



## Research on selectivity removing SO<sub>2</sub> from flue gas with a novel absorbent

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### Abstract:

Compared with the traditional methods of removal SO<sub>2</sub> from flue gas, the organic solvent absorption has some advantages as low investment, high SO<sub>2</sub> absorption efficiency and desorption efficiency. For the industrial application of organic solvent absorption as soon as possible, some laboratory research on selectively removing SO<sub>2</sub> and NO<sub>x</sub> from flue gas in the presence of CO<sub>2</sub> and an enlarged experiment has been done with a novel absorbent of Mn (II) + DMSO. The effect on desulfurization selectivity for absorbents is studied. And the regeneration capacities for absorbent are researched. The result shows that the novel absorbent has not only strong desulfurization efficiency, but also good selectivity for SO<sub>2</sub> and CO<sub>2</sub>, the feasibility of desulfurization absorbent has been proved.

**Keyword:** selectivity; desulfurization; flue gas; the enlarged experiment; DMSO +Mn (II)

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## 1 Introduction

Removal of SO<sub>2</sub>, CO<sub>2</sub> and NO<sub>x</sub> from flue gas is one of very important research projects in the world. In the previous literature [1], some laboratory research on removing SO<sub>2</sub> from flue gas with DMSO absorbent have been studied. But removing SO<sub>2</sub> and NO<sub>x</sub> from flue gas in the presence of CO<sub>2</sub> had not been reported. As we know, there is not only sulfur dioxide, but also a lot of CO<sub>2</sub> in flue gas from industrial emissions. Because SO<sub>2</sub> and CO<sub>2</sub> are all acidic gases, and the concentration of sulfur dioxide is much lower than carbon dioxide in the flue gas. If the absorbent has not selectivity for acid gas such as SO<sub>2</sub> and CO<sub>2</sub>, it will lead to a serious decrease for desulfurization efficiency because the absorbent absorbs a large number of CO<sub>2</sub>, therefore the good selectivity for SO<sub>2</sub> and CO<sub>2</sub> is necessary in order to reach higher desulfurization efficiency. Therefore, in this paper, based on our selected organic absorbent DMSO [2-7], a small amount of Mn (II) catalyst is added in the system of organic solvent absorption, it is found that the desulfurization efficiency with a small amount of Mn (II) catalyst has been much improved compared with pure physical solvent of DMSO, Mn (II) played a significant catalytic role [8]. In order to further verify the selectively desulfurization effect of Mn (II) + DMSO. Some laboratory research on selectively removing SO<sub>2</sub> and NO<sub>x</sub> from flue gas with Mn (II) + DMSO absorbent have been carried out. The research provides a basis for the industrial application of flue gas desulfurization technology.

## 2 Experimental Sections

The experiment includes two sections: (1) the selectivity absorption experiment of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> and NO<sub>x</sub> in the presence of CO<sub>2</sub>. (2) the enlarged experiment.

### 2.1 The selectively absorption experiment of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> and NO<sub>x</sub> in the presence of CO<sub>2</sub>.

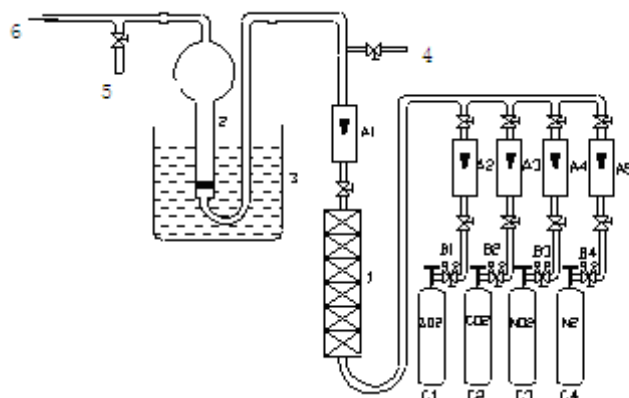
The technological process of the selectivity SO<sub>2</sub> removal experiment is shown in Figure 1. Sulfur dioxide with a mole fraction purity  $x(\text{SO}_2) \geq 0.997$ , carbon dioxide with a mole fraction purity  $x(\text{CO}_2) \geq 0.99$ , NO and NO<sub>2</sub> with a mole fraction purity  $x(\text{NO}_x) \geq 0.99$ , nitrogen of purity  $x(\text{N}_2) \geq 0.99999$  are obtained from cylinders. Sulfur dioxide, carbon dioxide, NO<sub>x</sub> and nitrogen are mixed by a gas mixer with static agitation to simulate the flue gas. A flowrator is used to control the gas flow. CO<sub>2</sub> is analyzed by CO<sub>2</sub> analyzer (type RD-7AG, Nanjing, China). Nitrogen Oxides is analyzed by NO<sub>x</sub> analyzer (type NA-721 Nitrogen Oxides Analyzer, China). O<sub>2</sub> is analyzed by AUS-gas analyzer with gas absorption method. A SO<sub>2</sub> analyzer (type NTS 100, Nanjing, China) is used for the SO<sub>2</sub> analysis of the gas phase from the inlet and tail gases. The SO<sub>2</sub> concentration in the gas phase before and after absorption is analyzed by SO<sub>2</sub> analyzer, and then the removal efficiency is calculated.

