Deep oxidative desulfurization of model fuels catalysed by immobilized ionic liquid on MIL-100(Fe)

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Deep desulfurization of fossil fuels has become urgently required because of the serious pollution by the large-scale use of fossil fuels. In this study, [PrSO₃HMIm]HSO₄@MIL-100(Fe) was synthesized by wet-impregnation of the ionic liquid (IL) of [PrSO₃HMIm]HSO₄ on MIL-100(Fe). The construction of [PrSO₃HMIm]HSO₄@MIL-100(Fe) was then confirmed by X-ray powder diffraction, N₂ adsorption-desorption experiments, infrared spectroscopy and elemental analysis, and then applied in the oxidative desulfurization of model fuels. In comparison with the corresponding IL, [PrSO₃HMIm]HSO₄@MIL-100(Fe) showed an enhanced performance in the desulfurization rate of model fuels due to the increase of the mass transfer rate. Under the optimized conditions (oxidant to sulphur ratio = 25, oil to acetonitrile ratio = 1, and temperature = 60 °C), a sulphur removal rate of 99.3% was observed (initial sulphur concentration = 50 ppm). The sulphur removal of three sulphur compounds by catalytic oxidation and extraction followed the order of dibenzothiophene (DBT) > thiophene (T) > benzothiophene (BT).

1 Introduction

The rapid development of the automotive industry has greatly exacerbated the use of fossil fuels. After combustion, the organic compounds in gasoline are converted to carbon oxides (COx), nitrogen oxides (NOx) and sulphur oxides (SOx), among which, sulphur oxides are one of the important reasons for haze and acid rain. Sulphide emissions from burning gasoline can cause harm to the human lungs and respiratory system, resulting in throat spasms and suffocation. Currently, the sulphur content in gasoline in China must be from 50 μg g⁻¹ to 10 μg g⁻¹ according to the National standard of gasoline for motor vehicles (GB17930-2013 and GB17930-2016). Above all, it is necessary to reduce the sulphur content of gasoline and reduce the emission of sulphur oxides into the atmosphere.

The oxidative desulfurization (ODS) technology has many advantages such as mild reaction conditions, high desulfurization rate, simple process and low equipment investment, compared with traditional hydrodesulfurization methods. ODS has been widely used for the development of ultra-low sulphur products. The catalysts used for oxidative desulfurization are very critical. In recent years, IL catalysts has been used in the oxidative desulfurization reaction in gasoline. However, although these IL catalysts are efficient in desulfurization, the ILs cannot be directly separated and recovered because they are easily dissolved in extracting agents, such as acetonitrile, which is not conducive to industrialization. In this study, a new catalyst, [PrSO₃HMIm]HSO₄@MIL-100, was synthesized by wet impregnation. The effect of several important factors, including the loading amount, reaction temperature, desulfurization system, catalyst dosage, and the recycling of catalyst, on the desulfurization efficiency was studied.

2 Experimental

2.1 Materials

Iron(III) chloride hexahydrate (FeCl₃·6H₂O) and ethanol were obtained from Damao Chemical Reagent Factory. Acetonitrile, n-octane, 1,3,5-benzenetricarboxylic acid (H₃BTC), thiophene (T), benzothiophen (BT) and dibenzothiodiphen (DBT) were purchased from J&K Scientific Ltd. 1-Sulfopropyl-3-methylimidazolium hydrosulphate ([PrSO₃HMIm]HSO₄) was obtained from Lanzhou Greenchem ILs. All solvents and reagents were used as received without further purification.

2.2 Synthesis of MIL-100(Fe)

MIL-100(Fe) was synthesized according to the previous report under the following methods. A mixture of FeCl₃·6H₂O (0.010 mol, 2.70 g) and BTC (0.066 mol, 1.39 g) was dispersed in 50 mL distilled water under weak sonication and was then reacted for 72 h at 130 °C in a Teflon lined autoclave. After being cooled down to room temperature, the light orange solid product was further purified by centrifugation at 6000 rpm for 10 min with ethanol (50 mL × 3). Then the highly purified solid was dried under vacuum at 60 °C overnight.
2.3 Synthesis of [PrSO₃HMIm]HSO₄@MIL-100(Fe)

[PrSO₃HMIm]HSO₄@MIL-100(Fe) was prepared by wet impregnation. A certain amount of anhydrous [PrSO₃HMIm]HSO₄ was dissolved in 50 mL ethanol. After complete dissolution, 15 g activated MIL-100(Fe) was added to the solution and subjected to continuous magnetic stirring for 24 h. Then, filtration was employed to remove the redundant solvent, followed by drying in vacuum at 80 °C overnight. The orange powder product was donated as X-[PrSO₃HMIm]HSO₄@MIL-100 (X mmol g⁻¹ [PrSO₃HMIm]HSO₄@MIL-100), where X means the adding amount of [PrSO₃HMIm]HSO₄ in the solution (X = 0.7, 0.9, 1.1, 1.3, 1.5).

