Effects of water vapor, CO$_2$ and SO$_2$ on the NO reduction by NH$_3$ over sulfated CaO

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Abstract—Gas effects on NO reduction by NH$_3$ over sulfated CaO have been investigated in the presence of O$_2$ at 700-850°C. CO$_2$ and SO$_2$ have reversible negative effects on the catalytic activity of sulfated CaO. Although H$_2$O alone has no obvious effect, it can depress the negative effects of CO$_2$ and SO$_2$. In the flue gas with CO$_2$, SO$_2$, and H$_2$O co-existing, the sulfated CaO still catalyzed the NO reduction by NH$_3$. The in situ DRTFTS of H$_2$O adsorption over sulfated CaO indicated that H$_2$O generated Bronsted acid sites at high temperature, suggesting that CO$_2$ and SO$_2$ competed for only the molecularly adsorbed NH$_3$ over Lewis acid sites with NO, without influencing the ammonia ions adsorbed over Bronsted acid sites. Lewis acid sites shifting to Bronsted acid sites by H$_2$O adsorption at high temperature may explain the depression of the negative effect on NO reduction by CO$_2$ and SO$_2$.

Key words: Sulfated CaO, NO, CO$_2$, SO$_2$, H$_2$O

INTRODUCTION

The nitric oxide (NO) reduction by ammonia (NH$_3$) over sulfated CaO has been investigated extensively because of its potential application in the simultaneous removal of SO$_2$ and NO$_x$ in circulating fluidized bed by flue gas desulphurization technique (CFB-FGD) [1-3]. Previous pilot-scale CFB-FGD results have indicated that the SO$_2$ removal efficiency could be as high as 85-95% at the Ca/S molar ratio of 2 in the temperature range of 700-800°C [4]. For NO$_x$ removal, the previous investigation indicated that sulfated CaO has a catalytic effect on NO reduction by NH$_3$ [1-2,5] in NH$_3$+NO+O$_2$+N$_2$ atmosphere. However, little investigation has been done on the effects of flue gas components, such as CO$_2$, SO$_2$, and H$_2$O, which may have significant influences on the catalyst activity [6-10]. Previous study has already found that CaO has high catalytic effect on NH$_3$ oxidation to NO [11-13], but greatly suppressed by H$_2$O [14-16] and SO$_2$ [5,17]. CO$_2$ can also affect the catalytic activity of CaO in NH$_3$ oxidation [14,18]. The effects of the major gas components on the catalytic activity of sulfated CaO in NO reduction are of great interest. For a catalyst to be used in the post-combustion process, it must have satisfactory activity in the presence of high concentration of CO$_2$ and H$_2$O, and sufficient resistance to the poisonous effect of SO$_2$. For the coal-fired electrical utility plant, the typical flue gas might contain 150-1,000 ppm NO, 5% O$_2$, 13% CO$_2$, 8% H$_2$O, and 200-2,000 ppm SO$_2$ [19]. Hence, 14% CO$_2$, 8% H$_2$O, and 2,000 ppm SO$_2$ were selected in this study to investigate the flue gas effect. In situ analytical techniques, such as in situ diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS), were used to determine the intermediates over the catalyst surface to analyze the flue gas affecting mechanisms. Previous study has indicated that the surface-species over sulfated CaO produce acidic sites and favor ammonia adsorption [2]. The acidic gas of SO$_2$ and CO$_2$ may not directly affect the surface acid sites, whilst the adsorbed H$_2$O over sulfates can generate Bronsted acid sites [20] or modifying the site structure [21-23]. Therefore, the influence of water vapor, SO$_2$, and CO$_2$ on the catalytic activity of sulfated CaO were comprehensively investigated in this study to understand their affecting mechanism under different atmospheres.

EXPERIMENTAL SECTION

1. Activity Measurement

The effect of flue gas species on the catalytic activity of sulfated CaO (prepared by CaO sulfation in 2,000 ppm SO$_2$+5%O$_2$+N$_2$ at 850°C, CaO sulfation extent (X$_{Ca}$) was 36.5%) has been tested in a quartz fixed-bed reactor system, as shown in Fig. 1. The quartz fixed-bed reactor system consists of a quartz tube with a reactor section (8 mm in diameter and 200 mm in length), a temperature controller (TC), and a flow control system (MFC). The gas mixture containing NH$_3$, CO$_2$, SO$_2$, and H$_2$O was supplied to the reactor through a mass flow controller (MFC). The exhaust gas was analyzed using an FTIR spectrometer (FTIR spectrometer).