$$\text{The removal efficiency, \%} = \frac{C_0 - C}{C_0} \times 100\% \quad (1)$$

In which, C<sub>0</sub> - The SO<sub>2</sub> or NO<sub>x</sub> concentration before absorption;

C - The SO<sub>2</sub> or NO<sub>x</sub> concentration after absorption

All the chemical reagents used are AR grade. Purified and deionized water of conductivity <0.06 μS·cm<sup>-1</sup> is used in the measurements.



**Fig. 1 The technological process of SO<sub>2</sub> absorption**

C1: SO<sub>2</sub> cylinder, C2: CO<sub>2</sub> cylinder, C3: NO<sub>x</sub> cylinder, C4: N<sub>2</sub> cylinder, B1-B4: valve, A1-A5: flowrator,  
 1- a gas mixer with static agitation unit, 2-absorption tube 3- constant temperature bath.  
 4-inlet gas to SO<sub>2</sub> analyzer, 5- outlet gas to SO<sub>2</sub> analyzer, 6- to fume hood

## 2.2 The enlarged experiment

The enlarged experimental apparatus is shown in Figure 2, the absorption tower is a packed tower, diameter is 25 mm, inside packing diameter is  $\Phi = 5 \text{ mm} \times 10 \text{ mm}$ . The regenerative tower is a packed tower of 25 mm  $\times$  78 mm, with heat preservation jacket. The flow and the direction of the absorption solution at rectifier bottom are controlled by the recycle pumps. SO<sub>2</sub>, CO<sub>2</sub>, O<sub>2</sub> and N<sub>2</sub> from cylinders are mixed by a gas mixer with static agitation to simulate the flue gas. A flowrator is used to control the gas flow. The mixed gas is absorbed through the saturation flask in absorption tower, A SO<sub>2</sub> analyzer (type NTS 100, Nanjing, China) is used for the SO<sub>2</sub> analysis of the gas phase. The SO<sub>2</sub> concentration of solution was measured by iodometric method.

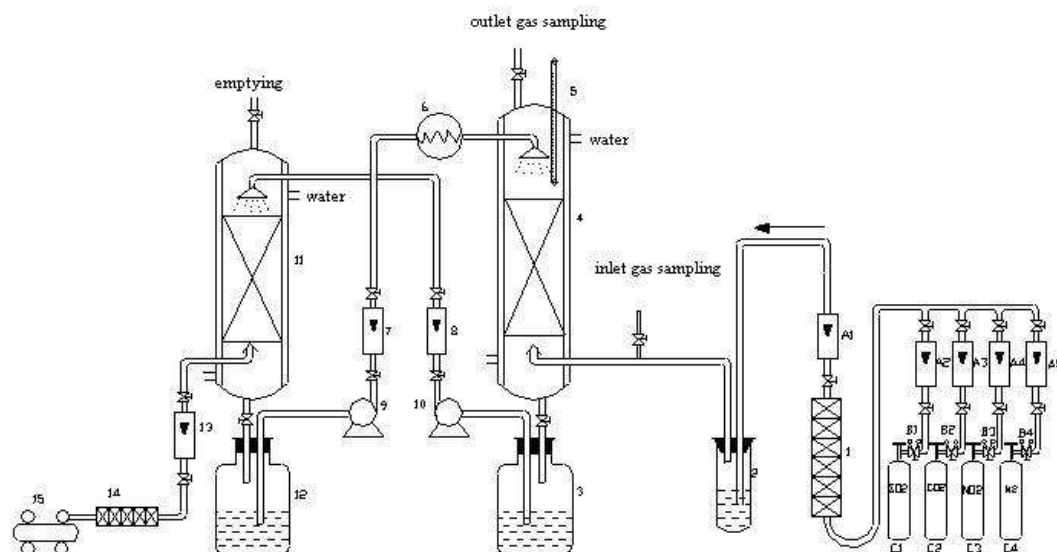
The effect of ratio gas to liquid, rate of flow, temperature and inlet gas composition on the desulfurization selectivity is determined, and the absorption and desorption efficiency was measured.

