

Air Quality Test Report

Secondary Crusher Baghouse Vent:

NSPS OOO Particulate Test

2017

St. Mary's Cement Inc. Charlevoix, Michigan



To the best of our knowledge, this report and the information provided herein is authentic and accurate and follows all statutory requirements relevant to the test program which was carried out.

Certified by:

Project Manager



INTRODUCTION

On December 8th, 2017 AccuAir, LLC (ACCUAIR) was onsite at St. Marys Cement, Inc. to perform air testing at their facility in Charlevoix, Michigan. ACCUAIR was contracted to perform compliance emission testing on the Secondary Crusher Baghouse Vent. The test was conducted in accordance with all appropriate United States Environmental Protection Agency (USEPA) methodologies, and the requirements outlined in the facility's Renewable Operating Permit (ROP) MI-ROP-B1559-2014.

The objective of the program was to demonstrate the compliance status of the site according to NSPS OOO and the requirements of the facility's Federally Enforceable State Operating Permit. Testing on the Vent was performed for the determination of emission rates for Particulate Matter (PM). Mr. Geoff Resney was the onsite project manager, and was assisted by Mr. Keith Woofter. Mr. Robert Dickman of the Michigan Department of Environmental Quality was onsite to observe a portion of the program.

Testing and analysis procedures used for this project are presented in the United States Environmental Protection Agency (USEPA) document Title 40, Code of Federal Regulations, Part 60 (40 CFR 60), Appendix A; the Quality Assurance Handbook for Air Pollution Measurement Systems, Volume III, Stationary Source Specific Methods, and on the USEPA Technology Transfer Network Measurement Center (EPA-TTN-EMC) Emission website. Test Methods Section (http://www.epa.gov/ttn/emc/). The methods are discussed in the Performance Test Procedures section of this report. Included in this test report are sections detailing the Test Program, including Executive Summary, Introduction, Sampling Procedures, Source Information, Sample Calculation and Test Results Sections; with Appendices which include calibrations, field data, calculated data, and laboratory data.

The test procedures referenced in this report are in accordance with the Approved Test Plan for this project. Deviations from the Test Plan or published Reference Methods are documented in the Problems, Deviations and/or Exceptions section of this report. A copy of the Test Plan is provided in Appendix H of this report.

Provided in Table 1-1 are the results of the test program.

Source Unit	Analytes, Units	Three Run Average	Emissions Limit	PASS?
Secondary Crusher Vent	PM gr/dscf	0.002	0.014	YES

Table 1-1. Summary of Performance Test Results



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SUMMARY OF RESULTS

Testing was performed using a Method 5 train for the determination of PM emitted from the secondary crusher vent. Particulate testing consisted of three (3) test runs in accordance with NSPS OOO, the USEPA Methods 1, 2, 3, 4, and 5 and the approved test protocol. Provided in Tables 1-2 are the results of the sampling and operating parameters monitored during testing.

During testing, the processes were operated at maximum normal rates. Process operation was monitored and recorded as motor amperage during the testing. The recorded data is included in the appendices of this report.

Airflow was determined using USEPA Methods 1 and 2. Molecular weight was determined in ambient air and in accordance with Method 3 The moisture content of the stack gas was measured using Method 4, which was performed inclusively with Method 5. Testing procedures are discussed in more detail in the test methods section. All calculated reference method data is provided in Appendix E, and examples of all calculations are presented in the Sample Calculation section of this report.

Problems, Deviations and/or Exceptions

Sampling issues required that the main testing day be delayed one day. NSPS OOO requires that 60dscf be sampled for each run which resulted in the need to sample for approximately 1.5 hours. Production limitations resulted in a shorter than expected final test run that was 0.4 cubic feet short of the required 60.



