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Title:	Source Emission Tests At Stark Ceramics, Inc., East Canton, Ohio, No. 3 Kiln Stack, September 16, 1993, Custom Stack Analysis Company, Alliance, OH, October 1993.

AP-42 Section 11.7
Reference 8
Report Sect. _____
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SOURCE EMISSION TESTS
AT
STARK CERAMICS, INC
EAST CANTON, OHIO
NO. 4 KILN STACK
SEPTEMBER 16, 1993



CONTENTS

BRIEF OF TESTS	1
RESULTS	1
TEST METHODS	1
PARTICULATE TEST APPARATUS	2
PARTICULATE TEST PROCEDURES	3
SULFUR DIOXIDE TEST APPARATUS	4
SULFUR DIOXIDE TEST PROCEDURES	4
FLOURIDE EMISSIONS APPARATUS AND PROCEDURES	5
MISCELLANEOUS	5
COMPILED DATA (PARTICULATE)	6
COMPILED DATA (FLOURIDE)	7-8
LOCATION OF SAMPLE PORTS AND POINTS	9
SCHEMATIC OF METHOD 5 APPARATUS	10
SCHEMATIC OF INTEGRATED SAMPLER	11
SCHEMATIC OF METHOD 6 APPARATUS	12
SCHEMATIC OF FLORIDE SAMPLING PROBE	13
APPENDIX I	
COMPUTER NOMENCLATURE	14
CALCULATION FORMULA	15
PROCESS WEIGHT SHEET	15-20
FLOURIDE ANALYSIS	21
TEST 1 RAW DATA AND CALCULATIONS (M-5)	22-23
TEST 2 " " " "	24-25
TEST 3 " " " "	26-27
TEST 1 RAW DATA AND CALCULATIONS (M-6)	28-31
TEST 2 " " " "	32-35
TEST 3 " " " "	36-39
TEST 1 RAW DATA AND CALCULATIONS (M-13B)	40-41
TEST 2 " " " "	42-43
TEST 3 " " " "	44-45
APPENDIX II	
INTENT TO TEST NOTIFICATION	46-56

SOURCE EMISSION TESTS

AT

STARK CERAMICS, INC.

EAST CANTON, OHIO

NO.3 KILN STACK

SEPTEMBER 16, 1993

Brief of Tests

Tests were performed on the stack of the brick kiln as per EPA Federal Register methods for particulate matter, sulfur dioxide and flouride emissions for stationary sources.

The tunnel kiln is a continuous operation where the green product is loaded on cars that travel through the various heat zones and discharge as a finished product at the other end. The kiln is heated with natural gas with the products of combustion and the emissions from the curring process discharging to a refractory stack via induced draft fan. The samples were extracted from the stack above the roofline.

Results (See tables 1, 2 & 3)

- A) The particulate emission rate was 5.04 pounds per hour.
- B) The sulfur dioxide emission rate was 22.6 pounds per hour.
- C) The flouride emission rate was 1.86 pounds per hour or 55.50 milligrams per cubic meter.
- D) The process weight rate was 5,842 pounds per hour.

Test Methods

A twelve point traverse was selected for the thirty five inch square brick stack as per method 1. Each point was sampled for five minutes for a total test time of 60 minutes. (See figure 1) The CO₂ and O₂ analysis were conducted by an orsat flue gas analyzer from an integrated sample taken at the test ports to provide data for method 3.

Method 5 was used to determine the particulate quantity, moisture determination by method 4 and gas velocity by method 2 using a Research Appliance Corp. "Stacksamplr".

Method 6 was used for the determination of sulfur dioxide.

Particulate Test Procedures

The probe, filter and glassware was assembled and leak tested in our lab before transporting to the job site. Three sets of equipment were used. At the job site a preliminary pitot traverse was performed to select the proper nozzle size. The nozzles were measured with an inside vernier caliper and micrometer calibrated with a one inch micrometer standard.

The first and second impingers were filled with 100 ml of distilled water and the "Drierite" columns were connected just prior to elevating the probe into position.

After leak testing the apparatus at 10 inches of mercury the probe was inserted at the first sample point to start the test.

The isokinetic sampling rates were determined using a portable desk top computer programmed to calculate the proper ΔH setting at the flue gas temperature, pressure, density and the assumed moisture along with the temperature, ΔP and an assumed ΔH of the test module. The sampling rate (ΔH) can be determined in less than 5 seconds using this technique.

The apparatus was leak tested after the test was completed at a vacuum exceeding that encountered during the test.

The moisture content was determined from the amount of condensate collected in the impingers and the difference of the tare and gross weight of the "Drierite" desiccant column. The desiccant column was weighed on an Ohaus 5 Kg electronic toploader balance to the nearest tenth gram.

The filter media was dried in a desiccator at room temperature to dryness before tare and gross weighing on a Stanton CL4D analytical balance.

The probe liner, nozzle, cyclone bypass and front half of the filter holder were washed with acetone and brush after testing and evaporated to dryness at room temperature in 500 ml beakers. The beakers were dried and tare weighed prior to sample evaporation and gross weighed after allowing the samples to desiccate to dryness. These weights were accomplished with the Stanton Balance.

The contents of the impingers were measured by weighing the three impingers and contents before and after testing. The total

moisture was determined by adding net weight of the impingers to the net weight of the drierite column.

The integrated gas sampler was started at the beginning of the method 5 test with about two cubic foot of gas collected in the tedlar bag at the end of the test. The sampling rate was maintained constant during the test by maintaining a constant reading on the rotometer.

The orsat analysis was performed on the gases contained in the bag shortly after the test was completed.

Sulfur Dioxide Test Apparatus

A schematic of the test apparatus is shown in figure 4. The gases were drawn through a heated pyrex probe with a fiberglass wad attached to the sampling end of the probe. The probe was connected to the impinger train where it first passed through a midget bubbler containing 15 ml of 80% isopropyl alcohol. A wad of glass wool was packed in the top of the bubbler to collect any acid mist carryover. The gases then passed through two midget impingers, each containing 15 ml of 3% hydrogen peroxide solution followed by an empty midget impinger to collect the carryover from the preceeding impingers. The midget bubbler and impingers were contained in an ice bath to condense acid and water vapor. After leaving the dry trap the gases passed through a "Drierite" column containing about 500 grams of "Drierite" via rubber hose to remove the remaining water vapor. The dry gas was transmitted to the module through "polyflow" tubing. In the module the gas was moved through the system by a leakless air pump connected to the rotometer to measure the gas flow rate and a Rockwell type "S" dry test meter to measure the gas volume. The dry test meter was equipped with a thermometer located in the top half of the gas meter to measure the temperature of the meter. The gas from the meter was discharged to the atmosphere.

Sulfur dioxide test procedures

Two method 6 runs of twenty minute duration comprised one test. A single point near the center of the duct using one of the method 5 sample ports was used for the sample location.

The sample train was leak checked using a calibrated vacuum gage. The vacuum gage was connected to the inlet of the sample

train and the system was brought to a ten inch mercury vacuum and held for one minute before starting the test. After the test was completed the system was purged for fifteen minutes at the same flow rate using a separate vacuum pump and rotometer. After purging the system, the contents of the midget bubbler were discarded and the contents of the midget impingers along with the washings were placed in a marked polyethylene bottle. The samples were titrated at our Alliance lab as per EPA Federal Register methods using barium perchlorate titrant and thorin indicator. The barium perchlorate was standardized against a purchased .01 normal sulfuric acid standard. The dry gas volumetric flow rate (Q-CFH) from method 5 was used to calculate the pounds per hour of sulfur dioxide emission.

Fluoride emissions test apparatus and procedures.

The apparatus was the same as the method 5 apparatus except the probe liner and nozzle were made of one piece construction as shown in figure 5. A teflon coated filter back-up and a Watman[™] No.1 filter were used in place of the fiberglass filter and fritted glass back-up plate.

The preparation of the sample train was identical to the method 5 train.

The sampling procedure was the same as the method 5 procedure.

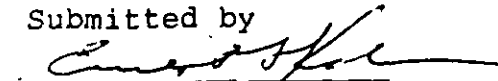
The cleanup differed from the method 5 cleanup by using distilled water instead of acetone.

The filter was placed in the contents of the probe wash combined with the impinger contents and impinger wash. The total volume of the combined sample was measured and a distilled water blank of the same volume with a filter included was transported to another lab for analysis. The analysis were performed as per method 13B.

Miscellaneous

The raw data and calculations are shown in appendix I and the Intent to Test Notification is shown in Appendix II.

Submitted by



Ernest L. Kolm

Method 13B was used for flouride emissions.

Particulate Test Apparatus (See figure 2)

The dust laden gases were passed through a pyrex lined probe and heated glass cyclone separator bypass followed by a four inch filter holder containing Gelman Type A-E fiberglass filter media. The gases leaving the filter were cooled in a series of three impingers packed in ice. The first and third impinger were the modified Greenburg-Smith type and the second one was a standard Greenburg-Smith. The First and second impingers were filled with 100 ml of distilled water with the third one used as a dry trap. After leaving the dry trap, the gases passed through a "Drierite" column containing about 500 grams of calcium sulfate (CaSO_4) desiccant to remove the remaining water vapor. The dry gas passed through the hose portion of the umbilical cord to a Research Appliance Corp. model 2343 "Stacksamplr" module. In the module the gas was moved through the system by a leakless air pump to a rockwell 175-S dry test meter. The dry test meter exhausted to a calibrated orifice to measure the flow rate of the gases passing through the sampling apparatus. A type "S" pitot tube was attached to the sheath of the heated probe to measure the velocity head of the flue gases near the tip of the probe nozzle. The orifice pressure taps and the pitot tube were connected to a Dwyer dual 10 inch combination inclined well type manometer. One half of the manometer measured the orifice differential (ΔH) and the other half measured the flue gas velocity head (ΔP).

The temperature of the flue gas was measured by a type "K" thermocouple connected to a PyroMation digital temperature indicator.

The CO_2 and O_2 were measured with a Burrell "Industro" Model B orsat from an integrated sample taken by withdrawing a constant flow rate of gas from the stack and injecting it into a Tedlar bag. This was done by drawing the gas through an in-stack filter via neoprene tubing to a condenser and condensate collector ahead of a leakless diaphragm vacuum-pressure pump. The pump discharged to the rotometer and the Tedlar bag. The apparatus is equipped with valves to by-pass the rotometer and bag when clearing the sample line as shown in figure 3.

Test	1	2	3	Average
Date	9/16/93	9/16/93	9/16/93	
Time	9:42/11:01	11:51/13:05	13:55/15:07	

FLOURIDE EMISSIONS

Grains/f3 Dry STP	0.02438	0.02395	0.02428	0.02420
Pounds/Hr.	1.8758	1.8080	1.8872	1.8570

STACK GAS CONDITIONS

Temperature -dg. F	583.0	588.0	595.0	588.7
Static Pressure-in H2O	0.00	0.00	0.00	0.00
CO2 - %	3.4	3.1	3.2	3.2
O2 - %	16.4	16.6	16.6	16.5
H2O - %	6.2	6.3	6.2	6.2
Velocity - FPS	37.92	37.40	38.74	38.02
Stack Area - sqr. ft.	8.50	8.50	8.50	8.50
Gas Flow - ACFM	19337	19073	19757	19389
Gas Flow (DSTP) CFH	537395	527140	542883	535806

SAMPLE TRAIN CONDITIONS

Pitot Delta P in H2O	.222	.214	.228
Orifice Delta P in H2O	2.77	2.73	2.96
Temp. Meter - dg. F	89	101	105
Gas Volume - CF Dry STP	48.26	48.60	49.35
Barometer - in Hg	29.08	29.08	29.08
Probe Tip Dia. - in.	0.391	0.385	0.383
Isokinetic Var. - %	91.9	97.4	97.0

STARK CERAMICS, INC.
EAST CANTON, OHIO
NO. 3 KILN STACK
COMPILED DATA

Table 2

Test	1	2	3	Average
Date	9/16/93	9/16/93	9/16/93	
Time	9:42/11:01	11:51/13:05	13:55/14:17	
PARTICULATE QUANTITY				

Grains/f3 Dry STP	0.07375	0.05182	0.06315	0.06290
Pounds/Hr.	5.8452	4.1293	5.1344	5.0363
GASEOUS EMISSIONS				

SO2 Pounds/Hr	20.75	23.82	23.26	22.61
STACK GAS CONDITIONS				

Temperature -dg. F	583.0	588.0	595.0	588.7
Static Pressure-in H2O	0.00	0.00	0.00	0.00
CO2 - %	3.4	3.1	3.2	3.2
O2 - %	16.4	16.6	16.6	16.5
H2O - %	6.7	6.2	6.4	6.4
Velocity - FPS	39.27	39.45	40.59	39.77
Stack Area - sqr. ft.	8.50	8.50	8.50	8.50
Gas Flow - ACFM	20027	20121	20702	20283
Gas Flow (DSTP) CFH	553572	556565	567866	559334
SAMPLE TRAIN CONDITIONS				

Pitot Delta P in H2O	.237	.238	.251	
Orifice Delta P in H2O	1.09	1.15	1.22	
Temp. Meter - dg. F	83	93	97	
Gas Volume - CF Dry STP	31.68	32.13	34.39	
Barometer - in Hg	29.08	29.08	29.08	
Probe Tip Dia. - in.	0.313	0.313	0.313	
Isokinetic Var. - %	91.4	92.2	96.7	

STARK CERAMICS, INC.
EAST CANTON, OHIO
NO 3 KILN STACK
COMPILED DATA

Table 1

FLOURIDE CONCENTRATION

$Cs = Ft/Vm_{stp}$ $Ft = \text{mg Flouride}, Vm_{stp} = \text{cubic meters (m}^3\text{)}$

