Molten Salt Synthesis of Iron Oxide Modified Attapulgite for Catalytic Oxidation of Elemental Mercury

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(Received on 13th July 2012, accepted in revised form 12th November 2012)

Summary: Iron oxide modified attapulgite (Fe$_2$O$_3$-ATP) catalysts, with special network structure, was prepared by molten salt method, and was applied to investigate Hg$^0$ oxidation ability in simulated flue gas from 100 to 260°C. Scanning electron microscopy (SEM), X-ray diffraction (XRD) and Fourier-transform infrared (FTIR) spectroscopy were employed to characterize the catalysts. The results showed that Fe$_2$O$_3$-ATP exhibited about 90% average Hg$^0$ oxidation efficiency at 220°C. With the decrease of the Fe$_2$O$_3$ in Fe$_2$O$_3$-ATP, the Hg$^0$ oxidation efficiency reduced. Furthermore, the flue gas components seriously affected the Hg$^0$ oxidation. The presence of Cl$_2$ and O$_2$ could promote the Hg$^0$ oxidation, while SO$_2$ would inhibit Hg$^0$ oxidation. Cl$_2$ was the most effective component of the three for Hg$^0$ oxidation.

Keywords: Fe$_2$O$_3$-ATP, Hg$^0$, SEM, XRD, FTIR.

Introduction

The pollution of trace metal, especially mercury, in coal-fired flue gas has aroused wide concern. Due to its high toxicity, strong volatility, and lasting stability, mercury is poisonous to both human and animals. There exists three forms of mercury in coal-fired flue gas: elemental(Hg$^0$), oxidized(Hg$^{2+}$), and particulate(Hg(p)) [1]. Among them, Hg(p) and Hg$^{2+}$ are easier to be removed than Hg$^0$. Hg(p) can be captured by existing air pollution control devices such as electrostatic precipitators (ESP) and bag dust collector. Hg$^{2+}$ can be removed by wet flue gas desulfurization (FGD) due to its water solubility. In contrast, Hg$^0$ can’t be removed by FGD because it is almost insoluble in water. Consequently, if Hg$^0$ could be oxidized to water-soluble Hg$^{2+}$ by catalytic oxidation of catalysts, it will have important significance for the control of mercury emission in coal-fired flue gas.

Currently, several metal oxides such as Fe$_2$O$_3$ [2], CeO$_2$ [3], Mn$_2$O$_3$ [4], CuO [5] and V$_2$O$_5$ [6] have been demonstrated to have oxidation activities toward Hg$^0$, especially at relative low temperatures. Among the above catalysts, Fe$_2$O$_3$ has the advantages of abundance, strong toxicity resistance and inexpensive. However, the pure Fe$_2$O$_3$ catalysts are harmful to the environment, and its catalytic activity is not high. By using attapulgite as carrier, Fe$_2$O$_3$ is uniformly dispersed on the surface of attapulgite for preventing agglomeration, and enhancing catalyst activity.

Attapulgite (ATP) is a natural structural hydrated magnesium aluminum silicate with 1-D fibrillar morphology [7], which has large surface area and zeolite-like channels, and can be used as absorbent, catalyst and catalyst support [8-11]. At present, the methods of metal modified ATP mostly focus on impregnation, ion exchange, co-precipitation and sol-gel. These methods are complex and limited. Molten salt synthesis is one of the simplest ways to prepare oxides or complex oxides. When the temperature is above the melting point of the salts, the liquid phase is formed to be used as reaction medium or promoter [13-15]. This method enables shorter reaction time and lower formation time because of the high diffusivities of the components, moreover it can control the particle morphology and stabilize different polymorphs [14].

In this work, the netting-shaped iron oxide modified attapulgite (Fe$_2$O$_3$-ATP) catalysts were synthesized by molten salt method. The study focused on the preparation of catalysts, the effects of Fe$_2$O$_3$/ATP ratio, reaction temperature and individual flue gas components for Hg$^0$ oxidation efficiency.