Fig. 1. Schematic of the reaction system.

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bed reactor is the main body of the reaction system, with an inner diameter of 20 mm. The reactor is placed in an electrically heated furnace capable of maintaining the reaction zone at a constant temperature. The bed material is placed on a sintered porous quartz disc located in the middle section of inner reactor. Blank tests without bed materials were conducted at 800 and 850 °C to determine the effect of SNCR (selective non-catalytic reduction). The results indicated that SNCR contributed less than 4% of NO reduction at 800 °C and about 30% of NO reduction at 850 °C.

The CaO sulfation extent (X_{Ca}) was calculated from the mass increase of the solid sample defined as (1):

$$X_{Ca} = \frac{n_C}{n_{CaO}}\times 100$$  \hspace{1cm} (1)

where \(n_C\) was the molar of sulfur element in the sulfated CaO, and \(n_{CaO}\) was the molar of calcium element in the sulfated CaO. The catalytic activity of the sulfated CaO in the presence/absence of H\(_2\)O, SO\(_2\), and CO\(_2\) was mainly measured over 4.0 g sulfated CaO. The inlet flow gas was carefully controlled by mass flow controllers to obtain the desired gas concentration and flow rate. The total flow rate was set at 1,000 ml/min (273.15 K, standard atmospheric pressure), then the gas velocity was 83,000 h\(^{-1}\) at 800 °C, and 87,000 h\(^{-1}\) at 850 °C over the 4.0 g sulfated CaO. The gas mixture at inlet was 500 ppm NO+500 ppm NH\(_3\)+5%\(\text{O}_2\)+N\(_2\) (as balanced gas), and 2,000 ppm SO\(_2\), 14% CO\(_2\), 8% H\(_2\)O would be added when their effects were investigated. The concentration of H\(_2\)O vapor was controlled by regulating the flow rate of the N\(_2\) through a 55 °C water bath to prevent the water vapor condensation and the potential secondary effects were investigated. The concentration of H\(_2\)O vapor was continuously monitored by a pre-calibrated FTIR spectrometer (Nicolet Corporation, NEXUS 670) aided by a liquid N\(_2\)-cooled MCT detector and a 2 m gas cell, collecting 32 scans at a resolution of 0.5 cm\(^{-1}\). The gas cell temperature was maintained at 150 °C. The pipeline between the reactor and the gas analyzer was heated to 90 °C to prevent the water vapor condensation and the potential secondary reactions in gas mixtures. The measurement accuracy has been estimated to be ±2% for a single gas, and ±2% (NO), ±3% (NH\(_3\)), ±3% (SO\(_2\)) for simultaneous monitoring of multiple gas species, as tested by Li et al. [24]. During the catalyst activity test, the NO conversion (X_{NO}) and NH\(_3\) conversion (X_{NH3}) were calculated from their inlet and outlet concentration. These parameters were defined as Eqs. (2)-(3):

$$X_{NO} = \left(1 - \frac{C_{NO}}{C_{NO}^{in}}\right)\times 100$$  \hspace{1cm} (2)

$$X_{NH3} = \left(1 - \frac{C_{NH3}}{C_{NH3}^{in}}\right)\times 100$$  \hspace{1cm} (3)

where \(C_{NO}^{in}\) was the NO concentration at the inlet of the quartz reactor, \(C_{NO}\) was the NO concentration at the outlet of the quartz reactor; \(C_{NH3}^{in}\), was the NH\(_3\) concentration at the inlet of the quartz reactor, and \(C_{NH3}\) was the NH\(_3\) concentration at the outlet of the quartz reactor.

2. In Situ DRIFTS Study

In situ DRIFTS was used to study the H\(_2\)O adsorption over sulfated CaO in this study. The in situ DRIFTS spectra were obtained using an FTIR (Thermo Nicolet Corporation, NEXUS670) equipped with an MCT detector and diffuse reflectance kit (HARRICK) including the Praying Mantis and a high temperature reaction chamber. The chamber was enclosed with a dome opened three windows, two ZnSe windows for spectrometer radiation, and one UV quartz window for sample inspection. The chamber was connected to the gas flow apparatus which passed gas over the solid samples, and to the water cooling system which controlled the chamber and window temperatures during high temperature operation. The spectra were obtained with the collecting parameter of 64 scans at a resolution of 8 cm\(^{-1}\). Before H\(_2\)O adsorption, the catalysts were activated in N\(_2\) under 700 °C for 30 min to remove impurities in the catalysts and then cooled within the chamber with N\(_2\) purge to the desired experimental temperature 100 °C. During the cooling process of 700 °C to 100 °C, the spectra of the sample were collected at 500, 400, 300, 200 and 100 °C as the background spectra of corresponding temperature, which could be used later to generate H\(_2\)O adsorption spectra at different temperatures. For H\(_2\)O adsorption, the 100 °C was chosen to avoid water condensation in the reaction chamber; also, the connecting pipeline was heated to 100 °C.

