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FOR SO₂ AND PARTICULATE REMOVAL IN A MARBLE BED SCRUBBER



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LIME/LIMESTONE SCRUBBING FOR SO₂ AND PARTICULATE REMOVAL IN A MARBLE BED SCRUBBER

bу

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ABSTRACT

The Environmental Protection Agency (EPA) awarded a contract to Combustion Engineering, Inc. (C-E) to conduct research and development work on SO₂ scrubber systems using the C-E test equipment and facilities.

Sixteen once-through soluble system tests using sodium carbonate scrubbing solution were conducted. The results showed that the marble bed scrubber is a very good liquid-gas contacting device for SO_2 removal from flue gases with an overall efficiency of 90 to 95 percent. Liquid to gas ratio and scrubber liquid composition significantly affected the SO_2 removal while other variables had little or no effect on SO_2 removal.

Six limestone furnace injection systems tests were conducted using boiler calcined limestone and flyash mixture. The results also showed that solids concentration in the spray slurry and liquid to gas ratio significantly affected the SO₂ removal.

Six limestone tail-end system tests were conducted using commercial limestone in a dual marble bed scrubber. It was determined that the SO_2 removal efficiencies of the low and upper beds are the same, based on the SO_2 concentrations entering the respective beds.

It was demonstrated that scale-free operation of both the furnace injection and tail-end systems can be achieved in a closed loop system without employing the liquid blowdown by maintaining 8 to 10% solids in the spray slurry.

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SECTION 1

PROGRAM OVERVIEW

1.1 INTRODUCTION

Combustion Engineering (C-E) has developed an Air Pollution Control System (APCS) employing lime/limestone wet scrubbing. At the time of contract negotiations between C-E and EPA in 1970, the C-E APC systems at Union Electric (St. Louis) and Kansas Power and Light (Lawrence) were experiencing calcium sulfite and sulfate scaling problems.

The purpose of the contract was to analyze the previous C-E APCS data and to conduct research and development on small pilot scale (KDL pilot plant), large pilot scale (KDL prototype), and full plant scale versions of C-E lime/limestone wet scrubbing process (limestone furnace injection system with single marble bed scrubbers) in order to accelerate its commercial development by solving the calcium sulfite and sulfate scaling problems. The original purpose of the contract was later revised to include the following:

- (1) Confirm the adequacy of the methods developed by C-E to control calcium sulfate and sulfite scaling in the system.
- (2) Obtain the vapor-liquid and solid-liquid mass transfer rate data that could be used in the design of the APCS.
- (3) Predict two marble bed scrubber system performance from the performance of a single marble bed scrubber system.

Experimental work under the contract was carried out on the prototype scrubber system at the Kreisinger Development Laboratory (KDL) of Combustion Engineering in Windsor. Three kinds of systems were studied:

- (1) once through Soluble System using sodium carbonate scrubbing solution,
- (2) Limestone Furnace Injection System using boiler calcined limestone and flyash mixture as the additive, and (3) Limestone Tail-End System.

The Soluble System tests were run to obtain data pertaining to the absorption characteristics of the marble bed scrubber. Since no solids were present in the system, material balances could be made more accurately for the marble bed and overall system. This information was then used to calculate the stage efficiency of the marble bed for various test conditions.

In order to develop a better understanding of the Limestone Furnace Injection System and Limestone Tail-End System, detailed material balances were carried out for all of the tests run. These material balances permitted the calculation of dissolution and precipitation rates for important chemical species in the marble bed and associated equipment. Supersaturation of sulfur compounds was also investigated using these test data and the equilibrium computer program.

1.2 SUMMARY OF RESULTS AND CONCLUSIONS

1.2.1 Previous C-E APCS Data

EPA was supplied with the technical information generated by

C-E in previous and current APCS development work. This information consisted of reports covering the work on the KDL Prototype and the field units at Detroit Edison Company, Union Electric Company, and Kansas Power and Light Company. An oral presentation was made on October 13, 1971 in Windsor by C-E personnel to EPA and Radian Corporation personnel covering C-E's experience with Air Pollution Control Systems both in the field and in the laboratory.

The objective was to analyze previous C-E APCS data and use this data in the development of a test program to be carried out on the KDL Prototype. The purpose of this program was to determine a set of optimum operating conditions for improving the operation of C-E APCS field units. Radian and EPA concluded that the previous C-E APCS data were incomplete and could not be used in the development of the KDL Prototype test program.

1.2.2 <u>Collection and Storage of Boiler Calcined Material</u>

A mixture of boiler calcined limestone and flyash was used in six tests conducted on the KDL prototype. Considerable effort and funds were expended in the collection and storage of the boiler calcined material. The boiler calcined material and flyash were collected from unit No. 2 of the Meramec plant of Union Electric Company, St. Louis. About 30 tons of boiler calcined limestone and flyash mixture, 65 tons of boiler calcined dolomite and flyash mixture, and 50 tons of flyash was stored in 50 lb bags in the warehouse of Pozament Corporation in Milford, Connecticut. About 135 tons of boiler calcined limestone and flyash mixture was initially stored in North Haven, Connecticut for six months in a silo rented from Guyott Co. (owned by

Connecticut Highway Equipment Co.) and was later transferred to Pozament Corporation. Only a small fraction of this boiler calcined material was used and the leftover material was disposed of as instructed by EPA personnel.

1.2.3 <u>Soluble System Tests</u>

Soluble System experiments were performed to determine the vapor-liquid mass transfer characteristics (overall tray efficiency) of the marble bed scrubber using once-through sodium carbonate scrubbing solution. These tests showed that:

- (1) The marble bed scrubber is a good liquid-gas contacting device with an overall tray efficiency of 90 to 95 percent.
- (2) The ${\rm SO}_2$ removal in the marble bed scrubber is limited by the vapor-liquid equilibrium.

Liquid to gas ratio (L/G) and the scrubber liquor composition strongly influence the SO_2 removal. For example, increasing L/G from 15 to 20 GPM per 1000 CFM raised SO_2 removal from 60% to 77%. Increasing the sodium carbonate concentration in the scrubber from 25 to 120 millimoles per liter raised the SO_2 removal from 64% to 95%. Variables such as gas and liquid temperatures, scrubber feed location (above or below the bed) and gas flow do not seem to affect the SO_2 removal. No NO_X removal can be obtained with sodium carbonate scrubbing solution.

1.2.4 Limestone Furnace Injection System Tests

Limestone Furnace Injection System experiments were performed to determine the system performance and the solid-liquid mass transfer characteristics in the marble bed scrubber and the hold tank (reaction tank) using boiler calcined limestone and flyash mixture as the additive.

These tests showed that the major parameters influencing the ${\rm SO}_2$ removal of the system are liquid to gas ratio and the solids content of the spray slurry. An increase in L/G from 20 to 35 GPM/1000 CFM improved ${\rm SO}_2$ removal from 60 percent to 70 percent with other factors held constant. ${\rm SO}_2$ removal was improved from 36 percent to 68 percent by increasing the solids content of the spray slurry from 0.7 percent to 3.5 percent. Further increases in slurry concentration up to 8 percent did not result in additional improvement in ${\rm SO}_2$ removal.

Calcium sulfate scaling in the Furnace Injection System can be prevented by maintaining the relative supersaturation of this material below 1.3. This was achieved in a closed loop system with no liquid blowdown by maintaining 8 percent total solids (including flyash) in the spray slurry. Calcium sulfite scaling, on the other hand, occurs in the scrubber when the spray slurry pH reaches 11 with CaO or Ca(OH)₂ solids entering the scrubber.

1.2.5 <u>Limestone Tail-End System Tests</u>

The Limestone Tail-End System tests were performed in order to determine whether two marble bed scrubber performance (SO₂ removal and scaling) can be predicted by extrapolating the single marble bed scrubber performance of the C-E scrubber at Shawnee (EPA test facility). Information concerning the solid-liquid mass transfer characteristics in the marble bed scrubber and the hold tank was also desired.

The tests revealed that the SO_2 removal efficiency and scaling tendencies of a scrubber with two marble beds can be predicted by extrapolating single bed test results at Shawnee. The SO_2 removal efficiencies of the lower and upper beds appear to be the same based on the SO_2 concentrations entering the respective beds. SO_2 removal can be improved significantly (from 76 percent

removal to 87 percent removal) by increasing L/G from 15 to 25 GPM/1000 CFM with other factors held constant. Limestone feed rates above 100% stoichiometry have little or no effect on SO_2 removal efficiency in high solids systems. In these tests more than half the additive dissolution occurs in the marble bed in spite of the short residence time there.

Calcium sulfate scaling can be controlled in the system by maintaining the relative supersaturation level below 1.7. This can be achieved in a closed loop system without employing liquid blowdown by maintaining 8 to 10 percent solids (excluding flyash) in the spray slurry.

1.3 TEST EQUIPMENT

1.3.1 General Description

The Prototype is located at Kreisinger Development Laboratory of Combustion Engineering, Inc. in Windsor, Connecticut. A schematic of the system is shown in Figure 1-1. The system consists of all components of the C-E field units.

The flue gas from an oil fired boiler (30,000 to 40,000 pounds of steam per hour) passes through a heat extractor in which the gas can be cooled down to any desired temperature between 150 and 300°F before entering the scrubber. The flue gas from the heat extractor passes through the scrubber inlet section, marble beds, demister, and reheater before entering the stack. The scrubber inlet section is about 8 ft. long and converges towards the scrubber. Provisions are made for introducing either flyash or additive or both into the inlet either to simulate coal firing or furnace injection. The inlet is kept from plugging with deposits by the periodic operation of a soot blower.

CE AQCS PROTOTYPE

The marble bed consists of a 5 ft. by 5 ft. perforated, stainless steel plate supporting 3/4 inch diameter glass spheres (marbles) 3 inches deep; five overflow pots of 10 inch diameter; and five downcomers of 3 inch pipe. The overflow and pot height controls the turbulent layer height and is usually set at 9 inches from the perforated plate, but can be varied by making some minor changes. A stainless steel perforated plate with 3/8 inch holes and 35 percent open area supports the marbles. There are 36 commercial spray nozzles under the bed and 8 nozzles consisting of 1 inch pipes with splash plates above the bed. The spray slurry or spray liquid can be introduced either under or above the bed or both. There are two marble beds in the scrubber. The upper bed can be removed from the scrubber when it is not needed.

The chevron type demister made of stainless steel separates the entrained liquid from the gas and prevents the reheater from plugging. The gas leaving the scrubber is heated 25 to 50°F in the reheater to protect the I.D. fan.

Clarified liquid from the clarifier or the reaction tank (hold tank) effluent can be used as spray water or spray slurry. Make-up water and additive for the soluble and tail-end systems are added to the hold tank.

1.4.2 Flow Measurement

Liquid flow in the system is measured by magnetic flow meters which are calibrated both electrically and by manually measuring the flow. For the manual calibration, the hold tank is filled with water and the flow through the flow meter is set at a particular value. Water levels in the hold tank at the beginning and end of the calibration procedure are noted and the flow

meter reading is checked against the flow rate obtained from the difference in water level in the hold tank.

Additive feed rate is controlled by using Wallace & Tiernan feeders. These feeders are calibrated both by using the calibration weights and by weighing a collected sample from the feeder. The manual sample is checked against the feeder reading.

Gas flow is measured with a pitot tube located at the center of the duct in the stack after the I.D. fan. The single point pitot tube gas flow measuring technique was checked against the multiple point pitot tube traverse and the SO_2 tracer gas method. The gas flow check is given in detail in Appendix A.

 ${\rm SO}_2$ concentrations were measured using both the manual method and the Dupont 400 Photometric Analyzer. The manual method consists of absorbing ${\rm SO}_2$ gas into 3 weight percent ${\rm H}_2{\rm O}_2$ solution and titrating with 0.1N (for the inlet sample) and 0.01N (for the outlet sample) NaOH solution. The Dupont Analyzer was calibrated with ${\rm SO}_2$ gas from standard gas cylinders. The gas cylinder concentrations were also verified by the manual method described.

SECTION 2

ONCE-THROUGH SOLUBLE SYSTEM TESTS

2.1 TEST DESCRIPTION

The objective of the once-through soluble system tests was to determine the vapor-liquid mass transfer characteristics (overall tray efficiency) of the marble bed scrubber using sodium carbonate scrubbing solution.

The operation of the KDL Prototype for the once-through soluble system tests is schematically represented in Figure 2-1. Flue gas from the package boiler (burning oil) entered the scrubber after it passed through a heat extractor. The flue gas was cooled to any desired temperature between 150 and 300°F in the heat extractor. Sulfur dioxide (SO_2) gas was added to the flue gas in order to increase the scrubber inlet SO_2 concentration to approximately 2,000 PPM (0.2 mole %). The flue gas passed through the marble bed and the turbulent layer where it was in contact with the scrubbing liquor. The flue gas left the system after passing through a demister and a reheater.

Scrubbing liquor was prepared by mixing solid sodium carbonate (Na_2CO_3) and well water in the hold tank. The hold tank of approximately 6,000 gallon capacity represented an average residence time of 30 to 40 minutes for most of the soluble system tests. Thus, fluctuations in the scrubber liquor composition due to minor fluctuations in the solid Na_2CO_3 feed to the hold tank could be assumed negligible. Scrubber liquor $(Na_2CO_3$ solution) was introduced into the scrubber through 36 spray nozzles under the bed and/or through eight pipes with splash plates at the end above the bed. The bed reject was drained through the scrubber bottom while the liquid from the turbulent layer was drained through the overflow pot-downcomer arrangement.

Liquid from the downcomers and the scrubber bottoms was pumped out of the system through the clarifier which was used as a liquid disposal tank during the once-through soluble system tests. Scrubber bed height or turbulent layer height was varied by varying the overflow pot height.

The test program for the soluble system tests was designed to study the effect of operating parameters such as gas flow rate, liquid flow rate, liquid to gas ratio, scrubbing liquor composition and temperature, scrubber inlet gas temperature and scrubber bed height on the vapor-liquid mass transfer characteristics of the marble bed scrubber. The proposed test program is shown in Table 2-1. Actual test conditions and test data are given in detail in Appendix B and are very nearly the same as the proposed test conditions given in Table 2-1.

2.2 DATA EVALUATION

2.2.1 <u>System Performance</u>

The test data and results are given in detail in Appendix B. Several runs were repeated because the total sulfur material balance did not close within \pm 10%. The data and results of the runs for which the material balance did not close within \pm 10% are not given in this report. Gas flow checks, as described in Appendix A, and liquid flow calibration checks were made periodically.

In all the runs except run 10R the overflow pot height was set at 9 inches from the perforated plate. At this setting the bed drained normally with most of the water draining through the overflow pots and downcomers, and with very little water draining through the bed itself. But, in run 10R, the overflow pot height was set at 15 inches from the perforated plate. Seepage through the bed was excessive; about 95% of the water drained through the bed while only 5% drained through the overflow pots. This was

2-1

TABLE 2-1. TEST PARAMETERS FOR THE SOLUBLE SYSTEM

	Experiment Number	Comments	FG Rate (ACFM)	SF Rate (GPM)	SF Composition M Moles/Lit.	SW Rate (GPM)	SW Composition M Moles/Lit.	Inlet Gas Temperature (°F)	Scrubber Bed Height (inches)	Hold Tank Temp. (°F)
	1R	Low Gas Temp.	11,000	55	25	110	25	225	9	110
	2R	SF Only	11,000	165	25	C	25	300	9	110
	3R	SW Only Amb. Liq. Temp.	11,000	0	25	165	25	300	9	70
	4R	SW Only	11,000	0	25	165	25	300	9	110
	5R	High Gas Flow	13,000	55	25	110	25	300	9	110
	6R	Low Gas Flow	9,000	55	25	110	25	300	9	110
	7R	High L/G	11,000	70	25	150	25	300	9	110
	8R	Low L/G	11,000	35	25	75	25	300	9	110
•	9R	Base Cond.	11,000	55	25	110	25	300	9	110
•	10R	High Bed Ht.	11,000	55	25	110	25	300	15	110
	118	SW Only Amb. Liq. Temp.	11,000	0	120	165	120	300	9	70
	12R	Base Cond.	11,000	55	120	110	120	300	9	110
	1 3R	Base Cond.	11,000	55	35	110	35	300	9	110
	14R	Low L/G	11,000	35	35	75	35	300	9	110
	15R	High Gas Flow	13,000	55	35	110	35	300	9	110
	16R	Low Gas Flow	11,000	55	35	110	35	225	9	110

Note: Inlet SO₂ is 2000 PPM

because the gas could not support a high enough turbulent layer to facilitate bed drainage through the overflow pots. Also, it was observed that the seepage through the bed increased when the scrubbing liquor was introduced into the scrubber above the marble bed rather than under the marble bed.

The results show that liquid to gas ratio (L/G) and scrubber liquor composition significantly affect the SO_2 removal in the scrubber. For example in experiments 4R and 7R while keeping other conditions the same, an increase in L/G from 15 to 20 GPM/1000 CFM resulted in SO_2 removal increase from 60 to 77%. Furthermore, an increase in liquor composition from 25 to 120 m moles/lit result in an increase in SO_2 removal from 64 to 95%. The other variables such as gas and scrubbing liquor temperatures, gas flow and feed location do not seem to have significant effect on SO_2 removal in the marble bed scrubber. The inlet and outlet NOx concentrations given in Appendix B are approximately the same within the accuracy of the experimental measurements. Therefore, it can be concluded that no NOx removal can be obtained with sodium carbonate scrubbing solution.

2.2.2 Sampling and Analytical Methods

The pH measurements were made with a SS-3 Zeromatic Beckman pH meter, which was carefully standardized and temperature compensated. The pH meter was always located 1 to 2 feet away from the sample points for better pH representation of the sample. The sample temperatures were measured with a mercury thermometer during sampling.

Samples were pumped through a Millipore filter holder (142 mm in diameter) and filtered through a l_μ Millipore membrane. The filtering equipment was set up 4 to 5 feet away from the sampling point to minimize the transport time between the sample point and filtration. The greater the

transport time the greater the chance for oxidation of sulfite to sulfate. A screw type pump was used to transport the sample isokinetically.

The analytical methods are given in detail in Appendix D.

The analytical procedures used by Combustion Engineering (C-E) and Radian

Corporation are different. At the instructions of the EPA project officer,

the sulfite analysis was made using Radian's procedure although it is more

laborious and time consuming than C-E's method. Sodium and total sulfur

analyses were made using C-E's method. In a few runs, sulfite analysis was

made using both the C-E and the Radian methods to compare the accuracy of the

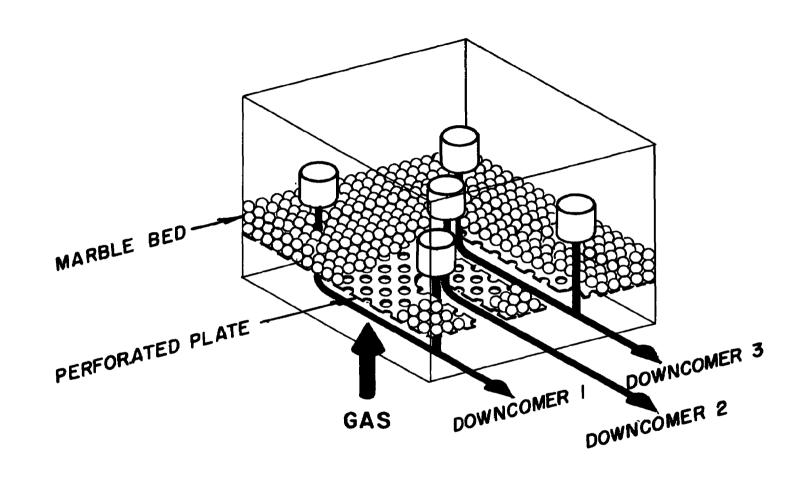
two methods. The liquid phase analytical results given in Appendix C show

that the C-E and Radian methods are comparable in results.

2.2.3 Analytical Results

The liquid phase analytical results and pH's are given in Appendix C. The preliminary tests showed that the marble bed liquor composition was not uniform over the entire cross-section of the marble bed. This probably resulted from non-uniform gas distribution to the marble bed.

The scrubber liquor discharges through three downcomers as shown in Figure 2-2. In order to determine which downcomers should be sampled to get a reasonable value for the concentration of sulfite $(SO_3^- + HSO_3^-)$ in the bed, all the three downcomers were sampled for a few runs. Based on the results of these runs, it was decided to sample the downcomers from both sides of the bed for sulfite and average the results to obtain the values for the concentration of sulfite in the bed. All other analyses for the bed liquor composition were made on pump discharge from the surge tank. The pump discharge is a good average of the three downcomers, since all three downcomers discharge into the surge tank, from which liquor is pumped to the clarifier tank for disposal.



OVERFLOW POT AND DOWNCOMER ARRANGEMENT

2.2.4 Material Balance

The results of the total sulfur material balance for the scrubber are shown in Table 2-2. The sulfur removed from the gas ΔS_{G} , and the sulfur absorbed by the liquid, ΔS_{L} , are calculated, for all the liquid streams entering and leaving the scrubber, from the following relationships:

 ΔS_G = (Gas Flow Rate X SO_2 concentration) in - (Gas Flow Rate X SO_2 concentration) out ΔS_L = Σ (Flow X Total Sulfur Concentration) out - Σ (Flow X Total Sulfur Concentration) in

The inlet gas flow rates were calculated from the measured outlet gas flow rates by correcting for liquid evaporated or condensed in the scrubber and for the 6.7% air leakage into the system between the inlet and outlet sampling points.

The results are within the accuracy of the experimental errors indicating adequacy of the flow measurements, sampling and analytical techniques.

2.3 CALCULATION OF STAGE EFFICIENCIES

Of the sixteen experiments run with sodium carbonate as the additive on the KDL APCS Prototype, five were chosen from which to calculate stage efficiencies based on maximum theoretical absorption of SO_2 . Experiments 1R, 5R, 8R, 12R, and 14R were evaluated since they represented variations in L/G, stoichiometry, inlet SO_2 , and other operating parameters.

The theoretical maximum amount of SO_2 which could have been absorbed in each experiment was determined by the following method. Soluble analyses from the marble bed effluent were the input to an equilibrium

Determined by Orsat measurement of oxygen in flue gas entering and leaving scrubber.

TABLE 2-2. TOTAL SULFUR MATERIAL BALANCE

Experiment		Gas Flow (CFM)	Liquor Flow (GPM)	SO ₂ Conc	entration PM)		Sulfur Removed From Gas	Sulfur Absorbed By the Liquid	AC . AS . W 100
Number	Date	(@ 130°F)	Below/Above	In	Out	% SO ₂ Removal	ΔSG (g Moles/Min)	ΔS _L (g Moles/Min)	ΔS _G - ΔS _L x 100
1R Set 1	10/29/71	10,960	107/54	2,018	880	61.4	11.92	12.63	- 6.0
Set 2	10/29/71	10,960	107/54	2,018	860	62.3	11.92	12.38	- 3.9
2R Set 1	10/27/71	10,750	165/0	2,050	75 0	63.5	12.78	13.23	- 3.5
Set 2	10/27/71	10,800	165/0	2,050	75 0	63.5	12.57	13.18	- 3.1
3R Set 1	10/14/71	11,200	170/0	2,095	860	59.0	14.18	14.73	- 3.9
Set 2	10/14/71	11,200	170/0	2,095	860	59.0	14.18	14.45	- 1.9
4R Set 1	10/28/71	10,800	170/0	2,030	800	60.7	12.10	12.21	- 0.9
Set 2	10/28/71	10,800	170/0	2,030	79 0	61.2	12.21	12.10	+ 0.9
5R Set 1	11/ 2/71	12,980	106/55	2,275*	1,020*		14.72	14.35	2.5
Set 2	11/ 2/71	12,980	107/55	2,290*	1,048*	54.3	-	-	-
6R Set 1	11/ 2/71	9,180	110/55	-	-	-	-	-	-
Set 2	11/ 2/71	9,180	110/55	2,050	480	76.5	14.06	13.84	- 1.6
7R Set 1	11/ 3/71	11,240	152/69	2,000	450	77.5	16.28	16.58	- 1.8
Set 2	11/ 3/71	11,240	152/69	2,000	460	77.0	16.16	16.61	- 2.8
8R Set 1	11/ 3/71	11,200	73/36	-	-	-	-	-	-
Set 2	11/ 3/71	11,190	73/36	1,782*	829*		9.39	9.65	- 2.8
9R Set 1	10/29/71	11,000	116/54	2,050	70 0	65.9	13.76	14.38	- 4.5
Set 2	10/29/71	10,910	116/54	2,010	732	63.7	12.96	14.27	-10.1
10R Set 1	11/ 9/71	10,680	112/53	1,980	540	72.6	14.22	14.08	1.3
Set 2	11/ 9/71	10,690	112/53	1,960	520	73.5	14.32	14.36	- 0.3
11R Set 1	10/14/71	11,500	165/0	1,980	120	94.0	22.25	22.40	- 0.7
Set 2	10/14/71	11,400	169/0	1,980	120	94.0	22.06	24.75	- 7.7
12R Set 1	11/ 9/71	11,210	110/53	2,020	80	94.4	20.61	20.10	2.5
Set 2	11/ 9/71	11,200	110/53	1,980	100	94.3	20.15	19.35	4.0
13R Set 1	11/ 4/71	11,330	110/54	2,000	420	81.5	17.78	18.06	- 7.6
Set 2	11/ 4/71	11,400	110/54	1,980	320	86.3	19.11	18.23	- 3.0
14R Set 1	11/ 5/71	11,300	75/36	2,070	780	62.4	13.20	12.93	2.0
Set 2	11/ 5/71	11,360	75/36	2,040	780	61.7	12.91	12.80	0.9
15R Set 1	11/ 5/71	12,980	110/55	2,040	500	75.6	18.97	18.85	0.6
Set 2	11/ 5/71	12,980	110/55	2,040	500	75.6	18.97	18.79	1.0
16R Set 1	11/ 5/71	11,500	110/55.5	2,010	350	82.5	18.03	17.86	1.1
Set 2	11/ 5/71	11,500	110/55.5	2,020	350	83.2	18.14	17.90	1.3

^{*}Manual ${\rm SO}_2$ Readings

computer program (obtained from EPA and modified by C-E) which calculated the partial pressure of SO_2 in equilibrium with the downcomer liquid at the scrubber operating conditions. To obtain an equilibrium $\underline{\mathrm{line}}$, the computer calculation was repeated for incremental amounts of total SO_2 (SO_3^- and HSO_3^-) added to the liquid over what was actually present in the analysis. For Experiment IR - Set 1 (Figure 2-3) two variations in this approach were tried: in the first case, the amount of total SO_2 which had oxidized in the liquid to sulfate was held constant as additional amounts of total SO_2 (liquid) were input to the computer program, while in the second case the ratio of sulfate to total SO_2 in the actual analysis was held constant as additional sulfur was added to the liquid. As can be seen in Figure 2-3, keeping the ratio constant caused only a small change in the equilibrium line and subsequently only a very small change in the stage efficiency calculation. For this reason, the equilibrium line for the other experiments was obtained by the first method described above.

Following construction of the equilibrium line on axes of mole fraction SO_2 in the gas versus mole fraction total sulfur in the liquid, an operating line was derived for each experiment and plotted on the same diagram. This operating line was obtained from the material balance equation

$$L (X_{out} - X_{in}) = G (Y_{in} - Y_{out})$$

where L = liquid flow rate entering and leaving stage

G = gas flow rate

 X_{out} = liquid composition leaving stage

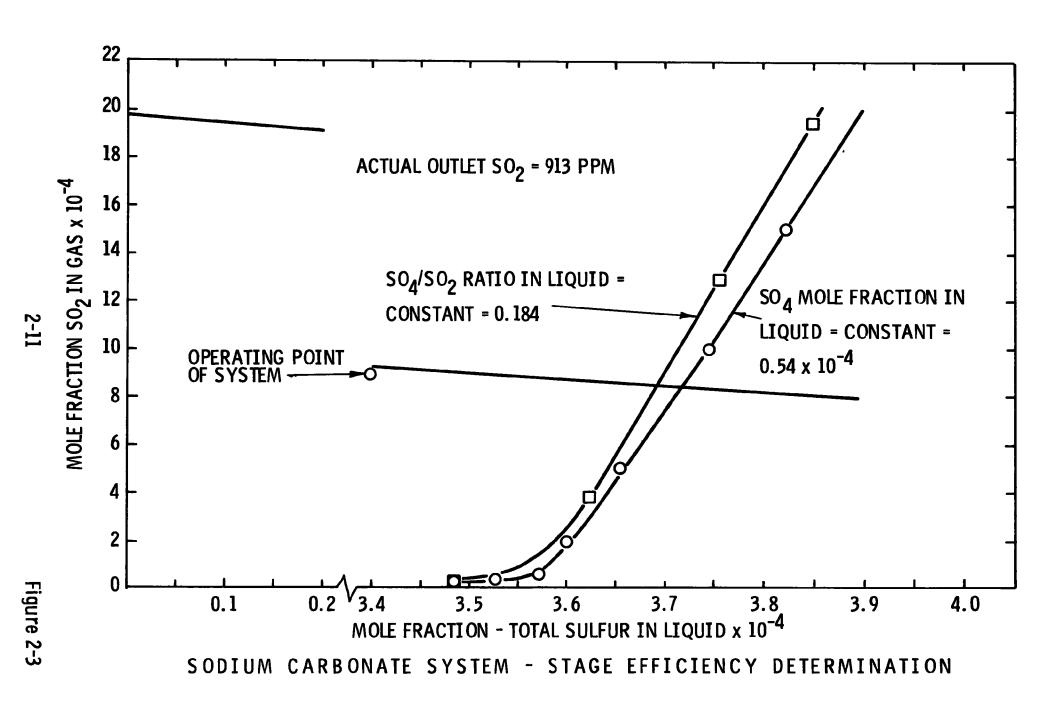
 X_{in} = liquid composition entering stage

 Y_{out} = gas composition leaving stage

 Y_{in} = gas composition entering stage

rearranged to

$$Y_{out} = (-L/G) X_{out} + [(L/G) X_{in} + Y_{in}]$$



Corrections in SO_2 concentrations and gas flow rates were made for air leakage and changes in humidity across the marble bed. The point of maximum theoretical SO_2 absorption was obtained from the intersection of the operating and equilibrium lines.

The diagrams for the other experiments done are presented in Appendix E. Efficiencies were obtained from

Inlet
$$SO_2$$
 - Outlet SO_2 (Actual)

Inlet SO_2 - Outlet SO_2 (Theoretical)

and the results tabulated along with operating parameters in Table 2-3.

The actual operating point of the marble bed for experiment 1R Set 1 was plotted in Figure 2-3 by averaging the analysis of the downcomer and bottoms streams and using the corrected SO_2 outlet concentration. If the material balance for this experiment had closed completely, the point would have fallen on the operating line. As can be seen, some deviation exists and causes a small error in the efficiency calculation. In the other experiments, the point fell either slightly above or below the operating line causing a maximum + 5% error to be introduced into the efficiency calculation.

The values for the stage efficiencies in Table 2-3 are very close for all of the experiments except 8R. This low value is caused by errors in material balance and analysis and is not due to any operating condition. In fact, no conclusions can be made regarding which film, gas or liquid, controls mass transfer based on these test data since too many conditions are varied from test to test.

In general, the data indicates that the marble bed is an efficient SO_2 contractor and that SO_2 removal was limited by vapor-liquid equilibria in the test run. Assuming that the bed is well-mixed, the rate of mass transfer is controlled by the composition of the bulk liquid which determines the rate of product and reactant diffusion through the liquid film.

TABLE 2-3. SUMMARY OF STAGE EFFICIENCY CALCULATIONS

Experiment Number/ Set Number	L/G	Moles Na ₂ CO ₃	Inlet*	Outlet* SO2	Equilibrium Outlet SO ₂	Stage Efficiency
1R/Set 1	15.2	0.284	1980	913	850	94.5
1R/Set 2	15.2	0.291	1980	902	840	94.5
5R/Set 1	12.8	0.255	2175	1055	1010	96
8R/Set 2	10.0	0.295	1680	857	690	83
12R/Set 1	15.2	1.89	1920	83	0	95.5
14R/Set 1	10.1	0.307	1940	805	720	93

^{*}Inlet and outlet SO₂ values represent concentrations immediately before and after the marble bed, not at the points where they were actually measured. Corrections for humidity were made for both inlet and outlet values while corrections for leakage were made for inlet but not outlet values.

Calculation of gas-phase mass transfer coefficients (K_ga) for the experiments run was not attempted because of the difficulty in determining the gas phase driving force. By examining the plot of Experiment 1R in Figure 2-3 it can be seen that small errors in the scrubber effluent analysis can cause large errors in the calculated SO_2 partial pressure of the sample. For example, if the actual mole fraction of total sulfur in the liquid is 3.65×10^{-4} corresponding to partial pressure of 500 ppm SO_2 , an error of \pm 3% in the analysis would cause the calculated SO_2 partial pressure to fluctuate from 100 to 1000 ppm SO_2 .

If we use the relationship

$$\frac{G}{K_g}$$
a X N_{og} T.U. = constant

to calculate K_{a} ;

where G = gas flow rate

 K_{ga} = gas phase mass transfer coefficient N_{oq} T.U. = number of gas phase transfer units

 N_{og} T.U. is determined from

$$N_{og}$$
 T.U. = $\frac{Y}{SO_2}$ in - $\frac{Y}{SO_2}$ out $\frac{Y}{Y}$ - $\frac{Y*}{Im}$

which reduces to

$$N_{og}$$
 T.U. = $ln \frac{y}{y} \frac{SO_2 in - y*}{SO_2 out - y*}$

for a well-mixed reactor where Y* is the ${\rm SO}_2$ partial pressure over the liquid. From these equations it can be seen that determination of ${\rm SO}_2$ partial pressure over the liquid is an important step in obtaining Kga's and any error in partial pressure calculations would be reflected in the Kga values.

SECTION 3

LIMESTONE FURNACE INJECTION SYSTEM TESTS

3.1 SYSTEM CHEMISTRY

The process of removing SO_2 from the flue gas using boiler calcined limestone (CaO) as the additive in the limestone furnace injection system consists of the following reactions:

$$CaO + H_2O \longrightarrow Ca(OH)_2 \tag{1}$$

$$Ca(OH)_2 + 2SO_2 + H_2O \longrightarrow Ca(HSO_3)_2 + H_2O$$
 (2)

$$CaSO_3 + SO_2 + H_2O \rightarrow Ca(HSO_3)_2$$
 (3)

$$Ca(HSO_3)_2 + Ca(OH)_2 \rightarrow 2CaSO_3 + 2H_2O$$
 (4)

$$CaSO_3 + 1/2 O_2 \rightarrow CaSO_4$$
 (5)

The CaO coming from the furnace is first hydrated as shown in reaction 1. Removal of SO_2 in the limestone furnace injection system depends upon the formation of calcium bisulfite by reaction of suspended calcium sulfite (reaction 3) and calcium hydroxide (reaction 2) with sulfur dioxide and water.

The reactions in which soluble bisulfite is converted to insoluble calcium sulfite (reaction 4) and sulfite is oxidized to sulfate (reaction 5) account for the water products as well as the regeneration of the solid calcium sulfite reactant that is recirculated to the scrubber.

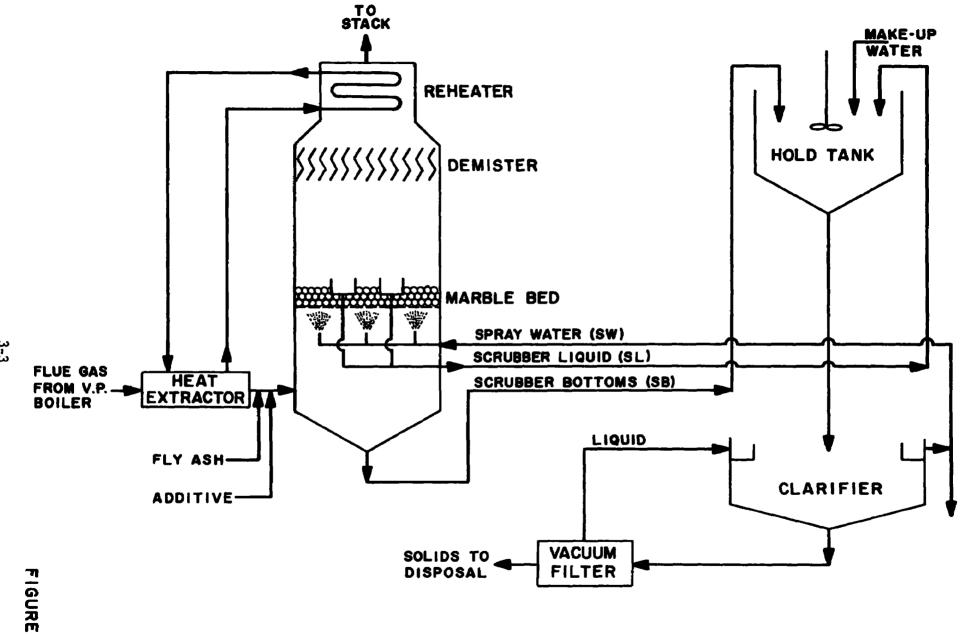
3.2 TEST DESCRIPTION

The purpose of the furnace injection test series was to determine the solid-liquid mass transfer characteristics in the scrubber and the hold tank, and to define a range of satisfactory operating conditions for application to the field units. In addition the following information was determined because of its importance in designing furnace injection SO_2 scrubbing systems:

- (1) Rate of hydration and dissolution of calcined limestone.

 This determines the alkalinity in the scrubber bed and the size of the reaction tank. This rate can be determined either using a material balance or using equilibrium methods.
- (2) Rate of precipitation of calcium sulfate, calcium sulfite and calcium carbonate. These rates assist in designing the reaction tank so that the exiting stream will be close enough to saturation to prevent calcium sulfate scaling in the scrubber.
- (3) Rate of oxidation to sulfate. This determines the incremental increase in supersaturation of calcium sulfate in the marble bed and determines the liquid to gas ratio (L/G) and the limit on supersaturation entering the scrubber needed to prevent calcium sulfate scaling in the scrubber system.
- (4) A correlation between the reactivity of the calcined limestone and its rate of hydration and dissolution. The design of the SO_2 scrubbing system will not only depend on the above measurements but will very strongly depend on the reactivity of the additive entering the system. The reactivity in turn usually depends on the following variables: type of limestone, temperature of calcination and place of injection in the furnace.

Figure 3-1 is the flow arrangement for experiment 17R and Figure 3-2 is for experiments 18R, 19R, 20R, 21R and 22R. The test conditions are shown in Table 3-1. In these tests, the furnace injection APC System was simulated by blowing a mixture of boiler calcined limestone and flyash mixture into the scrubber inlet gas stream. Liquid SO_2 was vaporized using steam and then injected into the flue gas, FG (generated from an oil fired boiler) to increase the SO_2 concentration to 0.15 to 0.2 mole percent, depending on the test requirement. In Experiment 17R, the slurry from the marble bed turbulent



LIMESTONE FURNACE INJECTION SYSTEM-WITH RECYCLE

3-2

TABLE 3-1. TEST PARAMETERS FOR THE LIMESTONE FURNACE INJECTION SYSTEM

Test No.	1 <i>7</i> R	18R	19R	20R	21R	22R
Gas Flow Rate, ACFM @ 120°F	11,000	11,000	10,000	10,000	10,000	10,000
Inlet SO ₂ , PPM	1,500	1,500	2,000	2,000	2,000	2,000
Additive (Flyash &						
calcined limestone)						
feed rate (% of						
stoichiometry)	75	75	75	75	75-100	75-100
Underbed Slurry (GPM)	110	198	205	205	200	350
Liquid to Gas Ratio, L/G						
(GPM/1000 CFM)	10	18	20	20	20	35
Overbed Spray, GPM	0	0	0	0	0	0
Excess 0, %	5	5	-	-	8	5
Inlet Gas Temp (°F)	300	300	300	300	300	300
Liquid Blowdown (GPM)	55	0	-	-	0	0
Clarifier Liquid (GPM)	0	25	-	-	15	15
Hold Tank, Tank Capacity (GAL)	6,000	6,000	6,000	6,000	3,000	5,200
Hold Tank Stirring	Max.	Max.	Max.	Max.	Max.	Max.
Make up Water (GPM)	55	5	-	-	5	5
Solid Concentration in						
Spray Slurry (%)	0	3	2	1	8	8

layer (SL) left the scrubber through the overflow pots into the downcomers and then discharged into the hold tank. The turbulent layer provided gas liquid contacting for SO₂ absorption. The scrubber bottom slurry (SB), which is rejected spray water, flyash and additive, was also discharged into the hold tank. The hold tank provides good solid liquid contacting and thus allows for hydrolysis and subsequent dissolution of calcined limestone. The slurry entering the hold tank had a pH of 4-6; the slurry leaving had a pH of 10-11. The hold tank effluent was discharged into the clarifier where the solids were settled, and the clarifier underflow was sent to the vacuum filter where the solids were further concentrated and then sent to disposal. Most of the clear liquid (165 gallons per minute) was carried to the scrubber as spray water (pH - 10.5 -11) and the remaining clarifier liquid of about 50 gallons per minute was blowdown.

In experiments 18R, 21R and 22R, part of the hold tank effluent was used as spray water, while the rest was sent to the clarifier, and the liquid returned to the hold tank. In these tests the solid concentration in the slurry was maintained between 3 and 8 percent (30% to 60% flyash, see Table F-7).

In experiments 19R and 20R, the solid concentration in the slurry was about 1 and 2 percent, and therefore a larger portion of the hold tank effluent was sent to the clarifier. Part of the clarifier liquid separated in the clarifier was removed from the system as "blowdown" and the rest was returned to the hold tank.

The flow rates for all the streams in experiments 17R to 22R are shown in Table F-1 in Appendix F.

To determine when the system reached steady state, samples were taken from the spray water (SW), and the clarifier liquid (CL) and analyzed for calcium, sulfite and sulfate. Steady state in these tests was defined as the point when the calcium and total sulfur concentration in the filtrate of the clarifier liquid (CL) and the spray water (SW) were reasonably close. Depending upon the test conditions, steady state was usually reached after 6-20 hours of operation. These analyses of the samples to determine steady state are shown in Tables F-2 to F-4 in Appendix F.

Spot checks of the liquid and gas flowmeters were made on a regular basis before every test. These checks showed that the original calibration curves prepared during the soluble tests were still valid. A listing of these procedures is available in the soluble system section and in Appendix A. Major mechanical modifications which were made between experiments are listed in Appendix G.

3.3 DATA EVALUATION

3.3.1 <u>System Performance</u>

Table 3-2 summarizes the limestone furnace injection results. In experiments 21R and 22R, while holding other conditions the same, the SO_2 removal efficiency increased from 59 percent to 72 percent with an increase of L/G from 20 to 36 GPM/1000 CFM. Therefore, the SO_2 removal efficiency tends to increase with liquid to gas ratio. In experiments 20R, 19R and 18R, as the solids concentration in the slurry increased from 0.7 to 3.5 the SO_2 removal efficiency increased from 36 to 68 percent, while in experiments 18R and 21R no further increase in the SO_2 removal efficiency was observed as the solid concentration in the slurry was increased from 3.5 to 7.4 percent. Therefore, it can be concluded that no improvement in SO_2 removal efficiency can be obtained by increasing the solid concentration in the spray water

TABLE 3-2. SUMMARY OF LIMESTONE FURNACE INJECTION TESTS PERFORMANCE RESULTS

Experiment No.*	1 <i>7</i> R	20R	19R	18R	21R	22R
Gas Flow ACFM @ 130°F	11,000	10,020	10,000	11,000	9,800	9,900
L/G, GPM/100 CFM	10.0	20.3	20.2	18.6	20.4	36
Inlet SO, Conc. (PPM)	1,456	1,950	1,882	1,471	1,992	2,020
Solid in Underbed Slurry (%)	0	0.7	1.3	3.5	7.4	8.9
Solid Recycle (%)	0	72	85	89	95	95
Spray Water pH	11.2	5.8	5.5	10.6	8.6	6.0
Stoichiometry (%)	71.0	85.5	89.1	72.8	90.1	88.1
SO ₂ Removal Efficiency (%)	43.0	35.7	43.6	67.6	59.2	72.5
Liquid Blowdown to						
Control Calcium						
Sulfate Scale	Yes	Yes	Yes	No	No	No
Calcium Sulfate Scaling	No	Yes	Yes	Yes	No	No
Calcium Sulfite Scaling	Yes	No	No	No	No	No

^{*} The listing of experiments is based on the order in which they were conducted.

beyond 3.5 percent for this system. Table 3-2 also shows that the solid recycle increase is accompanied by an increase in spray water pH and SO_2 removal efficiency. This is believed to be the result of increased retention time of the solids in the system, which allows the hydration and dissolution of CaO to near completion and thus provide greater alkalinity and consequently results in greater pH and SO_2 removal efficiency.

In experiment 17R the spray water pH was about 11 and minor calcium sulfite scaling resulted. Calcium sulfate scaling did not occur anywhere above a solids concentration in the slurry of about 8 percent, but did occur at solids concentrations below 3.5 percent. This leads to the conclusion that liquid blowdown is not needed to control calcium sulfate scaling when high solid concentration in the slurry is utilized.

The problems associated with the furnace injection system test are listed in Appendix G.

3.3.2 <u>Analytical Results and Sampling Methods</u>

A solid-liquid separation device consisting of a Millipore filter and filter holder was used to obtain solid and liquid samples. The samples were drawn such that the residence time in the sampling system was much smaller than in the vessel from which the sample was drawn. Since the marble bed slurry discharge had to flow a long distance before entering the hold tank, samples from both the marble bed and the scrubber liquid at the hold tank were taken to determine if any change had taken place while flowing in the pipe. The same technique was used with the scrubber bottom, where samples were taken at both the scrubber and the hold tank.

Since Radian Corp. was performing most of the solid and liquid chemical analyses, C-E decided to analyze the liquid samples for soluble calcium, sulfite and sulfate mainly for control purposes. In experiments 17R and 18R, C-E used the same method used by Radian for the sulfite

analysis, namely the Arsenite method. In experiments 21R and 22R, C-E used the sodium thiosulfate back titration method, while in experiments 19R and 22R, no analyses were made by C-E.

A summary of each analytical procedure is given in Appendix D. The C-E and Radian Analytical results in experiments 17R, 18R, 21R, and 22R are within 10 percent of each other, except for the marble bed samples from experiments 21R and 22R which differed by about 40 percent. The difference between the C-E and Radian results in the marble bed is attributed to difficulties in sampling.

Results of the individual liquid and solid analysis made by C-E and Radian are listed in tables F-5 through F-16 in Appendix F. The chemical analysis of the additive is listed in Tables F-17 and F-18 in Appendix F.

3.3.3 Total Sulfur Material Balance

Detailed calculations of the total sulfur material balances of the limestone furnace injection experiments are listed in Table H-1 through H-4 in Appendix H. The results showed unexpectedly good material balance closure. The purpose of performing sulfur material balances was to check the reliability of the flow measurements and the analytical results and as a criterion for determining the reliability of the tests. Of the experiments with high solid concentration in the slurries, only experiment 18R was used to perform a total sulfur material balance. The closure errors between the total sulfur in entering and leaving streams were relatively low within 9 percent in the hold tank and within 13 percent in the marble bed.

3.3.4 Rate Calculations with Slurries of Low Solid Concentrations

It was found that in order to successfully and completely characterize the streams in the system, only slurries with low solids

concentration could be used. This is because in the case of high solid slurries, a difference in the calculated rates resulting from a slight change in solid concentration was masked by that resulting from the error in solid concentration measurement. For example, when the error involved in the solid concentration measurement is ±5%, then a change in solid concentration of ±2% due to precipitation or dissolution will be completely masked by that error. With zero or low solid concentration in the slurry, the rates of formation, dissolution and oxidation were successfully calculated without making any significant assumptions. These calculations which are listed in Table H-5 to H-7 in Appendix H were made for experiments 17R, 19R and 20R. While experiment 17R gave consistent results, experiments 19R and 20R gave very inconsistent results.

3.3.5 Rate Calculations With Slurries of High Solid Concentrations

As mentioned in the previous section, a detailed species material balance cannot be performed successfully in experiments with high solids concentrations due to high experimental errors. Therefore a slightly different approach had to be taken in calculating the rate of precipitation, dissolution and oxidation. Based on the results obtained in the experiment with low solids concentration (17R), the following assumptions were made:

- (1) All of the oxidation in the system occurred in the marble bed.
- (2) Total oxidation in the system is the ratio of the total sulfate to total sulfur in both the solid and liquid streams leaving the system.

 $0xidation = \frac{(S0_4^{=}) Liquid + (S0_4^{=}) Solid}{Total Sulfur}$

- (3) Formation of $CaCO_3$ in the scrubber is negligible.
- (4) The amount of ${\rm CO}_2$ transferred to the hold tank from the atmosphere is negligible.

These assumptions were applied to data from experiments 17R, 19R and 20R, as well as to experiments 18R, 21R and 22R. The rate calculations using a liquid material balance and the above assumptions are shown in Table H-8 to H-13 in Appendix H.

Tables 3-3a and 3-3b summarize all the rate results obtained for all the experiments. The following criteria were used to determine the reliability of the results:

- (1) Total calcium hydroxide dissolution rate in the system should not exceed the total calcium feed rate to the system with the additive.
- (2) The rate of SO_2 removal from the gas should always be greather than the calcium sulfite and calcium sulfate precipitation rates in the whole system.
- (3) The rate of $S0_2$ oxidation anywhere in the system cannot be negative.
- (4) The rate of $Ca(OH)_2$ dissolution anywhere in the system cannot be negative.
- (5) The error in the total sulfur material balance around both the hold tank and the marble bed should not exceed 10 percent.