Both SO<sub>2</sub> removal efficiency and desulfurization selectivity are used as the evaluation index. The desulfurization selectivity is defined as follows,

$$\text{Desulfurization selectivity} = \frac{[SO_2]_l}{[CO_2]_l} \quad (2)$$

[SO<sub>2</sub>]<sub>l</sub>: SO<sub>2</sub> concentration of rich solution

[CO<sub>2</sub>]<sub>l</sub>: CO<sub>2</sub> concentration of rich solution



**Fig. 2 The experimental apparatus of removing SO<sub>2</sub>**

1 a gas mixer with static agitation unit 2.saturation flask 3.absorption SO<sub>2</sub> flask 4.absorption tower 5.thermometer 6.heater 7. flowmeter 8. flowmeter 9.pump 10.pump 11.regenerative tower 12. absorption flask after desorption SO<sub>2</sub> 13. flowmeter 14.filter 15. air compressor

### 3 Results and discussion

#### 3.1 The selectively absorption experiment of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> and NO<sub>x</sub> in the presence of CO<sub>2</sub>

##### 3.1.1 The effect of Mn (II)+DMSO on the removal efficiency of SO<sub>2</sub>

Experiment condition: temperature 303.15 K, gas flow: 80ml/min, inlet SO<sub>2</sub> concentration: 0.1792%. The effect of Mn (II)+DMSO absorbent on the removal efficiency of SO<sub>2</sub> is listed in Table 1, Mn (II) concentration is 0.03 mol/L.

**Table 1 The effect of absorbent of Mn (II)+DMSO on the removal efficiency of SO<sub>2</sub>**

No	inlet SO <sub>2</sub> concentration, %	outlet SO <sub>2</sub> concentration, %	SO <sub>2</sub> removal efficiency, %
1	0.1806	0.0034	98.12
2	0.1670	0.0024	98.56
3	0.2608	0.0095	96.36

From Table 1, it can be seen that Mn (II)+DMSO absorbent have high the removal efficiency for SO<sub>2</sub>, and the removal efficiency of SO<sub>2</sub> increases with the decrease of inlet SO<sub>2</sub> concentration.

##### 3.1.2 The selectively absorption effect of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> in the presence of CO<sub>2</sub>

Experiment condition: temperature 301.15 K, gas flow: 140 mL·min<sup>-1</sup>, inlet SO<sub>2</sub> concentration: 0.1893 % SO<sub>2</sub>, inlet CO<sub>2</sub> concentration: 11.3 % CO<sub>2</sub>. The result is listed in Table 2


**Table 2 Selectivity absorption effect of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> in the presence of CO<sub>2</sub>**

t, min	5		10		15	
	outlet concentration, 10 <sup>-6</sup>	removal efficiency, %	outlet concentration, 10 <sup>-6</sup> ,	removal efficiency, %	outlet concentration, 10 <sup>-6</sup> ,	removal efficiency, %
SO <sub>2</sub>	2	99.89	5	99.44	5	99.44
CO <sub>2</sub>	8.9	21.23	9.2	18.58	9.4	16.81

t, min	20		25		30	
	outlet concentration, 10 <sup>-6</sup> ,	removal efficiency, %	outlet concentration, 10 <sup>-6</sup>	removal efficiency, %	outlet concentration, 10 <sup>-6</sup>	removal efficiency, %
SO <sub>2</sub>	7	99.22	8	99.10	12	98.66
CO <sub>2</sub>	9.6	15.04	9.8	13.27	10.2	9.73

From Table 2, it can be shown that the removal efficiencies of DMSO+Mn (II) absorbent for SO<sub>2</sub> are over 98% in the presence of CO<sub>2</sub> within the absorption time of 0 to 30 min, but only 9-21% removal efficiency for CO<sub>2</sub>, it reveals that DMSO+Mn (II) absorbents have a good selective removal for SO<sub>2</sub> in the presence of CO<sub>2</sub>.