TEST 1

48.26 Ft^3 Dry stp X 0.02831 $m^3/Ft^3 = 1.3666 m^3$
76.4 mg/1.3666 $m^3 = 55.91 \text{ mg Flouride}/m^3$

TEST 2

48.60 Ft^3 Dry stp X 0.02831 $m^3/Ft^3 = 1.3760 m^3$
75.6 mg/1.3760 $m^3 = 54.94 \text{ mg Flouride}/m^3$

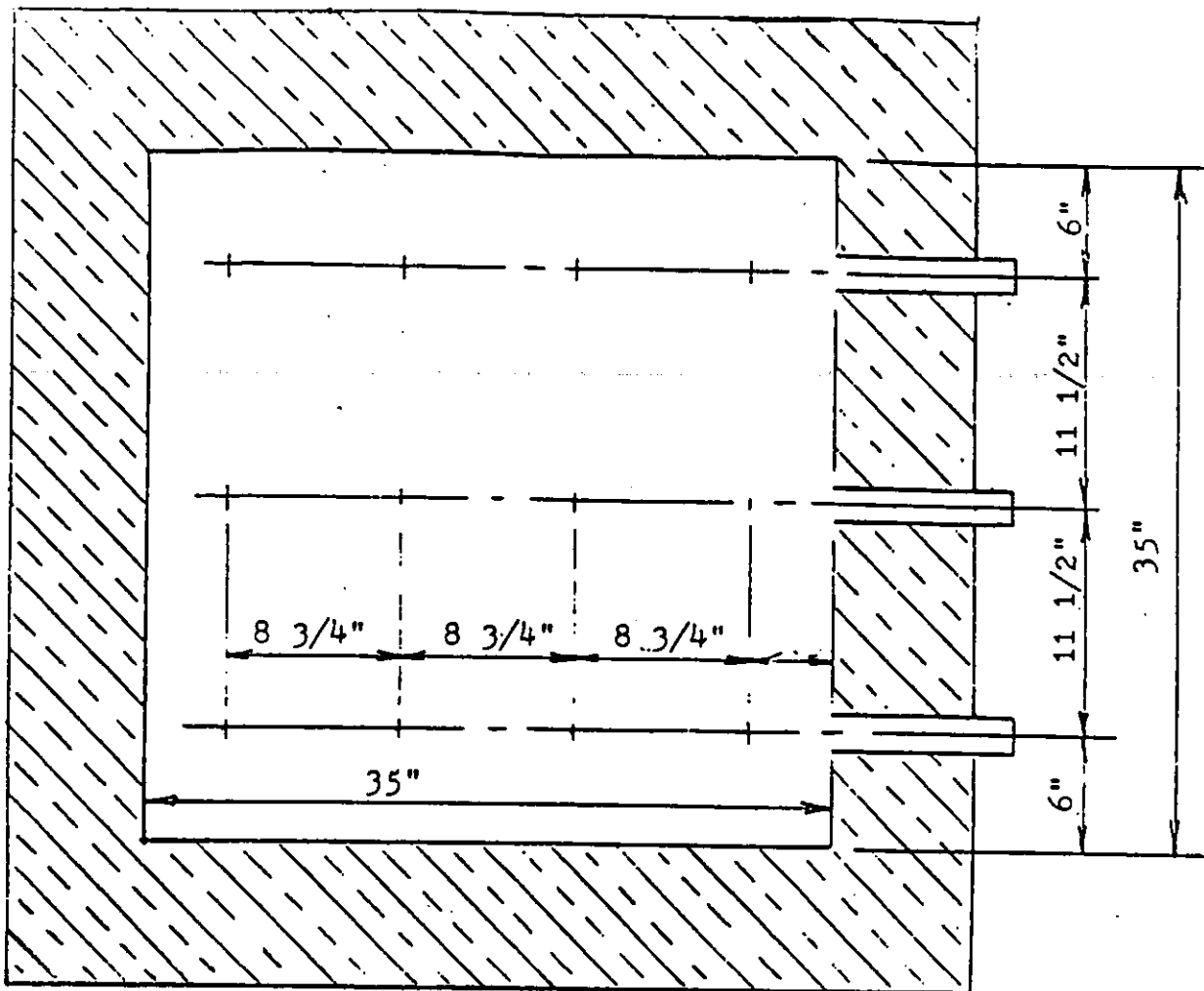
TEST 3

49.35 Ft^3 Dry stp X 0.02831 $m^3/Ft^3 = 1.3974 m^3$
77.8 mg/1.3974 $m^3 = 55.67 \text{ mg Flouride}/m^3$

AVERAGE OF 3 TESTS = 55.50 mg/m³

STARK CERAMICS, INC.
EAST CANTON, OHIO
NO. 3 KILN STACK
COMPIED DATA

TABLE 3

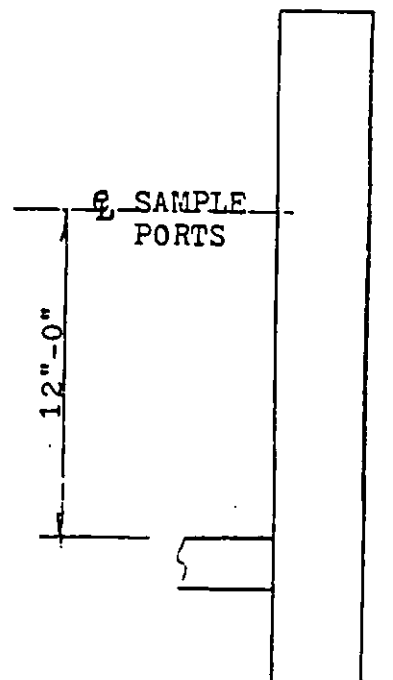


$$\text{AREA} = 8.5 f^2$$

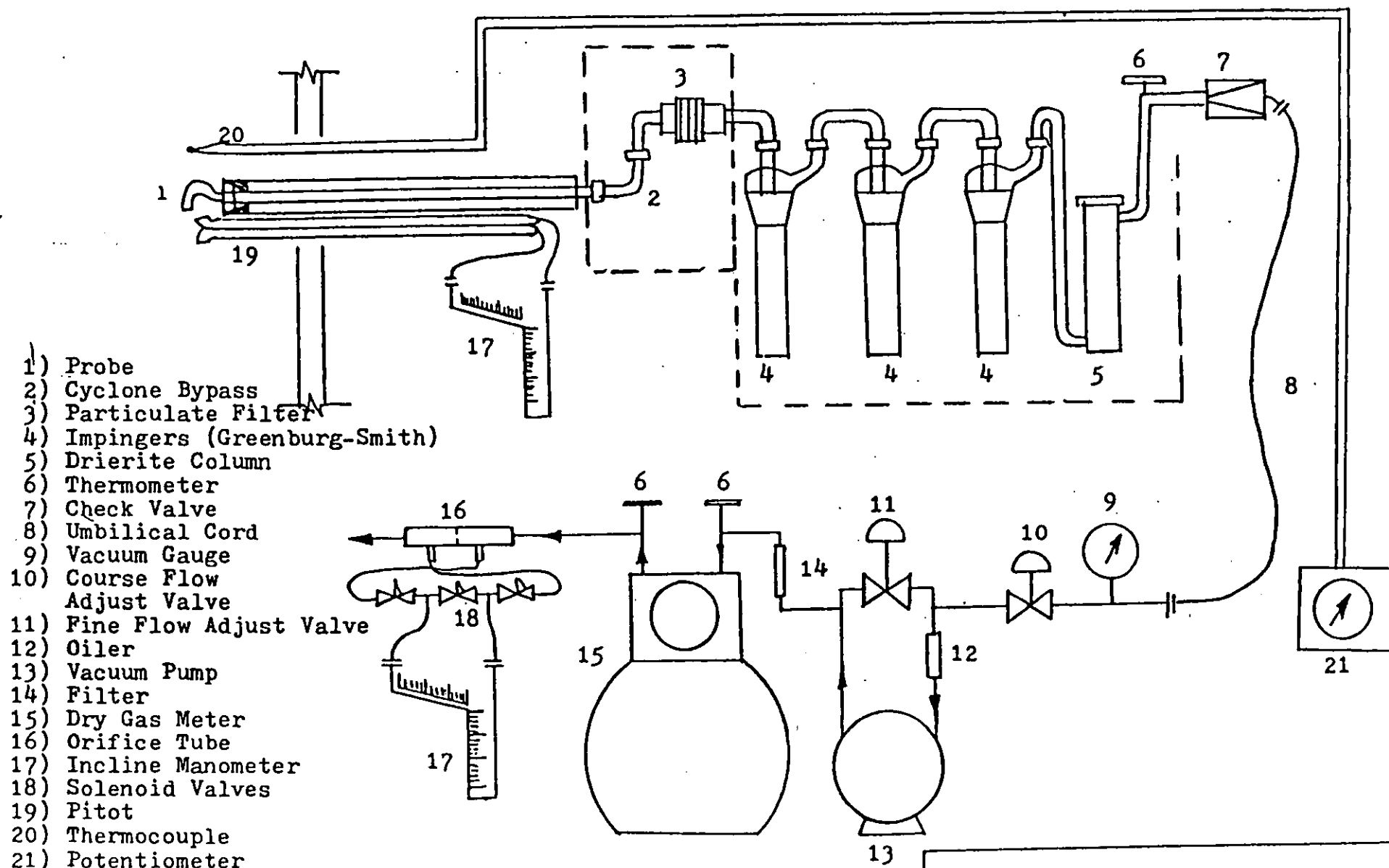
$$\text{EQUIVALENT DIA.} = 35"$$

$$\text{EQUIV. DIA. UPSTREAM} = 4.1$$

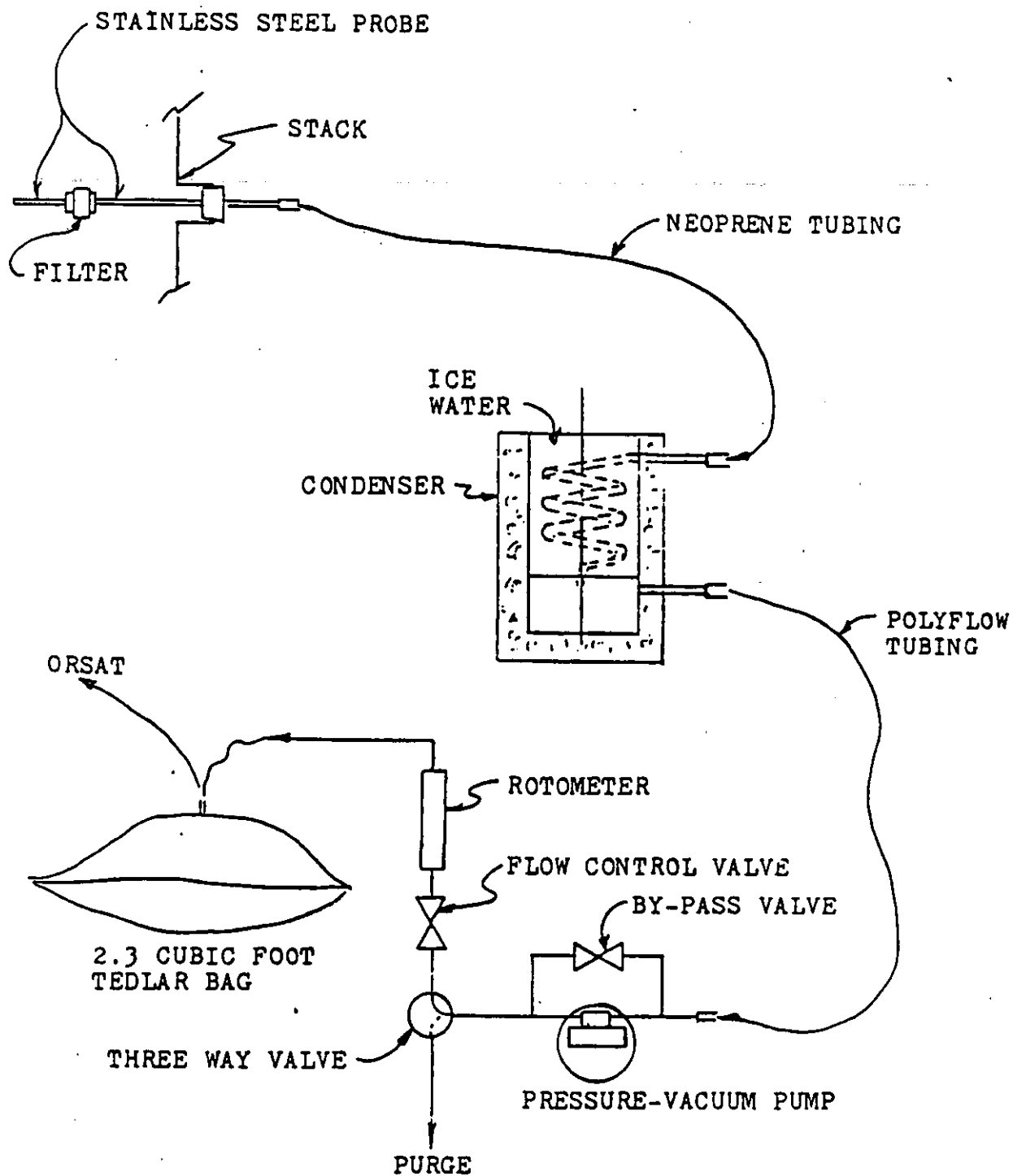
$$\text{EQUIV. DIA. DOWNSTREAM} = 2 +$$



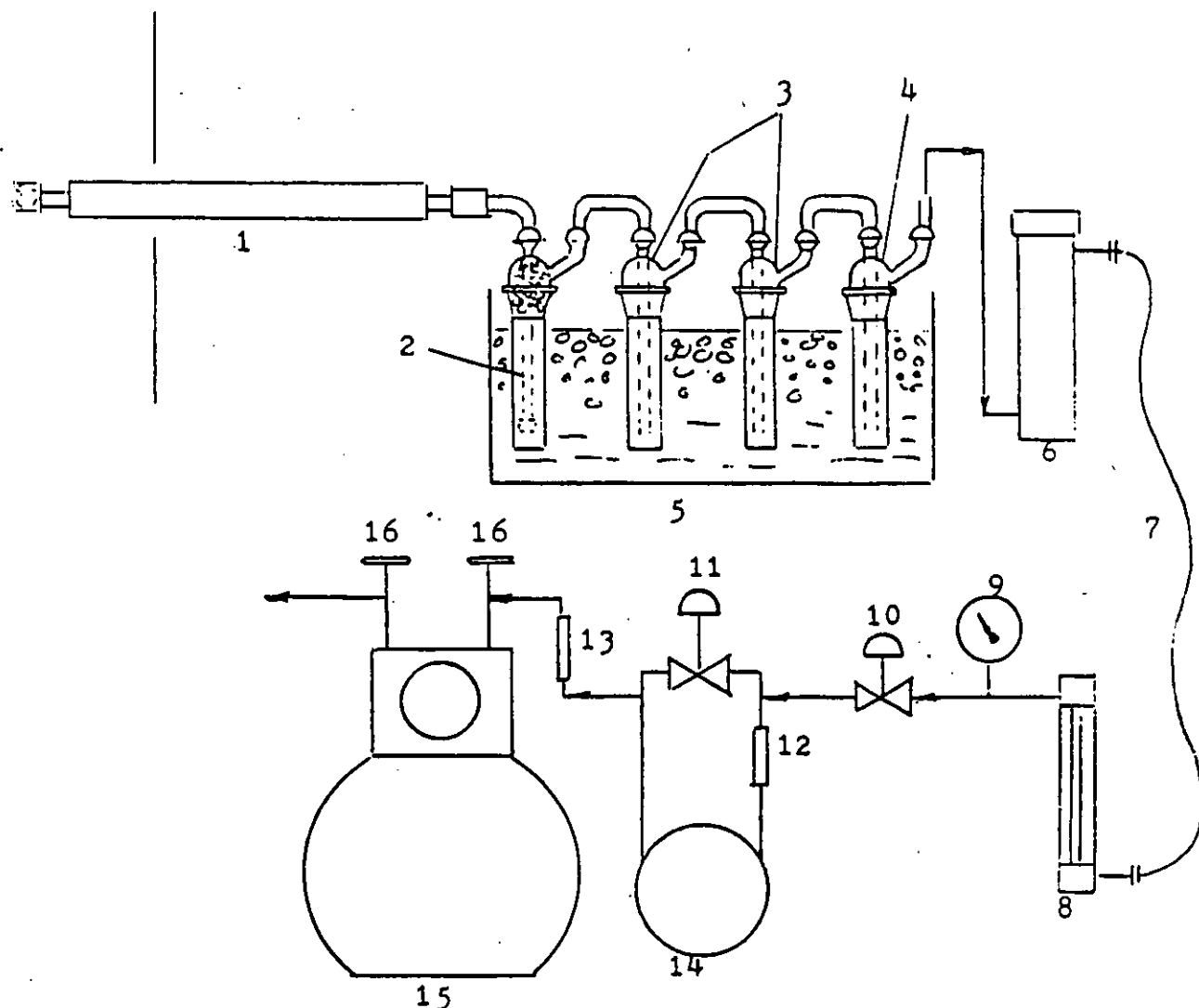
STARK CERAMICS



C.S.A. Co.
MODIFICATION OF R.A.C. SAMPLE TRA
USING DRIERITE COLUMN IN PLACE OF
IMPINGER DESICANT

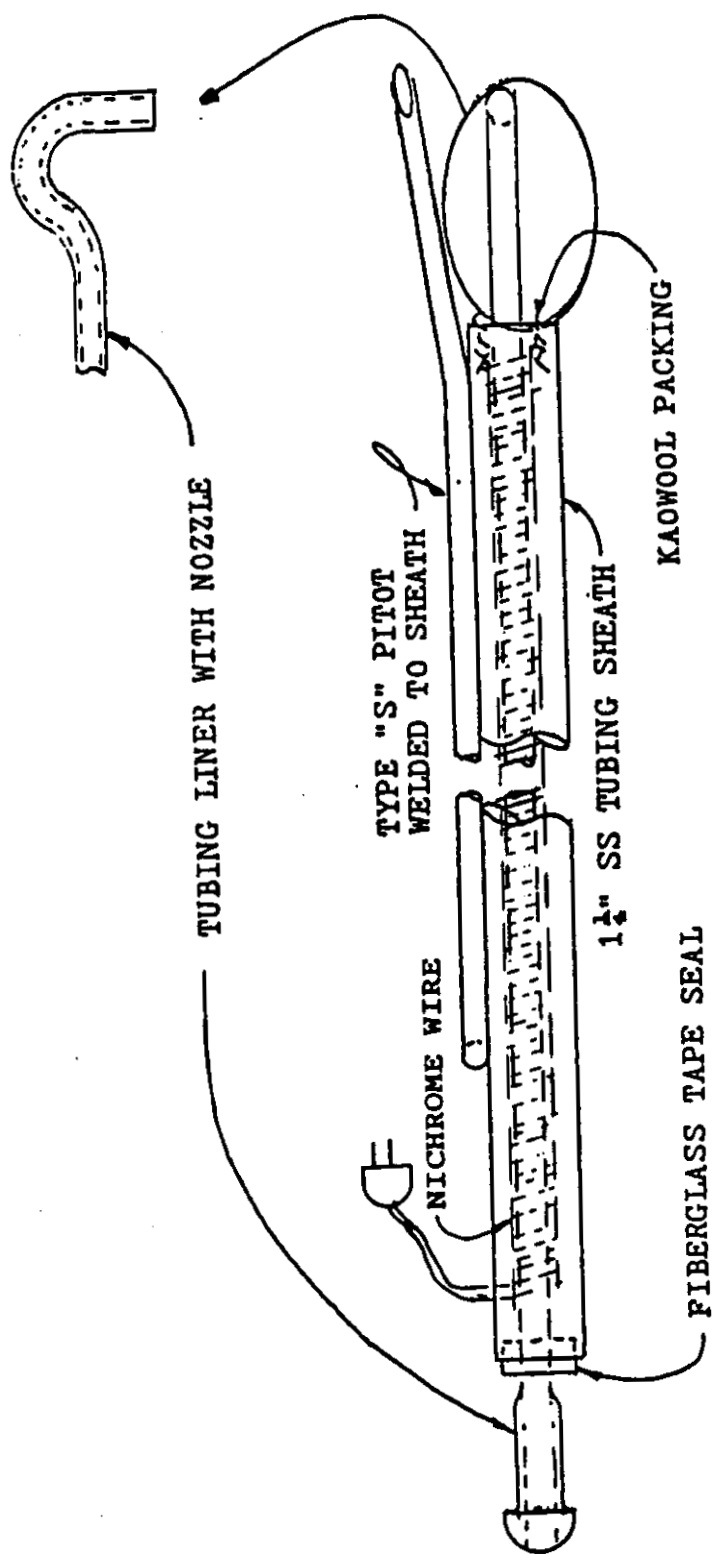


INTEGRATED SAMPLER



- 1) HEATED PYREX LINED PROBE. END PACKED WITH QUARTZ WOOL
- 2) MIDGET BUBBLER. TOP PAKED WITH GLASS WOOL
- 3) MIDGET IMPINGERS. EACH FILLED WITH 15 ml OF H_2O_2 (3%)
- 4) MIDGET IMPINGER. DRY TRAP
- 5) ICE BATH
- 6) DRIERITE COLUMN. FILLED WITH CALCIUM SULFATE DESICANT
- 7) UMBILICAL CORD
- 8) ROTAMETER
- 9) VACUUM GAGE
- 10) VALVE FOR COARSE FLOW ADJUSTMENT
- 11) VALVE FOR FINE FLOW ADJUSTMENT
- 12) OILER
- 13) OIL FILTER
- 14) VACUUM PUMP
- 15) DRY GAS TEST METER
- 16) THERMOMETER

E.P.A. METHOD 6 SO_2 SAMPLING TRAIN



HEATED ONE PIECE PROBE LINER/NOZZLE

APPENDIX I

METHOD 5

<u>SYMBOL</u>	<u>COMPUTER NOMENCLATURE</u>	<u>UNITS</u>
<u>DESCRIPTION</u>		
VM, FT ³	METER VOLUME	CUBIC FEET
Pb, IN.H2O	BAROMETER	INCHES MERCURY
ΔH, IN H2O	ORIFICE DIFFERENTIAL	INCHES WATER
PMA, IN Hg	ABSOLUTE METER PRESSURE	INCHES MERCURY
TM, F	METER TEMPERATURE	DEGREES FARENHEIT
TS, F	STACK TEMPERATURE	DEGREES FARENHEIT
PG, IN H2O	STACK STATIC PRESSURE	INCHES WATER
PSA, IN Hg	ABSOLUTE STACK PRESSURE	INCHES MERCURY
CP,	PITOT COEFFICIENT	DIMENTIONLESS
√ΔP, IN H2O	AVERAGE SQUARE ROOT OF ΔP	INCHES WATER ^{1/2}
%CO2	CARBON DIOXIDE	PERCENT
%O2	OXYGEN	PERCENT
%CO	CARBON NONOXIDE	PERCENT
%N2	NITROGEN	PERCENT
MD	MOLECULAR WEIGHT DRY	DIMENTIONLESS
VLC,ML	VOLUME OF CONDENSATE	MILLILITERS
AS, FT ²	STACK AREA	SQUARE FEET
MN, MG	TOTAL PARTICULATE CATCH WEIGHT	MILLIGRAMS
O, MIN	TEST TIME	MINUTES
DN, IN	NOZZLE DIAMETER	INCHES
AN, FT ²	NOZZLE AREA	SQUARE FEET
L, #/HR	EMISSION RATE	POUNDS PER HOUR
I, %	ISOKINETIC VARIATION	PERCENT
VMSTP, FT ³	METER VOLUME @ STANDARD TEMP. & PRESS. (DRY)	CUBIC FEET
VWSTP	METER VOLUME @ STANDARD TEMP. & PRESS. (WET)	CUBIC FEET
BWO%	MOISTURE	PERCENT
MS	MOLECULAR WEIGHT @ STACK CONDITIONS	DIMENTIONLESS
VS, FT/SEC	STACK VELOCITY	FEET PER SECOND
QS, FT ³ /HR	STACK GAS FLOW	CUBIC FEET PER HOUR
CS, #/SCF	EMISSION RATE	LB./DRY F3 @ STP
C"S,GR/SCF	EMISSION RATE	GRAINS/DRY F3 @ STP

