table 1-1. Detailed results can be found appended to this report.

Table 1-1 EU-BABBITTLINE Summary of Test Results and Permit Limits

PM ₁₀ Emissions	Permit Limit	Units
0.6	1.3	lbs/hr
0.003	0.01	gr/dscf

2.0 PROCESS DESCRIPTION

The Babbitt operation (EU-BABBITTLINE) consists of casting, milling, skiving and wire brushing of alloy on steel strip. Coiled steel strips are unwound and enter an induction heat box. From the heat box, the steel strip enters a furnace along with a mixture of hydrogen and nitrogen gases. As the steel strip leaves the furnace, depending on the desired product, gaseous HCI is introduced to the strip for bonding purposes of the Babbitt alloy to the strip. The steel strip then enters a pot of molten tin or Babbitt, depending on the end product, and is re-introduced to gaseous HCI (for bonding purposes of Babbitt only). After this process, the steel strip is applied with molten Babbitt, and then cooled with water. After the steel strip is cooled, it is then wire brushed, skived, milled, and wound into a coil. Emissions from EU-BABBITTLINE are controlled by a ductwork and hood system and a lime-injected Torrit cartridge collector.



3.0 REFERENCE METHODS AND PROCEDURES

Triplicate 120-minute test runs were conducted for PM at each location. The following United States Environmental Protection Agency (U.S. EPA) reference Test Methods were utilized for emissions testing.

- Method 1: Sampling and Velocity Traverses for Stationary Sources
- Method 2: Determination of Stack Gas Velocity and Volumetric Flow Rate
 (Type "S" Pitot Tube)
- Method 3A: Determination of Oxygen and Carbon Dioxide Concentrations in Emissions
 from Stationary Sources
- Method 4: Determination of Moisture Content in Stack Gases
- Method 5: Determination of Filterable Particulate Matter Emissions from Stationary Sources
- Method 202: Dry Impinger Method for Determining Condensable Particulate Emissions from
 Stationary Sources

3.1 Traverse Points

The number of traverse points for exhaust gas velocity and cyclonic air flow was determined in accordance with U.S. EPA Method 1. The EU-Babbitt exhaust duct measured 40 inches at the test location. A total of 12 measurement points were selected at each location utilizing two ports which resulted in 24 traverse points per test port. Diagrams depicting the sampling points and port locations are presented in Figure 1.

3.2 Velocity and Temperature

The exhaust stack gas velocity and temperature measurements were conducted in accordance with U.S. EPA Method 2. The exhaust stack pressure differential (delta P) was measured at each traverse point using a calibrated S-type Pitot tube connected to an appropriately sized inclined water column manometer. Temperatures were recorded in conjunction with delta P determinations using a calibrated Type "K" thermocouple attached directly to the pitot tube.

3.3 Molecular Weight

The exhaust gas composition was determined using U.S. EPA Reference Method 3A. The oxygen and carbon dioxide concentrations were used to determine exhaust gas composition and molecular weight.



3.4 Moisture

The exhaust gas moisture content was determined in accordance with U.S. EPA Reference Method 4. The sample was passed through a series of four impingers, with the first two containing deionized water, the third empty, and the fourth containing silica gel. The impingers were immersed in an ice bath to ensure condensation of the flue gas stream moisture. The amount of water collected was measured gravimetrically to determine moisture content.

3.5 Filterable Particulate Matter

Filterable particulate matter (PM) concentrations were determined following the guidelines of U.S. EPA Method 5. The sample apparatus consisted of a glass nozzle, a heated glass lined probe, a heated 83 mm glass fiber filter, four chilled impingers, and a metering console. The PM sample was collected in the nozzle, probe, and filter. At the conclusion of each test run, the filter was removed from the filter holder, visually inspected and placed into a petri dish. The front half of the filter holder was rinsed with acetone into a separate sample bottle. Acetone blanks were collected during the PM testing.