Results and Discussion

Catalysts Characteristics

Fig. 1 presents the SEM images of ATP (calcined at 500°C), Fe$_2$O$_3$(molten salt method) and
Fe₂O₃-ATP(2:1) respectively. As shown in Fig. 1a and 1b, ATP showed the long and narrow fiber-shape [16], and Fe₂O₃ displayed the disordered sheeted structure. However, after synthesizing Fe₂O₃-ATP(2:1) by molten salt method, the Fe₂O₃ sheets aggregated together orderly forming netting(Fig. 1c, 1d), and no special fiber structure of ATP existed, which might be due to Fe₂O₃ was widely dispersed on the ATP surface. It’s propitious to increase the active sites on catalysts for mercury oxidation.

Fig. 1: SEM images of ATP (calcined at 500°C), Fe₂O₃ (molten salt method) and Fe₂O₃-ATP(2:1).

The XRD patterns of ATP(calcined at 500°C), Fe₂O₃-ATP(1:1), and Fe₂O₃-ATP(2:1) are shown in Fig. 2. The typical diffraction peaks at 2θ=19.96°, 27.58° were corresponding to the primary diffraction of the (040) and (400) planes for ATP [17,18], which could be detected in the X-ray diffraction pattern of ATP and Fe₂O₃-ATP(1:1). However, these characteristic peaks of ATP weren’t found over Fe₂O₃-ATP(2:1), which indicated that Fe₂O₃ was highly dispersed on the surface of ATP. The peaks at 2θ=33.14°, 35.60°, 49.40° and 54.26° ascribed to the specific peaks of Fe₂O₃, which could also demonstrate that Fe₂O₃ had dispersed on the ATP surface for both Fe₂O₃-ATP(1:1) and Fe₂O₃-ATP(2:1).

Fig. 2: XRD patterns of ATP(calcined at 500°C), Fe₂O₃-ATP(1:1) and Fe₂O₃-ATP(2:1).

Fig. 3 shows the FT-IR results of ATP(calcined at 500°C), Fe₂O₃, Fe₂O₃-ATP (1:1) and Fe₂O₃-ATP (2:1) composite material. It was observed that the FT-IR spectra of Fe₂O₃-ATP (1:1) and Fe₂O₃-ATP (2:1) are almost the same. The absorbance bands at 463.74cm⁻¹ and 573.16cm⁻¹ ascribed to the symmetry and asymmetry stretching vibration of Fe-O [19], which could be detected in the curves of Fe₂O₃, Fe₂O₃-ATP(1:1) and Fe₂O₃-ATP (2:1). After adding ATP, a new band at 1029.07cm⁻¹ of Si-O characteristic peak for ATP emerged in the FT-IR spectrum of Fe₂O₃-ATP (1:1) and Fe₂O₃-ATP (2:1). The result indicated that Fe₂O₃ had successfully dispersed on the ATP surface. The result was in compliance with the result of Fig. 1 and Fig. 2.

Fig. 3: FT-IR spectra of ATP(calcined at 500°C), Fe₂O₃, Fe₂O₃-ATP(1:1) and Fe₂O₃-ATP(2:1).
The wide and strong absorbance bands at 3424.38 cm\(^{-1}\) and 1632.73 cm\(^{-1}\) were corresponding to the stretching and bending vibrations of the absorbed water.

**Catalytic Activity**

Hg\(^0\) oxidation efficiency over ATP (calcined at 500°C), Fe\(_2\)O\(_3\) (molten salt method), Fe\(_2\)O\(_3\)-ATP(1:1) and Fe\(_2\)O\(_3\)-ATP(2:1) at various temperatures is displayed in Fig. 4. It was observed that the Hg\(^0\) oxidation efficiency improved remarkably for Fe\(_2\)O\(_3\)-ATP (2:1), which presented approximate 90% oxidation efficiency at 220°C. Nevertheless, with the decrease of Fe\(_2\)O\(_3\), the activity of Fe\(_2\)O\(_3\)-ATP(1:1) for Hg\(^0\) oxidation obviously reduced. In this process, the adsorption of Hg\(^0\) by ATP played the leading role, because ATP wasn’t completely coated by Fe\(_2\)O\(_3\). Almost no obvious Hg\(^0\) oxidation was observed over pure ATP, the Hg\(^0\) oxidation efficiency was less than 5%. However, the Hg\(^0\) removal efficiency of ATP could reach to 10% through adsorption mainly. The adding of Fe\(_2\)O\(_3\) provided more surface chemisorbed oxygen, and the special network structure supplied more active sites for Hg\(^0\) oxidation. For all catalysts above, the Hg\(^0\) oxidation efficiency enhanced when temperature was from 100 to 220°C, and with the continued increase of the temperature to 260°C, the Hg\(^0\) oxidation efficiency decreased.