CSA CO.
PARTICULATE CALCULATIONS FOR CFR METHOD 5

Dry Molecular Weight lb/lb-mole

$$M_d = 0.44 (\%CO_2) + 0.32 (\%O_2) + 0.28 (\%N_2 + \%CO)$$

Dry Gas Volume (standard conditions), ft³

$$V_{mstd} = 17.64 V_m Y (P_{bar} + \Delta H/13.6) / T_m$$

Volume of Water (standard conditions), ft³

$$V_{wstd} = 0.04707 V_{lc}$$

Moisture Content (proportion by volume)

$$B_{wo} = V_{wstd} / (V_{mstd} + V_{wstd})$$

Molecular Weight of Stack Gas (wet basis), lb/lb-mole

$$M_s = M_d(1 - B_{wo}) + 18 B_{wo}$$

Stack Gas Velocity ft/sec

$$V_s = 85.49 C_p \sqrt{\Delta P_{avg}} \sqrt{T_s(avg)} / P_s M_s$$

Stack Gas Volumetric Flow Rate (dry standard conditions), ft³/hr

$$Q_s = 3600(1 - B_{wo}) V_s A_s (528 / T_{savg}) (P_{sa} / 29.92)$$

Concentration (particulate in gas), lb/f³_{std dry}

$$C_s = 2.205 \times 10^{-6} M_n / V_{mstd}$$

Load (particulate in stack gas), lb/hour

$$L = Q_s C_s$$

Concentration (particulate in gas), grains/f³_{std dry}

$$C''_s = 0.0154 M_n V_{mstd}$$

Isokinetic Variation, %

$$I = 100 T_s [0.002669 V_{lc} + (V_m Y / T_m) (P_{bar} + \Delta H/13.6)] / 600 V_s P_s A_n$$

Dry wt in # 156

		<u>Shade</u>	<u>Shape</u>	<u>Pcs.</u>	<u>(4)</u>		
142	FTX	516	8WC80D	220	-	9,042	
(5)		333	8W5	40	-	1,004	(10,046) ✓
370	FTX	516	8WC80D	280		11,508	(11,508) ✓
262	FTX	516	8WC80D	244		10,028.4	
		904	8W54L	17		311.1	
(3)		904	8W54R	4		13.2	10,472 ✓
		815	S-453	3	2204	60.0	
264	FTX	516	8WC80D	244		10,028.4	10,028 ✓
(2)		815	S-453	36			
249	FTX	516	8WC80D	232		9,535.2	
(1)		333	8W5	29		127.9	10,324 ✓
		815	S-453	3	2205	60.0	
282	FTX	516	8WC80D	244		10,028.4	
		815	S-453	36	220	720.0	10,748 ✓
285	FTX	516	8WC80D	41		1,685.1	
		1022	412-4	1188	x10.9	12,949.2	15,701
		904	6TT20D-K	84	12.7	1066.8	

	<u>made</u>	<u>Shape</u>	<u>Pcs</u> (3)	
#357 (14)	195	8WCA	880	(13,200)
#304 (13)	195	8WCA	880	(13,200)
#275 (12)	127	8WCA	880	(13,200)
#272 (11)	127	8WCA	880	(13,200)
#372 (10)	195 127	8WCA 8WCA	7 873	105 (13,200) 13,295
#265 (9)	127 FTX 516	8WCA 8WC80D	472 112	7,080 (11683.3) 4,603.2
#330 (8)	FTX 516 333	8WC80D 8W5	220 40	9,042 (10,046) 1,004
#346 (7)	FTX 516 333	8WC80D 8W5	220 40	9,042 (10,046) 1,004
#257 (6)	FTX 516 333	8WC80D 8W5	220 40	9,042 (10,046) 1,004

Page 3 Dry wt
in lbs

Rock	Shape	Pcs	(2)		
281	564	8WCV80BT	134	C44.4	5950
(23)	904	6TT20D-K	30	@12.7	381
	177	8WCV80BT	131	@44.4	58166
#278	*177	8WCV80BT	256		11366
(24)	904	6TT20D-K	30		381
(25)					11,747
#359	177	8WCV80BT	280		12,432
#239	*177	8WCV80BT	250		11,100
(26)	904	8W54R	20		366
					11,466
#352	177	8WCV80BT	20		888
(19)	195	8WCA	796		11,940
					12,828
#353	195	8WCA	880		13,200
(18)					
#259	195	8WCA	880		13,200
(17)					
#116	195	8WCA	880		13,200
(16)					
#324	195	8WCA	880		13,200
(15)					

	Size	Shape	Pcs.	(1)	
343	904	8WCBK	466	10,311.2	
(31)	031	4W83-S484	12 @ 31.5	378	
	516	8W24AL	20	288	(11,524)
	516	4T58NR	6 @ 7.7	47	
*335	564	8WCV80BT	244	10,834	
(30)	031	4W83-S483	18 @ 31.5	378	11,212
*289	564	8WCV80BT	232	10,301	
(29)	516	4T58NR	4 } @ 7.8	109	10,786
	516	4T58NL	10 }		
	516	8W24CR	16	316	
*277	564	8WCV80BT	232	10,301	
(28)	516	8W24CR	6	141	10,955
	031	4W83-S484	7 @ 31.5	220	
	516	8W54L	16	292.8	
*276	564	8WCV80BT	244	10,834	
(27)	516	6T54R	19 @ 10.8	205	11,240
	516	8W54L	8	146.4	
	904	8W54L	3	54.9	
*371	564	8WCV80BT	256	11,366	
(26)	904	8W54L	16	292.8	11,659
*302	904	8WCBK	425	9,860	
(25)	904	6TT20D-K	46 @ 12.7	584	12,975
	564	8WCV80BT	57	2531	
*320	564	8WCV80BT	250	11,100	11,760
(24)	904	6TT20D-K	52	660	

	<u>Shade</u>	<u>Shape</u>	<u>Pcs.</u>			
53	904	8WCBK	460	@13.2	10811	
(36)	031	4W83-5484	30	@31.5	945	11756

234	904	8WCBK	516		11971	
(35)	904	6TT20D-K	78	@12.7	991	12962

338	904	8WCBK	516		11971	
34	904	6TT20D-K	74		940	12911

354	904	8WCBK	516		11971	
33	904	6TT20D-K	64		813	12784

371	904	8WCBK	474		10997	
(34)	772	5W8	40	@13.5	540	11,893
	904	6TT20D-K	28		356	

Total for 37 KCs = 438783

Arg Dry wt per KC = 11,859 lbs

5,842 #/HR

or 5.93 TONS

AMERICAN ANALYTICAL LABORATORIES, INC.

Pg. 1

INDUSTRIAL HYGIENE AND ENVIRONMENTAL SCIENCES

840 S. MAIN STREET
AKRON, OHIO 44311-1516
(216) 535-1300

WORK ORDER #: 93-09-229

SAMPLES RECEIVED: 09/17/93
ANALYSIS REPORTED: 09/27/93

REPORT ISSUED TO:

Ernie Kolm
CSA Company
Custom Stack Analysis
P.O. BOX 3750
Alliance, Ohio 44601

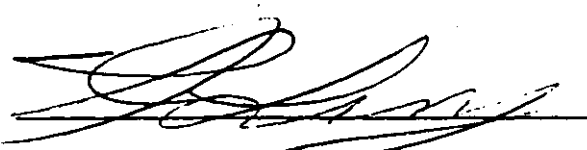
WORK ID: Flouride Analysis

SAMPLED BY: Client
SAMPLE TYPE: Water/Air

SAMPLE ANALYSIS REPORT

SAMPLE ID AAL LAB #	DATE COLLECTED		METHOD(S)
	PARAMETER(S)	RESULT(S) UNITS	
SC Test 1 9309229-01	09/16/93 Total Flouride Emmissions	76.4 mg/Sample	EPA 13B
SC Test 2 9309229-02	09/16/93 Total Flouride Emmissions	75.6 mg/Sample	EPA 13B
SC Test 3 9309229-03	09/16/93 Total Flouride Emmissions	77.8 mg/Sample	EPA 13B
SC Blank 1 9309229-04	09/16/93 Total Flouride Emmissions	<0.5 mg/Sample	EPA 13B
SC Blank 2 9309229-05	09/16/93 Total Flouride Emmissions	<0.5 mg/Sample	EPA 13B
SC Blank 3 9309229-06	09/16/93 Total Flouride Emmissions	<0.5 mg/Sample	EPA 13B

ANALYSIS REVIEWED AND APPROVED BY



CSA CO. DATA SHEET (METHOD 5)

TEST NO. 1 PLANT STARK CERAMICS											DATE 9/16/93
LOCATION NE 3 KILN STACK											BY GT M/M
BAROMETER (P _B) 29.08 AMBIENT TEMP											ASSUMED MOISTURE "Y" 1.001
MODULE NO. 2031 PROBE NO. 1-2 FILTER NO. 12 HEATER NO. 3 NOZZLE DIA.											
TEST POINT	METER VOLUME (Vm)	METER TEMP (Tm) IN OUT		COND TEMP °F	FILTER HEATER TEMP °F	STACK TEMP °F (Ts)	STACK PRESS. (Ps)	VACUUM "Hg	ORIFICE "H ₂ O (ΔH) (Pm)	VEL HEAD (ΔP)	TIME
1	222.1	66	69	60	252	586	0	3-	1.3	.28	9.42
1	225.0	70	67	52	251	576	0	3-	1.3	.28	47
2	227.8	79	70	52	251	577	0	3-	1.0	.24	52
2	230.8	84	72	48	252	584	0	3+	1.6	.35	
4	234.7	88	74	44	250	587	0	4-	2.1	.45	10.02
1	234.7	88	78	49	251	583	0	3-	.9	.19	10.10
1	230.7	88	79	54	252	582		3-	.9	.19	.19
2	240.0	90	80	55	250	580		3-	.9	.19	.24
3	242.6	91	82	58	250	581		3-	.8	.18	.29
4	245.6	91	84	58	251	583		3-	1.4	.29	.34
1	245.6	89	85	57	250	588		3+	.8	.17	.41
1	248.2	94	86	56	252	583		3-	.8	.17	.46
2	250.5	95	87	58	251	582		3-	.7	.14	.51
3	252.9	97	88	58	250	582		3+	.8	.17	.56
4	255.6	98	88	58	251	589		3+	1.1	.28	11.01

E.P.A. METHOD 5

DATA INPUT

Pb, In Hg	Barometer -----	29.08
VM, ft3	Meter Volume -----	33.5
ΔH, In H2O	Orifice Differential -----	1.09
PG, In H2O	Stack Static Pressure -----	0
Tm, F	Meter Temperature -----	83
Ts, F	Stack Temperature -----	583
CP	Pitot Coefficient -----	.84
ΔP, In H2O	Average Square Root Of Delta P -----	.4870
% CO2	Carbon Dioxide -----	3.4
% O2	Oxygen -----	16.4
% N	Nitrogen -----	80.2
VCL, Ml	Volume Of Condensate -----	48.5
AS, ft2	Stack Area -----	8.5
MN, Mg	Total Particulate Catch Weight -----	151.7
Ø, Min	Test Time -----	60
DN, In	Diameter Of Nozzle -----	.313

RESULTS

Pma, In Hg	Absolute Meter Pressure -----	29.16
PSA, In Hg	Absolute Stack Pressure -----	29.08
MD	Molecular Weight Dry Gas -----	29.20
MS	Molecular Weight @ Stack Conditions -----	28.45
VMSTP, ft3	VM Standard Temp. & Press. Dry -----	31.68
VWSTP, ft3	VM Standard Temp. & Press. Wet -----	2.28
BWO, %	Moisture -----	6.72
I, %	Isokeimetic Variation -----	91.43
VS, ft/Sec	Stack Velocity -----	39.27
QA, ASCFM	Stack Gas Flow (Actual) -----	20027
QS, ft3/HR	Stack Gas Flow (Dry STP) -----	553573
L, #/Hr	Emission Rate -----	5.85
CS, #/SCF	Particulate Emission -----	1.055898E-05
CSS, Grn/SCF	Particulate Emission -----	0.07375

STARK CERAMICS
EAST CANTON OH
NO 1 KILN STACK
9/16/93
TEST 1

CSA CO. DATA SHEET (METHOD 5)

[illegible]

E.P.A. METHOD 5

DATA INPUT

Pb.In Hg	Barometer -----	29.08
VM.ft3	Meter Volume -----	34.6
ΔH.In H2O	Orifice Differential -----	1.15
PG.In H2O	Stack Static Pressure -----	0
Tm.F	Meter Temperture -----	93
Ts.F	Stack Temperture -----	588
CP	Pitot Coefficient -----	.84
ΔP.In H2O	Average Square Root Of Delta P -----	.4883
% CO2	Carbon Dioxide -----	3.1
% O2	Oxygen -----	16.6
% N	Nitrogen -----	80.3
VCL.M1	Volume Of Condensate -----	45.2
AS.ft2	Stack Area -----	8.5
MN.Mg	Total Particulate Catch Weight -----	108.1
Ø.Min	Test Time -----	60
DN.In	Diameter Of Nozzle -----	.313

RESULTS

Pma.In Hg	Absolute Meter Pressure -----	29.16
PSA.In Hg	Absolute Stack Pressure -----	29.08
MD	Molecular Weight Dry Gas -----	29.16
MS	Molecular Weight @ Stack Conditions -----	28.47
VMSTP.ft3	VM Standard Temp.& Press. Dry -----	32.13
VWSTP.ft3	VM Standard Temp.& Press. Wet -----	2.13
BWO.%	Moisture -----	6.21
I.%	Isokeimetic Variation -----	92.23
VS.ft/Sec	Stack Velocity -----	39.45
QA,ASCFM	Stack Gas Flow (Actual) -----	20122
QS.ft3/HR	Stack Gas Flow (Dry STP) -----	556565
L.#/Hr	Emission Rate -----	4.13
CS,#/SCF	Particulate Emission -----	7.419186E-06
C'S.Grns/SCF	Particulate Emission -----	0.05182

STARK CERAMICS
EAST CANTON OH
NO 1 KILN STACK
9/16/93
TEST 2

CSA CO. DATA SHEET (METHOD 5)

TEST NO. 3 PLANT Stack Ceramics							DATE 9/16/07				
LOCATION No 3 Kiln Stack							BY Am				
BAROMETER (P _B) 21.08 AMBIENT TEMP ASSUMED MOISTURE "Y" 1.001											
MODULE NO. F-021 PROBE NO. 6 FILTER NO. 9 HEATER NO. 6 NOZZLE DIA.											
TEST POINT	METER VOLUME (Vm)	METER TEMP (Tm) IN	METER TEMP (Tm) OUT	COND TEMP °F	FILTER HEATER TEMP °F	STACK TEMP °F (Ts)	STACK PRESS. (Ps)	VACUUM "Hg	ORIFICE "H ₂ O ΔH (Rm)	VEL HEAD ΔP	TIME
1	290.7	89	89	58	250	591	0	2-	1.0	.20	13:55
1	293.5	91	87	60	252	594		3+	1.0	.20	14:00
2	296.7	96	87	60	251	592		3+	1.4	.30	05
2	300.2	100	90	60	252	596		S+	1.9	.38	10
4	304.0	102	91	60	250	598		S+	1.9	.38	15
1	304.0	96	92	61	252	596		3-	1.0	.21	14:12
1	306.8	100	93	61	250	595		3-	1.0	.21	27
2	309.5	102	94	62	252	593		3-	.9	.19	32
2	312.4	104	94	62	251	593		3-	1.1	.22	37
4	315.5	105	95	62	252	595		3+	1.3	.25	42
1	315.5	100	95	61	251	599		3+	1.1	.22	14:47
1	318.3	103	96	58	250	599		3+	1.1	.22	52
1	321.7	105	97	58	252	593		3-	1.2	.24	57
2	324.7	106	97	58	250	595		3-	1.1	.21	12
4	328.0	107	98	58	251	596		4-	1.3	.25	17
	290.7										
(Vm)	37.3	100.5	93.0								
AVERAGE		97		****	*****	595	0	MAX 5	1.22	.251	60 MINUTE
INTEGRATED GAS SAMPLE				AVG	N ₂ =80.2	CONDENSATE COLLECTED					
CO ₂		3.2		3.2	DRIERITE	FINAL 992.1	INITIAL 984.7	TOTAL 7.4			
O ₂		16.6		16.6	IMPINGER	242.3	200	42.3			
FILTER WT		PROBE WASH WT		IMPINGER WASH WT		GRAND TOTAL 49.7					
GROSS .7838	169.4020						LEAK RATE @ 11 "Hg ± .02				
TARE .6824	169.3624						STACK AREA (As) f _s				
NET .1014	.0396						AVG SQ RT Δ P .5007				
TOTAL PARTICULAT WT (Mn) 141.0 Mg											

E.P.A. METHOD 5

DATA INPUT

Pb, In Hg	Barometer -----	29.08
VM, ft3	Meter Volume -----	37.3
ΔH, In H2O	Orifice Differential -----	1.22
PG, In H2O	Stack Static Pressure -----	0
Tm, F	Meter Temperature -----	97
Ts, F	Stack Temperature -----	595
CP	Pitot Coefficient -----	.84
ΔP, In H2O	Average Square Root Of Delta P -----	.5007
% CO2	Carbon Dioxide -----	3.2
% O2	Oxygen -----	16.6
% N	Nitrogen -----	80.2
VCL, Ml	Volume Of Condensate -----	49.7
AS, ft2	Stack Area -----	8.5
MN, Mg	Total Particulate Catch Weight -----	141.0
Ø, Min	Test Time -----	60
DN, In	Diameter Of Nozzle -----	.313

RESULTS

Pma, In Hg	Absolute Meter Pressure -----	29.17
PSA, In Hg	Absolute Stack Pressure -----	29.08
MD	Molecular Weight Dry Gas -----	29.18
MS	Molecular Weight @ Stack Conditions -----	28.46
VMSTP, ft3	VM Standard Temp. & Press. Dry -----	34.39
VWSTP, ft3	VM Standard Temp. & Press. Wet -----	2.34
BWO, %	Moisture -----	6.37
I, %	Isokeimetic Variation -----	96.75
VS, ft/Sec	Stack Velocity -----	40.59
QA, ASCFM	Stack Gas Flow (Actual) -----	20702
QS, ft3/HR	Stack Gas Flow (Dry STP) -----	567866
L, #/Hr	Emission Rate -----	5.13
CS, #/SCF	Particulate Emission -----	9.041635E-06
CSS, Grn/SCF	Particulate Emission -----	0.06315

STARK CERAMICS
EAST CANTON OH
NO 1 KILN STACK
9/16/93
TEST 3

CSA CO.

EPA METHOD 6 (SULFUR DIOXIDE) DATA AND CALCULATION SHEET

TEST NO 1A PLANT JARK CERAMICS DATE 9/16/93
 LOCATION NO KILN STACK BY DR G AMP

Vm	Tmi	ROTOMETER	TIME
162.678	60	4	10:43
162.938	66		48
163.198	69		53
163.457	72		58
163.717	74		11:03
Vm 1.039	avg 68		

Pb 29.08

20 min

$$Vm_{std} = K_1 Y(Vm)(Pb)/(Tm)$$

$$Vm_{std} = 17.64 (.948)(1.039)(29.08)/(460 + 68) = 1.007$$

	1st TITRATION	2nd TITRATION	3rd TITRATION	AVERAGE	V _b TITRATION
Vt	11.4	11.4		11.4	0

Cso₂ = Concentration of sulfur dioxide - lb/dry standard cubic foot
 N = Normality of barium perchlorate titrant
 Pb = Barometric pressure - inches Hg
 Tm = Temperature of meter - °F
 Va = Volume of sample aliquot - ml
 Vm = Volume of gas at meter temperature and pressure - f³
 Vm_{std} = Vm corrected to standard temperature and pressure - f³
 Vsoln = Total volume of sample solution - ml
 Vb = Volume of titrant used for blank - ml
 Y = dry gas meter calibration factor - dimensionless
 K₁ = 17.64
 K₂ = 7.061 X 10⁻⁶

METHOD 6

N= NORMALITY OF BARIUM PERCHLORATE TITRATE
 VT= VOLUME OF TITRANT, ML
 VTB= VOLUME OF TITRANT FOR BLANK, ML
 VA= VOLUME OF ALQUOT SAMPLE, ML
 VM= VOLUME OF GAS SAMPLE FROM METHOD 5, VM(DSCF)
 VSOLN= TOTAL VOLUME OF SAMPLE SOLUTION, ML
 QS= DSCF/HOUR FROM METHOD 5 CALCULATION
 DSCF= DRY STANDARD CUBIC FOOT
 MM= 1.000.000

WHEN MMBTU/HR = 1, THEN LB/MMBTU = LB/HR

SULFUR DIOXIDE CONCENTRATION = SO₂ LB/DSCF = S2

N= .0096
 VT= 11.4
 VTB= 0
 VA= 20
 VM= 1.007
 VSOLN= 100
 QS= 553572
 MMBTU/HR= 1

$S2 = (.00007061 \times N \times (VT - VTB) \times VSOLN) / (VM \times VA) = 3.836922E-05 \text{ LB SO}_2/\text{DSCF}$

$SO \text{ PPM} = 6.037 \times MM \times S2 = 231.635$

$SO_2 \text{ LB/MMBTU} = QS \times S2 / \text{MMBTU/HR} = 21.24012 = \text{LB/HR}$

STARK CERAMICS
 EAST CANTON OH
 NO 1 KILN STACK
 9/16/93
 TEST 1A

CSA CO.

