

Note: This is a reference cited in *AP 42, Compilation of Air Pollutant Emission Factors, Volume I Stationary Point and Area Sources*. AP42 is located on the EPA web site at www.epa.gov/ttn/chief/ap42/

The file name refers to the reference number, the AP42 chapter and section. The file name "ref02_c01s02.pdf" would mean the reference is from AP42 chapter 1 section 2. The reference may be from a previous version of the section and no longer cited. The primary source should always be checked.

Background Report Reference

AP-42 Section Number: 12.20

Background Chapter: 4

Reference Number: 41

Title: Source Test Report of a CM&E
Demonstration Unit Serving a Hard
Chromic Plating Tank at Electrolyzing
Inc., 1947 Hooper Avenue, Los
Angeles, CA 90011

PES

January 1989

ELECTROPLATING

REF. 4-41



PACIFIC ENVIRONMENTAL SERVICES, INC.

WASHINGTON, D.C. • RESEARCH TRIANGLE PARK, NC • LOS ANGELES, CA

Source Test Report of a CM&E Demonstration Unit
Serving a Hard Chrome Plating Tank at
Electrolizing Inc., 1947 Hooper Avenue
Los Angeles, CA 90011

Conducted for the
Metal Finishing Association of Southern California
5000 Van Nuys Blvd
Sherman Oaks, CA 91403

Prepared by
Pacific Environmental Services, Inc.
150 E. Foothill Blvd
Arcadia, CA 91006
1/25/89

Source Test Report of a CM&E Demonstration Unit
Serving a Hard Chrome Plating Tank at
Electrolizing Inc., 1947 Hooper Avenue
Los Angeles, CA 90011

INTRODUCTION

Pacific Environmental Services, Inc. (PES) was contracted by the Metal Finishing Association of Southern California (MFASC), to perform a screening test on the control efficiency of a CM&E demonstration unit used to collect hexavalent chromium emissions from a hard chrome plating operation. Set up and velocity determinations were conducted on January 18, and the source testing was done on January 19, 1989.

In order to evaluate the effectiveness of the CM&E unit, it was necessary to have a hard chrome plating tank that was outfitted with an emission collection system. Electrolizing Inc. in Los Angeles, CA, was the site chosen for the testing. Susan Grant, General Manager of Electrolizing Inc., arranged for the hookup of the demonstration unit and the tank operation during source testing.

The exhaust was taken from hard chrome plating tank No. 10 to supply the CM&E unit. The plating tank was covered by a prefabricated dome enclosure and a flexible hose was attached to the enclosure outlet above the tank. The flexible hose ran across ceiling joints then vertically to the roof, then across the roof and down to the inlet of the CM&E unit (Figure 1). The length of the hose and pipe was approximately 100 feet between the tank and the control unit. The outlet of the tank, the inlet to the CM&E unit, and the outlet of the CM&E unit were all 6" I.D..

On January 18th, velocity traverses were taken at 10 different ports to determine the velocities of the emission system. Two traverses were near the tank exhaust, four traverses were made on the inlet near the unit and four on the outlet (Figure 2). The preliminary velocity traverses above the tank measured 9.4 ft/sec, at the inlet to the CM&E unit it was 8.7 ft/sec, and at the outlet it was 9.8 ft/sec.

Figure 1: Diagram of the Exhaust Line From Plating Tank No. 10 to the CH&E Demonstration Unit.