2.4 Desulfurization

Dibenzothiophene (0.05 g, DBT) was completely dissolved into n-octane to prepare model oil with the sulphur content of 50 ppm. Similarly, the model oils with different sulphur compounds, including thiophene and benzothiophenen, was prepared. The catalyst, acetonitrile and 10 mL model oil were added into a flask. Then the flask was in an oil bath at the desired temperature and stirred for 15 min. H₂O₂ was injected into the flask. After the start of the reaction, the stirring was stopped at regular intervals, the samples from the upper oil phase were withdrawn and filtered with an organic filter, then sulphur content in the samples were determined by the WK-2D micro-coulometer (Jiangfen Electroanalytical instrument Co., Ltd, China, detection limit: 0.1 ng mL⁻¹). And the conversion rate of the sulphur was calculated according to eqn (1):

\[
R = \frac{C_0 - C_e}{C_0} \times 100\% \tag{1}
\]

where, R is the desulfurization rate, C₀ is the initial sulphur content of the oil, and Cₑ is the sulphur content of the oil after desulfurization.

2.5 Characterization

X-Ray Diffraction (XRD) patterns were collected with a Bruker D8 ADVANCE diffractometer using Cu Kα radiation at a scanning rate of 1° min⁻¹ between 5° and 40°. Infrared spectra were recorded in KBr discs with a Bruker Fourier-transform infrared (FT-IR) spectrometer. The specific surface areas and pore size distributions were obtained from N₂ adsorption-desorption isotherms measured on a Micromeritics ASAP-2010 instrument at −196 °C. The contents of C, H, N, and S in the composite material were measured using Elementar's Vario EL III elemental analyzer.

3 Results and discussion

3.1 X-ray diffraction (XRD) analysis

The powder XRD patterns of the virgin MOFs and a series of ILs@MOF are exhibited in Fig. 1. These diffraction patterns agreed with those reported in the literature. The peak at 10.9° indicated that the pattern corresponded to the reported MIL-100(Fe). The XRD patterns of the ILs@MIL-100(Fe) materials were similar to that of the parent material MIL-100(Fe), which indicated that the crystal structure of MIL-100(Fe) was preserved after the incorporation of [PrSO₃HMIm]HSO₄, and adding IL did not change the crystal form of MIL-100.

3.2 Fourier transform infrared (FT-IR)

The FT-IR spectra of MIL-100(Fe), X-[PrSO₃HMIm]HSO₄@MIL-100(Fe) and [PrSO₃HMIm]HSO₄ were recorded in the spectral range of 4000–400 cm⁻¹ (Fig. 2). The peaks of [PrSO₃HMIm]HSO₄ at 1224 and 1046 cm⁻¹ were related to ν(O=O) asymmetric and symmetric stretching vibrations of the sulfonic acid group. The characteristic peaks of MIL-100(Fe) at 1714 and 1383 cm⁻¹ attributed to ν(C=O) stretching vibration and ν(C=O) stretching vibration in the MIL-100(Fe) matrix could be observed. The characteristic vibrational bands of the C=C and C=N group at 1573 and 1457 cm⁻¹ were also observed corresponding to the imidazole ring. It was concluded that [PrSO₃HMIm]HSO₄ was successfully supported on MIL-100(Fe).

3.3 Elemental analysis

The S and N elements of the X-[PrSO₃HMIm]HSO₄@MIL-100(Fe) (X = 0.7, 0.9, 1.1, 1.3, 1.5) composites were analysed to calculate the amount of IL in the hybrid materials. It could be seen from the Table 1 that the S and N elements in the hybrid material showed an increasing trend with the increase of the
adding amount of IL, indicating that more IL groups were
grafted into the pores of MIL-100(Fe) when more IL was added.
The actual loading of [PrSO₃HMIm][HSO₄] could be calculated by
the percentage content of the S elements in the hybrid material.
According to the results, it can be seen from Table 1 that there
were some certain differences between the actual loading and
the theoretical one, because a part of ionic liquids on the
surface of the materials were washed away during the process of
synthesizing the composite materials.