Run No.	· 1 ·	2	' 3	
Date	12/8/17	12/8/17	12/8/17	
Start Time	9:50	11:44	14:07	
Stop Time	<u>11:31</u>	13:55	15:31	AVG
Nozzle Diameter, in.	0.188	0.188	0.188	0.188
Barometric Pressure, in. Hg	28.40	28.40	28.40	28.40
Net Sampling Time, minutes	96.0	84.0	74.0	84.7
Volume Metered, cf	70.367	63.211	56.514	63.364
Avg. DGM Temp, F	48.42	47.50	49.08	48.33
AVG Delta H, in of H2O	1.61	1.67	1.65	1.64
AVG Delta H, in of Hg	0.1181	0.1225	0.1214	0.1207
DGM Calibration Factor	1.0033	1.0033	1.0033	1.0033
Volume of Gas Collected, dscf	69.883	62.900	56.058	62.947
Total Water Collected, mL	21.0	19.0	18.0	19.3
Volume of Water Vapor, scf	0.990	0.896	0.849	0.912
Moisture, %	1.4	1.4	1 .5	1.4
Dry Mole Fraction, 100-%M	0.9860	0.9860	0.9851	0.9857
CO2 at Stack, % dry	0.50	0.50	0.50	0.50
O2 at Stack, % dry	20.90	20.90	20.90	20.90
CO + N2, % dry	78.60	78.60	78.60	78.60
Dry Molecular Weight, lb/lb mole	28.92	28.92	28.92	28.92
Wet Molecular Weight, lb/lb mole	28.76	28.76	28.75	28.76
Stack Diameter, in.				
Stack Area, sq. in.	400.0	400.0	400.0	400
Static Pressure, in. of H2O	0.50	0.50	0.50	0.50
Stack Pressure, in. of Hg	28.44	28.44	28.44	28.44
Avg. Stack Temp., F	41.0	41.0	41.0	41.0
Avg. Sqroot of Delta P	<u>1.1284</u>	1.1510	1.1434	1.1409
SDE Average	25.257	25.763	25.593	25.538
Pitot Coefficient	0.84	0.84	0.84	0.84
Stack Gas Velocity, afpm	3805.5	3881.9	3856.8	3848.1
Stack Flowrate, wet acfm	10,571	10,783	10,713	10,689
Stack Flowrate, dry scfm	10,440	10,649	10,571	10,554
sokinetics, %	1010	101.9	103.8	102.2

PARTICULATE DATA FRONT HALF ONLY						
Front Half Rinse (FHACE), g	0.007	0.007	0.009	0.0077		
Particulate Filter (PF), g	0.003	0.001	0.001	0.0018		
Total Disolved Solids, g						
Organics, g						
Total Particulate Collected, g	0_0	0.0	۳ 0.0	0.0095		
Grain Loading, gr/dscf	0.0023	0.0020	0.0027	0.0023		

Table 1-2. Secondary Crusher Vent Summary Results



OPERATING PARAMETERS AND SOURCE INFORMATION

The facility is a dry process operation and acquires shale and limestone materials form a nearby quarry, as well as other locations, and purchases mill scale, slag, bottom ash, bauxite, sand, iron ore, and gypsum.

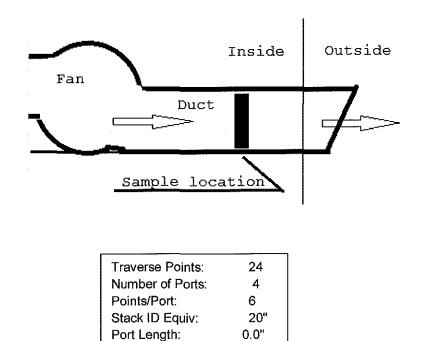
SMC prepares the raw materials through pyro-processing that takes place in a kiln and raw mill (in-line kiln/raw mill). The raw materials are dried in the raw mill by recirculated gas from the kiln system. The materials are fed through a preheater/precalciner countercurrent to the hot gas stream from flash furnace. After passing through the kiln, the raw materials are in the form of clinker, which is milled with other constituents to form Portland cement.

