

# AIR EMISSION TEST REPORT FOR THE VERIFICATION OF AIR POLLUTANT EMISSIONS FROM AN ELECTRIC ARC FURNACE BAGHOUSE CONTROL SYSTEM

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# **Report Certification**

# AIR EMISSION TEST REPORT FOR THE VERIFICATION OF AIR POLLUTANT EMISSIONS FROM AN ELECTRIC ARC FURNACE BAGHOUSE CONTROL SYSTEM

# Ervin Industries – Amasteel Division Adrian, Michigan

The material and data in this document were prepared under the supervision and direction of the undersigned.

Impact Compliance and Testing, Inc.

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This test report has been reviewed by Ervin Amasteel representatives and is approved for submittal to EGLE-AQD. A signed ROP certification form (EQP-5736) accompanies this document at the beginning of the report.



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# 1.0 Introduction

Ervin Industries – Amasteel Division (Ervin Amasteel) State Registration Number (SRN) B1754 retained Impact Compliance & Testing, Inc. (ICT) to measure carbon monoxide (CO) emissions from the exhaust of an electric arc furnace (EAF) processes operated at the Adrian, Michigan facility.

The facility is regulated by Michigan Department of Environmental, Great Lakes, and Energy – Air Quality Division (EGLE-AQD) Renewable Operating Permit (ROP) MI-ROP-B1754-2018 and National Emission Standards for Hazardous Air Pollutants (NESHAP) for Area Sources: Electric Arc Furnaces (40 CFR Part 63 Subpart YYYYY). EAF processes are collectively referred to as FG-0009 in MI-ROP-B1754-2018.

Testing was conducted July 11-12, 2023, by ICT personnel Max Fierro and Christian Smith. Assistance and process coordination was provided by Richard Payne, Plant Engineer, Ervin Amasteel.

The exhaust gas sampling and analysis was performed using procedures specified in the approved Test Protocol prepared by ICT dated June 27, 2023. Mrs. Regina Angellotti of EGLE-AQD was on site to observe portions of the test program.



# 2.0 Summary of Test Results and Operating Conditions

### 2.1 Purpose and Objective of the Tests

Conditions of the NESHAP for Area Sources: Electric Arc Furnaces (40 CFR Part 63 Subpart YYYY) require Ervin Amasteel to test initial compliance after any modifications. Subpart YYYYY also states that previous emissions tests may be used to demonstrate compliance provided that the test was conducted within 5 years of the compliance date. MI-ROP-B1754-2018 requires annual CO monitoring in lieu of operating a continuous emissions monitoring system (CEMS).

### 2.2 Operating Conditions During the Compliance Tests

The process operated normally during the triplicate heat-length (approximately 80 to 90 minute) CO test periods. Each heat processed approximately 23.8 tons of scrap steel per hour (ton/hr) during the CO test periods. CO test runs were ended once the facility process performed a tap out where steel production was paused, and the melt cycle (heat) was completed.

Process data and production rates are provided in Appendix 1.

#### 2.3 Summary of Air Pollutant Sampling Results

The gases exhausted from the EAF were sampled for three heat lengths (batch cycles) for determination of carbon monoxide (CO) emission rates and factors.

Table 2.1 presents a summary of the measured exhaust gas flowrate, and CO emission rates compared to the emission limits in the ROP and NESHAP.

The data presented in Table 2.1 is the average of the three test periods. Data for individual test periods is presented at the end of this report in Table 6.1.



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Table 2.1 Summary of measured exhaust gas flowrate and CO emission	rates
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Parameter	Electric Arc Furnace Three-Test Average Result	Permitted Limit
Exhaust Flowrate (dscfm)	213,503	
Scrap Melted (tons)	23.8	30
Melt Cycle Duration (minutes)	104	
CO Emission Factor (lb/ton)	0.88	3
CO Emission Rate (lb/hr)	21.16	90
CO Emission Rate (TpY)	92.7	322.5