Meanwhile another experiment is also carried out to examine the absorption of SO<sub>2</sub> in the presence of CO<sub>2</sub> and SO<sub>2</sub> with pure DMSO and the similar results are obtained.

### 3.1.3 The effect of absorbent of Mn (II)+DMSO on the removal efficiency of NO<sub>x</sub>

The effect of absorbent of Mn (II)+DMSO on the removal efficiency of NO<sub>x</sub> is listed in Table 3.

**Table 3 The effect of absorbent of Mn (II)+DMSO on the removal efficiency of NO<sub>x</sub>**

	inlet NO concentration, 10 <sup>-6</sup>	outlet NO concentration, 10 <sup>-6</sup>	NO removal efficiency, %	inlet NO <sub>2</sub> concentration, 10 <sup>-6</sup>	outlet NO <sub>2</sub> concentration, 10 <sup>-6</sup>	NO <sub>2</sub> removal efficiency, %
DMSO	185	16	91.35	59	8	86.44
DMSO+0.03MMnSO	480	32	93.33	60	6	90.00

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The results indicate that absorbent of Mn (II)+DMSO has high the removal efficiency for NO<sub>x</sub>, and the addition of Mn (II) in DMSO can increase removal efficiency of NO<sub>x</sub>.

### 3.1.4 The selectively absorption effect of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> and NO<sub>x</sub> in the presence of CO<sub>2</sub>

Experiment condition: temperature 301.15 K, gas flow: 130 mL·min<sup>-1</sup>. The results are listed in Table 4.



**Table 4 The selectivity absorption effect of Mn (II) + DMSO on removal efficiency of SO<sub>2</sub> and NO<sub>x</sub> in the presence of CO<sub>2</sub>**

t, min	inlet	outlet	removal	inlet	outlet	removal	inlet	outlet	removal efficiency, %
	concentration, 10 <sup>6</sup>	concentration, 10 <sup>6</sup>	efficiency, %	concentration, 10 <sup>6</sup>	concentration, 10 <sup>6</sup>	efficiency, %	concentration, 10 <sup>6</sup>	concentration, 10 <sup>6</sup>	
SO <sub>2</sub>	718	17	97.63	675	14	97.92	1274	21	98.35
CO <sub>2</sub>	12%	9.5%	20.83	10.1%	8.5%	15.84	9.7%	7.8%	19.58
NO <sub>2</sub>	85	8	90.59	70	6	91.42	50	5	90.00
NO	403	38	91.00	445	43	90.33	436	40	90.82

The results indicate that the removal efficiencies of DMSO+Mn (II) absorbent for SO<sub>2</sub> are over 97% in the presence of CO<sub>2</sub>, and for NO<sub>x</sub> are over 90% in the presence of CO<sub>2</sub>.

### 3.2 Result of the enlarged experiment

#### 3.2.1 The effect of ratio of gas/liquid on the desulfurization selectivity

The suitable gas /liquid (G/L) ratio is the key factors influencing removal efficiency of SO<sub>2</sub>. The effect of ratio of G/L on the selectivity removal efficiency of SO<sub>2</sub> is listed in Table 5. The results showed the suitable gas /liquid ratio is 1:40-1:500.