The raw material conversion process area, referred to as the in-line kiln/raw mill, contains two process stacks identified as the SVMAIN and SVBYPASS stacks. Two baghouses control the particulate matter generated from the process operations.

Figures 1-1 show the stack and the test location.



It is a 20" square duct with a slot cut from the side for sampling access. The duct is at platform level inside a building and vents out the side of that building. The sampling location is 48" downstream and 35" upstream of the nearest disturbance (ID fan) and the exit respectively.





Performance Test Procedures

USEPA Reference Methods

This section provides a detailed description of the individual USEPA Reference Methods employed in this test (40 CFR 60, Appendix A). Schematics of the various sampling systems used to perform the test program on the sources can be found in Figures 1-1 through 1-4. Specifics for the test equipment utilized in this program are presented in Appendix A.

USEPA Method 1: Sample and Velocity Traverses for Stationary Sources

Sampling traverse points were determined based on the ratio of the stack diameter to the upstream and downstream distances of the sampling plane to the closest disturbances. The minimum number of traverse points on the sampling plane is determined from Figure 1-2 and Table 1-2 of 40CFR60, Appendix A, Method 1. (ACCUAIR SOP: GSC-TM002)

USEPA Method 2: Determination of Stack Gas Velocity and Volumetric Flow Rate (Type S Pitot Tube)

Velocity and volumetric flow rates were determined from the measurement of the stack gas velocity head with a Type-S Pitot tube and inclined manometer. The Pitot tube was constructed per USEPA Method 2 design specifications. Based on the face opening alignments, external tubing diameter, and base-to-opening plane distances, a coefficient value of 0.84 was assigned to the Pitot tube. (ACCUAIR SOP: GSC-TM003)

USEPA Method 3: Determination of Stack Gas Molecular Weight

The composition of the stack gas was determined to be essentially ambient air. Therefore, the dry molecular weight of 29.0 lb/lb-mole from Section 8.6 from Method 2 was used in the calculations. (ACCUAIR SOP: GSC-TM012)

USEPA Method 4: Determination of Moisture Content

USEPA Method 4 was performed inclusively with Method 5/202 to measure the moisture content of the stack gas for calculating the volumetric flow rate and molecular weight on a dry basis. The impingers for each high-flow sample train were weighed before and after each run. (ACCUAIR SOP: GSC-TM016)



USEPA heated Method 5: Filterable Particulate Matter

Filterable particulate matter (FPM) sampling was conducted utilizing Modified USEPA Method 5 where the probe and filter were separate from, but connected to the impinger train by a heated line. Samples were collected over a 60-minute period. The gas was drawn from the outlet of the source through a nozzle and heated probe liner enclosed in a stainless steel sheath, through a pre-tared heated filter, and into an impinger train for collecting moisture. The probe was equipped with type S pitot tubes for measuring gas velocity and a thermocouple sensor for measuring stack gas temperature. The thermocouple sensor was connected to a digital thermocouple indicator, which was used to measure the stack gas temperature at each sample point. The temperature of the filter exit was monitored to assure that it remained at the proper temperature. A temperature gauge capable of measuring temperature to within 3°C (5.4°F) is installed.

The sample train was connected to a meter box by means of an umbilical cord, which contained a vacuum hose, pitot lines, thermocouple wires and a 4-wire electrical cord. The meter box was used to monitor stack conditions and to facilitate isokinetic sampling. The meter box consists of a leak-free pump used to pull the stack gas through the sample train, fine, and coarse metering valves to control the sampling rate, a vacuum gauge that measures the pressure drop from the sampling nozzle to the metering valves, and a calibrated dry gas meter readable to 0.001 cubic feet (ft³). The dry gas meter inlet and outlet temperatures were monitored by thermocouples, which were connected to a The dry gas meter calibration factor, Y, was multichannel thermocouple indicator. determined by calibrating the meter against a wet test meter or calibrated dry gas meter. At the outlet of the dry gas meter was a calibrated orifice that was used to monitor the flow of aas through the metering system to assure that samples were collected isokinetically. The pressure drop across the orifice was monitored at each sample point. The pitot tubes utilized to measure stack gas velocity were connected to the control box via the umbilical cord. The control box contained either low and high range magnehelic gauges or an incline manometer, which were used for the velocity measurement.