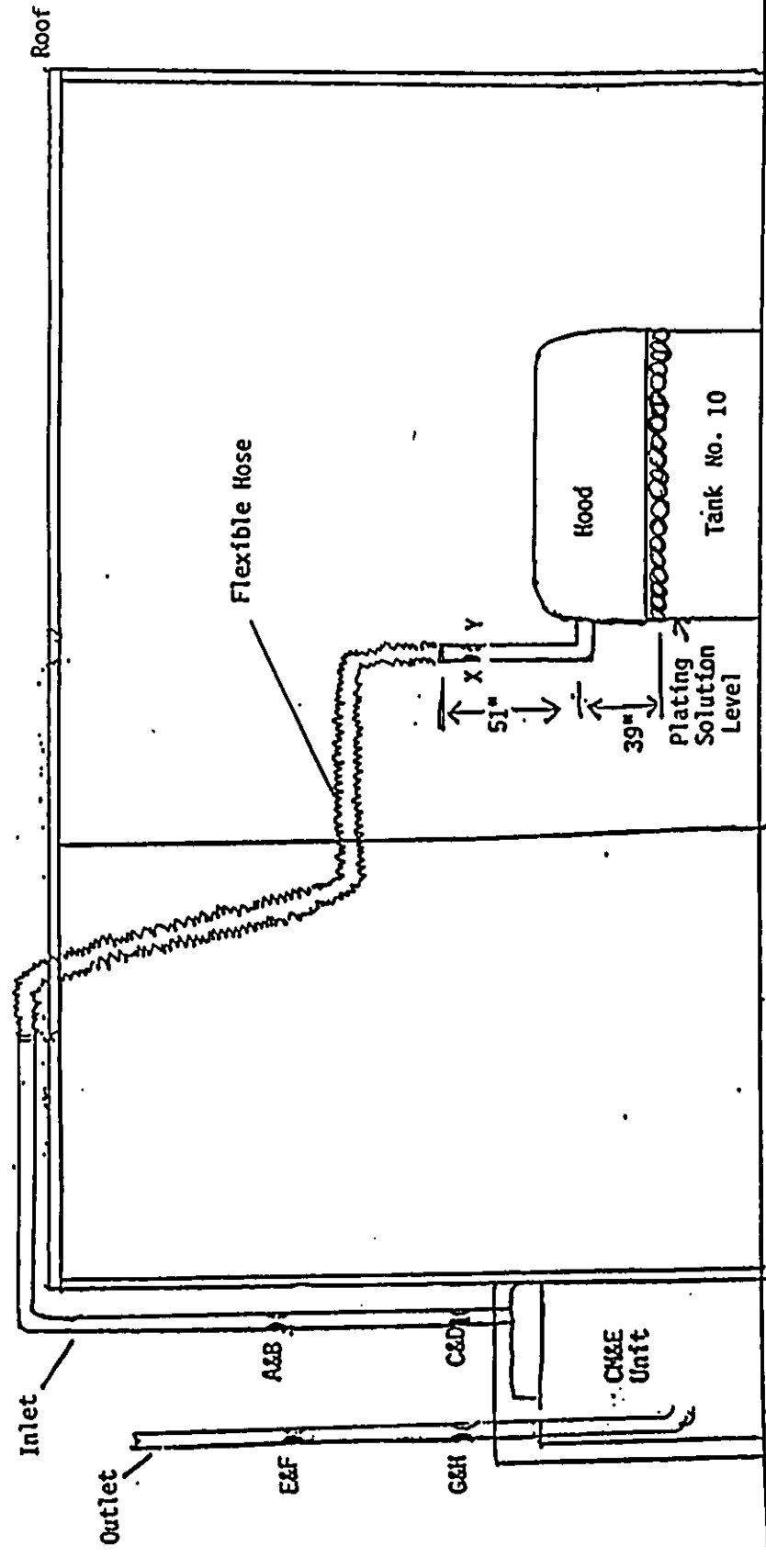
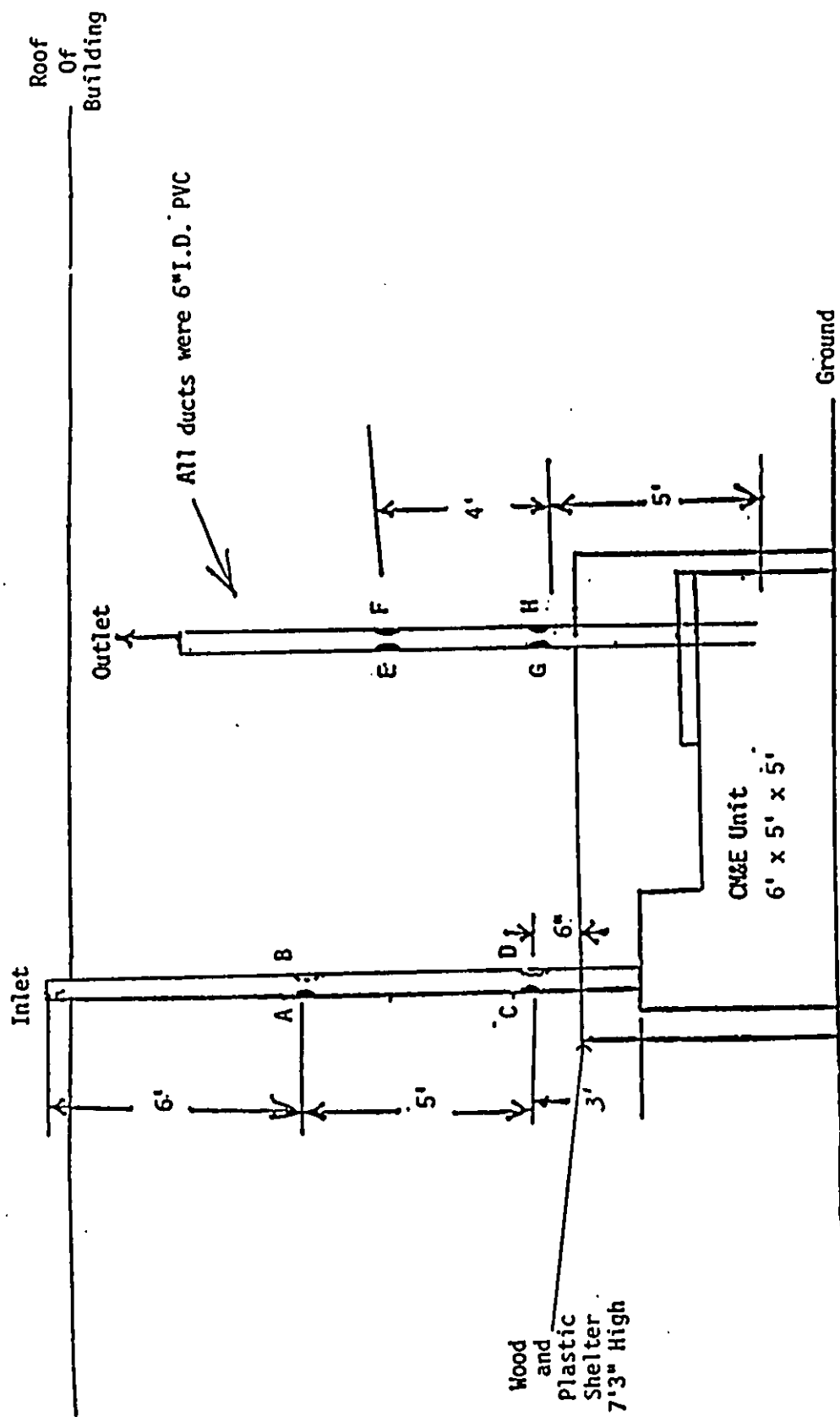