**Table 5. The experimental result of removing SO<sub>2</sub> by Mn (II) + DMSO**

No.	rate of flow/m <sup>3</sup> .h <sup>-1</sup>	ratio of G/L	temper ature /°C	absorption solution concentration, mol/kg		inlet gas composition, %			outlet gas composition, %		removal efficiency, %		selectivity
				SO <sub>2</sub>	CO <sub>2</sub>	SO	CO	O <sub>2</sub>	SO <sub>2</sub>	CO <sub>2</sub>	SO <sub>2</sub>	CO <sub>2</sub>	
						2	2						
1	0.04	40	25	0.0485	0.000910	0.45	9.8	2.3	0.0010	5.0	99.78	48.98	53.27
2	0.1	100	25	0.0610	0.001100	0.45	9.8	2.3	0.0060	5.2	99.67	46.94	55.45
3	0.2	200	25	0.1295	0.001300	0.45	9.8	2.3	0.0194	5.8	95.69	40.82	99.46
4	0.25	250	25	0.1325	0.001310	0.45	9.8	2.3	0.0274	7.0	93.9	28.57	100.92
5	0.04	40	30	0.0379	0.000801	0.45	9.8	2.3	0.0015	5.2	99.67	46.94	47.31
6	0.1	100	30	0.0551	0.000900	0.45	9.8	2.3	0.0072	5.5	98.4	43.37	61.20
7	0.2	200	30	0.1295	0.001000	0.45	9.8	2.3	0.0205	6.0	95.56	38.78	128.60



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8	0.25	250	30	0.130	0.00110	0.4	9.8	2.3	0.041	7.2	90.88	26.53	118.45
						5							
9	0.04	40	40.5	0.025	0.00060	0.4	9.8	2.3	0.001	5.4	99.64	46.94	41.67
				0	0	5			6				
10	0.1	100	40.5	0.038	0.00070	0.4	9.8	2.3	0.024	5.8	94.67	40.82	54.29
				0	0	5							
11	0.2	200	40.5	0.108	0.00089	0.4	9.8	2.3	0.032	7.2	92.71	26.53	120.11
					9	5			8				
12	0.04	40	25	0.041	0.00070	0.4	7.4	2.2	0.000	3.8	99.86	48.65	59.09
				4	1	2			6				
13	0.1	100	25	0.054	0.00090	0.4	7.4	2.2	0.002	4.2	99.53	43.24	60.00
				0	0	2			0				
14	0.2	200	25	0.086	0.00110	0.4	7.4	2.2	0.004	5.0	99.03	32.43	78.55
				4		2			1				
15	0.04	40	30	0.035	0.00060	0.4	7.4	2.2	0.000	3.8	99.83	48.65	58.42
				1	1	2			7				
16	0.1	100	30	0.051	0.00080	0.4	7.4	2.2	0.002	4.8	99.46	33.33	63.78
				0	0	2			3				
17	0.2	200	30	0.110	0.00090	0.4	7.4	2.2	0.004	5.2	98.91	27.78	122.33
					0	2			6				
18	0.04	40	40	0.021	0.00060	0.4	7.4	2.2	0.001	4.3	99.55	40.28	35.00
				0	0	2			9				
19	0.1	100	40	0.035	0.00080	0.4	7.4	2.2	0.006	5.1	98.46	29.17	43.75
				0	0	2			5				
20	0.2	200	40	0.089	0.00090	0.4	7.4	2.2	0.010	6.3	97.63	12.50	98.89
				0	0	2							
21	0.04	40	25	0.038	0.00110	0.4	19.	2.0	0.000	11.6	99.90	38.95	35.00
				5		1	0		4				
22	0.1	100	25	0.051	0.00130	0.4	19.	2.0	0.002	15.1	99.41	20.53	39.23
				0		1	0		4				
23	0.2	200	25	0.079	0.00160	0.4	19.	2.0	0.003	16.3	99.14	14.21	49.50
				2		1	0		5				
24	0.1	100	30	0.030	0.00070	0.4	19.	2.0	0.000	12.0	99.85	58.33	43.03
				1	0	1	0		6				
25	0.2	200	30	0.048	0.00100	0.4	19.	2.0	0.002	14.6	99.34	23.16	48.50
				5		1	0		7				

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26	0.3	300	30	0.098 0	0.00140	0.4 1	19. 0	2.0	0.024 3	16.0	94.04	15.79	70.00
27	0.1	100	40	0.020 0	0.00050 0	0.4 1	19. 0	2.0	0.001 6	15.5	99.61	18.42	40.00
28	0.2	200	40	0.036 4	0.00080 0	0.4 1	19. 0	2.0	0.006 0	15.9	98.53	16.32	45.50
29	0.3	300	40	0.079 0	0.00100	0.4 1	19. 0	2.0	0.038 3	16.6	90.61	12.63	79.00

From Table 5, it can be found that removal efficiency of SO<sub>2</sub> is influenced by gas /liquid ratio. If liquid flow is kept constant, the alteration of gas /liquid ratio has very little influence on removal efficiency of SO<sub>2</sub>, but the effect on the removal efficiency of CO<sub>2</sub> is obvious.