3.4 N₂ adsorption–desorption experiment

Fig. 3 show N₂ adsorption–desorption isotherms of MIL-100(Fe)
and X-[PrSO₃HMIm][HSO₄]@MIL-100(Fe). The BET surface area
and pore volume were obtained from nitrogen physisorption
isotherms at 77 K, and the results are given in Table 2.
It could be seen that the N₂ adsorption–desorption
isotherms of the series of composites showed a type IV
isotherms. There were also hysteresis loops, which were charac-
teristic of mesoporous or microporous materials. In addition,
the pore sizes of the samples were greater than 2 nm, indicating
that they were both mesoporous materials, and the structure
of MIL-100(Fe) were not significantly altered by loading the ionic
liquids by the impregnation method. The BET surface area and
pore volume for the prepared MIL-100(Fe) was 1168.1 m² g⁻¹
and 0.74 cm³ g⁻¹, respectively, close to the values reported in
the literature.²⁸,²⁹ According to the results, the specific surface
area and pore volume decreased after the impregnation process.

Table 1  Elemental analysis of MIL-100(Fe) and X-ILs@MIL-100(Fe)

| Samples                | S (wt%) | N (wt%) | Actual loading (mmol g⁻¹) |
|------------------------|---------|---------|--------------------------|
| MIL-100(Fe)            | —       | 1.019   | —                        |
| 0.7-ILs@MIL-100(Fe)   | 3.515   | 1.516   | 0.549                    |
| 0.9-ILs@MIL-100(Fe)   | 3.990   | 1.642   | 0.623                    |
| 1.1-ILs@MIL-100(Fe)   | 4.474   | 1.800   | 0.699                    |
| 1.3-ILs@MIL-100(Fe)   | 4.921   | 1.940   | 0.769                    |
| 1.5-ILs@MIL-100(Fe)   | 5.492   | 2.173   | 0.858                    |

The decrease could be related to the pore blocking of the
support by the attachment of ILs.³⁵

3.5 Catalytic activities for ODS

3.5.1 Effects of time and loading amount on oxidative
desulfurization. The ILs loading amount had a significant effect
on the reaction. It was evident from Fig. 4 that ODS was cata-
ysed by various catalysts. The desulfurization efficiency of the
[PrSO₃HMIm][HSO₄]@MIL-100(Fe) systems was higher than that
of the virgin MIL-100(Fe) and [PrSO₃HMIm][HSO₄] under the
same catalysis conditions. The activity was improved by
increasing the contents of [PrSO₃HMIm][HSO₄] in the [PrSO₃-
HMIm][HSO₄]@MIL-100(Fe) system from 0.7 to 1.1 mmol g⁻¹,
and the sulphur removal ratio of 1.1-[PrSO₃HMIm][HSO₄]@MIL-
100(Fe) reached 99.31%, showing a beneficial effect of [PrSO₃-
HMIm][HSO₄] during the catalysis. However, the sulphur
removal of 1.3-ILs@MIL-100(Fe) and 1.5-ILs@MIL-100(Fe) was
lower than that of 1.1-ILs@MIL-100(Fe). We speculated that the
increased acid ion loading resulted in a reduced MIL-100(Fe)
pore volume and reaction specific surface area, which reduced
the mass transfer efficiency.³⁶,³⁷ Thus, the loading amount of
1.1 mmol g⁻¹ and the optimal time of 180 min were proper for
the following investigation, respectively.

Table 2  Structural properties of MIL-100(Fe) and X-ILs@MIL-100(Fe)

| Samples                | BET surface area (m² g⁻¹) | Pore volume (cm³ g⁻¹) |
|------------------------|--------------------------|----------------------|
| MIL-100(Fe)            | 1168.1                   | 0.74                 |
| 0.7-ILs@MIL-100(Fe)   | 785.6                    | 0.47                 |
| 0.9-ILs@MIL-100(Fe)   | 674.5                    | 0.42                 |
| 1.1-ILs@MIL-100(Fe)   | 525.7                    | 0.33                 |
| 1.3-ILs@MIL-100(Fe)   | 418.2                    | 0.26                 |
| 1.5-ILs@MIL-100(Fe)   | 170.0                    | 0.13                 |

Fig. 3  N₂ adsorption–desorption isotherms for MIL-100(Fe) (1) and X-
[PrSO₃HMIm][HSO₄]@MIL-100(Fe) (2–6) (X = 0.7, 0.9, 1.1, 1.3, 1.5).