Figure 2: Diagram of Testing Ports for the CM&E Unit



On January 19th, two locations were chosen for sampling. The first sample, CME-I, was taken from the inlet to the unit at sampling port A. This sample was used to determine the quantity of emissions in the gas stream before entering the CM&E unit. The second sample, CME-O, was taken from the outlet of the CM&E unit at sampling port G, in order to determine the emission rate and subsequently the reduction efficiency of the CM&E unit for hexavalent chromium emissions. Continuous velocity measurements were observed in ports C and F.

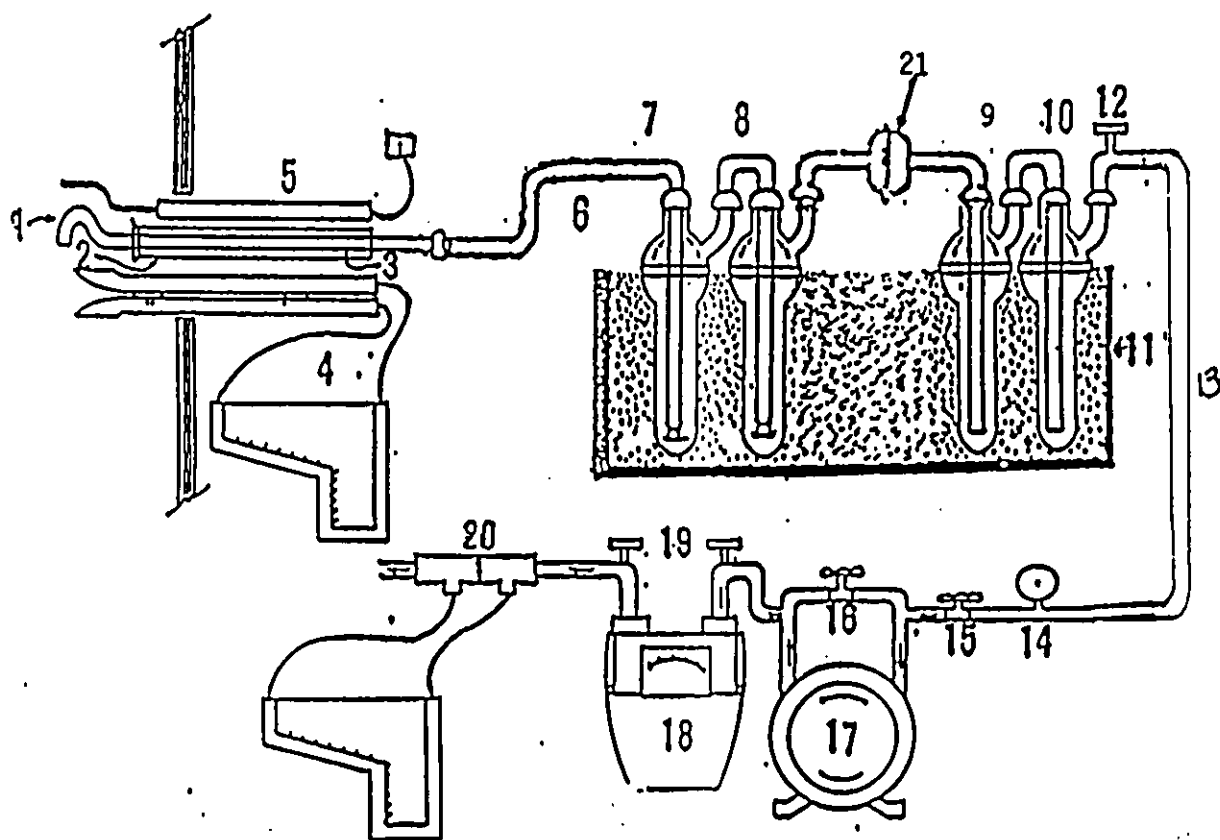
EQUIPMENT AND PROCESS DESCRIPTION

Electrolizing Inc., 1947 Hooper Avenue, Los Angeles, CA 90011, is engaged in the business of hard chrome plating for the aircraft and various other industries. During the test, the CM&E unit served hooded hard chrome plating tank No. 10. The dimensions of the tank were approximately 3.5' x 8' x 3' deep. The tank had a 3" freeboard and had a 3/4" polyball coverage of 150%. During the test the temperature of the tank was maintained at 140 F, the voltage averaged 6.5 volts, and the amperage averaged 1065 amps. The concentration of chrome in the tank was 143,500 mg/l for total chrome and 140,000 mg/l for hexavalent chrome.

The CM&E demonstration unit was approximately 6' x 5' x 5'. It was mounted on a trailer that was 5' x 9' and weighed 1300 pounds. The full name of the unit is the SVS/MECA, which stands for its two stages, the Smart Ventilation System and the Multipurpose Emission Control Assembly. The power requirements were 230 V at 20 amps or 460 V at 10 amps. The demonstration unit was sized for application on tanks 20 sq.ft. or smaller.