Table 3-4 lists these criteria for all the furnace injection experiments, and indicates whether or not each of these crtieria is satisfied. Calculated rate data for the following tests are considered reliable:

17R, (17R), 18R, (19R), (20R)

TABLE 3-3a. SUMMARY OF RATE RESULTS FROM LIMESTONE FURNACE INJECTION TEST

			ent 17R		Experim	ent 19R	Experiment 20R		
		e Bed	Hold Tank		le Bed	Hold Tank	Marble Bed	Hold Tank	
Location	<u>Set #1</u>	<u>Set #2</u>	Set #1 Set	2 Set #1	Set #2	Set #1 Set #2	Set #1 Set #2	Set #1 Set #2	
Outlet Flue Gas Flow (CFM @ 130°F)	11,000	11,000		10,000	9,940		10,020 10,020		
Liquid to Gas Ratio - (GPM/1000 CFM)	10	10		20	20		20 20	t.	
CaO Feed Rate (M Moles/Min)	11,306	11,306		15,750	15,750		15,750 15,750	ı	
Stoichiometry based on inlet SO_2 (%)	71.0	71.0		88.0	89.5		85.0 86.0	I	
SO ₂ Removal Eff. (%)	43.0	43.0		43.7	43.6		36.1 35.3	(
Solid Conc. in Spray Slurry (wt. %)	NONE	NONE		1.14	1.46		0.69 0.7	,	
ΔS _G (Amount of SO ₂ absorbed (M Moles/Min)	7,077	7,077		5,766	5,368		6,616 6,400	ı	
Ca(OH) ₂ Dissolution (M Moles/Min)	2,196 (5,007)	2,698 (5,358)	1,900 1,48 (1,151) (7	38 3,006 17) (268)	7,844 (1,965)	3,687 3,006 (2,579) (2,577			
Sulfite Oxidation (M Moles/Min)	3,130 (3,107)	4,305 (2,951)	224 28	31 - 600 - (3,775)	-2,994 (3,275)	530 508 	557 1,339 (3,652) (3,488		
CaSO ₂ -1/2 H ₂ O formation (M ³ Moles/Min)	1,307 (1,355)	791 (2,144)	1,848 1,96 (2,083) (2,19		1,358 (-4,200)	5,329 4,804 (5,858) (5,312	1,209 395) (-1,887) (-1,741		
CaSO ₄ ·2 H ₂ O formation (M Moles/Min)	- 955 (1,592)	- 539 (784)		57 - 899 19) (- 105)	- 550 (- 273)	2,198 1,223 (383) (186			
CaCO ₃ formation (M'Moles/Min)	- 216	16	790 51 (226) (1	77 - 985 58)	565	142 - 32 (320) (68	- 612 - 660)	- 34 210 (159) (170)	
Error in Total Sulfur Material balance (<u>In - Out</u>) x 100 (%)	- 10.7	- 12.4	1.5 - 2	.1 6.0	9.0	5.0 2.3	5.2 - 0.7	3.6 5.2	

TABLE 3-3b. SUMMARY OF RATE RESULTS FROM LIMESTONE FURNACE INJECTION TESTS

			ent 18R			Experin	ment 21R			Experiment 22R		
		e Bed	Hold	Tank	Marbl	e Bed		Tank	Marbl	e Bed	Hold Tank	
Location	Set #1	Set #2	Set #1	Set #2	Set #	Set #2	Set #1	Set #2	Set #1	Set #2	Set #1 Set	#2
Outlet Flue Gas Flow (CFM @ 130°F)	11,000	11,000			9,670	10,000			9,940	9,900		
Liquid to Gas Ratio - (GPM/1000 CFM)	18.6	18.6			20.6	20			36	35.5		
CaO Feed Rate (M Moles/Min)	11,503	11,503			16,599	16,599			16,599			
Stoichiometry based on inlet $S0_2$ (%)	72.8	72.8			91.5	88.8			87.8	88.4		
SO ₂ Removal Eff. (%)	67.6	67.6			57.7	60.7			70.8	74.2		
Solid Conc. in Spray Slurry (wt. %)	3.67	3.35			8.02	6.67			8.58			
ΔSG (Amount of SO ₂ absorbed) (M Moles/Min)	10,717	10,717			13,917	14,703			16,760			
Ca(OH) ₂ Dissolution (M Mole/Min)	6,137	6,868	3,387	3,286	5,179	9,422	-10,894	- 6,564	14,745		15,167	
Sulfite Oxidation (M Moles/Min)	2,990	3,097			4,968	3,910			5,799			
CaSO ₃ •1/2 H ₂ O formation (M Moles/Min)n)	5,943	5,027	1,435	1,849	2,563	4,380	- 280	556	4,664	17,265		
CaSO ₄ •2 H ₂ O formation (M Moles/Min)	- 1,339	- 560	1,711	1,069	1,150	459	- 1,391	- 3,985	4,664		9,442	
CaCO ₃ formation (M Moles/Min)							- 72	594			398	
Error in Total Sulfur Material balance (<u>In - Out</u>) x 100 (%)	13.5	12.2	0.3	- 9.2								

TABLE 3.4. CRTIERIA FOR DETERMINATION OF LIMESTONE FURNACE INJECTION TEST RELIABILITY

Test No.	Ca(OH) ₂ Dissolution > Ca++ in Additive		ΔS _G > CaSO ₂ ·1/2H ₂ O +CaSO ₄ ·2H ₂ O		Oxidation Anywhere > 0		Ca(OH)2 Dissolution in Hold Tank and Marble Bed		in Error in Total Sulfued > 0 Material Balance > 1	
	Set #1	Set #2	Set #1	Set #2	Set #1	Set #2	Set #1	Set #2	Set #1	Set #2
17R	X	X	X	X	X	X	x	x	X	*
(17R)	X	X	X	X	x	X	X	x	X	*
18R	X	X	x	X	x	x	X	x	X	X
19R	X	X	x	X	*	*	X	x	X	X
(19R)	X	x	X	X	x	x	X	x	X	X
20R	X	X	x	X	X	X	X	x	X	X
(20R)	X	X	x	X	x	X	x	x	X	X
21R	X	X	x	X	x	X	*	*	-	-
22R	*	-	X	-	X	-	X	-	-	-

KEY: () Calculated by assuming all the oxidation occurs in marble bed
X Good
* Bad_

⁻ No Data

The parentheses indicate that the rate calculations for the high solid slurry experiments were determined assuming all the oxidation occurred in the marble bed.

Table 3-3a and 3-3b summarize the rates of $CaSO_3^{-1}/2H_2^{-0}$ and $CaSO_4^{-2}H_2^{-0}$ precipitation, sulfite oxidation and $Ca(OH)_2$ dissolution. In the low slurry solid concentration experiments, most of the $CaSO_3^{-1}/2H_2^{-0}$ precipitation occurred in the hold tank, while in the high solid concentration experiments, most of the calcium sulfite precipitated in the marble bed. Calcium sulfate precipitation rate data were inconsistent, and therefore it was almost impossible to detect the trend and location of its precipitation. The dissolution rate of $Ca(OH)_2$ in the marble bed was always greater than 50 percent of the total dissolution except in experiment 19R, where it was 30 percent.

This leads to the conclusion that most of the additive dissolves in the marble bed. It is important however, to point out that the percent of total dissolution in the marble bed should be controlled so that calcium sulfite scaling will not occur.

It should also be noted that in most experiments the rate of $CaCO_3$ formation was negligible in both the marble bed and the hold tank. In addition, out of the total SO_2 absorbed in the system, the fraction that underwent oxidation was higher (40-55 percent) in the low solid slurry experiments than in the high solid slurry experiments (25-35 percent).

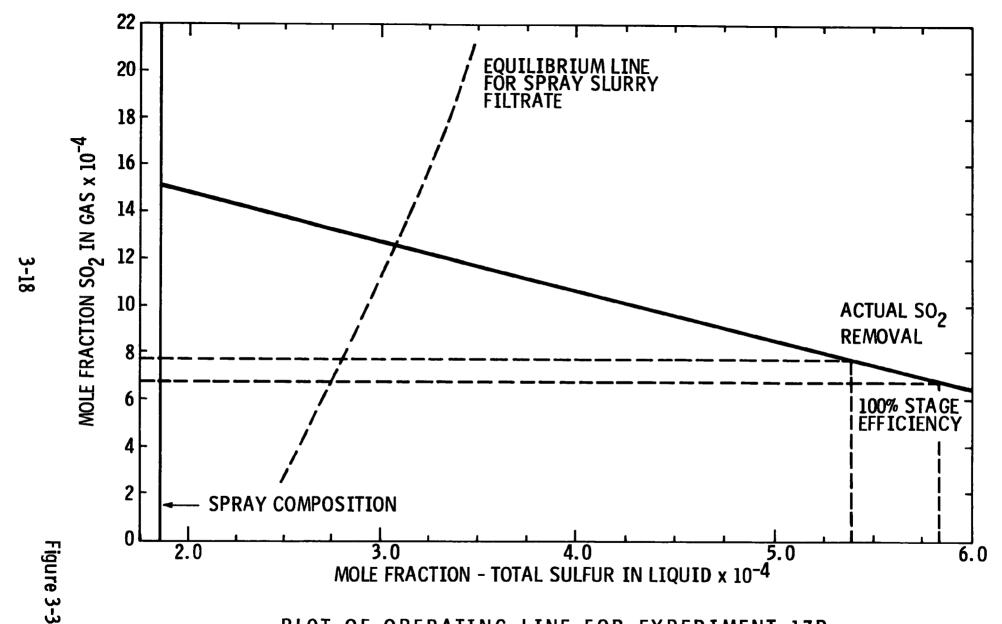
3.3.7 <u>Calculation of Additive Dissolution Rate From Equilibrium Data</u>

Based on the results of the stage efficiency calculations for the soluble system tests, additive dissolution rates were determined for the furnace injection tests by using vapor-liquid equilibria. Two major assumptions were made in these calculations:

- (1) Stage efficiency remained constant at 90% (from the soluble system tests).
- (2) The marble bed operated as a well mixed reactor and additive dissolved at a fixed rate to maintain a constant partial pressure of SO_2 exerted by the liquid in the bed.

Figure 3-3 is an operating and equilibrium line diagram of experiment 17R. The equilibrium line on the diagram represents the soluble portion of the scrubber feed. The abscissa is constructed so that the total sulfur concentration in the liquid entering the scrubber is represented by the vertical line at the far left of the graph. The point on the abscissa corresponding to the actual SO₂ removal is the total sulfur concentration in the liquid leaving the scrubber. This concentration does not exist in the actual data taken but is equal to the weighted average of the downcomers and scrubber bottoms concentrations assuming no precipitation of sulfur compounds in the scrubber.

If no additive dissolution had occurred, the SO_2 outlet concentration could have been no lower than 1260 ppm as represented by the intersection of the operating and equilibrium lines. Since the actual SO_2 outlet concentration obtained during the test was 770 ppm, some additive dissolution had to occur. Assuming the stage to be 90% efficient for 770 ppm outlet SO_2 , the outlet SO_2 equivalent to 100 percent stage efficiency would be 680 ppm (according to the previously stated assumption, the rate of additive dissolution should be sufficient to maintain 680 ppm SO_2 partial pressure over the liquid). Even if the stage efficiency is less than 100 percent, the rate of additive dissolution would be the same. In order to calculate the dissolution, the simplest method would be to determine what quantity was necessary to maintain 680 ppm SO_2 over the liquid if SO_2 removal equal to 100 percent stage efficiency was obtained.



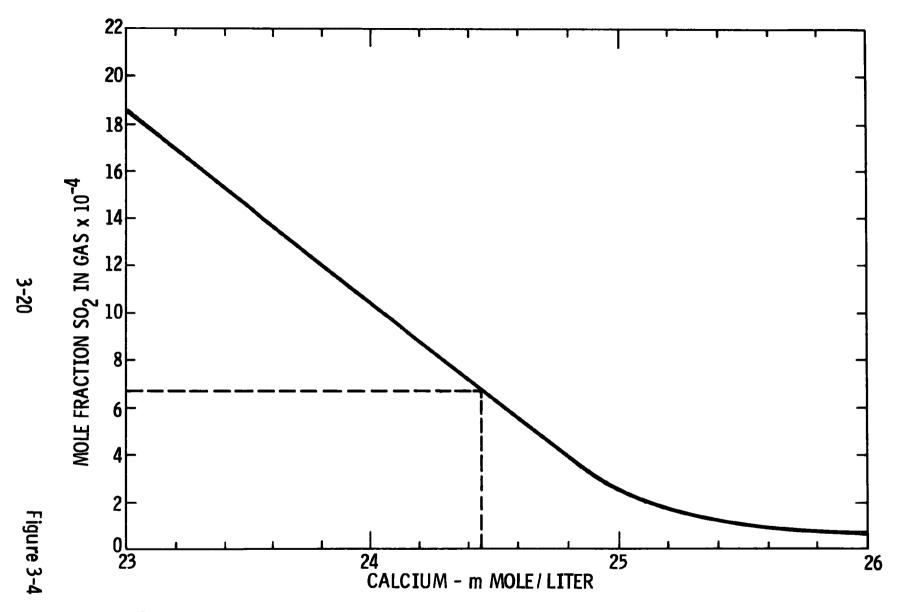
PLOT OF OPERATING LINE FOR EXPERIMENT 17R

To quantitatively determine the additive dissolution, varying amounts of calcium were input to the computer equilibrium program along with most of the spray water composition. Only soluble carbon dioxide values were taken from the downcomer analysis, and sulfite and sulfate were those derived from Figure 3-3 for 100% stage efficiency. Total system oxidation was used to determine the ratio of sulfate and sulfite at this point. Partial pressure of SO_2 over the liquid for varying amounts of soluble calcium is plotted for experiment 17R in Figure 3-4. The concentration of calcium producing a partial pressure of 680 ppm SO_2 was found to be 24.45 $\frac{\mathrm{m}\ \mathrm{moles}}{\mathrm{liter}}$. By subtracting the amount of soluble calcium entering the scrubber from this value and multiplying the difference by the flow rate, a dissolution rate of 4950 $\frac{\mathrm{m}\ \mathrm{moles}}{\mathrm{min}}$ was obtained. This procedure was carried out for experiments 18R to 22R; the graphs are presented in Appendix I. Table 3-5 contains a summary of these results along with the results obtained by material balance methods.

Agreement of the dissolution rates obtained by equilibrium data with those done by liquid species material balance is reasonable in most cases. Experiments 18R and 22R exhibit the most deviation. Comparing the dissolution rates to the SO_2 absorption rates indicates that additive dissolution is responsible for an average of two-thirds of the SO_2 removal.

3.3.8 Supersaturation of Calcium Sulfate and Sulfite

Using the soluble chemical analyses from marble bed samples and the equilibrium computer program, supersaturation values of calcium sulfate (ratio of the activity product to solubility product) were calculated for experiments 17R through 22R. Table 3-6 contains these results along with an indication of any calcium sulfate scaling which occurred during the tests. A supersaturation value of approximately 1.3 appears to be the threshold for calcium sulfate scaling in these tests.



PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 17R

TABLE 3-5. SUMMARY OF CALCIUM HYDROXIDE DISSOLUTION CALCULATIONS FOR MARBLE BED

All values are in $\frac{M \ Moles}{Min.}$

Experiment No.	Set No.	Liquid Species Material Balance Using Assumed Oxidation*	Solids Species Material Balance	Results From Equilibrium Diagrams	Amount of SO ₂ absorbed
17R	1 2	5000 5400	2200 2700	5000	7100 7100
18R	1 2	6100 6900		2800	10700 10700
19R	1 2	3000 7800	300 2000	5200	5800 5400
20R	1 2	6100 2600	3900 3700	4200	6600 6400
21R	1 2	5200 9400		8200	13900 14700
22R	1	14700		8800	16800

^{*} Oxidation value obtained for entire system was assumed to occur only in marble bed.

TABLE 3-6. COMPARISON OF CALCIUM SULFATE SUPERSATURATION AND SCRUBBER PERFORMANCE

Test No.	Relative Supersaturation Of Calcium Sulfate*	Did Calcium Sulfate Scaling Occur?
17R	0.97	No
18R	1.33	Yes
19R	1.84	Yes
20R	2.03	Yes
21R	1.18	No
22R	1.32	No

Values of supersaturation for calcium sulfite in the marble bed were not calculated because of the large fluctuations with pH. Although a supersaturation value can be obtained for the marble bed samples, this probably does not represent the actual value in the bed itself. Any dissolution of additive while the sample was being taken would raise the pH and increase the calcium sulfite supersaturation.

3.4 CONCLUSIONS

In summary, the following conclusions can be drawn from the limestone furnace injection system tests.

- (1) SO₂ removal can be improved significantly (8 to 10 percentage points) by increasing the liquid to gas ratio from approximately 20 to 35 GPM/1000 CFM.
- (2) SO_2 removal can be improved significantly by increasing the solids concentration in the spray slurry from 0 to 3.5%.
- (3) No improvement in SO_2 removal could be obtained by increasing the solids in the spray slurry beyond 3.5%.
- (4) Calcium sulfate scaling was controlled in a closed loop system without employing liquid blowdown by maintaining 8% total solids (30%-60% flyash) in the spray slurry. (Based on a maximum continuous run time of 10 hours).
- (5) Calcium sulfate scaling can be controlled by maintaining the supersaturation level below 1.3.
- (6) Calcium sulfite scaling in the scrubber can be controlled by maintaining the spray slurry pH below 11 to insure that no CaO or $Ca(OH)_2$ solids enter the scrubber.
- (7) More than half the additive dissolves in the scrubber bed in spite of the short residence time in the bed.

SECTION 4

LIMESTONE TAIL-END SYSTEM TESTS

4.1 SYSTEM CHEMISTRY

The process of removing SO₂ from the flue gas using limestone in the limestone tail-end system consists of the following reactions:

$$CaCO_3 + CO_2 + H_2O \longrightarrow Ca(HCO_3)_2$$
 (1)

$$2SO_2 + Ca(HCO_3)_2 \rightarrow Ca(HSO_3)_2 + 2CO_2$$
 (2)

$$CaSO_3 + SO_2 + H_2O \rightarrow Ca(HSO_3)_2$$
 (3)

$$Ca(HSO3)2 + CaCO3 \longrightarrow CaSO3 + Ca(HCO3)2$$
 (4)

$$CaSO_3 + 1/2 O_2 \longrightarrow CaSO_4$$
 (5)

Reactions 1, 2 and 3 are the principal absorption reactions. Sulfur dioxide reacts with relatively soluble bicarbonate to form calcium bisulfite. In addition, solid calcium sulfite recycled from the reaction tank or hold tank reacts with SO_2 to form calcium bisulfite.

The reactions in which sulfite is oxidized to sulfate (reaction 5) and soluble bisulfite is converted to insoluble calcium sulfite (reaction 4) account for the waste products as well as the regeneration of the solid calcium sulfite reactant that is recirculated to the scrubber. The ratio of calcium sulfite to calcium sulfate found in the APCS solid waste depends upon the extent to which these reactions go to completion.

4.2 TEST DESCRIPTION

The C-E scrubber at EPA's alkali Scrubbing Test Facility (at TVA's Shawnee steam plant) has only one marble bed, while current commercially offered C-E scrubbers have two marble beds. On the other hand, the other two scrubbers at the facility are similar to current commercially offered designs. Therefore, it was felt that comparing the

Performance of the C-E scrubber with the other two test scrubbers at Shawnee might be difficult. Therefore, the one and two marble bed tailend limestone tests using the KDL prototype were designed to assist EPA in extrapolating the single marble bed results from Shawnee to predict the performance of the C-E scrubber with two marble beds. The tests also provided data for determining the solid-liquid mass transfer rates in the scrubber and the hold tanks. In addition, the following information was sought which would greatly assist in designing a limestone tail-end system for scrubbing SO_2 :

- (1) Rate of dissolution of limestone $(CaCO_3)$. This would determine the alkalinity in the scrubber bed and the proper size of the reaction tank. The rate can be determined either using a material balance or using equilibrium methods.
- (2) Rate of precipitation of calcium sulfate and calcium sulfite. The rates would assist in designing the reaction tank such that the exiting stream will be close enough to saturation to prevent calcium sulfate scaling in the scrubber.
- (3) Rate of oxidation to sulfate. This would determine the incremental increase in supersaturation of calcium sulfate in the marble bed and therefore would determine both the liquid to gas ratio (L/G) and the limit on supersaturation entering the scrubber needed to prevent calcium sulfate scaling in the scrubber system.

Contrary to the furnace injection tests where the additive was introduced into the scrubber along with the flue gas, the additive in the tail-end system was introduced into the hold tank as shown in Figure 4-1. Liquid SO_2 was vaporized using steam and then injected into the flue gas (FG) in order to increase its SO_2 concentration to about 0.25 mole percent. The slurry (SL) from the marble beds' turbulent layers (pH of

5-5.5) left the scrubber through the overflow pots into the downcomers and was then discharged into the hold tank (10 feet diameter by 10 feet high). The scrubber bottom slurry (SB) which was rejected spray water was also discharged into the hold tank. The hold tank provided good solid liquid contacting and allowed for the limestone dissolution. The test conditions for the six tests are listed in Table 4-1 and the operating data is given in Table J-1 in Appendix J.

In experiments 25R, 26R and 27R, only one marble bed was used, while in experiments 28R, 29R and 30R, two marble beds were used. Most of the slurry leaving the hold tank was introduced under the marble bed through the spray nozzles. The spray water pH varied between 6.0 and 6.5. The marble bed together with the turbulent layer the volume of which is about 20 cubic feet, provided good mixing where the absorption of S02 took place. A portion of the hold tank effluent was pumped to the clarifier to maintain about 8 percent solid concentration in the slurry. The solids were settled in the clarifier and the clarified liquid (CL) was returned to the hold tank. The hold tank was maintained at full capacity of about 6000 gallons at all times. The reaction tank residence time at a 500 gpm pumping rate is about 12 minutes.

To determine when the system reached steady state, samples taken from the spray water (SW) and the clarifier liquid (CL) were analyzed for calcium, sulfite and sulfate. Steady state in these tests is defined as the point when the calcium and total sulfur concentration in the filtrate of the clarified liquid (CL) and the spray water (SW) were reasonably close. The analyses of the samples used to determine steady state are shown in Table J-2 in Appendix J.

TABLE 4-1. TEST PARAMETERS FOR THE LIMESTONE TAIL-END SYSTEM

Experiment No.	25R	26R	27R	28R	29R	30R
Number of beds	1	1	1	2	2	2
Gas Flow Rate						
ACFM @ 120°F	10,000	10,000	10,000	10,000	10,000	10,000
ft/Min	450	450	450	450	450	450
Inlet SO ₂ , PPM	2,400	2,400	2,400	2,400	2,400	2,400
Limestone Feed Rate	100	150	150	150	150	100
(% Stoichiometry)						
Fly Ash Feed Rate (Gr/SCFM)	0	0	0	0	0	0
Underbed Slurry						
GPM/bed	250	250	150	150	250	250
L/G, GPM/1000 CFM	25	25	15	15	25	25
Overbed Spray, GPM/bed	0	0	0	0	0	0
Solid Conc. in Slurry, wt. %	8	8	8	8	8	8

4.3 DATA EVALUATION

4.3.1 System Performance

The results of the tail-end tests are summarized in Table 4-2. While holding all other conditions the same in single bed experiments 26R and 27R, the SO_2 removal efficiency increased by 8 percentage points when the liquid to gas ratio (L/G) was increased from 15 to 25 GPM/1000 CFM. In double bed experiments 28R and 29R, the overall SO_2 removal increased by 11 percentage points when the L/G was increased from 15 to 25 GPM/1000 CFM per bed. It can be concluded therefore, that the liquid to gas ratio has a significant effect on the SO_2 removal efficiency.

The overall SO_2 removal efficiency remained unchanged in both single and double marble bed tests while the additive feed rate was increased from about 100 to 150 percent stoichiometry with all the other conditions kept the same. This leads to the conclusion that in high solids systems when the additive feed rate is increased beyond 100 percent, the SO_2 removal efficiency remains unchanged and as a result, the additive utilization tends to decrease.

While the solid concentration in the slurry in all tailend experiments was maintained between 6.5 and 8.5 percent, no calcium
sulfite or calcium sulfate scaling was observed. Liquid blowdown was not
used during these tests to control calcium sulfate scaling. Therefore,
it can be concluded that calcium sulfate scaling in a limestone tail-end
system can be controlled by maintaining a solid concentration in the slurry
of about 8 percent (excluding fly ash).

The SO_2 removal efficiency in the upper and the lower marble beds based on the SO_2 concentrations in the gas entering the respective marble beds, was the same. For example, in experiment 27R, the lower marble bed removed about 50 percent of about 2400 ppm SO_2 entering

TABLE 4-2. SUMMARY OF LIMESTONE TAIL-END TESTS

PERFORMANCE RESULTS

Experiment No.	25R	26R	27R	28R	29R	30R
No. of Marble Beds Used	1	1	1	2	2	2
L/G (GPM/1000 CFM) - Lower Bed	24.5	24.0	15.0	16.0	24.5	25.0
L/G (GPM/1000 CFM) - Upper Bed	-	-	-	15.0	22.5	23.5
Solids in Spray water (%)	7.4	6.5	7.5	6.7	8.6	-
Inlet SO ₂ (PPM)	2345	2505	2315	2420	2435	2380
Solid Recycle - (%)	96	94	92	95	97	96
Stoichiometry (%)	98	145	156	152	148	94
SO ₂ removal efficiency (%)*	55.5	57	49	76	87	86
Additive Utilization - %	57	39	31	50	59	91
Calcium Sulfite Scale	None	None	None	None	None	None
Calcium Sulfate Scale	None	None	None	None	None	None
Liquid Blowdown	None	None	None	None	None	None

The gas flow was maintained at about 10,000 CFM @ 130°F, 1 atm in all tests.

^{*}Corrected for Air leakage

it, and in experiment 28R, both marble beds removed about 75 percent of the SO_2 (2400 ppm) entering the lower bed. This means that the lower bed removed 50% of the 2400 ppm SO_2 entering it while the upper bed removed another 50% of the remaining 1200 ppm SO_2 entering the upper bed. This resulted in an overall removal of 75%. Therefore, it can be concluded that single marble bed SO_2 removal efficiency for the range of inlet SO_2 concentration studied can be extrapolated to predict two marble bed SO_2 removal efficiency.

4.3.2 Analytical Results and Sampling Methods

The furnace injection sampling method was also used in the tail-end tests, and the corresponding samples from the upper marble bed were added to the list of samples used in the furnace injection tests. C-E, however, did not perform chemical analysis on any of the tail-end test series in order not to duplicate Radian Corporation's effort. The results of the solid analyses are listed in Tables J-3 through J-8, and those of the liquid analysis are listed in Tables J-9 through J-14, in Appendix J. In their July, 1972 Progress Report to EPA, Radian indicated a significant error in the liquid sulfite analysis in experiment 25R. This caused an error in the sulfate results, since sulfate is obtained by the difference between total sulfur and sulfite. The remainder of the analyses were fairly accurate except for the marble bed sulfite and sulfate results which are slightly in error due to a relatively large residence time in the sampling lines thus allowing further time for reaction and oxidation. The samples for obtaining solid concentration in the slurry in experiment 30R were accidentally discarded, thus preventing total sulfur material balance calculation for that test.

4.3.3 Total Sulfur Material Balance

Detailed calculations of the total sulfur material balance of the limestone tail-end experiments are summarized in Tables K-1 through K-5 in Appendix K, and the results are summarized in Tables 4-3a and 4-3b. The results show low errors considering the analytical and sampling problems, and the inaccurancy in sampling for high solid concentration in the slurry. In arriving at the amount of total sulfur in the slurry streams, the specific gravity of the slurry used was assumed to be 1.0 in order to convert the slurry flow rate from GPM to pounds per minute.

4.3.4 Rate Calculations

Since the solid concentration in all of the limestone tailend tests was high, a material balance calculation to determine the rates was not feasible without making certain assumptions. In the case of high solid slurries, a difference in a rate resulting from a slight change in solid concentration was masked by that resulting from the error in the solid concentration measurement. Therefore, in order to obtain the rates of precipitation and dissolution, the following assumptions and observations were used:

- (1) All of the oxidation in the system occurred in the marble bed. (From experiment 17R, furnace injection.)
- (2) Total oxidation in the system is the ratio of the total sulfate to total sulfur in both the solid and liquid streams leaving the systems.

Oxidation = $\frac{(SO_4^{-}) \text{ Liquid + } (SO_4^{-}) \text{ Solid}}{\text{Total Sulfur}}$

(3) The amount of CO₂ transferred to the hold tank from the atmosphere is negligible.

TABLE 4-3a. SUMMARY OF RATE RESULTS FROM LIMESTONE TAIL END TESTS

			Experime				Experime				Experime		
		Marbl	e Bed	<u>Syst. R</u>	emainder	Marbl	e Bed	Syst. R	emainder	Marbl	e Bed	Syst. Re	mainder
	Location	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>
	Inlet Blue Gas Flow (CFM @130°F)	9,950	9,900			10,060	10,100		10,250	10,180			
	Liquid to Gas Ratio - Upper per Bed	-	-	-	-	-	-	-	-	-	-	-	-
	(GPM/1000 CFM) - Lower	24.1	25.2			24,3	23.3		14.6	14.7			
	Additive Feed Rate (M Moles/Min)			24,062	24,062			38,590	38,590			39,044	39,044
	Stoichiometry based on inlet $S0_2(%)$	98.5	97.6			144.5	.46.1		156.7	156.6			
	SO ₂ Removal Eff. (%)	56.2	54.7			56.5	57.0		48.2	49.4			
	Solid Conc. in Spray Slurry (wt. %)	7.55	-			-	6.57		7.18	7.69			
4-10	ΔS ₆ (Amount of SO ₂ absorbed) (M Moles/Min)	12,770	13,501			14,081	11,911		11,237	11,438			
	Sulfite Oxid. (%)	25.0	23.4			26.1	23.8		26.4	24.9			
	Sulfite Oxidation (M Moles/Min)	3,192	3,159			3,675	2,834		2,966	2,848			
	CaSO ₃ ·1/2 H ₂ O formation (M Moles/Min)	745		10,497	10,127	-3,122	-1,146	13,769	13,098	-2,907	247	11,279	8,255
	CaSO ₄ ·2 H ₂ O formation (M Moles/Min)	4,997		1,233	-2,201	2,898	-274	1,568	3,000	3,160	885	-742	1,854
	CaCO ₃ dissolution (M Moles/Min)	4,832		5,586	4,281	6,769	5,413	6,801	7,647	5,092	6,954	5,937	7,565
	Error in Total Sulfur material balance												
	$(\frac{In - Out}{In}) \times 100$	-3.1		-0.2	-0.5		-20.7		3.7	-9.2	-13.1	-3.3	-1.9

^{*}System remainder includes hold tank, surge tanks and clarifier.

TABLE 4-3b. SUMMARY OF RATE RESULTS FROM LIMESTONE TAIL END TESTS

		Experiment 28R					Experime	nt 29R			Experime	nt 30R	
		Harb1	e Bed	Syst. R	emainder*	Marble	e_Bed	Syst. R	emainder*	Marbl	e_Bed	Syst. Rer	
	Location	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	<u>Set #2</u>	<u>Set #1</u>	Set #2	<u>Set #1</u>	<u>Set #2</u>	Set #1	<u>Set #2</u>
	Inlet Flue Gas Flow (CFM @ 130°F)	10,160	10,400			10,200	10,400		10,280	10,280			
	Liquid to Gas Ratio - Upper per Bed (GPM/1000 CFM) - Lower	14.8 15.5	14.4 15.3			22.0 24.0	21.6 23.5		22.8 24.3	22.8 24.3			
	Additive Feed Rate (M Moles/Min)			39,044	39,044			39,044	39,044			24,213	24,213
	Stoichiometry based on inlet $SO_2(%)$	152.4	151.7			147.8	147.5		97.1	90.0			
	SO ₂ Removal Eff. (%)	75.5	76.0			87.1	87.6		84.4	87.3			
	Solid Conc. in Spray Slurry (wt.%)	6.40	6.71			8.41	8.72		-	-			
4-11	ΔS _G (Amount of SO ₂ absorbed) (M Moles/Min)	17,002	18,057			21,489	21,050		19,643	20,463			
	Sulfite Oxid. (%)	24.4	25.9			27.9	28.4		30.0	30.5			
	Sulfite Oxidation (M Moles/Min)	4,150	4,680			5,995	5,978		5,893	6,241			
	CaSO ₃ ·1/2 H ₂ O formation (M Moles/Min)	1,319	10	13,009	12,991	2,825	3,076	11,687	11,562	-3,306	-5,086	23,284	19,592
	CaSO ₄ ·2 H ₂ O formation (M Moles/Min)	1,430	1,721	3,695	3,346	-676	-262	6,839	6,373	161	1,120	6,154	2,205
	CaCO ₃ dissolution (M Moles/Min)	9,503	10,102	7,893	7,744	12,601	12,152	7,782	9,124	9,861	9,394	15,975	8.199
	Error in Total Sulfur material balance $(\frac{\text{In} - \text{Out}}{\text{In}}) \times 100$	-14.8	-3.9	-5.6	2.2	-16.0	-20.9	-31.2	-4.5				

^{*}System remainder includes hold tank, surge tanks and clarifier

A summary of the rate calculations are shown in Tables K-6 through K-11 in Appendix K. Tables 4-3a and 4-3b summarize the rate results obtained from the limestone tail-end tests. In order to determine the reliability of rate data from these experiments, the following criteria were used:

- (1) The rate of SO_2 removal from the gas should always be greater than the sum of calcium sulfite and calcium sulfate precipitation rates in the system (marble bed and system remainder).
- (2) The rate of SO_2 removal from the gas should be equal to the rate of limestone dissolution in the system (marble bed and system remainder).
- (3) The sulfite analysis should be reliable (discussed in section 4.3.2).

Table 4-4 lists these criteria for the limestone tail-end experiments and indicates whether or not each criterion is satisfied. The following tests were considered reliable:

26R(1), 27R(1, 2), 28R(2), 29R(1, 2), 30R(2).

The parentheses indicate the set number. The negative calcium sulfite and calcium sulfate precipitation rates in the marble bed and the hold tank could be attributed to one or both of the following:

- (1) Errors resulting from the liquid sulfite analysis and/or liquid sampling techniques.
- (2) The assumption that all of the oxidation in the system occurred in the marble bed may not be true. The calculation of the rate of limestone dissolution, however, should not be affected by the calcium sulfite and calcium sulfate precipitation rates, since the rate of limestone dissolution depends on the total sulfur analysis which in this case is relatively accurate.

TABLE 4-4. CRITERIA FOR DETERMINATION RELIABILITY OF LIMESTONE TAIL-END TEST

Test No.	ΔSG> CaSO ₃ .1/2H ₂ O) + CaSO ₄ .2H ₂ O	$\Delta S_G = CaCO_3$	Dissolution	Sulfite	Analysis
	Set 1	Set 2	Set 1	Set 2	Set 1	Set 2
25R	*	-	*	-	*	X
26R	X	*	X	X	X	X
27R	X	X	X	X	X	X
28R	*	X	X	X	X	X
29R	X	X	X	X	χ -	X
30R	*	X	*	X	X	X

X - Good

^{* -} Bad

^{- -} No Data

In all the limestone tail-end experiments, most of the calcium sulfite precipitation occurred in the hold tank. The fraction of total precipitation occurring in the marble bed could not be determined due to the error involved in the determination of sulfite concentration in the liquid.

Most of the calcium sulfate precipitation occurred in the hold tank when two marble beds were used. However, when one marble bed was used, the calculated precipitation rates fluctuated appreciably. This inconsistency prevented drawing any conclusions regarding where most of the calcium sulfate precipitation took place. The calcium sulfate precipitation rates, like the calcium sulfite precipitation rates, were in error for the reasons discussed above.

During the experiments with one marble bed, the rates of limestone dissolution in the marble bed and the hold tank were approximately equal. The dissolution rates in the marble bed basically did not change with the liquid to gas ratio. When two marble beds were employed, the fractional rate of dissolution of limestone in the marble bed increased from 50 percent to about 60 percent. This was expected since it was accompanied by an increase in SO₂ removal efficiency.

4.3.5 <u>Calculation of Additive Dissolution Rate From Equilibrium Data</u>

The dissolution rate of calcium carbonate in the marble bed was calculated from an equilibrium approach in the same manner as described in the furnace injection tests. For the first single bed calcium carbonate test, 25R, the operating line is plotted in Figure 4-2. As was determined in the soluble system tests, a stage efficiency of 90% was used in making dissolution rate calculations. The amount of calcium needed in solution in the marble bed to maintain the partial pressure corresponding to 100% stage

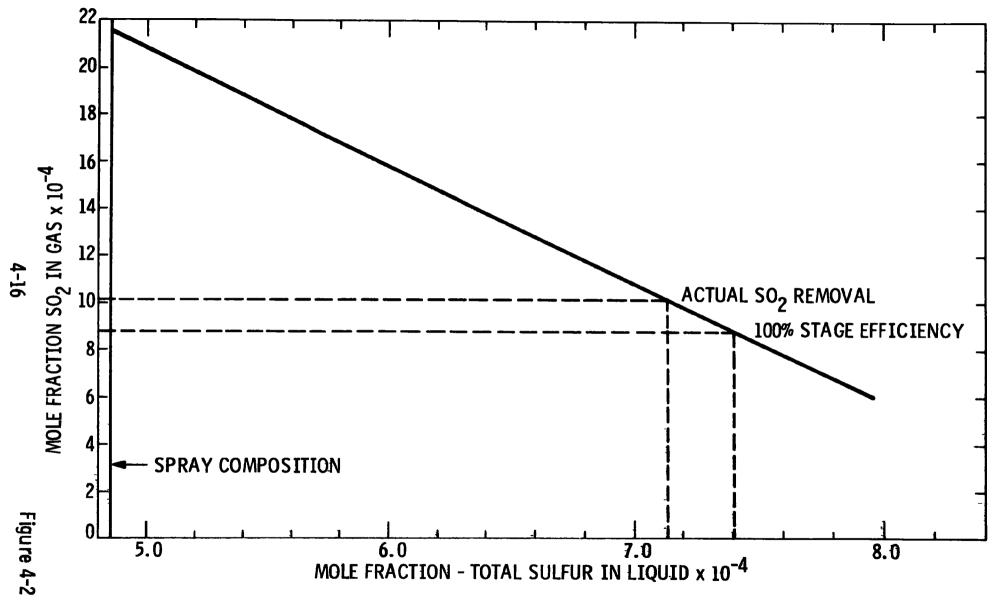
efficiency was determined for experiment 25R from Figure 4-3.* To calculate the dissolution rate, the concentration of soluble calcium in the scrubber feed was subtracted from the calcium concentration determined above and the difference was multiplied by the inlet liquor flow rate.

These results are summarized in Table 4-5.

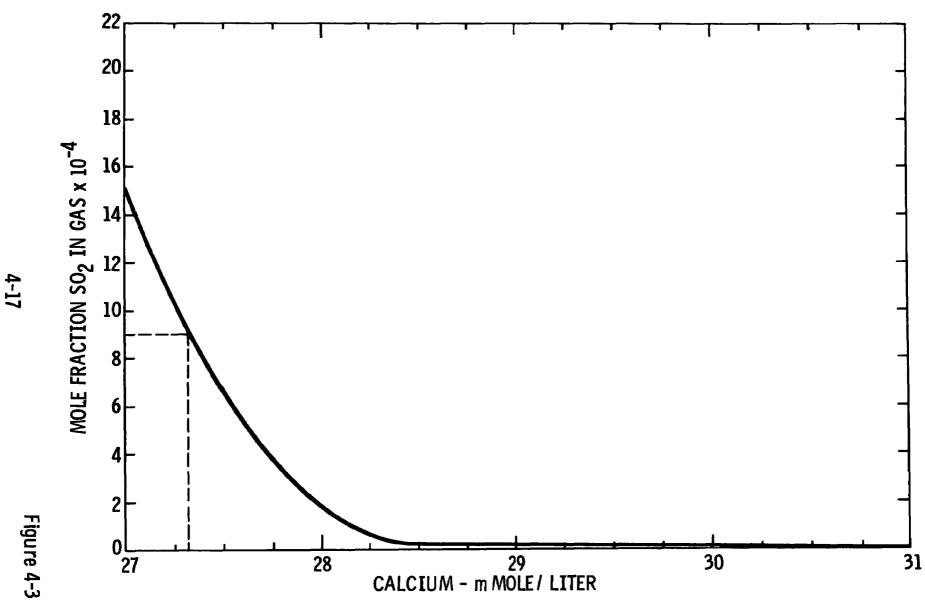
Calculation of dissolution rates for the tests with two marble beds presented one major problem: no measured sulfur dioxide concentration between the first and second beds. If liquid analyses were used, corrections would have been necessary to account for amounts of calcium sulfite and sulfate precipitated in the scrubber or in the lines before the sample was taken. Since gas-side analysis is needed to determine the amounts of calcium sulfate and sulfite precipitated, a value for the amount of sulfur dioxide removed in the first bed could not be obtained from liquid analyses either. For these reasons, assumptions for the amount of sulfur dioxide removed in the first bed were made based on results from the single bed tests. These values were correlated to the single bed tests according to the L/G in each test; included in the L/G for the bottom bed in the two bed tests was the amount of slurry rejected from the top bed. These calculated sulfur dioxide values are presented in Table 4-6.

Figure 4-4 is the operating line diagram for the upper bed in experiment 28R; the diagram for the determination of calcium concentration is shown in Figure 4-5. Appendix L contains the diagrams for tests 29R and 30R. The rates for dissolution of calcium carbonate for the upper bed are listed in Table 4-5 along with a total dissolution rate for both beds using average dissolution values from the single-bed tests. Calcium carbonate dissolution values by material balance are also listed in Table 4-5.

^{*}All diagrams for experiments 26R and 27R are given in Appendix L.



PLOT OF OPERATING LINE FOR EXPERIMENT 25R



PLOT OF CALCIUM VS PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 25R

TABLE 4-5. SUMMARY OF CALCIUM CARBONATE DISSOLUTION CALCULATIONS

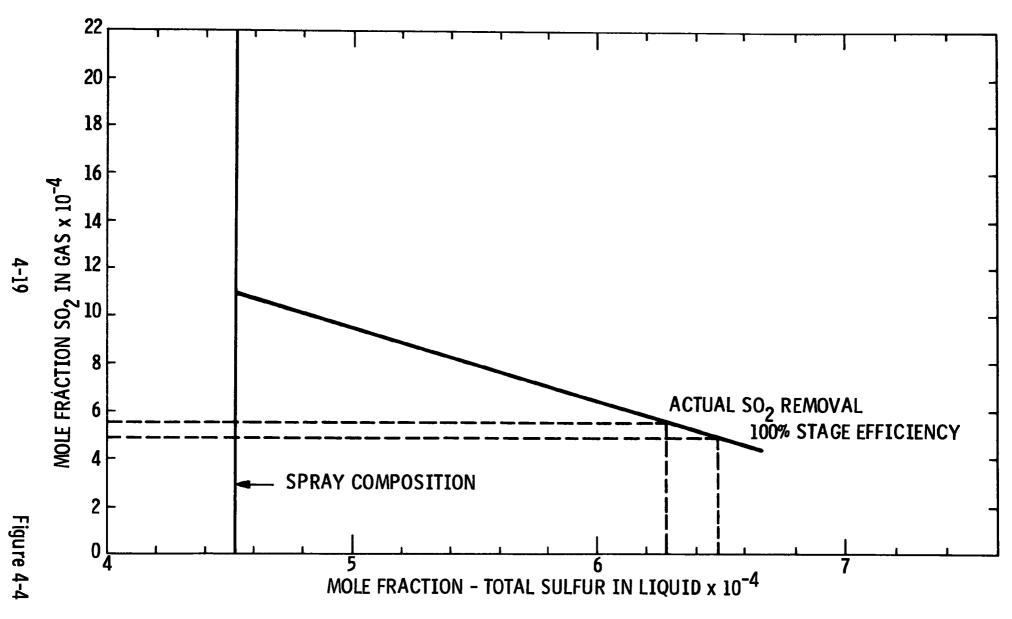
All values are in $\frac{M \text{ Moles}}{Min.}$

	Rate From Eg	uilibrium !	Method	Rate From	<u>Material Bala</u>					
Experiment	Bottom Bed	Top Bed	Total for Both Beds	Bottom Bed	Total for Both Beds	Amount of SO ₂ absorbed				
25R	3030			4832		12770				
26R	6880			6769		14081				
27R	5700			5092		11237				
28R	5700	2380	8080		9503	17002				
29R	6880	2520	9400		12601	21489				
30R	6880	3080	9960		9861	19643				

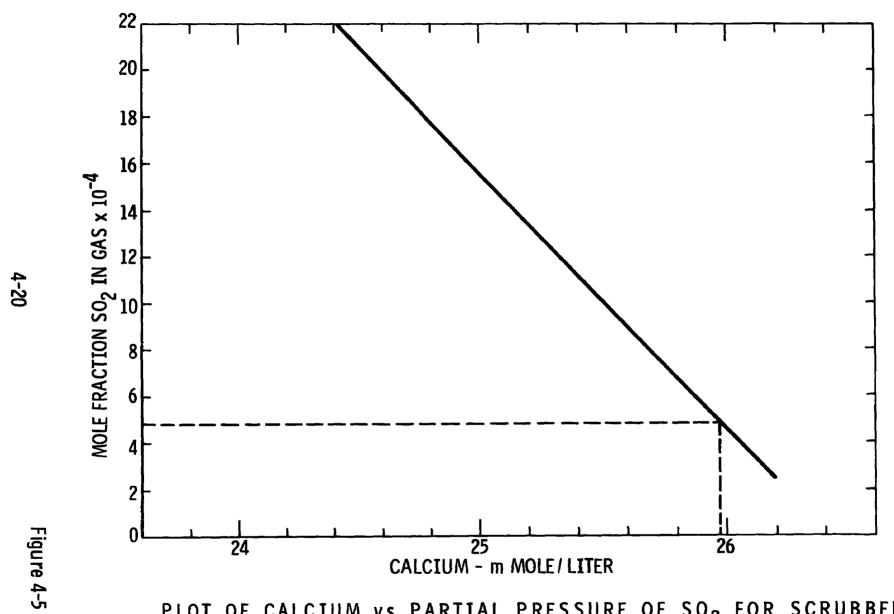
TABLE 4-6. ASSUMED ${
m SO}_2$ REMOVAL FOR LOWER BED IN TWO-BED CALCIUM CARBONATE TESTS

Experiment	28R	29R	30R
SO ₂ entering lower bed	2223	2248	2240
SO ₂ leaving lower bed	1091	943	940
SO ₂ leaving top bed	563	299	377

SO₂ values are corrected for humidity and air leakage.



PLOT OF OPERATING LINE FOR EXPERIMENT 28R



PLOT OF CALCIUM VS PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 28R

Overall, the agreement between the two methods of calculation is very good. Due to the inaccurate sulfite analyses in experiment 25R, the dissolution rates obtained are probably not reliable. In general, one half of the total system dissolution took place in the marble bed, indicating that half of the SO_2 removal was because of additive dissolution and half by the soluble alkali in the scrubber feed.

4.3.6 Supersaturation of Calcium Sulfate and Sulfite

Values for the degree of supersaturation of calcium sulfate in the marble bed were calculated for Tests 25R-30R. The soluble analyses from the marble bed samples were input to a computer equilibrium program to determine the activities of calcium and sulfates. Table 4-7 presents the results of these calculations.

No calcium sulfate scaling was observed for any of the six tests, although supersaturation values are higher than the threshold values of 1.3 obtained in the furnace injection tests. No definite explanation can be given at this time for the fact that supersaturation values as high as 1.7 did not cause calcium sulfate scaling in the calcium carbonate tests.

Values of supersaturation for calcium sulfite were not calculated for the same reasons as given in Section 3.3.8 (Furnace Injection Tests). No calcium sulfite scaling was observed during the tests.

4.4 <u>CONCLUSIONS</u>

The following conclusions can be drawn from the tail-end limestone system tests:

(1) The performance (SO₂ removal efficiency and scaling) of the scrubber with two marble beds can be predicted by extrapolating the single marble bed test results of the C-E scrubber at Shawnee (EPA

TABLE 4-7. CALCIUM SULFATE SUPERSATURATION

Experiment No.	Bed	Temp(°C)	Relative S	upersaturation
				RADIAN*
25R	Bottom	49	0.92	0.87
26R	Bottom	48	1.62	1.51
27R	Bottom	47	1.40	1.32
28R	Bottom	44	1.73	1.34
	Тор	47	1.70	1.44
29R	Bottom	46	1.53	1.40
	Тор	46	1.61	1.40
30R	Bottom	43	1.54	1.37
	Тор	47	1.57	1.37

^{*}Determined by Radian Corporation under EPA Contract 68-02-0023

test facility). The ${\rm SO}_2$ removal efficiencies of the lower and upper beds appear to be the same based on the ${\rm SO}_2$ concentrations entering the respective beds.

- (2) Above 100% stoichiometry, limestone feed rate has little or no effect on SO_2 removal efficiency in high solids systems.
- (3) Calcium sulfate scale can be controlled in a closed loop system without employing liquid blowdown by maintaining 8 to 10% solids (excluding flyash) in the spray slurry. (Based on a maximum continuous run time of 10 hours).
- (4) SO₂ removal can be improved significantly (8 to 10 percentage points) by increasing the liquid to gas ratio from approximately 15 to 25 GPM/1000 CFM.
- (5) Calcium sulfate scaling can be controlled by maintaining the supersaturation below 1.7.
- (6) More than half the additive dissolution occurs in the marble bed in spite of the short residence time in the bed.

APPENDIX A GAS FLOW CHECK

APPENDIX A

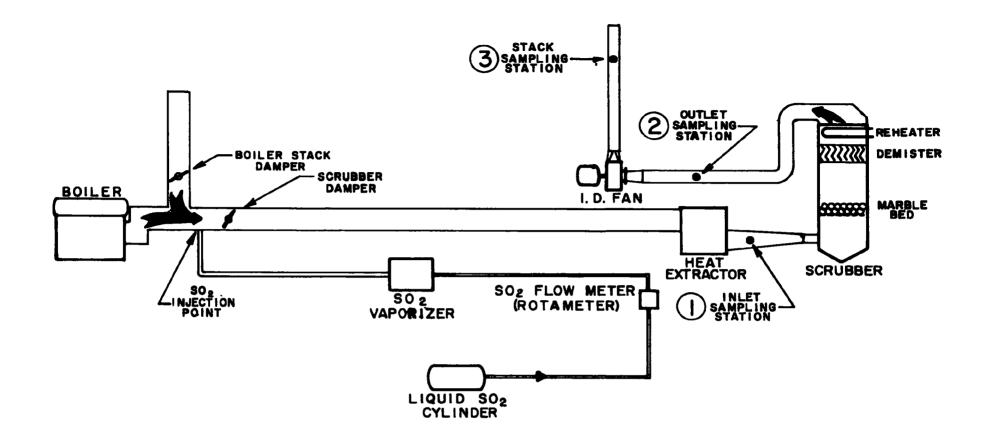
GAS FLOW CHECK

The gas flow check was divided into two parts.