**RESULTS AND DISCUSSION**

1. Effect of H\(_2\)O, SO\(_2\), and CO\(_2\) on SCR Activity

Fig. 2 shows NO conversion over 1.0 g sulfated CaO (particle diameter, 0.30-0.45 mm) in 1,000 ppm NH\(_3\)+1,000 ppm NO+5%\(\text{O}_2\) with various SO\(_2\) concentration at 800 °C. At first, the NO conversion was 15.1% without SO\(_2\) in the inlet flue gas. Then the experiment were done with 2,000, 1,500, 1,000, 500, 250, 100 and 50 ppm SO\(_2\) added successively, and the NO conversion was 4%, 4.4%, 4.8%, 5.5%, 6.5%, 7.4% and 8%, accordingly, which indicated the obvious restraining effect of SO\(_2\) even when only 50 ppm SO\(_2\) was added. Finally, the SO\(_2\) was switched off, and the NO conversion increased to 13.5% immediately, which verified the restraining effect of SO\(_2\) again. However, the little difference between the NO conversion of 13.5% (at last, without SO\(_2\)) and 15.1% (at first, without SO\(_2\)) was caused by the long term introducing of SO\(_2\) over CaO sulfation products making the sulfation extent change, and the reason that the NO conversion decreased as the CaO sulfation increased.

![Fig. 2. The effect of SO\(_2\) with different concentration on NO conversion at 800 °C.](image-url)
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after a certain CaO conversion has been described and analyzed in detail in our previous work [1,2]. Besides, the increase of the NO conversion to 13.5% at last time when SO₂ was 0 ppm also verified the gas restraining effect of SO₂ on the catalytic activity when SO₂ was added during the experiments, without considering the CaO sulfation effect.

Fig. 3 shows the NO conversion over 1.0 g sulfated CaO (particle diameter, 0.30-0.45 mm) in the atmosphere of 500 ppm NH₃+500 ppm NO+5% O₂+N₂ with or without 2,000 ppm SO₂ at 800°C. The results revealed that the NO outlet concentration increased when 2,000 ppm SO₂ was added into the main gas stream (during the time of 20-37 and 72-87 min), which again indicated the restraining effect of SO₂ on the NO reduction by NH₃. However, when SO₂ was cut off after the 2,000 ppm (during the time of 38-64 min and 88-105 min), the NO outlet concentration recovered to about 440 ppm as appeared during 0-19 min. Since the introducing of SO₂ was not for a long time, and the CaO sulfation extent almost did not change, then the sulfation products catalytic activity for deNOx reaction did not change after SO₂ removal, which indicated that the gas effect of SO₂ was reversible for a certain CaO sulfation product.

The combined effects of H₂O, SO₂ and CO₂ on the SCR reaction were investigated at 800°C (under which temperature the SNCR effect could be neglected) and 850°C (under which temperature the SNCR effect was apparent) to see the effects of gas composition more obviously. In these tests, the sulfated CaO sample was 4.0 g.

The NO and NH₃ conversions of the reaction Sequence 1-5 over 4.0 g sulfated CaO at 850°C are shown in Fig. 4. In Sequence 1, there were just NO, NH₃ and O₂, and the conversions of NO and NH₃ were 47% and 85.2%, respectively. When 14% CO₂, 8% H₂O and 2,000 ppm SO₂ were all presented in Sequence 2, the NO conversion decreased to 20.8%, while the NH₃ conversion decreased to 30%, indicating the overall effect of H₂O, SO₂ and CO₂ was to suppress the NO and NH₃ conversion, and consequently, the catalyst activity was reduced. In Sequence 3, only H₂O was added upon Sequence 1; the NO conversion was 48.2%, and NH₃ conversion was 59.8%. The NO conversion in Sequence 3 was almost the same with that in sequence 1, while the NH₃ conversion in Sequence 3 was lower than that in Sequence 1, which indicated that H₂O did not decrease the catalytic activity of sulfated CaO, but enhanced the selectivity of NH₃ for NO reduction. The combined effect of H₂O and SO₂ on the SCR reaction is shown in Sequence 4, and the individual effect of SO₂ is shown in Sequence 5. Comparing the NO and NH₃ conversions in Sequences 4 and 5, it could be found

Fig. 4. NO, NH₃ conversion over 4.0 g sulfated CaO at different atmosphere at 850°C (The inlet gas composition: Sequence 1, base gas: 500 ppm NO+500 ppm NH₃+5% O₂+N₂; Sequence 2, base gas+14% CO₂+2,000 ppm SO₂+8% H₂O; Sequence 3, base gas+8% H₂O; Sequence 4, base gas+8% H₂O+2,000 ppm SO₂; Sequence 5, base gas+2,000 ppm SO₂).