TESTING METHODOLOGY

The samples at each location were extracted with modified CARB method 425 source testing trains as described in a 1/20/89 CARB protocol (Figure 3). The source sampling trains consisted of a 0.52" I.D. glass nozzle, a teflon lined stainless steel probe, two impingers each with 100 ml of 0.1N sodium hydroxide, an 89mm teflon coated glass fiber filter placed after the second wet impinger, an empty impinger, an impinger filled with silica gel, a vacuum pump, a dry gas meter, and a calibrated orifice connected to an inclined manometer.



1. Glass or stainless steel sampling nozzle
2. Stainless steel probe sheath
3. Stainless steel, glass, or Teflon-lined probe
4. Type S pitot tube and manometer for ΔP readings
5. Stack temperature sensor
6. 15 ft. teflon probe line with ball and socket connectors
7. Greenburg-Smith impinger with 100 ml. of .1 N NaOH solution
8. Greenburg-Smith impinger with 100 ml. of .1 N NaOH solution
9. Modified Greenburg-Smith impinger - dry
10. Modified Greenburg-Smith impinger filled with 200 grams of silica gel
11. Impinger case filled with ice
12. Impinger exit gas temperature sensor
13. Umbilical line to meter box
14. Vacuum gauge
15. Coarse adjustment valve
16. By-pass valve to adjust ΔH
17. Leak free vacuum pump
18. Dry gas meter
19. Dry gas meter inlet and outlet temperature sensors
20. Orifice meter with manometer for ΔH readings
21. Filter Holder with teflon-lined glass fiber filter.

FIGURE 3
Diagram Of Sampling Train

Sampling at the two ports was conducted simultaneously at a single stack velocity location for a period of just over five hours. Velocity and temperature of the emission streams were measured and recorded every 30 minutes during the test. The meter temperature and the volume of gas through the meter also were measured every 30 minutes during the test. Applied voltage and amperage to the chrome plating tanks and plating bath temperature was recorded by CM&E personnel. All information was recorded on field data sheets (Appendix).

Laboratory analyses were conducted by Thermo Analytical Laboratory in Monrovia, CA. Total chrome concentration and hexavalent chrome concentration were determined for the two stack samples, a blank, and the plating bath solution. Chrome analyses and pH were measured for the scrubbing solution of the CM&E unit. Analyses for total chrome used atomic absorption spectrophotometry and analyses for hexavalent chrome used diphenylcarbazide colorimetric techniques.

RESULTS OF TESTING

After running 5 hours and 7 minutes the test was stopped immediately when water was observed in the silica gel impinger on the outlet train. The bottom of the impinger had broken off and allowed water to be sucked into the impinger. A post sampling leak check of the sample collection system passed at 6" Hg vacuum so we believe that the test results were not seriously effected. A leak in the silica gel impinger would allow the pump to suck ambient air, thus providing a lower concentration of chrome than would be correct. Results of the test should not be used quantitatively but can be used to indicate the general concentration of chrome in the inlet and outlet ducts.

Because of the above difficulty, no moisture determination could be made on the outlet train. It was decided to estimate moisture on both the inlet and outlet assuming saturated air at the measured stack temperature and pressure. The estimate was 2% moisture content.

Calculations were made from the field data sheets to determine sample volume, molecular weight, velocities, flow rates, and isokinetic testing for the gas at each sample location. Stack gas volumes were calculated from the velocity measurements taken during sample extraction. The isokinetic sampling rates were 94.9% for the inlet and 98.4% for the outlet. Results are shown on the emission test calculation sheets (Appendix).

The inlet loading to the CM&E unit was 0.4579 mg/amp-hour for total chrome and 0.3271 mg/amp-hour for hexavalent chrome. The outlet was 0.0091 mg/amp-hour for total chrome and 0.0059 for hexavalent chrome. The control efficiency for total chrome was calculated at 98.02% and for hexavalent chrome was calculated at 98.19% (Tables 1 and 2).

TABLE 1: CONTROL EFFICIENCY OF THE CM&E PYS/MSCA DEMONSTRATION UNIT ON TOTAL CHROME EMISSIONS.

CM&E-I Sample liquid from the impingers in the inlet sample train.
 CM&E-FI Sample caught on the filter in the inlet sample train.
 CM&E-O Sample liquid from the impingers in the outlet sample train.
 CM&E-FO Sample caught on the filter in the outlet sample train.

SAMPLE RUN NUMBER	SAMPLE VOLUME (ml)	Conc. Cr T (mg/l)	Impinger		Total Catch Cr T (mg)	SAMPLE METER VOL (dscf)	EMISSIONS		STACK FLOW RATE (dscfm)	ACTUAL EMISSIONS		AVERAGE AMP-HOURS	EMISSIONS		CONTROL EFFICIENCY Cr T %
			Catch Cr T (mg)	Filter Catch Cr T (mg)			Cr T (mg/scf)	Cr T (mg/hour)		Cr T (mg/amp-hour)					
Inlet					17.7215	233.004	0.0761	106.87	487.69	1065	0.4578				
CM&E-I	545	32.2	17.549												
CM&E-FI				0.1725											
Outlet					0.3655	275.343	0.00133	121.31	9.66	1065	0.0091				98.02
CM&E-O	650	0.46	0.289												
CM&E-FO				0.0665											

TABLE 2: CONTROL EFFICIENCY OF THE CH&E PYS/MECA DEMONSTRATION UNIT ON HEXAVALENT CHROME EMISSIONS.