- a. Determination of the duct coefficient which is defined as the ratio of the average velocity to the center line velocity.
- b. Comparison of gas flows obtained by using the pitot tube located at the center of the duct and the SO_2 tracer gas method.

The flow arrangement used for gas flow check is shown in Figure A-1. The gas flow was set at approximately 12,500 CFM at 130°F and 1 atmosphere. Six sets of pitot tube traverses were taken in two perpendicular directions using a standard pitot tube and the equal area method. These results are shown in Table A-1. The duct coefficient is defined as the ratio of the average velocity obtained by multiple point pitot tube traverse to the center line velocity. Once the duct coefficient is determined, the need for a multiple point velocity traverse is eliminated. The center line velocity obtained by using the pitot tube fixed at the center of the duct could be converted to the average velocity that could have been obtained with the multiple point pitot tube traverse by multiplying the center line velocity by the average duct coefficient. The average duct coefficient was found to be 0.932.

A measured quantity of gaseous $\rm SO_2$ was introduced into the duct right after the boiler and long before it gets to the scrubber. The $\rm SO_2$ concentrations at the inlet, outlet and stack sampling stations measured simultaneously with the center line velocity taken at the stack sampling station were used to calculate the gas flow rate and the leakage into the system. The center line velocity was converted to the average velocity of



CE AQCS PROTOTYPE - GAS FLOW CHECK

TABLE A-1. C-E APCS - GAS FLOW CHECK
DUCT COEFFICIENT* - VELOCITY TRAVERSE

Ditot	Set ! Velocity:	No. 1 , Ft/Sec.	Set I Velocity	io. 2 Ft/Sec.	Set I Velocity	
Pitot Tube Location	N-W Direction	S-W Direction	N-W Direction	S-W Direction	N-W Direction	S-W <u>Direction</u>
1	63.8	69.6	62.9	67.4	62.5	69.0
2	70.7	73.7	71.9	74.8	71.1	74.7
3	74.7	75.8	74.8	76.8	74.6	75.6
4	77.5	77.7	76.9	77.3	77.3	77.3
5	76.8	74.9	77.4	74.9	77.6	76.6
6	74.9	70.8	74.9	69.6	74.9	69.7
7	67.7	64.9	67.5	64.0	70.3	63.5
Average	71.7	71.8	71.9	71.6	71.8	71.7
Duct* Coefficient	0.925	0.925	0.935	0.925	0.93	0.927
	Set 1	No. 4			Set 1	No. 6
1	61.7	60.9	63.3	59.6	63.3	59.6
2	69.1	67.3	68.0	66.4	68.7	66.4
3	70.6	70.2	70.6	68.7	71.0	68.7
4	71.0	70.2	71.0	71.0	70.6	69.5
5	69.1	68.7	69.1	69. 1	68.7	68.4
6	66.4	65.3	66.9	65.7	66.1	65.7
7	61.3	60.5	61.3	59.6	59.6	61.3
Average	66.3	65.6	66.4	66.5	66.2	65.1
Duct* Coefficient	0.934	0.935	0.936	0.936	0.936	0.936

Average Duct Coefficient = 0.932

^{*}Duct Coefficient = Average velocity/center line velocity

the multiple point velocity traverse by multiplying it with the average duct coefficient. No liquid was introduced into the scrubber. A mixture of air and flue gas from the boiler, to which gaseous ${\rm SO}_2$ was added, passed through the scrubber. The boiler was operated on natural gas so that the SO_2 added to the duct at the ${\rm SO}_2$ injection point shown in Figure A-1 was the only source of ${\rm SO}_2$. The boiler operation was necessary to provide the steam needed to vaporize SO_2 in the SO_2 vaporizer. Liquid SO_2 taken from the SO_2 cylinder was measured using a rotameter before it reached the ${\rm SO}_2$ vaporizer. Manual SO_2 method was used to measure the SO_2 concentrations at the inlet, outlet and stack sampling stations, because the DuPont SO₂ analyzer cannot measure ${\rm SO}_2$ concentration at more than one location at any given time. The results shown in Table A-2 show that there is very good agreement in gas flow obtained using pitot tube and SO₂ tracer gas methods. Since the gas flow measured with the pitot tube checked very well with that obtained using ${\rm SO}_2$ as tracer gas, an independent method, it could be concluded that pitot tube is a reasonably good instrument to measure gas flow on the KDL prototype scrubber.

 ${
m SO}_2$ concentrations at the stack sampling station should be less than or equal to those at the outlet sampling station depending upon whether or not there is leakage into the system between the stack and the outlet sampling stations. Slightly higher ${
m SO}_2$ concentrations at the stack compared to the outlet shown in Table A-2 are within the accuracy limits (\pm 4.25%) of the manual ${
m SO}_2$ method. Since the leakage into the system given in Table A-2 is calculated from the ${
m SO}_2$ concentrations at the inlet, outlet and stack, the leakage between the inlet and outlet sampling stations appears to be slightly higher than the leakage between the inlet and stack sampling stations. The air leakage into the system was found to be 5 to 8 percent of the gas flow measured at the stack sampling station.

TABLE A-2. COMPARISON OF PITOT TUBE AND SO_2 TRACER GAS METHODS

Date of Run	10/8/71	10/8/71	10/11/71	11/1/71	11/11/71
Gas Flow, CFM @ 1 atm. and 130°F					
Center Line Velocity Converted to Multiple Point Pitot Tube Traverse using Duct Coefficient Measured at the Stack Sampling Station	11,950	11,900	11,510	11,900	12,285
SO ₂ as Tracer Gas - Measured at the Stack Sampling Station	12,150	12,010	11,540	-	-
SO ₂ as Tracer Gas - Measured at the Outlet Sampling Station	-	-	-	12,590	12,315
SO ₂ Concentration in the Gas, PPM					
Inlet Sampling Station	1,659	2,241	2,930	2,182	1,704
Outlet Sampling Station	1,548	2,122	2,710	2,013	1,561
Stack Sampling Station	1,581	2,121	2,770	-	-
Leakage into the System, % of Gas Flow Measured at the Stack Sampling Station					
Between Inlet and Stack Sampling Stations	5.0	5.3	5.5	-	-
Between Inlet and Outlet Sampling Stations	7.2	5. 3	7.5	7.8	8.4

APPENDIX B

SOLUBLE SYSTEM TEST DATA

AND RESULTS

APPENDIX B
SOLUBLE SYSTEM TEST DATA AND RESULTS

Experiment No. Date of Run		R 9/71	2R 10/27/71		
Set No.	1	2	1	2	
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SB Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SB Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Analyzer Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	10,960 Dilute	10,960 Dilute	10,750 Dilute 0 165 88 77 112 129 129 118 123 294 130 9 2,025 2,050 796 750 63.5 40.4 12.78	10,800 Dilute 0 165 85 80 111 129 129 118 123 292 130 9 2,045 2,050 830 750 63.5 40.6 12.57	
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	-6.0	-3.9	-3. 5	-3.1	
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet		- .7 .4	3	.9 .1 .0	
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet	12 11		12 12		
NO _X , ppm Inlet Outlet		-		-	

Experiment No. Date of Run		BR 4/71		4R 28/71
Set No.	1	2	1	2
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SL Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO2, ppm - Manual Inlet SO2, ppm - Analyzer Outlet SO2, ppm - Analyzer % SO2 Removal % Stoichiometry Sulfur removed from Flue Gas ΔSG, gmole/min. Sulfur absorbed by Na2CO3 ΔSL, gmole/min.	11,200 Dilute 170 0 160 15.0 52 105 102 112 91 318 108 9 2,215 2,095 880 860 59.0 35.6	11,200 Dilute 170 0 160 15.0 52 105 102 112 91 318 108 9 2,215 2,095 880 860 59.0 33.4 14.18	10,800 Dilute	10,800 Dilute 170 0 150 15.5 117 129 126 118 121 309 130 9 2,080 2,030 875 790 61.2 36.9 12.21
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	-3.9	-1.9	-0.9	+0.9
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet		- - -		3.8 3.6 5.1
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet		-		3.1 2.7
NO _X , ppm Inlet Outlet		<u>-</u>		<u>-</u>

Experiment No. Date of Run		R 2/71	6R 11/2/71		
Set No.	1	2	1	2	
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SL Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Analyzer Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	12,980 Dilute	12,980 Dilute	9,180 Dilute 110 555 135 29 115 125 122 121 303 127 9 - *	9,180 Dilute 110 55 135 24 115 125 125 122 121 304 127 9 2,110 2,050 450 480 76.5 52.3 14.06	
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	2.5	•	-	-1.6	
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet	3	3.6 - -	3	.8 .1 .5	
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet		:		.4 .8	
NO _x , ppm Inlet Outlet		143 159		62 70	

Experiment No. Date of Run		'R '3/71	8R 11/3/71		
Set No.	1	2	1	2	
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SL Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Manual Outlet SO ₂ , ppm - Analyzer Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	11,240 Dilute 152 69 185 34.5 112 120 122 116 123 291 122 9 1,815 2,000 310 450 77.5 51.3	11,240 Dilute 152 69 185 34.5 112 121 123 116 123 298 122 9 1,770 2,000 670 460 77.0 52.4 16.16	11,200 Dilute 73 36 93 15 109 121 124 116 123 312 125 9 *	11,190 Dilute 73 36 90 15.5 110 125 125 116 123 300 125 9 1,782 - 829 - 53.5 34.9 9.65	
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	-1.8	-2.8	-	-2.8	
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet	3. 6. 7.	.3	4. 5. 7.	2	
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet	10. 10.		11. 10.		
NO _X , ppm Inlet Outlet	17 19			-	

Experiment No. Date of Run	9 10/2		10R 11/9/71		
Set No.	1	2	1	2	
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SB Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SB Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Manual Outlet SO ₂ , ppm - Manual Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	11,000 Dilute 116 54 155 14.5 110 121 120 115 119 292 124 9 - 700 65.9 35.7 13.76 14.38	10,910 Dilute 116 54 155 14.5 110 121 120 115 119 295 124 9 2,010 - 732 63.7 36.5 12.96	10,680 Dilute 112 53 154 9.9 110 120 120 114 120 302 120 15 2,040 1,980 620 540 72.6 41.2 14.22	10,690 Dilute 112 53 153 10.7 111 120 121 114 120 304 121 15 2,070 1,960 615 520 73.5 41.4 14.32	
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	-4.5	-10.1	1.3	-0.3	
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet	3	.1 .9 .0	3	.9 .7 .0	
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet	13 12		12 11		
NO _X , ppm Inlet Outlet		-		59 53	

Experiment No. Date of Run		11R /14/71		12R /9/71
Set No.	1	2	1	2
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SB Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SB Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Analyzer Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	11,500 CONC 165 0 155 14 52 105 100 113 90 288 108 9 2,084 1,980 118 120 94.0 146	11,400 ENTRATED 169 0 160 13 52 105 105 105 108 90 289 108 9 2,084 1,980 118 120 94.0 155		11,200 ENTRATED 110 53 150 17 110 121 123 114 118 296 122 9 2,080 1,980 113 100 94.3 192 20.15
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	-0.7	-7.7	2.5	4.0
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet		- - -		3.9 4.1 5.3/
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet		<u>-</u>		11.0 10.6
NO _X , ppm Inlet Outlet		-		-

Experiment No. Date of Run	13 11/4		14R 11/5/71		
Set No.	1	2	1	2	
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SB Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SB Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Analyzer Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	11,330 INTERM 110 54 153 54 110 120 120 116 123 298 122 9 2,050 2,000 380 420 81.5 46.5 17.78 18.06	11,400 EDIATE 110 54 153 54 110 119 120 116 123 298 122 9 2,040 1,980 280 320 86.3 46.5	11,300 INTERM 75 36 95 15 115 122 122 113 122 300 129 9 2,100 2,070 - 780 62.4 31.5	11,360 EDIATE 75 36 95 14 115 122 122 113 122 299 129 2,070 2,040 775 780 61.7 32.8 12.91	
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	-7.6	-3.0	2.0	0.9	
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet	3	.2 .7 .3		4.2 4.4 5.3	
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet	14 12			2.8 2.8	
NO _X , ppm Inlet Outlet		52 48		-	

Experiment No. Date of Run	15I 11/5,		16R 11/5/71		
Set No.	1	2	1	2	
FG Rate (CFM @ 130°F) Feed Composition SW Rate (GPM) SF Rate (GPM) SL Rate (GPM) SB Rate (GPM) SW, SF Temp. (°F) SL Temp. (°F) SB Temp. (°F) Inlet Gas Dew Point (°F) Outlet Gas Dew Point (°F) H. Extractor Outlet Gas Temp. (°F) Reheater Inlet Gas Temp. (°F) Overflow Pot Height (inches) Inlet SO ₂ , ppm - Manual Inlet SO ₂ , ppm - Analyzer Outlet SO ₂ , ppm - Analyzer % SO ₂ Removal % Stoichiometry Sulfur removed from Flue Gas ΔS _G , gmole/min. Sulfur absorbed by Na ₂ CO ₃ ΔS _L , gmole/min.	12,980 INTERME 110 55 145 16.5 110 120 115 118 302 120 9 2,100 2,040 528 500 75.6 43.7 18.97		11,500 INTERMI 110 55.5 145 20 110 115 116 111 118 223 116 9 2,042 2,010 - 350 82.5 48.6 18.03	11,500 EDIATE 110 55.5 145 20 110 114 115 111 118 222 115 9 2,052 2,020 389 350 83.2 48.6 18.14	
$\frac{\Delta S_{G} - \Delta S_{L}}{\Delta S_{G}} \times 100$	0.6	1.0	1.1	1.3	
O ₂ in the Flue Gas, % Boiler Outlet Scrubber Inlet Scrubber Outlet	3.8 4.1 5.1		4.0 4.4 5.5		
CO ₂ in the Flue Gas, % Scrubber Inlet Scrubber Outlet		2.8 2.8		2.8 2.8	
NO _X , ppm Inlet Outlet		-		-	

APPENDIX C

SOLUBLE TESTS - ANALYTICAL RESULTS

APPENDIX C

SOLUBLE TESTS - ANALYTICAL RESULTS

M Mole/Liter

			M Mole/	Liter		
10/29/71 Experiment No. 1R	Downcomer 1 SL	Downcomer 2 SL	Downcomer 3 SL	Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1 Time Total Sulfur as SO ₄	1052 -	1055 -	1058 -	1100 20.3	1110 0.3	1106 18.0
1)* S0 ⁼ + HS0 ⁻ 3	15.5	14.4	14.8	15.0	0.12	13.0
2) SO ₃ + HSO ₃	16.7 -	16.3	16.4	16.2 21.1	- 21.0	15.8 21.0
pH, initial pH, final	5.2 4.5	5.5 5.4	5.5 5.35	5.38 5.1	10.3	5.65
Set 2 Time Total Sulfur as SO ₄	1115 -	1117	1120 -	1125 19.7	1117 0.3	1130 18.4
1) $S0_3^2 + HS0_3^-$	15.1	14.0	14.8	14.8	-	13.4
2) SO ₃ + HSO ₃	16.0 -	16.2 -	15 . 2	16.7 21.1	- 21.0	15.8 21.1
pH, initial pH, final	5.28 5.19	5.42 5.35	5.1 5.02	5.1 5.02	10.30	5.92 6.0
10/27/71 Experiment No. 2R						
Set 1 Time Total Sulfur as SO ₄	1400 21.8	1409 22.7	1411 22.9	1416 20.6	1418 0.3	1414 22.3
1) $S0_3^2 + HS0_3^-$	17.7	13.3	16.9	15.2	0.12	15.8
2) $S0_3^{=} + HS0_3^{-}$ Na+ pH, initial pH, final	15.4 28.6 5.95 5.9	13.2 28.6 6.30 6.20	15.7 27.8 6.05 5.95	15.3 27.8 6.05 5.95	- 27.6 10.45 -	14.6 27.8 6.05 5.95
Set 2 Time Total Sulfur as SO ₄	1420 21.8	1422 22.3	1424 22.9	1429 20 . 9	1430 0.3	1428 21.8
1) $50\frac{1}{3} + HSO\frac{1}{3}$	17.0	13.5	16.3	14.4	0.12	15.5
2) $S0_3^2 + HS0_3^-$ Na+ pH, initial pH, final	15.5 27.8 6.0 5.95	12.6 27.8 6.3 6.25	13.1 27.8 6.05 6.00	27.6 6.05 6.00	- 27.3 10.45 -	- 27.6 5.95 5.90

^{*1)} C-E Method 2) Radian Method

			M Mole/	'Liter		
10/14/71 Experiment No. 3R	Downcomer 1 SL	Downcomer 2 SL	Downcomer 3 SL	Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1						
Time Total Sulfur as $S0_4^=$	1200 22.91				1200 0.34	1200 18.75
SO ₃ + HSO ₃ Na+ pH, initial pH, final	22.70 26.96 4.65 4.40				26.96 10.93	18.70 26.08 6.00 5.75
Set 2						
Time Total Sulfur as $S0_4^=$	1200 22.59				1200 0.34	1200 19.27
SO ₃ + HSO ₃ Na+ pH, initial pH, final	21.80 25.21 4.0 3.68				- 25.21 10.9 -	18.90 25.65 5.9 5.62
10/28/71 Experiment No. 4R						
Set 1						
Time Total Sulfur as SO ₄	1215 20.6	1216 19.7	1218 20.5	1219 19.7	0.3	1220 17.1
SO ₃ + HSO ₃ Na+ pH, initial pH, final	17.6 18.6 24.5 3.75	15.5 16.9 24.7 5.20	16.0 17.7 24.5 5.29	16.8 17.3 24.5 4.38	0.12 - 24.3	13.70 15.1 24.5 5.90
Set 2						
Time Total Sulfur as SO ₄	1221 20 . 8	1222 20.3	1223 19.7	1224 20.3	1227 0.3	1225 16.6
SO ₃ + HSO ₃ Na+ pH, initial pH, final	17.5 18.0 25.2 3.93	15.4 16.6 25.0 5.35	16.2 17.8 25.0 5.36	16.5 18.2 25.0 4.50	0.12 - 24.1 -	13.7 14.8 25.0 6.02

11/2/71 Experiment No. 5R	Downcomer 1 SL	Downcomer 3 SL	M Mole/Liter Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1					
Time Total Sulfur as SO ₄	1548 -	1551 -	1553 23.75	- 0.28	1554 19.79
S0 ⁼ + HS0 ⁻ Na ⁺	20.72	19.28	- 25.00	0 24.56	21.38 24.56
pH, initial pH, final	5.48 5.32	5.55 5.54	5.57 5.48	10.37	5.82 5.85
Set 2					
Time Total Sulfur as SO ₄	1601 -	1603 -	1604 23.95	- 0.28	- 19.79
S03 + HS03	21.01	19.37	- 24.34	0 24.34	21.36 24.34
pH, initial pH, final	5.30 5.35	5.55 5.52	5.58 5.43	10.38	5.85 5.85
11/2/71 Experiment No. 6R					
Set 1					
Time Total Sulfur as $S0_4^=$	1320 -	1322 -	1324 24.01	1332 0.28	1330 18.75
S03 + HS03 Na+	21.02	20.40	- 26 . 95	0 26.95	14.92 26.95
pH, initial pH, final	5.95 5.92	6.05 -	6.12 6.08	10.35	6.44 6.32
Set 2					
Time Total Sulfur as $S0_4^{=}$	1348 -	1351 -	1356 24.06	1353 0.28	1359 18.95
S03 + HS03 Na+	20.41	20. 51	- 26.72	0 26.86	16.67 26.60
pH, initial pH, final	5.95 5.88	6.08 5.99	6.12 6.02	10.35	6.43 6.38

11/3/71 Experiment No. 7R	Downcomer 1 SL	Downcomer 3 SL	M Mole/Liter Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1					
Time Total Sulfur as SO ₄	1200 -	1201 -	1204 20.20	1207 0.28	1205 20.31
SO ₃ + HSO ₃ Na +	20.13	19.55	- 27.17	0 27.17	18.55 26.95
pH, initial pH, final	6.15 6.05	6.31 6.25	6.40 6.21	10.70	6.20 6.15
Set 2					
Time Total Sulfur as SO ₄	1210 -	1212 -	1220 20.31	1227 0.28	1225 20.20
SO ₃ + HSO ₃ Na+ pH, initial pH, final	18.43 - 6.20 6.06	16.91 - 6.35 6.326	- 26.73 6.22 6.20	0 26.52 10.68	18.60 26.52 6.18 6.06
11/3/71 Experiment No. 8R					
Set 1					
Time Total Sulfur as SO ₄	1527 -	1529 -	1532 27 . 60	1536 0.27	1533 20.05
SO3 + HSO3 Na4 pH, initial pH, final	20.75 - 6.00 5.87	25.05 - 5.87 5.78	- 33.04 5.92 5.84	0 32.39 10.46 -	17.62 31.52 6.48 6.52
Set 2					
Time Total Sulfur as SO ₄	1645 -	1647 -	1650 25.52	1654 0.27	1652 19 . 27
SO ₃ + HSO ₃ Na+ pH, initial pH, final	21.63 - 5.64 5.56	22.04 - 5.65 5.55	- 29 . 13 5.62 5.62	0 29.13 10.38	18.49 29.34 6.38 6.37

	M Mole/	M Mole/Liter				
10/29/71 Experiment No. 9R	, Downcomer 1 SL	Downcomer 2 SL	Downcomer 3 SL	Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1						
Time Total Sulfur as SO ₄	1602 -	1607 -	1610 -	1613 23.1	1610 0. 3	1615 20.1
1)* S0 ⁼ + HS0 ⁻ 3	18.4	16.0	17.0	16,8	0.12	15.4
2) $S0_3^{=} + HS0_3^{-}$ Na+ pH, initial pH, final	18.4 - 5.65 5.69	16.9 - 5.95 5.97	18.2 - 5.90 5.85	17.7 24.3 5.82 5.75	- 24.3 -	18.2 24.3 6.02 6.00
Set 2						
Time Total Sulfur as SO ₄	1621 -	1625 -	1627 -	1628 22.8	1625 0.3	1633 20.1
1) $S0_3^2 + HS0_3^-$	18.0	15.6	16.4	16.7	0.12	15.1
2) SO3 + HSO3 Na+ pH, initial pH, final	19.4 - 5.45 5.52	16.3 - 5.87 5.92	16.9 - 5.84 5.79	17.8 23.9 5.71 5.72	- 24.1 10.38	18.7 23.9 5.97 5.98

^{*1)} C-E Method 2) Radian Method

11/9/71 Experiment No. 10R	Downcomer 1 SL	Downcomer 3 SL	M Mole/Liter Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1					
Time Total Sulfur as SO ₄	1305 -	1308 -	1312 22.91	1314 0.27	1311 24.06
S0 ⁼ ₃ + HS0 ⁻ ₃ Na ⁺	20.78	19.41 -	- 26.52	0 26.73	20.44 26.73
pH, initial pH, final	5.62 5.62	5.62 5.55	5.80 5.73	10.38 10.35	6.04 5.88
Set 2					
Time Total Sulfur as SO ₄	1316 -	1318 -	1321 23.43	1322 0.27	1320 23.95
SO ₃ + HSO ₃ Na ² pH, initial pH, final	20.53 - 5.75 5.65	18.73 - 5.85 5.62	- 26.73 5.85 5.77	0 26.73 10.37 10.32	19.88 26.73 6.08 5.95
10/14/71 Experiment No. 11R					
Set 1					
Time Total Sulfur as SO ₄	1430 34.37			1430 0.44	1430 41.66
SO ₃ + HSO ₃ Na+ pH, initial pH, final	34.25 109.56 7.4 7.2			0.13 110.86 11.5	34.75 110.43 7.8 7.55
Set 2					
Time Total Sulfur as SO ₄	1430 38.02			1430 0.51	1430 42.18
SO ₃ + HSO ₃ Na+ pH, initial pH, final	33.25 114.78 7.5 7.28			0.38 115.65 11.44 -	34.85 114.78 7.5 7.25

11/9/71 Experiment No. 12R	Downcomer 1 SL	Downcomer 3 SL	M Mole/Liter Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
<u>Set 1</u>					
Time Total Sulfur as $S0_4^{=}$	1510 -	1512 -	1513 31.66	1517 0.39	1515 37.29
SO3 + HSO3 Na [∓]	30.10	27.23 -	- 130.43	0 130.43	34.15 130.43
pH, initial pH, final	7.85 7.78	8.00 7.95	8.02 7.95	10.80 10.75	7.75 7.62
Set 2					
Time Total Sulfur as SO ₄	1520 -	1522 -	1523 30.46	1526 0.39	1525 35.62
SO ₃ + HSO ₃ Na+ pH, initial pH, final	29.15 - 8.00 7.97	26.97 - 8.15 8.12	- 133.91 8.12 8.05	0 133.47 10.80 10.70	32.86 133.91 7.95 7.82
11/4/71 Experiment No. 13R					
Set 1					
Time Total Sulfur as $50\frac{1}{4}$	1218 -	1220 -	1223 28 . 64	1225 0 . 27	1224 26.56
SO ₃ + HSO ₃ Na+ pH, initial pH, final	24.88 - 6.05 5.95	23.51 - 6.18 6.13	- 33.91 6.22 6.12	0 34.13 10.48	22.05 34.34 6.42 6.38
Set 2					
Time Toțal Sulfur as SO ₄	1228 -	1230 -	1237 28 . 90	1234 0.27	1233 26.82
SO ² + HSO ² Na ² pH, initial pH, final	25.65 - 6.18 6.08	24.30 - 6.35 6.22	- 36.30 6.35 6.10	0 36.73 10.52	23.10 36.30 6.45 6.42

Experiment No.	Downcomer 1 SL	Downcomer 3 SL	M Mole/Liter Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1					
Time Total Sulfur as SO ₄	1511 -	1513 -	1514 32.70	1518 0.27	1516 22.91
$S0_3^{-} + HSO_3^{-}$ Na+ pH, initial pH, final	26.79 5.87 5.75	26.50 - 5.75 5.63	- 33.47 5.87 5.77	0 33.69 10.37	21.23 33.47 6.55 6.48
Set 2					
Time Total Sulfur as SO ₄	1521 -	1523 -	1524 32.50	1527 0.27	1526 23.17
SO ₃ + HSO ₃ Na+ pH, initial pH, final 11/5/71 Experiment No. 15R	25.73 - 5.78 5.78	25.45 - 5.75 5.72	33.04 5.81 5.72	0 32.60 10.45 10.37	20.18 32.60 6.53 6.47
Set 1					
Time Total Sulfur as SO ₄	1324 -	1326 -	1327 31.25	1330 0.27	1329 30.00
SO ₃ + HSO ₃ Na+ pH, initial pH, final	26.38 - 5.97 5.42	25.91 - 6.18 6.04	- 36.08 6.18 6.08	0 36.08 10.38	25.72 36.30 6.35 6.32
Set 2					
Time Total Sulfur as SO ₄	1335 -	1337	1338 31.14	1341 0.27	1340 29.89
SO ⁼ ₃ + HSO ⁻ ₃ Na ⁺ pH, initial pH, final	26.58 - 6.08 5.96	25.10 - 6.05 6.08	- 36.08 6.25 6.08	0 35.86 10.38	25.40 35.86 6.25 6.22

11/5/71 Experiment No. 16R	Downcomer 1 SL	Downcomer 3 SL	M Mole/Liter Pump Discharge SL	Scrubber Feed (SW, SF)	Scrubber Bottom SB
Set 1					
Time Total Sulfur as $S0_4^=$	1137 -	1139 -	1141 29.58	1144 0.27	1143 26.30
SO ₃ + HSO ₃ Na+ pH, initial pH, final	25.30 - 6.05 6.03	24.57 - 6.35 6.22	35.24 6.42 6.28	0 35.43 10.38	18.81 34.95 6.45 6.37
Set 2					
Time Total Sulfur as SO ₄	1148 -	1149 -	1151 29.39	1153 0.27	1154 26.22
SO3 + HSO3 Na4 pH, initial pH, final	27.14 - 6.05 6.02	22.99 - 6.32 6.18	35.21 6.35 6.25	0 35.52 10.38	22.49 35.21 6.55 6.44

APPENDIX D

ANALYTICAL PROCEDURES
FOR APCS SAMPLES

APPENDIX D

ANALYTICAL PROCEDURES FOR APCS SAMPLES

The procedures for quantitatively analyzing solutions containing sulfite, sulfate, calcium, magnesium and sodium used in the APCS testing follow. These methods are time-tested and generally follow currently accepted analytical practices. It should be stressed that the instability of scrubber samples produces obvious variations and that sampling techniques are extremely important. The methods referred to are for control testing and in the hands of a good analyst produce results within 5% of the actual concentration.

APCS SULFITE*

Analysis (Liquid)

1. Transfer 20 ml of standard potassium iodide-iodate (0.0125 N = 0.5 mg SO_3/ml) to a 150 ml beaker.

Note: If sample contains solids it must be pressure filtered at the point of sampling prior to Step 2. The sample bottle (250 ml or less) should be filled to the top and capped immediately. A nitrogen purge can be maintained on samples during titration where sulfite values are below 200 ppm and precision is necessary. Samples must be analyzed within 10 minutes from time of sampling.

- 2. Pipet a sample volume (freshly taken-middle of sample bottle) containing less than 10 mg $\rm SO_3$ beneath the surface of the KI-KIO $_3$ (if a 5 ml sample volume is used, procedure is good to 2000 ppm).
- 3. To the 150 ml beaker containing the sample and KI-KIO $_3$ add 1 ml of 1 + 1 HCL, and 1 ml of starch solution. Stir gently and immediately back titrate to a colorless end point with 0.100 N sodium thiosulfate.

Calculations:

(ml of KI-KIO₃) (Normality) - (ml of Na₂S₂O₃) (Normality)
= meq SO₃/sample
meq SO₃ x
$$\frac{1000}{\text{Sample Vol}}$$
 x 40 = ppm SO₃
SO₃x 0.800 = SO₂

*The EPA Arsenite method was used for comparison. This method fixes the sulfite at the point of sampling. It may be slightly more precise but requires considerably more time in preparation.

APCS SULFATE

Analyses (Soluble SO₂)

INTRODUCTION

This procedure is for total sulfur as SO_4 . All sulfite is oxidized to sulfate in this method.

1. Pipet a filtered sample volume (see note) into a l liter volumetric flask containing 2 ml of 1 + 1 HCl, 1 ml of 30% H_2O_2 and approximately 300 ml of D. I. water. Stir on magnetic stirrer and dilute to 1 liter mark.

NOTE: For best sulfate results when using turbidmetric method, the diluted volume should contain between 20 and 80 ppm. Run at least two standards near level expected and check curve routinely.

- 2. ASTM 516-68 modified sulfate method:
- A. Pipet 25 ml of sample into four separate beakers.
- B. Place beakers on magnetic stirrers and insert plastic magnets. Adjust to reasonable mixing speed.
- C. Add simultaneously to all beakers 10 ml glycerol and 5 ml of sodium chloride solution.
- D. Add 0.3g of barium chloride dihydrate crystals to only three samples. (Triplicate analyses) Leave one sample as reagent blank.
- E. Begin timing. Stir all solutions for 1 minute and then remove beakers.
- F. Fill a 2 cm cell with reagent (blank) not containing barium chloride and prepare spectrophotometer as shown below.

Model DU

Wavelength - 400 nm Sensitivity - 2
Slit - 0.1 Phototube - Blue

- G. Let solutions containing barium chloride stand for 4 minutes \pm 30 seconds and return to stirrer for 15 seconds.
- H. Set spectrophotometer at zero absorbance on the reagent blank not containing the barium chloride. Then read.

CALCULATIONS

- I. Record optical density of triplicate samples. If O. D. varies on any one sample more than 0.007 units, repeat analyses.
- J. Run at least two standards in area of expected concentration to determine slope of curve. Whenever new reagents are made or if the room temperature varies more than 2°F, check curve with standards before running samples.

REAGENTS: Barium Chloride - Crystals of barium chloride (BaCl₂ 2H₂O screened to 20 to 30-mesh).

Sodium Chloride Solution (240 g per liter) - Dissolve 240g of sodium chloride (NaCl) in water containing 20 ml of concentrated hydrochloric acid (HCl, sp gr 1.19), and dilute to 1-liter with water. Filter the solution if turbid.

Sulfate, Standard Solution (ml = 0.100 mg SO_4 --). Dissolve 0.1479 g of anhydrous sodium sulfate (Na₂SO₄) in water, and dilute with water to 1 liter in a volumetric flask. Standardize by the procedure prescribed in Section 2.

APCS SODIUM ANALYSIS

- 1. Take the diluted sample used in the sulfate determination and determine the sodium concentration by flame emission.
- 2. Prepare the necessary calibration curves using the following settings:

MODEL B	<u>0</u>	- 10 ppm	10 - 25 ppm	<u>20 - 50 ppm</u>
$\lambda = 589 \text{ nm}$	Photomultiplier	D	D	C
H ₂ = 3.5 psi	Sensitivity	2	1	1
0 ₂ = 10.5 psi				

- 3. In the concentration range expected adjust the slit to approximately 50% transmittance with the maximum curve standard. Peak out wavelength for optimum sensitivity and then readjust slit opening to 100% t.
- 4. Continually flush distilled water through aspirator when not running sample.
- 5. When reading sample or standard allow 30 seconds for meter reading to stabilize. Record transmittance of sample and read from graph.
- 6. Routinely check the transmittance of at least two standards during sample analyses.

REFERENCE: ASTM D 1428

REAGENTS: <u>Sodium Chloride</u> (0.2542 g per liter). Dry sodium chloride (NaCl) to constant weight at 105 C. Dissolve 0.2542 g of NaCl in water and dilute to 1 liter with water. Dilute 10 ml of this solution to 1 liter and store in polyethylene or equally alkali-metal-free containers. The latter solution contains 1000 ppb sodium. This solution can be further diluted to provide a known standard of any sodium concentration less than 1000 ppb Na+.

APCS CALCIUM

Analysis (Liquid)

- 1. Take a portion of the diluted sample prepared under the sulfate determination and read on atomic absorption. Dilute sample so that % absorption falls between 15 and 75% (absorbance 0.070 to 0.600).
- 2. With A/A warmed up and set properly according to
 P. E. book run at least three standards of a similar matrix salt to
 prepare a calibration curve. (5, 10 and 15 ppm with 3 slot burner should
 be adequate). Use 422.7 nm wave length.

Calculations:

Read off curve or set up ratio of absorbance to nearest standard. Then multiply by dilution factor to obtain ppm of Ca.

 $Ca \times 1.40 = Ca0$

X-RAY METHOD FOR CALCIUM SULFATE SOLUTIONS ANALYSIS

The x-ray fluorescence method is advantageous when rapidity of analysis is an important factor. Analytical results in most cases are comparable and in some cases superior to those attainable by "wet chemical" methods.

Presently the technique has been applied to the analysis of sulfur (40 ppm and greater) and calcium (2 ppm and greater) in solutions.

- 1. Procedure
- a) Dilute 10 ml of unknown 1:1 with 10% $\rm HNO_3$ (by volume) spiked with $\rm H_2O_2$ prepared from conc. $\rm HNO_3$ (sp. gr. = 1.42).
- b) Transfer 5 ml of sample to specimen cup and obtain count at S K α (1) and Ca K α (1) peaks. Correct count for background.

APCS MAGNESIUM

Analysis (Liquid)

- 1. Take a portion of the diluted sample prepared under the sulfate determination and read on atomic absorption. Dilute sample so that % absorption falls between 15 and 75% (absorbance 0.070 to 0.600).
- 2. With A/A warmed up and set properly according to P. E. book run at least three standards of a similar matrix salt to prepare a calibration curve. (0.5, 1.0, 2.0 ppm with 3 slot burner should be adequate). Use 285.2 nm wave length.

Calculations:

Read off curve or set up ratio of absorbance to nearest standard. Then multiply by dilution factor to obtain ppm of Mg.

 $Mg \ X \ 1.646 = MgO$

PROCEDURE FOR THE DETERMINATION OF TOTAL SO, IN AQUEOUS SOLUTIONS

Introduction

This method is intended to give an accurate determination of total SO_2 in aqueous solutions taken from limestone injection - wet scrubbing processes and containing interfering substances such as nitrite ion. The sample is added to an excess of buffered iodine solution. The iodine remaining after the stoichiometric SO_2 oxidation is titrated with standard sodium arsenite solution using an amperometric dead-stop method for endpoint detection.

The iodine solution is generated as needed for each determination using standard iodate solution and excess iodide ion at low pH (\sim 1-2). This method is more convenient and reliable than using standard iodine solutions. The iodine solution is buffered to pH 6.0-6.2 to inhibit sulfite-nitrite and nitrite-iodine (iodide) reactions. This also inhibits the air oxidation of iodide. Arsenite solutions give more accurate results than thiosulfate solutions in the presence of nitrite and are also more stable under ordinary conditions. The deadstop end-point detection method gives more reliable and accurate results than starch indicators, etc. In practice, the deadstop method is also convenient and simple.

In this method the order of addition of reagents and other procedures are critical and the procedure given should be followed closely.

Apparatus

- 1) 50 ml burette (preferably an automatic burette)
- 2) magnetic stirrer

- 3) pipets (including 2 ml and 20 ml sizes) and bulb pipet fillers
 - 4) 400 ml beakers (preferably graduated)
 - 5) deadstop apparatus:
 - a) two platinum electrodes
- b) 1.5 volt dry cell battery (a #735 "hobby battery works fine)
 - c) electrometer or 0-15 or 20 μA microammeter
 - d) voltage divider: 1.5 v to 0.1 v

Connect one electrode to one terminal of the voltage divider output (0.1 volt). Place the meter in series with the second electrode and the other voltage divider terminal.

Reagents and Solutions

Use distilled water and reagent grade chemicals for all solutions.

CAUTION: Arsenic is toxic and care should be taken when preparing and handling these solutions.

l) Sodium arsenite stock solution (0.100 mole/liter) - Weigh out accurately 9.893g arsenic trioxide (As_2O_3 , primary standard, 99.99%) and dissolve in about 100 ml 2M NaOH (8g NaOH/100 ml) with stirring. Adjust the pH to about 7 to 8 with first concentrated, then 1N HCl. Add about 1g NaHCO $_3$, transfer the solution quantitatively to a 1 liter volumetric flask and dilute to the mark with distilled water. This solution will be used to make up the standard solution for iodine titration.

- 2) Sodium arsenite standard solution (0.0100 mole/liter) Pipet 100.0 ml of the 0.100 mole/l sodium arsenite solution prepared in 1) into a l liter volumetric flask and dilute to the mark with distilled water. This solution is most conveniently and safely handled in an automatic burette assembly.
- 3) Potassium iodide (50 f/liter) Dissolve 50g of iodate free KI and 0.5g of NaHCO₃ in freshly boiled and cooled distilled water and dilute to one liter. Alternatively, 45g of iodate-free NaI may be used in place of the KI. The water is boiled or otherwise deaerated to remove dissolved oxygen which might oxidize iodide to iodine causing errors in the determination.
- 4) Potassium iodate standard solution (0.0833) mole/liter)-Dissolve 8.917g of potassium iodate (KIO_3), dried at 120°C, and 0.5g of sodium bicarbonate ($NaHCO_3$) in distilled water and dilute to 500 ml in a volumetric flask.
- 5) 1N HC1 Dilute 86 ml of concentrated reagent grade hydrochloric acid (36%) to 1 liter with distilled water.
- 6) pH 6.0 buffer (1 mole/liter sodium acetate, 0.05 mole/liter acetic acid) This solution should contain 82 g/liter anhydrous sodium acetate (or 136 g/liter of the trihydrate) and 2.9 ml/liter glacial acetic acid. It is convenient to prepare several liters of this solution at a time since about 175 ml is used for each determination. The water used to prepare this solution should be deaerated by boiling or bubbling nitrogen through it.

Procedure

The iodine-buffer solution should be prepared just prior to addition of the sample. The order of the following steps is important and should be maintained.

- a) Place 20 ml of KI solution (3) in a 400 ml beaker containing a magnetic stirring bar.
- b) Accurately pipet 2.00 ml of the ${\rm KIO}_3$ solution (4) into the beaker.
- c) Add 2.0 ml 1N HCl and swirl the beaker for about 10-15 seconds to ensure complete mixing and reaction.
- d) Immediately add about 175 ml pH 6.0 buffer (6) being careful not to lose any solution due to splashing. Stir the solution with the magnetic stirrer at low speed.
- e) Place the platinum electrodes in the solution and hook up the deadstop equipment (see apparatus). The current should be about 10-15 microamps.
- f) Accurately pipet a volume of sample containing between 5×10^{-5} and 5×10^{-4} mole total SO_2 into the iodine-buffer solution. For the in-house test about 10 ml of sample should probably be used (also see j).
- g) Increase the stirrer speed and begin the titration with the 0.01 mole/liter sodium arsenite solution (2) using a 50 ml burette. The color of the solution may be used as a rough guide to the approach of the end point. The iodine-red color changes to yellow about 10 ml before the end point. When the solution becomes light yellow the titration should be continued dropwise. The solution usually becomes colorless a few drops before the end point. The current will decrease about 0.2 to 1 microamp immediately before the end point and then decrease about 8-9 microamp at the end point.

- h) When the large current drop described above occurs stop the titration and record the ml of sodium arsenite used. Rinse the electrodes with distilled water after removing them from the solution.
- i) Run blanks each day using the above procedure without the addition of a sample. Two blanks should agree within 0.1 ml. Record the volume of titrant used in the blanks (it should be about 50 ml).
- j) If the iodine color is completely discharged (the solution becomes colorless) when the sample is added, prepare a fresh iodine-buffer solution and use a smaller sample volume (5 ml). If the volume of titrant taken for the blank and sample are within 5 ml of each other, repeat the determination, using a correspondingly larger sample. If very low SO₂ concentrations are encountered modify the reagent concentrations for the determination.

Calculation and Accuracy

The concentration of total ${\rm SO}_2$ in the sample can be calculated using the following equation.

$$C = \frac{(B-S)M}{V}$$

where:

C = concentration of total SO₂ (mole/liter)

B = volume in milliliters of arsenite solution needed to titrate the blank

 $\mbox{S = volume in milliliters of arsenite solution needed to} \\ \mbox{titrate the sample}$

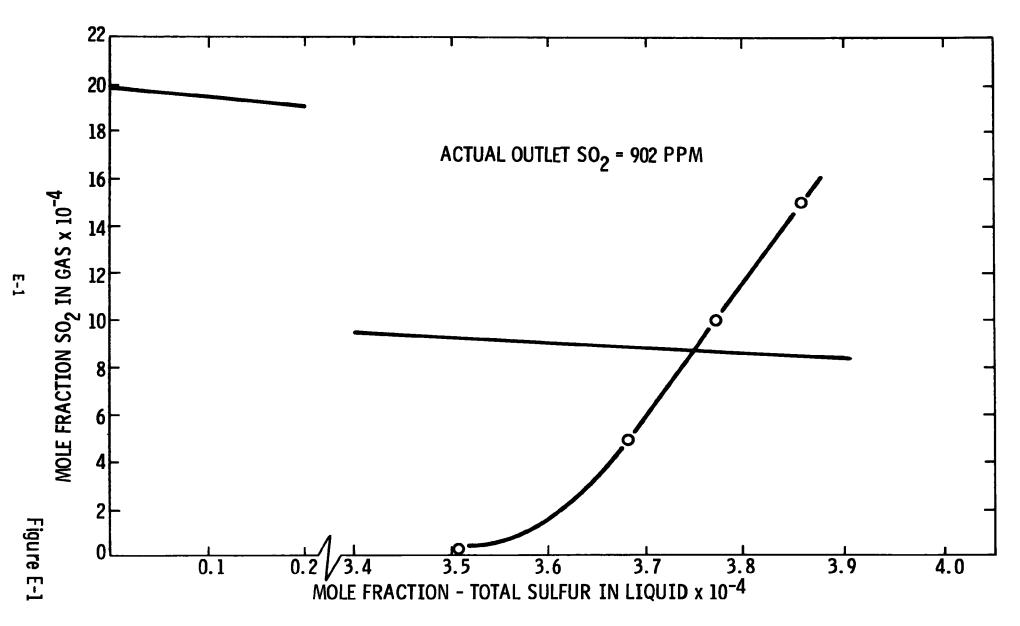
M = molarity of the arsenite solution, mole/liter (normally
0.0100)

V = volume of sample used, milliliters.

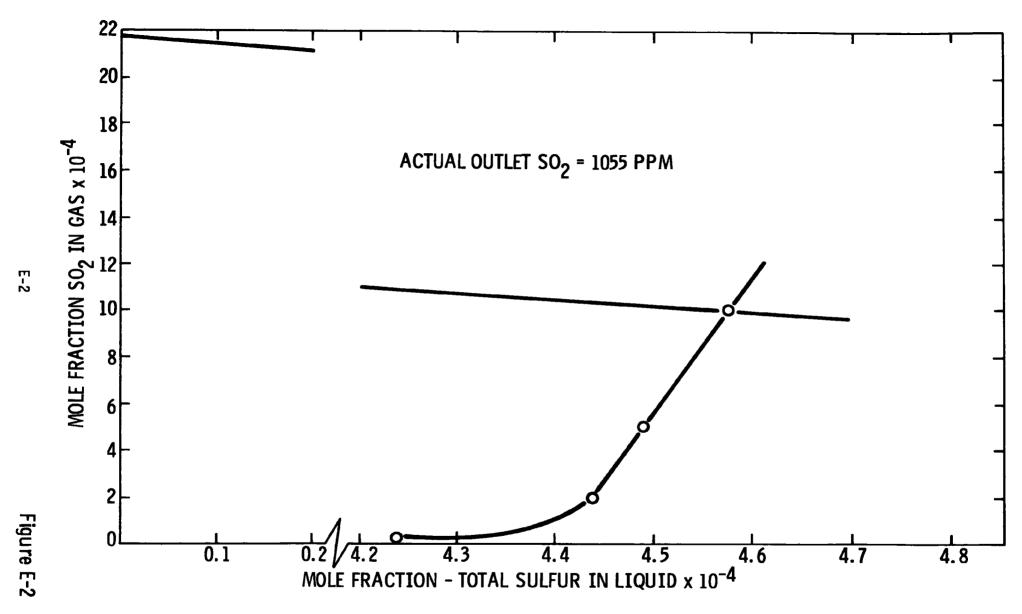
Using a $NO_2^-:SO_2$ mole ratio of 50:1 the SO_2 determination has been made with a 1-3% error using this procedure. Five determinations of K_2SO_3 without nitrite added gave a relative deviation of 0.25%.