At the laboratory, Method 5 analytical procedures were used to analyze the samples for PM. The acetone rinses were evaporated and desiccated to dryness and the residue weighed to determine the amount of PM collected. The filters were also desiccated to remove uncombined water and then weighed. A diagram of the PM sampling apparatus is presented in Figure 4.

3.6 Condensable Particulate Matter

The condensable particulate matter concentrations were measured using U.S. EPA Reference Method 202. The exhaust gas was extracted from the sample stream isokinetically through a heated glass lined probe, a glass coil type condenser, a dropout impinger and a modified Greenburg-Smith impinger with an open tube tip, a condensable particulate matter filter holder containing a Teflon_c membrane filter, one impinger containing 100 mL of water and one impinger containing silica gel for moisture collection. All glassware used in the Method 202 sampling train was cleaned prior to testing according to method specifications. During the testing, the condensable particulate matter filter temperatures were monitored and maintained at the method appropriate temperatures through the use of a recirculation pump attached to the condenser, and chilled water surrounded the impinger apparatus. Figure 3 shows the Method 202 apparatus.



4.0 QUALITY ASSURANCE

Each promulgated U.S. EPA reference method described above is accompanied by a statement indicating that to obtain reliable results, persons using these methods should have a thorough knowledge of the techniques associated with each. To that end, NTH attempts to minimize any factors in the field that could increase error by implementing our quality assurance program into every testing activity segment.

The pitot tubes and thermocouples used to measure the exhaust gas during this test program were calibrated according to the procedures outlined in the *Quality Assurance Handbook for Air Pollution Measurement Systems: Volume III, Stationary Source-Specific Methods,* Method 2, *Type S Pitot Tube Inspection,* and Calibration Procedure 2E *Temperature Sensor.*

5.0 SUMMARY OF RESULTS

Plant operations appeared normal throughout the testing event. A leak rate below the allowable limit was unable to be obtained so the sample volume was corrected to account for the ambient air. This leak correction methodology was approved onsite by Tom Gasloli of MDEQ. The leak correction caused isokinetic percentage to fall below 90%, this was discussed and approved by Mr. Gasloli. Results are appended in Table 1 and Appendix B. Operating data was collected by Federal-Mogul and can be found in Appendix C.





TABLE 1

Federal Mogul

Summary of Particulate Matter Emissions

Babbit Line

U.S. EPA Methods 5 and 202

3/5/.	14 -	3/6/	14
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Run No.		2 - A - A	3	Averngo
Date	March 5, 2014	March 5, 2014	March 6, 2014	
Run Time	935-1152	1249-1500	1215-1420	
Sample Duration (Minutes)	120	120	120	
Sample Volume (dscf)	73.9	55.1	91.5	73.5
Volumetric Flow Rates				
Actual Cubic Feet Minute	27,926	20,664	30,925	26,505
Standard Cubic Feet Minute	27,150	19,958	30,020	25,709
Dry Standard Cubic Feet Minute	26,970	19,929	29,927	25,609
Fixed Gases				
Oxygen, % by volume, dry	20.9	20.9	20.9	20.9
Carbon dioxide, % by volume, dry	0.00	0.00	0.00	0.0
Moisture, % by volume	0.7	0.1	0.3	0.4
Emission Rate:				
Filterable Particulate Matter (gr/dscf)	0.001	0.003	0.002	0.002
Filterable Particulate Matter (lb/hr)	0.2	0.4	0.5	0.4
Condensable Particulate Matter (gr/dscf)	0.001	0.001	0.001	0.001
Condensable Particulate Matter (lb/hr)	0.2	0.2	0.1	0.2
PM ₁₀ (lb/hr):	0.5	0.7	0.6	0.6
PM ₁₀ (gr/dscf):	0.002	0.004	0.002	0.003

dscf: gr/dscf : lb/hr: dry standard cubic feet grains per dry standard cubic feet of sample volume collected

: pounds per hour



FIGURES





Figure 2 Method 5 Sampling Apparatus