**Effect of O\(_2\), Cl\(_2\) and SO\(_2\) on Hg\(^0\) Oxidation Efficiency**

To better understand the Hg\(^0\) oxidation capacity of Fe\(_2\)O\(_3\)-ATP(2:1), it’s necessary to discuss the effect of individual flue gas components on Hg\(^0\) oxidation efficiency, especially O\(_2\), Cl\(_2\) and SO\(_2\). The experiments were examined at 220°C.

The average Hg\(^0\) oxidation efficiency of different O\(_2\) content (0, 6%, 12%) in simulated flue gas is shown in Fig. 5. With the increase of O\(_2\), Hg\(^0\) oxidation efficiency rose slowly. Under the absence of O\(_2\), Hg\(^0\) oxidation efficiency was about 24.8%. When the concentration of O\(_2\) was up to 6%, no obvious increase was observed, and only about 29.7% of Hg\(^0\) was oxidized even if the concentration is increased to 12%. The results revealed that the Fe\(_2\)O\(_3\)-ATP(2:1) possessed large oxygen storage capacity and consumed only a small part of the stored oxygen in mercury oxidation [3]. Cl\(_2\) has promotional effect for Hg\(^0\) oxidation due to HgCl\(_2\) was the main oxidized mercury species in coal combustion flue gas [20]. As described in Fig. 6, the impacts of Cl\(_2\) were researched in the range of 0-50 ppm. Comparing with O\(_2\), Hg\(^0\) oxidation ability in the present of Cl\(_2\) enhanced largely. When the concentration of Cl\(_2\) was transferred from 0 to 50 ppm, the Hg\(^0\) oxidation efficiency of Fe\(_2\)O\(_3\)-ATP(2:1) was enhanced from 32.5% to 93.2%. Whereas, as the Cl\(_2\) continued to increase, no evident increase of Hg\(^0\) oxidation efficiency was detected. Impeditive effect of SO\(_2\) on Hg\(^0\) oxidation over Fe\(_2\)O\(_3\)-ATP(2:1) was summarized in Fig. 7. When SO\(_2\)(400, 800, 1600 ppm) was added into the simulated flue gas in the absence of O\(_2\), almost no Hg\(^0\) oxidation was observed. Hailong Li et al [3] studied the competition of Hg\(^0\) and SO\(_2\) for the active sites on catalysts, and the result illustrated that the affinity between SO\(_2\) and catalyst was stronger than that between Hg\(^0\) and catalyst. Therefore, the presence of SO\(_2\) restrained the Hg\(^0\) oxidation capacity of Fe\(_2\)O\(_3\)-ATP(2:1).

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**Fig. 4:** Effect of the temperature on Hg\(^0\) oxidation.

**Fig. 5:** Effect of O\(_2\) on Hg\(^0\) oxidation.

**Fig. 6:** Effect of Cl\(_2\) on Hg\(^0\) oxidation.
Experimental

Catalysts Preparation

Attapulgite (ATP) was supplied by Jiangsu Dongda Heating and Mechanical Manufacture Company (China). Hydrochloric acid (HCl), silver nitrate (AgNO₃), ferric nitrate (Fe(NO₃)₃·9H₂O), sodium chloride (NaCl) and absolute ethyl alcohol (EtOH) used in this study were analytical grade.

ATP was firstly treated with 37% hydrochloric acid at 110°C for 2 h, followed by washing with distilled water until no Cl⁻ detected (tested with AgNO₃), and then dried at 80°C for 8 h before use.

The molten salt method was used to synthesize Fe₂O₃-ATP catalysts. 0.01 mol Fe(NO₃)₃·9H₂O, 0.1 mol NaCl and the purified ATP (the mass ratio of ATP/Fe(NO₃)₃·9H₂O was 1:1 and 1:2, respectively) were mixed, and ground homogenously. The mixtures were heated to a target temperature of 500°C for 8 h, and then air-cooled. The products was thoroughly washed with 50% aqueous ethanol (v/v) at room temperature, and then dried overnight in air at 80°C.