A stack sample was drawn isokinetically from the stack through a nozzle and unheated probe liner, through an unheated and tared filter. At the exit of the filter housing the sample was then passed through a Method 4 moisture train.

The sample train was recovered with acetone per USEPA Method 5 procedures and the filterable particulate matter quantified gravimetrically after evaporation of the acetone and weighing of the filter. The moisture train impingers were weighed onsite after each run to determine the amount of water collected during sampling.

USEPA Method 9 (not performed)

An observer momentarily observes the emission point at 15 second intervals. After a few seconds, the observation is recorded on a form. Observations are recorded in 5-percent opacity intervals. This continues until the required number of observations are completed. Opacity is determined as an average of consecutive observations over a 30 minute period.

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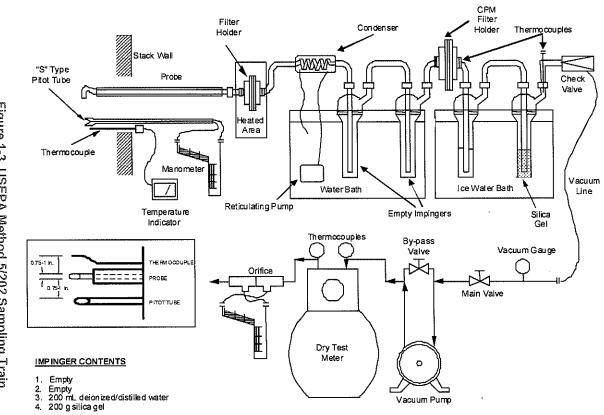


Figure 1-3. USEPA Method 5/202 Sampling Train

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SAMPLE CALCULATIONS

The following calculations were used in the determination of emission rates for the unit's exhaust.

Absolute Stack Gas Pressure (P_s)

$$P_s = P_{bar} + \frac{P_{static}}{13.6}$$

Where:

P_{bar} = Barometric pressure (in. Hg) P_{static} = Static pressure of stack gas (in. H₂O)

Gas Volume Sampled at Standard Conditions (V_{m(std)})

$$V_{m(std)} = \left(\frac{528}{29.92}\right) x V_m x Y \left[\frac{P_{bar} + \left(\frac{\Delta H}{13.6}\right)}{T_m}\right]$$

Where:

 V_m = Actual gas volume sampled (ft³) Y = Gas meter calibration factor P_{bar} = Measured barometric pressure (in. Hg) ΔH = Average differential pressure (in. H₂O) T_m = Absolute average meter temperature (°R)

Water Vapor Collected at Standard Conditions (Vw(std))

$$V_{w(std)} = 0.04715 \ x \ V_{lc}$$

Where:

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Measured Stack Gas Moisture Content (Bws)

$$B_{ws} = \frac{V_{w(std)}}{\left(V_{w(std)} + V_{m(std)}\right)}$$

Wet Molecular Weight of Stack Gas

$$M_{s} = M_{d}(1 - B_{ws}) + 18B_{ws}$$

Stack Gas Velocity

$$v_s = (85.49)(C_p)(avg\sqrt{\Delta P})\sqrt{\frac{T_s}{(P_s)(M_s)}}$$

Where:

 C_p = Pitot coefficient (0.84) T_s = Average stack temperature (°R) P_s = Absolute stack gas pressure (in. Hg) M_s = Molecular weight of stack gas (lb/lb-mole)

Volumetric Flow Rate (Actual cubic feet per min)

$$Q_{aw} = v_s \, x \, A_s \, x \, 60$$

Where:

 Q_{aw} = Volumetric flow rate (acfm) v_s = Stack Gas Velocity (ft/sec) A_s = Stack Area (ft²)



Volumetric Flow Rate (Standard Conditions, dry basis)

$$Q_{sd} = \left(\frac{528}{29.92}\right) x \left(Q_{aw}\right) x \left(\frac{P_s}{T_s}\right) x \left(1 - B_{ws}\right)$$

Where:

 Q_{sd} = Volumetric flow rate (dscfm) P_s = Absolute stack gas pressure (in. Hg) T_s = Average stack temperature (°R) B_{ws} = Stack moisture content

Particulate Matter Sample and Emission Rate Calculations

Isokinetic Sampling Rate

$$I = \left(\frac{0.0945 \, x \, T_s \, x \, V_{m(std)}}{P_s \, x \, v_s \, x \, A_n \, x \, \theta \, x \, (1 - B_{ws})}\right)$$

Where:

 $T_{s} = Stack \ temperature \ (°R)$ $V_{m(std)} = Sample \ volume \ at \ standard \ conditions \ (ft^{3})$ $P_{s} = Absolute \ stack \ gas \ pressure$ $v_{s} = Stack \ gas \ velocity \ (fps)$ $A_{n} = Area \ of \ nozzle \ opening \ face \ (ft^{3})$ $\theta = Run \ time \ (minutes)$ $B_{ws} = Stack \ gas \ moisture \ content$

Stack Gas Particulate Concentration (grains per dscf)

$$C_s = 0.01543 \ x \frac{m_n}{V_{m(std)}}$$

Where:

 C_s = Particulate concentration (gr/dscf) m_n = Total particulate collected in sample (mg) $V_{m(std)}$ = Sample volume at standard conditions (ft³)

Stack Gas Particulate Emission Rate (lb/hr)

$$E = \frac{C_s}{7,000} x Q_{std} x 60$$

Where:

E = Particulate emission rate (lb/hr) C_s = Particulate concentration (gr/dscf) Q_{std} = Stack gas flow rate (dscfm) 7,000 = Grains per pound

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Stack Gas Particulate Emission Rate (tons/yr)

$$E_{tons} = \frac{E x 8,760}{2,000}$$

Where:

E_{tons} = Particulate emission rate (tons/hr) *E* = Particulate emission rate (lb/hr) 8,760 = Hours per year 2,000 = Pounds per ton RECEIVED FEB 06 2018

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Process Data Calculations

Total Solids Used (lb/hr)

$$TS = CA x \frac{CSC}{100} x CD$$

Where:

TS = Total Solids Used (lb/hr) CA = Coating Applied (gal/hr) CSC = Coating Solids Content (% by WT) CD = Coating Density (lb/gal)

Solids Applied to Can (lb/hr)

 $SA = CC \times AFW \div 1,000 \div 453.59$

Where:

SA = Solids Applied to Cans (lb/hr) CC = Cans Coated per Hour AFW = Applied Film weight (mg/can) 1,000 = Milligrams per gram (mg/g) 453.59 = Grams per pound (g/lb)

Solids Exhausted to Control (lb/hr)

$$SE = TS - SA$$

Where:

SE = Solids Exhausted to Control (lb/hr) TS = Total Solids Used (lb/hr) SA = Solids Applied to Cans (lb/hr)



Particulate Collection Efficiency (%)

 $CE = (SE - FPM) \div SE \times 100$

Where:

CE = Particulate Collection Efficiency (%)

SE = Solids Exhausted to Control (lb/hr)

FPM = Filterable Particulate Matter emission rate (lb/hr)

Particulate lbs per 1000 lbs Exhaust Gas

lbs/1000lbs = (E/EGE)x1000

Where:

lbs/1000lbs =Lbs Particulate Per 1000Lbs Exhaust Gas E = PM Emission Rate (lb/hr) EGE = Exhaust Gas Emission Rate (lb/hr)