CH&E-I Sample liquid from the impingers in the inlet sample train.
 CH&E-FI Sample caught on the filter in the inlet sample train.
 CH&E-O Sample liquid from the impingers in the outlet sample train.
 CH&E-FO Sample caught on the filter in the outlet sample train.

SAMPLE RUN NUMBER	SAMPLE VOLUME (ml)	SAMPLE CONC. Cr 6 (ug/l)	Impinger		Filter Catch Cr 6 (ug)	Total Catch Cr 6 (ug)	SAMPLE METER VOL (dscf)	EMISSIONS Cr 6 (ug/scf)		STACK FLOW RATE (dscfm)	ACTUAL EMISSIONS Cr 6 (ug/hour)		AVERAGE AHP-HOURS	EMISSIONS Cr 6 (ug/amp-hour)		CONTROL EFFICIENCY Cr 6 %
			Catch Cr 6 (ug)	Catch Cr 6 (ug)												
Inlet						12.6582	233.004	0.0543	106.87	348.35	1065	0.3271				
CH&E-I	545	23	12.535													
CH&E-FI					0.1232											
Outlet						0.2384	275.343	0.0009	121.31	6.30	1065	0.0059				98.19
CH&E-O	650	0.3	0.195													
CH&E-FO					0.0434											

Plant: Electrolizing Inc
 Date: 1-19-1989
 Source/Sample Number: CNE-T

$$1. V_m(\text{std}) = 17.64 V_m Y \left[\frac{P_{\text{bar}} + (\Delta H / 13.6)}{T_m} \right]$$

$$V_m(\text{std}) = 17.64 (243.005)(1.003) \left[\frac{(29.90) + (2045 / 13.6)}{(554.5)} \right]$$

$$V_m(\text{std}) = \underline{233.004} \text{ dsof}$$

2. Volume Water Vapor Collected, Standard Conditions

$V_{10} = \underline{\hspace{2cm}}$ condensate from impingers and Silica Gel

$$V_H(\text{std}) = .04707 V_{10} = \underline{\hspace{2cm}} \text{ sof}$$

$$V_H(\text{std}) = \underline{\hspace{2cm}} \text{ sof}$$

3. Percent Moisture, by volume

$$B_{ws} = \frac{V_H(\text{std})}{V_H \text{ std} + V_m(\text{std})} = \frac{(\hspace{1cm})}{(\hspace{1cm}) + (\hspace{1cm})} = \underline{0.020} = \text{maximum saturation content}$$

$$B_{ws} = \underline{0.020}$$

4. Molecular Weight, Stack gas

Dry Molecular Weight

$$M_d = 0.440 (\% \text{ CO}_2) + 0.320 (\% \text{ O}_2) + 0.280 (\% \text{ N}_2 + \% \text{ CO})$$

$$M_d = 0.440 (\hspace{1cm}) + 0.320 (\hspace{1cm}) + 0.280 (\hspace{1cm})$$

$$M_d = \underline{29.0} \text{ lb/lb-mole} \quad [40 \text{ CFR, Part 60, Method 2, App A, section 3.6}]$$

$$M_s = M_d + B_{ws} (18 - M_d) = (29.0) + (0.020)(18 - 29.0)$$

$$M_s = \underline{28.78} \text{ lb/lb-mole}$$

Plant: Electrolizing Inc.
 Date: 1-19-1989
 Source/Sample Number: CME-I

5. Stack Gas Velocity Average

$$V_s(\text{avg}) = 85.49 C_D (\sqrt{\Delta P}) \text{avg} \sqrt{\frac{T_s}{P_s M_s}}$$

$$V_s(\text{avg}) = 85.49 (0.84) (0.1656) \sqrt{\frac{(533.6)}{(29.88)(28.78)}}$$

$$V_s(\text{avg}) = \underline{9.37} \text{ - ft/sec.}$$

6. Stack Volumetric Flow Rate, Actual Conditions (Stack Temperature and Pressure)

$$Q_s = 60 V_s A = 60 (9.37) (0.1963)$$

$$Q_s = \underline{110.39} \text{ acfm}$$

7. Stack volumetric flow rate, standard conditions (68° F, 29.92 Hg)

$$Q(\text{std}) = 17.64 Q_s (1 - B_{ws}) \frac{P_s}{T_s} =$$

$$Q(\text{std}) = 17.64 (110.39) (1 - 0.02) \frac{(29.88)}{(533.6)}$$

$$Q_{\text{std}} = \underline{106.870} \text{ dsocfm}$$

8. Isokinetic Variation

$$\%I = K \frac{T_s V_m(\text{std})}{P_s V_s A_n \theta (1 - B_{ws})}$$

$$= 0.0945 \frac{(533.6)(233.004)}{(29.88)(9.37)(0.1963)(307)(1 - 0.020)}$$

$$\%I = \underline{94.9} \%$$

Plant: Electrolizing Inc.
 Date: 1-17-1999
 Source/Sample Number: CME-0

1. $V_m(\text{std}) = 17.64 V_m Y \left[\frac{P_{\text{bar}} + (\Delta H / 13.6)}{T_m} \right]$

$V_m(\text{std}) = 17.64 (284.897) (0.993) \left[\frac{(29.90) + (26.14 / 13.6)}{(515.4)} \right]$