# 3.0 Source and Sampling Location Description

# 3.1 General Process Description

Ervin Amasteel manufactures cast steel abrasives using a 30-megawatt (MW) electric arc furnace and heat-treating furnaces. Steel scrap is charged into the furnace and the furnace roof is then closed. Large electrodes are arced within the scrap bringing it to a molten state, which meets quality standards of the facility. The melt rate (scrap to molten metal) is controlled by regulation of amperage and voltage inputs to the EAF electrodes. When in a molten state, approximately one percent (1%) by weight of carbon, manganese, and silicon and a fraction of a percent of aluminum are added as alloys. The molten metal is then poured into a ladle and the melt process is repeated. The facility performs the melt cycles, called "heats", during the evening (off peak) hours.

# 3.2 Rated Capacities and Air Emission Controls

The facility processes and melts approximately 25 tons of scrap steel per hour, or approximately 40 tons per melt cycle (heat). The scrap steel is melted to approximately 3,100 degrees Fahrenheit (°F) prior to being poured into the ladle.

Emissions from melting the scrap metal are directed, prior to discharge to the atmosphere, to a positive-pressure fabric filter baghouse. The emissions are directed to the baghouse via an inline dirty air fan to a water-cooled duct system that terminates into dry ducting. Dry ducting tempers the furnace fume with fugitive emissions captured from furnace charging, tapping, and casting operations.

Appendix 2 presents sampling locations

### 3.3 Sampling Locations

Inlet gas velocity was measured at the baghouse inlet duct which has an inner diameter of 113.5 inches.

Exhaust gas CO concentration and CO<sub>2</sub>/O<sub>2</sub> content was measured inside the inlet duct downstream of where stack velocity is measured, which is at ground level. Due to the variable nature of the EAF exhaust CO concentration and the ground level sampling location not meeting USEPA Method 1 criteria the exhaust gas cannot be classified as unstratified using the guidelines (i.e., the results indicate stratification pursuant to the Method 7E guidelines due the time-dependent variability of the CO concentration). Therefore, the maximum number of sampling points, determined in accordance with USEPA Method 1, were sampled throughout each test period (i.e., twelve points were sampled).

Sampling location diagrams are provided in Appendix 2.



# 4.0 Sampling and Analytical Procedures

A test protocol for the air emission testing was reviewed and approved by the EGLE-AQD. This section provides a summary of the sampling and analytical procedures that were used during the testing periods.

## 4.1 Summary of Sampling Methods

USEPA Method 1	Velocity and sampling locations were selected based on physical stack measurements in accordance with USEPA Method 1.
USEPA Method 2	Exhaust gas velocity pressure and temperature using a Type-S Pitot tube connected to a red oil incline manometer; temperature was measured using a K-type thermocouple connected to the pitot tube.
USEPA Method 3A	Exhaust gas O <sub>2</sub> and CO <sub>2</sub> content determined using paramagnetic and infrared instrumental analyzers, respectively.
USEPA Method 4	Exhaust gas moisture content was determined using the wet bulb/dry bulb technique.
USEPA Method 10	Exhaust gas CO concentration measured using an infrared instrumental analyzer.

Appendix 3 provides sample train drawings and detailed sampling procedures



# 4.2 Sampling Location and Exhaust Gas Velocity Determination (USEPA Methods 1 and 2)

A single inlet duct contributes to the total air volume introduced into the baghouse. The gas velocity and volumetric flowrate for the inlet duct were measured using USEPA Methods 1 and 2.

Velocity measurement points were determined in accordance with the procedures specified in USEPA Method 1. The Pitot tube was positioned at each of the velocity traverse points with the planes of the face openings of the Pitot tube perpendicular to the stack crosssectional plane. The Pitot tube was then rotated to determine the null angle (rotational angle as measured from the perpendicular, or reference, position at which the differential pressure is equal to zero). Velocity pressure measurements were performed at each traverse point using an S-type Pitot tube and red-oil manometer. Temperature measurements at each traverse point were conducted using a K-type thermocouple and a calibrated digital thermometer.

Volumetric flowrate measurements were performed prior to and after each heat-length CO test run with the flowrate measurement used to calculate CO mass emissions.