APPENDIX E

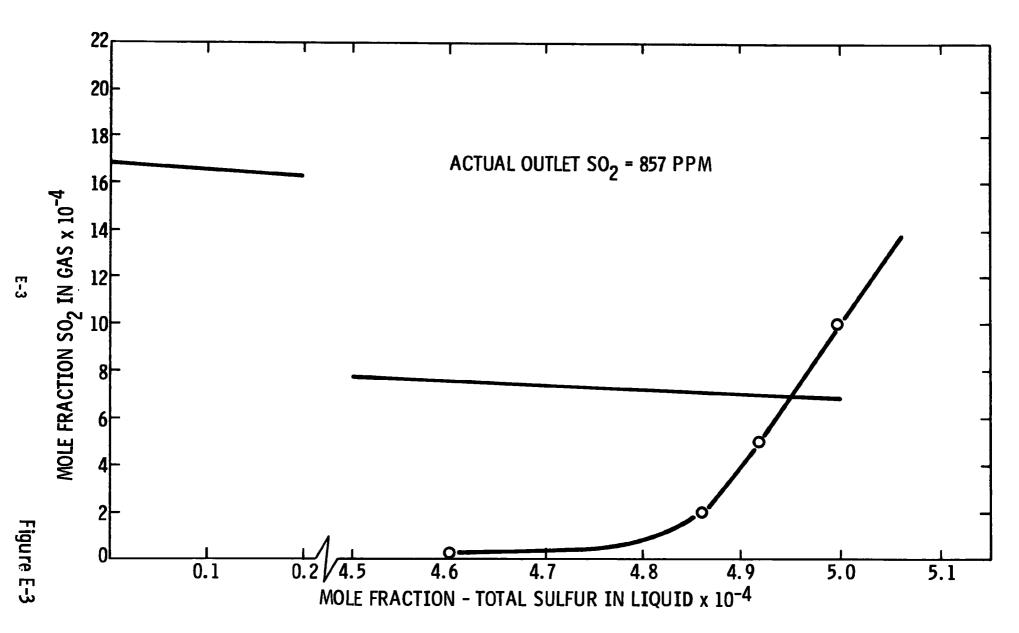
SOLUBLE SYSTEM
STAGE EFFICIENCY CALCULATION DIAGRAMS



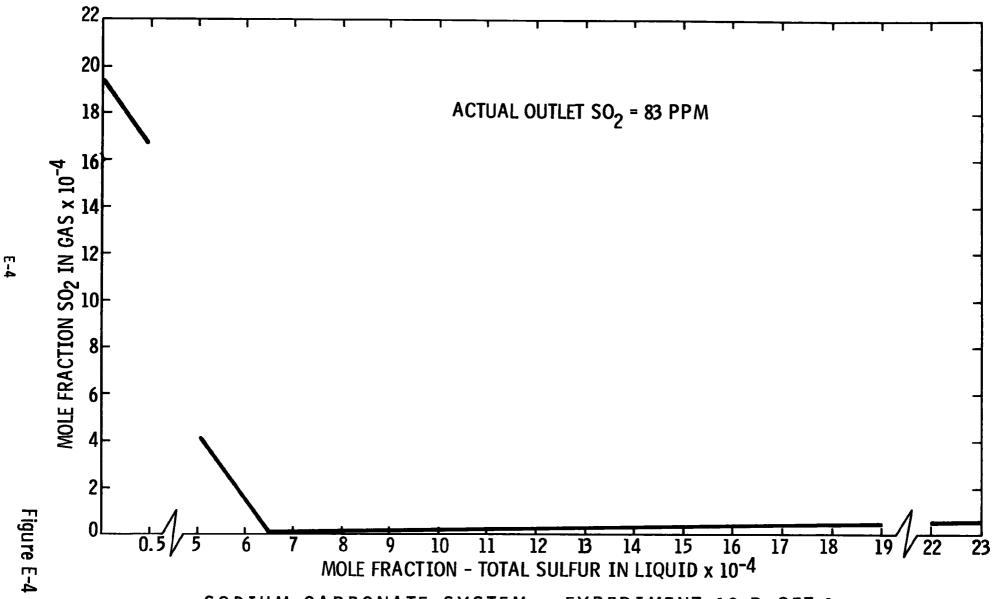
SODIUM CARBONATE SYSTEM - EXPERIMENT 1 R-SET 2



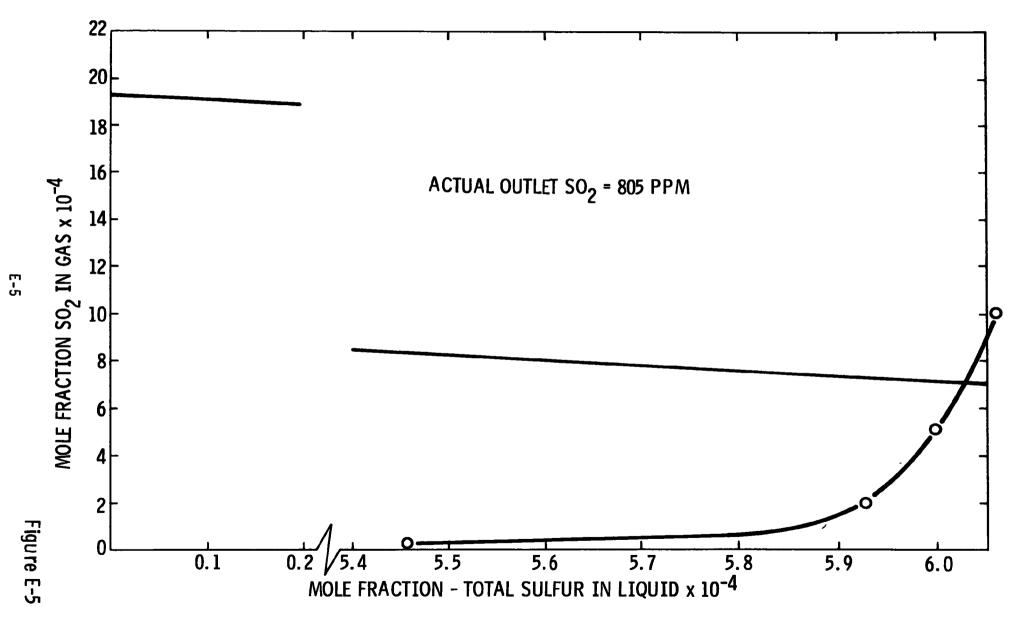
SODIUM CARBONATE SYSTEM - EXPERIMENT 5 R-SET 1



SODIUM CARBONATE SYSTEM - EXPERIMENT 8 R-SET 2



SODIUM CARBONATE SYSTEM - EXPERIMENT 12 R-SET 1



SODIUM CARBONATE SYSTEM - EXPERIMENT 14 R-SET 1

APPENDIX F

LIMESTONE FURNACE INJECTION SYSTEM
OPERATING DATA

AND

ANALYTICAL RESULTS

TABLE F-1. C-E APCS PROTOTYPE FURNACE INJECTION TESTS

Experiment No. Date of Run Set No.	17 12/2	R 8/71		8R 9/72	19 4/2	9R 1/72
Time	1	2	1	2	1	2
	1320-1405	1550-1650	1800-1900	1900-2030	1130-1200	1250
Flue Gas (FG) Rate (CFM @ 130°F)	11,000	11,000	11,000	11,000	10,000	9,940
Fly Ash + Additive Feed Rate (1b/hr.)	270	270	270	270	370	370
Spray Water SW (gpm)	110	110	205	205	201	201
Scrubber Liquid SL (gpm)	89	87	178	182	175	175
Scrubber Bottom SB (gpm)	25	28	30	30	20	20
Clarifier Liquid CL (gpm)	165	165	20	20	0	0
Liquid Blowdown LB (gpm)	55	55	0	0	35	35
Clarifier Feed CF (gpm)	165	165	25	25	35	35
Clarifier Bottom CB (gpm)	4	4	3	3 3	3	3 2
Filter Liquid FL (gpm)	4	4	3	3	2	2
Spray Water SW (°F)	100	100	120	120	106	108
Scrubber Liquid SL (°F)	123	121	122	123	115	115
Scrubber Bottom SB (°F)	121	119	122	123	116	118
Inlet Gas Dew Point (°F)	127	126	122	122	112	_
Outlet Gas Dew Point (°F)	122	121	121	121	106	_
Reheater Inlet Gas Temp. (°F)	128	124	125 ·	125	112	112
Heat Extractor Outlet Gas Temp. (°F)	304	305	285	285	293	289
Inlet SO ₂ (ppm)	1,456	1,456	1,471	1,471	1,883	1,881
Outlet SO ₂ (ppm)*	764	764	447	447	925	925
Inlet 02 (%)	5.6	5.6	4.5	4.5	10.7	_
Outlet 02 (%)	6.1	6.1	5.8	5.8	12.0	-
Inlet CO ₂ (%)	9.3	9.3	6.2	6.2	7.0	-
Outlet CO ₂ (%)	7.8	7.8	6.2	6.2	6.0	-
SO ₂ Removal Efficiency (%)	43.0	43.0	67.6	67.6	43.7	43.6
Stoichiometry (%)	71.0	71.0	72.8	72.8	88.8	89.5
Solid Concentration in Spray Water (%.)		-	3.67	3.35	1.14	1.46
Average Air Leakage (%)**	6.7	6.7	6.7	6.7	15.0	15.0
Dust Loading on the Outlet (gr./SCFM)	0.033	0.033		• • • • • • • • • • • • • • • • • • • •	0.002	0.003
bust Luading on the outlet (gr./3cm)	0.000	0.000			V. UUL	0.000

⁺High soot concnetration due to low $\mathbf{0}_2$ conc. in the boiler *Uncorrected for air leakage. **Average air leakage.

TABLE F-1. C-E APCS PROTOTYPE FURNACE INJECTION TESTS (Continued)

Experiment No.	20			1R	22R		
Date of Run	_ 4/20		4/2	6/72	_ 4/2	8/72	
Set No.	1	2	1	2	1_	2	
Time	1035-1130	1245-1345	1330-1345	1530-1420	1315-1630	1515-1600	
Flue Gas (FG) Rate (CFM @ 130°F)	10,020	10,020	9,670	10,000	9,940	9,900	
Fly Ash + Additive Feed Rate (1b/hr.)	370	370	390	390	390	390	
Spray Water SW (gpm)	204	205	200	200	358	355	
Scrubber Liquid SL (gpm)	180	180	180	180	258	260	
Scrubber Bottom SB (gpm)	20	20	20	20	100	95	
Clarifier Liquid CL (gpm)	40	40	10	10	10	10	
Liquid Blowdown LB (gpm)	40	40	0	0	0	0	
Clarifier Feed CF (gpm)	80	80	10	10	10	10	
Clarifier Bottom CB (gpm)	-	-	3	3	3	3	
Filter Liquid FL (gpm)	-	-	2	2	-	-	
Spray Water SW (°F)	108	109	120	120	116	119	
Scrubber Liquid SL (°F)	120	119	125	123	117	120	
Scrubber Bottom SB (°F)	120	119	125	123	116	119	
Inlet Gas Dew Point (°F)	104.5	104.5	110	-	108.2	107.5	
Outlet Gas Dew Point (°F)	113.2	113	114.5	-	112.5	117	
Reheater Inlet Gas Temp. (°F)	114	114	121	120	117	119	
Heat Extractor Outlet Gas Temp. (°F)	300	299	342	342	291	299	
Inlet SO ₂ (ppm)	1,962	1,939	2,000	1,985	2,022	2,019	
Outlet SÕ ₂ (ppm)*	1,091	1,090	735	677	514	453	
Inlet 0 ₂ (%)	_	10.5	10.4	_	10.4	-	
Outlet 02 (%)	_	11,9	12.0	-	11.9	-	
Inlet CO ₂ (%)	_	7.0	7.0	_	6.9	-	
Outlet CO2 (%)	-	6.0	6.0	-	6.0	-	
SO ₂ Removal Efficiency (%)	36.1	35.1	57.7	60.7	70.8	74.2	
Stoichiometry (%)	85.0	86.0	91.5	88.8	87.8	88.4	
Solid Concentration in Spray Water (%.)	0.69	0.70	8.02	6.67	8.58	_	
Average Air Leakage (%)**	15.0	15.0	15.00	15.00	15.00	15.00	
Dust Loading on the Outlet (gr./SCFM)	0.002	0.003	0.005	0.025+	-	-	

⁺High soot concnetration due to low $\mathbf{0}_2$ conc. in the boiler *Uncorrected for air leakage.

^{**}Average air leakage.

TABLE F-2. ANALYSIS OF FILTRATE SAMPLES FOR STEADY STATE DETERMINATION EXPERIMENT 17R

	12-27-71															
Time	#	Ca	Na	EB Mg	C1	so ₄	Ca	Na	SL Mg	C1	so ₄	Ca	Na	Mg	C1	s0 ₄
0920	1	909		<1		1699	1140		17		2982	1078		<1		1986
1145	2	650		16		1516	761		17		2230	540		3		1566
1315	3	603		3		1434	768		10		2322	525		<1		1132
1430	4	568		6		1208	747		12		2175	569		<1		1192
1540	5	567				1033	831				2200	632				1161
								12-	-28-7	l						
0020	6	628		< 1		1628						490		~ 1		1700
0820			10		00		75.0	19	13	110	2193	588	18	<1	100	1172
0930	7	513	18	<1 •	90	1160	752									1128
1030	8	504	18	<1	90	1100	828	18	21	100	2041	579	18	< 1	90	
1250	9	549				1051	824				2154	610				1020
1400	10	528				1007	734				1872	560				1036
1750	11	485				1012	759				2071	517				974
1835	12	424				919	608				1745	513				915
1915	13	408	14		70	969	623	14		70	1693	479	14		60	1019

1066

598

1718

473

990

EB - Hold Tank Effluent

420

SR - Scrubber Liquid

2015 14

SW - Spray Water Feed

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TABLE F-3. ANALYSIS OF FILTRATE SAMPLES FOR STEADY STATE DETERMINATION EXPERIMENT 18R

2-1-72

			Cl	(ppm)			SL	(ppm)			SV	(ppm)	
Time	#	Ca++	so ₌	s0 ₄ =	рH	Ca++	so ₌	so <mark>=</mark>	рН	Ca++	so ₃ =	s0 ₄ =	рН
1305	1	524	-	1170	7.20	1469	1232	1461	5.45	908	0	1820	5.75
						2.	-2-72						
0848 1030 1130 1200 1230 1300 1330	2 3 4 5 6 7 8	520 566 598 599 613 636 664	140 20 44 32 40 52 40	1172 1346 1477 1271 1342 1407 1472	7.28 8.62 7.38 8.43 9.31 9.25 9.38	741 1209 1221 922 1151 1136 917	1352 1232 800 796 712 320	2217 3021 2260 2564 2425 2066	5.51 5.45 5.65 5.82 6.21 8.29	756 830 807 914 929 909	100 160 56 56 40 48	1960 2348 2232 2422 2312 2142	8.32 6.78 8.52 9.49 9.73 10.01
1130 1300 1500 1530 1600 1630 1700	9 10 11 12 13 14	700 736 790 632 740 794 792	60 60 40 32 40 32 34	1538 1508 1682 1475 1510 1553 1557	10.87 11.16 10.91 11.22 11.18 11.22 11.18	800 826 660 794 816 828	60 60 192 232 256 12 252	- 1658 1999 1677 1862 1755	11.48 8.11 6.52 6.49 9.75 6.39 6.24	1052 1000 926 748 912 850 918	56 60 56 56 32 32 32	1312 1528 1922 1598 1679 1741 1679	12.01 11.65 10.95 10.92 10.88 11.19

TABLE F-4. FILTRATE ANALYSIS AND SOLID CONCENTRATION VALUES FOR STEADY STATE DETERMINATION

			CL						SW			
Experiment	Time	Ca++ (PPM)	SO ₃ (PPM)	Total Sulfur As SO ₄ PPM	S04 (PPM)	рН	Ca++ (PPM)	SO ₃ (PPM)	Total Sulfur As SO4 PPM	SO4 (PPM)	рН	Percent Solids
			4-18-72						4-18-72			
20R 20R	1100 1600	594 659	352 412	1860 1920	1437 1420	6.2 6.0	6 6 0 784	644 552	2130 2150	1357 1487	5.8 5.7	0.4 0.3
			4-19-72						4-19-72			
20R 20R 20R 20R 20R	1200 1300 1400 1500 1600	79 <i>2</i> 663 811 800 823	380 552 320 316 304	2290 1950 2340 2230 2260	1834 1287 1956 1850 1859	6.0 6.1	882 790 878 879 855	592 132 376 324 268	2540 2230 2470 2390 2200	1829 2071 2018 2001 1878	6.7 6.3 6.3	0.5 0.5 0.6 0.8 1.1
			4-20-72						4-20-72			
20R 20R 20R 20R 20R 20R 20R	0700 0730 0830 0900 0930 1000	745 894 893 857 877	252 240 312 336 380 432	2040 2280 2550 2440 2650	1737 1992 2175 2036 2194	6.25 6.2 6.3 6.3	902 952 886 829 813	180 440 356 352 300 320	2140 2630 2370 2440 2340	1924 2102 1942 2017 1980	9.00 6.4 5.8 5.8 5.8	1.1 1.1 1.1 0.9 0.7
			4-21-72						4-21-72			
19R 19R 19R 19R 19R 19R 19R 19R	0800 0830 0900 0930 1000 1030 1100	904 892 890 906 900 876 880 855	460 444 452 472 444 452 452 448	2850 2760 3000 2890 2940 2710 2690 2660	2298 2227 2457 2323 2407 2167 2147 2122	5.7 4.8 4.9	1034 983 978 965 875 841 830 836	72 472 632 700 572 624 632 612	2100 3050 3340 3420 2890 2730 2790 2820	2013 2483 2581 2580 2203 1981 2031 2085	5.7 5.4 5.7 5.5	0.8 1.6 2.4 1.5 1.7 1.4 1.7

TABLE F-4. FILTRATE ANALYSIS AND SOLID CONCENTRATION VALUES FOR STEADY STATE DETERMINATION (Continued)

			CL						SW			
Experiment	Time	Ca++ (PPM)	S03 (PPM)	Total Sulfur As SO ₄ PPM	SO ₄ (PPM)	рН	Ca++ (PPM)	SO3 (PPM)	Total Sulfur As SO ₄ PPM	SO4 (PPM)	рН	Percent Solids
			4-25-72					•	4-25-72			
21R 21R 21R 21R 21R 21R	1300 1400 1500 1530 1600						778 679 733 744 712	900 240 120 - 120	3450 2580 2590 2720 2870	2370 2292 2446 - 2726	6.1 7.0 7.3	4.4 6.1 7.9 8.5
			4-26-72						4-26-72			
21R 21R 21R 21R 21R 21R 21R 21R 21R	0630 0700 0730 0830 0900 1000 1100 1200 1300	789 808 818 849 853 844 833	20 60 60 80 80 92 64 120	1960 1940 2190 2030 2110 2150 2140	1936 1868 2118 1934 3014 2039 2063		1297 1297 686 791 721 825 853 866 776	140 40 120 112 80 128 140 108 88	1470 1550 2280 2260 2640 2550 2820 2700 2780 4-28-72	1302 1502 2136 2125 2544 2396 2652 2570 2674	6.7 6.2 6.2 9.15	7.8 6.3
22R 22R 22R 22R 22R 22R 22R 22R 22R	0750 0830 0900 0930 1000 1110 1210 1315	860	56	1990	1992		791 857 913 848 817 756 729 719	44 60 40 52 64 336 124	2060 2290 2320 2250 2440 3340 3960 3020	2067 2218 2272 2187 2363 2936 2811 2972	9.4 7.4 6.7 9.5	7.1 10.2 10.5 9.0 8.4 9.5

TABLE F-5. RESULTS OF SOLID PHASES ANALYSES EXPERIMENT 17R

Composition in Millimole/Gram Solid Wt % Solids Weight S0₃ Sample Location in Slurry Date Total S Μg S0₂ CO₂ % Undissolved Ca Set 1 51.3 Marble Bed Front 12/28/71 0.211 0.869 3.22 .522 0.80 0.07 0.220 12/28/71 0.316 Marble Bed Back Scrubber Bottom "S" 35.5 12/28/71 1.02 0.677 0.09 5.68 .402 0.59 0.362 12/28/71 .0078 Spray 1.51 0.650 41.4 Hold Tank Effluent 12/28/71 0.327 1.74 4.57 0.479 0.23 Set 2 49.6 0.274 0.939 0.42 0.227 Scrubber Liquid "T" 1 12/28/71 3.40 . 448 0.52 53.7 Scrubber Liquid "T" 2 12/28/71 0.251 3.68 .488 0.55 0.45 0.236 1.002 0.224 51.7 0.269 0.914 Scrubber Liquid "T" 3 12/28/71 3.57 .506 0.49 0.42 41.9 1.012 0.566 Scrubber Bottom "T" 1 12/28/71 0.850 0.88 0.13 5.52 .430 0.23 41.5 1.012 0.554 Scrubber Bottom "T" 2 12/28/71 0.904 5.59 .435 0.78 0.650 41.4 12/28/71 1.74 4.56 .479 1.51 0.23 Hold Tank Effluent 0.327 55.7 0.296 12/28/71 0.765 0.30 0.46 Marble Bed Front 0.163 3.28 .498 Marble Bed Back 12/28/71 0.293 41.2 0.511 Scrubber Bottom "S" 12/28/71 1.11 5.80 .416 0.50 0.14 0.692 12/28/71 0.0103 Spray 0.755 40.5 12/28/71 1.11 2.10 3.87 .516 1.76 0.34 Clarifier Bottom Filter Liquid 12/28/71 0.0139 0.595 40.5 0.45 4.15 .868 0.09 79.6 Filter Solid 12/28/71 0.538

TABLE F-6. RESULTS OF SOLID PHASES ANALYSES EXPERIMENT 18R

Sample Location	Date	Wt % Solids in Slurry	Total S	Ca	Mg	so ₂	so ₃	_co ₂	Weight % Undissolved
Set 1									
Scrubber Liquid: T1 2 3	2/ 3/72	4.45 4.46 5.28	3.15 3.10 3.04	4.50 4.57 4.60	0.34 0.34 0.34	2.26 2.27 2.18	0.87 0.83 0.86	0.58 0.55 0.50	34.6 32.1 34.8
Scrubber Bottoms: T1 2 3	2/ 3/72	5.83 6.00 6.15	2.50 2.62 2.60	4.81 4.62 4.69	0.41 0.41 0.40	1.57 1.63 1.73	0.93 0.99 0.87	0.52 0.61 0.73	35.1 35.9 36.0
Clarifier Liquid Hold Tank Effluent Marble Bed: Front Back Scrubber Bottoms S Scrubber Spray Additive	2/ 3/72 2/ 3/72 2/ 3/72 2/ 3/72 2/ 3/72 2/ 3/72 2/ 3/72	.017 4.50 4.17 4.10 5.39 3.67	3.08 3.33 3.34 2.51 3.01 0.51	4.47 4.56 4.52 4.85 4.59 5.97	0.40 0.31 0.32 0.41 0.41	2.22 2.31 2.34 1.73 2.17 0.06	0.86 1.02 1.00 0.785 0.84 0.45	0.63 0.59 0.60 0.49 0.58 0.39	34.1 33.3 34.2 36.0 34.9 46.3
Set 2	- , -, -								1010
Scrubber Liquid: T1 2 3	2/ 3/72	4.07 4.12 3.93	3.28 3.16 3.20	4.59 4.54 4.49	0.33 0.33 0.33	2.26 2.32 2.36	1.02 0.84 0.84	0.51 0.54 0.58	34.0 34.5 34.6
Scrubber Bottoms: T1 2 3	2/ 3/72	5.26 5.61 5.79	2.36 2.42 2.66	4.80 4.85 4.64	0.41 0.41 0.41	1.63 1.67 1.89	0.73 0175 0.77	0.62 0.63 0.63	36.3 36.5 36.3
Hold Tank Effluent Marble Bed: Front Back	2/ 3/72 2/ 3/72	3.66 4.10 3.91	2.98 3.18 3.19	4.57 4.50 4.54	0.41 0.32 0.32	2.12 2.32 2.36	0.865 0.86 0.83	0.64 0.49 0.52	34.0 33.2 34.6
Scrubber Bottoms S Scrubber Spray Clarifier Bottoms	2/ 3/72 2/ 3/72	4.99 3.35	2.43 3.08 2.98	4.93 4.47 4.33	0.40 0.40 0.43	1.69 2.20 2.13	0.745 0.88 0.85	0.59 0.54 0.54	35.1 34.6 35.3

TABLE F-7. RESULTS OF SOLID PHASES ANALYSES EXPERIMENT 19R

Sample Location	Date	Wt % Solids in Slurry	Total S	Ca	Mg	so ₂	so ₃	co ₂	Weight % Undissolved
Set 1									
Scrubber Liquid Tk 1 Scrubber Liquid Tk 2 Scrubber Liquid Tk 3 Scrubber Bottoms Tk 1 Scrubber Bottoms Tk 2 Scrubber Bottoms Tk 3 Hold Tank Effluent Marble Bed: Front Marble Bed: Back Scrubber Bottoms S Scrubber Spray	4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72	1.22 1.22 1.23 2.61 2.73 3.28 1.39 1.50 1.38 3.21	1.96 1.90 1.95 2.06 1.64 1.88 2.26 1.83 1.90 1.53 2.21	3.30 3.31 3.45 4.10 4.93 4.53 3.85 3.68 3.47 4.88 3.73	0.224 0.222 0.228 0.329 0.321 0.330 0.200 0.244 0.221 0.341 0.197	1.32 1.29 1.34 1.57 1.28 1.52 1.60 1.24 1.28 1.47	0.64 0.61 0.61 0.49 0.36 0.36 0.66 0.59 0.62 0.06	0.175 0.166 0.177 0.271 0.240 0.237 0.172 0.171 0.171 0.280 0.215	53.0 50.2 50.5 44.5 42.3 45.1 47.7 51.1 51.7 45.1
Set 2 Scrubber Liquid Tk 1 Scrubber Liquid Tk 2 Scrubber Liquid Tk 3 Scrubber Bottoms Tk 1 Scrubber Bottoms Tk 2 Scrubber Bottoms Tk 3 Hold Tank Effluent Marble Bed: Front Marble Bed: Back Scrubber Bottoms S Scrubber Spray Fly Ash and Lime Fly Ash and Lime	4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72 4/21/72	1.21 1.35 1.34 3.27 3.54 3.10 1.49 1.67 1.52 4.19	1.85 1.84 1.77 2.13 1.85 1.90 2.05 1.66 1.79 1.42 2.01 0.45 0.54	3.39 3.52 3.50 4.19 4.72 4.16 3.84 3.98 3.89 5.01 4.01 5.81 5.78	0.240 0.243 0.235 0.336 0.334 0.215 0.265 0.255 0.350 0.224 0.500 0.498	1.22 1.23 1.15 1.58 1.37 1.45 1.45 1.18 1.15 1.33 1.41	0.63 0.61 0.62 0.55 0.48 0.45 0.60 0.48 0.64 0.09	0.161 0.209 0.201 0.333 0.349 0.335 0.189 0.198 0.243 0.402 0.180 0.455	51.1 51.9 50.5 44.0 42.0 43.4 45.5 48.6 48.9 42.4 45.4 45.8 47.6

TABLE F-8. RESULTS OF SOLID PHASES ANALYSES EXPERIMENT 20R

Sample Location	Date	Wt % Solids in Slurry	Total S	Ca	Mg	so ₂	so ₃		Weight % Undissolved
Set 1									
Scrubber Liquid Tk 1	4/20/72	0.735	1.67	3.18	0.300	0.98	0.69	0.205	50.3
Scrubber Liquid Tk 2	4/20/72	0.743	1.57	3.34	0.307	0.98	0.59	0.209	51.2
Scrubber Liquid Tk 3	4/20/72	0.691	1.58	3.79	0.370	1.03	0.55	0.172	52.2
Scrubber Bottoms Tk 1	4/20/72	2.67	1.72	4.61	0.483	1.37	0.35	0.384	40.8
Scrubber Bottoms Tk 2	4/20/72	1.87	1 .9 8	4.73	0.578	1.53	0.45	0.199	42.2
Scrubber Bottoms Tk 3	4/20/72	2.27	1.83	4.41	0.482	1.51	0.32	0.327	44.1
Clarifier Liquid	4/20/72	0.013				0.93			
Hold Tank Effluent	4/20/72	0.738	2.05	3.73	0.241	1.44	0.61	0.201	41.9
Marble Bed: Front	4/20/72	0.983	1.53	3.67	0.315	1.05	0.48	0.210	51.4
Marble Bed: Back	4/20/72	0.868	1.47	3.40	0.290	0.97	0.50	0.150	54.1
Scrubber Bottoms S	4/20/72	2.53	1.36	4.87	0.364	1.20	0.16	0.249	44.3
Scrubber Spray	4/20/72	0.694	2.00	3.77	0.251	1.42	0.58	0.181	48.8
Set 2									
Scrubber Liquid Tk 1	4/20/72	0.72	1.46	2.72	0.251	0.93	0.53	0.163	55.4
Scrubber Liquid Tk 2	4/20/72	0.703	1.42	3.08	0.290	0.95	0.47	0.204	58.3
Scrubber Liquid Tk 3	4.20/72	0.697	1.41	4.46	0.366	0.94	0.47	0.158	58.8
Scrubber Bottoms Tk 1	4/20/72	2.03	1.79	3.05	0.287	1.26	0.53	0.278	45.2
Scrubber Bottoms Tk 2	4/20/72	2.06	1.73	4.53	0.367	1.44	0.29	0.310	45.2
Scrubber Bottoms Tk 3	4/20/72	2.06	1.59	4.28	0.362	1.24	0.35	0.346	44.0
Hold Tank Effluent	4/20/72	0.665	1.93	3.61	0.249	1.30	0.63	0.217	47.9
Marble Bed: Front	4/20/72	0.843	1.26	3.70	0.318	1.03	0.23	0.223	54.2
Marble Bed: Back	4/20/72	0.756	1.21	3.21	0.320	0.88	0.33	0.192	55.7
Scrubber Bottoms S	4/20/72	2.26	1.21	4.86	3.84	1.16	0.05	0.299	49.]
Scrubber Spray	4/20/72	0.699	1.79	3.70	0.250	1.28	0.51	0.192	49.1
SCIUDDEI SPIUS	1, 20, 12								

TABLE F-9. RESULTS OF SOLID PHASES ANALYSES EXPERIMENT 21R

Sample Location	Date	Wt % Solids in Slurry	Total S	Ca	Mg	s0 ₂		co ₂	Weight % Undissolved
Set 1									
Hold Tank Effluent	4/26/72	8.02	3.53	4.37	. 245	2.27	1.26	0.250	36.2
Marble Bed: Front	4/26/72	8.09	3.46	4.46	. 216	2.26	1.20	0.235	36.7
Marble Bed: Back	4/26/72	9.10	3.65	4.50	. 200	2.40	1.25	0.227	34.7
Clarifier Bottoms	4/26/72		2.65	4.21	. 295	1.66	0.99	0.394	32.9
Set 2									
Hold Tank Effluent	4/26/72	6.67	3.20	4.11	.186	2.35	0.85	0.200	35.9
Marble Bed: Front	4/26/72	7.65	3.38	4.11	.184	2.28	1.10	0.241	36.5
Marble Bed: Back	4/26/72	7.83	3.35	4.51	. 194	2.22	1.13	0.211	36.0
HAIDIE DEG. DACK	., _0, , _								

TABLE F-10. RESULTS OF SOLID PHASES ANALYSES EXPERIMENT 22R

Sample Location	Date	Wt % Solids in Slurry	Total S	Ca	Mg	s0 ₂	s0 ₃		Weight % Undissolved
Set 1									
Hold Tank Effluent Marble Bed: Front Marble Bed: Back	4/28/72 4/28/72 4/28/72	8.58 7.84 8.51	3.55 3.61 3.83	4.37 4.44 4.52	.190 .200 .188	2.32 2.35 2.52	1.23 1.26 1.31	0.205 0.196 0.243	35.3 35.7 35.1
Set 2									
Marble Bed: Front Marble Bed: Back Additive	4/28/72 4/28/72	9.18 9.82	3.75 3.85 0.50	4.44 4.53 6.09	.172 .155 .483	2.40 2.51 0.08	1.35 1.34 0.42	0.146 0.133 0.469	34.9 34.6 47.4

TABLE F-11. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 17R

	C-4			Cat	+				Tot	tal S	so ₃			T-4	pН	Temp
	Set No.	Time	Sampling Point	RAD.	C-E	Mg++	Na+	co_=	RAD.	C-E	RAD.	C-E	C1-	Tot. N	Low / High	.C
	1	1331	Marble Bed Back	18.6	19.3				25.5	25.84	8.57	8.9				
	1	1322	Marble Bed Front	22.0	18.8	0.20	0.76	1.10	22.2	21.98	8.95	9.0	2.65	0.73	4.55/ 5.5	49.5
	1	1347	Scrubber Bottoms (Scrubber)	16.8	16.8			0.27	12.9	12.61	1.26	1.0	2.81	0.73	10.6/ 10.8	47.0
	1	1400	Spraywater	12.6	13.7	0.01	0.74	0.13	10.3	11.27	0.84	1.0	2.13		11.18	37.5
	1	1445	Scrubber Liquid at Tank		19.9					23.75		8.9			5.78/ 5.86	
	1	1457	Scrubber Bottom at Tank		17.4					11.30		1.0			11.15/11.22	
	1	1537	Hold Tank Effluent	12.6	11.7	0.07	0.68	0.26	10.0	10.90	1.30	1.5	2.06		10.75	37.5
	1		Make Up Water	1.08			0.45									
F-13																
Ξ	2	1701	Marble Bed Back	21.45	20.0		0.75		25.2	24.06	9.23	8.6			4.45/ 4.75	47.5
	2	1650	Marble Bed Front	19.0	18.2	0.27	0.68	0.87	21.6	21.08	7.57	5.8	2.29		4.52/ 5.08	46.5
	2	1712	Scrubber Bottom (Scrubber)	16.0	16.3		0.74	0.20	12.6	13.00	1.30	1.7	2.39		9.9/ 10.5	46.5
	2	1725	Spray Water	13.5	12.8	0.02	0.68	0.10	9.6	10.68	0.89	1.3	11.02			37.0
	2	1505	Scrubber Liquid at Tank	19.8	19.6	0.35	0.77	0.96	22.3	21.95	8.06	9.2	2.63		5.78/ 5.86	50.5
	2	1517	Scrubber Bottoms at Tank	17.5	17.6	0.01	0.77	0.06	10.7	12.74	1.00	1.1	2.81		11.15/11.22	48.0
	. 2	1537	Hold Tank Effluent	12.6	11.7	0.07	0.68	0.26	10.0	10.90	1.30	1.5	2.06		10.75	36.5
	2	1605	Clarifier Bottoms	16.2	17.2		0.66	0.12	8.5	10.00	0.58	0	0.73		11.85	25.0
	2		Make Up Water	1.08			0.45									

TABLE F-12. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 18R

Set			Ca+	+				so ₄	; 	so ₃ =				рН	_
No.	Time	Sampling Point	RAD.	C-E	Mg++	Na+	co=	RAD.	C-E	RAD.	C-E	C1-	Tot. N	Low / High	Temp °C
1	1905	Marble Bed Back	24.4	24.80	3.60	1.10		19.47	24.64	7.43	3.85			5.75	45.0
1	1854	Marble Bed Front	22.8	24.10	3.70	1.05		21.25	24.04	2.55	2.06	6.44		6.23	43.0
1	1913	Scrubber Bottoms (Scrubber)	25.7	24.42		1.14		17.41	19.09	0.74	0.45	6.89		10.6	47.0
1	1923	Spray Water	23.5	22.52		1.10		17.56	18.15	0.64	0.35	6.11		10.75	46.0
1	1757	Scrubber Liquid at Tank	20.6	21.25	3.41	1.10		18.56	19.98	3.64	2.57	6.33		6.45	47.9
1	1805	Scrubber Bottom at Tank	33.2	32.72		1.11		17.20	17.20	1.10	0.89	6.91		11.4	47.9
1	1838	Hold Tank Effluent	23.3	22.97		1.06		17.5	17.35	0.80	0.53	6.17		10.75	46.0
ויק	1833	Clarifier Liquid	20.9	20.60		0.93		16.38	16.49	0.57	0.56	3.91		11.05	37.5
F-14															
2	2022	Marble Bed Back	24.7	26.12	3.40	1.11		22.26	22.33	5.14	5.75			6.0	45.0
2	2030	Marble Bed Front	26.0	26.87	3.42	1.13		22.73	27.97	7.47	2.60	6.49		6.05	45.0
2	2040	Scrubber Bottoms (Scrubber)	25.5	24.95		1.17		18.60	19.02	0.92	0.42			10.45	47.0
2	2050	Spray Water	23.2	23.25		1.09		18.6	19.67	0.68	0.38			10.4	47.0
2	1933	Scrubber Liquid at Tank	21.8	22.17	3.40	1.12		19.78	20.46	4.92	3.25	6.45		5.90	47.8
2	1939	Scrubber Bottom at Tank	34.30	34.30		1.14		17.69	18.51	0.91	0.72	7.15		11.46	47.5
2	2011	Hold Tank Effluent	23.9	24.20		1.11		19.0	18.68	1.37	0.55	6.28		10.6	46.0
2	2067	Clarifier Liquid	22.0	22.35		0.94		17.1	16.96	0.73	0.40	4.40		11.2	37.9

TABLE F-13. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 19R

Set			Ca++	·				so ₄ =		so ₃			Tot.	рН	Temp
No.		Sampling Point	RAD.	C-E	Mg++	Na+	$\overline{\omega_{z}^{3}}$	RAD.	C-E	RAD.	C-E	C1-	N N	Low / High	.c.
1	1215	Marble Bed Rock	34.4		4.52		1.83	26.8		26.6				4.5	41.5
1	1208	Marble Bed Front	34.4		4.49		1.86	26.1		27.1		2.32		4.7	47.0
1	1225	Scrubber Bottoms (Scrubber)	27.7		4.20		0.83	24.45		14.75		3.24		5.6	44
1	1230	Spray Water	25.7		4.67		1.12	20.35		15.55		2.13		5.5	39
1	1125-1145	Scrubber Liquid at Tank	35.3		4.50			25.1		28.6		1.33		4.98	43.4
1	1130-1152	Scrubber Bottom at Tank	23.8		4.38		0.75	22.5		7.9		3.36		5.88	43.8
1	1200	Hold Tank Effluent	24.5		4.30		1.10	20.6		15.7		2.10		5.43	39.0
1	1157	Clarifier Liquid	22.5		3.41		0.24	23.1		6.0		2.00		5.30	23.5
1		Make Up Water	1.08			.45									
F-15															
⁵⁵ 2	1340	Marble Bed Back	34.8		4.37		1.97	27.45		25.45				4.65	42.0
2	1 330	Marble Bed Front	34.5		4.40		1.83	25.55		26.35		2.30		4.90	41.50
2	1345	Scrubber Bottom (Scrubber)	276		4.24		0.83	23.65		13.65		3.25		5.6	44.0
2	1 325	Spray Water	25.2		4.23		1.00	21.45		15.15		2.08		5.5	37.0
2	1243-1305	Scrubber Liquid at Tank	35.3		4.41		1.36	25.2		27.31		2.26		5.02	44
2	1248-1310	Scrubber Bottom at Tank	22.2		4.07		0.17	21.9		4.5		3.17		6.15	44
2	1 320	Hold Tank Effluent	25.5		4.23		0.97	20.85		15.05		2.03		5.50	39
2	1315	Clarifier Liquid	22.6		3.45		0.22	22.85		5.95		2.03		5.60	23.5

TABLE F-14. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 20R

Set			Ca++	+				so <mark>=</mark>		so ₃			Tot.	рН	Temp
No.	Time	Sampling Point	RAD.	C~E	Mg++	Na+	co ₃	RAD.	C-E	RAD.	C-E	C1-	N N	Low / High	, c.
1	1105	Marble Bed Back	32.5		3.59		1.69	31.25		15.25				4.53	42
1	1140	Marble Bed Front	32.0		3.44	0.40	1.49	30.15		14.75		2.23		4.7	42.5
1	1200	Scrubber Bottom at Scrubber	26.1		3.32		1.38	25.3		8.1		3.01		5.93	44
1	1208	Scrubber Spray	23.4		3.18		0.93	24.7		7.9		1.89		5.72	39.0
1	1036-1105	Scrubber Liquid at Tank	31.6		3.20	0.38		30.1		15.6				5.21	44
1	1048-1110	Scrubber Bottom at Tank	21.6		3.30		0.06	18.9		1.01		3.11		10.9	45.3
1	1125	Hold Tank Effluent	23.6		3.24	0.38	0.95	24.45		7.45		2.03		5.75	40
1		Make Up Water	1.08			. 45									
2	1340	Marble Bed Back	33.8		3.73		1.27	29.15		17.05				4.55	42.5
2	1350	Marble Bed Front	33.1		3.80		1.74	30.25		16.65		2.38		4.70	40
2	1357	Scrubber Bottom at Scrubber	30.0		3.62		0.70	26.6		12.0		3.13		5.70	42.5
2	1330	Scrubber Spray	26.6				0.98	25.5		10.2		2.03		5.87	44.5
2	1232-1300	Scrubber Liquid at Tank	33.9		3.65		1.42	29.1		18.3		2.04		5.10	43.5
2	1240-1307	Scrubber Bottom at Tank	24.7		3.58		0.48	23.3		2.85		3.12		6.30	44.5
2	1323	Hold Tank Effluent	25.9		3.57		0.87	24.65		10.25		2.13		5.70	40.0
2	1320	Clarifier Liquid	22.9		3.28		0.57	20.9		7.4		1.95		5.80	39.0

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TABLE F-15. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 21R

C-+			Ca+	+				S02	; ↓	so ₃			Tot.	рН	Temp
Set No.	Time	Sampling Point	RAD.	C-E	Mg++	Na+	∞ <u>³</u>	RAD.	C-E	RAD.	C-E	C1-	N	Low / High	°C
1	1430	Marble Bed Back	22.6	24.91	16.0		1.94	33.6	31.15	7.60	10.9			5.48	46.0
1	1422	Marble Bed Front	23.6	23.05	16.8		1.98	27.8	29.30	6.70	9.50	9.22		5.69	44.0
1	1440	Scrubber Bottom at Scrubber	20.0	21.75	10.5			25.9			1.60			8.18	47.5
1	1500	Scrubber Spray	16.8	21.79	13.9		0.23	28.1	24.82	1.28	0.90	8.70		8.30	45.5
1	1320-1350	Scrubber Liquid at Tank	24.7	30.20	16.0	1.11	0.18	27.4	25.50	5.05	0.70	9.19		6.01	46.8
1	1327-1355	Scrubber Bottom at Tank	24.3	28.75	1.29		0.23	21.6	21.30	0.9	1.00	10.7		9.77	45.0
1	1405	Hold Tank Effluent	18.1		11.7	1.06	0.28	26.7		1.10		8.85		8.52	45.0
1	1400	Clarifier Liquid	19.4	20.8	4.30		0.3	21.3	22.40	0.85	1.10	3.94		9.85	29.0
ח															
7 2	1642	Marble Bed Back	23.6	25.45	16.6		2.04	34.4	32.50	1.20	11.65			5.38	46.5
2	1633	Marble Bed Front	20.5	24.72	16.8		1.89	26.8	29.40	1.29	9.14	9.13		5.50	44.0
2	1652	Scrubber Bottom at Scrubber	19.4	20.90	12.0		0.52	27.9	25.60	1.12	1.25	10.0		6.42	47.8
2	1700	Scrubber Spray	16.9	18.59	13.9		0.15	27.8	25.82	1.32	1.00	8.56		8.90	46.0
2	1528-1605	Scrubber Liquid at Tank	19.8	21.17	17.3		1.42	34.4	24.70	7.6	2.60	9.00		5.37	47.2
2	1535-1610	·	21.1	25.80	9.8		0.19	27.8	16.33	1.2	1.35	10.3		9.48	46.0
2	1623	Hold Tank Effluent	16.9		15.1	1.09	0.50	28.8		1.73		8.73	0.3	7.03	45.0
2	1618		17.6	20.7	4.72		0.29	20.8	22.35	0.98	1.50	4.60		9.90	29.9

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TABLE F-16. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 22R

Set			Ca	++				so <mark>-</mark>	: 	SO.	= 3		T-4	рН	T
No.	Time	Sampling Point	RAD.	C-E	Mg++	Na+	co ₃	RAD.	C-E	RAD.	C-E	C1~	Tot. N	Low / High	Temp °C
1	1605	Marble Bed Back		22.10	19.9		0.90		35.05		8.05			5.44	42.5
1	1555	Marble Bed Front		21.75	19.9		1.44		32.10		9.64	9.81		5.35	44.0
1	1610	Scrubber Bottom at Scrubber		22.05	19.0		2.09		27.6		7.19	10.4		5.67	45.5
1	1620	Scrubber Spray		17.98	19.8		0.62		31.10		3.75	9.75		6.05	45.0
1	1335	Scrubber Liquid at Tank		27.20	18.5		0.79		39.50		18.55	9.30		5.27	46.0
1	1330	Scrubber Bottom at Tank		19.58	15.5		1.38		30.43		4.75	9.92		6.50	46.5
1		Hold Tank Effluent		16.4	17.9	1.15	0.65		29.77		1.93	9.56	0.3	6.81	45.5
<u>ן</u>	1340	Clarifier Liquid		21.10	5.20		0.23	22.55	21.50		0.75	5.58		8.88	28.0

TABLE F-17. CHEMICAL ANALYSIS OF CANNED SAMPLES OF ADDITIVE FROM ST. LOUIS (Wt%)

Boiler Calcined Limestone and Flyash

Sample #	SiO ₂	A1203	Fe ₂ 0 ₃	CaO	Mg0	Na ₂ 0	K ₂ 0	TiO ₂	<u>so</u> 3
LF1	29.1	10.3	9.0	39.2	0.6	0.3	1.1	0.5	4.6
LF2	31.6	11.8	10.0	29.6	2.8	0.7	1.6	1.1	5.0
LF3	33.9	11.3	9.3	34.6	3.0	0.5	1.4	0.5	3.6
LF4	31.4	10.5	8.3	36.1	3.0	0.5	1.1	1.3	4.0
LF5	27.4	10.1	8.0	36.6	3.1	0.7	1.1	0.9	4.4
LF6	15.8	7.4	7.5	44.9	2.8	0.4	0.9	0.7	4.4
LF7	20.8	10.4	7.4	37.2	3.6	0.8	1.2	0.7	4.6
		<u>Boiler</u>	Calcined	<u>Dolomit</u>	e and F	lyash*			
DF1	52.0	24.8	6.4	1.5	1.5	0.3	3.0	1.3	0.3
DF2	39.1	17.8	10.5	3.8	1.0	0.3	1.9	1.1	1.2
DF3									
	50.2	25.8	6.4	1.9	1.4	0.5	2.7	1.8	0.4
DF4	50.2 28.4	25.8 16.0	6.4 6.6	1.9 25.7	1.4 7.4	0.5	2.7 2.0	1.8	0.4 2.3
DF4			6.6						
DF4			6.6	25.7					

^{*} Although these samples were marked as Boiler Calcined Dolomite and Flyash, there appears to be very little dolomite in DF1, DF2, and DF3.

These samples were taken randomly and the sample numbers are for C-E's use only.

TABLE F-17 (Continued). CHEMICAL ANALYSIS OF BAGGED BOILER CALCINED ADDITIVE AND FLYASH (Wt%)

Sample #	Ca0	Mg0	SiO ₂	A1203	Fe ₂ 0 ₃	Na ₂ 0	S0 ₃	Ti0 ₅
04012	33.6	2.6	26.1	14.3	6.9	0.3	trace	0.4
04009	31.9	3.3	20.0	13.0	7.7	0.7	3.6	0.6
05011	31.2	2.4						
05012	30.9	2.5						
05013	29.7	2.3						
05014	30.9	2.5						
05015	30.9	2.4						
05016	31.8	2.4						
05017	32.4	2.4						

TABLE F-18. CHEMICAL ANALYSIS OF ADDITIVE SAMPLES FROM UNION ELECTRIC

Canned Boiler Calcined Limestone and Flyash (m mole/g)

Sample Number	CaO	S0 ₃
		
LF1	7.00	0.58
LF2	5.29	0.625
LF3	6.18	0.45
LF4	6.45	0.50
LF5	6.54	0.55
LF6	8.02	0.55
LF7	6.64	0.54

Bagged Boiler Calcined Limestone and Flyash (m mole/g)

04012	6.00	-
04009	5.70	0.45
05011	5.57	-
05012	5.52	-
05013	5.30	-
05014	5.52	_
05015	5.52	-
05016	5.68	_
05017	5.79	-

APPENDIX G

LIMESTONE FURNACE INJECTION SYSTEM TESTS
PROBLEMS AND MODIFICATIONS

APPENDIX G

LIMESTONE FURNACE INJECTION SYSTEM TESTS -- PROBLEMS AND MODIFICATIONS

Mechanical Performance: During the furnace injection tests several problems were encountered, resulting in lost down and correction time. The following summarizes these problems:

Additive Injection System: The additive feed into the system could not be maintained initially due to constant tripping of the fan motor used to blow the additive into the inlet gas stream. Since there was a vacuum being pulled on the discharge side of the blower, a high gas flow resulted which overloaded the motor. The problem was solved by installing an orifice on the suction side of the blower.

Scrubber Bottom Tank: The outlet line was frequently plugged as a result of solids settling due to low flow rate and lack of mixing. The problem was solved by installing a mixer and a recycle line to increase the flow rate.

 ${
m SO}_2$ Analyzer: Dust from the gas inlet leaked into the ${
m SO}_2$ analyzing system despite the availability of a filter. This problem was corrected first by introducing the sampling probes ahead of the point of additive injection, and then by shielding it with a piece of pipe with the pipe cross-sectional area perpendicular to the gas flow.

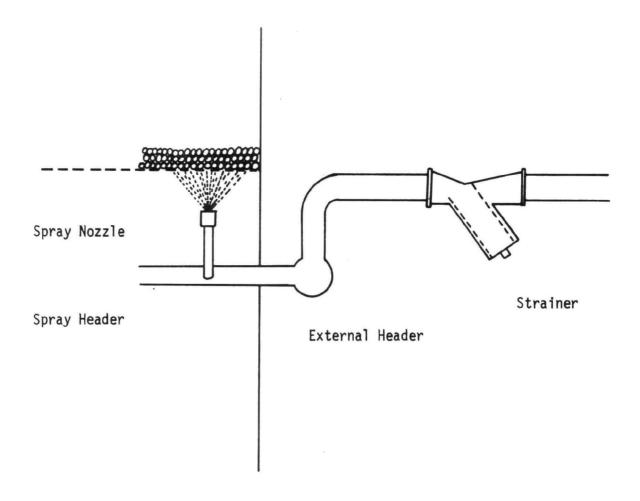
Nozzle Erosion and Plugging: Nozzle erosion was an operating problem since the nozzle material (brass) was not a good erosion resistant material. Nozzle erosion resulted in a change in the spray pattern, and therefore a disruption of liquid distribution. This problem was corrected by replacing the eroded nozzles.

Spray nozzle plugging was a persistent problem during the last test (18R) with boiler calcined limestone. This nozzle plugging was suspected to be the result of either or both of the following:

- a. Presence of particles larger than the nozzle orifice diameter.
- b. Maldistribution of fine particles in the three spray headers.

The first type of plugging was caused by particles with sizes larger than the spray nozzle orifice diameter (1/8"). When the nozzles were blocked with large particles, subsequent buildup of fine particles resulted. The source of large particles was suspected to be the additive itself, and chips peeling off both the Hold Tank and pipe walls and then carried to the nozzles in the slurry. A screen was installed on the outlet of the additive feeder to prevent the large particles in the additive from getting into the system. The second source was eliminated by installing a strainer in the slurry feedline to the scrubber as shown in Figure G-1. The strainer prevented large particles from the peeling of previous scale in both the Tank and pipes from reaching the spray nozzles.

The second type of nozzle plugging was a result of maldistribution of fine additive particles in the three headers. As the feed slurry, a, in the main external header turned 90° to enter spray header No. 1 (shown in Figure G-2), a centrifugal force would pull some of the solids toward spray headers No. II and III and hence increase the solid concentration in Stream b.



Large Particle Eliminator in the Spray Water

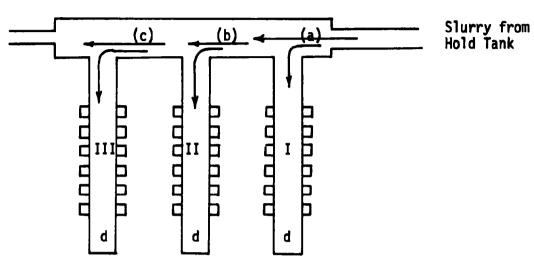
By the same token Stream c was higher in solid concentration than b and c. Thus the solid concentration in spray header III was greater than II, which in turn was greater than I. This was evident in the nozzle plugging pattern. To eliminate this maldistribution in the headers, the external spray header was modified to receive the slurry feed in two locations rather than one as shown in Figure G2. This improved the solids distribution in the three spray headers, and eliminated plugging. Also the dead space labeled (d) in Figure G-2 at the end of each spray header was removed to avoid buildup which eventually would plug the nozzles.

Heat Extractor Plugging: After the last EPA test series, deposit formation was noticed on the gas and liquid sides of the heat extractor tubes. The deposit on the gas side was mechanical (oil ash) in nature, and was cleaned by means of high pressure water. The deposit on the water side was chemical in nature and was cleaned using dilute acid. The heat Extractor and the duct cleanup was made as part of a maintenance repair to keep the equipment in good enough condition to carry out test work.