Characterization of Catalysts

The catalysts were characterized by X-ray diffraction (DX-2700 diffractometer), using Cu Kα as radiation over the 2θ range of 10°~70°. SEM images were obtained by using QUANTA200 scanning electron microscope (FEI Company) to observe the morphology of catalysts. FTIR spectra of the samples were taken with NEXUS-670 FT-IR spectroscopy (NICOLET) in the range of 400-4000 cm⁻¹.

Activity Tests

The apparatus used in this work is illustrated in Fig. 8. The catalytic oxidation activities of Hg⁰ over Fe₂O₃-ATP were carried out in a fixed-bed flow reactor system under a total flow of 1 L/min and a gas hourly space velocity (GHSV) of about 37700 h⁻¹. In each experiment, 0.5 g of the catalyst was placed in the fixed-bed reactor (13 mm inner diameter) with temperature controlling from 100 to 260°C. The simulated flue gas was composed of 6% O₂, 12% CO₂, 300 ppm NO, 1500 ppm SO₂, 50 ppm Cl₂, and balance gas N₂. The N₂ flow was distributed into two branches: one branch (100 ml/min) passed through a Hg⁰ permeation tube (QMG-6-6) to introduce the saturated Hg⁰ vapor (20 µg/m³) into the simulated flue gas system, and the other branch converged with the individual streams of NO, CO₂, SO₂ and O₂, and formed the main gas flow. The gases were adjusted by mass flow controllers, and mixed, pre-heated before entering the reactor.

The Hg⁰ concentrations at both the inlet and outlet of reactor were measured by the mercury analyzer (QM201H Hg⁰ analyzer). After analysis, the exhaust gas must pass through the 0.1 M KMnO₄ solution before being introduced into the air. However, the Hg⁰ analyzer can only measure the concentration of Hg⁰. Therefore, in order to measure the concentration of total mercury including Hg⁰ and oxidized Hg⁰, the SnCl₂ solution (the oxidized Hg⁰ could be reduced to Hg⁰ via the reducing action of SnCl₂ solution) was located between the fixed-bed reactor and the Hg⁰ analyzer. The Hg⁰ oxidation efficiency (η) was quantified by the difference of Hg⁰ concentration after and before passing the SnCl₂ solution. The efficiency of Hg⁰ oxidation over Fe₂O₃-ATP catalysts can be evaluated as follows:

\[ \eta_{\text{Hg oxid}(\%) \} = \frac{\text{[Hg]}_{\text{total}} - \text{[Hg⁰]}}{\text{[Hg]}_{\text{total}}} \times 100 \]

where \( \eta_{\text{Hg oxid}(\%)} \) is the Hg⁰ oxidation efficiency, \([\text{Hg]}_{\text{total}}\) is the concentration of total mercury after passing the SnCl₂ solution, and \([\text{Hg⁰}]\) is the concentration of Hg⁰ before passing the SnCl₂ solution.
Conclusion

In this work, the Hg\(^0\) oxidation efficiency of Fe\(_2\)O\(_3\)-ATP catalysts were studied in a fixed-bed flow reactor system at 100-260°C. As a result, Fe\(_2\)O\(_3\)-ATP with special netting structure were effective catalysts for Hg\(^0\) oxidation, the Hg\(^0\) oxidation efficiency of Fe\(_2\)O\(_3\)-ATP(2:1) arrived about 90% at 220°C. The decrease of Fe\(_2\)O\(_3\) would lessen the Hg\(^0\) oxidation efficiency of Fe\(_2\)O\(_3\)-ATP. In addition, the flue gas components have significant effects to Hg\(^0\) oxidation. Both O\(_2\) and Cl\(_2\) promoted Hg\(^0\) oxidation ability of Fe\(_2\)O\(_3\)-ATP(2:1), especially Cl\(_2\). When 50ppm Cl\(_2\) was added, the Hg\(^0\) oxidation efficiency reached 93.2%. However, SO\(_2\) went against Hg\(^0\) oxidation, almost no Hg\(^0\) oxidation was discovered in the presence of SO\(_2\) alone. In summary, Fe\(_2\)O\(_3\)-ATP were feasible catalysts for Hg\(^0\) oxidation in flue gas.

Acknowledgements

This work was supported financially by the Open Project of Key Laboratory for Advanced Technology in Environmental Protection of Jiangsu Province(AE201026, AE201165), the Innovation Fund Project of Yancheng Institute Of Technology(YKB201106),

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