EPA METHOD 6 (SULFUR DIOXIDE) DATA AND CALCULATION SHEET

TEST NO 1B PLANT STARKE CERAMICS DATE 9/16/93
 LOCATION NR KILN STACK BY LOKAL

Vm	Tmi	ROTOMETER	TIME
163.717	72	3.5	9:42
163.949	78		
164.181	81		
164.413	83		
164.645	85		
Vm .928	avg 80		

Pb 29.08

$$Vm_{std} = K_1 Y(Vm)(Pb)/(Tm)$$

$$Vm_{std} = 17.64 (.998) (.928) (29.08) / (460 + 80) = .880$$

	1st TITRATION	2nd TITRATION	3rd TITRATION	AVERAGE	V _b TITRATION
Vt	9.5	9.4	9.5	9.5	0

$$\frac{1A + 1B}{2} = \text{Ave TEST 1}$$

$$\frac{21.24 + 20.25}{2} = 20.75 \text{ \#/HR}$$

Cso₂ = Concentration of sulfur dioxide - lb/dry standard cubic foot

N = Normality of barium perchlorate titrant

Pb = Barometric pressure - inches Hg

Tm = Temperature of meter - °F

Va = Volume of sample aliquot - ml

Vm = Volume of gas at meter temperature and pressure - f³

Vm_{std} = Vm corrected to standard temperature and pressure - f³

Vsoln = Total volume of sample solution - ml

Vb = Volume of titrant used for blank - ml

Y = dry gas meter calibration factor - dimensionless

K₁ = 17.64

K₂ = 7.061 X 10⁻⁶

METHOD 6

N= NORMALITY OF BARIUM PERCHLORATE TITRATE
 VT= VOLUME OF TITRANT, ML
 VTB= VOLUME OF TITRANT FOR BLANK, ML
 VA= VOLUME OF ALQUOT SAMPLE, ML
 VM= VOLUME OF GAS SAMPLE FROM METHOD 5, VM(DSCF)
 VSOLN= TOTAL VOLUME OF SAMPLE SOLUTION, ML
 QS= DSCF/HOUR FROM METHOD 5 CALCULATION
 DSCF= DRY STANDARD CUBIC FOOT
 MM= 1.000,000
 WHEN MMBTU/HR = 1, THEN LB/MMBTU = LB/HR

SULFUR DIOXIDE CONCENTRATION = SO_2 LB/DSCF = S2

N= .0096
 VT= 9.5
 VTB= 0
 VA= 20
 VM= .88
 VSOLN= 100
 QS= 553572
 MMBTU/HR= 1

$S2 = (.00007061 \times N \times (VT - VTB) \times VSOLN) / (VM \times VA) = 3.658883E-05 \text{ LB SO}_2/\text{DSCF}$

$\text{SO PPM} = 6.037 \times MM \times S2 = 220.8867$

$\text{SO}_2 \text{ LB/MMBTU} = QS \times S2 / \text{MMBTU/HR} = 20.25455 = \text{LB/HR}$

STARK CERAMICS
 EAST CANTON OH
 NO 1 KILN STACK
 9/16/93
 TEST 1B

CSA CO.

EPA METHOD 6 (SULFUR DIOXIDE) DATA AND CALCULATION SHEET

TEST NO 2A PLANT STARK CERAMICS DATE 9/16/91
 LOCATION NE KILN STACK BY LEE ANN KIP

Vm	Tmi	ROTOMETER	TIME
164.645	76	3.5	11:51
164.855	77		56
165.064	78		12:01
165.773	81		06
165.483	84		11
Vm .838	avg 79		

Pb 29.08

$$Vm_{std} = K_1 Y(Vm)(Pb)/(Tm)$$

$$Vm_{std} = 17.64 (.998) (.838) (29.08) / (460 + 79) = .796$$

	1st TITRATION	2nd TITRATION	3rd TITRATION	AVERAGE	V _b TITRATION
Vt	10.5	10.5		10.5	0

Cso₂ = Concentration of sulfur dioxide - lb/dry standard cubic foot
 N = Normality of barium perchlorate titrant
 Pb = Barometric pressure - inches Hg
 Tm = Temperature of meter - °F
 Va = Volume of sample aliquot - ml
 Vm = Volume of gas at meter temperature and pressure - f³
 Vm_{std} = Vm corrected to standard temperature and pressure - f³
 Vsoln = Total volume of sample solution - ml
 Vb = Volume of titrant used for blank - ml
 Y = dry gas meter calibration factor - dimensionless
 K₁ = 17.64
 K₂ = 7.061 X 10⁻⁶

METHOD 6

N= NORMALITY OF BARIUM PERCHLORATE TITRATE
 VT= VOLUME OF TITRANT, ML
 VTB= VOLUME OF TITRANT FOR BLANK, ML
 VA= VOLUME OF ALQUOT SAMPLE, ML
 VM= VOLUME OF GAS SAMPLE FROM METHOD 5, VM(DSCF)
 VSOLN= TOTAL VOLUME OF SAMPLE SOLUTION, ML
 QS= DSCF/HOUR FROM METHOD 5 CALCULATION
 DSCF= DRY STANDARD CUBIC FOOT
 MM= 1,000,000

WHEN MMBTU/HR = 1, THEN LB/MMBTU = LB/HR

SULFUR DIOXIDE CONCENTRATION = SO₂ LB/DSCF = S2

N= .0096
 VT= 10.5
 VTB= 0
 VA= 20
 VM= .796
 VSOLN= 100
 QS= 556565
 MMBTU/HR= 1

$S2 = (.00007061 \times N \times (VT - VTB) \times VSOLN) / (VM \times VA) = 4.470785E-05 \text{ LB SO}_2/\text{DSCF}$

$SO \text{ PPM} = 6.037 \times MM \times S2 = 269.9013$

$SO_2 \text{ LB/MMBTU} = QS \times S2 / \text{MMBTU/HR} = 24.88282 = \text{LB/HR}$

STARK CERAMICS
 EAST CANTON OH
 NO 1 KILN STACK
 9/16/93
 TEST 2A

CSA CO.

EPA METHOD 6 (SULFUR DIOXIDE) DATA AND CALCULATION SHEET

TEST NO 2 B PLANT STARK CERAMICS DATE 9/16/91
 LOCATION US KILN STACK BY DAK AH GIMP

Vm	Tmi	ROTOMETER	TIME
165.483	83	3.5	12:20
165.713	85		23
165.944	87		30
166.174	89		35
166.404	90		12:40
Vm .921	avg 87		

Pb 29.08

$$Vm_{std} = K_1 Y(Vm)(Pb)/(Tm)$$

$$Vm_{std} = 17.64 (.994) (.921) (29.08) / (460 + 87) = .862$$

	1st TITRATION	2nd TITRATION	3rd TITRATION	AVERAGE	V _b TITRATION
Vt	10.4	10.5	10.4	10.4	0

Cso₂ = Concentration of sulfur dioxide - lb/dry standard cubic foot
 N = Normality of barium perchlorate titrant
 Pb = Barometric pressure - inches Hg
 Tm = Temperature of meter - °F
 Va = Volume of sample aliquot - ml
 Vm = Volume of gas at meter temperature and pressure - f³
 Vm_{std} = Vm corrected to standard temperature and pressure - f³
 Vsoln = Total volume of sample solution - ml
 Vb = Volume of titrant used for blank - ml
 Y = dry gas meter calibration factor - dimensionless
 K₁ = 17.64
 K₂ = 7.061 X 10⁻⁶

METHOD 6

N= NORMALITY OF BARIUM PERCHLORATE TITRATE
 VT= VOLUME OF TITRANT, ML
 VTB= VOLUME OF TITRANT FOR BLANK, ML
 VA= VOLUME OF ALQUOT SAMPLE, ML
 VM= VOLUME OF GAS SAMPLE FROM METHOD 5, VM(DSCF)
 VSOLN= TOTAL VOLUME OF SAMPLE SOLUTION, ML
 QS= DSCF/HOUR FROM METHOD 5 CALCULATION
 DSCF= DRY STANDARD CUBIC FOOT
 MM= 1.000.000

WHEN MMBTU/HR = 1, THEN LB/MMBTU = LB/HR

$$\frac{2A + 2B}{2} = \text{Avg SO}_2$$

$$\frac{24.882 + 22.759}{2} = 23.82 \text{ \#/hr}$$

SULFUR DIOXIDE CONCENTRATION = SO₂ LB/DSCF = S2

N= .0096
 VT= 10.4
 VTB= 0
 VA= 20
 VM= .862
 VSOLN= 100
 QS= 556565
 MMBTU/HR= 1

$$S2 = (.00007061 \times N \times (VT - VTB) \times VSOLN) / VM \times VA = 4.089155E-05 \text{ LB SO}_2/\text{DSCF}$$

$$SO \text{ PPM} = 6.037 \times MM \times S2 = 246.8623$$

$$SO_2 \text{ LB/MMBTU} = QS \times S2 / \text{MMBTU/HR} = 22.75881 = \text{LB/HR}$$

STARK CERAMICS
 EAST CANTON OH
 NO 1 KILN STACK
 9/16/93
 TEST 2B

CSA CO.

EPA METHOD 6 (SULFUR DIOXIDE) DATA AND CALCULATION SHEET

TEST NO 3A PLANT STARK CERAMICS DATE 9/16/93
 LOCATION NE Kiln Stack BY AK G/HM

Vm	Tmi	ROTOMETER	TIME
166.404	78	3.5	13:55
166.207	80		14:00
166.838	82		05
167.054	84		10
167.271	85		15
			20 min
Vm .867	avg 82		

Pb 29.08

$$Vm_{std} = K_1 Y(Vm)(Pb)/(Tm)$$

$$Vm_{std} = 17.64 (.998) (.867) (29.08) / (460 + 82) = .5189$$

	1st TITRATION	2nd TITRATION	3rd TITRATION	AVERAGE	V _b TITRATION
vt	8.9	9.0	9.0	9	0

Cso₂ = Concentration of sulfur dioxide - lb/dry standard cubic foot

N = Normality of barium perchlorate titrant

Pb = Barometric pressure - inches Hg

Tm = Temperature of meter - °F

Va = Volume of sample aliquot - ml

Vm = Volume of gas at meter temperature and pressure - f³

Vm_{std} = Vm corrected to standard temperature and pressure - f³

Vsoln = Total volume of sample solution - ml

Vb = Volume of titrant used for blank - ml

Y = dry gas meter calibration factor - dimensionless

K₁ = 17.64

K₂ = 7.061 X 10⁻⁶

METHOD 6

N= NORMALITY OF BARIUM PERCHLORATE TITRATE
 VT= VOLUME OF TITRANT, ML
 VTB= VOLUME OF TITRANT FOR BLANK, ML
 VA= VOLUME OF ALQUOT SAMPLE, ML
 VM= VOLUME OF GAS SAMPLE FROM METHOD 5, VM(DSCF)
 VSOLN= TOTAL VOLUME OF SAMPLE SOLUTION, ML
 QS= DSCF/HOUR FROM METHOD 5 CALCULATION
 DSCF= DRY STANDARD CUBIC FOOT
 MM= 1,000,000

WHEN MMBTU/HR = 1, THEN LB/MMBTU = LB/HR

SULFUR DIOXIDE CONCENTRATION = SO₂ LB/DSCF = S2

N= .0096
 VT= 9
 VTB= 0
 VA= 20
 VM= .8789
 VSOLN= 100
 QS= 567866
 MMBTU/HR= 1

$S2 = (.00007061 \times N \times (VT - VTB) \times VSOLN) / (VM \times VA) = 3.470648E-05 \text{ LB SO}_2/\text{DSCF}$

$SO \text{ PPM} = 6.037 \times MM \times S2 = 209.523$

$SO_2 \text{ LB/MMBTU} = QS \times S2 / \text{MMBTU/HR} = 19.70863 = \text{LB/HR}$

STARK CERAMICS
 EAST CANTON OH
 NO 1 KILN STACK
 9/16/93
 TEST 3A

CSA CO.

EPA METHOD 6 (SULFUR DIOXIDE) DATA AND CALCULATION SHEET

TEST NO 3 B PLANT STARIC CERAMICS DATE 7/16/97
 LOCATION N^o KICK STACK BY SKGTW/MP

Vm	Tmi	ROTOMETER	TIME
167.271	85	3.0	14:22
167.436	87		127
167.601	88		132
167.767	89		137
167.932	90		14:42
Vm .661	avg 88		

Pb 29.08

$$Vm_{std} = K_1 Y(Vm)(Pb)/(Tm)$$

$$Vm_{std} = 17.64 (.998) (.661) (29.08) / (460 + 88) = .6115$$

	1 st TITRATION	2 nd TITRATION	3 rd TITRATION	AVERAGE	V _b TITRATION
Vt	8.6	8.6	-	8.6	0

Cso₂ = Concentration of sulfur dioxide - lb/dry standard cubic foot
 N = Normality of barium perchlorate titrant
 Pb = Barometric pressure - inches Hg
 Tm = Temperature of meter - °F
 Va = Volume of sample aliquot - ml
 Vm = Volume of gas at meter temperature and pressure - f³
 Vm_{std} = Vm corrected to standard temperature and pressure - f³
 Vsoln = Total volume of sample solution - ml
 Vb = Volume of titrant used for blank - ml
 Y = dry gas meter calibration factor - dimensionless
 K₁ = 17.64
 K₂ = 7.061 X 10⁻⁶

METHOD 6

N= NORMALITY OF BARIUM PERCHLORATE TITRATE
 VT= VOLUME OF TITRANT, ML
 VTB= VOLUME OF TITRANT FOR BLANK, ML
 VA= VOLUME OF ALQUOT SAMPLE, ML
 VM= VOLUME OF GAS SAMPLE FROM METHOD 5, VM(DSCF)
 VSOLN= TOTAL VOLUME OF SAMPLE SOLUTION, ML
 QS= DSCF/HOUR FROM METHOD 5 CALCULATION
 DSCF= DRY STANDARD CUBIC FOOT
 MM= 1.000.000

WHEN MMBTU/HR = 1, THEN LB/MMBTU = LB/HR

$$\frac{3A + 3B}{2} = \text{Avg Test}$$

$$\frac{19.709 + 26.803}{2} = 23.26 \text{ #/HR}$$

SULFUR DIOXIDE CONCENTRATION = SO₂ LB/DSCF = S2

N= .0096
 VT= 8.6
 VTB= 0
 VA= 20
 VM= .6175
 VSOLN= 100
 QS= 567866
 MMBTU/HR= 1

$$S2 = (.00007061 \times N \times (VT - VTB) \times VSOLN) / (VM \times VA) = 4.720294E-05 \text{ LB SO}_2\text{/DSCF}$$

$$SO \text{ PPM} = 6.037 \times MM \times S2 = 284.9641$$

$$SO_2 \text{ LB/MMBTU} = QS \times S2 / \text{MMBTU/HR} = 26.80494 = \text{LB/HR}$$

STARK CERAMICS
 EAST CANTON OH
 NO 1 KILN STACK
 9/16/93
 TEST 3B

FLORAIDE

CSA CO. DATA SHEET (METHOD 5)

B

A

164

FLOURIDE

DATA INPUT

Pb,In Hg	Barometer -----	29.08
VM,ft3	Meter Volume -----	51.6
ΔH,In H2O	Orifice Differential-----	2.77
PG,In H2O	Stack Static Pressure -----	0
Tm,F	Meter Temperture -----	89
Ts,F	Stack Temperture -----	583
CP	Pitot Coefficient -----	.84
ΔP,In H2O	Average Square Root Of Delta P -----	.4707
% CO2	Carbon Dioxide -----	3.4
% O2	Oxygen -----	16.4
% N	Nitrogen -----	80.2
VCL,Ml	Volume Of Condensate -----	68
AS,ft2	Stack Area -----	8.5
MN,Mg	Total Fluorine Weight -----	76.4
Ø,Min	Test Time -----	60
DN,In	Diameter Of Nozzle -----	.391

RESULTS

Pma,In Hg	Absolute Meter Pressure -----	29.28
PSA,In Hg	Absolute Stack Pressure -----	29.08
MD	Molecular Weight Dry Gas -----	29.20
MS	Molecular Weight @ Stack Conditions -----	28.50
VMSTP,ft3	VM Standard Temp.& Press. Dry -----	48.26
VWSTP,ft3	VM Standard Temp.& Press. Wet -----	3.20
BWO,%	Moisture -----	6.22
I,%	Isokeimetic Variation -----	91.95
VS,ft/Sec	Stack Velocity -----	37.92
QA,ASCFM	Stack Gas Flow (Actual) -----	19338
QS,ft3/HR	Stack Gas Flow (Dry STP) -----	537395
L,#/Hr	Emission Rate -----	1.88
CS,#/SCF	Fluorine Emission -----	3.49058E-06

STARK CERAMICS
EAST CANTON OH
NO 1 KILN STACK
9/16/93
TEST 1

Flouride

CSA CO. DATA SHEET (METHOD 5)