### 3.2.2 The effect of rate of gas flow on the desulfurization selectivity

The effect of rate of gas flow on the selectivity removal efficiency of SO<sub>2</sub> was studied with four rate of gas flow between 0.04 m<sup>3</sup>/h -0.25 m<sup>3</sup>/h, and the result is listed in Table 5. From Table 5, it indicates that the removal efficiency of CO<sub>2</sub> is lower; the selectivity is higher with the increase of the gas flow rate. Therefore, when the flue gas has high content of CO<sub>2</sub>, we can decrease the absorption for CO<sub>2</sub> and increase the absorption of SO<sub>2</sub> by using the increase of the gas rate.

### 3.2.3 The effect of temperature on the desulfurization selectivity

The effect of temperature on the selectivity removal efficiency of SO<sub>2</sub> is listed in Table 5. The results indicate that the removal efficiency of SO<sub>2</sub> is high with the decrease of temperature; therefore high removal efficiency of SO<sub>2</sub> is obtained when SO<sub>2</sub> absorption is kept at suitable low temperature.

### 3.2.4 The effect of inlet gas composition on the desulfurization selectivity

Three different gases, such as SO<sub>2</sub>, CO<sub>2</sub> and O<sub>2</sub> was mixed in the process of the experiments. Inlet SO<sub>2</sub> concentration is about 0.4 %, O<sub>2</sub> concentration is 2% or so, the inlet CO<sub>2</sub> concentration is the change, the effect on the selectivity removal efficiency of SO<sub>2</sub> in the presence of CO<sub>2</sub> is studied. The result is listed in Table 5. From Table 5, it can be found that the removal efficiency of SO<sub>2</sub> of DMSO+ Mn (II) absorbents are over 90% when the concentration of CO<sub>2</sub> changed, therefore the selectivity for SO<sub>2</sub> removal does not decrease only increasing CO<sub>2</sub> concentration. i.e. absorbents still have high purification efficiencies for SO<sub>2</sub> in the presence of high content of CO<sub>2</sub>.

### 3.2.5 The regeneration capacities of Mn (II) + DMSO

The solution of being absorbed was desorbed at room temperature at the rate of 100 ml/min, the desorption result of Mn (II) + DMSO is listed in Table 6 , it showed that the absorbents of Mn (II) + DMSO have good desorption efficiency , and the desorption efficiency are over 97% at room temperature.

**Table 6 The desorption result of Mn (II) + DMSO**

SO <sub>2</sub> concentration before desorption, mol/kg	desorption temperature, °C	rate of flow, ml/min	SO <sub>2</sub> concentration after desorption, mol/kg	desorption efficiency , %
0.1081	25	100	0.0025	97.67
0.1101	25	100	0.0028	97.46



### 3.3. Mechanism Analysis

The absorption process for  $\text{SO}_2$  containing  $\text{DMSO} + \text{Mn(II)}$  is a complicated mechanism coexisting of physical and chemical absorptions.

#### 3.3.1. Physical absorption

$\text{DMSO}$  is a polar aprotic organic solvent having a larger dipole moment (4.03 D). Its molecular structure is shown in Figure 3.  $\text{SO}_2$  and  $\text{NO}_2$  are polar gas molecules of V-type molecular configuration and their molecular structure are shown in Figure 3. However  $\text{CO}_2$  is a symmetrical linear non-polar molecule, and its molecular configuration is shown in Figure 3.