Appendix 2 provides drawings for the inlet duct sampling locations. Flowrate calculations and field data sheets are presented in Appendix 4.

#### 4.3 Exhaust Gas Molecular Weight Determination (USEPA Method 3A)

Carbon dioxide (CO<sub>2</sub>) and oxygen (O<sub>2</sub>) concentrations were measured concurrently with the CO test runs and the PM test runs using an instrumental analyzer in accordance with Method 3A. A Servomex 1440D single beam single wavelength infrared (SBSW) Gas Analyzer was used to measure the CO<sub>2</sub> content in the exhaust gas. A Servomex 1440D Gas Analyzer equipped with a paramagnetic sensor was used to measure the O<sub>2</sub> content in the exhaust gas.

The flue gas was withdrawn continuously from the inlet duct of the baghouse using a heated Teflon sample line and sample pump. Moisture was removed from the sampled gas stream using a condenser and the conditioned (dried) gas samples were delivered to the instrumental analyzers.

Appendix 4 provides  $O_2$  and  $CO_2$  calculation sheets. Raw instrument data are provided in Appendix 6.



# 4.4 Moisture Content (USEPA Method 4)

Exhaust gas moisture content for the CO test runs was determined by using the wet bulb/dry bulb technique. The moisture content determination worksheet uses two equations to provide the percentage of moisture in an exhaust gas stream.

The following Equation was used to determine moisture content based on the wet bulb temperature and the dry bulb temperature.

$$\%H_2O = \frac{e'' - \frac{(P_a - e'') * (t_d - t_w)}{2,800 - 1.3 * t_w}}{P_a} * 100$$

e" vapor pressure of water at the wet bulb temperature (in. Hg)

- P<sub>a</sub> absolute barometric pressure (in. Hg)
- t<sub>d</sub> dry bulb temperature (°F)
- t<sub>w</sub> wet bulb temperature (°F)

The vapor pressure (e") of water is required in the equation above, and can be determined using the following equation:

$$e'' = (6.07864 * 10^{-6}) * (t_w)^3 - (1.00431 * 10^{-3})(t_w)^2 + (0.075602) * t_w - 1.69343$$

These equations are limited to stack temperatures between 50°F and 200°F. The stack temperatures during each flowrate were within this range.

Appendix 4 provides moisture catch recovery field data sheets.

### 4.5 CO Concentration Measurements (USEPA Method 10)

Exhaust gas CO concentrations were determined during each sample period using a Thermo Environmental Inc. Model 48i Non-Dispersive Infrared (NDIR) Gas Analyzer in accordance with USEPA Method 10.

Exhaust gas was withdrawn continuously from the inlet duct of the baghouse using a heated Teflon sample line, conditioned, and delivered to the CO instrumental analyzer. Sampling was conducted at twelve points within the stack cross-section for a minimum of 5 minutes per point to satisfy stratification requirements.

Appendix 4 provides CO calculation sheets. Raw instrument response data are provided in Appendix 6.



# 5.0 Quality Assurance/Quality Control Activities

Appendix 6 provides sampling equipment quality assurance and calibration data. A summary of these procedures is provided in this section.

### 5.1 Sampling Location and Flow Measurement Equipment

The representative flowrate locations were determined in accordance with USEPA Method 1 based on the measured distance to upstream and downstream disturbances. The flowrate location was determined to be acceptable based on the absence of significant cyclonic flow, which was measured and recorded on field data sheets. The inlet duct diagram is provided in Appendix 2.

Prior to performing the initial velocity traverse each day, the S-type Pitot tube and manometer lines were leak-checked. These checks were made by blowing into the impact opening of the Pitot tube until 3 or more inches of water were recorded on the manometer, then capping the impact opening and holding it closed for 15 seconds to ensure that it was leak free. The static pressure side of the Pitot tube was leak-checked using the same procedure.