TEST NO. 2 PLANT SPARK CERAMICS								DATE 9/16/03			
LOCATION #23 KILN STACK								BY AKG/MMP			
BAROMETER(P _B) 29.09				AMBIENT TEMP		ASSUMED MOISTURE		"Y" 1.001			
MODULE NO. 203		PROBE NO. 29		FILTER NO.		HEATER NO. 8		NOZZLE DIA. .385			
TEST POINT	METER VOLUME (Vm)	METER TEMP(Tm) IN OUT		COND TEMP °F	FILTER HEATER TEMP°F	STACK TEMP °F(Ts)	STACK PRESS. (Ps)	VACUUM "Hg	ORIFICE "H ₂ O ΔH)(Pm)	VEL HEAD ΔP)	TIME
1	153.2	88	88	62	248		0	7	2.0	.15	11:51
1	157.3	94	88	60	250			7	2.0	.16	SC
2	161.3	102	88	56	251			11	2.6	.21	12:01
3	166.1	110	89	58	250			18-	3.3	.28	OG
4	171.0	115	90	56	250			19+	3.5	.28	11
1	171.0	98	89	58	250			7	1.6	.12	12:19
1	174.6	109	92	58	252			7	1.6	.12	24
2	178.7	114	94	56	250			11	2.0	.15	29
3	182.6	116	95	56	250			13	2.4	.18	34
4	186.6	120	96	56	251			19+	3.6	.27	39
1	186.6	104	94	58	252			18	3.2	.24	12:43
1	191.2	113	97	56	251			18	3.2	.24	50
2	195.7	118	97	58	250			18+	3.3	.24	55
3	200.3	122	98	58	250			18+	3.3	.24	13:00
4	206.3	124	101	60	248			18+	3.4	.24	13:05
	153.2										
(Vm)	53.1	109.9	93.0								
AVERAGE		101		****	*****	589	0	MAX 20	2.73	.214	60 MINUTES
INTEGRATED GAS SAMPLE		Z		AVG		N ₂ = 80.3		CONDENSATE COLLECTED			
CO ₂				3.1				FINAL	INITIAL	TOTAL	
O ₂		Fnd - M-5		16.6		DRIERITE		1011.6	1000.2	11.4	
CO						IMPINGER		357.9	300	57.9	
FILTER WT		PROBE WASH WT		IMPINGER WASH WT		GRAND TOTAL 69.3					
GROSS						LEAK RATE @ 20"Hg = <.02					
TARE						STACK AREA (A _S) f2					
NET						AVG SQ RT Δ P .4628					
TOTAL PARTICULAT WT (Mn) 75.6 (MF) Mg											

FLOURIDE

DATA INPUT

Pb.In Hg	Barometer -----	29.08
VM.ft3	Meter Volume -----	53.1
H.In H2O	Orifice Differential-----	2.73
PG.In H2O	Stack Static Pressure -----	0
Tm.F	Meter Temperture -----	101
Ts.F	Stack Temperture -----	588
CP	Pitot Coefficient -----	.84
P.In H2O	Average Square Root Of Delta P -----	.4628
% CO2	Carbon Dioxide -----	3.1
% O2	Oxygen -----	16.6
% N	Nitrogen -----	80.3
VCL.M1	Volume Of Condensate -----	69.3
AS.ft2	Stack Area -----	8.5
MN.Mg	Total Particulate Catch Weight -----	75.6
Ø.Min	Test Time -----	60
DN.In	Diameter Of Nozzle -----	.385

RESULTS

Pma.In Hg	Absolute Meter Pressure -----	29.28
PSA.In Hg	Absolute Stack Pressure -----	29.08
MD	Molecular Weight Dry Gas -----	29.16
MS	Molecular Weight @ Stack Conditions -----	28.46
VMSTP.ft3	VM Standard Temp.& Press. Dry -----	48.60
VWSTP.ft3	VM Standard Temp.& Press. Wet -----	3.26
BWO.%	Moisture -----	6.29
I.%	Isokeimetic Variation -----	97.36
VS.ft/Sec	Stack Velocity -----	37.40
QA,ASCFM	Stack Gas Flow (Actual) -----	19074
QS.ft3/HR	Stack Gas Flow (Dry STP) -----	527140
L,#/Hr	Emission Rate -----	1.81
CS,#/SCF	Particulate Emission -----	3.429823E-06
C'S,Grn/SCF	Particulate Emission -----	0.02395

STARK CERAMICS
EAST CANTON OH
NO 1 KILN STACK
9/16/93
TEST 2

HYDROGEN FLUORIDE
CSA CO. DATA SHEET (METHOD 5)

TEST NO. 3 PLANT STACK CERAMICS						DATE 9/16/90						
LOCATION NE KILN STACK								BY EK AG CT MP				
BAROMETER(PH) 29.08 AMBIENT TEMP								ASSUMED MOISTURE "Y" 1.001				
MODULE NO. 203 PROBE NO. 3 FILTER NO.								HEATER NO. 9 NOZZLE DIA. .305				
TEST POINT	METER VOLUME (Vm)	METER TEMP (Tm)		COND TEMP °F	FILTER HEATER TEMP °F	STACK TEMP °F(Ts)	STACK PRESS. (Ps)	VACUUM "Hg	ORIFICE "H ₂ O ΔH (Pm)	VEL HEAD ΔP	TIME	
		IN	OUT									
1	206.6	87	87	58	250		0	14	2.4	.19	13:55	
1	211.0	97	90	58	251			14	2.4	.19	14:00	
2	214.6	106	91	60	250			14	2.4	.19	05	
3	219.1	112	92	60	250			16	2.6	.20	10	
4	223.7	116	94	60	252			19	3.7	.28	15	
1	223.7	103	92	60	252			14+	2.3	.18	14:22	
1	228.0	112	94	60	251			14+	2.3	.18	27	
2	232.8	117	96	61	248			13	2.1	.16	32	
3	237.3	122	98	60	252			18	2.8	.22	37	
4	242.1	125	100	60	250			18+	2.8	.22	42	
1	242.1	110	98	58	249			19	3.5	.27	14:47	
1	246.8	119	100	58	250			19	3.5	.27	53	
2	251.4	121	101	58	250			19	3.6	.27	57	
3	255.8	125	103	58	250			19+	3.8	.28	15:03	
4	260.9	126	104	58	251			20-	4.2	.31	15:07	
	200.6											
(Vm)	54.3	113.2	96.0									
AVERAGE		125	***	*****	59.5	0	MAX 20	2.96	.278	60 MINUTES		
INTEGRATED GAS SAMPLE				AVG N ₂ =80.2	CONDENSATE COLLECTED							
CO ₂				3.2	DRIERITE	FINAL 1031.4	INITIAL 1016.2	TOTAL 15.2				
O ₂				16.6	IMPINGER	354.2	300	54.2				
CO												
FILTER WT		PROBE WASH WT			IMPINGER WASH WT		GRAND TOTAL 69.4					
GROSS						LEAK RATE @ 20"Hg = <.02						
TARE						STACK AREA (As) f2						
NET						AVG SQ RT Δ P _s = 4780						
TOTAL PARTICULAT WT (Mn) 77.8 (MF) Mg												

FLOURIDE

DATA INPUT

Pb.In Hg	Barometer -----	29.08
VM.ft3	Meter Volume -----	54.3
PH.In H2O	Orifice Differential-----	2.96
PG.In H2O	Stack Static Pressure -----	0
Tm.F	Meter Temperture -----	105
Ts.F	Stack Temperture -----	595
CP	Pitot Coefficient -----	.84
AP.In H2O	Average Square Root Of Delta P -----	.4780
% CO2	Carbon Dioxide -----	3.2
% O2	Oxygen -----	16.6
% N	Nitrogen -----	80.2
VCL.M1	Volume Of Condensate -----	69.4
AS.ft2	Stack Area -----	8.5
MN.Mg	Total Particulate Catch Weight -----	77.8
Ø.Min	Test Time -----	60
DN.In	Diameter Of Nozzle -----	.383

RESULTS

Pma.In Hg	Absolute Meter Pressure -----	29.30
PSA.In Hg	Absolute Stack Pressure -----	29.08
MD	Molecular Weight Dry Gas -----	29.18
MS	Molecular Weight @ Stack Conditions -----	28.48
VMSTP.ft3	VM Standard Temp.& Press. Dry -----	49.35
VWSTP.ft3	VM Standard Temp.& Press. Wet -----	3.27
BWO.%	Moisture -----	6.21
I.%	Isokeimetic Variation -----	97.00
VS.ft/Sec	Stack Velocity -----	38.74
QA,ASCFM	Stack Gas Flow (Actual) -----	19758
QS.ft3/HR	Stack Gas Flow (Dry STP) -----	542883
L,#/Hr	Emission Rate -----	1.89
CS,#/SCF	Particulate Emission -----	3.47624E-06
CSS,Grn/SCF	Particulate Emission -----	0.02428

STARK CERAMICS
EAST CANTON OH
NO 1 KILN STACK
9/16/93
TEST 3

Agency Use Only
Date Received _____
No. Assigned _____
Premise No. _____

INTENT TO TEST NOTIFICATION

Proposed Test Date SEPT 16, 1993
Pre-Test Meeting Desired? Yes ☐ No ☒

A. Facility Information

Name STARK CERAMICS
Contact Person SURINDER M. HESHNARY

Address 600 WEST CHURCH STREET, EAST CANTON, OH 44730
Telephone Number (216) 488-1211

B. Testing Firm Information

Name CSA CO
Contact Person ERNIE KOLM

Address 24385 CENTER ROAD, ALLIANCE, OH 44601
Telephone Number (216) 525-5119

C. Source Sampling Information: Identify all sources and pollutants to be sampled.

Source	Control Equipment	Monitoring Equipment	Pollutant to be Tested	EPA Test Method	Filter Box Temperature	Number of Sampling Points	Total Time for Test Run	Number of Sampling Runs
<u>NK 3</u> <u>BRICK KILN</u>	<u>NONE</u>	<u>NONE</u>	<u>PM</u>	<u>5</u>	<u>250°F</u>	<u>12</u>	<u>10 min</u>	<u>3</u>
			<u>SO₂</u>	<u>6</u>	<u>-</u>	<u>2</u>	<u>2 Runs @ 10 min ea</u>	<u>3</u>

Any modifications to USEPA Reference Method(s)? Yes ☐ No ☐ ; If yes explain _____

Sampling Location(s): Inlet ☐ Outlet ☐ Simultaneous ☐ Will cyclonic flow check(s) be conducted? Yes ☐ No ☐

Fuel Sampling: Coal - Proximate ☐ Ultimate ☐ Other (specify) _____

Emission Rate to be calculated using: F-Factor ☐ Ultimate Coal Analysis ☐ Other (specify) _____

Are concurrent Method 9 readings to be performed? Yes ☐ No ☐

D. Sample Train Calibration: All affected measuring and metering equipment should be calibrated within 60 days of the scheduled testing.

Plant SHAW CERAMICS

Date 9/16/93

Location EAST CANYON

Starting time 9 AM

Plant process BRICK KILN

For sample port and sample point locations, see sketch(s)

Calibration sheet(s) are attached.

GENERAL

The sample train is a Research Appliance Corp. model 2043 using a PYREX lined probe heated to 250 °F. The filter media is A/E FIBERGLASS enclosed in pyrex holders heated to 250 °F.

The impingers contain;

1st 100 ml DW

4th

2nd 100 ml DW

5th

3rd Empty

6th

A drierite column containing Drierite® (calcium sulfate) is used for final moisture removal in the impinger train.

The contents of the impingers will be used for moisture determination and _____

The front half glassware will be washed using acetone and brush or _____

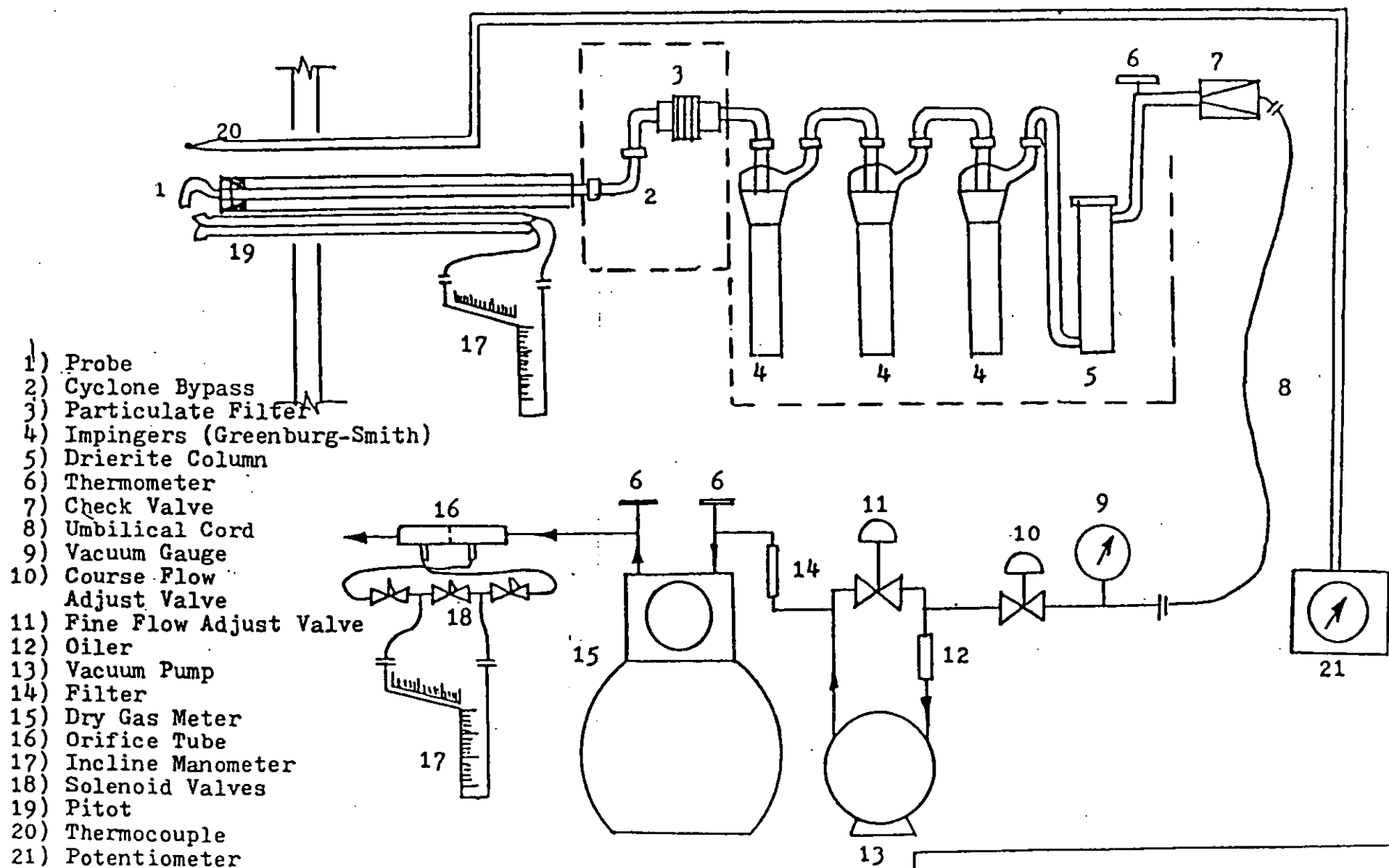
The back half will be washed using DW _____

The test and analytical procedures are as per latest Federal Register or local requirement.

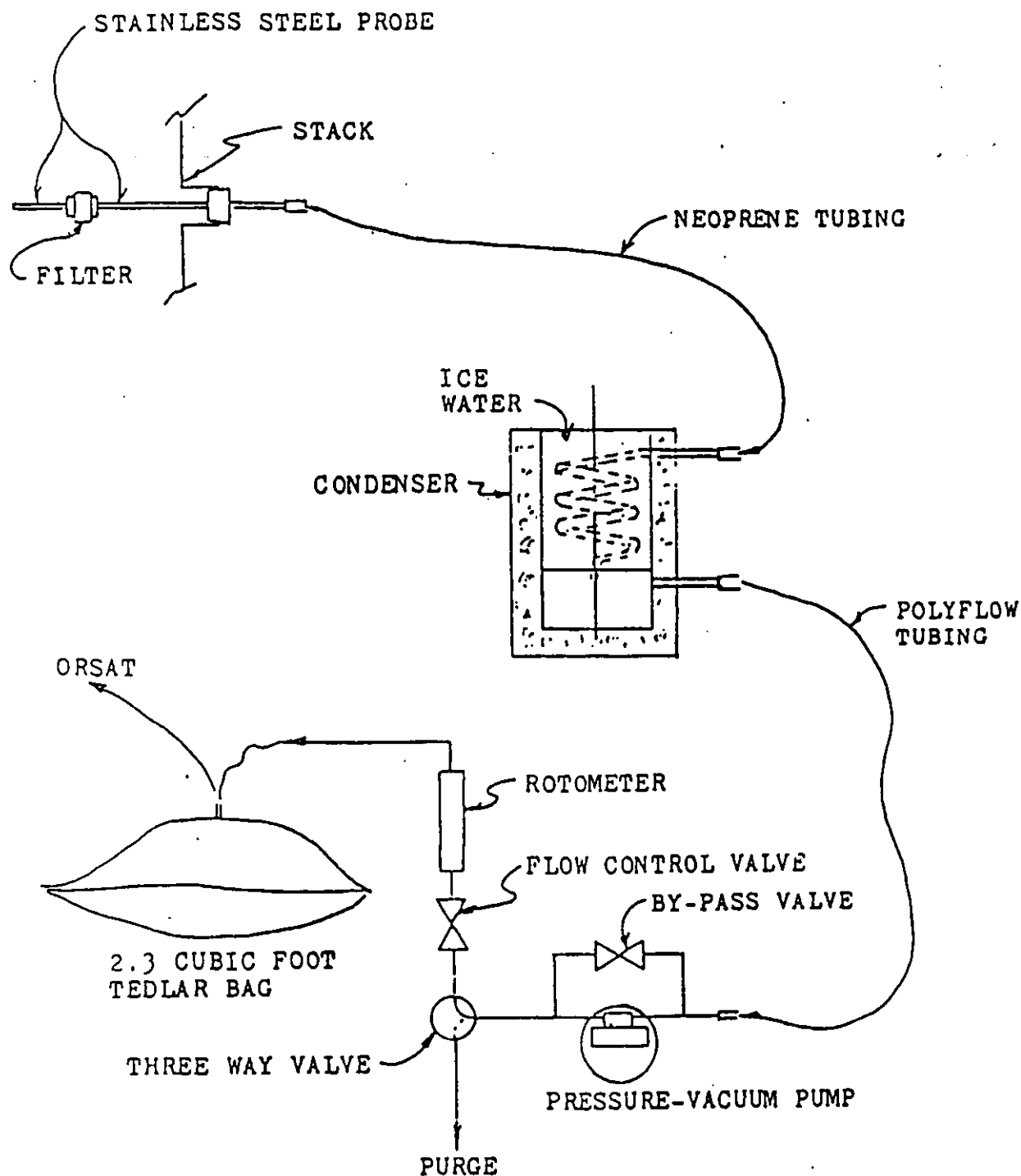
~~CO measurement by method 10-10A using the MSA Lira 3000. See sketch.~~

CO₂ and O₂ measurement by orsat from an integrated sample. See sketch.