$V_m(\text{std}) = \underline{275.343} \text{ dsof}$

2. Volume Water Vapor Collected, Standard Conditions

$V_{10} = \underline{\hspace{2cm}}$ condensate from impingers and Silica Gel

$V_W(\text{std}) = .04707 V_{10} = \underline{\hspace{2cm}} \text{ sof}$

$V_W(\text{std}) = \underline{\hspace{2cm}} \text{ sof}$

3. Percent Moisture, by volume

$B_{Ws} = \frac{V_W(\text{std})}{V_W \text{ std} + V_m(\text{std})} = \frac{(\hspace{1cm})}{(\hspace{1cm}) + (\hspace{1cm})} = \underline{0.020} \text{ max saturation level.}$

$B_{Ws} = \underline{0.020}$

4. Molecular Weight, Stack gas

Dry Molecular Weight

$M_d = 0.440 (\% \text{ CO}_2) + 0.320 (\% \text{ O}_2) + 0.280 (\% \text{ N}_2 + \% \text{ CO})$

$M_d = 0.440 (\hspace{1cm}) + 0.320 (\hspace{1cm}) + 0.280 (\hspace{1cm})$

$M_d = \underline{29.0} \text{ lb/lb-mole (40 CFR, Section 60, Method 2, App A, ext 3.6)}$

$M_s = M_d + B_{Ws} (18 - M_d) = (29.0) + (0.020)(18 - 29.0)$

$M_s = \underline{28.78} \text{ lb/lb-mole}$

Plant: Electrolizing Inc.
 Date: 1-19-1989
 Source/Sample Number: CME-O

5. Stack Gas Velocity Average

$$V_s(\text{avg}) = 85.49 C_D (\sqrt{\Delta P}) \text{avg} \sqrt{\frac{T_s}{P_s M_s}}$$

$$V_s(\text{avg}) = 85.49 (0.84) (0.1903) \sqrt{\frac{(546.5)}{(29.90)(28.78)}}$$

$$V_s(\text{avg}) = \underline{10.89} \text{ - ft/sec.}$$

6. Stack Volumetric Flow Rate, Actual Conditions (Stack Temperature and Pressure)

$$Q_s = 60 V_s A = 60 (10.89) (0.1963)$$

$$Q_s = \underline{128.26} \text{ acfm}$$

7. Stack volumetric flow rate, standard conditions (68° F, 29.92 Hg)

$$Q(\text{std}) = 17.64 Q_s (1-B_{ws}) \frac{P_s}{T_s} =$$

$$Q(\text{std}) = 17.64 (128.26) (1-0.02) \left(\frac{29.90}{546.5} \right)$$

$$Q_{\text{std}} = \underline{121.31} \text{ dsofm}$$

8. Isokinetic Variation

$$\%I = K \frac{T_s V_m(\text{std})}{P_s V_s A_n \theta (1-B_{ws})}$$

$$= 0.0945 \frac{(546.5)(275.343)}{(29.90)(10.89)(307)(1-0.02)}$$

$$\%I = \underline{98.42} \%$$

plant Electrolyzing Inc.

Date 1-17-89 J
Sampling Location INLET
Sample Type CHROME
Run Number DATE 42 CME-I
Operator SIMON/COTTON
Ambient Temperature 75 F
Barometric Pressure _____
Static Pressure (P_s) -0.3
Filter Number(s) 75 Flan - NA
Pretest Leak Rate = 0.002 cfm @ 15 in. Hg
Pretest Pitot Leak Check _____
Pretest Orsat Leak Check NA
Read and Record all Data Every 30 Minutes

Schematic of Traverse Point Layout

Probe Length and Type 3' Teflon Lined
 Pitot Tube I.D. No. _____
 Nozzle I.D. 0.520" 31955
 Assumed Moisture, % 1.0
 Temp. Readout S/N _____
 Meter Box Number NUTECH
 Meter Δ lg 1.92
 C Factor _____
 Meter Gamma 1.003
 Heater Box Setting NONE
 Reference Δp 0.021
 Post Test Leak Rate = 0.002 cfm @ 10 in. Hg
 Post Test Pitot Leak Check _____
 Post Test Orsat Leak Check N.A.

[illegible]

243.005

Plant	Electrodeizing Inc
Date	1-19-89
Sampling Location	OUTLET - Stack
Sample Type	EPA ANALYSE
Run Number	547-1A 10 ME-0
Operator	CONANT
Ambient Temperature	75
Barometric Pressure	29.9
Static Pressure (P _s)	± 0.02
Filter Number(s)	NA-1552
Pretest Leak Rate	0.081 cfm @ 15 in. Hg
Pretest Pitot Leak Check	
Pretest Orsat Leak Check	1-00204
Read and Record all Data Every	Minutes

Probe Length and Type 3 Teflon
Pitot Tube I.D. No. _____
Nozzle I.D. 0.520 GLASS
Assumed Moisture, % 1
Temp. Readout S/N NUTZLA
Meter Box Number 204
Meter A11g 1.735
C Factor 1.05
Meter Gamma 0.993
Diastere Box Setting NA
Reference Δp 0.26
Post Test Leak Rate = 0.002cfm @ 6 in. Hg
Post Test Pitot Leak Check _____
Post Test Orsat Leak Check _____

Schematic of Traverse Point Layout

[illegible]

SAMPLE RECOVERY DATA

Plant: Electrolyzing Inc.
 Date: 1-19-89
 Sampling Location: SCRUBBER INLET & OUTLET
 Sample Type: EPA CHROME
 Run Number: CME-I and CME-O
 Sample Box Number: SEE FIELD DATA FORM FOR MORE INFO
 Clean-up Man: SIMON/COTTONE
 Job Number: 3625
 Comments: These Samples have filters