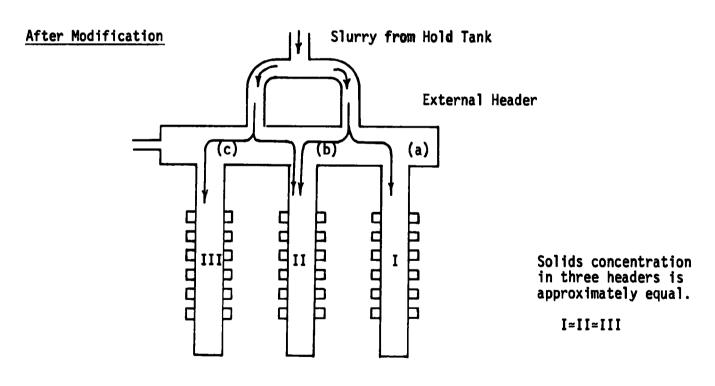
Air Leakage Due the Additive Injection System: The 0_2 concentration at the scrubber inlet is considerably higher than at the boiler because of the air leakage into the system between the scrubber and the boiler. Thus, with a minimum excess 0_2 of 5% in the boiler, the 0_2 concentration in the scrubber inlet was about 10%. Therefore, under these conditions, Test 21R had to be eliminated since it called for 5 percent 0_2 in the flue gas at the scrubber inlet.

Before Modification

External Header



Dead Zone Accumulating
Fine Particles
Solids Concentration in I<II<III



Dead Zone Eliminated

System Modification to Eliminate Fine Particle Nozzle Plugging

APPENDIX H

LIMESTONE FURNACE INJECTION SYSTEM

MATERIAL BALANCE

AND

RATE CALCULATIONS

TABLE H-1. TOTAL SULFUR MATERIAL BALANCE

Experiment 17R

		FI	ow Rate	Solid Content	Total S in Solid (m mole/g)	Total S in Liquid (m mole/1)	Total S in Gas (ppm)	Total S (m mole/ min)
Marble Bed (Set #1) Entering Streams	Spray Water Additive Gas In	2,010	l/min g/min g mole/min	0	0.55	11.27	1,456	4,576 1,106 15,943
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	329	g mole/min l/min l/min	2.635 10.2	0.869 0.677	23.41 12.61	764	8,866 8,455 1,991
	Total	Sul fur	In = 21,625	- Total S	ulfur Out = 19	9,312		
Marble Bed (Set #2) Entering Streams	Spray Water Additive Gas In	2,010	l/min g/min g mole/min	0	0.55	10.68	1,456	4,603 1,106 15,943
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	329	g mole/min l/min l/min	2.28 11.1	0.765 0.692	22.57 13.0	764	8,866 7,999 2,109
	Total	Sul fur	In = 21,652	- Total S	ulfur Out = 18	3,974		
Hold Tank (Set #1) Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	102	l/min l/min l/min	2,635 10.20	0.869 0.677	23.75 11.3		8,567 1,857
Leaving Streams	Hold Tank Eff.	639	1/min	3.27	1.73	10.9		10,580
	Total	Sul fur	In = 10,424	- Total S	ulfur Out = 10	,580		
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	102	l/min l/min l/min	2.65 8.77	0.95 1.012	21.95 12.74		8,050 2,205
Leaving Streams	Hold Tank	631	1/min	3.27	1.74	10.9		10,468

TABLE H-2. TOTAL SULFUR MATERIAL BALANCE

Experiment 18R

		Flow Rate	Solid Content g/l	Total S in Solid (m mole/g)	Total S in Liquid (m mole/l)	Total S in Gas (ppm)	Total S (m mole/ min)
Hold Tank (Set #1) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	682 l/min 114 l/min 76 l/min	47.3 60.0 0.17	3.09 2.57	22.55 18.09 17.05		115,058 19,641 1,295
Leaving Streams	Hold Tank Eff.	872 1/min	45.0	3.08	17.88		136,450
	Total	Sulfur In = 135,99	94 - Total	Sulfur Out = 1	136,450		
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	682 l/min 114 l/min 76 l/min	40. 4 55.5	3.21 2.49	23.71 19.23 17.36		104,614 17,946 1,319
Leaving Streams	Hold Tank Eff.	872 1/min	36.6	2.98	19.99		112,538
	Total	Sulfur In = 123,87	79 - Total	Sulfur Out =	112,538		
Marble Bed (Set #1) Entering Streams	Inlet Gas Scrubber Spray Additive	10,820 g mole/min 796 l/min 2,045 g/min	36.7	3.01 0.51	18.50	1,471	15,920 102,658 1,043
Leaving Streams	Outlet Gas Scrubber Liquid* Scrubber Bottom	11,630 g mole/min 682 l/min 114 l/min	41.4 53.9	3.34 2.51	27.30 19.54	447	5,200 112,923 17,650
	Total	Sulfur In = 119,62	21 - Total	Sulfur Out = '	135,773		
Marble Bed (Set #2) Entering Streams	Inlet Gas Scrubber Spray Additive	10,820 g mole/min 796 l/min 2,045 g/min	33.5	3.08 0.51	20.05	1,471	15,916 98,091 1,042
Leaving Streams	Outlet Gas Scrubber Liquid Scrubber Bottom	11,630 g mole/min 682 l/min 114 l/min	40.5 49.9	3.18 2.435	29.30 19.44	447	5,198 107,817 16,067
	Total	Sulfur In = 115,04	19 - Total	Sulfur Out =	129,082		

^{*}Average of Back and Front Marble Bed

TABLE H-3. TOTAL SULFUR MATERIAL BALANCE

Experiment 19R

		Flow Ra	Solid Conten te g/l		Total S in Liquid (m mole/l)	Total S in Gas (ppm)	Total S (m mole/ min)
Marble Bed (Set #1) Entering Streams	Gas In Scrubber Spray Additive	9,200 g mo 738 l/mi 2,800 g/mi	n 13.9	2.21 0.55	35.90	1,883	17,324 40,165 1,540
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	10,555 g mo 662 1/mi 76 1/mi	n 14.4	1.87 1.83	53.30 39.20	1,095	11,558 53,111 7,444
	Total	Sulfur In =	68,029 - Total	Sulfur Out = 7	2,113		
Marble Bed (Set #2) Entering Streams	Gas In Scrubber Spray Additive	9,120 g mo 738 1/mi 2,800 g/mi	n 14.60	2.0 0.55	36.60	1,881	17,155 48,560 1,540
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	10,487 g mo 662 l/mi 76 l/mi	n 15.90	1.73 1.82	52.40 37.30	1,124	11,787 52,898 8,630
	Total	Sulfur In =	67,255 - Total	Sulfur Out = 7	3,315		
Hold Tank (Set #1) Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	662 l/mi 76 l/mi 133 l/mi	n 28.70	1.68 2.17	53.7 30.4		49,118 7,044
Leaving Streams	Hold Tank Eff.	871 1/mi	n 13.90	2.26	36.3		58,979
	Total	Sulfur In =	56,162 - Total	Sulfur Out = 5	8,979		
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	662 l/mi 76 l/mi 133 l/mi	n 33.0	1.69 2.09	52.51 26.4		49,306 7,248
Leaving Streams	Hold Tank Eff.	871 1/mi	n 14.90	2.05	35.90		57,873
	Total	Sulfur In =	56,554 - Total	Sulfur Out = 5	,873		

TABLE H-4. TOTAL SULFUR MATERIAL BALANCE
Experiment 20R

	was the state of t	Flow Rate	Solid Content g/l	Total S in Solid (m mole/g)	Total S in Liquid (m mole/l)	Total S in Gas (ppm)	Total S (m mole/ min)
Marble Bed (Set #1) Entering Streams	Gas In Scrubber Spray Additive	9,400 g mole/min 757 l/min 2,800 g/min	6.94	2.0 0.50	32.6	1,962	18,443 35,185 1,400
Leaving Streams	Scrubber Liquid Scrubber Bottom Gas Out	681 1/min 76 1/min 10,850 g mole/min	9.25 25.3	1.50 1.53	45.7 33.4	1,090	40,571 5,480 11,827
	Total	Sulfur In = 55,02	8 - Total S	ulfur Out = 57	,878		
Marble Bed (Set #2) Entering Streams	Gas In Scrubber Spray Additive	9,400 g mole/min 757 l/min 2,800 g/min	6.99	1.78 .50	35.7	1,939	18,227 36,444 1,400
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	10,850 g mole/min 681 l/min 76 l/min	8.00 22.6	1.23 1.48	46.55 38.60	1,090	11,827 38,402 5,476
	Total	Sulfur In = 56,07	1 - Total S	ulfur Out = 55	,705		
Hold Tank (Set #1) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid Make Up Water	681 1/min 76 1/min 151 1/min 151 1/min	7.33 22.7 0.13	1.61 1.80	45.7 19.9 28.40		39,158 4,618 4,288
Leaving Streams	Hold Tank Eff.	1,059 l/min	7.38	2.05	31.90		49,804
	Total	Sulfur In = 48,064	4 - Total S	ulfur Out = 49	,804		
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid Make Up Water	681 1/min 76 1/min 151 1/min 151 1/min	7.07 20.5	1.43 1.69	47.4 26.1 28.3		39,164 4,617 4,273
Leaving Streams	Hold Tank Eff.	1,059 1/min	6.65	1.93	34.90		50,551
	Total	Sulfur In = 48,054	4 - Total S	ulfur Out = 50	,551		

TABLE H-5. RATE CALCULATION USING SOLID MATERIAL BALANCE

Experiment 17R

			Stream	Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/1)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
Mar	ble Bed (Set #1)								
1.	CaSO ₃ 1/2 H ₂ O Fo	ormation							
	Entering Streams		-		-	•	-	-	•
	Leaving Streams	Scrubber Liquid* Scrubber Bottom		l/min l/min	2.635 10.200	0.80 0.59			693 614
		Rate of CaSO ₃ 1	/2 H ₂ 0 I	Formation =	CaSO ₃ 1/2 H ₂ O	(Out) - CaS	ю ₃ 1/2 н ₂ 0 (In)	
				=	693 + 614 = 1	307 m mole/m	in.		
2.	SO ₂ Oxidation								
	Entering Streams	Inlet Flue Gas Scrubber Spray		g mole/min l/min			1.0	1,456	15,943 406
	Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	329	g mole/min l/min l/min			8.95 1.0	764	8,866 2,944 102
		Rate of SO ₂ Oxid	ation =	SO ₂ (In) -	so ₂ (Out) - 0	CaSO ₃ 1/2 H ₂ O	Formation R	ate	
			=	16,349 - 11	,912 - 1,307	= 3,130 m mc	le/min.		
١.	Ca SO ₄ 2 H ₂ O Form	mation							
	Entering Streams	(Additive)	2,010	g/min		0.55			1,105
	Leaving Streams	Scrubber Liquid* Scrubber Bottom		l/min l/min	2,635 10.2	0.069 0.087			59.8 90.5
		Rate of Ca SO ₄ 2	H ₂ O For	mation = Ca	350 ₄ 2 H ₂ 0 (Ou	it) - CaSO ₄ 2	2H ₂ O (In)		
				= 9	90.5 + 59.8 -	1,105 = 150	- 1,105		
				= -9	955 m mole/min	ı			
١.	CaCO ₃ Formation								
	Entering Streams	(Additive)	2,010	g/min		0.39**			784.0
	Leaving Streams	Scrubber Liquid Scrubber Bottom		l/min l/min	2.635 10.2	0,220 0,362			191 377
		Rate of CaCO ₃ Fo	rmation	= CaCO ₃ (Ou	ut) - CaCO ₃ (1	in)			
				= 191 + 377	7 - 784 = -216	s m mole/min			
.	Ca(OH) ₂ Dissolut	ion							
	Entering Stream	Scrubber Spray	431	1/min ⁺			13.7		5,905
	Leaving Streams	Scrubber Liquid Scrubber Bottom		l/min l/min			19.0 16.8		6,251 1,714
		Rate of Ca(OH) ₂	Dissolut	tion = Ca(1i	iq.) Out - Ca((liq.) In + (CaCO ₃ + CaSO ₄	2 H ₂ 0 +	CaSO ₃ 1/2 H ₂ 0
		-		= 6.25	1 + 1.714 - 5,	,905 + (1.307	7 - 955 - 216	5)	- -
				- 2,196	5 m moles/min.	•			

^{*}Average of marble bed front and back
**From Radian Corp.
+Subtotal of Scrubber Bottom + Scrubber Liquid

		Stream Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. Species In Gas Flow Rate (PPM) m mole/min
Hold Tank (Set #1)						
1. CaSO ₃ 1/2 H ₂ O Fo						
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 l/min 102 l/min 208 l/min	2.635 10.20 -	0.80 0.59 -		693 614
Leaving Streams	Hold Tank Eff.°	639 1/min [†]	3.27	1.51		3,155
	Rate of Ca $S0_3$	1/2 H ₂ O Formation =	CaSO ₃ 1/2 H	₂ 0 (Out) - C	aso ₃ 1/2 H ₂	0 (In)
		=	3,155 - 693	- 614		
		-	1,848 m mole	s/min.		
2. SO ₂ Oxidation						
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min			8.9 1.0	2 , 928 102
Leaving Streams	Hold Tank Eff.°	639 1/min ⁺			1.5	958
	Rate of SO ₂ Oxid	ation = SO ₂ (liq.)	In - SO ₂ (liq	.) Out - CAS	о _з 1/2 н ₂ 0	Formation Rate
		= 2,928 + 102	- 958 - 1,84	8		
		= 224 m mole/	min.			
3. CaSO ₄ 2H ₂ O Form	ation					
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min	2.635 10.2	0.069 0.087		60 90
Leaving Stream	Hold Tank Eff.	639 1/min	3.27	0.23		481
	Rate of CaSO ₄ 2	H ₂ O Formation = CaS	0 ₄ 2H ₂ 0 (Out) - CaSO ₄ 2	H ₂ 0 In	
		= 481	- 90 - 60 = 3	331 m mole/m	in.	
4. CaCO ₃ Formation						
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min	2.635 10.2	0.220 0.362		191 377
Leaving Stream	Hold Tank Eff.	639 1/min	3.27	0.650		1,358
	Rate of CaCO ₃ For	rmation = CaCO ₃ (Ou	t) - CaCO ₃ (Ir	n)		
		= 1,358 - 3	77 - 191 = 790	O m mole/min	•	
5. Ca(OH) ₂ Dissoluti	on					
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min			19.9 17.4 1.08	6,547 1,774 224
Leaving Stream	Hold Tank Eff.	639 1/min			11.7	7,476
	Rate of Ca(OH) ₂	Dissolution = Ca(lie	q.) Out - Ca(1	liq.) In + Ca	aCO ₃ + CaSO ₄	2H ₂ 0 + CaSO ₃ 1/2 H ₂ 0
		= 7,476	- 6,547 - 1,7	774 - 224 +	1,848 + 331	+ 790

= 1,900 m moles/min

⁺Subtotal of Scrubber Bottom + Scrubber Liquid oStream characterization is assumed to be the same as in Set 2 since the sample was taken in between the two sets.

TABLE H-5. RATE CALCULATION USING SOLID MATERIAL BALANCE (Continued)

44-4		Stream	Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
Marble Bed (Set #2))							
1. CaSO ₃ 1/2 H ₂ O Fo	ormation							
Entering Stream		-		-	-			
Leaving Streams	Scrubber Liquid Scrubber Bottom		l/min l/min	2.28 11.10	0.30 0.50			225 566
	Rate of CaSO ₃ 1/	2 H ₂ 0 Fc	ormation = (CaSO ₃ 1/2 H ₂ O	(Out) - CaSO	3 1/2 H ₂ 0 (1	in)	
			= 7	'91 m mole/min	ı			
2. SO ₂ Oxidation								
Entering Stream	Inlet Flue Gas Scrubber Spray	431	g moles/mir l/min moles/min	1		1.3	1,456 764	15,943 560 8,866
	Outlet Flue Gas Scrubber Liquid Scrubber Bottom	329	nores/min 1/min 1/min			7.2 1.7	704	2,368 173
	Rate of SO ₂ Oxid	ation =	SO ₂ (In) -	SO ₂ (Out) - 0	aso ₃ 1/2 H ₂ 0	Formation F	late	
		=	15,943 + 56	50 - 8,866 - 2	,368 - 173 -	791 = 4,305	m mole/n	nin
3. CaSO ₄ 2H ₂ O Forma	ntion							
Entering Stream	Additive	2,010	g/m1n		0.55			1,105
Leaving Streams	Scrubber Liquid Scrubber Bottoms		l/min l/min	2.28 11.1	0.465 0.192			349 217
	Rate of CaSO ₄ 2H	2 ^{0 Forma}	ation = CaS(0 ₄ 2H ₂ O (Out)	- CaSO ₄ 2H ₂ O	(In)		
			= 217	+ 349 - 1,105	5 = - 539 m m	nole/min		
4. CaCO ₃ Formation								
Entering Stream	Additive	2,010	g/min		0.39			784
Leaving Stream	Scrubber Liquid Scrubber Bottom		l/min l/min	2.28 11.1	0.296 0.511			222 578
	Rate of CaCO ₃ Fo	rmation	= CaCO ₃ (Ou	ıt) CaCO ₃ (In	1)			
			= 222 + 578	3 - 784 = 16 m	n mole/min			
5. Ca(OH) ₂ Dissolut	tion							
Entering Stream	Scrubber Spray	431	1/min			12.8		5,516
Leaving Stream	Scrubber Liquid Scrubber Bottom		l/min l/min			19.1 16.3		628 1 , 662
	Rate of Ca(OH) ₂	Dissolut	tion = Ca(1	iq.) Out - Ca((liq.) In = (CaCO ₃ + CaSO ₄	1 2H ₂ 0 + 1	CaSO ₃ 1/2 H ₂ C
			= 6,28	1 + 1,662 - 5,	,516 + 791 -	539 + 16		

= 2,698 m mole/min

TABLE H-5. RATE CALCULATION USING SOLID MATERIAL BALANCE (Continued)

		Stream Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
Hold Tank (Set #2)							
I. CaSO ₃ 1/2 H ₂ O For	mation						
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min	2.65 8.77 -	0.52 0.83			453 742 -
Leaving Streams	Hold Tank Eff.°	639 1/min	3.27	1.51			3,155
	Rate of CaSO ₃ 1/	2 H ₂ O Formation = 0	CaSO ₃ 1/2 H ₂ O	(Out) - CaSO	3 1/2 H ₂ 0 (I	n)	
		= 3	3,155 - 453 -	742 = 1,960	m mole/min		
2. SO ₂ Oxidation							
Entering Stream	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min			9.2 1.7		3,026 173 -
Leaving Stream	Hold Tank Eff.	639 1/min			1.5		958
	Rate of SO ₂ Oxid	lation = SO ₂ (1iq)	In - SO ₂ (liq)) Out - CaSO ₃	1/2 H ₂ 0 For	mation R	ate
	•	_	3 - 958 - 1,96				
3. CaSO ₄ 2H ₂ O Format	tion						
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	329 1/min 102 1/min 208 1/min	2.65 8.77	0.43 0.182			375 163
Leaving Stream	Hold Tank Eff.	639 1/mi n	3.27	0.23			481
	Rate of CaSO ₄ 2H	1 ₂ 0 Formation = CaS	0 ₄ 2H ₂ O (In)				
		= 481	- 375 - 163	= 57 m mole/m	nin		
. CaCO ₃ Formation							
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	329 l/min 102 l/min 208 l/min	2.635 10.2	0.229 0.560			198 583
Leaving Streams	Hold Tank Eff	639 1/min	3.27	0.650			1,358
	Rate of CaCO ₃ Fo	ormation = CaCO ₃ (O	ut) - CaCO ₃ (In)			
		= 1,358 -	198 - 583 = 5	77 m mole/mir	ı		
6. Ca(OH) ₂ Dissoluti	ion —						
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up Water	329 l/min 102 l/min 208 l/min			19.6 17.6 1.08		6,448 1,795 225
Leaving Streams	Hold Tank Eff.	639 1/min			11.7		7,476
	Rate of Ca(OH) [issolution = Ca (1	iq.) Out - Ca	(liq.) In +	CaCO ₃ + CaSo	0 ₄ 2H ₂ O +	CaSO ₃ 1/2 H
		= 7,476	- 6,448 - 1,	795 - 225 + 3	,960 - 57 +	577 = 1,	488 m mole/m

		Stream Flow Rate	Slurry Salid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/1)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
Marble Bed (Set #1)				·			
1. CaSO ₃ 1/2 H ₂ O Form	nation						
Entering Streams	Scrubber Spray	738 1/min	13.94	1.62			16,618
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 1/min 76 1/min	14.4 32.10	1.26 1.47			12,011 3,586
	Rate of Ca SO ₃ 1/2	H ₂ 0 Formation = Cas	50 ₃ 1/2 Н ₂ 0 (0	ut) - CaSO ₃	1/2 H ₂ O (In)		
		= ((12,011 + 3,58	6) - 16,618			
		= 1	5,597 - 16,61	8			
		= -	-1,021 m mole/	min			
2. SO ₂ Oxidation							
Entering Streams	Inlet Flue Gas Scrubber Spray	9,200 g mole/min 738 l/min			15.55	1,883	17,324 11,476
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,555 g mole/min 662 l/min 76 l/min			26.8 14.75	1,095	11,558 17,742 1,121
	Rate of SO ₂ Oxidat	ion = SO ₂ (In) - SO ₂	, (Out) - CaSO	3 1/2 H ₂ 0 Fo	rmation Rate	!	
	-	= 28,800 - 30,	,421 + 1 , 021 =	-600 m mole	/min		
3. CaSO ₄ 2 H ₂ O Format	ton						
Entering Streams	(Additive) Spray Water	2,800 g/min 738 l/min	13.9	.55 .59			1,540 6,052
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 l/min 76 l/min	14.4 32.1	.61 .36			5,815 878
	Rate of CaSO ₄ 2H ₂ O	Formation = $Caso_4$ 2	2H ₂ O (Out) - C	aso ₄ 2H ₂ 0 (I	n)		
		= 6,693 -	7,592				
		= - 899 m	mole/min				
. CaCO ₃ Formation							
Entering Streams	(Additive) Spray Water	2,800 g/min 738 l/min	13.9	.39 .215			1,092 2,206
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 1/min 76 1/min	14.4 32.1	.171 .280			1,630 683
	Rate of CaCO ₃ Form	ation = CaCO ₃ (Out)	- CaCO ₃ (In)				
	·	= 2,313 - 3,	•				
		= -985 m mol	e/min				
. Ca(OH) ₂ Dissolutio	n						
Entering Stream		738 l/min			25.7		18,967
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 1/min 76 1/min			34.4 27.7		22,773 2,105
	Rate of Ca(OH) ₂ Dis	ssolution = Ca(liq) = 24,878	Out - Ca(11q) 3 - 18,967 + -		, ,	+ CaSO ₃	1/2 H ₂ 0 Form

ΔSame as the hold tank effluent +Subtotal of scrubber bottom and scrubber liquid

	Stream Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
ation						
Scrubber Liquid	662 1/min	12.20	1.32			10,661
Make Up Water Scrubber Bottom	133 1/min 76 1/min	28.70	1.55			3,381
Hold Tank Eff.	871 1/min	13.9	1,60			19,371
Rate of CaSO ₂ 1/2 H	,0 Formation = CaS	0 ₃ 1/2 H ₂ 0 (Ou	t) - CaSO ₃ 1	/2 H ₂ O (In)		
3	-	_				
	= 5,3	29 m moles/min	1			
Scrubber Liquid Scrubber Bottom	662 l/min 76 l/min 133 l/min			28.6 7.9		18,933 600
•	•			15.7		13,674
		- SO ₂ (liq.)	Out - CaSO ₂	1/2 H ₂ 0 Form	nation Rat	e
2	-	_	3	L		
	= 530 m moles/m	in				
on						
Scrubber Bottom T Scrubber Liquid T	76 l/min 662 l/min 133 l/min	28.70 12.20	.36 .62			785 5 , 008
Hold Tank Eff.	871 1/min	13.90	.66			7,991
Rate of CaSO ₄ 2H ₂ O	Formation = CaSO ₄	2H ₂ O (Out) CaS	50 ₄ 2H ₂ 0 (In))		
, .						
	= 2,198	m mole/min				
Scrubber Bottom T Scrubber Liquid T Make Up Water	76 l/min 662 l/min 133 l/min	28.70 12.20	.249 .173			543 1 , 397
Hold Tank Eff.	871 1/min	13.90	.172			2,082
Rate of CaCO ₃ Forma	ition = CaCO ₃ (Out)	CaCO ₃ (In)				
Ū	= 2,082 - 1,9	940				
	= 142 m mole/	'min				
n						
 Scrubber Bottom T Scrubber Liquid T Make Up Water	76 l/min 662 l/min 133 l/min			23,8 35.3 1.08		1,809 23,369 144
Hold Tank Eff.	871 1/min			24,5		21,340
Rate of Ca(OH), Dis	ssolution = Ca(lig.	.) Out - Ca(110	q.) In + (Ca	CO ₃ + CaSO ₄	2H ₂ O + Ca	50 ₃ 1/2 H ₂ 0
- · · L				•		
- 2		- 25,322 + (14			_	
	Make Up Water Scrubber Bottom Hold Tank Eff. Rate of CaSO ₃ 1/2 H Scrubber Bottom Make Up Water Hold Tank Eff. Rate of SO ₂ Oxidati on Scrubber Bottom T Scrubber Liquid T Make-Up Water Hold Tank Eff. Rate of CaSO ₄ 2H ₂ O Scrubber Bottom T Scrubber Liquid T Make Up Water Hold Tank Eff. Rate of CaSO ₃ Formation Scrubber Bottom T Scrubber Liquid T Make Up Water Hold Tank Eff. Rate of CaCO ₃ Formation Scrubber Bottom T Scrubber Liquid T Make Up Water Hold Tank Eff.	Scrubber Liquid 662 1/min Make Up Water 133 1/min 76 1/min Hold Tank Eff. 871 1/min Rate of CaSO ₃ 1/2 H ₂ O Formation = CaS = 19,	Scrubber Liquid 662 1/min 12.20 Make Up Water 133 1/min 28.70 Hold Tank Eff. 871 1/min 13.9 Rate of CaSO ₃ 1/2 H ₂ O Formation = CaSO ₃ 1/2 H ₂ O (Ou = 19,371 - 14,042 = 5,329 m moles/min Scrubber Liquid 662 1/min Make Up Water 133 1/min Hold Tank Eff. 871 1/min Rate of SO ₂ Oxidation = SO ₂ (1iq.) In - SO ₂ (1iq.) = 19,533 - 13,674 - 5,329 = 530 m moles/min On Scrubber Bottom 7 76 1/min 28.70 Scrubber Liquid 7 662 1/min 12.20 Make-Up Water 133 1/min - Hold Tank Eff. 871 1/min 13.90 Rate of CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O Formation = CaSO ₄ 2H ₂ O (Out) CaSO ₄ 2H ₂ O (Out) CaSO ₆ 2 1/min 12.20 Make Up Water 133 1/min - 13.90 Rate of CaCO ₃ Formation = CaCO ₃ (Out) CaCO ₃ (In) = 2,082 - 1,940 = 142 m mole/min Scrubber Bottom T 76 1/min Scrubber Liquid T 662 1/min Make Up Water 133 1/min Scrubber Bottom T 76 1/min Scrubber Liquid T 662 1/min Make Up Water 133 1/min Hold Tank Eff. 871 1/min 133 1/min Hold Tank Eff. 871 1/min	Stream Flow Rate	Stream Flow Rate Solid Cont. (g/1) In Solid m mole/g In Liquid m mole/g In Liquid In Liquid In Liquid In Make Up Mater 133 1/min 28.70 1.55	Stream Flow Rate Solid In Solid In Liquid In Game In Solid In Manager In Solid In Solid In Manager In Solid In Solid In Manager In Solid In Manager In Solid In Solid In Manager In Solid In Solid In Manager In Solid In Manager In Solid In Solid In Manager In Solid In Solid In Solid In Manager In Solid In Solid

= 3,687 m moles/min H-10

		Stream Flow Rate	Slurry Solid Cont. (g/1)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. In Gas (PPM)	Total Species Flow Rate <u>m mole/min</u>
Marble Bed (Set #2)							
1. CaSO ₃ 1/2 H ₂ O For		700 17 1	14.60	3 43			15,192
Entering Streams	Scrubber Spray	738 1/min	14.60	1.41			12,315
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 l/min 76 l/min	15.9 41.90	1.17 1.33			4,235
	Rate of CaSO ₃ 1/2	Formation = $Caso_3$ 1,	/2 H ₂ O (Out) -	CaSO ₃ 1/2 H	1 ₂ 0 (In)		
		= 16,550	- 15,192				
		= 1,358 m	mole/min				
2. SO ₂ Oxidation							
Entering Streams	Inlet Flue Gas Scrubber Spray	9,120 g mole/min 738 l/min			15.15	1,881	17,154 11,180
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,487 g mole/min 662 l/min 76 l/min			25.90 13.65	1,124	11,787 17,146 1,037
	Rate of SO ₂ Oxidat	ion = SO ₂ (In) - SO	₂ (Out) - CaSO) ₃ 1/2 H ₂ 0 Fe	ormation Rate	•	
•	_	= 28,334 - 29,9	70 - 1,358				
		= -2,994 m mole	/min				
3. CaSO ₄ 2H ₂ O Format	ion						
Entering Streams	Additive Spray Water	2,800 738 l/min	14.60		0.55 .60		1,540 6,464
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 l/min 76 l/min	15.90 41.90		.56 .49°		5,894 1,560
	Rate of CaSO ₄ 2H ₂ O	Formation = $Caso_4$	2H ₂ O (Out) - (Caso ₄ 2H ₂ 0 (In)		
		= 7,454	- 8,004				
		= - 550 1	nim/elom m				
4. CaCO ₃ Formation							
Entering Streams	Additive Spray Water	2,800 g/min 738 l/min	14.60	.39 .18			1,092 1,939
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 1/min 76 1/min	15.90 41.90	.22 .402			2,316 1,280
	Rate of CaCO ₃ Form	$ation = CaCO_3$ (Out)	- CaCO ₃ (In)				
		= 3,596 - 3,0	31				
		= 565 m mole/	min				
5. Ca(OH) ₂ Dissolution	o n 						
Entering Stream	Spray Water	738 1/min			25,2		18,598
Leaving Stream	Scrubber Liquid* Scrubber Bottom	662 1/m 76 1/min			34.7 27.6		22,971 2,098
	Rate of Ca(OH) ₂ Di	ssolution = Ca(liq.) Out - Ca(11	q.) In + CaC	0 ₃ + CaSO ₄ 2	H ₂ 0 + CaS	0 ₃ 1/2 H ₂ 0 For
		= 25,069	- 18,598 + (5	65 - 550 + 1	,358)		
		= 25,069	- 18,598 + (1	,373)			
		= 7,844 m	mole/min				

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TABLE H-6. RATE CALCULATION USING SOLID MATERIAL BALANCE (Continued)

Scrubber Liq. T Scrubber Bottom Make Up Water Hold Tank Eff. Rate of CaSO ₃ 1/2 H ₂		13.0 33.0 - 14.90	1,20 1,47			10,327
Scrubber Liq. T Scrubber Bottom Make Up Water Hold Tank Eff.	76 l/min 133 l/min 871 l/min 0 Formation = CaSC	33.0				
Scrubber Bottom Make Up Water Hold Tank Eff.	76 l/min 133 l/min 871 l/min 0 Formation = CaSC	33.0				
	O Formation = CaSC	14,90				3,687
Rate of CaSO ₃ 1/2 H ₂			1,45			18,818
) ₃ 1/2 H ₂ O (Ou	t) - CaSO ₃ 1	/2 H ₂ D (In)		
	= 18,0					
	= 4,80	04 m mole/min				
Scrubber Liquid T Scrubber Bottom T Make Up H ₂ O	562 1/min 76 1/min 133 1/min			27.31 4.5		18,079 342
Hold Tank Eff.	871 1/min			15.05		13,109
Rate of SO ₂ Oxidation	n = 50 ₂ (1iq) In -	· SO ₂ (liq) Ou	t - CaSO ₃ 1/	2 H ₂ 0 Form R	ate	
-	= 18,421 - 13,10	9 - 4,804				
	= 508 m mole/min	1				
on						
Scrubber Bottom T Scrubber Liq. T Nake Up Water	76 l/min 662 l/min 133 l/min	33.0 13.0	.49 .62			1,228 5,336
Hold Tank Eff	871 1/ min	14,90	.60			7,787
Rate of CaSO ₄ 2H ₂ O Fo	ormation = CaSO ₄ 2	2H ₂ O (Out) - C	aSO ₄ 2H ₂ O (I	n)		
	· · · · · · · · · · · · · · · · · · ·		. –			
	= 1,223 m	mole/min				
Scrubber Bottom Scrubber Liquid Make Up Water	76 l/min 662 l/min 133 l/min	33.0 13.0	.339 .19			850 1,635
Hold Tank Eff.	871 1/min	14.9	.189			2,453
Rate of CaCO ₃ Format	ion = CaCO ₃ (Out)	- CaCO ₃ (In)				
·	ū	•				
	= - 32 m mole/	min				
า						
Scrubber Liq T Scrubber Bottom T Make Up Water	662 1/min 76 1/min 133 1/min			35.3 22.2 1.08		23,369 1,687 144
Hold Tank Eff.	871 1/min			25.5		22,211
Rate of Ca(OH) ₂ Disso	olution = Ca(liq)	Out - Ca(liq)	In + (CaCO ₃	+ CaSO ₄ 2H ₂	0 + CaSO ₃	1/2 H ₂ 0)
_	= 22,211 -	- 25,200 + (-3	2 + 1,223 +	4,804)	,	_
	Scrubber Bottom T Make Up H ₂ O Hold Tank Eff. Rate of SO ₂ Oxidation Scrubber Bottom T Scrubber Liq. T Nake Up Water Hold Tank Eff Rate of CaSO ₄ 2H ₂ O Fo Scrubber Liquid Make Up Water Hold Tank Eff. Rate of CaCO ₃ Format Scrubber Bottom T Make Up Water Hold Tank Eff.	= 18,8 = 4,80 Scrubber Liquid T	= 18,818 - 14,014 = 4,804 m mole/min Scrubber Liquid T	= 18,818 - 14,014 = 4,804 m mole/min Scrubber Liquid T	Scrubber Liquid T 562 1/min 27.31 4.5 562 1/min 4.5 564 1/min 562 1/min 562 1/min 563 1/min 563 1/min 565 1/min	= 18,818 - 14,014 = 4,804 m mole/min Scrubber Liquid T

TABLE H-7. RATE CALCULATION USING SOLID MATERIAL BALANCE

Experiment 20R

		Stream Flow Rate	Slurry Solid Cont. (g/1)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/1)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
Marble Bed (Set #1)							
1. CaSO ₃ 1/2 H ₂ O For	rmation						
Entering Stream	Scrubber Spray	757 1/min	6.94	1.42			7,460
Leaving Stream	Scrubber Spray Scrubber Bottom	681 1/min 76 1/min	9.25 25.3	1.01 1.20			6,362 2,307
	Rate of CaSO ₃ 1/	2 H ₂ O Formation = Ca	SO ₃ 1/2 H ₂ O	(Out) - Caso	3 1/2 H ₂ 0 (1	n)	
	•	= 86	69 - 7460		-		
		= 12	09 m mole/min				
2. SO ₂ Oxidation							
Entering Stream	Inlet Flue Gas Scrubber Spray	9,400 g mole/min 757 l/min			7.9	1,962	18,443 5,980
Leaving Stream	Outlet Flue Gas Scrubber Liquid Scrubber Bottom	10,850 g mole/min 681 l/min 76 l/min			15.00 8.1	1,090	11,826 10,215 616
	Rate of Oxidation	of $SO_2 = SO_2$ (In) -	SO ₂ (Out) - (CaSO ₃ 1/2 H ₂	,0 Formation	Rate	
		= 24,423 - 23	2,657 - 1,209	•	-		
		= 557 m mole,	/min				
3. CaSO ₄ 2 H ₂ O Form	ation						
Entering Stream	Additive Scrubber Spray	2,800 g/min 757 l/min	6.94	0,5 0.58			1,400 3,047
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 1/min 76 1/min	9.25 25.3	0.49 0.33			3,087 635
	Rate of CaSO ₄ 2 I	i_2 0 Formation = CaSO	2 H ₂ O (Out)) - CaSO ₄ 2	H ₂ 0 (In)		
		= 3,721	- 4,447				
		= = 72!	m mole/min				
. CaCO ₃ Formation					-		
Entering Stream	Additive Scrubber Spray	2.800 g/min 757 l/min	6.94	0.455 0.181			1,274 951
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 I/min 76 I/min	9.25 25.3	0.180 0.249			1,134 479
	Rate of CaCO ₃ Form	nation = CaCO ₃ (Out)	- CaCO ₃ (In)				
		= 1,613 - 2,22	! 5				
		= - 612 m male	e/min				
. Ca(OH) Dissolution	<u>n</u>						
Entering Stream	Scrubber Spray	757 1/min			· 23 . 4		17,714
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 1/min 76 1/min			32.3 26.1		21,996 1,984
	Rate of Ca(OH) ₂ Di	ssolution = Ca(liq)	Out - Ca(11q)	In + (CaCO ₃	+ CaSO ₄ 2	H ₂ 0 + Cas	0 ₃ 1/2 H ₂ 0)
		= 23,980 -	17,714 + - 6	512 + - 725 +	1209		

TABLE H-7. RATE CALCULATION USING SOLID MATERIAL BALANCE (Continued)

		Stream Flow Rate	Slurry Solid Cont. (g/1)	Concen. In Solid) (m mole/g)	Concen. In Liquid (m mole/1)	Total Concen. Species In Gas Flow Rate (PPM) m mole/min
Hold Tank (Set #1)						
1. ÇaSO ₃ 1/2H ₂ O						
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 1/min 76 1/min 151 1/min 151 1/min	7.33 22.7 0.13	1.00 1.47 0.93		4,992 2,536 19
Leaving Stream	Hold Tank Eff.	1,059 l/min	7.38	1.44		11,254
	Rate of CaSO ₃ 1/2	Formation = $Caso_3$	1/2 H ₂ 0 (Out	:) - CaSO ₃ 1,	/2 H ₂ O (In)	
		= 11,254	- 7,547			
		= 3,707	m mole/min			
2. SO ₂ Oxidation						
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 l/min 76 l/min 151 l/min 151 l/min			15.6 1.01 5.05	10,624 77 763
Leaving Stream	Hold Tank Effluent	1,059 l/mtn			7.45	7,890
	Rate of SO ₂ Oxidat	ion = S0 ₂ (In) - S0	2 (Out) - Cas	ю _з 1/2 н ₂ 0 і	Form Rate	
		= 11,464 - 7,89	0 - 3,707			
		= - 133 m mole/	min			
3. CaSO4 2 H ₂ O For	mation		•			
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 1/min 76 1/min 151 1/min 151 1/min	7.33 22.7 0.13	0,61 .33		3,045 569 20
Leaving Stream	Hold Tank Effluent	1,059 1/min	7.38	0.61		4,767
	Rate of CaSO ₄ 2 H ₂	O Formation = CaSO	4 2 H ₂ 0 (Out	:) - CaSO ₄ 2	H ₂ 0 (In)	
		= 4,76	7 - 3,634			
		= 1,13	3 m mole/min			
4. CaCO ₃ Formation						
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 1/min 76 1/min 151 1/min 151 1/min	7.33 22.7 0.13	0,195 0.355		973 612 20
Leaving Stream	Hold Tank Effluent	1,059 1/min	7.38	0.201		1,571
	Rate of CaCO ₃ Forma	tion = CaCO ₃ (Out)	- CaCO ₃ (In)			
		= 1,571 - 1,60	05			
		= -34 m mole/i	nin			
Ca(OH) Dissolut	ion					
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 1/min 76 1/min 151 1/min 151 1/min			31.6 21.6 22.1 1.08	21,520 1,642 3,337 163
Leaving Stream	Hold Tank Effluent				23.6	24,992
	Rate of Ca(OH) ₂ Dis				_	CO ₃ + CaSO ₃ 1/2 H ₂ O)
		= 24,992	- 26,662 + 1,	133 - 34 + 3	707	

TABLE H-7. RATE CALCULATION USING SOLID MATERIAL BALANCE (Continued)

Marble Bed (Set #2)		Stream Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
1. CaSO ₃ 1/2 H ₂ O Fo	ormation						
Entering Stream	Scrubber Spray	757 l/min	6.99	1.28			6,773
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 l/min 76 l/min	8.00 22.6	0.95 1.16			5,176 1,992
	Rate of CaSO ₃ 1/2	H ₂ O Formation = CaS	0 ₃ 1/2 H ₂ 0	(Out) - CaSO ₃	1/2 H ₂ 0 (In)	
		= 7,1	68 - 6,773				
		= 395	m mole/min				
2. SO ₂ Oxidation							
Entering Stream	Inlet Flue Gas Scrubber Spray	9,400 g mole/min 757 l/min			10.2	1,939	18,227 7,721
Leaving Stream	Outlet Flue Gas Scrubber Liquid Scrubber Bottom	10,850 g mole/min 681 l/min 76 l/min			16.85 12.00	1,090	11,827 11,475 912
	Rate of SO ₂ Oxidat	ion = SO ₂ ltq (In) -	SO ₂ 11q (Ou	t) - CaSO ₃ 1	/2 H ₂ O Forma	tion	
		= 25,948 - 24,21	4 - 395				
		= 1,339 m mole/m	iîn				
3. CaSO ₄ 2 H ₂ O Form	ation						
Entering Stream	Additive Scrubbing	2,800 g/min 757 l/min	6.99	0.50 0.51			1,400 2,699
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 1/min 76 1/min	8.00 22.6	0.28 0.32°			1,525 550
	Rate of CaSO ₄ 2 H	₂ O Formation = CaSO ₄	2 H ₂ O (Out) - CaSO ₄ 2	H ₂ 0 (In)		
		•	- 4,099				
		= - 2,0	24 m mole/mi	n			
4. CaCO ₃ Formation							
Entering Stream	Additive Scrubber Spray	2,800 g/min 757 l/min	6.99	0.455 0.192			1,274 1,016
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 l/min 76 l/min	8.00 22.6	0,205 0,299			1,116 514
	Rate of CaCO ₃ Form	ation = CaCO ₃ (Out)	- CaCO ₃ (In)				
	•	= 1,630 - 2,29	0				
		= - 660 m mole	s/min				
5. Ca(OH) ₂ Dissoluti	on						
Entering Stream	Scrubber Spray	757		26,6			20,136
Leaving Stream	Scrubber Liquid Scrubber Bottom	681 76		33.4 30.0			22,745 2,280
	Rate of Ca(OH) ₂ Di	ssolution = Ca(11q)	Out - Ca(liq) In + (CaCO ₃	+ CaSO ₃ /1/	'2 H ₂ 0 +	CaSO ₄ 2 H ₂ O)
		= 25,025 -	20,136 + -	660 + 395 + -	2,024		
		= 2,600 m	moles/min				

[°]Scrubber bottom of tank was used

TABLE H-7. RATE CALCULATION USING SOLID MATERIAL BALANCE (Continued)

Hold Tank (Set #2)		Stream Flow Rate	Slurry Solid Cont. (g/l)	Concen. In Solid (m mole/g)	Concen. In Liquid (m mole/l)	Concen. In Gas (PPM)	Total Species Flow Rate m mole/min
1. CaSO ₃ 1/2 H ₂ O Fo	mmation						
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 1/min 76 1/min 151 1/min 151 1/min	7.07 20.5	0.94 1.25			4,526 1,948
Leaving System	Hold Tank Effluent	1,059 1/min	6.65	1.30			9,155
	Rate of CaSO ₃ 1/2	H ₂ O Formation = Ca	s0 ₃ (Out) - C	aso ₃ In			
		= 9,	155 - 6,474				
		= 2,	681 m mole/mi	n			
2. SO ₂ Oxidation							
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 l/min 76 l/min 151 l/min 151 l/min			18.3 2.85 7.4		12,462 217 1,117
Leaving Stream	Hold Tank Effluent	1,059 1/min			10.25		10,855
	Rate of SO ₂ Oxidati	on = SO ₂ (liq) In	- SO ₂ (1iq) 0	ut - CaSO ₃ 1	/2 H ₂ O Forma	tion	
		= 13,796 - 10,8	55 - 2,681				
		= 260 m mole/mi	n				
3. CaSO ₄ 2 H ₂ O Form	ation						
Entering Streams	Scrubber Liquid Scrubber Bottom	681 1/min 76 1/min	7.07 20.5	.49 .44			2,359 685
Leaving Stream	Hold Tank Effluent	1,059 1/min	6.65	.63			4,435
	Rate of CaSO ₄ .2 H ₂	O Formation = CaSO	4 .2 H ₂ 0 (Out	.) - CaSO ₄ .2	H ₂ 0 (In)		
		•	6 - 3,044				
		= 1,39	2 m mole/min				
4. CaCO ₃ Formation							
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 l/min 76 l/min 151 l/min 151 l/min	7.07 20.5	.173 .311			833 485
Leaving Stream	Hold Tank Effluent	1,059 1/min	6.65	0.217			1,528
	Rate of CaCO ₃ Forma	tion = $CaCO_3$ (Out)	- CaCO ₃ (In)				
		= 1,528 - 1,3	18				
		= 210 m moles	/min				
5. Ca(OH) ₂ Dissolution	on						
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid Makeup Water	681 1/min 76 1/min 151 1/min 151 1/min			33.9 24.7 22.9 1.08		23,086 1,877 3,458 163
Leaving Stream	Hold Tank Effluent	1,059 l/min			25.9		27,428
	Rate of Ca(OH) ₂ Dis	solution = Ca(liq)	Out - Ca(liq) In + CaCO ₃	+ CaSO ₄ .2 H	1 ₂ 0 + CaSO	₃ .1/2 н ₂ 0
		= 27,428	- 28,584 + 21	0 + 1,392 + 2	,681		

= 3,127 m mole/min

TABLE H-B. RATE CALCULATIONS USING LIQUID MATERIAL BALANCE Experiment 17R

·		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #1)					
1. CaSO ₃ . 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	10,950 g mole/min 431 l/min	1.0	1 ,456	15 , 943 431
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottoms	11,605 g mole/min 329 l/min 102 l/min	8.95 1.0	764	8,866 2,944 102
	Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ IN - SO	O ₂ OUT - Oxidation Rai	te	
		= 15,943 + 43	31 - 8,866 - 2,944	.439 (15,943 - 8	.866) - 102
		= 4,462 - 3,1	107		
		= 1,355 m mol	łe/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray	431 1/min	10.27		4,426
Leaving Streams	Scrubber Liquid* Scrubber Bottom	329 l/min 102 l/min	14.46 11.61		4.757 1.184
	Rate of CaSO ₄ 2	: H ₂ 0 = S0 ₄ (Liq.) II	N - 50 ₄ (Liq.) OUT + (xidation Rate	
		= 4,426 - 5,94	1 + 3,107		
		= 1,592 m mole,	/min		
3. Ca(OH) ₂ Dissolut	ion				
Entering Streams	Scrubber Spray	431 1/min	13.7		5,905
Leaving Streams	Scrubber Liquid* Scrubber Bottom	329 1/min 102 1/min	19.0 16.8		6,251 1,714
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	- Ca IN + Ca Prec. Ra	ite	
		= 7,965 -	- 5,905 + 2,947		
		= 5,007 m	n mole/min		

^{*}Ave. of Marble Bed Front and Back

TABLE H-8. RATE CALCULATIONS USING LIQUID MATERIAL BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	329 l/min 102 l/min 208 l/min	8.9 1.0 -		2,928 102
Leaving Streams	Hold Tank Eff.	631 1/min	1.5		947
	Rate of CaSO ₃ 1,	/2 H ₂ 0 = SO ₂ IN - S	0 ₂ OUT		
	•	= 3,030 - 94	7		
		= 2,083 m mo	le/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	329 1/min 102 1/min 208 1/min	14.85 10.3		4,886 1,051
Leaving Streams	Hold Tank Eff.	631 1/min	9.4		5,931
	Rate of CaSO ₄ 2f	4 ₂ 0 = SO ₄ (IN) - SO	4 (OUT)		
	·	= 5,937 - 5,931			
		= 6 m mole/min			
3. CaCO ₃					
Entering Streams	Scrubber Liquid ⁺ Scrubber Bottom ⁺⁺ Make Up H ₂ O	329 1/min 102 1/min 208 1/min	1.1		362 28
Leaving Streams	Hold Tank Eff.	631 1/min	.26		164
	Rate of CaCO ₃ = C	3 IN - CO3 OUT			
	= 39	90 - 164			
	= 22	26 m mole/min			
4. Ca(OH) ₂ Dissoluti	on				
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	329 l/min 102 l/min 208 l/min	19.9 17.4 1.08		6,547 1,775 225
Leaving Streams	Hold Tank Eff.	631 1/min	17.7		7,383
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	- Ca IN + Ca Prec. R	ate	
	_	= 7,383	- 8,547 + (226 + 6 +	2,083)	
		= 1,151	m mole/min		

⁺Values from Marble Bed Front and Back ++Values from Scrubber Bottom at Scrubber

TABLE H-8. RATE CALCULATIONS USING LIQUID MATERIAL BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	10,950 g mole/min 431 l/min	1.3	1,456	15 ,943 560
Leaving Streams	Outlet Flue Gas Scrubber Liquid Scrubber Bottom	11,605 g mole/min 329 l/min 102 l/min	7.2 1.7	764	8,866 2,369 173
	Rate of CaSO ₃	1/2 H ₂ 0 = SO ₂ IN - SO	O ₂ OUT - OXID. Rate		
		= 16,503 - 11	.408417 (7,077)		
		= 16,503 - 11	,408 - 2,951		
		= 2,144 m mol	le/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Spray	431]/min	9.38		4,043
Leaving Streams	Scrubber Liquid Scrubber Bottom	329 l/min 102 l/min	15.37 11.3		5,057 1,153
	Rate of CaSO ₄	$2H_20 = SO_4 IN - SO_4 O$	OUT + OXID. Rate		
		= 4,043 - 6,210	+ 2,951		
		= 784 m mole/min	1		
3. Ca(OH) ₂ Dissolution	on 				
Entering Streams	Scrubber Spray	431 1/min	12.8		5,517
Leaving Streams	Scrubber Liquid Scrubber Bottom	329 l/min 102 l/min	19.1 16.3		6,284 1,663
	Rate of Ca(OH) ₂ O	dissolution = Ca OUT	- Ca IN + Ca Prec. Ra	te	
		= 7,947 -	5,517 + (784 + 2,144)	
		= 5,358 m	mole/min		