Fig. 5. NO, NH₃ conversion over 4.0 g sulfated CaO at different atmosphere at 800°C (The inlet gas composition: Sequence a, base gas: 500 ppm NO+500 ppm NH₃+5% O₂+N₂; Sequence b, base gas+14% CO₂+2,000 ppm SO₂+8% H₂O; Sequence c, base gas+8% H₂O+2,000 ppm SO₂; Sequence d, base gas+2,000 ppm SO₂; Sequence e, base gas+8% H₂O+2,000 ppm SO₂).
that SO2 had a strong inhibiting effect on NO conversion with the NO conversion decreased from 47% (in Sequence 1) to 5.4% (in Sequence 5), and H2O could depress the inhibiting effect of SO2 with the NO conversion increased from 5.4% (in Sequence 5) to 28.4% (in Sequence 4). The results in Sequence 2 and 4 suggested that CO2 also had a negative effect on the SCR reaction. The results in Sequence 2 and 3 indicated that, when SO2 and CO2 were switched off, their negative effects disappeared, implying that the effects of CO2 and SO2 were both reversible.

The NO and NH3 conversions of the reaction Sequence a-e over 4.0 g sulfated CaO at 800°C are shown in Fig. 5. In Sequence a, NO conversion was 18.6% and NH3 conversion was 24.4%. In Sequence b, CO2, H2O and SO2 were simultaneously added, and the NO conversion decreased to 10.6% from 18.6%, while NH3 conversion decreased from 24.4% to 13%, which agreed with the results of Sequence 1 and 2 at 850°C in Fig. 4. Upon Sequence b, CO2 was switched off in Sequence c, the NO conversion and NH3 conversion both increased, which confirmed the negative effect of CO2 on the catalyst activity. Upon Sequence c, H2O was switched off and only SO2 was added in Sequence d, and then the NO conversion significantly decreased. When H2O was added again in Sequence c, NO conversion recovered to that in Sequence c. The results in Sequence c-e once again verified that H2O depressed the negative effect of SO2 on the catalytic activity.

The flue gas effect investigation at 800 and 850°C both indicated that CO2 and SO2 had a negative effect on the catalytic activity, and their negative effect was reversible. H2O can depress the negative effect of SO2 and the catalyst still promotes NO reduction when CO2, SO2 and H2O coexisted.

2. DRIFTS of H2O Adsorption

To understand the improving mechanism of H2O, H2O adsorption over sulfated CaO was investigated. Fig. 6 shows the DRIFTS of H2O adsorption with increasing temperature over sulfated CaO after passing N2 with 8% H2O over the sample at 100°C. Many H2O adsorption bands exist and the distinctive positive broad peak at 2,600-3,700 cm\(^{-1}\) has been assigned as the hydroxyl group from H2O [25]. As the temperature increased from 100 to 400°C, the peaks of the hydroxyl group from H2O disappeared gradually, leaving two obvious peaks of freedom OH stretching band at 3,633 cm\(^{-1}\) and associated OH bending band at 1,475 cm\(^{-1}\) [26]. When the temperature increased to 500°C, the obvious peak of the associated OH at 1,475 cm\(^{-1}\) still existed, and the OH associated to sulfate would be good for ammonia ions forming during the ammonia adsorption, which indicated the Brønsted acidity formed by water adsorption at high temperature over the active sites of sulfated CaO.

