		Na
2	S	An I

Acture Environment and Pollution Technology n International Quarterly Scientific Journal

Vol. 9 No. 3

2010

pp. 559-562

Original Research Paper

# Removal of Sulphur Dioxide from Exhaust Sulphurous Flue Gases at Shree Cement Ltd., Beawar for Environment Cleaning

#### Rashmi Sharma, Shveta Acharya and Arun Kumar Sharma

Department of Chemistry, S.D. Govt. College, Beawar-305 901, Rajasthan, India

Nat. Env. Poll. Tech. ISSN: 0972-6268 www.neptjournal.com

# ABSTRACT

Laboratory studies were conducted to know about the effect of pH of sodium hydroxide solution and its temperature for absorption of SO<sub>2</sub> contained in flue gases. it was found that the pH of the solution should be alkaline for good absorption of SO<sub>2</sub>. Because of the acidic nature of SO<sub>2</sub>, the reaction is restricted in acidic solution and maximum absorption of SO<sub>2</sub> was found in temperature range between 20-25°C. At higher temperature reversible reaction may take place and partially formed product may be changed back into initial reactants.

Key Words: Flue gases Sulphur dioxide Desulphurization Shree Cement Ltd.

## INTRODUCTION

Shree Cement Ltd., having a coal based power plant, is one of the top ten cement producers in India and as a leading player in north Asia. The main pollutants from thermal power plants are dust and objectible gases like CO,  $CO_2$ ,  $SO_2$ ,  $NO_2$ , etc. Sulphur dioxide is a major constituent in air pollution (Arthur et al. 1975). Coal is re-emerging as a dominant fuel for power generation in various power plants. Various coals such as petcock, lignite, bituminous, etc. are used in power plants in which sulphur is 6.0 %, 4.0 % and 3.8 % respectively. The common element in fuels is carbon, which is principle combustible constitute of all fossils. Oxygen, nitrogen, hydrogen and sulphur are not combustible elements (Ram 2005). Sulphur in coal reacts with oxygen and forms  $SO_2$  and  $SO_3$ . Sulphur dioxide is a major constituent in air pollution, which affect the environment in number of ways like acid rain, corrosion and severe damage to health (Maohong 2002). A 22.5 MW power generation unit emits 1690 tons of  $SO_2$  per year (Sheth et al. 2006, Shih-Wu Sung et al. 2002).

Flue gas desulphurization (FGD) is the technique used for removal of sulphur dioxide from the exhaust flue gases in power plants. The aim of this study is to reduce the  $SO_2$  in environment and to produce a byproduct with SO<sub>2</sub> to control pollution.

In accordance with the invention, flue gases containing sulphur dioxide are passed through a solution rich in sodium ions.  $SO_2$  reacts with these ions to produce sodium sulphate. Most complete removal of  $SO_2$  in flue gases has been observed using this process in the power plant of Shree Cement Ltd, Beawar, Rajasthan.

## MATERIALS AND METHODS

**Experimental A: Effect of pH of solution for SO<sub>2</sub> absorption:** All experiments were conducted on stack monitoring kit (Model No. and Make VSS1, 141 DTH 2005, Vayubodhan). First of all, SO<sub>2</sub> monitoring kit for SO<sub>2</sub> measurement was set up at chimney inlet of Boiler No. 4 of the industry. Ten

percent sodium hydroxide solution was taken in first impinger and flue gases containing  $SO_2$ were passed through it using a flexible pipe connected to the  $SO_2$  monitoring kit. The  $SO_2$  flow was controlled using an inlet line rotameter and was maintained at 3 L/min.

One end of flexible pipe was connected to chimney inlet for suction of  $SO_2$  and other end was connected to  $SO_2$  monitoring kit which has impingers of 10 cm diameter and 100 cm length. The impinger was filled with 100 mL of scrubbing media i.e., NaOH solution. Samples of 10 mL were collected from the bottom at intervals of every 15 minutes and each sample was

Table 1: Operating conditions for  $SO_2$  absorption in Sodium hydroxide solution.

S. No.	Operating Condition	Value
1	Initial Concentration of Sodium	10 %
2	pH of solution	12.57
3	Total liquid hold up	100 mL
4	Temperature of solution	varying
5	Time period for reaction	0.5 hr
5	Flow of flue gas in impinger	3 LPM
7	SO <sub>2</sub> load in flue gas	3000-3200 ppm
8	Flue gas Temperature	<b>135</b> ℃
9	Flue gas flow in duct of ESP O/L	150522 m <sup>3</sup> /hr
10	Pet Coke Feeding Rate	13 ton/ hr
11	Lime Stone Feeding Rate	1.0 ton/hr

analyzed for pH and also titrated with 1 M oxalic acid to determine fall in concentration of NaOH. Three parameters, % SO<sub>3</sub> (gravimetric), % SO<sub>2</sub> (volumetric) and % alkalinity were analyzed by the methods of BIS (8-13).

S. No.	S. No. Time(Min.) pH of solution		Volume of 1 M Oxalic acid consumed in titration using phenolphthalein indicator (mL)	Conc. of NaOH (%)	
1	0	12.57	20.05	80.06	
2	15	10.62	15.56	62.2	
3	30	8.82	3.5	14.2	
4	45	7.95	1.23	4.8	
5	60	5.62	0.56	2.2	
6	75	4.75	0.32	1.2	

Table 2: Effect of pH of NaOH solution for absorption of SO<sub>2</sub>



Fig. 1: Experimental setup by using SO<sub>2</sub> monitoring kit for absorption of SO<sub>2</sub>.