TABLE H-8. RATE CALCULATIONS USING LIQUID MATERIAL BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m_mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	329 1/min 102 1/min 208 1/min	9.2 1.1		3,027 112
Leaving Streams	Hold Tank Eff.	631 1/min	1.5		946
	Rate of CaSO ₃	1/2 H ₂ 0 = SO ₂ IN - S	0 ₂ 0 UT		
		= 3,139 - 94	6		
		= 2,193 m mo	le/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	329 1/min 102 1/min 208 1/min	12.75 11.64		4,195 1,187
Leaving Streams	Hold Tank Eff.	631 1/min	9.4		5,931
	Rate of CaSO ₄ 2	$2 H_2 0 = SO_4 IN - SO_4$	OUT		
		= 5,382 - 5,93	1		
		≃ ~ 549 m mole,	/min		
3. <u>CaCO₃</u>					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	329 1/min 102 1/min 208 1/min	.96 .06		316 6
Leaving Streams	Hold Tank Eff.	631 1/min	. 26		164
	Rate of CaCO ₃ = C	0 ₂ IN - CO ₂ OUT			
	= 3	22 - 164			
	= 1	58 m mole/min			
4. Ca(OH) ₂ Dissoluti	on				
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H _Z O	329 1/min 102 1/min 208 1/min	19.6 17.6 1.08		6,448 1,795 225
Leaving Streams	Hold Tank Eff.	631 1/min	11.7		7,383
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	Ca IN + Ca Prec. Rate	es .	
		= 7,383 -	- 8,468 + (158 - 549 +	- 2,193)	
		= 717 m n	nole/min		

TABLE H-9. RATE CALCULATIONS USING LIQUID BALANCE

Experiment 18R

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #1)	2.				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	10,820 g mole/min 796 l/min	.35	1,471	15,916 279
Leaving Streams	Scrubber Liquid* Scrubber Bottom Outlet Flue Gas	682 l/min 114 l/min 11,630 g mole/min	2.95 .45	447	2,012 51 5,199
	Rate of CaSO ₃ 1,		O ₂ OUT - Oxidation Ra	te	
	3		262279 (15,916 -		
		= 8,933 - 2,9	390 -		
		= 5,943 m mol	le/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Spray	796 1/min	18.15		14,447
Leaving Streams	Scrubber Liquid* Scrubber Bottom	682 l/min 114 l/min	24.34 19.09		16,600 2,176
	Rate of CaSO ₄ 2	1 ₂ 0 = SO ₄ IN - SO ₄ (OUT + Oxidation Rate		
		= 14,447 - 18,77	76 + 2,990		
		= -1,339 m mole/	/min		
3. Ca(OH) ₂ Dissolution	on 				
Entering Streams	Scrubber Spray	796 1/m1n	22,52		17,926
Leaving Streams	Scrubber Liquid* Scrubber Bottom	682 1/min 114 1/min	24.45 24.42		16,675 2,784
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	- Ca IN + Ca Prec. Ra	ites	
		= 19,459	- 17,926 + (-1,339 +	5,943)	
		= 6,137 m	n mole/min		

^{*}Ave. of Marble Bed Front and Back

TABLE H-9. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	682 1/min 114 1/min 76 1/min	2.57 .89 .56		1,753 101 43
Leaving Streams	Hold Tank Eff.	872 1/min	.53		462
	Rate of CaSO ₃ 1	/2 H ₂ 0 Form = SO ₂ I	N - SO ₂ OUT		
		= 1,897	~ 462		
		= 1,435	m mole/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	682 I/min 114 1/min 76 1/min	19.98 17.20 16.49		13,626 1,961 1,253
Leaving Streams	Hold Tank Eff.	872 1/min	17.35		15,129
	Rate of CaSO ₄ 2	H ₂ 0 = SO ₄ IN - SO ₄ (DUT		
		= 16,840 - 15,1	29		
		= 1,711 m mole/u	nin		
3. CaCO ₃ (No CO ₂ Dat	a are Available)				
4. Ca(OH) ₂ Dissoluti	on				
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	682 1/min 114 1/min 76 1/min	21.25 32.72 20.6		14,493 3,730 1,566
Leaving Streams	Hold Tank Eff.	872 1/min	22.97		20,030
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	- Ca IN + Ca Prec. Ra	ites	
	-	= 20,030	- 19,789 + (3,146)		

= 3,387 m mole/min

TABLE H-9. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set 2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	10,820 g mole/min 796 l/min	.38	1,471	15,916 302
Leaving Streams	Outlet Flue Gas Scrubber Liquid Scrubber Bottom	11,630 g mole/min 682 l/min 114 l/min	4.175 .42	447	5,199 2,847 48
	Rate of CaSO ₃ 1,	/2 H ₂ 0 = SO ₂ IN - SO	0 ₂ OUT - Oxidation Rai	te	
		= 16,218 - 8	,094289 (10,717)		
		= 16,218 - 8	,094 - 3,097		
		= 5,027 m mol	le/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Spray	796 l/min	19.67		15,657
Leaving Streams	Scrubber Liquid* Scrubber Bottom	682 1/min 114 1/min	25.15 19.02		17,152 2,168
	Rate of CaSO ₄ 21	1 ₂ 0 = SO ₄ IN - SO ₄ (OUT + Oxidation Rate		
		= 15,657 - 19,32	20 + 3,097		
		= -566 m mole/mi	in		
3. Ca(OH) ₂ Dissolution	on 				
Entering Streams	Scrubber Spray	796 1/min	23,25		18,507
Leaving Streams	Scrubber Liquid* Scrubber Bottom	682 1/min 114 1/min	26.495 24.95		18,070 2,844
	Rate of Ca(OH) ₂ Di	ssolution = Ca QUT	- Ca IN + Ca Prec. Ra	te	
		= 20,914	- 18,507 + (4461)		
		= 6,868 m	ı mole/min		

^{*}Ave. of Marble Bed Front and Back

TABLE H-9. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

_			Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hol	ld Tank					
1.	CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	682 1/min 114 1/min 76 1/min	3.25 .72 .40		2,217 82 30
	Leaving Streams	Hold Tank Eff.	872 1/min	.55		480
		Rate of CaSO ₃ 1/	/2 H ₂ 0 Form = SO ₂ I	N - SO ₂ OUT		
			= 2,329	- 480		
			= 1,849	m mole/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	682 1/min 114 1/min 76 1/min	20.46 18.51 16.96		13,954 2,110 1,289
	Leaving Streams	Hold Tank Eff.	872 1/min	18.68		16,289
		Rate of CaSO ₄ 2H	1 ₂ 0 Form = SO ₄ IN -	SO4 OUT		
			= 17,353 -	16,289		
			= 1,064 m r	mole/min		
3.	CaCO ₃ (No CO ₂ Data	Was Available)				
4.	Ca(OH) ₂ Dissolutio	on				
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	682 1/min 114 1/min 76 1/min	22.17 34.30 22.35		15,120 3,910 1,699
I	Leaving Streams	Hold Tank Eff.	872 1/min	24,20		21,102
		Rate of Ca(OH) ₂ Dt	ssolution = Ca OUT	- Ca IN + Ca Prec. Ra	ites	
			= 21,102	- 20,729 + 2,913		

= 3,286 m mole/min

TABLE H-10. RATE CALCULATIONS USING LIQUID BALANCE

Experiment 19R

	····	Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #1)	1				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	9,200 g mole/min 738 l/min	15.55	1,884	17,333 11,476
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,555 g mole/min 662 l/min 76 l/min	26.85 14.75	1,060	11,188 17,775 1,121
	Rate of CaSO ₃ · 1/	/2 H ₂ 0 = SO ₂ IN - SO ₂	₂ OUT - OXID. Rate		
		= 28,809 - 30,	,084533 (17,333 -	11,188)	
		= - 4,550 m mc	ole/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray	738 1/min	20.35		15,018
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 1/min 76 1/min	26.45 24.45		17,510 1,858
	Rate of CaSO ₄ = SC	04 IN - SO4 OUT + OXI	ID. Rate		
	= 15	5,018 - 19,368 + 3,27	75		
	= -	1,075 m mole/min			
3. Ca(OH) ₂ Dissolut	ion 				
Entering Streams	Scrubber Spray	738 1/min	25.7		18,967
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 l/min 76 l/min	34.4 27.7		22,773 2,105
	Rate of Ca(OH) ₂ Di	ssolution = CA OUT -	- Ca IN + Ca Prec. Rat	es	
		= 24,878 -	18,967 + - 4,550 - 1	,075	
		= 286 m mo	le/min		

^{*}Ave. of Marble Bed Front and Back

TABLE H-10. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

	· 	Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 76 1/min 133 1/min	28.6 7.9		18,933 600
Leaving Streams	Hold Tank Eff.	871 1/min	15.7		13,675
	Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ IN - SO ₂	₂ OUT		
		= 19,533 - 13	,675		
		= 5,858 m mole	e/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 76 1/min 133 1/min	25.1 22.5		16,616 1,710
Leaving Streams	Hold Tank Eff.	871 1/min	20.6		17,943
	Rate CaSO ₄ 2 H ₂	0 = SO4 IN - SO4 OUT			
		= 18,326 - 17,943			
		= 383 m mole/min			
3. CaCO ₃					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ 0	662 1/min 76 1/min 133 1/min	1,845 ,75		1,221 57
Leaving Streams	Hold Tank Eff.	871 1/min	1,10		958
	Rate of CaCO ₃ = Co	0 ₂ IN - CO ₂ OUT			
	= 1:	278 - 958			
	= 32	20 m mole/min			
4. Ca(OH) ₂ Dissoluti	ion				
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 76 1/min 133 1/min	35.3 23.8 1.08		23,369 1,809 144
Leaving Streams	Hold Tank Eff.	871 1/min	24.5		21,340
	Rate of Ca(OH) ₂ D	issolution = Ca OUT -	- Ca IN + Ca Prec, Rat	te	
		= 21,340 -	- 25,322 + (320 + 383	+ 5,858)	

= 2,579 m mole/min

TABLE H-10. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set 2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	9,120 g mole/min 738 l/min	15.15	1,881	17,155 11,181
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,487 g mole/min 662 l/min 76 l/min	25.9 13.65	1,060	11,116 17,146 1,037
	Rate of CaSO ₃ = SO	₂ IN - SO ₂ OUT - OX	ID. Rate		
	= 28	,336 - 29,29953	6 (17,155 - 11,116)		
	= 28	,336 - 29,29953	6 (6,039)		
	= 28	,336 - 29,299 - 3,2	37		
	= -4	,200 m mole/min			
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray	738 1/min	21,45		15,830
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 1/min 76 1/min	26.5 23.65		17,543 1,797
	Rate of CaSO ₄ 2	H ₂ 0 = SO ₄ IN - SO ₄	OUT + OXID. Rate		
		= 15,830 - 19,3	40 + 3,237		
		= -273 m mole/m	t n		
3. Ca(OH) ₂ Dissoluti	lon				
Entering Streams	Scrubber Spray	738 1/min	25,2		18,598
Leaving Streams	Scrubber Liquid* Scrubber Bottom	662 l/min 76 l/min	34.65 27.6		22,938 2,098
	Rate of Ca(OH) ₂ Di	ssolution = Ca OUT	- Ca IN + Ca Prec. Ra	te	
		= 25,036	- 18,598 + (- 273 - 4	,200)	
		= 25,036	- 18,598 + -4,473		
		= 1,965 m	mole/min		

TABLE H-10. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 76 ⁻ 1/min 133 1/min	27.31 4.5		18 , 079 342
Leaving Streams	Hold Tank Eff.	871 1/min	15.05		13,109
	Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ (IN) -	SO ₂ (OUT)		
		= 18,421 - 13	,109		
		= 5,312 m mol	e/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 76 1/min 133 1/min	25.2 21.9		16,682 1,664
Leaving Streams	Hold Tank Eff.	871 1/min	20,85		18,160
	Rate of CaSO ₄ 2	$H_20 = SO_4 (IN) - SO_4$	4 (OUT)		
		= 18,346 - 18,1	60		
		= 186 m mole/mi	n		
3. <u>CaCO₃</u>					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 133 1/min	1.36 0.17		900 13
Leaving Streams	Hold Tank Eff.	871 1/min	0.97		845
	Rate of CaCO ₃ = C	O ₂ (IN) - CO ₂ (OUT)			
	= 9	13 - 845			
	= 68	3 m mole/min			
4. Ca(OH) ₂ Dissolut	ion				
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	662 1/min 76 1/min 133 1/min	35.3 22.2 1.08		23,369 1,687 144
Leaving Streams	Hold Tank Eff.	871 1/min	25.5		22,211
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	- Ca IN + Ca Prec. Ra	tes	
		= 22,211	- 25,200 + (5,566)		
		= 2,577 m	mole/min		

TABLE H-11. RATÉ CALCULATIONS USING LIQUID BALANCE
Experiment 20R

_			Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Mai	rble Bed (Set #1)				
1.	CaSO ₃ 1/2 H ₂ O	·				
	Entering Stream	s Inlet Flue Gas Scrubber Spray	9,400 g mole/mi 757 l/min	n 7 . 9	1,962	18,443 5,980
	Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,850 g mole/mi 681 l/min 76 l/min	n 15.0 8.1	1,090	11,827 10,215 616
		Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ (IN)	- SO ₂ (OUT) - Oxidation	Rate	
			= 24,423 - 3	22,658552 (18,443 -	11,827)	
			= 24,423 = 3	22,658552 (6,616)		
			= 24,423 - 3	22,658 - 3,652		
			= -1,887 m r	mole/min		
2.	CaSO ₄ 2H ₂ O					
	Entering Streams	s Scrubber Spray	757 1/min	24.7		18,698
	Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 76 1/min	30.7 25.3		20,907 1,923
		Rate of CaSO ₄ = SO	0 ₄ (IN) - SO ₄ (OUT)) + Oxidation Rate		
		= 18	8,698 - 22,830 + 3,	,652		
		= -	480 m mole/min			
3.	Ca(OH) ₂ Dissolut	cion				
Į	Entering Streams	Scrubber Spray	757 1/m1n	23.4		17,714
ı	Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 76 1/min	32.25 26.1		21,962 1,984
		Rate of Ca(OH) ₂ Di	issolution = Ca (OU	IT) - Ca (IN) + Ca Prec.	Rate	
			= 23,946	5 - 17,714 + (-542 - 1,7	(69)	
			= 23,946	i - 17,714 + (-2,367)		
			= 3,865	m mole/min		

TABLE H-11. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2	H ₂ 0				
Entering Lic	quid Scrubber Liquid Scrubber Bottom Make Up H ₂ O Clarified Liquid	681 1/min 76 1/min 151 1/min 151 1/min	15.6 1.01 5.05		10,624 77 - 763
Leaving Stre	eams Hold Tank Eff.	1,059 1/min	7.45		7,890
	Rate of CaSO ₃ 1	/2 H ₂ 0 = S0 ₂ IN - S0	0 ₂ (OUT)		
		= 10,701 - 7	,890		
		= 2,811 m moi	le/min		
2. CaSO ₄ 2 H ₂	20				
Entering Lic	Scrubber Bottom Make-Up H ₂ 0	681 1/min 76 1/min 151 1/min 151 1/min	30.1 18.9 - 23.35		20,498 1,436 3,526
Leaving Stre	Clarified Liq.	1,059 1/min	24.45		25,893
LCGVING SCIE		H ₂ O = SO ₄ IN - SO ₄ (
		= 25,460 - 25,89			
		= -433 m mole/mi			
3. CaCO ₃					
Entering Lic	quid Scrubber Liquid Scrubber Bottom Make-Up H ₂ O Clarified Liquid	681 1/min 76 1/min 151 1/min 151 1/min	1.59∆ .06 _ .51		1,083 5 77
Leaving Stre	eams Hold Tank Eff.	1,059 1/min	.95		1,006
	Rate of $CaCO_3 = C$	0 ₂ IN - CO ₂ OUT			
	= 1	,165 - 1,006			
	= 1.	59 m mole/min			
4. Ca(OH) ₂ Diss	solution				
Entering Str	reams Scrubber Liquid Scrubber Bottom Make-Up H ₂ O Clarified Liquid	681 1/min 76 1/min 151 1/min 151 1/min	31.6 21.6 1.08 22.1		21,520 1,642 163 3,337
Leaving Stre	eams Hold Tank Eff.	1,059 1/min	23.6		24,992
	Rate of Ca(OH) ₂ D	issolution = Ca OUT	- Ca IN + Ca Prec. Ra	te	
		= 24,992	- 26,662 + (2,811 - 4	33 + 159)	
		= 24,992	- 26,662 + 2,537		
		= 867 m r	mole/min		

 $\Delta From\ marble\ bed\ front\ and\ back$

TABLE H-11. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set 2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	9,400 g mole/min 751 l/min	10.2	1,939	18,227 7,721
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,850 g mole/min 681 l/min 76 l/min	16.85 12.0	1,090	11,827 11,475 912
	Rate CaSO ₃ 1/2 H	l ₂ 0 Form = SO ₂ IN -	SO ₂ OUT - OXID. Rate		
		= 25,948 - 3	24,214543 (18,227	- 11,827)	
		= 1,734 - 3	, 475		
		= -1,711 m ı	mole/min		
2. CaSO ₄ 2H ₂ O					
Entering Stream	Scrubber Spray	757 1/min	25.5		19,304
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 l/min 76 l/min	29.7 26.6		20,226 2,022
	Rate of CaSO ₄ 2H	$_{2}0 = S0_{4} IN - S0_{4} OI$	UT + OXID. Rate		
		= 19,304 - 22,248	3 + 3,488		
		≈ 544 m mole/min			
3. Ca(OH) ₂ Dissoluti	on				
Entering Streams	Scrubber Spray	757 1/min	26.6		20,136
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 76 1/min	33.45 30.0		22,779 2,280
	Rate of Ca(OH) ₂ Di	ssolution = Ca OUT -	- Ca IN + Ca Pre. Rate	!	
		= 25,059 -	- 20,136 + (544 - 1,74	11)	
		= 25,059 -	- 20,136 + (-1,197)		
		= 3,726 m	mole/min		

TABLE H-11. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O	681 l/min 76 l/min 151 l/min	18.3 2.85		12,462 217
	Clarified ² Liquid	151 1/min	7.4		1,117
Leaving Streams	Hold Tank Eff.	1,059 1/min	10.25		10,855
	Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ IN - SO	2 OUT		
		= 13,796 - 10	,855		
		= 2,941 m mol	e/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams		681 1/min	29.1		19,817
	Scrubber Bottom Make Up H ₂ O	76 l/min 151 l/min	23.3		1,771
	Clarified_Liquid	151 1/min	20.9		3,156
Leaving Streams	Hold Tank Eff.	1,059 1/min	24.65		26,104
	Rate of CaSO ₄ 2	$H_20 = S0_4 \text{ IN} - S0_4 \text{ O}$	υT		
		= 24,744 - 26,10	4		
		= - 1,360 m mole	/m1n		
3. <u>CaCO</u> 3					
Entering Streams	Scrubber Liquid Scrubber Bottom Make Up H ₂ O Clarified ² Liquid	681 1/min 76 1/min 155 1/min	1.42		967 36
lasudas Stucias	•	155 1/min	.57		88
Leaving Streams	Hold Tank Eff.	1,059 1/min	.87		921
	Rate of CaCO ₃ = CC				
		091 - 921			
4. Ca(OH) ₂ Dissoluti		0 m mole/min			
		CO1 1 (-1-			
Entering Liquid	Scrubber Liquid Scrubber Bottom Make Up H ₂ O Clarified ² Liquid	681 l/min 76 l/min 155 l/min 155 l/min	33.9 24.7 1.08 22.9		23,086 1,877 167 3,550
Leaving Streams	Hold Tank Eff.	1,059 1/min	25.9		27,428
	Rate of Ca(OH) ₂ Di	ssolution = Ca OUT -	- Ca IN + Ca Prec. Rat	es	
		= 27,428 -	- 28,680 + 1,751		
		= 499 m mc	ole/min		

^{*}Ave. of Marble Bed Front and Back

TABLE H-12. RATE CALCULATIONS USING LIQUID BALANCE

Experiment 21R

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #1)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	10,200 g mole/min 758 l/min	.9	2,000	20,400 682
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	8,820 g mole/min 681 l/min 76 l/min	10.2 1.6	735	6,483 6,946 122
	Rate of CaSO ₃ 1	/2 H ₂ 0 Form = SO ₂ IN	- SO ₂ OUT - Oxid. Ra	te	
		= 21,082	- 13,551357 (20,	400 - 6,483)	
		= 21,082	- 13,551 - 4,968		
		= 2,563	m mole/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Spray	758 1/min	24.82		18,814
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 l/min 76 l/min	30.27 26.55		20,614 2,018
	Rate of CaSO ₄ 2	H ₂ O Form = SO ₄ IN -	SO ₄ OUT - Oxid. Rate		
		= 18,814 -	22,632 + 4,968		
		= 1,150 m m	ole/min		
3. Ca(OH) ₂ Dissolut	ion				
Entering Streams	Scrubber Spray	758 1/min	21,79		16,517
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 l/min 76 l/min	23.98 21.75		16,330 1,653
	Rate of Ca(OH) ₂ D	issolution = Ca (OUT) - Ca (IN) + Ca Prec	. Rate	
		= 17,983	- 16,517 + (2,563 + 1	,150)	
		= + 5,179	m mole/min		

^{*}Average of marble bed front and back

TABLE H-12. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Steams	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	.7 1.0 1.10		477 76 42
Leaving Streams	Hold Tank Eff.	795 1/min	1.10		875
	Rate of CaSO ₃ 1,	/2 H ₂ 0 = S0 ₂ IN - S	0 ₂ OUT		
		= 595 - 875			
		= -280 m mol	e/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 l/min 76 l/min 38 l/min	25.5 21.3 22.4		17,366 1.619 851
Leaving Streams	Hold Tank Eff.	795 1/min	26.7		21,227
	Rate of CaSO ₄ 2	H ₂ 0 Form = SO ₄ IN -	SO4 OUT		
		= 19,836 -	21,227		
		= - 1,391 :	m mole/min		
3. CaCO ₃					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	.18 .23 .3		123 17 11
Leaving Streams	Hold Tank Eff.	795 1/min	.28		223
	Rate of CaCO ₃ Form	$m = CO_2 IN - CO_2 OU$	т		
		= 151 - 223			
		= 72 m mole/min			
4. Ca(OH ₂) Dissolut	ion				
Set #1 Marble Be	d				
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	30.2 28.75 20.8		20,566 2,185 790
Leaving Stream	Hold Tank Eff.	795 l/min	18.1		14,390
	Rate of Ca(OH ₂) D	issolution = Ca OUT	+ Ca IN + Ca Prec. Ra	te	
		= 14,390	- 23,541 + (-72 -1391	-280)	
		= - 10,8	94 m mole/min		

TABLE H-12. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Set #2 Marble Bed					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet flue Gas Scrubber Spray	10,550 g mole/min 758 l/min	1.00	1,985	20,941 758
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	9,200 g mole/min 681 l/min 76 l/min	10.39 1.25	678	6,238 7,076 95
	Rate of CaSO ₃ 1,	/2 H ₂ O Form = SO ₂ IN	- SO ₂ (OUT) - Oxid. 1	Rate	
		= 21,699	- 13,409 - 0.266 (14	,703)	
		= 4,380 n	n mole/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Spray	758 1/min	25.82		19,572
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 76 1/min	30.95 25.6		21,077 1,946
	Rate of CaSO ₄ 2H	1 ₂ 0 = SO ₄ IN - SO ₄ OU	IT + Oxd. Rate		
		= 19,572 - 23,023	+ 3,910		
		= 459 m mole/min			
3. Ca(OH) ₂ Dissoluti	on				
Entering Streams	Scrubber Spray	758 1/min	18.59		14,091
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 76 1/min	25.09 20.9		17,086 1,588
	Rate of Ca (OH) ₂	dissolution = Ca OUT	- Ca IN + Ca Prec. Ra	ite	
		= 18,674	- 14,091 + (459 + 4,3	80)	
		= 9,422 m	mole/min		

^{*}Average of marble bed front and back

TABLE H-12. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Hold Tank					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	2.6 1.35 1.50		1,771 103 57
Leaving Streams	Hold Tank Eff.	795 1/min	1.73		1,375
	Rate of CaSO ₃ 1,	/2 H ₂ 0 Form = SO ₂ IN	- so ₂ out		
	-	= 1,931	- 1,375		
		= 556 m i	mole/min		
2. CaSO ₄ 2H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	24.7 16.33 22.35		16,821 1,241 849
Leaving Streams	Hold Tank Eff.	795 1/min	28.8		22,896
	Rate of CaSO ₄ 2	120 = SO4 IN - SO4 0	UT		
		= 18,911 - 22,899	5		
		= -3,985 m mole/r	nin		
3. CaCO ₃					
Entering Stream	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	1.42 .19 .29		967 14 11
Leaving Streams	Hold Tank Eff.	795 1/min	.50		398
	Rate of CaCO ₃ = CC	o ₂ IN - CO ₂ OUT			
	= 99	2 - 398			
	= 59	4 m mole/min			
4. Ca(OH) ₂ Dissoluti	on				
Entering Stream	Scrubber Liquid Scrubber Bottom Clarified Liquid	681 1/min 76 1/min 38 1/min	21.17 25.80 20.7		14,417 1,961 787
Leaving Streams	Hold Tank Eff.	795 l/min	16.9		13,436
	Rate of Ca(CH ₂) Di	ssolution = Ca OUT	- Ca IN + Ca Prec. Ra	te	
		= 13,436	- 17,165 - 2,835		

= -6,564 m mole/min

TABLE H-13. RATE CALCULATIONS USING LIQUID BALANCE

Experiment 22R

_			Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Ma	rble Bed					
١.	CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Inlet Flue Gas Scrubber Spray	10,466 g mole/min 1,355 l/min	3,75	2,021	21,152 5,081
	Leaving Streams	Outlet Flue Gas Scrubber Bottom Scrubber Liquid*	9,075 g mole/min 380 l/min 977 l/min	7.19 8.85	484	4,392 2,732 8,646
		Rate of CaSO ₃ 1,	/2 H ₂ 0 = SO ₂ IN - SO	2 OUT - OXID. Rate		
			= 26,233 - 15	,770346 (21,152 -	4,392)	
			= 26,233 - 15	,770346 (16,760)		
			= 26,233 - 15	,770 - 5,799		
			= 4,664 m mol	e/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Scrubber Spray	1,355 1/min	31.10		42,140
	Leaving Streams	Scrubber Liquid* Scrubber Bottom	977 1/min 380 1/min	33.58 27.6		32,807 10,488
		Rate of CaSO ₄ 2	$H_20 = S0_4 IN - S0_4$	OUT + OXID. Rate		
			= 42,140 - 43,2	95 + 5,799		
			= 4,644 m mole/	min		
3.	Ca(OH) ₂ Dissoluti	on				
	Entering Streams	Scrubber Spray	1,355 1/min	17.98		24,362
	Leaving Streams	Scrubber Liquid* Scrubber Bottom	977 1/min 380 1/min	21.925 22. 0 5		21,420 8,379
		Rate of Ca(OH) ₂ Di	issolution = Ca OUT	- Ca IN + Ca Prec. Ra	tes	
			= 29,799	- 24,362 + (4,664 + 4	,644)	
			= 14,745	m mole/min		

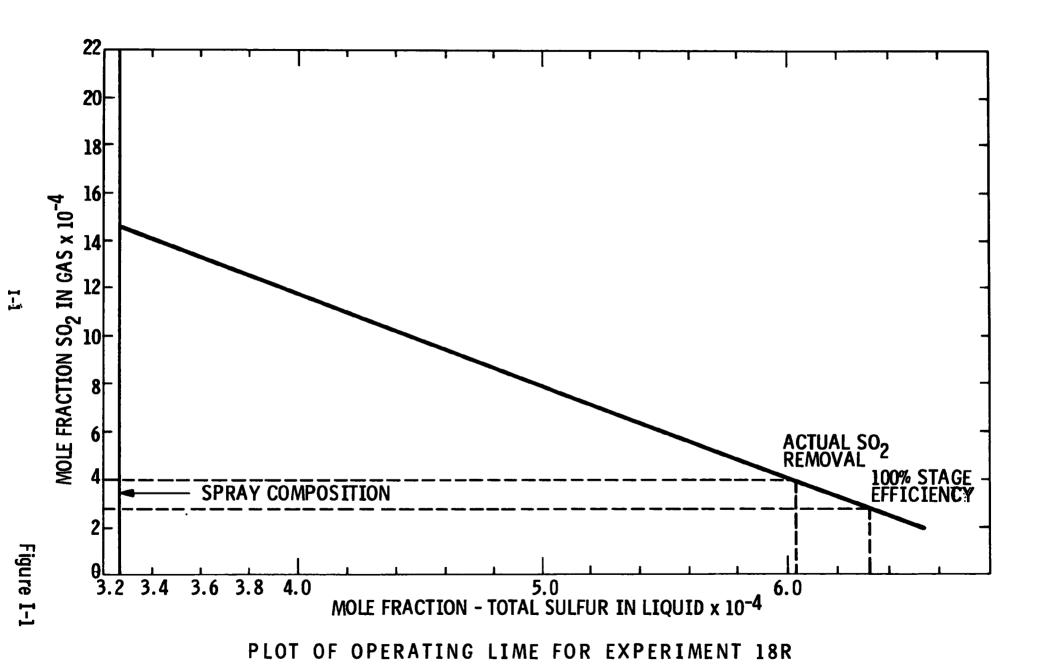
^{*}Ave. of Marble Bed Front and Back

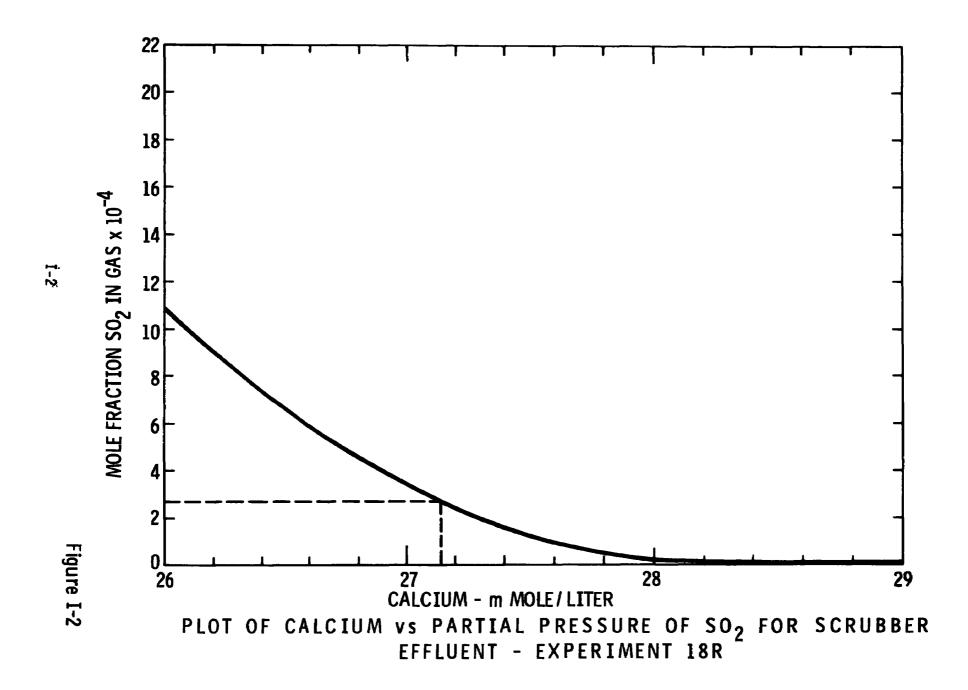
TABLE H-13. RATE CALCULATIONS USING LIQUID BALANCE (Continued)

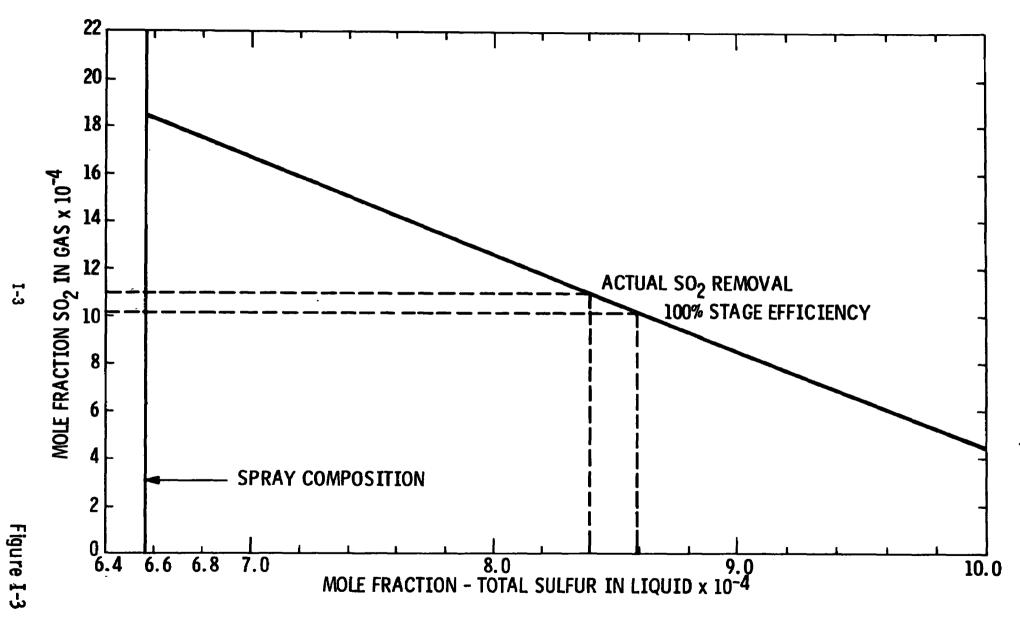
_			Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Нс	old Tank					
1.	. CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Clarified Liquid Scrubber Liquid Scrubber Bottom	38 1/min 977 1/min 380 1/min	.75 18.55 4.75		29 18,123 1,805
	Leaving Streams	Hold Tank Eff.	1,395 1/min	1.93		2,692
		Rate of CaSO ₃ 1,	/2 H ₂ 0 = SO ₂ IN - S	so ₂ out		
			= 19,957 - 2	2,692		
			= 17,265 m r	mole/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	977 1/min 380 1/min 38 1/min	39.5 30.43 21.5		38,591 11,563 817
	Leaving Streams	Hold Tank Eff.	1,395 1/min	29.77		41,529
		Rate of CaSO ₄ 2	$H_20 = S0_4 \text{ IN} - S0_4 = 50,971 - 41,971}$			
			= 9,442 m mole	e/min		
3.	CaCO ₃					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	977 1/min 380 1/min 38 1/min	.79 1.38 .23		772 524 9
	Leaving Streams	Hold Tank Eff.	1,395 l/min	.65		907
		Rate of CaCO ₃ = CC	0 ₂ IN - CO ₂ OUT			
		= 1,	305 - 907			
		= 39	98 m mole/min			
4.	Ca(OH) ₂ Dissoluti	on				
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarified Liquid	977 l/min 380 l/min 38 l/min	27.2 19.58 21.1		26,574 7,440 802
	Leaving Streams	Hold Tank Eff.	1,395 1/min	16.4		22,878
		Rate of Ca(OH) ₂ Di	ssolution = Ca OUT	Γ - Ca IN + Ca Prec. Rat	te	
			= 22,878	3 - 34,816 + (17,265 + 9	9,442 + 398)	
			= 15,167	m mole/min		

APPENDIX I

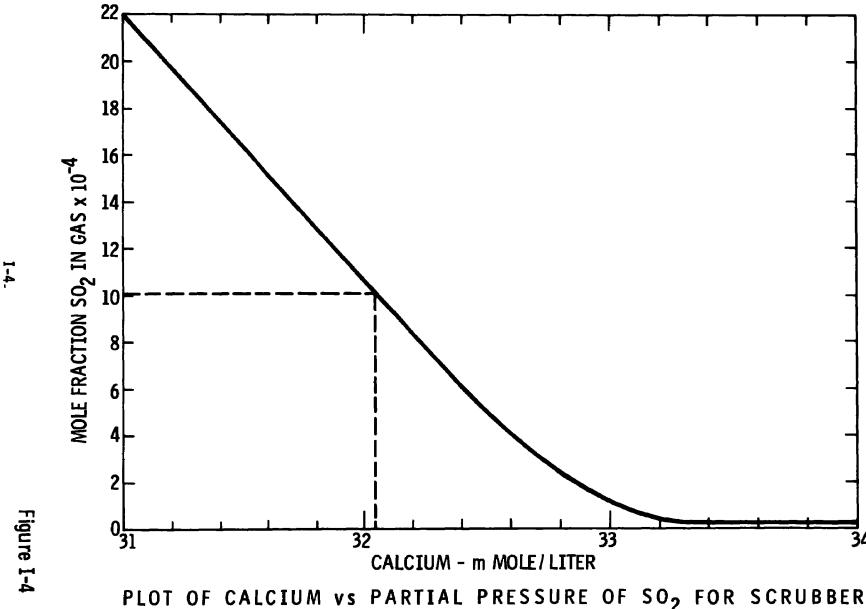
LIMESTONE FURNACE INJECTION SYSTEM
ADDITIVE DISSOLUTION RATE DETERMINATION DIAGRAMS



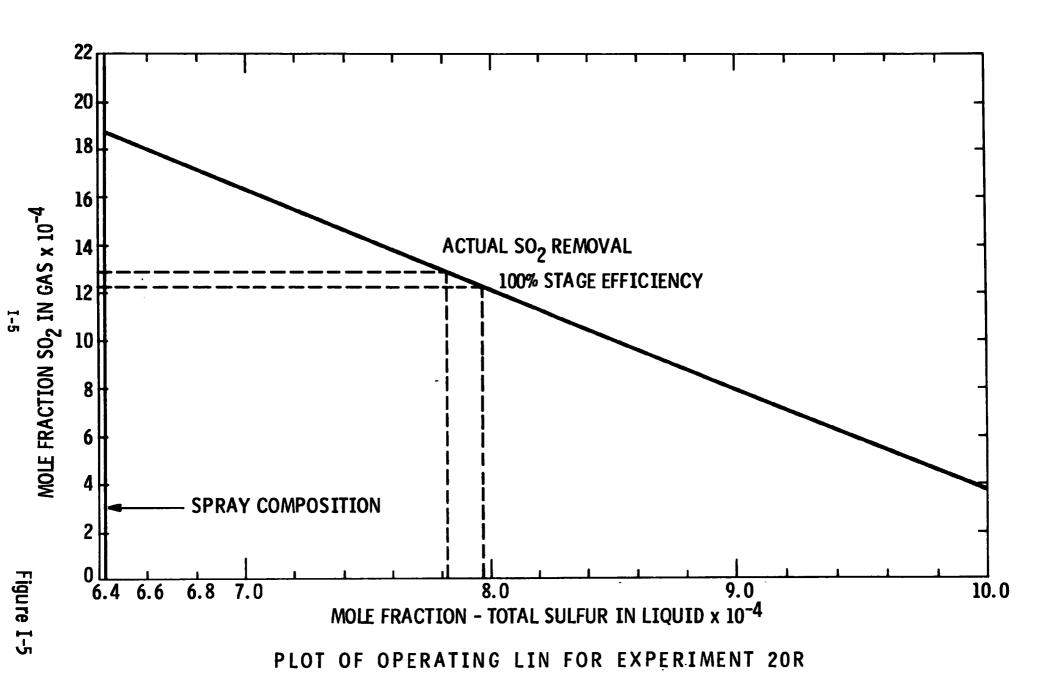


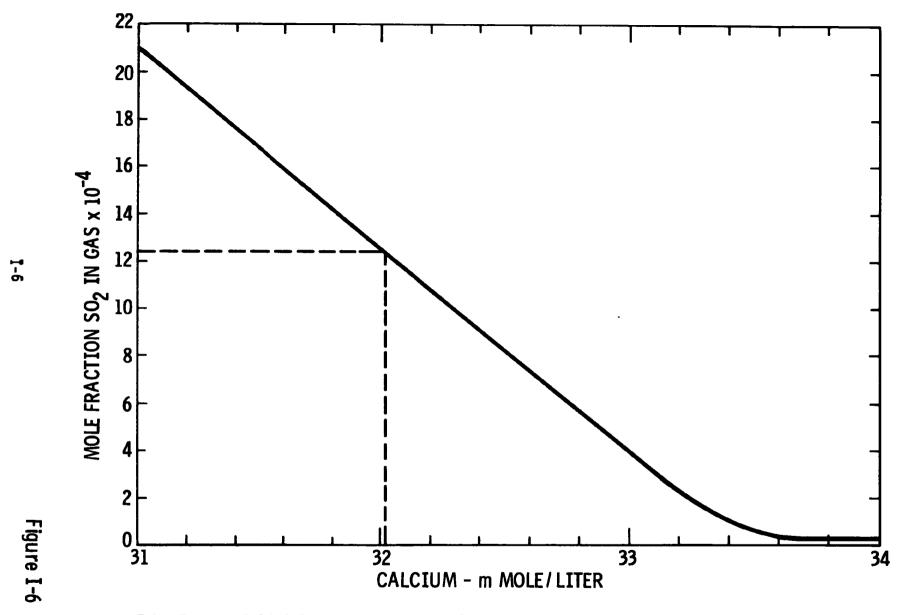


PLOT OF OPERATING LINE FOR EXPERIMENT 19R

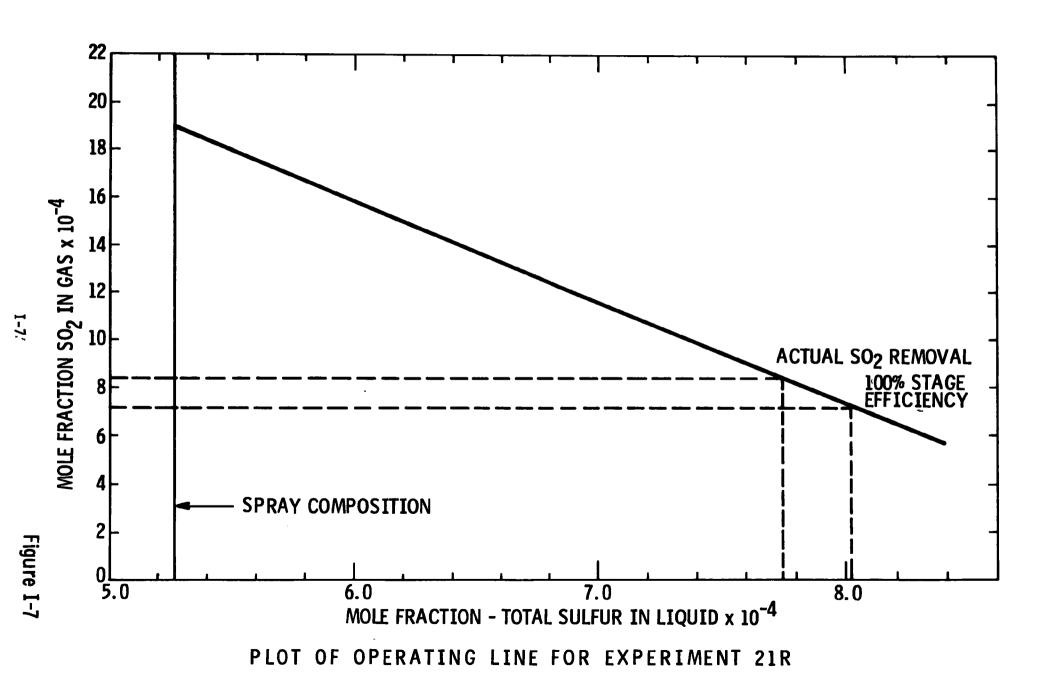


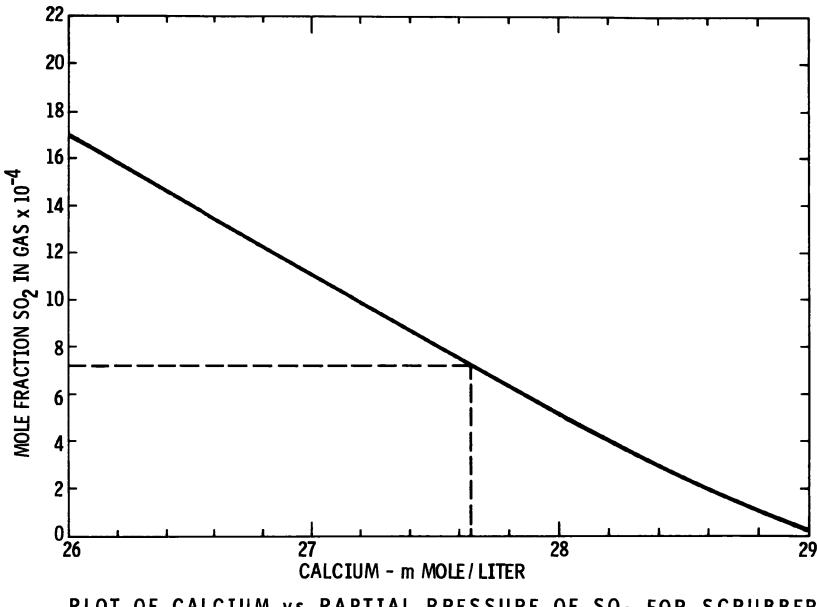
PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 19R



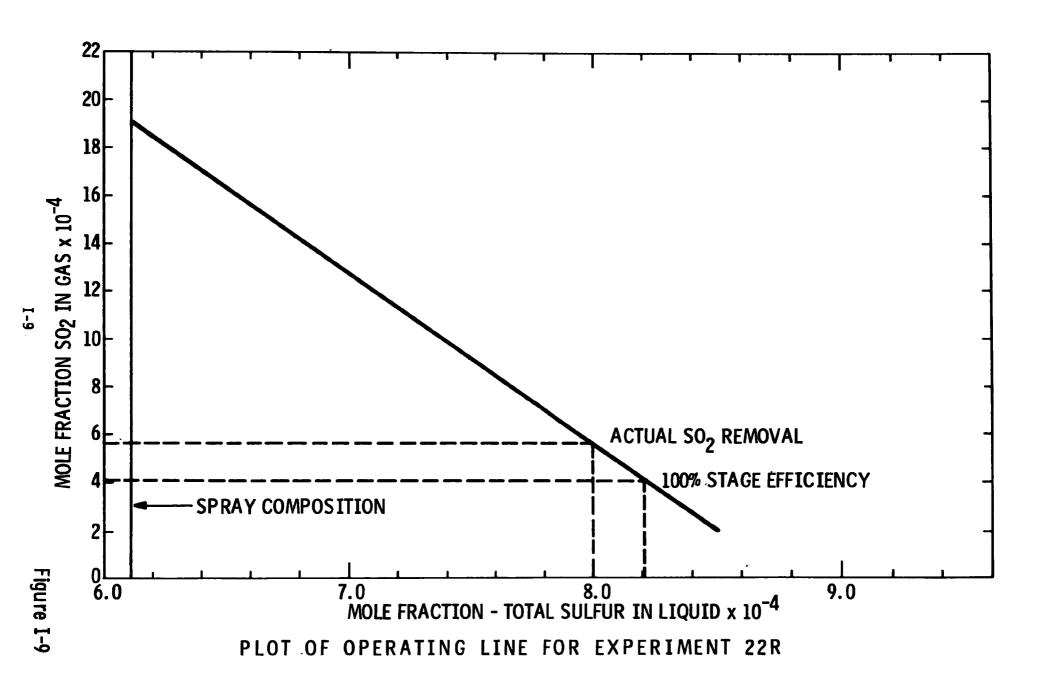


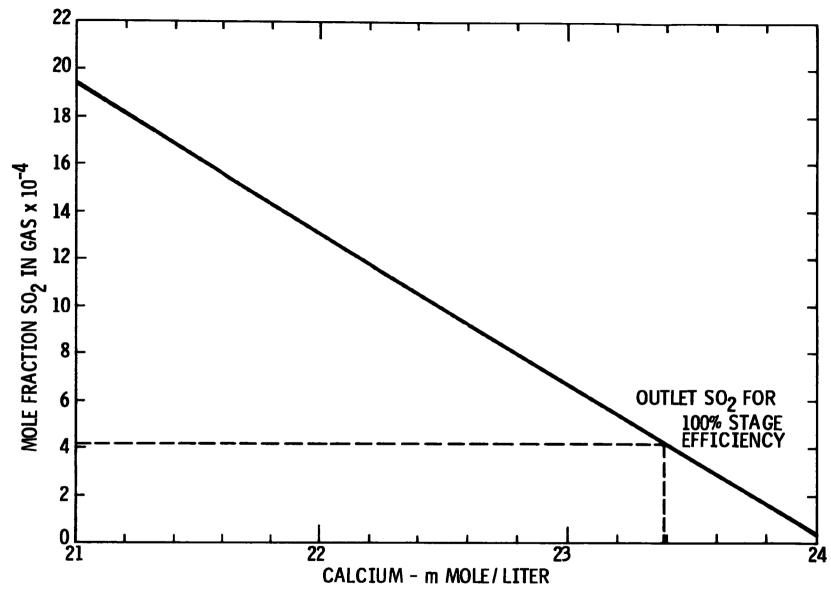
PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 20R





PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 21R





PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 22R

APPENDIX J

OPERATING DATA

AND

ANALYTICAL RESULTS

TABLE J-1. C-E APCS PROTOTYPE TAIL-END LIMESTONE TESTS

Experiment No. Date of Run	25 7/7	R /72	26R 7/10/72		27R 7/11/72	
Set Number	1 ′′′	,,,	1 "	2	1 // '	2
Time	1045-1200	1230-1315	1400-1500	1515-1600	0920-1030	1030-1130
Flue Gas (FG) Rate (cfm @ 130°F)	9,950	9,900	10,060	10,100	10,250	10,180
Additive Feed Rate (lb/hr)	318	318	510	510	516	516
Spray Water Lower (SWL) Rate (gpm)	240	250	245	235	150	150
Spray Water Upper (SWU) Rate (gpm)		-			-	-
Scrubber Liquid Lower (SLL) Rate (gpm)	180	195	180	178	135	135
Scrubber Liquid Upper (SLU) Rate (gpm)	-	-	-	-	-	-
Scrubber Bottom (SB) Rate (gpm)	60	55	65	58	15	15
Clarifier Liquid (CL) Rate (gpm)	10	10	15	15	12	12
Liquid Blowdown (LB) Rate (gpm)	Ö	0	Ō	0	0	0
Clarifier Feed (CF) Rate (gpm)	10	10	15	15	12	12
Clarifier Bottom (CB) Rate (gpm)	_	_	3	3	-	-
Filter Liquid (FL) Rate (gpm)	-	-	_	_	_	-
Spray Water (SW) Temp. (°F)	122	121	120	120	120	120
Scrubber Liquid (SL) Temp. (°F)	122	121	122	122	121	121
Scrubber Bottom (SB) Temp. (°F)	134	135	131	1 35	131	131
Inlet Gas Dew Point (°F)	114	114	113	113	104.5	104.5
Outlet Gas Dew Point (°F)	121.5	121.5	119	119	117.5	117.5
Reheater Inlet Gas Temp. (°F)	125	124	121	122	122	122
Heat Extractor Outlet Gas Temp. (°F)	295	298	315	315	298	305
Inlet SO ₂ (ppm)	2,329	2,362	2,519	2,490	2,306	2,323
Outlet SO ₂ (ppm)	956	997	1,023	999	1,114	1,099
Inlet 0 ₂ (%)*	10.6	10.6	9.8	9.8	10.0	10.0
Outlet O2 (%)	_	-	10.3	10.3	10.6	10.6
Inlet CO ₂ (%)*	8.0	8.0	8.2	8.2	8.4	8.4
Outlet CO2 (%)	-	_	7.8	7.8	7.5	7.5
Outlet SO2 corrected for air leakage (ppm)	1,022	1,070	1,097	1,072	1,195	1,177
SO ₂ Removal Efficiency (%)	56.2	54.7	56.5	57.0	48.2	49.4
Stoichiometry (%)	98.5	97.6	114.5	146.1	156.7	156.6
Solid Concentration in Spray Water	7.56	7.14	-	6.57	7.18	7.69

^{*} Average Air Leakage 7.2%

TABLE J-1. (Continued)

Experiment No.	2	8R		9R	3	30R		
Date of Run		7/12/72 7/13/72				2 7/14/72		
Set Number	7	2	1	2	1	2		
Time (50) Parks (50) 2007)	1445-1545	1605-1 <i>7</i> 05	1100-1200	1245-1315	0950-1050	1125-1225		
Flue Gas (FG) Rate (cfm @ 130°F)	10,160	10,400	10,200	10,400	10,280	10,280		
Additive Feed Rate (1b/hr)	516	516	516	516	320	320		
Spray Water Lower (SWL) Rate (gpm)	158	160	245	245	250	250		
Spray Water Upper (SWU) Rate (gpm)	150	150	225	225	235	235		
Scrubber Liquid Lower (SLL) Rate (gpm)	168	170	210	200	215	215		
Scrubber Liquid Upper (SLU) Rate (gpm)	112	110	170	180	180	180		
Scrubber Bottom (SB) Rate (gpm)	28	30	90	90	90	90		
Clarifier Liquid (CL) Rate (gpm)	15	15	15	15	10	10		
Liquid Blowdown (LB) Rate (gpm)	0	15	0	0	0	0		
Clarifier Feed (CF) Rate (gpm)	15	15	15	15	10	10		
Clarifier Bottom (CB) Rate (gpm)	-	-	-	-	-	-		
Filter Liquid (FL) Rate (gpm)	-	-	_	_	_	_		
Spray Water (SW) Temp. (°F)	125	126	126	125	124	123		
Scrubber Liquid (SL) Temp. (°F)	125	121	137	134	130	136		
Scrubber Bottom (SB) Temp. (°F)	125	132	127	126	125	125		
Inlet Gas Dew Point (°F)	119.3	119.3	114	114	115	115		
Outlet Gas Dew Point (°F)	125	125	125.5	125.5	122	122		
Reheater Inlet Gas Temp. (°F)	125	126	125	125	125	125		
Heat Extractor Outlet Gas Temp. (°F)	308	312	315	305	340	340		
Inlet SO ₂ (ppm)	2,392	2,432	2,456	2,415	2,300	2,457		
Outlet SO ₂ (ppm)	546	543	297	280	334	405		
Inlet 0 ₂ (%)*	9.3	9.3	10.6	10.6	_	-		
Outlet O2 (%)	9.8	9.8	11.6	11.6	_	_		
Inlet CO2 (%)*	8.4	8.4	7.4	7.4	_	_		
Outlet CO2 (%)	8.1	8.1	6.7	6.7	_	_		
Outlet SO ₂ corrected for air leakage (ppm)	585	582	318	300	358	434		
SO ₂ Removal Efficiency (%)	75.5	76.0	87.0	87.6	84.4	87.3		
Stbichiometry (%)	152.4	151.7	147.8	147.5	97.1	90.94		
Solid Concentration in Spray Water	6.39	6.97	8.40	8.75	-	- -		

^{*} Average Air Leakage 7.2%

TABLE J-2. SPRAY WATER FILTRATE ANALYSIS AND SOLIDS CONCENTRATION DATA FOR STEADY STATE DETERMINATION

Data	Test No.	Sample No.	Time	Ca++ (ppm)	S03 (ppm)	Total Sulfur as SO ₄ (ppm)	SO ₄ (ppm)	pH	Solid Conc. (%)
7/5/72 17/6/72 11 11 11 11 11 11 11 11 11	25R " 25R " " " " "	1 2 3 4 5 6 7 8 9 10 11 12 13	1200 1300 1340 0900 1000 1100 1200 1300 1400 1500 1600 1700 1800	1272 1287 962 755 524 793 760 799 754 737 701 753 758	952 852 524 376 376 412 404 412 392 404 352 360 412 404 104	4460 4700 3080 2450 1740 2540 2400 2550 2650 2680 2350 2390 2490 2430	3317 3677 2451 1998 1317 2045 1915 2055 2179 2195 1927 1958 1995 1945 2495	6.15 6.05 6.05 6.13 6.19 6.18	1.3 1.8 2.6 3.3 4.7 5.2 5.8 7.4 7.9 7.5 7.4 7.2 7.3
7/7/72 " "	 H	15 16 17	0730 0800 0900 1000	786 761 758 Sampling	352 372 g for Te	2580 2620	2157 2173	6.20	6.4 6.8 6.9
7/7/72	26R " " " " " " " " "	18 19 20 21 22 23 24 25 26 27 28	1330 1400 1500 1600 1700 0800 1000 1030 1100 1200 1300 1630 1645 1700	777 741 525 787 750 796 756 780 700 761 780	396 376 328 408 112 272 92 220 192 188 192	2790 2410 1780 2740 2560 2780 2510 2780 2690 2640 2680	2314 1958 1386 2250 2425 2453 2399 2516 2459 2414 2449	6.10 6.10 6.0 6.02 6.10	7.2 7.5 7.0 6.4 6.2 6.5 6.6 7.3 6.5 6.1

Sampling for Test 26R

TABLE J-2. (Continued)

Data	Test No.	Sample No.	Time	Ca++ (ppm)	SO ₃ (ppm)	Total_Sulfur as SO ₄ (ppm)	SO ₄ (ppm)	рН	Solid Conc. (%)
7/11/72	27R	29 30	0800 0830	823 812	272 192	3010 2770	2683 2539	6.1	5.9 6.5
II	11	31	0900	802	184	2870	2649	6.1	6.0
			and I	Sampling nstalling	for Tes Upper I	t 27R Marble Bed			
7/12/72 "	28R	32	0900 0930	784	232	2700	2421	6.61	6.0 6.1
II II	U U	33	1000 1030	882	212	3070	2815	6.50	6.1 6.1
1) 1)	11 11	34	1100 1130	845	192	2860	2629	6.40	6.8 6.6
# #	., H	35 36	1200 1300 1330	824 772	216 172	3010 2880	2750 2673	6.50 6.50	6.9 12.8 6.6
H		37	1400	756	156	2840	2652	6.40	5.5
			9	Sampling	for Tes	t 28R			
7/13/72 "	29R "	38	0800 0840	781	252	2970	2667	6.1	6.3 6.1
11 11 11	11 11 11	39	0900 0930 1000 1030	806	268	3070	2748		6.2 6.0 9.6 6.8
				Sampling	for Test	t 29R			0,0
7/14/72	30R	40	0645 0715	716	292	3200	2849		7.4 7.7
II II	11 11	41	0745 0815	655	196	2850	2614		7.4 7.5
II .	H	42	0845	718	196	3030	2794		6.9

Sampling for Test 30R

TABLE J-3. RESULTS OF SOLID PHASE ANALYSES EXPERIMENT 25R

7/7/72

		U4 0 C-1:4-	Cor	npositi	on in Mi	llimole	s/Gram		Nodah t
Sample Location	Time	Wt % Solids in Slurry	Total S	Ca	Mg	s0 ₂	50 ₃	^{CO} 2	Weight % Undissolved
Set 1									
Scrubber Liquid Tk	1055	7.13	4.74	7.29	0.030	3.57	1.17	2.70	1.64
Scrubber Bottoms Tk	1100	7.74	4.42	7.38	0.031	3.27	1.15	3.09	2.18
Hold Tank Effluent	1120	7.79	4.28	7.46	0.032	3.21	1.07	3.24	2.18
Marble Bed: Front	1130	7.32	4.64	7.22	0.029	3.50	1.14	2.73	1.90
Marble Bed: Back	1142	6.32	5.39	7.32	0.023	4.02	1.37	2.08	2.05
Scrubber Bottoms S	1155	7.90	4.61	7.40	0.030	3.42	1.19	2.94	2.07
Scrubber Spray	1203	7.55	4.36	7.39	0.032	3.32	1.04	3.05	2.50
Set 2									
Scrubber Liquid Tk	1235	7.04	4.85	7.31	0.027	3.70	1.16	2.69	1.73
Scrubber Bottoms Tk	1240	7.29	4.67	7.48	0.028	3.51	1.16	2.81	2.09
Hold Tank Effluent	1 300	7.14	4.31	7.31	0.029	3.30	1.01	2.94	2.04
Lime Stone			0.002	9.50	0.13	-	0.002	9.24	2.60
Lime Stone			0.015	9.45	0.13	-	0.002	9.40	2.26

TABLE J-4. RESULTS OF SOLID PHASE ANALYSES EXPERIMENT 26R

7/10/72

		Wt % Solids	Co	mpositi	on in Mi	llimole	s/G ram		lie de mind
Sample Location	Time	in Slurry	Total S	Ca	Mg	so ₂	S03	co ₂	Weight % Undissolved
Set 1									
Scrubber Liquid Tk	1400	_	4.94	7.73	0.04	3.69	1.25	2.87	1.59
Scrubber Bottoms Tk	1405	-	4.92	7.69	0.04	3.76	1.16	2.86	1.43
Hold Tank Effluent	1420	-	4.59	7.83	0.04	3.39	1.20	3.23	1.80
Marble Bed: Front	1435	-	5.02	7.72	0.04	3.76	1.26	2.75	1.53
Marble Bed: Back	1445	-	5.26	7.67	0.03	4.02	1.24	2.57	1.43
Scrubber Bottom S	1450	_	4.54	7.92	0.04	3.40	1.03	3.40	2.08
Scrubber Spray	1 455	-	3.94	8.05	0.05	2.90	1.04	4.37	2.89
Set 2									
Scrubber Liquid Tk	1515	6.24	4.54	7.67	0.06	3.39	1.15	3.30	1.84
Scrubber Bottoms Tk	1525	6.34	4.38	7.84	0.05	3.30	1.08	3.40	1.97
Hold Tank Effluent	1547	6.43	4.12	7.71	0.06	3.14	0.98	3.61	1.93
Marble Bed: Front	1555	7.94	4.31	7.69	0.06	3.29	1.02	3.40	1.97
Marble Bed: Back	1602	6.65	4.45	7.60	0.05	3.38	1.07	3.38	1.95
Scrubber Bottoms S	1615	-	4.43	7.68	0.06	3.37	1.06	3.40	1.73
Scrubber Spray	1610	6.57	3.82	7.90	0.06	2.94	0.88	4.03	2.44

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TABLE J-5. RESULTS OF SOLID PHASE ANALYSES EXPERIMENT 27R

7/11/72

		Wt % Solids	Co	mpositi	on in Mi	llimole	s/Gram		11-2-L
Sample Location	Time	in Slurry	Total S	Ca	Mg	s0 ₂	so ₃	^{CO} 2	Weight % Undissolved
Set 1									
Scrubber Liquid Tk Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front Marble Bed: Back Scrubber Bottoms S Scrubber Spray	0935 0940 0953 1000 1007 1015 1020	6.70 5.94 7.43 6.46 7.52 6.92 7.18	4.02 4.53 3.60 4.22 3.98 4.07 3.57	8.04 7.95 8.13 7.84 7.95 7.93 8.08	0.5 0.5 0.06 0.06 0.06 0.06	3.05 3.53 2.65 3.26 3.04 3.17 2.70	0.97 1.00 0.95 0.96 0.94 0.90	3.82 3.22 4.36 3.81 4.02 3.79 4.54	2.37 1.79 2.70 2.24 2.91 1.64 3.16
Set 2									
Scrubber Liquid Tk Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front Marble Bed: Back Scrubber Bottoms S Scrubber Spray	1040 1045 1100 1110 1115 1122 1128	6.80 7.47 7.38 7.08 7.97 7.49 7.69	3.91 3.72 3.49 4.24 4.44 3.69 3.53	8.00 8.06 8.07 8.01 7.96 8.12 8.08	0.06 0.07 0.06 0.06 0.06 0.06	2.94 2.78 2.62 3.23 3.45 2.76 2.74	0.98 0.93 0.87 1.01 0.99 0.93 0.79	4.00 4.28 4.65 3.79 3.55 4.50 4.45	2.03 2.12 2.28 1.96 1.83 2.72 2.58

TABLE J-6. RESULTS OF SOLID PHASE ANALYSES EXPERIMENT 28R

7/11/72

		Wt % Solids	Co	mpositi	on in Mi	llimole	s/Gram		Madaba
Sample Location	Time	in Slurry	Total S	Ca	Mg	s0 ₂	50 ₃	co ₂	Weight % Undissolved
Set 1									
Scrubber Liquid Tk Lower Scrubber Liquid Tk Upper Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front-Upper Marble Bed: Front-Lower Marble Bed: Back-Lower Scrubber Bottoms S Scrubber Spray	0250 0255 0300 0312 0325 0345 0355 0330 0337	6.04 6.07 6.60 6.17 6.61 8.45 4.72 6.93 6.40	4.74 4.31 4.09 4.02 4.30 4.07 5.49 4.16 3.84	7.82 7.87 7.94 7.88 7.86 7.95 7.52 7.90 8.01	0.06 0.06 0.07 0.07 0.06 0.07 0.04 0.07	3.19 3.26 3.07 3.04 3.21 3.00 4.26 3.14 2.86	1.05 1.05 1.02 0.98 1.09 1.07 1.23 1.02 0.98	3.65 3.70 3.70 3.74 3.40 3.79 2.16 3.67 3.91	2.11 1.92 1.93 2.12 1.81 2.12 1.20 2.02 1.99
Set 2									
Scrubber Liquid Tk Lower Scrubber Liquid Tk Upper Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front-Upper Marble Bed: Front-Lower Marble Bed: Back-Lower Scrubber Bottoms S Scrubber Spray	0405 0410 0420 0440 0512 0447 0455 0505	6.05 6.17 6.60 6.47 6.13 6.61 4.46 7.67	4.46 4.52 4.22 4.06 5.07 4.18 5.90 4.50 4.21	7.82 7.83 7.85 7.77 7.67 7.76 7.45 7.84 7.92	0.04 0.05 0.06 0.04 0.05 0.03 0.05	3.34 3.36 3.15 3.01 3.90 3.11 4.71 3.39 3.13	1.12 1.16 1.07 1.05 1.17 1.07 1.19 1.11	3.30 3.20 3.49 3.90 2.88 3.53 1.87 3.00 3.48	1.99 1.80 1.98 1.97 1.02 1.96 0.83 2.04 1.64

TABLE J-7. RESULTS OF SOLID PHASE ANALYSES EXPERIMENT 29R

7/13/72

		Wt % Solids	Cor	mpositi	on in Mi	llimole	s/G ra m		Mad ah h
Sample Location	Time	in Slurry	Total S	Ca	Mg	s0 ₂	so ₃	C0 ₂	Weight % Undissolved
Set 1									
Scrubber Liquid Tk Lower Scrubber Liquid Tk Upper Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front-Upper Marble Bed: Front-Lower Marble Bed: Back-Lower Scrubber Bottoms S Scrubber Spray	1100 1105 1115 1130 1203 1141 1155 1145	7.21 7.32 7.59 10.05 11.31 7.86 6.18 7.09 8.41	4.75 4.69 4.56 4.51 5.32 4.55 6.09 4.80 4.53	7.72 7.85 7.80 7.62 7.36 7.57 7.19 7.54 7.55	0.05 0.05 0.06 0.08 0.06 0.05 0.05	3.42 3.40 3.28 3.25 3.88 3.37 4.54 3.45 3.27	1.32 1.28 1.28 1.26 1.44 1.18 1.55 1.35	3.00 2.95 3.04 2.93 2.13 2.92 1.32 2.73 2.90	1.60 1.44 1.75 1.76 1.14 1.77 0.66 1.49 1.83
Set 2									
Scrubber Liquid Tk Lower Scrubber Liquid Tk Upper Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front-Upper Marble Bed: Front-Lower Marble Bed: Back-Lower Scrubber Bottoms S Scrubber Spray	1225 1230 1235 1256 0135 0110 0118 0130 0140	7.21 8.19 10.27 8.85 12.34 8.99 6.87 9.17	5.04 4.75 4.65 4.61 5.60 4.68 6.05 5.07	7.45 77 7.54 7.47 7.30 7.46 7.22 7.43 7.46	0.07 0.07 0.07 0.07 0.04 0.06 0.03 0.05	3.59 3.36 3.33 3.30 4.10 3.32 4.50 3.58 3.63	1.45 1.39 1.32 1.31 1.50 1.36 1.55 1.49	2.59 2.75 2.85 2.93 2.06 2.97 1.34 2.40 2.67	1.47 1.66 1.75 1.68 0.98 1.79 0.76 1.42 1.62

TABLE J-8. RESULTS OF SOLID PHASE ANALYSES EXPERIMENT 30R

7/14/72

		U4 & C-114-	Coi	mpositi	on in Mi	llimole	s/ Gra m		المامة المامة
Sample Location	Time	Wt % Solids in Slurry	Total S	Ca	Mg	s0 ₂	S0 ₃	co ₂	Weight % Undissolved
Set 1									
Scrubber Liquid Tk-Lower Scrubber Liquid Tk-Upper Scrubber Bottoms Tk Hold Tank Effluent Marble Bed: Front-Upper Marble Bed: Front-Lower Marble Bed: Back-Lower Scrubber Bottoms S Scrubber Spray	1000 1005 1015 1031 1052 1038 1045 1058 1105		5.79 5.67 5.52 5.63 6.10 5.81 6.46 5.66 5.70	7.10 7.09 7.13 7.16 7.10 7.01 7.04 7.17 7.15	0.05 0.06 0.05 0.05 0.04 0.05 0.03 0.05	4.08 4.00 3.87 3.94 4.30 4.04 4.62 3.89 3.87	1.71 1.68 1.65 1.69 1.80 1.77 1.84 1.77	1.45 1.59 1.82 1.82 1.19 1.49 0.87 1.82 1.67	1.10 1.35 1.55 1.46 1.00 1.21 0.64 1.34
Set 2									
Scrubber Liquid Tk-Lower Scrubber Liquid Tk-Upper Scrubber Bottoms Tk 1 Hold Tank Effluent Marble Bed: Front-Upper Marble Bed: Front-Lower Marble Bed: Back-Lower Scrubber Bottoms S Scrubber Spray	1125 1130 1135 1150 1210 1200 1205 1218 1227		5.81 5.82 5.64 5.67 6.12 5.88 6.39 5.95 5.63	7.09 7.15 7.15 7.14 7.03 7.10 7.00 7.09 7.13	0.04 0.04 0.05 0.05 0.05 0.05 0.04 0.04	4.02 4.02 3.89 3.94 4.25 4.06 4.56 4.17 3.91	1.79 1.80 1.75 1.73 1.87 1.82 1.83 1.78	1.46 1.47 1.59 1.64 1.33 1.27 0.86 1.27	1.34 1.35 1.59 1.51 0.99 1.26 0.71 1.22

TABLE J-9. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 25R
7/7/72
Concentration in m moles/liter

Set No.	Time	Sampling Point	Ca++	Mg++	Na+	co ₃ _	s0 ₄	so ₃	C1-	Tot.	рН	Temp. °C
1	1142	Marble Bed Back	24.67	3.32	0.72	2.14	13.82	21.0	1.38	0.3	5.29	-
1	1130	Marble Bed Front	25.85	3.29	0.70	1.48	-	45.6	1.43	0.3	5.31	-
1	1155	Scrubber Bottoms (Scrubber)	24.35	3.26	0.71	3.15	24.67	9.95	1.32	0.3	5.23	49.0
1	1203	Spray Water	24.03	3.14	0.83	5.35	18.52	8.51	1.31	0.3	6.02	_
1	1055	Scrubber Liquid at Tank	22.55	3.23	0.69	6.52	8.54	21.55	1.44	0.3	5.50	50.5
1	1100	Scrubber Bottom at Tank	20.58	3.20	0.68	6.08	8.94	18.56	1.44	0.3	5.59	49.8
1	1120	Hold Tank Effluent	17.94	3.11	0.69	6.56	15.1	6.57	1.39	0.3	6.05	49.0
. 1	1115	Clarifier Liquid	15.83	0.90	0.90	2.01	12.8	4.02	1.36	0.3	6.99	27.5
4												
2		Marble Bed Back										
2		Marble Bed Front										
2		Scrubber Bottom (Scrubber)										
2		Spray Water										
2	1235	Scrubber Liquid at Tank	22.0	3.24	0.71	5.15	9.99	20.96	1.31	0.3	5.45	49.3
2	1240	Scrubber Bottom at Tank	21.32	3.21	0.70	5.29	8.95	19.90	1.30	0.30	5.56	49.0
2	1 300	Hold Tank Effluent	17.80	3.18	0.69	6.36	12.05	9.62	1.32	0.3	6.02	48.5
2	1255	Clarifier Liquid	15.98	0.96	0.91	2.12	10.02	6.98	1.15	0.3	6.96	28.0

TABLE J-10. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 26R
7/10/72
Concentration in m moles/liter

Set No.	Time	Sampling Point	Ca++	Mg++	Na+	co ₃ =	s0 ₄	so ₃	<u>C1-</u>	Tot.	рН	Temp. °C
1	1445	Marble Bed Back	28.5	3.52	0.88	3.09	24.1	15.8	1.30	0.5	5.13	48.0
1	1435	Marble Bed Front	30.7	3.56	0.89	3.80	22.5	21.8	1.35	-	4.97	48.0
1	1450	Scrubber Bottoms (Scrubber)	27.5	3.54	0.88	3.41	19.3	17.0	1.33	-	5.32	48.5
1	1455	Spray Water	21.5	3.45	0.87	5.67	21.4	3.73	1.37		5.97	47.0
1	1400	Scrubber Liquid at Tank	24.8	3.57	0.87	6.43	24.50	12.30	1.34	0.5	5.30	49.0
1	1405	Scrubber Bottom at Tank	24.2	3.73	0.88	5 .69	21.3	9.62	1.37	0.5	5.32	48.7
1	1420	Hold Tank Effluent	19.7	3.35	0.87	6.52	20.4	3.31	1.35	0.5	6.00	48.0
1	1425	Clarifier Liquid	17.5	1.84	0.98	3.39	18.0	0.74	1.14	0.5	7.19	31.0
2	1602	Marble Bed Back	29.4	3.52	88.0	3.93	23.8	18.6	1.18	-	5.24	47.0
2	1555	Marble Bed Front	32.1	3.60	0.87	3.59	24.3	19.5	1.28	-	5.19	46.0
2	1615	Scrubber Bottom (Scrubber)	27.9	3.54	0.87	3.68	23.8	15.5	1.22	0.5	6.05	47.0
2	1610	Spray Water	22.5	3.56	0.87	6.16	-	-	1.21	-	5.29	47.0
2	1515	Scrubber Liquid at Tank	25.3	3.54	0.86	6.48	21.86	12.60	1.30	0.5	5.39	49.0
2	1520	Scrubber Bottom at Tank	27.36	3.57	88.0	5.26	24.36	11.03	1.22	0.5	5.41	48.5
2	1547	Hold Tank Effluent	20.5	3.38	0.85	5 .65	20.60	3.36	1.19	0.5	6.07	47.0
2	1542	Clarifier Liquid	17.7	1.95	0.97	4.26	18.2	0.76	1.12	0.5	7.15	31.5

TABLE J-11. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 27R
7/11/72
Concentration in m moles/liter

Set No.	Time	Sampling Point	Ca++	Mg++	Na+ 	co ₌	so <mark>=</mark>	50 ₃	C1-	Tot.	рН	Temp.
1	1007	Marble Bed Back	27.70	4.04	0.88	3.01	20.00	22.60	1.30	-	5.28	47.4
1	1015	Scrubber Bottoms (Scrubber)	26.90	3.92	0.89	3.49	16.60	23.80	1.34	0.5	5.27	48.2
1	1020	Spray Water	19.10	3.80	0.88	7.77	20.00	3.04	1.29	0.5	6.10	47.7
1	0935	Scrubber Liquid at Tank	24.33	3.94	0.86	7.45	22.65	10.02	1.35	0.5	5.42	49.2
1	0940	Scrubber Bottom at Tank	22.06	3.92	0.88	6.60	21.62	7.17	1.43	0.5	5.68	48.3
1	0953	Hold Tank Effluent	19.40	3.86	0.86	6.63	20.80	2.65	1.33	0.5	6.05	48.0
1	0950	Clarifier Liquid	17.90	2.43	0.92	4.33	18.70	1.08	1.22	0.5	6.68	32.5
2	2022	Marble Bed Back	29.80	3.86	0.90	2.79	23.10	16.90	1.25	0.5	5.59	47.2
2	2030	Marble Bed Front	30.20	3.97	0.91	3.98	25.40	18.90	1.22	0.5	5.31	48.5
2	2040	Scrubber Bottoms (Scrubber)	28.50	3.92	0.92	3.20	24.30	16.20	1.26	0.5	5.49	47.7
2	2050	Spray Water	19.60	3.89	0.89	7.51	-	-	1.41	0.5	6.22	48.1
2	1040	Scrubber Liquid at Tank	24.46	4.10	0.86	7.65	22.96	10.44	1.40	0.5	5.52	49.8
2	1045	Scrubber Bottom at Tank	21.66	3.97	0.92	7.21	20.83	7.99	1.30	0.5	5.67	49.1
2	1100	Hold Tank Effluent	24.88	3.84	88.0	6.85	14.9	12.5	1.38	0.5	6.12	47.4
2	1056	Clarifier Liquid	17.60	2.45	0.95	5.34	18.4	1.12	1.24	0.5	6.81	33.9

TABLE J-12. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 28R 7/11/72 Concentration in m moles/liter

Set No.	Time	Sampling Point	Ca++	Mg++	Na+	co ₃	so <mark>=</mark>	so ₃ =	C1-	Tot.	рН	Temp.
1	0325	Marble Bed Front-Upper	27.22	4.53	0.65	2.85	25.15	11.40	1.24	0.5	5.76	46.6
1	0355	Marble Bed Back-Lower	26.41	4.48	0.65	2.75	21.38	16.12	1.17	0.5	5.61	50.0
1	0345	Marble Bed Front-Lower	28.86	4.72	0.64	3.86	25.49	16.14	1.22	0.5	5.66	44.2
1	0330	Scrubber Bottoms (Scrubber)	26.95	4.55	0.64	4.16	22.22	16.14	1.26	0.5	5.67	50.5
1	0377	Spray Water	21.80	4.51	0.62	7.66	21.18	3.98	1.20	0.5	6.42	50.0
1	0250	Scrubber Liquid at Tank-Lower	24.65	4.50	0.70	6.71	22.30	12.60	1.15	0.5	5.69	51.4
1	0255	Scrubber Liquid at Tank-Uper	22.55	4.34	0.70	6.90	21.62	8.19	1.11	0.5	5.96	51.1
1	0300	Scrubber Bottom at Tank	23.10	4.50	0.68	5.83	21.44	8.83	1.16	0.5	5.91	50.3
1	0312	Hold Tank Effluent	20.06	4.33	0.68	6.63	20.92	3.48	1.21	0.5	6.31	50.3
1	0309	Clarifier Liquid	19.68	3.19	0.90	4.59	19.61	2.12	1.19	0.5	7.02	37.0
2	0512	Marble Bed Front-Upper	26.90	4.84	0.70	3.79	26.07	11.25	1.22	0.5	5.95	47.5
2	0455	Marble Bed Back-Lower	26.35	4.83	0.68	3.32	24.90	12.68	1.25	0.5	5.80	49.0
2	0477	Marble Bed Front-Lower	28.23	4.82	0.70	2.91	21.41	18.88	1.28	-	5.81	46.0
2	0505	Scrubber Bottoms (Scrubber)	25.73	4.90	0.64	3.68	22.53	15.68	1.29	05	5.79	50.8
2	0520	Spray Water	20.03	4.75	0.65	6.87	21.65	3.06	1.27	0.5	6.49	50.8
2	0410	Scrubber Liquid at Tank-Lower	23.52	4.65	0.66	5.33	22.34	10.05	1.19	0.5	5.98	51.5
2	0415	Scrubber Liquid at Tank-Upper	24.25	4.72	0.67	5.38	22.87	10.41	1.20	0.5	5.96	51.4
2	0420	Scrubber Bottom at Tank	23.93	4.75	0.66	5.58	23.32	9.05	1.27	0.5	6.00	50.9
2	0440	Hold Tank Effluent	19.80	4.63	0.72	6.07	21.27	3.27	1.25	0.5	5.45	50.8
2	0436	Clarifier Liquid	18.85	3.37	0.88	4.57	20.04	1.21	1.22	0.5	7.10	38.0

TABLE J-13. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 29R
7/13/72
Concentration in m moles/liter

Set No.	Time	Sampling Point	Ca++	Mg++	Na+	co ³	so ₄	s0 ⁼	<u> </u>	Tot.	рН	Temp. °C
1	1203	Marble Bed Front-Upper	23.97	5.84	0.68	4.34	26.62	5.81	1.34	0.5	5.81	45.5
1	1155	Marble Bed Back-Lower	25.80	5.94	0.66	4.05	27.43	9.91	1.35	0.5	5.40	48.5
1	1141	Marble Bed Front-Lower	26.66	5.84	0.66	3.65	23.74	15.32	1.36	0.5	5.36	46.0
1	1145	Scrubber Bottoms (Scrubber)	26.63	5.83	0.66	3.09	22.13	15.51	1.37	0.5	5.29	50.5
1	1210	Spray Water	19.60	5.89	0.66	3.09	21.53	3.58	1.36	0.5	6.02	50.5
1	1100	Scrubber Liquid at Tank-Lower	23.87	5.65	0.65	6.54	23.36	10.57	1.36	0.5	5.50	51.5
1	1105	Scrubber Liquid at Tank-Upper	21.99	5.51	0.66	3.74	24.02	6.04	1.33	0.5	5.79	51.5
1	1115	Scrubber Bottom at Tank	23.26	5.80	0.65	4.87	23.72	9.54	1.35	0.5	5.49	51.3
7	1130	Hold Tank Effluent	19.40	5.67	0.66	4.91	21.41	4.03	1.36	0.4	6.04	51.2
- 1	1125	Clarifier Liquid	18.28	3.67	0.82	3.69	20.64	0.88	1.20	0.5	6.80	35.5
7												
2	0135	Marble Bed Front-Upper	23.85	6.13	0.70	3.34	26.48	6.50	1.35	0.5	5.68	46.0
2	0118	Marble Bed Back-Lower	24.57	6.07	0.68	3.28	26.79	9.96	1.36	0.5	5.45	45.8
2	0110	Marble Bed Front-Lower	25.85	6.25	0.70	3.75	24.48	12.87	1.33	0.5	5.59	43.0
_2	0130	Scrubber Bottoms (Scrubber)	24.13	6.28	0.70	3.59	23.14	14.29	1.37	0.5	5.31	50.7
2	0140	Spray Water	19.22	6.23	0.66	5.40	21.96	3.34	1.37	0.5	6.10	50.00
2	1225	Scrubber Liquid at Tank-Lower	23.01	6.10	0.67	6.22	23.10	10.42	1.46	0.5	5.50	50.9
2	1230	Scrubber Liquid at Tank-Upper	22.10	6.06	0.68	4.20	22.76	8.43	1.26	0.5	5.70	50.9
2	1235	Scrubber Bottom at Tank	22.10	6.25	0.66	6.97	24.49	7.50	1.39	0.5	5.54	50.5
2	1256	Hold Tank Effluent	18.91	6.13	0.70	5.38	21.85	3.50	1.36	0.5	6.03	50.4
2	1252	Clarifier Liquid	18.45	3.79	0.82	3.81	20.85	0.90	1.21	0.5	7.00	35.5

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TABLE J-14. LIQUID CHEMICAL ANALYSIS AT STEADY STATE EXPERIMENT 30R
7/14/72
Concentration in m moles/liter

Set No.	Time	Sampling Point	Ca++	Mg++	Na+ 	co ₃	so ₄ =	so ₃ =	<u>C1-</u>	Tot.	pH	Temp. °C
1	1052	Marble Bed Front-Upper	24.09	6.62	0.55	2.52	26.44	8.04	1.31	0.5	5.60	47.0
1	1045	Marble Bed Back-Lower	26.59	6.92	0.60	2.81	26.27	14.66	1.37	0.5	5.09	49.3
1	1038	Marble Bed Front-Lower	27.18	6.53	0.60	2.89	24.24	17.28	1.36	0.5	5.50	43.0
1	1158	Scrubber Bottoms (Scrubber)	26.92	6.89	0.52	2.92	23.95	18.26	1.34	0.5	5.11	49.6
1	1105	Spray Water	18.78	6.86	0.55	5.91	22.34	4.18	1.37	0.5	5.90	50.0
1	1000	Scrubber Liquid at Tank-Lower*	23.57	6.67	0.58	5.45	23.61	12.50	1.34	0.5	5.36	50.7
1	1005	Scrubber Liquid at Tank-Upper*	22.39	6.46	0.57	3.85	23.90	9.02	1.31	0.5	5.58	50.5
1	1020	Scrubber Bottom at Tank*	25.21	6.71	0.58	3 .5 8	25.24	11.74	1.37	0.5	5.30	50.0
1	1031	Hold Tank Effluent	18.53	6.45	0.56	4.77	22.06	4.40	1.33	0.5	5.91	50.0
1	1023	Clarifier Liquid	18.57	4.60	0.85	4.09	20.47	1.73	1.27	0.5	6.68	36.0
2	1210	Marble Bed Front-Upper	24.43	6.78	0.50	3.08	25.36	9.70	1.34	0.5	5.94	47.0
2	1205	Marble Bed Back-Lower	27.63	7.03	0.48	3.10	26.61	17.09	1.34	0.5	5.02	49.9
2	1200	Marble Bed Front-Lower	25.53	6.95	0.48	2.36	21.14	20.48	1.34	0.5	5.08	47.80
2	1218	Scrubber Bottoms (Scrubber)	26.98	7.09	0.45	2.90	23.55	20.03	1.35	0.5	5.09	50.00
2	1227	Spray Water	18.59	6.97	0.42	6.34	21.59	5.13	1.35	0.5	5.84	50.00
2	1125	Scrubber Liquid at Tank-Lower*	24.75	6.89	0.47	4.85	24.27	14.02	1.35	0.5	5.30	50.9
2	1130	Scrubber Liquid at Tank-Upper*	22.68	6.77	0.45	4.70	23.66	10.50	1.34	0.5	5.49	50.5
2	1135	Scrubber Bottom at Tank*	25.12	6.84	0.45	4.33	24.39	15.13	1.36	0.5	5.25	50.3
2	1150	Hold Tank Effluent	18.91	6.92	0.47	5.28	21.91	4.89	1.36	0.5	5.80	50.00
2	1145	Clarifier Liquid	18.26	4.70	0.90	4.75	20.38	2.16	1.25	0.5	6.60	37.0

APPENDIX K

LIMESTONE TAIL-END SYSTEM

MATERIAL BALANCES

AND

RATE CALCULATIONS

TABLE K-1. TOTAL SULFUR MATERIAL BALANCE Experiment 25R

		F1	ow Rate	Solid Content	Total S in Solid (m mole/g)	Total S in Liquid (m mole/1)	Total S in Gas (ppm)	Total S (m mole/ min)
Marble Bed (Set #1) Entering Streams	Spray Water Gas In		l/min g mole/min	75.5 g/l	4.36	27.03	2,329	323,438 22,805
Leaving Streams	Gas Out* Scrubber Liquid Scrubber Bottom	681	g mole/min l/min l/min	68.2 g/1 79.0 g/1	5.01 4.61	34.82 34.62	956	10,035 256,397 90,529
	Total Sulfur In =	346,243	m mole/min	- Total Sul	fur Out = 35	6,961 m mole/r	nin	
Hold Tank (Set #1) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	227	l/min l/min l/min	71.3 g/l 77.4 g/l	4.74 4.42	30.09 27.50 16.82		250,643 83,901 693
Leaving Streams	Hold Tank Eff.	946	l/min	77.9 g/1	4.28	21.67		335,907
	Total Sulfur In =	335,237	m mole/min	- Total Sul	fur Out = 33	5,907 m mole/	nin	
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	208	l/min l/min l/min	70.4 g/l 72.9 g/l	4.85 4.67	30.96 28.85 17.00		260,680 81,245 676
Leaving Streams	Hold Tank	984	1/min	71.4 g/1	4.73	21.67		344,298
	Total Sulfur In =	342,401	•	- Total Sul	fur Out = 34	4,298		

^{*}Average of marble bed front and back *From hold tank eff.

TABLE K-2. TOTAL SULFUR MATERIAL BALANCE Experiment 26R

		<u>F1</u>	ow Rate	Solid Content	Total S in Solid (m mole/g)	Total S in Liquid (m mole/l)	Total S in Gas (ppm)	Total S (m mole/ min)
Marble Bed (Set #1) Entering Streams	Spray Water Gas In							
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom							
Marble Bed (Set #2) Entering Streams	Spray Water Gas In		l/min g mole/min	65.7	3.82	23.96	2,490	244,416 24,751
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	674	g mole/min l/min l/min	72.9 63.4∆	4.38 4.43	43.10 39.20	999	10,644 244,258 70,094
	Total Sulfur In =	269,167	m mole/min	- Total Su	fur Out = 324	1,995 m moles,	/min	
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	219	l/min l/min l/min	62.4 63.4	4.54 4.38	34.5 35.4 19.0		214,194 68,567 1,083
Leaving Streams	Hold Tank	946	1/min	64.3	4.12	24.0		273,314
	Total Sulfur In =	283,844	,	- Total Su	 fur Out = 27:	3,314		

ΔScrubber bottom at tank

TABLE K-3. TOTAL SULFUR MATERIAL BALANCE

Experiment 27R

		Flow Rate	Solid Content	Total S in Solid (m mole/g)	Total S in Liquid (m mole/l)	Total S in Gas (ppm)	Total S (m mole/ min)
Marble Bed (Set #1) Entering Streams	Spray Water Gas In	568 1/min 10,100 g mole/min	71.8	3.57	23.04	2,306	158,657 23,290
Leaving Streams	Gas Out Scrubber Liquid Scrubber Bottom	10,820 g mole/min 511 1/min 57 1/min	69.9 69.2	4.1 4.07	42.5 40.4	1,114	12,053 168,216 18,357
	Total Su	lfur In = 181,947 -	Total Sulf	ur Out = 198,0	526		
Marble Bed (Set #2) Entering Streams	Spray Water Gas In	568 l/min 10,000 g mole/min	76.9	3.53	27.4*	2,323	169,750 23,230
Leaving Streams	Gas Out Scrubber Liquid- Scrubber Bottom	10,730 g mole/min 511 l/min 57 l/min	75.25 74.9	4.34 3.6 9	42.15 40.5	1,099	11,792 188,424 18,062
	Total Su	lfur In = 192,980 - 1	Total Sulf	ır Out = 218,2	278		
Hold Tank (Set #1) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	67.0 59.4	4.02 4.53	33.6 28.4 19.8		154,802 16,985 891
Leaving Streams	Hold Tank Eff.	613 1/min	74.3	3.60	23.5		178,371
	Total Su	lfur In = 172,678 - 1	Total Sulfu	ır Out = 178,3	371		
Hold Tank (Set #2) Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	68.0 74.8	3.91 3.72	33.4 28.8 19.5		152,932 17,502 878
Leaving Streams	Hold Tank	613 1/min	73.8	3.49	27.1		174,498
	Total Su	lfur In = 171,312 - 1	Total Sulfu	ır Out = 174,4	198		

^{*}Hold Tank Eff.

TABLE K-4. TOTAL SULFUR MATERIAL BALANCE Experiment 28R

		F1	ow Rate	Solid Content	Total S in Solid (m mole/g)	Total S in Liquid (m mole/1)	Total S in Gas (ppm)	Total S (m mole/ <u>min)</u>
Marble Bed (Set #1)								
Entering Streams	Spray Water- Lower Bed Spray Water-	613	3 1/min	64.00	3.84	25.16		166,074
	Upper Bed Gas In		l/min g mole/min	64.00	3.84	25,16	2,392	157,946 23,800
Leaving Streams	Gas Out	10,700	g mole/min				546	5,842
	Scrubber Liquid- Lower Scrubber Liquid-	653	3 1/min	65.85	4.48	39.51		218,440
	Upper	435	i 1/min	66.10	4.30	36.55		139,539
	Scrubber Bottom	109	1/min	69.30	4.16	38.36		35,604
	Total Sulfur In =	347,820	m moles/min	- Total S	Sulfur Out = 39	9,425 m moles	s/min	
Marble Bed (Set #2)								
Entering Streams	Spray Water- Lower Bed	622	1/min	67.1	4.21	24.71		191,079
	Spray Water- Upper Bed	502	1/min	67.1	4.21	24.71		179,098
	Gas In		g mole/min	07.1	4.21	24.71	2,432	24,004
Leaving Streams	Gas Out Scrubber Liquid-	10,950	g mole/min				543	5,946
	Lower	660	1/min	55.4	5.04	38.94		209,983
	Scrubber Liquid- Upper	427	1/min	61.30	5.07	37.32		148,643
	Scrubber Bottom		1/min	76.70	4.50	38.21		44,853
	Total Sulfur In =	394,181	m moles/min	- Total S	iulfur Out = 40	9,424 m moles	/min	
Hold Tank (Set #1)								
Entering Streams	Scrubber Liquid-			50. 4	4 74			
	Lower Scrubber Liquid-	653	l/min	60.4	4.74	34.90		209,740
	Upper		1/min	60.7	4.31	29.81		126,770
	Scrubber Bottom Clarifier Liquid		l/min l/min	66.0	4.09	38.36 21.73		33,604 1,260
tanda at a	•		•	40 -				
Leaving Streams	Hold Tank Eff.	1,254	1/min	61.7	4.02	24.40		341,632
	Total Sulfur In =	371,374	m moles/min	- Total S	ulfur Out = 34	1,632 m moles	/min	
Hold Tank (Set #2)								
Entering Streams	Scrubber Liquid-	560	1/min	60 E	A A6	22 41		100 470
	Lower Scrubber Liquid-	000	1701111	60.5	4.46	32,41		199,478
	Upper		1/min	61.7	4.52	33.28		133,294
	Scrubber Bottom Clarifier Liquid		l/min l/min	66.0	4.22	38.21 21.25		37,057 1,232
Leaving Streams	Hold Tank Eff.		1/min	64.7	4.06	24.54		362,761
		•	•					302,701
	Total Sulfur In =	371,061	m moles/min	- Total S	ulfur Out = 36	2,761 m moles	/min	

TABLE K-5. TOTAL SULFUR MATERIAL BALANCE
Experiment 29R

		F1	ow Rate	Solid Content	Total S in Solid (m mole/g)	Total S in Liquid (m mole/1)	Total S in Gas (ppm)	Total S (m mole/ _min)
Marble Bed (Set #1) Entering Streams	Spray Water-Lower Spray Water-Upper Gas In	852	l/min l/min g moles/min	84.1 84.1	4.53 4.53	25.11 25.11	2,456	376,439 345,983 24,924
Leaving Streams	Gas Out Scrubber Liquid-	10,750	g moles/min				297	3,193
	Lower Scrubber Liquid-	794	1/min	70.2	5.32	38.20		326,861
	Upper Scrubber Bottom		l/min l/min	113.1 70.9	5.32 4.80	32.43 37.64		407,740 128,884
	Total Sulfur In =	747,346	m moles/min	- Total S	Sulfur Out = 80	66,678 m mole	s/min	
Marble Bed (Set #2)								
Entering Streams	Spray Water-Lower Spray Water-Upper Gas In	852	l/min l/min g moles/min	87.2 87.2	5.00 5.00	25.40 25.40	2,415	427,717 393,112 24,754
Leaving Streams	Gas Out	10,980	g moles/min				280	3,074
	Scrubber Liquid- Lower	757	l/min	79.8	5.36	37.05		351,836
	Scrubber Liquid- Upper		l/min	123.4	5.60	32.98		492,333
	Scrubber Bottom		1/min	91.7	5.07	37,43	4	171,301
	Total Sulfur In =	845,583	m moles/min	- 10tal 5	ouitur out = 1	,018,544 m mo	ies/min	
Hold Tank (Set #1) Entering Streams	Scrubber Liquid-							
	Lower Scrubber Liquid-	/94	l/min	72.1	4.75	33.93		298,865
	Upper Scrubber Bottom		l/min l/min	73.2 75.9	4.69 4.56	30.06 33,26		240,075 129,363
	Clarifier Liquid		1/min	73.3	4.50	21.52		1,226
Leaving Streams	Hold Tank Eff.	1,836	1/min	100.5	4.51	25.44		878,884
	Total Sulfur In =	669,529	m moles/min	- Total S	iulfur Out = 87	78,884 m mole	s/min	
Hold Tank (Set #2) Entering Streams	Scrubber Liquid-							
	Lower Scrubber Liquid-		1/min	72.1	5.04	33.52		300,456
	Upper Scrubber Bottom	680 241	l/min l/min	81.9 102.7	4.75 4.65	31.19 31.99		285,746 173,745
	Clarifier Liquid		1/min	104.7	4,00	21.75		1,240
Leaving Streams	Hold Tank Eff.	1,836	1/min	88.5	6.61	25.35		795,603
	Total Sulfur In =	761,187	m moles/min	- Total S	Sulfur Out = 79	95,603 m mole	s/min	

TABLE K-6. RATE CALCULATIONS

Experiment 25R

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #1)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Flue Gas In Scrubber Spray	9,792 g mole/min 908 l/min	8.51	2,329	22,805 7,727
Leaving Streams	Flue Gas Out Scrubber Liquid Scrubber Bottom	10,497 g mole/min 681 l/min 227 l/min	21.0* 9.95	956	10,035 14,301 2,259
	Rate of CaSO ₃ 1	/2 H ₂ 0 = S0 ₂ In - S0 ₃	₂ Out - Oxid. Rate		
		= 30,532 - 26	,59525 (22,805 -	10,035)	
		= 30,532 - 26	,595 - 3,192		
		= 745 m mole/n	min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray	908 1/min	18.5		16,816
Leaving Streams	Scrubber Liquid Scrubber Bottom	681 l/min 227 l/min	13.82* 24.67		9,411 5,600
	Rate of CaSO ₄	$H_20 = \Sigma SO_4 In - \Sigma SO_6$	4 Out + Oxid. Rate		
		= 16,816 - 15,0	11 + 3,192		
		= 4,997 m mole/	min		
3. CaCO ₃ Dissolution	n _				
Entering Streams	Scrubber Spray	908 1/min	24.03		21,819
Leaving Streams	Scrubber Liquid Scrubber Bottom	681 1/min 227 1/min	25.26 24.35		17,202 5,527
	Rate of CaCO ₃ Dis	solution = ΣCa Out -	ΣCa In + ΣCa Prec. R	ates	
		= 21,819 -	22,729 + 4,997 + 745		
		= 4,832 m m	ole/min	•	

^{*}Value from marble bed back only

TABLE K-6. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	inks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	681 1/min 227 1/min 38 1/min	21.00 9.95 4.02		14,301 2,258 153
Leaving Streams	Hold Tank Eff.	946 1/min	6.57		6,215
	Rate of CaSO ₃ 1/	'2 H ₂ 0 = SO ₂ In - SO	2 Out		
		= 16,712 - 6,	215		
		= 10,497 m mo	le/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	681 1/min 227 1/min 38 1/min	13.82 24.67 12.8		9,411 5,620 486
Leaving Streams	Hold Tank Eff.	946 1/ min	15.1		14,284
	Rate of CaSO ₄ 2	H ₂ 0 = ΣSO ₄ In - ΣSO	4 Out		
		= 15,517 - 14,2	84		
		= 1,233 m mole/	min		
3. CaCO ₃ Dissolution					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	681 1/min 227 1/min 38 1/min	25.26 24.35 15.83		17,202 5,527 601
Leaving Streams	Hold Tank Eff.	958 1/min	17.94		17,186
	Rate of $CaCO_3 = \Sigma C$	a Out - ΣCa In + ΣC	a Formation Rates		
	= 17	,186 - 23,330 + 10,	497 + 1,233		

= 5,586 m mole/min

TABLE K-6. RATE CALCULATIONS (Continued)

	************	Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	nks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	738 1/min 208 1/min 38 1/min	21.0 19.9 6.98		14,700 4,378 265
Leaving Streams	Hold Tank Eff.	984 1/min	9.62		9,216
	Rate of CaSO ₃ 1,	$72 \text{ H}_2\text{O} = \text{SO}_2 \text{ In - SO}_2$	2 Out		
		= 19,343 - 9,2	216		
		= 10,127 m mo	le/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	738 1/min 208 1/min 38 1/min	9.99 8.95 10.02		6,993 1,969 381
Leaving Streams	Hold Tank Eff.	984 1/min	12.05		11,544
	Rate of CaSO ₄ 2	H ₂ 0 = ΣSO ₄ In - ΣSO ₄	4 Out		
		= 9,343 - 11,544	4		
		= -2,201 m mole/	/min		
3. CaCO ₃					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	738 l/min 208 l/min 38 l/min	22.0 21.32 15.98		15,400 4,690 607
Ļ Leaving Streams	Hold Tank Eff.	984 1/min	17.80		17,052
	Rate of $CaCO_3 = \Sigma C$	a In – ΣCa Out + ΣCa	a Formation Rate		
	= 17	,052 - 20,697 + 10,1	127 - 2,201		
	= 4,	281 m moles/min			