FRONT HALF

Filter Number: CME-FI CME-FO
 Description of Filter: 89mm Teflon Coated

MOISTURE

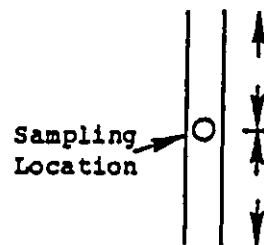
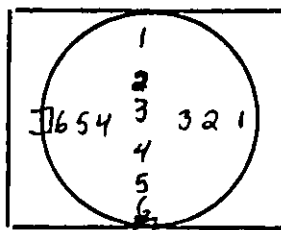
Impingers	<u>≈ 260 ml</u>		<u>280 ml</u>		
Final Volume:	<u>29</u> ml		<u>260 ml</u>	ml	
Initial Volume:	<u>200 ml</u>	ml	<u>200 ml</u>	ml	
Net Volume:	<u>60 ml</u>	ml	<u>90</u>	ml	
Total H ₂ O:					

Silica Gel			(BROKEN IMPINGER)		
Final Volume:	<u>290.9</u>	g	<u>29 LOST</u>	g	
Initial Volume	<u>250.9</u>	g	<u>250.9</u>	g	
Net Volume	<u>40.8</u>	g		g	
Total Moisture:					

Description of Impinger Catch: INLET - LIGHT yellow
OUTLET - clear

POST TEST PRELIMINARY VELOCITY TRAVERSE

Plant: Electrolyzing Inc
 Date: 1-14-1989
 Location: CME Scrubber Inlet & Outlet
 Stack I.D.: 0.5'
 Barometric Pressure, in. Hg: 29.9
 Stack Gauge Pressure, in. H₂O:
 Operators: Simon Cotton
 Pitot Tube I.D. Number: A (Type S) Cp = 0.94
 Temperature Readout I.D.:
 Pitot Tube Leak Check: ✓



Schematic of Traverse Point Layout

Inlet Stack Gauge = -0.3

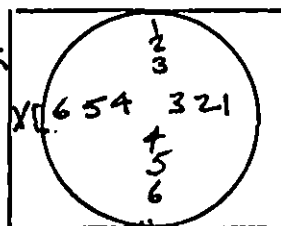
Outlet Stack Gauge = -0.02

Traverse Point Number	Velocity Head (ΔP_g) in. H ₂ O	Stack Temp. (T_g), °F	Cyclonic Flow Check ° from Null
C-1	0.028	73.55°	
C-2	0.031		
C-3	0.030		
C-4	0.029		
C-5	0.027		
C-6	0.012		
D-1	0.026		
D-2	0.026		
D-3	0.027		
D-4	0.027		
D-5	0.026		
D-6	0.025		
		$V = 85.4 (0.84) \sqrt{P_{avg}}$	
		$M_s = 28.62$	
		$P_s = 29.88, T_s = 583.85$	
	$(\sqrt{DP})_{avg}$		Velocity
Average	0.159		9.02 ft/s

Traverse Point Number	Velocity Head (ΔP_g) in. H ₂ O	Stack Temp. (T_g), °F	Cyclonic Flow Check ° from Null
G 1	0.034	545.45	
G 2	0.037	85.95°	
G 3	0.038		
G 4	0.037		
G 5	0.036		
G 6	0.034		
H 1	0.035		
H 2	0.037		
H 3	0.038		
H 4	0.037		
H 5	0.035		
H 6	0.032		
		$T_s = 545.45$	
		$M_s = 28.66$	
		$P_s = 29.9$	
	$\sqrt{DP}_{avg}$		Velocity
Average	0.189		10.83 ft/s

PRELIMINARY VELOCITY TRAVERSE

Plant: ELECTROLYZING INC.
Date: 11-8-80
Location: SCRUBBER INLET ABOVE TANK
Stack I.D.: CME-INLET = 6"
Barometric Pressure, in. Hg: 29.86
Stack Gauge Pressure, in. H₂O: N/A.
Operators: SIMON/PARK
Pitot Tube I.D. Number: "A" C=0.99
Temperature Readout I.D.: —
Pitot Tube Leak Check: ✓



Schematic of Traverse Point Layout

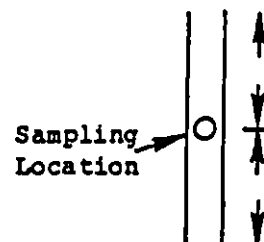
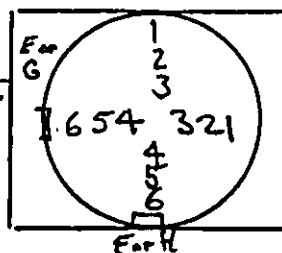
$$\text{Velocity} = 85.49 (0.99) (\sqrt{\Delta P})_{\text{avg}} \sqrt{\left(\frac{T_3}{P_3} \right) \frac{\rho}{M_3}}$$