**Experimental B: Effect of temperature of NaOH solution for maximum recovery of SO<sub>2</sub>:** Experiment was conducted by passing flue gases directly at different temperatures of 10 % NaOH at the rate of 3 L/min. The pH of the solution was around 12.57 and time for every experiment was 30 min. Experimental set up is shown in Fig. 1 and operating conditions for SO<sub>2</sub> absorption are given in Table 1.

### **RESULTS AND DISCUSSION**

Fig. 2 and Table 2 report the effect of pH of NaOH solution and absorption of  $SO_2$  and it is confirmed that when there is an increase in the time period for absorption of  $SO_2$  in NaOH solution, then there is a significant decrease in pH. Fig. 3 reports that with the increase of time period for absorption of  $SO_2$  in NaOH solution there is a significant decrease in concentration of NaOH solution.

Fig. 4 and Table 3 show that by increasing temperature of NaOH solution, there is significant decrease in recovery of  $SO_2$ . It can be confirmed from Figs. 5 and 6 and Table 4 that precipitate,

S. No.	Temperature of solution	Initial Conc. of SO <sub>2</sub> (ppm)	Conc. of SO <sub>2</sub> after formation of Sulphate (ppm)	Recovery (%)
1	20-25°C	3080	302	90.18
2	25-30 °C	3080	566	81.62
3	30-35 °C	3080	675	78.08

Table 3: Effect of temperature of sodium hydroxide solution for maximum recovery of SO2.

Table 4: Analysis results of precipitate which was prepared by varying the temperature of NaOH solution.

S.No	Temperature of NaOH solution	Yield(g)	%SO <sub>3</sub>	%SO <sub>2</sub>	%Na <sub>2</sub> SO <sub>4</sub>	% Alkalinity
1	20-25°C	9.77	0.62	38.72	1.100	1.68
2	25-30°C	9.25	0.42	31.92	0.745	1.80
3	30-35°C	9.06	0.22	17.87	0.390	1.95





Fig. 2: Figure depicting relation between pH of NaOH solution and absorption of SO<sub>2</sub>





which is formed by lower temperature of solution has maximum %  $SO_3$  (gravimetric) and %  $SO_2$  (volumetric). Fig. 7 reports that when %  $SO_3$  decreases then % alkalinity increases in precipitate.

### CONCLUSION

1. The change in colour of the solution can be seen easily, i.e., initial colour is white, and after passing SO, colour is yellow.

Nature Environment and Pollution Technology 

Vol. 9, No. 3, 2010



- 2. The pH should be more alkaline for good absorption of SO<sub>2</sub>.
- 3. Absorption and precipitisation of sulphate is a temperature sensitive reaction. At higher temperature the sodium sulphate is formed but remains soluble as solubility increases with the increasing temperature. The maximum recovery of  $SO_2$  was found at the temperature in the range of 20-25°C and this seems to be optimum temperature.
- At higher temperature, the reversible reaction may take place, and partially formed sodium sulphate may be changed back into NaOH.
- 5. Prepared Na<sub>2</sub>SO<sub>4</sub> can be used as a home laundry detergent, and also used in paper production. In the laboratory Na<sub>2</sub>SO<sub>4</sub> is used as an inert drying agent for removing traces of water from organic solutions.

#### ACKNOWLEDGEMENT

Authors are thankful to Principal and HOD of Chemistry, S.D. Govt. College for facilities provided. One of the authors Arun Sharma is highly thankful to Mr. V. Kapil and Mr. G.P. Kalwar of Shree Power for their support and contribution during the research.

#### REFERENCES

- Arthur, Kohl, Richard and B. Nielsen 1975. Gas Purification. Fifth edition, Gulf Publishing Company, Texas.
- BIS 1985. Method for Measurement of Emission From Stationary Sources. Indian Standard . IS 11255 (Part 2).
- BIS 2001. Method for Measurement of Emission From Stationary Sources. Indian Standard . IS 5182 (Part 2).
- BIS 1992. Indian Standard Methods of Chemical Analysis of Limestone, Dolomite and Allied Materials, Determination of Iron Oxide, Calcium Oxide and Magnesia. 1 <sup>st</sup> revision, IS No. 1760, Part 3, 1992.
- BIS 1987. Indian Standard Methods of Sodium bi Sulphite Technical. Fourth revision, IS 248: 1987
- BIS 1991. Indian Standard Methods of Caustic Soda, Pure and Technical. Third revision). IS 252:1991
- BIS 1964. Indian Standard Methods of Sampling and Test (Physical and Chemical) for Water Used in Industries. IS 3025-1964.
- Maohong Fan and Robert C. Brown 2002. A process for synthesizing polymeric ferric sulfate using sulphur dioxide from coal combustion. International Journal of Environment and Pollution, 17: 102-109.
- Ram, S. and Gosavi 2005. Flue gas desulphurization An overview of system and technologies. Environmental Pollution Control Journal, 8(5): 5-8.
- Sheth, K.N., Patel and Patel Neha, J. 2006. Effect of concentration in absorption of sulphur dioxide with sodium hydroxide. Environmental Pollution Control Journal, 9(2): 14-18.
- Shih-Wu Sung, Robert and C. Brown, 2002. Synthesis, characterization and coagulation performance of polymeric ferric sulphate. Journal of Environmental Engineering, 128: 483-490.
- Slack A. V. and Hollindon, G.A. 1975. Sulphur Dioxide Removal from Stake Gases. Second edition, Noyes Data Corporation, New Jersey.