To verify the effect of the H2O adsorption over the sulfated CaO catalyst, the ‘wet’ sulfated CaO was also prepared. The sulfated CaO was first put into a fixed bed reactor, passing N2 with 8% H2O through it at 650°C for 30 min, and then removed from the reactor and cool to the room temperature (RT) for DRIFTS experiments. The spectra of sulfated CaO (wet) shown in Fig. 7 were collected using the spectra of the sulfated CaO without H2O adsorption at RT as the background. The peaks around 1,430 and 1,500 cm\(^{-1}\) were the peaks of associated OH over sulfates [26], indicating the Brønsted acid sites formed at high temperature. The other curve in Fig. 7 is the spectra of the sample heated up to 500°C after H2O adsorption with the peak at 1,475 cm\(^{-1}\). Comparing the two curves in Fig. 7, the peaks of 1,430, 1,475 and 1,500 cm\(^{-1}\) could all be the associated OH bands, presenting Bronsted acid sites formed at high temperature, and the shifts between them could be caused by the temperature difference. Hence, it once again indicates that the H2O adsorption over sulfated CaO enhances Bronsted acidity at high temperature.

3. Influencing Mechanisms of Gas Species

The investigation of flue gas effect on the catalytic activity of sulfated CaO have indicated that CO2 and SO2 have a reversible negative effect on the catalytic activity, whereas H2O can depress the negative effect from SO2. Previous research has indicated that the SCR reaction over sulfated CaO could be the adsorbed NH3 reacting with gaseous NO to form N2 [2]. For NH3 adsorption, previous study on V2O5-based catalysts and other metal oxide catalysts indicated that there are two kinds of ammonia adsorption patterns: (i) molecularly adsorbed ammonia through Lewis-type interaction on

![Fig. 6. DRIFTS of H2O adsorption over sulfated CaO at different temperatures (a) at 100°C; (b) at 200°C; (c) at 300°C; (d) at 400°C; (e) at 500°C.](image-url)
suggested that H2O molecules could provide OH group to adsorb on Brønsted acidity at high temperature. Yang et al. [31] indicated that CO2 and SO2 may not influence the NH3 adsorbed over Brønsted acid sites over sulfated CaO; hence, it could be deduced that H2O can depress the inhibiting effect of CO2 and SO2 on the catalytic activity of sulfated CaO in Fig. 6 and 7 indicated that H2O adsorption forms Brønsted acidity at high temperature. Yang et al. [31] suggested that H2O molecules could provide OH group to adsorb ammonia and form ammonium ions required in SCR reaction. Furthermore, some investigations on H2O adsorption over V2O5-based catalyst also indicated the formation of Brønsted acidity in the presence of H2O [32-34]. Over sulfated CaO, the coordinate modes of SO42- have been presented in previous study [2]. Combining the coordinate structure of SO42- with the Brønsted acid sites generated from H2O adsorption over vanadyl centers [29], the generation schemes of Brønsted acid sites over sulfated CaO are proposed in Fig. 8.

As the inhibiting effect of CO2 and SO2 on the catalytic activity of sulfated CaO was reversible, the affecting mechanism was inferred to be their competition for active sites. CO2 and SO2 are acidic gases; they may compete for the adsorbed NH3 with NO over sulfated CaO, which would cause the decrease of NO reduction rate and NO conversion. However, the addition of H2O can depress the inhibiting effect of SO2. DRIFTS results have indicated that H2O can generate Brønsted acid sites over sulfated CaO; hence, it could be deduced that CO2 and SO2 may not influence the NH3 adsorbed over Brønsted acid sites, and just influence the molecularly adsorbed NH3 over Lewis acid sites. When H2O changed some Lewis acid sites to Brønsted acid sites and suppresses the inhibiting effect of CO2 and SO2.

CONCLUSIONS

The experimental results and discussions show that:

1. CO2 and SO2 inhibit the catalytic activity of sulfated CaO for NO reduction by NH3 in the presence of O2, whereas H2O alone has no obvious effect on the catalytic activity of the sulfated CaO. However, H2O can depress the inhibiting effect of CO2 and SO2 on the catalytic activity of sulfated CaO. Sulfated CaO can still catalyze NO reduction by NH3 when CO2, SO2, H2O, and O2 coexist in flue gas, which indicates the feasibility of using sulfated CaO as a NO removal catalyst in post-combustion process;

2. In situ DRIFTS experiments of H2O adsorption over sulfated CaO indicate that H2O generates Brønsted acid sites over sulfated CaO at high temperature;

3. Combining the activity tests with and without CO2, SO2, and H2O, the gas influencing mechanism was analyzed. CO2 and SO2 affect the catalytic activity of sulfated CaO by competing for the adsorbed NH3 over Lewis acid sites against NOx, and therefore decrease the NO reduction rate. CO2 and SO2 do not affect the ammonia ions over Brønsted acid sites. The addition of H2O converts Lewis acid sites to Brønsted acid sites and suppresses the inhibiting effect of CO2 and SO2.

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