Traverse Point Number	Velocity Head (Δp_g) in. H ₂ O	Stack Temp. (T_g), °F	Cyclonic Flow Check ° from Null
X-1	0.020	70°	
X-2	0.023	70°	
X-3	0.024	70°	
X-4	0.023	70°	
X-5	0.019	70°	
X-6	0.016	70°	
Y-1	0.014	70°	
Y-2	0.018	70°	
Y-3	0.020	70°	
Y-4	0.020	70°	
Y-5	0.020	70°	
Y-6	0.021	70°	
	$(\sqrt{\Delta P})_{AVG}$		VELOCITY
Average	0.1405		9.37 f/s

[illegible]

PRELIMINARY VELOCITY TRAVERSE

Plant: ELECTROLYZING INC.
 Date: 1-18-89
 Location: SCRUBBER OUTLET
 Stack I.D.: OUTLET FROM CMPE UNIT
 Barometric Pressure, in. Hg: 29.9 27.6
 Stack Gauge Pressure, in. H₂O: -0.30
 Operators: SIMON/PARKS
 Pitot Tube I.D. Number: "A" C=0.99
 Temperature Readout I.D.: -
 Pitot Tube Leak Check: 1



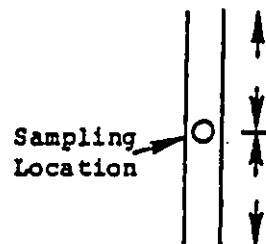
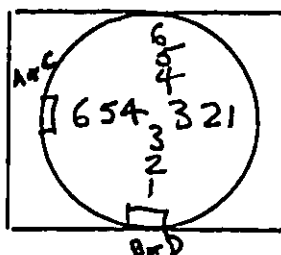
Schematic of Traverse Point Layout

Traverse Point Number	Velocity Head (Δp_g) in. H ₂ O	Stack Temp. (T_g), °F	Cyclonic Flow Check ° from Null
E-1	0.015		
E-2	0.020		
E-3	0.023		
E-4	0.024		
E-5	0.023		
E-6	0.022		
F-1	0.016		
F-2	0.018		
F-3	0.022		
F-4	0.024		
F-5	0.023		
F-6	0.023		
	(VAP) AVG.		VELOCITY
Average	0.1448		9.65 ft/s

Traverse Point Number	Velocity Head (Δp_g) in. H ₂ O	Stack Temp. (T_g), °F	Cyclonic Flow Check ° from Null
G-1	0.020		
G-2	0.023		
G-3	0.025		
G-4	0.024		
G-5	0.023		
G-6	0.020		
H-1	0.016		
H-2	0.020		
H-3	0.022		
H-4	0.024		
H-5	0.023		
H-6	0.022		
	($\sqrt{\Delta P}$) _{AVG}		VELOCITY
Average	0.1475		9.85 ft/s

PRELIMINARY VELOCITY TRAVERSE

Plant: Electrolyzing Inc.
Date: 1-18-89
Location: SCRUBBER INLET
Stack I.D.: INLET TO CASE = 6"
Barometric Pressure, in. Hg: 29.86
Stack Gauge Pressure, in. H₂O: +0.02
Operators: SIMON/PARS
Pitot Tube I.D. Number: A 0.99 = C
Temperature Readout I.D.: -
Pitot Tube Leak Check: ✓



Schematic of Traverse Point Layout

Traverse Point Number	Velocity Head (Δp_g) in. H ₂ O	Stack Temp. (T_g), °F	Cyclonic Flow Check ° from Null
A-1	0.013		
A-2	0.015		
A-3	0.019		
A-4	0.020		
A-5	0.019		
A-6	0.016		
B-1	0.017		
B-2	0.019		
B-3	0.020		
B-4	0.017		
B-5	0.014		
B-6	0.011		
	$(\sqrt{\Delta p})_{\text{AVG}}$		VELOCITY
Average	0.1286		8.57 ft/s

Traverse Point Number	Velocity Head (Δp_s) in. H ₂ O	Stack Temp. (T _a), °F	Cyclonic Flow Check ° from Null
C-1	0.015		
C-2	0.018		
C-3	0.020		
C-4	0.021		
C-5	0.018		
C-6	0.017		
D-1	0.012		
D-2	0.015		
D-3	0.018		
D-4	0.020		
D-5	0.018		
Average	(ΔP) _{AVG} 0.1326		VELOCITY 8.86 F/S

Page 2

Received: 01/20/89

TMA Inc.

REPORT

Work Order # 89-01-092

Results by Sample

SAMPLE ID CME-I

SAMPLE # 01 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR HEX 23. CR L 32.2 WC 545

mg/L mg/L ml

SAMPLE ID CME-O

SAMPLE # 02 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR HEX 0.3 CR L 0.46 WC 650

mg/L mg/L ml

SAMPLE ID CME-B

SAMPLE # 03 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CRHGAW 0.002 CR HEX <0.05 WC 475

mg/L (ppm) mg/L ml

SAMPLE ID CME-FI

SAMPLE # 04 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR S 182.

Total ug



Page 3

Received: 01/20/89

TMA Inc.

REPORT

Work Order # 89-01-092

Results by Sample

SAMPLE ID CME-FO

SAMPLE # 05 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR S

76.

Total ug

SAMPLE ID CME-FB

SAMPLE # 06 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR S

9.5

Total ug

SAMPLE ID CME-EL-PRI

SAMPLE # 07 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR HEX

140,000

CR L

143,500

mg/L

SAMPLE ID CME-SCR-ORIG

SAMPLE # 08 FRACTIONS: A

Date & Time Collected 01/19/89

Category

CR HEX

1.7

CR L

2.20

PH

7.93

mg/L

mg/L

pH units