TABLE K-7. RATE CALCULATIONS

Experiment 26R

	·	Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #1)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	9,900 g mole/min 927 l/min	3.73	2,519	24,938 3,457
Leaving Streams	Outlet Flue Gas Scrubber Liquid* Scrubber Bottom	10,613 g mole/min 681 l/min 246 l/min	18.8 17.0	1,023	10,857 12,803 4,182
	Rate of CaSO ₃ 1/	'2 H ₂ 0 = SO ₂ In - SO ₂	Out - Oxid. Rate		
		= 28,395 - 27,	,842261 (14,081)		
		= 28,395 - 27,	842 - 3,675		
		= 3,122 m mole	e/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray	927 1/min	21.4		19,837
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 246 1/min	23.3 19.3		15,867 4,747
	Rate of CaSO ₄ 2	H ₂ 0 = ESO ₄ In - ESO ₄	Out + Oxidation Rate	:	
		= 19,837 - 20,61	4 + 3,675		
		= 2,898 m mole/m	nin		
3. CaCO ₃ Dissolution	 -				
Entering Streams	Scrubber Spray	927 1/min	21.5		19,930
Leaving Streams	Scrubber Liquid* Scrubber Bottom	681 1/min 246 1/min	29.6 27.5		20,158 6,765
	Rate of CaCO ₃ Diss	olution = Σ Ca Out -	ΣCa In + ΣCa Prec. Ra	ites	
		= 26,923 - 1	9,930 + (-3,122 + 2,8	398)	
		= 6,769 m mo	ole/min		

^{*}Ave. of marble bed front and back

TABLE K-7. RATE CALCULATIONS (Continued)

			Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
	stem Remainder old Tank, Surge Ta	nks and Thickener)				
۱.	CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	681 1/min 246 1/min 57 1/min	18.8 17.0 .74		12,802 4,182 42
	Leaving Streams	Hold Tank Eff.	984 1/min	3.31		3,257
		Rate of CaSO ₃ 1/2	$2 H_2 0 = SO_2 (In) - 1$	SO ₂ (Out)		
			= 17,026 - 3,	257		
			= 13,769 m mo	le/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	681 1/min 246 1/min 57 1/min	23.3 19.3 18.0		15,867 4,748 1,026
	Leaving Streams	Hold Tank Eff.	984 1/min	20.4		20,073
		Rate of CaSO ₄ 2	1 ₂ 0 = ΣSO ₄ In - ΣSO	4 Out		
			= 21,641 - 20,0	73		
			= 1,568 m mole/n	πîn		
3.	CaCO ₃ Dissolution					
	Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	681 1/min 246 1/min 57 1/min	29.6 27.5 17.5		20,157 6,765 998
	Leaving Streams	Hold Tank Eff.	984 1/min	19.7		19,384
		Rate of CaCO ₃ Disso	olution = Σ Ca Out -	ΣCa In + ΣCa Pres. Ra	ites	
			= 19,384 - 2	27,920 + (13,769 + 1,5	668)	
			= 6,801 m ma	ole/min		

TABLE K-7. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Se	t #2)				
1. CaSO ₃ 1/2	H ₂ 0				
Entering St	reams Inlet Flue Gas Scrubber Spray	9,940 1/min 889 1/min	3.36+	2,490	24,750 2,987
Leaving Str	eams Scrubber Liquid Scrubber Bottom Outlet Flue Gas		19.05 15.5	999	12,839 3,394 10,644
	Rate of CaSO ₃	1/2 H ₂ 0 = SO ₂ In - SO	2 Out - Oxid. Rate		
		= 27,737 - 26	,877238 (24,750 -	12,839)	
		= 29,584 - 27	,896 - 2,834		
		= - 1,146 m m	pole/min		
2. CaSO ₄ 2 H	₂ 0				
Entering St	reams Scrubber Spray	889 1/min	20.6+		18,313
Leaving Str	eams Scrubber Liquid Scrubber Bottom		24.05 23.8		16,209 5,212
	Rate of CaSO ₄	2 H ₂ 0 = ΣSO ₄ In - ΣSO	4 Out + Oxid. Rate		
		= 18,313 - 21,4	21 + 2,834		
		= -274 m mole/m	nin		
3. CaCO ₃ Disso	lution				
Entering St	reams Scrubber Spray	889 1/min	22.5		20,002
Leaving Str	eams Scrubber Liquid Scrubber Bottom		30.75 27.9		20,725 6,110
	Rate of $CaCO_3$ D	issolution = Σ Ca Out -	· ΣCa In + ΣCa Prec. R	ates	
		= 26,835 -	20,002 + (-274 + - 1,	146)	
		= 5,413 m n	nole/min		

+Value taken from hold tank off

TABLE K-7. RATE CALCULATIONS (Continued)

~~~~		Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	ank and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	674 1/min 219 1/min 57 1/min	19.05 15.5 .76		12,839 3,394 43
Leaving Streams	Hold Tank Eff.	946 <b>1/mi</b> n	3 <b>.3</b> 6		3,178
	Rate of CaSO ₃ 1/	$2 H_2 0 = S0_2 In - S0$	2 Out		
		= 16,276 - 3,	178		
		= 13,098 m mo	le/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	674 1/min 219 1/min 57 1/min	24.05 23.8 18.2		16,209 5,212 1,037
Leaving Streams	Hold Tank Eff.	946 <b>l/mi</b> n	20.6		19,487
	Rate of CaSO ₄ 2	$H_20 = \Sigma SO_4 \text{ In } - \Sigma SO_4$	4 Out		
		= 22,458 - 19,4	87		
		= 3,000 m mole/	min		
3. CaCO ₃ Dissolution	1 -				
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	674 l/min 219 l/min 57 l/min	30.75 27.9 17.7		20,725 6,110 1,009
Leaving Streams	Hold Tank Eff.	946 <b>1/min</b>	20.5		19,393
	Rate of CaCO ₃ Diss	olution = ΣCa Out -	ΣCa In + ΣCa Prec. Ra	ates	
		= 19,393 -	27,844 + (13,098 + 3,0	000)	

= 7,647 m mole/min

#### TABLE K-8. RATE CALCULATIONS

# Experiment 27R

			Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Mar	ble Bed (Set #1)					
1.	CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Flue Gas In Scrubber Spray	10,100 g mole/min 568 l/min	3.04	2,306	23,290 1,727
	Leaving Streams	Flue Gas Out Scrubber Liquid* Scrubber Bottom	10,820 g mole/min 511 l/min 57 l/min	22.60 23.8	1,114	12,053 11,548 1,357
		Rate of CaSO ₃ 1/2	2 H ₂ 0 = SO ₂ In - SO ₂	Out - Oxid. Rate		
		-	= 25,017 - 24,	958 - 0.269 (23,290	- 12,053)	
			= 25,017 - 24,	958 - 2,966		
			= -2,907 m mol	e/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Scrubber Spray	568 1/min	20.0		11,360
	Leaving Streams	Scrubber Liquid Scrubber Bottom	511 1/min 57 1/min	20.0 16.6		10,220 946
		Rate of CaSO ₄ 2 H	$4_20 = \Sigma SO_4 \text{ In } - \Sigma SO_4$	Out + Oxid. Rate		
			= 11,360 - 11,16	6 + 2,966		
			= 3,160 m mole/m	in		
3.	CaCO ₃ Dissolution					
	Entering Streams	Scrubber Spray	568 1/min	19.1		10,849
1	Leaving Streams	Scrubber Liquid* Scrubber Bottom	511 1/min 57 1/min	27.7 26.9		14,155 1,533
		Rate of CaCO ₃ Disso	olution = ΣCa Out -	ΣCa In + ΣCa Prec. Ra	ites	
			= 15,688 - 1	0,849 + (3,160 - 2,90	07)	
			= 5,092 m mo	le/min		

^{*}Taken from marble bed back

TABLE K-8. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge T	anks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	22.60 23.8 1.08		11,548 1,356 48
Leaving Streams	Hold Tank Eff.	613 1/min	2.65		1,625
	Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ In - SO	2 Out		
		= 12,904 - 1,	625		
		= 11,279 m mo	le/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	20.0 16.6 18.7		10,220 946 842
Leaving Streams	Hold Tank Eff.	613 1/min	20.8		12,750
	Rate of CaSO ₄ 2	$H_20 = \Sigma SO_4 \text{ In } - \Sigma SO_4$	4 Out		
		= 12,008 - 12,7	50		
		= -742 m mole/m	in		
3. CaCO ₃					
Entering Stream	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	27.7 26.9 17.90		14,154 1,533 805
Leaving Streams	Hold Tank Eff.	613 1/min	19.40		11,892
	Rate of $CaCO_3 = \Sigma$	Ca In - ΣCa Out + ΣC	a Formation Rates		
	= ]	1,892 - 16,492 + 11,	279 - 742		

= 5,937 m mole/min

TABLE K-8. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set 2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas Scrubber Spray	10,000 g mole/min 568 l/min	3.04°	2,323	23,230 1,727
Leaving Streams	Outlet Flue Gas Scrubber Liquid Scrubber Bottom	10,730 g mole/min 511 l/min 57 l/min	17.9 16.2	1,099	11,792 9,147 923
	Rate of CaSO ₃	1/2 H ₂ 0 = SO ₂ In - S	O ₂ Out - Oxid. Rate		
		= 24,957 - 2	2,723 - (23,230 - 11,7	92) .249	
		= 24,957 - 2	2,723 - 2,848		
		= 247 m mole	/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray	568 1/min	20.0°		11,814
Leaving Streams	Scrubber Liquid Scrubber Bottom	511 1/min 57 1/min	24.25 24.3		12,392 1,385
	Rate of CaSO ₄	$2 H_2 0 = \Sigma SO_4 In - \Sigma S$	O ₄ Out + Oxid. Rate		
		= 11,814 - 13,	777 + 2,848		
		= 885 m mole/m	nin		
3. CaCO ₃ Dissolution	) -				
Entering Streams	Scrubber Spray	568 l/min	19.6		11,133
Leaving Streams	Scrubber Liquid ^A Scrubber Bottom	511 1/min 57 1/min	30.0 28.5		15,330 1,625
	Rate of CaCO ₃ Di	ssolution = ΣCa Out	- ΣCa In + ΣCa Prec. R	ate	
		= 16,955 -	11,133 + (247 + 885)		
		= 6,954 m	mole/min		

[°]Values taken from SS Set 1 ^Average of marble bed front and back

TABLE K-8. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	inks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	17.9 16.2 1.12		9,146 923 50
Leaving Streams	Hold Tank Eff.	613 <b>1/min</b>	3.04°		1,864
	Rate of CaSO ₃ 1	/2 H ₂ 0 = SO ₂ In - SO	2 Out		
		= 10,119 - 1,	864		
		= 8,255 m mol	e/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	24.25 24.3 18.4		12,391 1,385 828
Leaving Streams	Hold Tank Eff.	613 1/min	20.8°		12,750
	Rate of CaSO ₄ 2	$H_20 = \Sigma SO_4 \text{ In } - \Sigma SO_4$	4 Out		
		= 14,604 - 12,7	50		
		= 1,854 m mole/	min		
3. CaCO ₃ Dissolution	  -				
Entering Streams	Scrubber Liquid Scrubber Bottom Clarifier Liquid	511 1/min 57 1/min 45 1/min	30.00 28.5 17.6		15,330 1,624 792
Leaving Streams	Hold Tank Eff.	613 1/min	24.8		15,202
	Rate of CaCO ₃ Diss	solution = $\Sigma$ Ca Out -	ΣCa In + Ca Prec. Ra	tes	
		= 15,202 -	17,746 + (8,255 + 1,8	54)	
		= 7,565 m m	ole/min		

#### TABLE K-9. RATE CALCULATIONS

# Experiment 28R

			Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Mar	ble Bed (Set #1)					
۱.	CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Gas Inlet Spray Water-Lower Spray Water-Upper	9,950 g mole/min 613 l/min 583 l/min	3.98 3.98	2.392	23,800 2,439 2,320
	Leaving Streams	Gas Outlet	10,700 g mole/min		546	5,840
	-	Scrubber Liquid- Lower	653 <b>]/mi</b> n	16.13		10,532
		Scrubber Liquid- Upper Scrubber Bottom	435 l/min 109 l/min	11.40 16.14		4,959 1,759
		Rate of CaSO ₃ 1/	2 H ₂ O Formation = Σ	SO ₃ (1iq.) In - ΣSO ₃ (1	iq.) Out - Oxid.	Rate
			= 2	8,559 - 23,090 - 0.24	(17,002)	
			= 1	,319 m mole/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Spray Water-Lower Spray Water-Upper	613 1/min 583 1/min	21.8 21.8		13,363 12,709
	Leaving Streams	Scrubber Liquid- Lower	653 <b>1/min</b>	23.63		15,430
		Scrubber Liquid- Upper Scrubber Bottom	435 1/min 109 1/min	25.15 22.22		10,940 2,422
		Rate of CaSO ₄ 2	$H_2^0$ Formation = $\Sigma$ S0	4(liq.) In - ΣSO ₄ (liq	.) Out + Oxid. R	ate
			= 26,	072 - 28,792 + 4,150		
			= 1,4	30 m mole/min		
3.	CaCO ₃ Dissolution					
	Entering Streams	Spray Water-Lower Spray Water-Upper	613 1/min 583 1/min	21.80 21.80		13,363 12,709
	Leaving Streams	Scrubber Liquid- Lower	653 1/min	27.64		18,048
		Scrubber Liquid- Upper Scrubber Bottom	435 1/min 109 1/min	27.22 26.95		11,841 2,937
		Rate of CaCO ₃ Diss	solution = ΣCa(liq.)	Out - ΣCa(liq.) In -	· ΣCa Formation R	ates
			= 32,826 -	26,072 + 1,134 + 1,61	5	

= 9,503 m mole/min

TABLE K-9. RATE CALCULATIONS (Continued)

	· · · · · · · · · · · · · · · · · · ·	Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	nks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	653 1/min	16.13		10,532
	Upper Scrubber Bottom Clarifier Liquid	435 1/min 109 1/min 58 1/min	11.40 16.14 2.12		4,959 1,759 123
Leaving Stream	Hold Tank Eff.	1,254 l/min	3.48		4,364
	Rate of CaSO ₃ 1,	/2 H ₂ O Formation = :	ΣSO ₃ (1iq.) In - ΣSO ₃ (1	iq.) Out	
		= '	17,373 - 4,364		
		=	13,009 m mole/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid- Lower	653 <b>]/min</b>	23.63		15,430
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	435 1/min 109 1/min 58 1/min	25.15 22.22 19 <b>.</b> 61		10,940 2,421 1,137
Leaving Stream	Hold Tank Eff.	1,254 1/min	20.92		26,233
	Rate of CaSO ₄ 2	H ₂ O Formation = ES	0 ₄ (liq.) In - ΣSO ₄ (liq	.) Out	
	·	= 29	,928 - 26,233		
		= 3,	695 m mole/min		
3. CaCO ₃					
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	653 <b>l/min</b>	27.64		18,048
	Upper Scrubber Bottom Clarifier Liquid	435 l/min 109 l/min 58 l/min	27.22 26.95 19.68		11,840 2,937 1,141
Leaving Streams	Hold Tank Eff.	1,254 l/min	20.06		25,155
	Rate of CaCO ₃ Disc	solution = ΣCa(liq.	) Out - ΣCa(liq.) In +	ΣCa Formation	
		= 25,155 -	33,966 + 13,009 + 3,6	95	

= 7,893 m mole/min

TABLE K-9: RATE CALCULATIONS (Continued)

····		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set 2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Gas Inlet Spray Water-Lower Spray Water-Upper	9,870 g mole/min 622 l/min 583 l/min	3.06 3.06	2,432	24,003 1,903 1,784
Leaving Streams	Gas Outlet	10,950 g mole/min		543	5,946
	Scrubber Liquid- Lower	660 1/min	15.78		10,415
	Scrubber Liquid- Upper Scrubber Bottom	427 l/min 117 l/min	11.25 15.68		4,804 1,835
	Rate of CaSO ₃ 1/	2 $H_2$ 0 Formation = $\Sigma$	SO ₃ (1iq.) In - εSO ₃ (1	iq.) Out - Oxid.	Rate
		= 2	7,690 - 23,000 - 0.25	9 (24,003-5,946)	
		= ]	O m mole/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Spray Water-Lower Spray Water-Upper	622 1/min 583 1/min	21.65 21.65		13,466 12,622
Leaving Streams	Scrubber Liquid- Lower	660 l/min	23.15		15,279
	Scrubber Liquid- Upper Scrubber Bottom	427 l/min 117 l/min	26.07 22.53		11,132 2,636-
	Rate of CaSO ₄ 2	$H_2$ 0 Formation = $\Sigma$ 80	₃ (liq.) In - ΣSO ₃ (liq.	.) Out + Oxid. R	ate
	<u>.</u>	= 26,	088 - 29,047 + 4,680		
		= 1,7	21 m mole/min		
3. CaCO ₃ Dissolution	  -				
Entering Streams	Spray Water-Lower Spray Water-Upper	622 1/min 583 1/min	20.03 20.03		12,459 11,677
Leaving Streams	Scrubber Liquid- Lower	660 l/min	27,29		18,011
	Scrubber Liquid- Upper Scrubber Bottom	427 l/min 117 l/min	26.90 25.73		11,486 3,010
	Rate of CaCO ₃ Disse	olution = ECa(liq.)	Out - ΣCa(1iq.) In +	ΣCa Formation Ra	ates
	•	= 32,507 - 3	24,136 + 50 + 1,681		

= 10,102 m mole/min

TABLE K-9. RATE CALCULATIONS (Continued)

		Stream	Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	anks and Thickener)					
1. CaSO ₃ 1/2 H ₂ O						
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	660	l/min	15.78		10,414
	Upper Scrubber Bottom Clarifier Liquid	117	l/min l/min l/min	11.25 15.68 1.21		4,803 1,834 70
Leaving Streams	Hold Tank Eff.	1,263	1/min	3.27		4,130
	Rate of CaSO ₃ 1	/2 H ₂ 0 F	ormation = 2	:S0 ₃ (liq.) In - ΣS0 ₃ (l	iq.) Out	
			= ]	7,121 - 4,130		
			= 1	2,991 m mole/min		
2. CaSO ₄ 2 H ₂ O						
Entering Streams	Scrubber Liquid- Lower	660	1/min	23.15		15,279
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	117	l/min l/min l/min	26.07 22.53 20.04		11,131 2,636 1,162
Leaving Streams	Hold Tank Eff.	1,263	1/min	21.27		26,864
	Rate of CaSO ₄ 2	H ₂ 0 For	mation = $\Sigma SC$	) ₄ (liq.) In - ΣSO ₄ (li	q.) Out	
			= 30,	208 - 26,864		
			= 3,3	346 m mole/min		
3. CaCO ₃						
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	660	1/min	27.29		18,011
	Upper Scrubber Bottom Clarifier Liquid	117	1/min 1/min 1/min	26.90 25.73 18.85		11,486 3,010 1,093
Leaving Streams	Hold Tank Eff.	1,263	1/min	19.80		25,007
	Rate of CaCO ₃ Dis	solution	= ΣCa(liq.)	Out - ΣCa(liq.) In +	ΣCa Formation	
	-		= 25,007 -	33,600 + 12,991 + 3,3	46	

= 7,744 m mole/min

#### TABLE K-10. RATE CALCULATIONS

#### Experiment 29R

			Stream Flow Rate	Species Conc. in Lig. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Ma	rbie Bed (Set #1)					
1.	CaSO ₃ 1/2 H ₂ O					
	Entering Streams	Gas Inlet Spray Water-Lower Spray Water-Upper	10,050 g mole/min 927 l/min 852 l/min	3.58 3.58	2,456	24,682 3,318 3,050
	Leaving Streams	Gas Outlet	10,750 g mole/mir	1	297	3,193
		Scrubber Liquid- Lower	794 1/min	12.61		10,012
		Scrubber Liquid- Upper Scrubber Bottom	643 l/min 341 l/min	5.81 15.51		3,736 5,289
		Rate of CaSO ₃ 1/	2 H ₂ O Formation =	ΣSO ₃ (liq.) In - ΣSO ₃ (li	iq.) Out - Oxid.	Rate
			=	31,050 - 22,230 - 0.279	24,682 - 3,19	3)
			=	2,825 m mole/min		
2.	CaSO ₄ 2 H ₂ O					
	Entering Streams	Spray Water-Lower Spray Water-Upper	927 l/min 852 l/min	21.53 21.53		19,958 18,344
	Leaving Streams	Scrubber Liquid- Lower Scrubber liquid-	<b>794</b> 1/min	25.58		20,310
		Upper Scrubber Bottom	643 l/min 341 l/min	26.62 22.13		17,117 7,546
		Rate of Ca SO ₄ 2	$H_20$ Formation = $\Sigma$	SO ₄ (1iq.) In - ΣSO ₄ (1ic	.) Out + Oxid.	Rate
			= 3	8,302 - 44,973 + 5,995		
			≂ -	676 m mole/min		
3.	CaCO ₃ Dissolution					
	Entering Streams	Spray Water-Lower Spray Water-Upper	927 l/min 852 l/min	19.60 19.60		18,169 16,699
	Leaving Streams	Scrubber Liquid- Lower Scrubber Liquid-	794 1/min	26.23		20,827
		Upper Scrubber Bottom	643 1/min 341 1/min	23.97 26.63		15,412 9,081
		Rate of CaCO ₃ Disse	olution = ΣCa(liq.	) Out - ΣCa(liq.) In +	ΣCa Formation Ra	ates
			= 45,320 -	34,868 + 2,825 - 676		

TABLE K-10. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	anks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid- Lower	794 1/min	12.61		10,012
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	643 1/min 341 1/min 57 1/min	5.81 15.51 0.88		3,735 5,289 50
Leaving Streams	Hold Tank Eff.	1,836 1/min	4.03		7,399
	Rate of CaSO ₃ 1,	/2 $H_2$ 0 Formation = $\Sigma$	:50 ₃ (1tq.) In - E50 ₃ (1	iq.) Out	
		= ]	9,086 - 7,399		
		= l	1,687 m mole/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	794 1/min	25.58		20,310
	Upper Scrubber Bottom Clarifier Liquid	643 l/min 341 l/min 57 l/min	26.62 22.13 20.64		17,116 7,546 1,176
Leaving Streams	Hold Tank Eff.	1,836 1/min	21.41		39,309
	Rate of CaSO ₄ 2	$H_2O$ Formation = $\Sigma SO$	) ₄ (liq.) In - ESO ₄ (liq	.) Out	
		= 46,	148 - 39,309		
		= 6,8	39 m mole/min		
3. CaCO ₃					
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	794 l/min	26,23		20,826
	Upper Scrubber Bottom Clarifier Liquid	643 l/min 341 l/min 57 l/min	23.97 26.63 18.28		15,413 9,081 1,042
Leaving Streams	Hold Tank Eff.	1,836 1/min	19.40		35,618
	Rate of CaCO ₃ Diss	solution = ΣCa(liq.)	Out - ECa(liq.) In +	ΣCa Formation R	ates
	_	= 35,618 -	46,362 + 11,687 + 6,8	39	
		= 7,782 m m	nole/min		

TABLE K-10. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set 2)					
1. caso ₃ 1/2 H ₂ 0					
Entering Streams	Gas Inlet Spray Water-Lower Spray Water-Upper	.10,250 g mole/min 927 l/min 852 l/min	3.34 3.34	2,415	24,754 3,096 2,846
Leaving Streams	Gas Outlet	10,980 g mole/min		280	3,704
	Scrubber Liquid- Lower	757 1/min	11.42		8,645
	Scrubber Liquid- Upper Scrubber Bottom	680 1/min 341 1/min	6.50 14.29		4,420 4,873
	Rate of CaSO ₃ 1/2	2 H ₂ O Formation = Σ	SO ₃ (liq.) In - ΣSO ₃ (l [.]	iq.) Out - Oxid.	Rate
		= 3	0,696 - 21,642 - 0,284	4 (24,754-3,704)	
		= 3	0,696 - 21,642 - 5,978	3	
		= 3	,076 m mole/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Spray Water-Lower Spray Water-Upper	927 l/min 852 l/min	21.96 21.96		20,356 18,710
Leaving Streams	Scrubber Liquid- Lower	757 1/min	25.64		10 400
	Scrubber Liquid- Upper	680 1/min	26.48		19,409 18,006
	Scrubber Bottom	341 1/min	23.14		7,891
	Rate of CaSO ₄ 2 H	$1_2$ 0 Formation = $\Sigma$ S0	₄ (11q.) In - ESO ₄ (11q.	.) Out + Oxid. Ra	ate
		= 39,0	066 - 45,306 + 5,978		
		= -268	2 m mole/min		
3. CaCO ₃ Dissolution	1 -				
Entering Streams	Spray Water-Lower Spray Water-Upper	927 1/min 852 1/min	19.22 19.22		17,817 16,375
Leaving Streams	Scrubber Liquid- Lower	757 1/min	25.21		19,084
	Scrubber Liquid- Upper Scrubber Bottom	680 1/min 341 1/min	23,85 24,13		16,218 8,228
	Rate of CaCO ₃ Disso	Diution = $\Sigma Ca(1iq.)$	Out ~ ECa(liq.) In +	ΣCa Formation Ra	
	•	= 43,530 - 3	34,192 + 3,076 - 262		
		= 12,152 m n	nole/min		

# TABLE K-10. RATE CALCULATIONS (Continued)

····		Stream	Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	nks, and Thickener)					
1. CaSO ₃ 1/2 H ₂ O						
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	757	1/min	11.42		8,644
	Upper Scrubber Bottom Clarifier Liquid	341	l/min l/min l/min	6.50 14.29 0.90		4,420 4,873 51
Leaving Streams	Hold Tank Eff.	1,836	1/min	3.50		6,426
	Rate of CaSO ₃ 1/	2 H ₂ 0 Fc	ormation = $\Sigma$	SO ₃ (11q.) In - ΣSO ₃ (1	iq.) Out	
			= 1	7,988 - 6,426		
			= 1	1,562 m mole/min		
2. CaSO ₄ 2 H ₂ O						
Entering Streams	Scrubber Liquid- Lower	757	1/min	25.64		19,409
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	341	l/min l/min l/min	26.48 23.14 20.85		18,006 7,890 1,188
Leaving Streams	Hold Tank Eff.	1,836	1/min	21.85		40,116
	Rate of CaSO ₄ 2	H ₂ 0 = ΣS	50 ₄ (liq.) In	- ΣSO ₄ (liq.) Out		
	·	= 46	,489 <b>-</b> 40,1	16		
		= 6,	373 m mole/	min		
3. CaCO ₃ Dissolution						
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	757	1/min	25,21		19,083
	Upper Scrubber Bottom Clarifier Liquid	341	l/min l/min l/min	23.85 24.13 18.45		16,218 8,228 1,051
Leaving Streams	Hold Tank Eff.	1,836	1/min	18.91		34,718
	Rate of CaCO ₃ Diss	olution	= ΣCa(liq.)	Out - ΣCa(1iq.) In +	ΣCa Formation R	ate
			= 34,718 -	43,529 + 11,562 + 6,3	73	

= 9,124 m moles/min

# TABLE K-11. RATE CALCULATIONS

# Experiment 30R

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)		
Marble Bed (Set #1)							
1. CaSO ₃ 1/2 H ₂ O							
Entering Streams	Inlet Flue Gas	10,116 g mole/min		2,300	23,267		
-	Scrubber Spray- Lower	946 <b>1/mi</b> n	4.18		3,954		
	Scrubber Spray- Upper	889 1/min	4.18		3,716		
Leaving Streams	Outlet Flue Gas	10,845 g mole/min		334	3,624		
	Scrubber Liquid*- Lower	814 1/min	15.97		13,000		
	Scrubber Liquid*- Upper Scrubber Bottom	681 1/min 341 1/min	8.04 18.26		5,499 6,227		
	Rate of CaSO ₃ 1/	2 H ₂ 0 = ΣSO ₃ (liq.)	IN - ESO3 (114.) OUT	- Oxidation Rat	e		
	Precipitation	= 30,937 - 28	,350 - 0.3 (19,643)				
		= -3,306 m mo	le/min				
2. CaSO ₄ 2 H ₂ O							
Entering Streams	Scrubber Spray- Lower	946 <b>1/min</b>	22.34		21,134		
	Scrubber Spray- Upper	889 1/min	22.34		19,860		
Leaving Streams	Scrubber Liquid*-	814 1/min	25.25		20,553		
	Lower Scrubber Liquid*-	681 1/min	26.44		18,006		
	Upper Scrubber Bottom	341 1/min	23.95		8,167		
	Rate of $CasO_4$ 2 $H_2O = \Sigma SO_4$ (liq.) IN - $\Sigma SO_3$ (liq.) OUT + Oxidation Rate						
	Precipitation	= 40,994 - 46,7	26 + 5,893				
		= 161 m mole/mi	n				
3. CaCO ₃ Dissolution	<u>.</u>						
Entering Streams	Scrubber Spray-	946 1/min	18.78		17,766		
	Lower Scrubber Spray- Upper	889 1/min	18.78		16,695		
leaving Streams	Scrubber Liquid*-	003 17 11111	10170		,		
Leaving Streams	Lower Scrubber Liquid*-	814 1/min	26.88		21,880		
	Upper Scrubber Bottom	681 1/min 341 1/min	24.09 26.92		16,405 9,180		
	Rate of CaCO ₃ Diss	solution - ΣCa (liq.	) OUT - ΣCa (liq.) IN	ł + ΣCa Formation	Rates		
		= 47,465 -	34,461 - 3,145				
		= 9,861 m m	noles/min				

^{*}Average of Marble Bed Front and Back

# TABLE K-11. RATE CALCULATIONS (Continued)

		Stream	Flow Rate	Species Conc. in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	nnks, and Thickener	)				
1. CaSO ₃ 1/2 H ₂ O						
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	814	l/min	15 <b>.97</b>		12,999
	Upper Scrubber Bottom Clarifier Liquid	341	l/min l/min l/min	8.04 18.26 1.73		5,480 6,226 99
Leaving Streams	Hold Tank Eff.	1,892	1/min	4.40		8,325
	Rate of CaSO ₃ 1,	/2 H ₂ 0 =	ΣSO ₃ (1iq.)	IN - ΣSO ₃ (11q.) OUT		
	Precipitation	=	24,804 - 8,	325		
		=	16,479 m mo	le/min		
2. CaSO ₄ 2 H ₂ O						
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	814	l/min	25.25		20,553
	Upper Scrubber Bottom Clarifier Liquid	341	l/min l/min l/min	26.44 23.95 20.47		18,005 8,166 1,167
Leaving Streams	Hold Tank Eff.	1,892	1/min	22.06		41,737
	Rate of CaSO ₄ 2	H ₂ 0 = ΣS	0 ₄ (liq.) I	N - ΣSO ₄ (1iq.) OUT		
	Precipitation	= 47	,891 - 41,7	37		
		= 6,	154 m mole/	min		
3. CaCO ₃ Dissolution	  -					
Entering Streams	Scrubber Liquid- Lower Scrubber Liquid-	814	1/min	26.88		21,880
	Upper Scrubber Bottom Clarifier Liquid	341	l/min l/min l/min	24.09 26.92 18.57		16,405 9,179 1,058
Leaving Streams	Hold Tank Eff.	1,892	l/min	18.53		35,059
	Rate of CaCO ₃ Diss	solution	= ΣCa (liq.	) OUT - ΣCa (liq.) IN	+ ΣCa Formation	Rates
			= 35,059 -	48,522 + 16,479 + 6,19	54	

= 9,170 m mole/min

TABLE K-11. RATE CALCULATIONS (Continued)

		Stream Flow Rate	Species Conc. in Liq. (m mole/1)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
Marble Bed (Set #2)					
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Inlet Flue Gas	10,116 g mole/min		2,457	24,855
	Scrubber Spray- Lower	946 l/min	5.13		4,853
	Scrubber Spray- Upper	889 1/min	5.13		4,561
Leaving Streams	Outlet Flue Gas	10,845 g mole/min		405	4,392
	Scrubber Liquid- Lower	814 1/min	18.78		15,286
	Scrubber Liquid- Upper Scrubber Bottom	681 1/min 341 1/min	9.70 20.03		6,606 6,830
	Rate of CaSO ₃ 1,	/2 H ₂ 0 = ΣSO ₃ (1iq.)	IN - ΣSO ₃ (11q.) OUT	- Oxidation Rat	e
	Precipitation	= 34,269 - 33	,114 - 0.305 (24,855	- 4,392)	
		= <b>-</b> 5,086 m mo	le/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Spray- Lower	946 1/min	21,59		20,424
	Scrubber Spray- Upper	889 1/min	21.59		19,193
Leaving Streams	Scrubber Liquid- Lower	814 1/min	23,88		19,438
	Scrubber Liquid- Upper Scrubber Bottom	681 1/min 341 1/min	25,36 23,55		17,270 8,030
	Rate of CaSO ₄ 2	$H_20 = \Sigma SO_4$ (11q.) I	N - ΣSO ₄ (11q.) OUT +	Oxidation Rate	
	Precipitation	= 39,617 - 44,7	38 + 6,241		
		= 1,120 m mole/	min		
3. CaCO ₃ Dissolution	•				
·	Scrubber Spray- Lower	946 1/min	18,59		17,586
	Scrubber Spray- Upper	889 1/min	18.59		16,526
Leaving Streams	Scrubber Liquid- Lower Scrubber Liquid-	814 1/min	26.58		21,636
	Upper Scrubber Bottom	681 1/min 341 1/min	24.43 26 <b>.</b> 98		16,636 9,200
	Rate of CaCO ₃ Dis	solution = $\Sigma$ Ca (11q.	) OUT - ΣCa (liq.) IN	l + ΣCa Formation	Rate
		= 47,472 -	34,112 - 3,966		

= 9,394 m mole/min

TABLE K-11. RATE CALCULATIONS (Continued)

		Stream Flow	Species Conc. Rate in Liq. (m mole/l)	Species Conc. in Gas (ppm)	Species Flow Rate (m mole/min)
System Remainder (Hold Tank, Surge Ta	nks and Thickener)				
1. CaSO ₃ 1/2 H ₂ O					
Entering Streams	Scrubber Liquid- Lower	814 1/min	18.78		15,286
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	681 l/min 341 l/min 57 l/min	20.03		6,605 6,830 123
Leaving Streams	Hold Tank Eff.	1,892 1/min			9,252
	Rate of CaSO ₃ 1,	/2 H ₂ 0 = ΣSO ₃	(1iq.) IN - ΣSO ₃ (11q.) OUT		
	Precipitation	= 28,84	4 - 9,252		
		= 19,59	2 m mole/min		
2. CaSO ₄ 2 H ₂ O					
Entering Streams	Scrubber Liquid- Lower	814 1/min	23.88		14,438
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	681 l/min 341 l/min 57 l/min	23.55		17,270 8,030 1,167
Leaving Streams	Hold Tank Eff.	1,892 1/min	21.91		43,700
	Rate of CaSO ₄ 2	$H_20 = \Sigma SO_4$ (1	iq.) IN - ΣSO ₄ (11q.) OUT		
	Precipitation	= 45,905	- 43,700		
		= 2,205 m	mole/min		
3. CaCO ₃ Dissolution	  -				
Entering Streams	Scrubber Liquid- Lower	814 1/min	26.58		21,636
	Scrubber Liquid- Upper Scrubber Bottom Clarifier Liquid	681 1/min 341 1/min 57 1/min	26.98		16,636 9,200 1,041
Leaving Streams	Hold Tank Eff.	1,892 1/min	18.19		34,415
	Rate of CaCO ₃ Dis	solution = ΣCa	(11q.) OUT - ECa (11q.) IN	l + ΣCa Formation	n Rate
	_	= 34,	415 - 48,013 + 19,592 + 2,2	205	

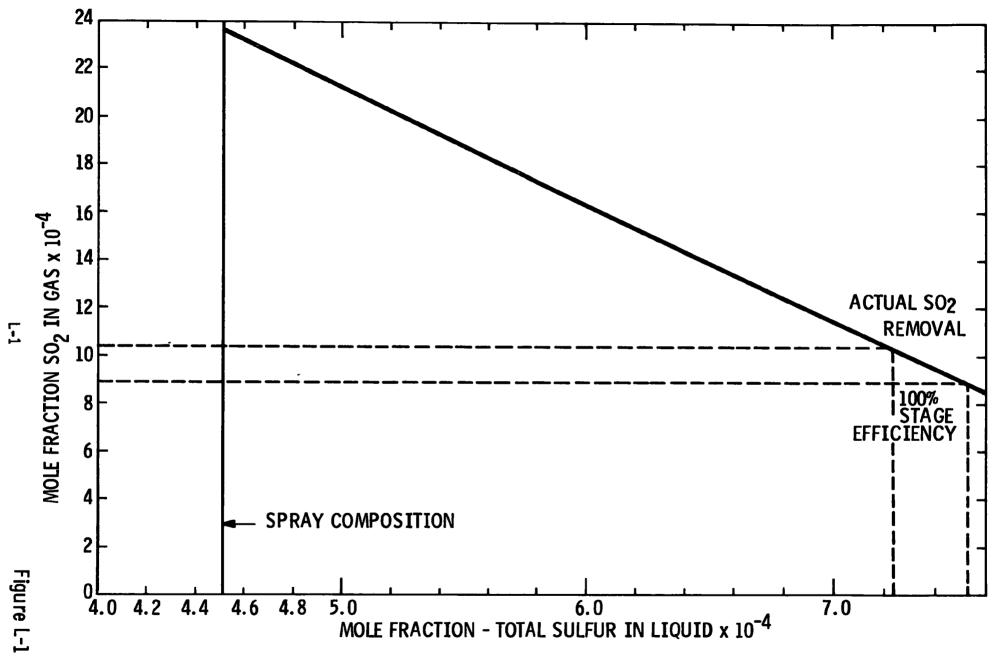
= 8,199 m moles/min

# APPENDIX L

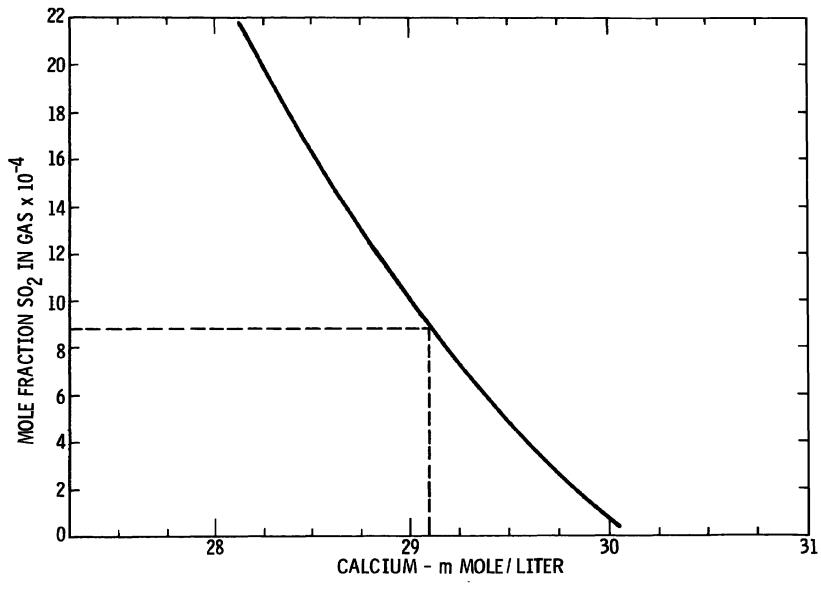
LIMESTONE TAIL-END SYSTEM

DISSOLUTION RATE DETERMINATION

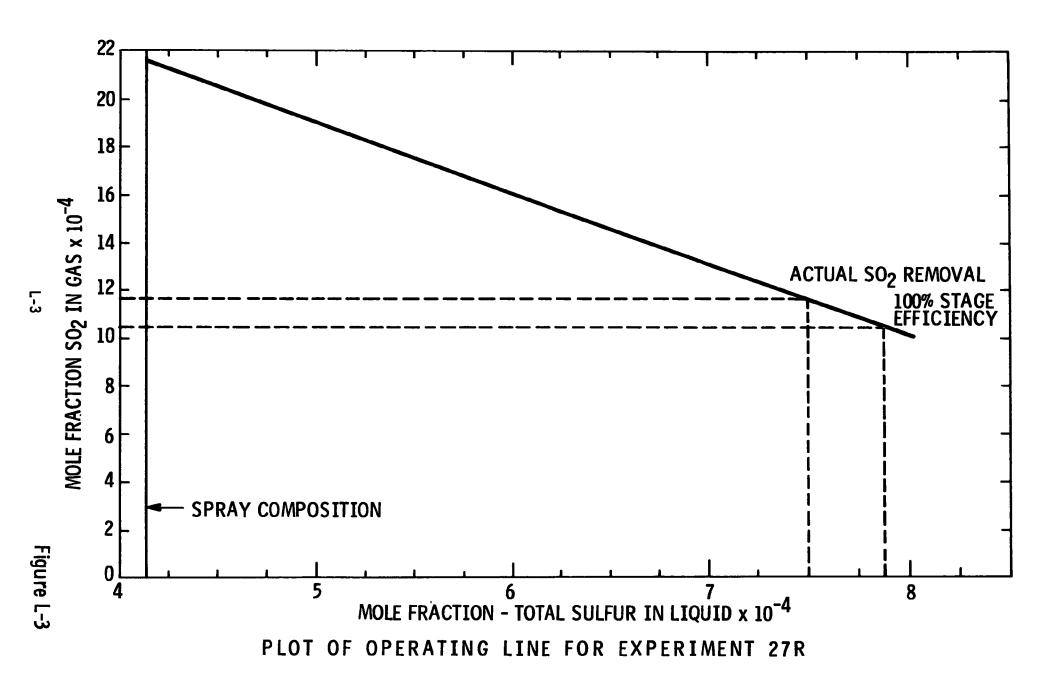
DIAGRAMS

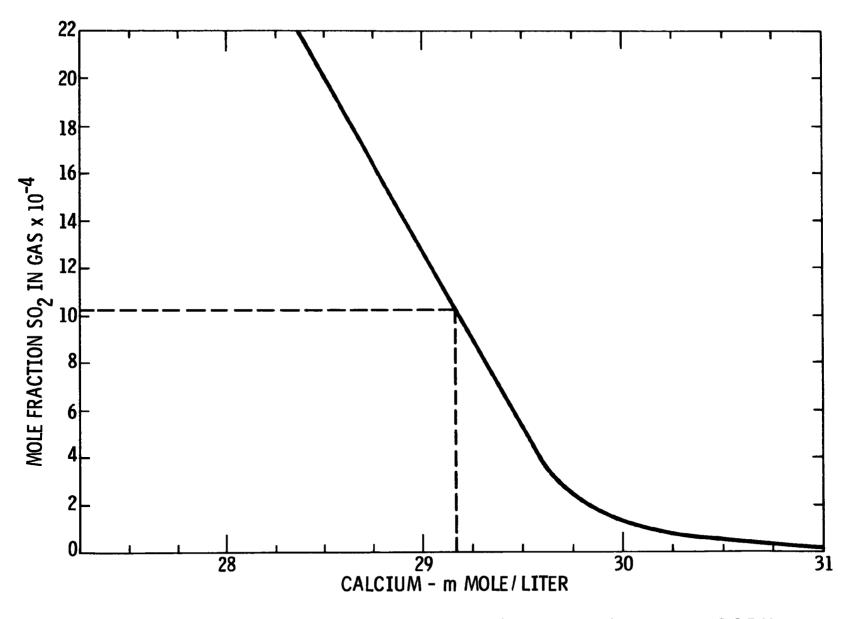


PLOT OF OPERATING LINE FOR EXPERIMENT 26R

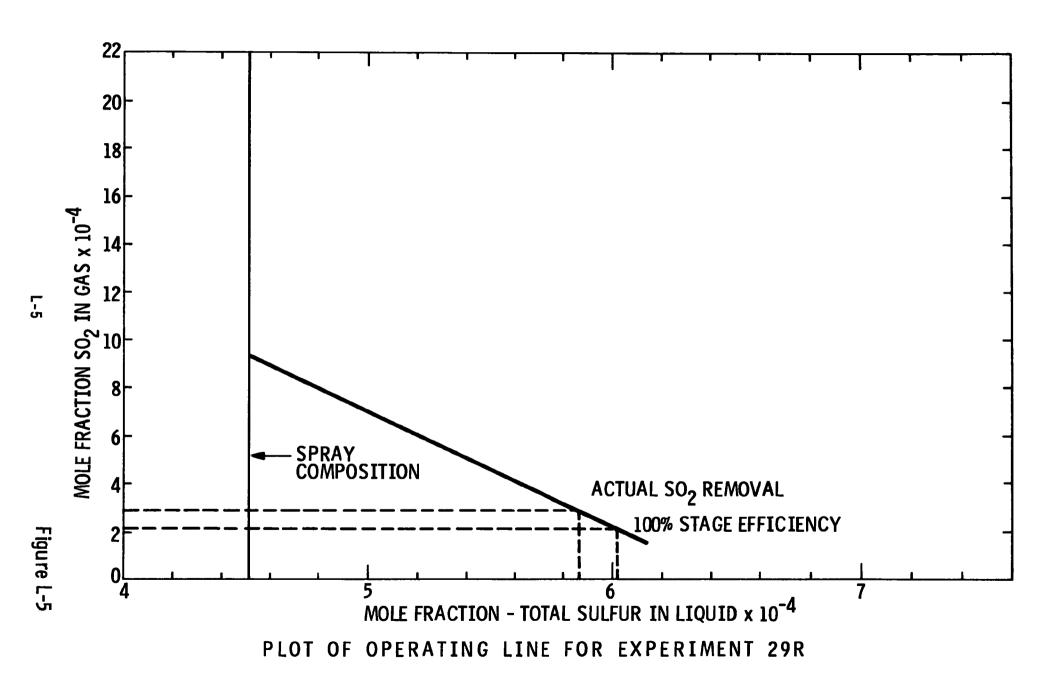


PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 26R

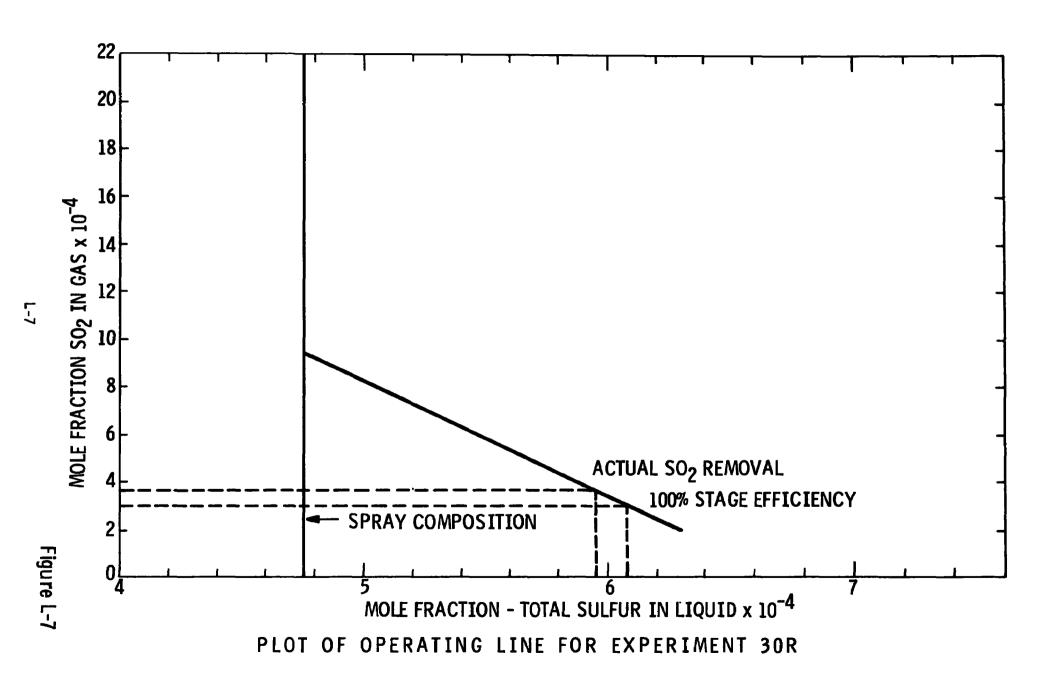




PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 27R



PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 29R



PLOT OF CALCIUM vs PARTIAL PRESSURE OF SO₂ FOR SCRUBBER EFFLUENT - EXPERIMENT 30R

TECHNICAL REPORT DATA (Picase read instructions on the reverse before completing)						
1. REPORT NO. 2. EPA-650/2-75-052	3. RECIPIENT'S ACCESSION NO.					
4. TITLE AND SUBTITLE Optimization of a Lime/Limestone Wet Scrubbing	June 1975					
Process for SO2 and Particulate Removal in a Marble Bed Scrubber	6. PERFORMING ORGANIZATION CODE					
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15. SUPPLEMENTARY NOTES

scrubber system. Results of 16 once-through soluble system tests, using sodium carbonate scrubbing solution, showed that the scrubber is a very good liquid/gas contacting device for SO2 removal from flue gases with an overall efficiency of 90 to 95 percent. Liquid/gas ratio and scrubber liquid composition significantly affected SO2 removal; other variables had little or no effect. Results of six limestone furnace injection system tests, using boiler calcined limestone and fly ash mixture, showed that solids concentration in the spray slurry and liquid/gas ratio significantly affected SO2 removal. Results of six limestone tail-end system tests, using commercial limestone in a dual marble bed scrubber, showed that the SO2 removal efficiencies of the lower and upper beds are the same, based on the SO2 concentrations entering the respective beds. It was demonstrated that scale-free operation of both the furnace injection and the tail-end systems can be achieved in a closed loop system, without employing liquid blowdown, by maintaining 8 to 10 percent solids in the spray slurry.

17.	KEY WORDS AN	D DOCUMENT ANALYSIS		
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