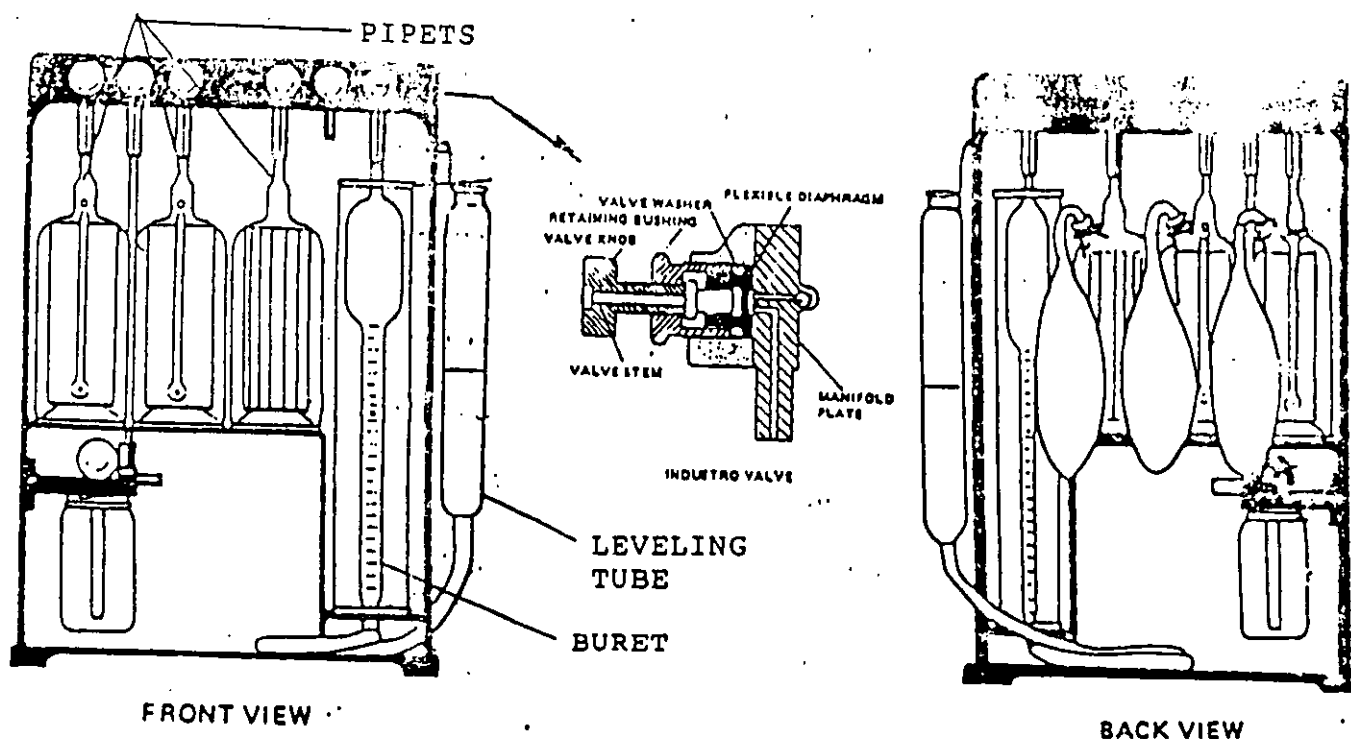
Other: SPA METHOD 6 - SEE SKETCH



C.S.A. Co.
MODIFICATION OF R.A.C. SAMPLE TR
USING DRIERITE COLUMN IN PLACE C
IMPINGER DESICANT



INTEGRATED SAMPLER



LEAK CHECKING

- 1) THE LEVEL OF THE SOLUTION IN THE PIPET WILL NOT BE MAINTAINED AT ITS PROPER LEVEL IF THERE IS A LEAK BETWEEN THE PIPET AND THE MANIFOLD VALVE OR IN THE MANIFOLD VALVE.
- 2) THE BURET SOLUTION IS SET TO ZERO WITH THE LEVELING TUBE. The LEVELING TUBE IS RAISED TO ABOUT TWENTY INCHES FOR TWO MINUTES TO APPLY PRESSURE TO THE MANIFOLD, VALVES AND REMAINING PART OF THE SYSTEM. THE LEVELING TUBE IS RETURNED TO ITS ORIGINAL ZERO POSITION. IF THERE IS A LEAK, THE BURET SOLUTION WILL BE ABOVE THE ZERO MARK. IT WILL BE ON THE ZERO MARK IF THERE ARE NO LEAKS.

THE LEAK CHECK IS PERFORMED BEFORE EACH GAS ANALYSIS OR SERIES OF ANALYSIS.

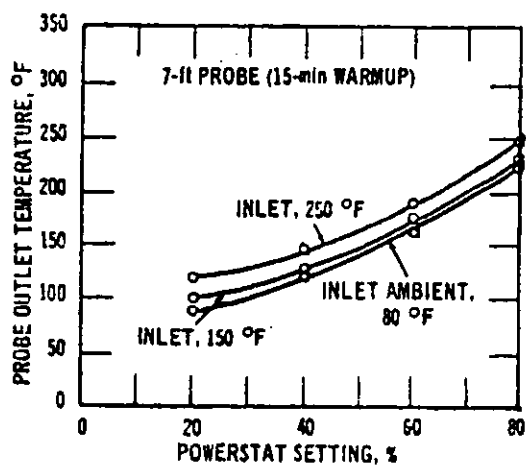
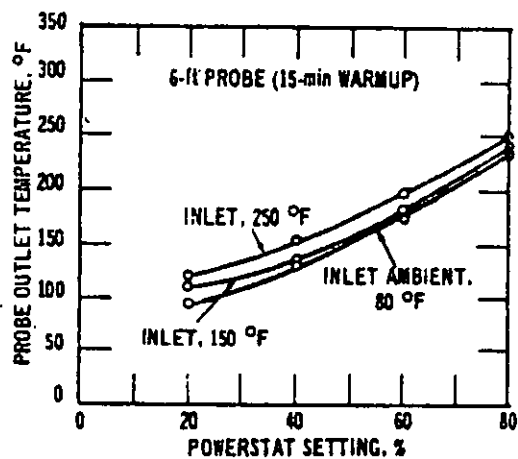
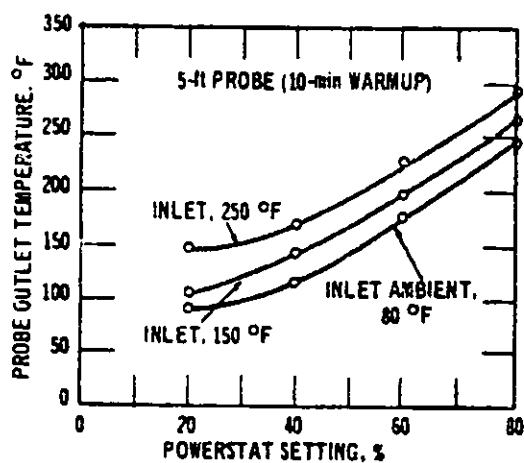
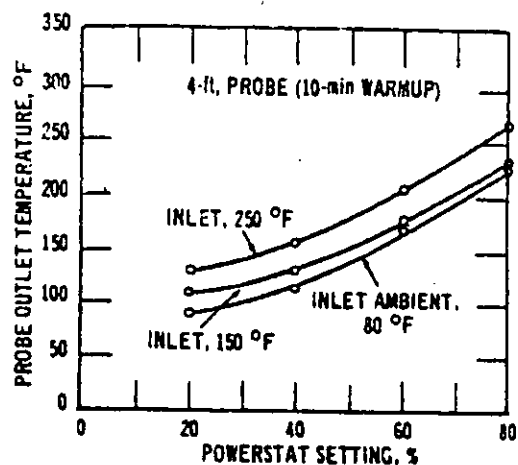
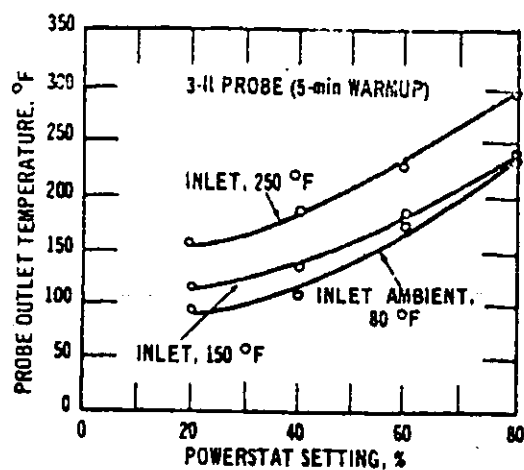
THE SOLUTIONS ARE PURCHASED FROM BURRELL CORP. THEY ARE IN THREE OUNCE BOTTLES THAT ARE THE PROPER SIZE FOR FILLING THE PIPET.

"DISORBENT" IS USED IN THE CONTACT PIPET FOR CO_2 .

"OXORBENT" IS USED IN THE FIRST AUTO-BUBBLER PIPET FOR O_2 .

"COSORBENT" IS USED IN THE SECOND AUTO-BUBBLER PIPET FOR CO .

THE BURRELL MODEL B "INDUSTRO" ORSAT GAS ANALYZER

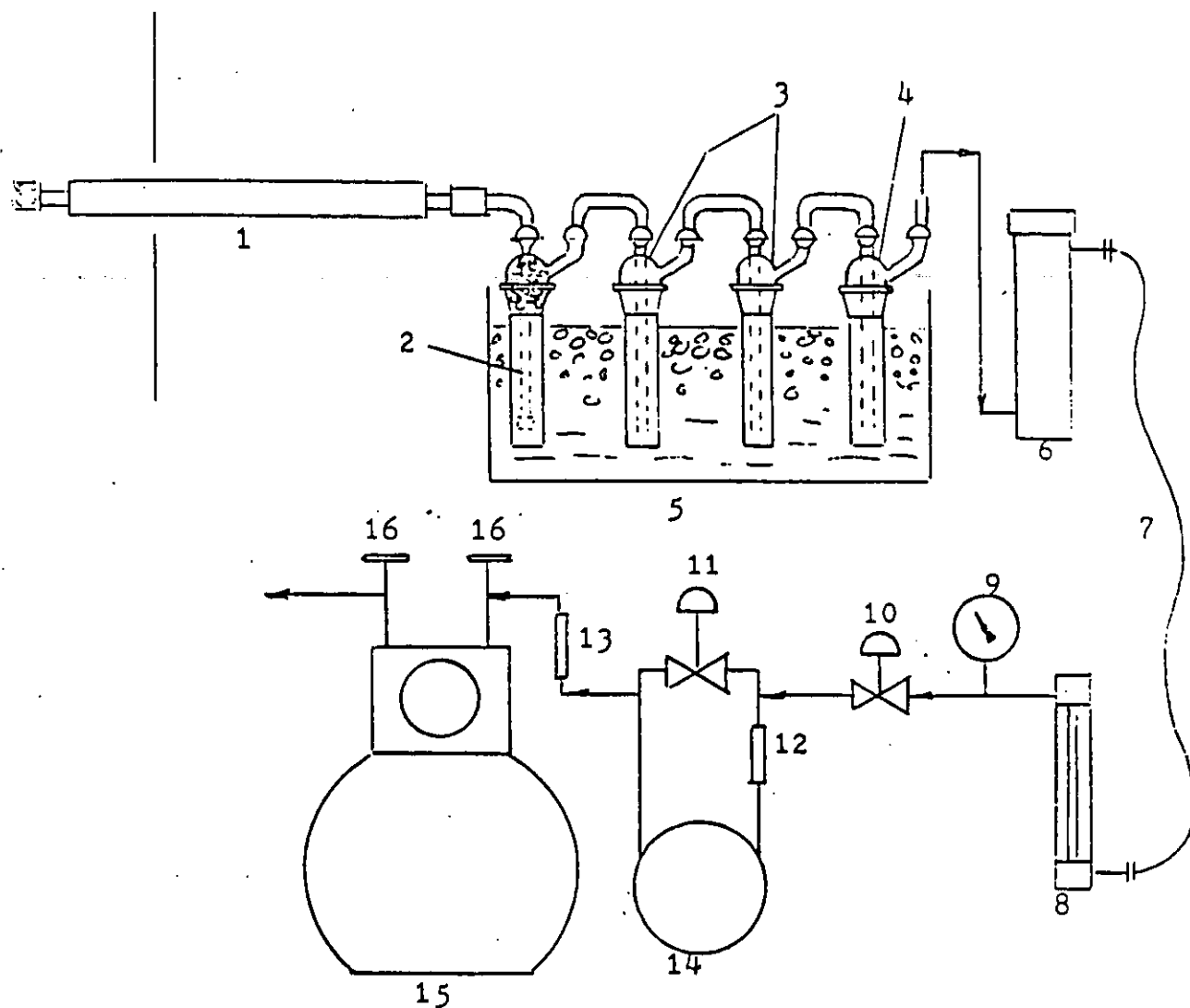


NOTE: Flow rate held constant at 0.75; 50% change in flow rate has little effect on probe temperature.

Figure 16
Probe Temperatures

CSA CO. DATA SHEET (METHOD 5)

[illegible]



- 1) HEATED PYREX LINED PROBE. END PACKED WITH QUARTZ WOOL
- 2) MIDGET BUBBLER. TOP PAKED WITH GLASS WOOL
- 3) MIDGET IMPINGERS. EACH FILLED WITH 15 ml OF H_2O_2 (3%)
- 4) MIDGET IMPINGER. DRY TRAP
- 5) ICE BATH
- 6) DRIERITE COLUMN. FILLED WITH CALCIUM SULFATE DESICANT
- 7) UMBILICAL CORD
- 8) ROTAMETER
- 9) VACUUM GAGE
- 10) VALVE FOR COARSE FLOW ADJUSTMENT
- 11) VALVE FOR FINE FLOW ADJUSTMENT
- 12) OILER
- 13) OIL FILTER
- 14) VACUUM PUMP
- 15) DRY GAS TEST METER
- 16) THERMOMETER

E.P.A. METHOD 6 SO_2 SAMPLING TRAIN

C.S.A. CO.
EPA METHOD 6 (SULFUR DIOXIDE)
DATA AND CALCULATION SHEET

TEST NO. _____ PLANT _____ DATE _____
LOCATION _____ BY _____

Vm	Tmi	Tmo	ROTOMETER	TIME
Vm	Tm			

Pbar _____

$$Vmstd = K_1 Y \frac{Vm Pb}{Tm}$$

$$Vmstd = 17.64 () \frac{() ()}{(460.+)} =$$

	1st TITRATION	2nd TITRATION	3rd TITRATION	AVERAGE	Vb TITRATION
Vt					

$$C_{SO_2} = K_2 \frac{(Vt - Vb)(N) \left(\frac{Vs_{oln}}{Va} \right)}{Vmstd}$$

$$= .00007061 \frac{(-) () ()}{()} = \text{_____ Lb/ft}^3 \text{ DSTP}$$

$$SO_2 \text{ (PPM)} = (C_{SO_2})(6.037)(10^6) =$$

$$SO_2 \text{ Lb/Hr} = (\text{ft}^3/\text{Hr DSTP}) (C_{SO_2}) =$$

C_{SO_2} = Concentration of sulfur dioxide - lb/dry standard cubic foot

N = Normality of barium perchlorate titrant

Pbar = Barometric pressure - inches Hg

Tmi = Temperature at meter inlet - °F

Tmo = Temperature at meter outlet - °F

Tm = Average meter temperature - °F

Va = Volume of sample aliquot titrated - ml

Vm = Dry gas volume as measured at meter - dcf

$Vm(std)$ = Vm corrected to standard conditions - dscf

Vs_{oln} = Total volume of sample solution - ml

Vt = Average volume of titrant used for sample aliquot - ml

Vb = Volume of titrant used for blank - ml

Y = Dry gas meter calibration factor

$K_1 = 17.64$ $K_2 = 7.061 \times 10^{-5}$

C.S.A. Co.
STACKSAMPLR CALIBRATION SHEET

Customer _____ Order No. _____
Date 6/11/93 Serial No. 2031 RAC Order No. _____

Pump OK Pump Oil OK Clean Quick Disconnects YES
Manometers OK Dry Test Meter OK Thermometers OK
Lights OK Electrical Check OK Variac OK
Vacuum Gauge OK Leak Check @ 27" Hg Vacuum NO LEAKS
Remarks PRESSURE CHECK @ 9"H2O - NO LEAKS

Barometer (Pb) 29.11

K	N	DH	CFw	CFd	Tw	ITd	OTd	TD	t	H	Y
.0158	.0368	0.5	5	5.10	68	80	75	78	12.78	1.848	0.998
.0317	.0737	1.0	5	5.12	68	85	81	83	9.05	1.838	1.002
.0634	.1470	2.0	10	10.33	68	90	86	88	12.88	1.845	1.000
.1268	.2490	4.0	10	10.30	68	88	90	89	9.14	1.844	1.001
.1902	.4310	6.0	10	10.28	68	92	90	91	7.47	1.848	1.000
.2536	.5880	8.0	10	10.25	68	96	91	94	6.48	1.851	1.003

Tolerances H=1.6-1.84-2.1 , Y=0.99-1.00-1.01

$H = (K / (Pb(OTd + 460))) * (((Tw + 460)t) / CFw)^2$

$Y = (CFw Pb (Td \text{ avg.} + 460)) / (CFd (Pb + N) (Tw + 460))$

DH= Orifice pressure drop - in. H₂O

CFw= Volume wet test meter - f³

CFd= Volume dry test meter - f³

Tw= Temp. wet test meter

ITd= Inlet temperature dry test meter

OTd= Outlet temperature dry test meter

Td avg.= Average temperature dry test meter

t= Time - minutes

Pb=Barometer press.

DIAL THERMOMETER CALIBRATION

Precision Lab Thermometer

Meter Thermometers

Impinger Outlet Thermometer (1)
(2)

ICE WATER

33

ITd 32 OTd 33

32

33

BOILING WATER

211

ITd 211 OTd 211

211

211

STACK DIGITAL TEMPERATURE INDICATORS

NBS TRACABLE FROM FACTORY. 2 UNITS CALIBRATED AGAINST EACH OTHER(PyroMation)

ALL THERMOCOUPLES ARE CALIBRATED BY OVEN FROM 100 TO 500°F AGAINST A CMS

LAB THERMOMETER (No. 227-934). ANY THERMOCOUPLE THAT IS MORE THAN ±2°F

FROM STANDARD IS DISCARDED.

C.S.A. Co.
STACKSAMPLR CALIBRATION SHEET

Customer _____ Order No. _____
Date 8/12/93 Serial No. 2031 RAC Order No. _____

Pump OK Pump Oil CHG Clean Quick Disconnects YES
Manometers CHG FLUID Dry Test Meter OK Thermometers SEE BELOW
Lights OK Electrical Check OK Variac OK
Vacuum Gauge OK Leak Check @ 27" Hg Vacuum NO LEAKS
Remarks PRESSURE CHECK @ 9" H2O - NO LEAKS

Barometer (Pb) 29.13

K	N	DH	CFw	CFd	Tw	ITd	OTd	TD	t	H	Y
.0158	.0368	0.5	5	5.07	72	82	76	79	12.80	1.877	0.998
.0317	.0737	1.0	5	5.07	72	86	80	83	9.08	1.881	1.004
.0634	.1470	2.0	10	10.24	72	89	86	88	12.91	1.880	1.001
.1268	.2490	4.0	10	10.19	72	88	85	86	9.11	1.876	0.999
.1902	.4310	6.0	10	10.18	72	91	88	90	7.46	1.877	1.001
.2536	.5880	8.0	10	10.15	72	94	89	92	6.47	1.879	1.002

Tolerances H=1.6-1.84-2.1 , Y=0.99-1.00-1.01

$H = (K / (Pb(OTd + 460))) * (((Tw + 460)t) / CFw)^2$

$Y = (CFw Pb (Td \text{ avg.} + 460)) / (CFd (Pb + N) (Tw + 460))$

DH= Orifice pressure drop - in. H2O

CFw= Volume wet test meter - f³

CFd= Volume dry test meter - f³

Tw= Temp. wet test meter

ITd= Inlet temperature dry test meter

OTd= Outlet temperature dry test meter

Td avg.= Average temperature dry test meter

t= Time - minutes

Pb=Barometer press.