### 5.2 Instrumental Analyzer Interference Check

The instrumental analyzers used to measure CO,  $O_2$  and  $CO_2$  have had an interference response test preformed prior to their use in the field, pursuant to the interference response test procedures specified in USEPA Method 7E. The appropriate interference test gases (i.e., gases that would be encountered in the exhaust gas stream) were introduced into each analyzer, separately and as a mixture with the analyte that each analyzer is designed to measure. All of analyzers exhibited a composite deviation of less than 2.5% of the span for all

### 5.3 Instrument Calibration and System Bias Checks

At the beginning of each day of the testing program, initial three-point instrument calibrations were performed for the CO, CO<sub>2</sub> and O<sub>2</sub> analyzers by injecting calibration gas directly into the inlet sample port for each instrument. System bias checks were performed prior to and at the conclusion of each sampling period by introducing the upscale calibration gas and zero gas into the sampling system (at the base of the stainless-steel sampling probe prior to the particulate filter and Teflon® heated sample line) and determining the instrument response against the initial instrument calibration readings.

The instruments were calibrated with USEPA Protocol 1 certified concentrations of CO<sub>2</sub>, O<sub>2</sub>, and CO in nitrogen and zeroed using nitrogen. A STEC Model SGD-710C ten-step gas divider was used to obtain intermediate calibration gas concentrations as needed.



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### 5.4 Gas Divider Certification

A STEC Model SGD-710C 10-step gas divider was used to obtain appropriate calibration span gases. The ten-step STEC gas divider was NIST certified (within the last 12 months) with a primary flow standard in accordance with Method 205. When cut with an appropriate zero gas, the ten-step STEC gas divider delivered calibration gas values ranging from 0% to 100% (in 10% step increments) of the USEPA Protocol 1 calibration gas that was introduced into the system. The field evaluation procedures presented in Section 3.2 of Method 205 were followed prior to use of gas divider. The field evaluation yielded no errors greater than 2% of the triplicate measured average and no errors greater than 2% from the expected values.



### 6.1 Test Results and Allowable Emission Limits

The CO mass emission rate was calculated based on the measured CO concentration in the baghouse inlet duct and the inlet duct volumetric flowrate. The CO emission rate per ton of steel tapped (lb/ton) was calculated based on the weight of scrap that was tapped during a period of time and the elapsed time for each included heat.

The average CO concentration for each test period was between 17.7 and 28.3 ppmvd, with concentration spikes up to around 92 ppm.

Test results in Table 7.1 indicate that Ervin Amasteel is operating within the following CO emission limits in MI-ROP-B1754-2018:

- 90 lb CO/hr on a three-hour average,
- 3.0 lb CO/ton of melted steel, and
- 322.5 tons CO/year.

#### 6.2 Variations from Normal Sampling Procedures or Operating Conditions

The testing was performed as described in the approved test protocol and specified USEPA test methods. During the test event the processes were operated normally, at or near normal maximum achievable capacity. The second heat was aborted due to an equipment malfunction of the sampling probe. The sampling probe was repaired, and a three-step instrument calibration check was performed prior to resuming sampling.



# Table 6.1 Measured CO emissions from FG-0009 exhaust

<b>Test No.</b> Test Date: Test Period:	<b>1</b> 7/11/2023 19:29-20:47	<b>2</b> 7/11-7/12/2023 23:59-01:21	<b>3</b> 7/12/2023 01:50-02:55	Three Test Average
Exhaust Gas Properties				
Exhaust gas flow (dscfm)	208,954	214,307	217,248	213,503
Moisture (% H2O)	1.81	2.23	2.06	2.09
CO2 (%)	0.30	0.30	0.30	0.30
O2 (%)	21.00	21.10	21.10	21.07
Tons scrap tapped per hour				
(Tph)	21.1	22.0	28.4	23.8
Carbon Monoxide Emissions				_
Concentration (ppmvd)	22.0	17.7	28.3	22.67
Emission Rate (Ib/hr)	20.1	16.5	26.8	21.16
CO Permit Limit (lb/hr)	-	-	-	90.0
Emission Rate (ton/yr)	88.0	72.4	118	92.7
CO Permit Limit (ton/yr)	-	-	-	322.5
Emission Factor (lb CO/ton steel)	0.95	0.75	0.95	0.88
Emission Factor Limit	-	-	-	3.0