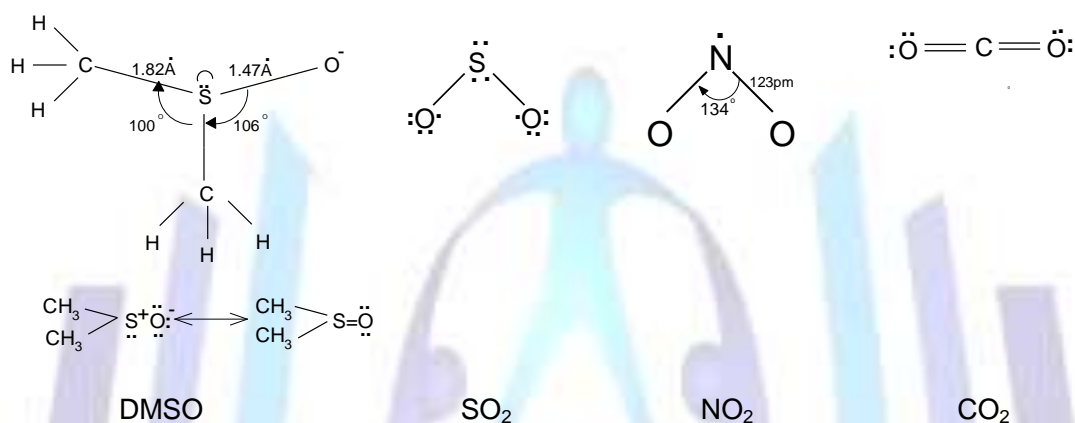


Fig. 3. Molecular structure for  $\text{DMSO}$ ,  $\text{SO}_2$ ,  $\text{NO}_2$  and  $\text{CO}_2$

$\text{DMSO}$  is a polar organic solvent with a rather large dipole matrix,  $\text{SO}_2$  and  $\text{NO}_2$  are also polar gas molecules, while  $\text{CO}_2$  is a non-polar gas molecule, therefore  $\text{DMSO}$  has a better absorption capacity on the polarity of the sulfur dioxide and nitrogen oxides, but less absorption capacity on the non-polar carbon dioxide. It agrees with the principle that like dissolves like.

Accordingly, the absorption model of  $\text{SO}_2$  in  $\text{DMSO}$  is shown in Figure 4 [9].

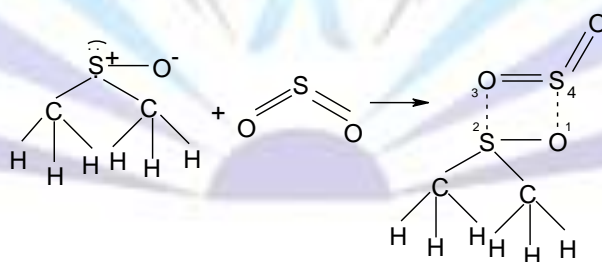
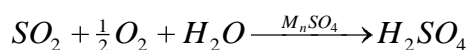


Fig. 4. The absorption model of  $\text{SO}_2$  in  $\text{DMSO}$

#### 3.3.2. Chemical absorption

After adding  $\text{Mn(II)}$  in  $\text{DMSO}$ ,  $\text{SO}_2$  is oxidized to  $\text{H}_2\text{SO}_4$  catalyzed by  $\text{MnSO}_4$ .

The chemical reaction is expressed as follows [10, 11, 12]:





The mixed absorbent has a good removal efficiency of SO<sub>2</sub> and a good regeneration. Therefore the selected absorbent has a good future in SO<sub>2</sub> removal.

#### 4 Conclusions

The desulfurization selectivity experiment of DMSO + Mn (II) is studied. The result shows that the desulfurization absorbent has not only high desulfurization efficiency, but also good selectivity for SO<sub>2</sub> and NO<sub>x</sub> in the presence of CO<sub>2</sub>, the feasibility of desulfurization absorbent has been proved.

The effect of gas/liquid ratio, flow rate, temperature and inlet gas composition on the removal efficiency was determined, and the absorption and desorption experiment was investigated. The results indicated when liquid flow is constant, the alteration of gas /liquid ratio has very little influence on removal efficiency of SO<sub>2</sub>, but the effect on the removal efficiency of CO<sub>2</sub> is obvious. The desulfurization absorbent still keep high desulfurization selectivity even if in high CO<sub>2</sub> concentration. The removal efficiency of SO<sub>2</sub> increases with the decrease of temperature; when the inlet gas composition changed, especially CO<sub>2</sub> concentrations vary from 7-19 %, there is no obvious influence on the removal efficiency of SO<sub>2</sub>, it shows that the desulfurization absorbents still keep high desulfurization efficiencies in the presence of high content of CO<sub>2</sub>.

The result provided a valuable reference for the industrial application of removal of SO<sub>2</sub> from flue gas.

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