DIAL THERMOMETER CALIBRATION

Precision Lab Thermometer

Meter Thermometers

Impinger Outlet Thermometer (1)

(2)

ICE WATER

33

ITd 33

33

33

BOILING WATER

211

ITd 210

211

210

STACK DIGITAL TEMPERATURE INDICATORS

NBS TRACEABLE FROM FACTORY. 2 UNITS CALIBRATED AGAINST EACH OTHER(PyroMation)

ALL THERMOCOUPLES ARE CALIBRATED BY OVEN FROM 100 TO 500°F AGAINST A CMS

LAB THERMOMETER (No. 227-934). ANY THERMOCOUPLE THAT IS MORE THAN ±2°F

FROM STANDARD IS DISCARDED.

#1

Visible Emission Observation Form

SOURCE NAME STARK CERAMICS			OBSERVATION DATE 9-16-93				START TIME 9:30		STOP TIME 11:00			
ADDRESS			SEC MIN	0	15	30	45	SEC MIN	0	15	30	45
CITY EAST CANTON STATE MA ZIP			1	5	5	5	5	31	5	5	5	5
PHONE			2	5	5	5	5	32	5	5	5	5
SOURCE ID NUMBER KIN No. 3			3	5	5	5	5	33	5	5	5	5
PROCESS EQUIPMENT KIN OPERATING MODE ON			4	5	5	5	5	34	5	5	5	5
CONTROLLER EQUIPMENT NONE OPERATING MODE			5	5	5	5	5	35	5	5	5	5
DESCRIBE EMISSION POINT STACK DISCH.			6	5	5	5	5	36	5	5	5	5
START STOP			7	5	5	5	5	37	5	5	5	5
HEIGHT ABOVE GROUND LEVEL START 50 STOP			8	5	5	5	5	38	5	5	5	5
HEIGHT RELATIVE TO OBSERVER START 0 STOP			9	5	5	5	5	39	5	5	5	5
DISTANCE FROM OBSERVER START 150 STOP			10	5	5	5	5	40	5	5	5	5
DIRECTION FROM OBSERVER START W STOP			11	5	5	5	5	41	5	5	5	5
DESCRIBE EMISSIONS GRAY SMOKE			12	5	5	5	5	42	5	5	5	5
START STOP			13	5	5	5	5	43	5	5	5	5
EMISSION COLOR GRAY PLUME TYPE: CONTINUOUS ☒			14	5	5	5	5	44	5	5	5	5
START STOP FUGITIVE ☐ INTERMITTENT ☐			15	0	5	5	5	45	5	5	5	5
WATER DROPLETS PRESENT: NO ☒ YES ☐			16	5	5	5	5	46	5	5	5	5
IF WATER DROPLET PLUME: ATTACHED ☐ DETACHED ☐			17	5	5	5	5	47	5	5	5	5
POINT IN THE PLUME AT WHICH OPACITY WAS DETERMINED START 1 DIA FROM STOP			18	5	5	5	5	48	5	5	5	5
DESCRIBE BACKGROUND GREEN TREES			19	5	5	5	5	49	5	5	5	5
START STOP			20	5	5	5	5	50	5	5	5	5
BACKGROUND COLOR GREEN SKY CONDITIONS 100%			21	5	5	5	5	51	5	5	5	5
START STOP START 0 CST STOP			22	5	5	5	5	52	5	5	5	5
WIND SPEED 1-2 MPH WIND DIRECTION SW			23	5	5	5	5	53	5	5	5	5
START STOP START STOP			24	5	5	5	5	54	5	5	5	5
AMBIENT TEMP. 65°F WET BULB TEMP. RH.percent			25	5	5	5	5	55	5	5	5	5
START STOP			26	5	5	5	5	56	5	5	5	5
Source Layout Sketch Draw North Arrow			27	5	5	5	5	57	5	5	5	5
			28	5	5	5	5	58	5	5	5	5
			29	5	5	5	5	59	5	5	5	5
			30	5	5	5	5	60	5	5	5	5
			AVERAGE OPACITY FOR HIGHEST PERIOD				NUMBER OF READINGS ABOVE % WERE					
COMMENTS			RANGE OF OPACITY READINGS				MAXIMUM					
			MINIMUM									
			OBSERVER'S NAME (PRINT) ANDY PASTO				OBSERVER'S SIGNATURE Andy Pasto					
I HAVE RECEIVED A COPY OF THESE OPACITY OBSERVATIONS			ORGANIZATION STARK CERAMICS				DATE 9/16/93					
			CERTIFIED BY				DATE					
			VERIFIED BY				DATE					
SIGNATURE			DATE				DATE					
TITLE			DATE				DATE					

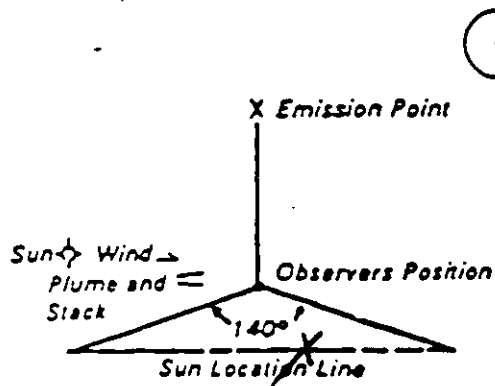
#2

Visible Emission Observation Form

SOURCE NAME STARK CERAMICS			OBSERVATION DATE 9-16-93				START TIME 11:35				STOP TIME 1:07			
ADDRESS			SEC MIN				SEC MIN							
			0 15 30 45				0 15 30 45							
CITY E. CANTON			STATE OHIO				ZIP							
PHONE			SOURCE ID NUMBER											
PROCESS EQUIPMENT KILN No. 3			OPERATING MODE on											
CONTROL EQUIPMENT None			OPERATING MODE											
DESCRIBE EMISSION POINT STACK DISCH.														
START			STOP											
HEIGHT ABOVE GROUND LEVEL START 50 STOP			HEIGHT RELATIVE TO OBSERVER START 8' STOP											
DISTANCE FROM OBSERVER START 150' STOP			DIRECTION FROM OBSERVER START W STOP											
DESCRIBE EMISSIONS GRAY-BLUE SMOKE														
START			STOP											
EMISSION COLOR GR-BL			PLUME TYPE: CONTINUOUS ☒ FUGITIVE ☐ INTERMITTENT ☐											
START			STOP											
WATER DROPLETS PRESENT: NO ☒ YES ☐			IF WATER DROPLET PLUME: ATTACHED ☐ DETACHED ☐											
POINT IN THE PLUME AT WHICH OPACITY WAS DETERMINED START 1 DIA. DOWN STOP														
DESCRIBE BACKGROUND GREEN TREES														
START			STOP											
BACKGROUND COLOR START GREEN STOP			SKY CONDITIONS 100%											
WIND SPEED 1-2 MPH			WIND DIRECTION SW											
START			STOP											
AMBIENT TEMP. 68°F			WET BULB TEMP.				RH. percent							
START			STOP											

Source Layout Sketch

Draw North Arrow



1	5	5	5	5	31	5	5	5	5
2	5	5	5	5	32	5	5	5	5
3	5	5	5	5	33	5	5	5	5
4	5	5	5	5	34	5	5	5	5
5	5	5	5	5	35	5	5	5	5
6	5	5	5	5	36	5	5	5	5
7	5	5	5	5	37	5	5	5	5
8	5	5	5	5	38	5	5	5	5
9	5	5	5	5	39	5	5	5	5
10	5	5	5	5	40	5	5	5	5
11	5	5	5	5	41	5	5	5	5
12	5	5	5	5	42	5	5	5	5
13	5	5	5	5	43	5	5	5	5
14	5	5	5	5	44	5	5	5	5
15	5	5	5	5	45	5	5	5	5
16	5	5	5	5	46	5	5	5	5
17	5	5	5	5	47	5	5	5	5
18	5	5	5	5	48	5	5	5	5
19	5	5	5	5	49	5	5	5	5
20	5	5	5	5	50	5	5	5	5
21	5	5	5	5	51	5	5	5	5
22	5	5	5	5	52	5	5	5	5
23	5	5	5	5	53	5	5	5	5
24	5	5	5	5	54	5	5	5	5
25	5	5	5	5	55	5	5	5	5
26	5	5	5	5	56	5	5	5	5
27	5	5	5	5	57	5	5	5	5
28	5	5	5	5	58	5	5	5	5
29	5	5	5	5	59	5	5	5	5
30	5	5	5	5	60	5	5	5	5

AVERAGE OPACITY FOR
HIGHEST PERIODNUMBER OF READINGS ABOVE
% WERERANGE OF OPACITY READINGS
MINIMUM

MAXIMUM

OBSERVER'S NAME
ANDY PASKOOBSERVER'S SIGNATURE
Andy PaskoDATE
9-16-96

ORGANIZATION

COMMENTS

I HAVE RECEIVED A COPY OF THESE OPACITY OBSERVATIONS
SIGNATURE

CERTIFIED BY

DATE
6/1

TITLE

DATE

VERIFIED BY

DATE

#3

Visible Emission Observation Form

SOURCE NAME			OBSERVATION DATE				START TIME		STOP TIME					
STARK CERAMICS			9-16-93				1:45		3:06					
ADDRESS			SEC	MIN	0	15	30	45	SEC	MIN	0	15	30	45
			1	5	5	5	5	31	5	5	5	5	5	5
CITY			2	5	5	5	5	32	5	5	5	5	5	5
EAST CANTON			3	5	5	5	5	33	5	5	5	5	5	5
STATE			4	5	5	5	5	34	5	5	5	5	5	5
OHIO			5	5	5	5	5	35	5	5	5	5	5	5
ZIP			6	5	5	5	5	36	5	5	5	5	5	5
PHONE			7	5	5	5	5	37	5	5	5	5	5	5
SOURCE ID NUMBER			8	5	5	5	5	38	5	5	5	5	5	5
PROCESS EQUIPMENT			9	5	5	5	5	39	5	5	5	5	5	5
KILN No. 3			10	5	5	5	5	40	5	5	5	5	5	5
OPERATING MODE			11	5	5	5	5	41	5	5	5	5	5	5
ON			12	5	5	5	5	42	5	5	5	5	5	5
CONTROL EQUIPMENT			13	5	5	5	5	43	5	5	5	5	5	5
NO			14	5	5	5	5	44	5	5	5	5	5	5
OPERATING MODE			15	5	5	5	5	45	5	5	5	5	5	5
-			16	5	5	5	5	46	5	5	5	5	5	5
DESCRIBE EMISSION POINT			17	5	5	5	5	47	5	5	5	5	5	5
STACK			18	5	5	5	5	48	5	5	5	5	5	5
START			19	5	5	5	5	49	5	5	5	5	5	5
STOP			20	5	5	5	5	50	5	5	5	5	5	5
HEIGHT ABOVE GROUND LEVEL			21	5	5	5	5	51	5	5	5	5	5	5
START 50' STOP			22	5	5	5	5	52	5	5	5	5	5	5
HEIGHT RELATIVE TO OBSERVER			23	5	5	5	5	53	5	5	5	5	5	5
START 8' STOP			24	5	5	5	5	54	5	5	5	5	5	5
DISTANCE FROM OBSERVER			25	5	5	5	5	55	5	5	5	5	5	5
START 150' STOP			26	5	5	5	5	56	5	5	5	5	5	5
DIRECTION FROM OBSERVER			27	5	5	5	5	57	5	5	5	5	5	5
START W STOP			28	5	5	5	5	58	5	5	5	5	5	5
DESCRIBE EMISSIONS			29	5	5	5	5	59	5	5	5	5	5	5
START GRAY-BLUE STOP SMOKE			30	5	5	5	5	60	5	5	5	5	5	5
EMISSION COLOR			AVERAGE OPACITY FOR HIGHEST PERIOD											
START GR-BLUE STOP			NUMBER OF READINGS ABC % WERE											
PLUME TYPE: CONTINUOUS ☒ FUGITIVE ☐ INTERMITTENT ☐			RANGE OF OPACITY READINGS											
WATER DROPLETS PRESENT: NO ☒ YES ☐			MINIMUM											
IF WATER DROPLET PLUME: ATTACHED ☐ DETACHED ☐			MAXIMUM											
POINT IN THE PLUME AT WHICH OPACITY WAS DETERMINED			OBSERVER'S NAME (PRINT)											
START 1 DIA. FROM STACK			ANDY PASKO											
DESCRIBE BACKGROUND			OBSERVER'S SIGNATURE											
GREEN TREES			Candy Pasko											
BACKGROUND COLOR			ORGANIZATION											
START GREEN STOP			I HAVE RECEIVED A COPY OF THESE OPACITY OBSERVATIONS											
SKY CONDITIONS			SIGNATURE											
START 100% 9:00 EST STOP			CERTIFIED BY											
WIND SPEED			DATE											
START 1-2 MPH STOP			9-16-93											
WIND DIRECTION			DATE											
START SW STOP			62											
AMBIENT TEMP.			DATE											
START 70°F STOP			DATE											
WET BULB TEMP.			DATE											
RH. percent			DATE											
Source Layout Sketch			DATE											
Draw North Arrow			DATE											
			DATE											
X Emission Point Sun ← Wind → Plume and Stack Observers Position 140° Sun Location Line			DATE											
COMMENTS			DATE											
			DATE											
I HAVE RECEIVED A COPY OF THESE OPACITY OBSERVATIONS			DATE											
SIGNATURE			DATE											

STARK CERAMICS (SC)
STACK TEST OF SEPTEMBER 16, 1993
15 76 06 0014

A compliance test was done on kiln No. 3 on September 16, 1993 by CSA (Ernie Kolm). The parameters evaluated were particulates, SO₂ and HF. Audit samples for SO₂ were given to Ernie for analysis.

It took about 1½ hours to get an hour's VE readings because of the time it took to transfer the testing probe and equipment to different testing traverses.

The manufacturing of ceramic bricks entails the following sequence of operations:

1. Compounded clays are processed through extruder dies for specific shape bricks. There is 5-6% moisture in the extruded bricks.
2. The freshly extruded bricks are set on a conveyor that transports the brick to a pigment spray area where the raw bricks are coated on one side with an aqueous mixture of color pigment.
3. The freshly sprayed bricks are loaded by hand on numbered cars that ride on rails. These numbered cars are next pushed into dryers where they remain for 55 hours at 180°F. In the dryer, the moisture content of the sprayed bricks very slowly diminishes to less than 1%.
4. From the dryer the car of bricks goes into the kiln preheater where the temperature is raised to 500°F over a 32 hour period. In the preheater are 16 cars from beginning to end. From the exit of the kiln preheater, the bricks enter into kiln No. 3 at about 450°F.
5. In the kiln (No. 3) 37 cars are there at all times but very slowly moving through different temperature zones from 450°F up to 2050°F then down to 400°F at the kiln exit. The exposure time in the kiln is 73 hours from entrance to exit.

The testing period covered from 9:30 a.m. to 3:06 p.m. All VE readings observed were 5% except one zero.

All the car identification tags scheduled for movement into the preheater and kiln were examined for the presence of any reburns on the cars. None indicated the presence of any reburn material on any car.

Bricks on several cars on tracks waiting to go into the preheater were closely examined. The bricks were dry but not hard. Those that had come out of the kiln and were cooling down were dry but very hard and the color pigment shiny and hard.

In the kiln control room on the table is a record of all cars in the kiln and exactly where they are located in the kiln. The kiln has a multiplicity of temperature zones from 450°F to 2050°F and down to 400°F. Instruments there provide a temperature digital readout anywhere in the kiln.

A record was made of the last 16 cars in the kiln (those put in the kiln last) on September 14, 1993 and the 16 cars in the preheater awaiting a turn to enter the kiln from the preheater. The same segments of data were collected on September 15, 1993 for the kiln and preheater.

On September 16, 1993, the day of the test, an inventory was made of those car numbers that were in the kiln. All cars that were supposed to be in the kiln were there. All number sequences of cars in the kiln were exactly what they were supposed to be based on car numbers recorded from September 14 and 15, 1993.

Surinder is providing CSA with the weight of all bricks coming out of the kiln during the 9:30 a.m. - 3:06 p.m. test period.

Photographs taken in April of emissions coming from the plant were reviewed with Surinder. He said they were coming from a clay loading dock area and that other emissions noted were probably in a baghouse area by the C plant. One of the photographs showed a blue haze in the area of the No. 3 kiln stack. Surinder made no comment regarding the blue haze. This agency did request a compliance test for kiln No. 3 because of the blue haze observed in the discharge area of that stack. The test was run to quantify the SO₂ and HF released from the stack. A blue haze is indicative of the presence of SO₂ or HF. The test will reveal the concentration of both gaseous parameters and the quantity released per hour.

The area where aqueous color glaze is sprayed on bricks before they enter the dryer was closely examined. The overspray hits an impingement plate, falls to a collecting trough and is pumped to a slurry collecting tank in another room.

Next to the slurry impingement plate is an opening to an exhaust duct that leads above the roof. The area on the roof surrounding the exhaust stack had a layer of clay (gray-white color) on the roof. The thickness and area of the clay deposits suggested that this accumulation may have been a product of escaping overspray for an extensive period of time.

Surinder said that there was a set of impingement baffles mounted in the two foot diameter exhaust stack from the glaze pigment spray area. He said he would provide me with a set of prints which depicted the stack and the impingement baffles therein. The size fan that pushes air past the baffle plates would be noted.

The slurry directed from the spraying area goes to a slurry tank from where it goes to a filter press that squeezes out the water and leaves a damp filter cake for disposal.

The matter of lead pigments being released from the baghouse in the C plant area was mentioned to Surinder. I suggested that he have a quantitative lead analysis made on these emissions so that a level of health risk could be established for the emissions from this baghouse. He appeared to resist the suggestion. He said that any lead would not be in a pure form but in a compound form. I reminded him that the toxicity was still there.

I think we will have to push more forcefully to get a lead test run from this source.



ANDY PASKO

AP0922SC.TL

64

Stark Ceramics, Inc.
600 West Church Street
East Canton, Ohio 44730
(216) 488-1211
FAX (216) 488-0333

FAX TRANSMISSION

DATE:	Sept. 2, 1993	TIME:	9:20 am
FAX #:	489-3335		
ATTN:	Mr. Bruce Blakenship		
FIRM:	Dir. of Air Pollution Control / C.C.H.P.		
FROM:	Suminder Maheshwary		
SUBJECT:	C Jet. Dust Collector / NO3 kiln Flame		
# OF PAGES (including this page):	3		

ADDITIONAL COMMENTS: Dear Mr. Blakenship:

Thank you for taking the time to visit our facility and walking through various stages of brick/terra Mfg. operations. The explanation for C Jet dust collector may be a bit more understandable from the account of events by Ken Kendall from his summary memo dated 9.1.93. It is WELL UNDER CONTROL FOR NOW. We'll follow up on your suggestions of (a) Manometer installation (b) keeping log of inspections & incidents (c) Repair records. I am also sending you the time-temp curve for NO3 kiln as per your instructions. If you have any questions, please call me.

Thanks.

Sincerely,
Sumeshwary

Stats

Stark Ceramics, Inc.