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# APPENDIX 1

PROCESS OPERATING DATA



Ervin Amasteel July 11-12, 2023 CO Testing Data

#### EAF Booth

Test#
Heat#
KWH
Charge 1 WT
Charge 2 WT
Alloy Weight
Final Weight
Start time
Sec. Chg Time
Ending Temp
Static Pres. On EAF
Tap #
Cycle Time
Pour Time
Pwr on Time in Cycle

(7/11/2023)	(7/11-12/2023)	(7/12/2023)
1	2	3
1	3	4
20,014	19,508	18,537
40,040	40,180	40,080
40,120	40,220	40,300
800	882	900
80960	81282	81280
19:26	23:56	1:47
19:45	0:18	2:09
3220	2948	3070
-0.05	-0.05	-0.05
16, 18, 15, 17	16,18,15,17	16,18,15,17
115	111	86
20:39	1:14	2:52
65.8	64.3	60.1

#### Dust Col. Cntrl Rm

D.C. Static Pres. Control Dampers % Temp @ Fan Fan Amps Pres. Drp Acrs Bgs

- 19 ·

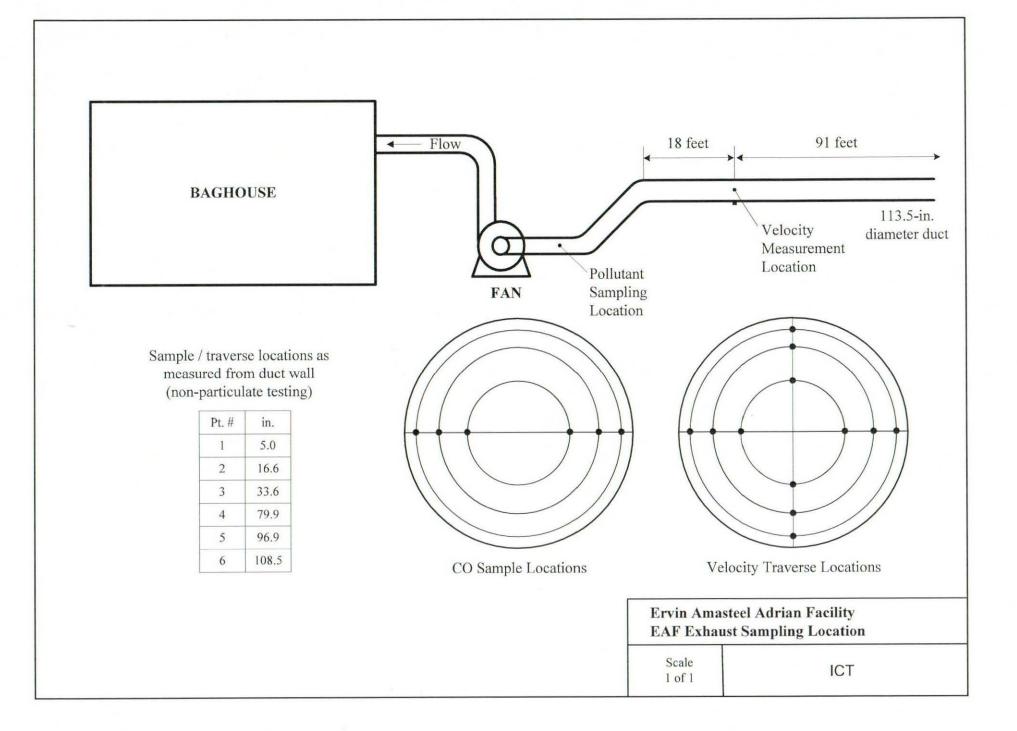
-5.3	-5.9	-6.1
96	96	96
165	139	146
163	159	161
7-1/4"	6-1/4"	6-3/4"

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# **APPENDIX 2**

EXHAUST STACK SAMPLING LOCATIONS





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# **APPENDIX 3**

SAMPLE TRAIN DIAGRAMS AND PROCEDURES