INTERNAL MEMORANDUM

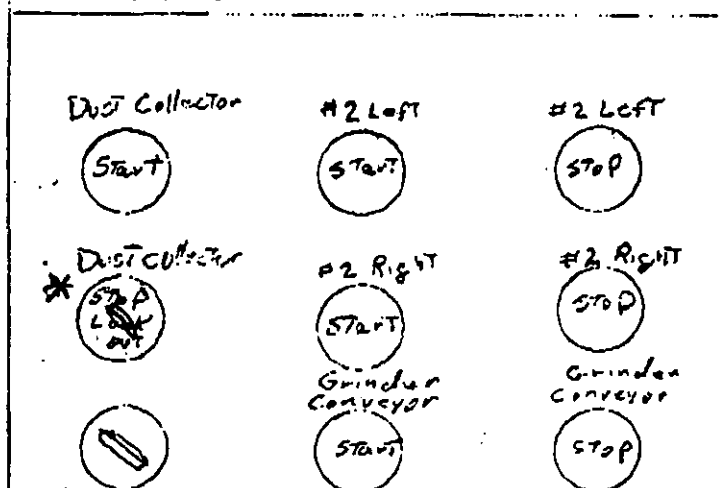
SPECIAL INSTRUCTIONS - WRITE IT DOWN!

SUBJECT:	C Plant Dust Collector	DATE	9/01/93
ATTENTION:	Surinder Maheshwary		
FROM:	Ken Kendall		

The results of the findings concerning the dust collector at C plant are as follows:

1. Maintenance was performed on the dust collector 8/28/93. Dust collector was cleaned and checked for holes in bags. Everything ok.
2. Approximately 10:00am 9/01/93, maintenance was called because of dust collector discharge. *was back in operation at 11:50 am.*
3. Shaker mechanism was tested and found to be operating properly.
4. Results of the finding are that the ^{*}lockout key switch was used to shut the system down instead of the stop button. When system is locked out no power is supplied to the shaker mechanism. This caused the bags to be overloaded and function improperly.
5. Conclusions: The supervisor and grinder ^{lead} operators were advised to the proper shut down procedure. The dust collector should be shut down properly at 9:00 break, 11:30 lunch and 4:00. This will allow proper cleaning of the dust collector bags by the shaker mechanism.

Control Panel for Grinders And Dust Collector



* It is my understanding from discussions with Dept. Foreman / Matinsipor / Electrician
Lock out key switch was "accidentally or otherwise" used at morning break or evening (K. P. D.)
causing shaker mechanism not to work. It was operational shortly before 12 noon
on 9-6-73 as per dept. foreman / electrician.

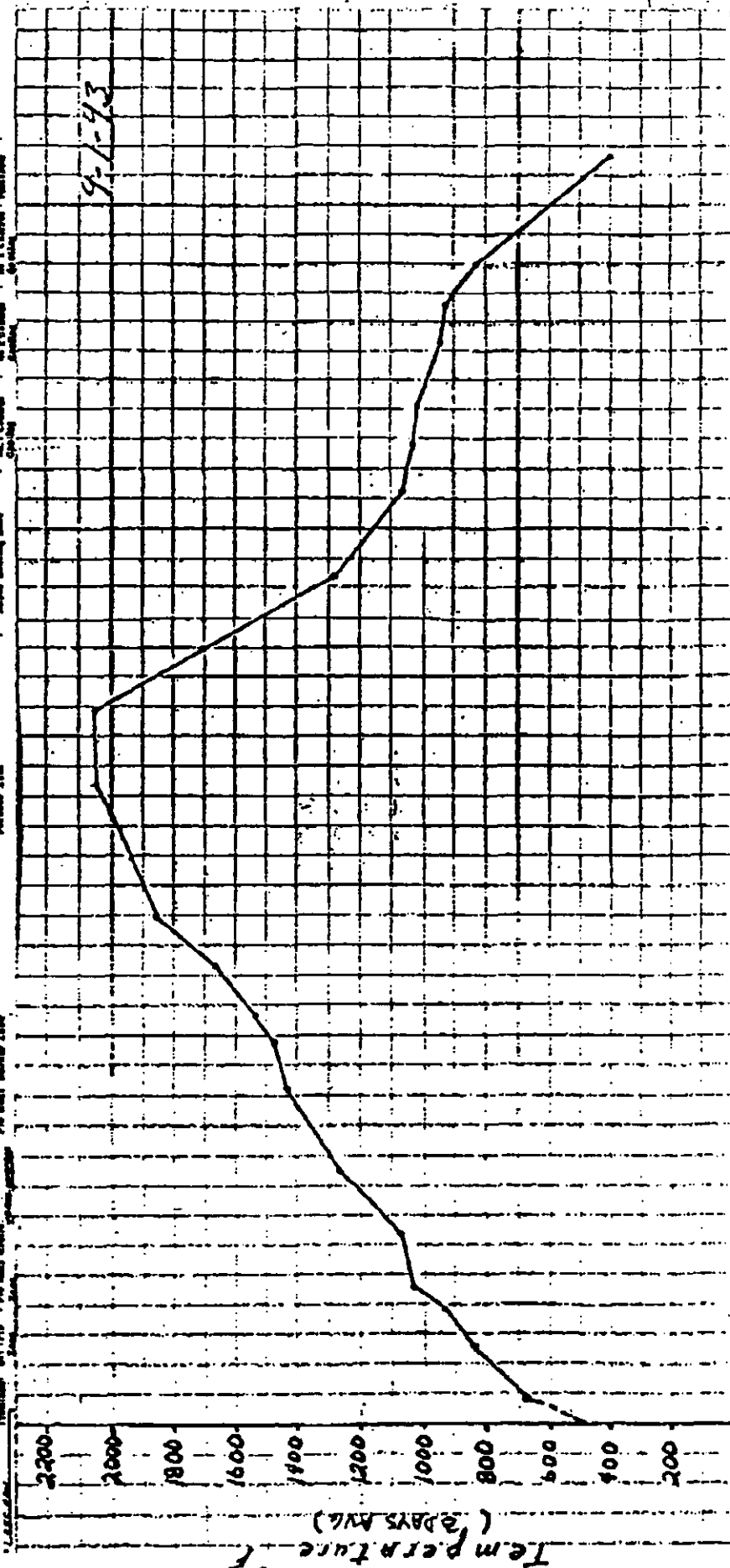
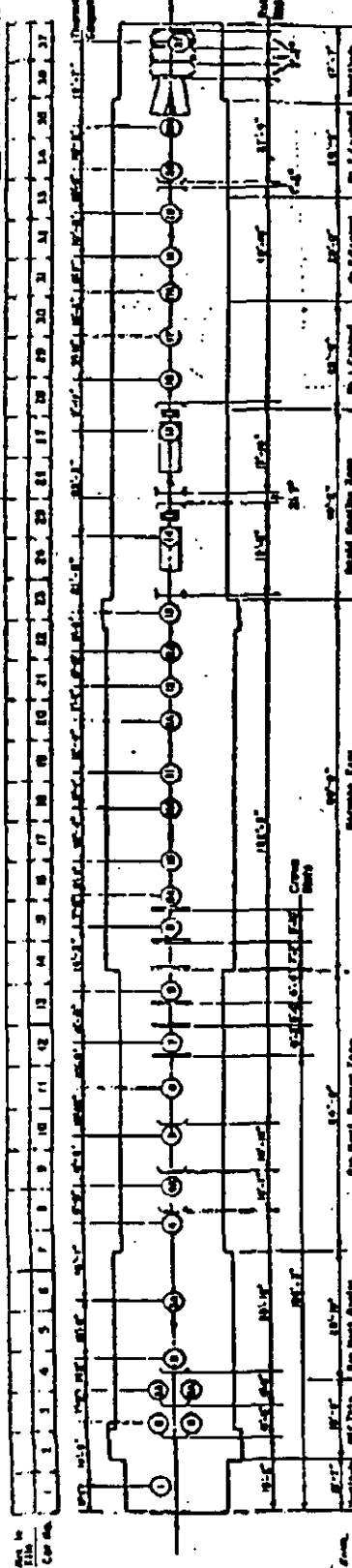
— Sunder Maheshwary 9.1.93

106

TIME-TEMPERATURE CURVE FOR #3 KILN

LONGITUDINAL ARRANGEMENT OF TUNNEL KILN #3

Cups Per 24 Hr Period



Thermocouple No. →

Thermocouple No.	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21
Temp in Kiln	0	777	167	346	740	790	793	793	793	378	378	445	482	433	537	537	537	537	537	537	537

TIME (Thermocouple)

APPENDIX K

DEFA STACK TEST REVIEW SUMMARY FORM

APPLICATION NUMBER 1576 06 0004 P005FACILITY NAME STARK CERAMICSSOURCE DESCRIPTION (OR SCC CODE) KILN No. 3CONTROL EQUIPMENT NONEDATE(S) OF TEST 9-16-93FINAL TEST REPORT RECEIVED ON 10-19-93POLLUTANT(S) TESTED PART. + SO₂TEST METHOD 5 - PART. ; 6 - SO₂TEST FIRM CSA

EMISSION RATES--

	PART	SO ₂
ACTUAL (lb(s)/hr)	5.04	22.6

	PART	SO ₂
ALLOWABLE--	9.49	46.30

OPERATING RATES--

DURING TEST-- 5842 LBS/HRMAXIMUM-- 5842 LBS/HR

EMISSION FACTOR--

COMMENTS: VISIBLE EMISSIONS DURING THE TEST WERE ALL 5% EXCEPT 2 READINGS AT ZERO

I HEREBY VERIFY THAT THE INFORMATION CONTAINED WITHIN THE STACK TEST REPORT HAS BEEN REVIEWED AND IT HAS BEEN DETERMINED THAT THE TEST PROCEDURES, ANALYSES AND CALCULATIONS ARE:

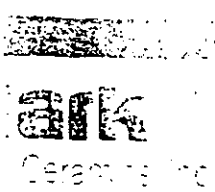
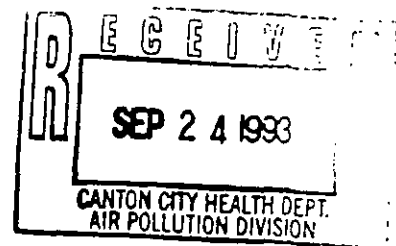
☒ AN ACCEPTABLE DEMONSTRATION OF CONFORMANCE WITH THE APPROVED TESTING METHODOLOGY.

☐ AN UNACCEPTABLE DEMONSTRATION OF CONFORMANCE WITH THE APPROVED TESTING METHODOLOGY.

10/20/93
DATE OF REVIEWAndy Parks
REVIEWED BY

-- BASED ON 3 RUN AVERAGE
-- SPECIFY APPLICABLE UNITS
-- SPECIFY IN UNITS OF MASS INPUT

68



September 23, 1993

Mr. Andy Pasko
Air Pollution Control Engineer
Canton City Health Department
420 Market Ave N
Canton, OH 44702

RE: Stack Test - Kiln #3

Dear Mr. Pasko:

As per your request, I am sending you the process weight sheets corresponding to the kiln cars that were in #3 tunnel kiln during the stack testing on September 16, 1993. The average, based on 37 kiln cars (#282 through #321), is 11,859 pounds on dry weight basis.

If you have any questions, please call me.

Sincerely,

STARK CERAMICS, INC.


Surinder K. Maheshwary
Director of Engineering, Maintenance, and QC

SKM/bjb

xc: Ernie Kolm, CSA Co.

giving 9-16-93
in film 9 AM IN

~~285~~ 285, 282, 249, 264 262 370

147 257 346 330 265 372

272 245 304 357 324 116

259 353 352 239 359 278

281 320 302 371 276 277

289 325 343 253 234 308

354 321 370ms

488 0333
FAX

Barb Coramio

Dry wt in # 148

S		shade	Shape	Fcs.	(4)		
*147	FTX	516	8WC80D	220	-	9042	
(5)		333	8W5	40	-	1,004	10,046 ✓
4							
*370	FTX	516	8WC80D	280		11,508	11,508 ✓
*262	FTX	516	8WC80D	244		10,028.4	
		904	8W54L	17		311.1	
(3)		904	8W54R	4		13.2	10,472 ✓
		815	S-453	3	C20#	60.0	
2							
*264	FTX	516	8WC80D	244		10,028.4	10,028 ✓
(2)		815	S-453	36			
*249	FTX	516	8WC80D	232		9,535.2	
		333	8W5	29		127.9	10,324 ✓
(1)		815	S-453	3	C20#	60.0	
382	FTX	516	8WC80D	244		10,028.4	
		815	S-453	36	C20	720.0	10,748 ✓
285	FTX	516	8WC80D	41		1,685.1	
		1022	412-4	1188	x10.9	12,949.2	15,701
		904	6TT20D-K	84	12.7	1066.8	

		<u>shade</u>	<u>shape</u>	<u>P.S.</u>	
	#357	195	8WCA	880	13,200
	(14)				
3	#304	195	8WCA	880	13,200
2	(13)				
	#245	127	8WCA	880	13,200
	(12)				
2	#272	127	8WCA	880	13,200
2	(11)				
1	#372	195	8WCA	7	105
1	(10)	127	8WCA	873	13,22
					13,095
	#265	127	8WCA	472	7,080
	(9)				11,683
#382 F	FTX	516	8WC80D	112	4,603.2
#330 FTX	516	8WC80D	220	9,042	10,04
(8)	333	8W5	40	1,004	
#346 FTX	516	8WC80D	220	9,042	10,01
(7)	333	8W5	40	1,004	
#257 FTX	516	8WC80D	220	9,042	10,04
(6)	333	8W5	40	1,004	

Page 3
Dry wt
in lbs

note	Shape	ics	Q		
#281	564	8WCV80BT	134	C44.4	5950
(23)	904	6TT20D-K	30	@12.7	381 12,146
	177	8WCV80BT	131	@44.4	58766
#278	*177	8WCV80BT	256		11366
(22)	904	6TT20D-K	30		381 11,747
(21)					
#359	177	8WCV80BT	280		12,432
#239	*177	8WCV80BT	250		11,100
(20)	904	8W54R	20		366 11,466
#352	177	8WCV80BT	20		888
(19)	195	8WCA	796		11,940 12,828
#353	195	8WCA	880		(13,200)
(18)					
#259	195	8WCA	880		(13,200)
(17)					
#116	195	8WCA	880		(13,200)
(16)					
#324	195	8WCA	880		(13,200)
(15)					

	Price	Shape	Qty.		
343	904	8WGBR	466	(1)	10,811.2
(31)	031	4W83-S484	12 @ 31.5		378
	516	8W24AL	20		288
	516	4T58NR	6 @ 7.77		47
#335	564	8WCV80BT	244		10834
(30)	031	4W83-S483	18 @ 31.5		378
#289	564	8WCV80BT	232		10301
(29)	516	4T58NR	4	@ 7.8	109
	516	4T58NL	10		10,786
	516	8W24CR	16		376
#277	564	8WCV80BT	232	10301	
	516	8W24CR	6		141
(28)	031	4W83-S484	7 @ 31.5		220
	516	8W54L	16		292.8
#276	564	8WCV80BT	244		10834
	516	6T54R	19 @ 10.8		205
(27)	516	8W54L	8		146.4
	904	8W54L	3		54.9
#371	564	8WCV80BT	256		11366
(26)	904	8W54L	16		292.8
#302	904	8WGBR	425		9,860
	904	6TT20D-K	46 @ 12.7		584
(25)	564	8WCV80BT	57		2531
#320	564	8WCV80BT	250		11100
(24)	904	6TT20D-K	52		660

ShadeShapePcs.

Page 5

#253	904	8WCBK	460	@23.2	10811	11756
(36)	031	4W83-5484	30	@31.5	945	

#234	904	8WCBK	516		11971	12962
(35)	904	6TT20D-K	78	@12.7	991	

#308	904	8WCBK	516		11971	12911
34	904	6TT20D-K	74		940	

#354	904	8WCBK	516		11971	12784
33	904	6TT20D-K	64		813	

#321	904	8WCBK	474		10997	11,893
	772	5W8	40	@13.5	540	
(32)	904	6TT20D-K	28		356	

Total for 37 KCs = 438783

Arg Dry wt per KC = 11,859 lbs

OR 5.93 TONS

Stark

Stark Ceramics, Inc.

September 8, 1994

Mr. Dan Schiltz
 Division of Air Pollution Control
 Canton City Health Department
 218 Cleveland Avenue SW
 Canton, OH 44702

RE: August Sulfur %

Dear Mr. Schiltz:

The 12 month rolling average % "S" from Stark's kiln operations are being faxed as per instructions. If you have any questions, please call me at 488-1211. Thanks.

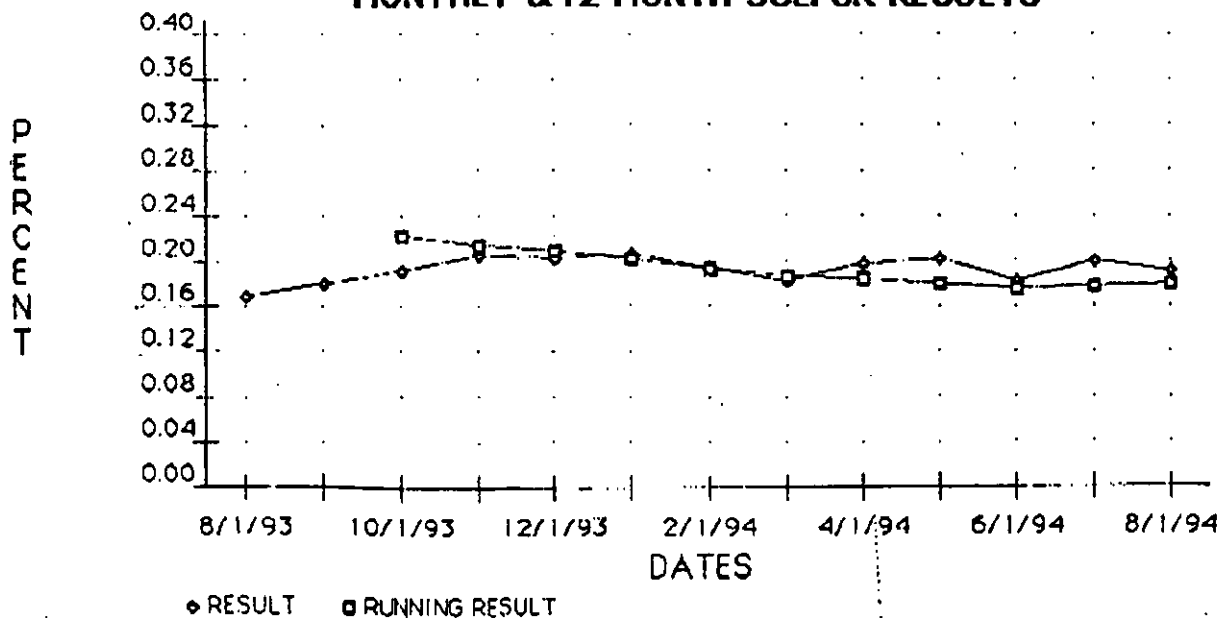
Sincerely,

STARK CERAMICS, INC.



Surlinder K. Maheshwary
 Director of Engineering, Maint., & QC

MONTHLY & 12 MONTH SULFUR RESULTS



MONTH	12/1/92	1/1/93	2/1/93	3/1/93	4/1/93	5/1/93	6/1/93	7/1/93	8/1/93	9/1/93	10/1/93	11/1/93	12/1/93	1/1/94	2/1/94
REGULATORY	0.304	0.271	0.291	0.288	0.269	0.278	0.267	0.221	0.170	0.179	0.190	0.205	0.202	0.200	0.195
ROLLING AVERAGE											0.275	0.214	0.209	0.202	0.194

3/1/94 0.185
 4/1/94 0.190
 5/1/94 0.203
 6/1/94 0.182
 7/1/94 0.199
 8/1/94 0.170