Fig. 4  Effects of reaction time and loading amount of IL on oxidative
desulfurization. (1) [PrSO₃HMIm][HSO₄], (2) MIL-100(Fe), (3–7) X-ILs@-
MIL-100(Fe) (X = 0.7, 0.9, 1.1, 1.3, 1.5), the following conditions were
used: catalyst, 0.05 g; model oil, 10 mL; acetonitrile, 10 mL; H₂O₂, 28
µL; t, 180 min; T, 60 °C.
3.5.2 Effects of catalyst dosage on oxidative desulfurization. The effects of catalyst dosage on the DBT oxidation were studied systematically. As shown in Fig. 5, the DBT removal was enhanced dramatically with the increasing catalyst usage from 0.03 g to 0.05 g, which can be attributed to the increase of the total amount of the catalytic active sites. Apparently, 0.05 g of catalyst had offered enough catalytic active sites to the conversion of DBT. However, the removal of DBT was decreased with the increasing catalyst usage from 0.05 g to 0.06 g. It may be that more catalysts would promote self-decomposition of H₂O₂, generating oxygen and water, which was not only detrimental to the progress of the oxidation reaction, but also cause waste of the catalysts. So, the optimal catalyst dosage was 0.05 g.

3.5.3 Effects of the H₂O₂/DBT molar ratios on oxidative desulfurization. To investigate the influence of the oxidant dosage, the experiment was carried out under various H₂O₂/DBT (O/S) molar ratios at 60 °C. As shown in Fig. 6, the removal of DBT increased with the increase of the H₂O₂ dosage (O/S ≤ 25), which can be attributed to the increasing production quantity of the catalytic activity species. When O/S = 25, the sulphur removal was 99.31%. However, when a higher amount of H₂O₂ was added, the sulphur removal decreased. On the one hand, this confirmed that water had a negative effect on the desulfurization as the amount of introduced water increased with the increasing amount of hydrogen peroxide. On the other hand, excess H₂O₂ enhanced the hydrogen bond between the catalyst particles in the acetonitrile phase, leading to transfer of the agglomerated catalyst into the oil phase, and thus its less availability for ODS which consequently resulted in lower desulfurization rate.

3.5.4 Removal of different sulphur substrates. To investigate the effects of [PrSO₃HMIm]HSO₄@MIL-100(Fe) on different sulphur compounds, ODS of DBT, BT and T were carried out under the same conditions. The removal rates of DBT, BT and T were shown in Fig. 7. The desulfurization reactivity followed the order: DBT > T > BT. The reactivity should follow the order of the electron density because the removal of sulphur substrates is mainly affected by the electron density on the sulphur atom. The unusual lowest reactivity of BT can be attributed to the steric hindrance. These results suggested that the reactivity order was affected by the simultaneous effects of the electron density on the sulphur atom and the steric hindrance of the sulphur compounds in this ODS system. Similar results were present in Lv’s studies, they found that sulphur removal selectivity also followed the order of DBT > BT in their ionic liquids extraction coupled with oxidation desulfurization systems (ECODS), and the catalytic activity of DBT removal could reach 98% at 30 °C in 3 h.

3.5.5 Recycling of catalysts. The catalysts were separated by centrifugation and dried under vacuum at 353 K for 12 h for regeneration. The reutilization for the removal of DBT was performed maintaining the same experimental conditions for each cycle. As is shown in Fig. 8, the catalytic performance of the catalyst decreased to 70% after five cycles, while the sulphur
removal in the ECODS systems could remain above 85% after recycling six times in Lv’s studies. The rapid decrease could mainly be attributed to the partial loss of the ionic liquid during the reaction. This shows that the carrier with the proper pore structure had a great influence on the catalytic oxidative desulfurization performance.

3.5.6 Proposed mechanism of coupled oxidation extraction desulfurization. Several studies have shown that high desulfurization rate can be achieved with \( \text{H}_2\text{O}_2 \) or molecular oxygen as oxidants and different kinds of ionic liquids as catalysts or extractants. In this paper, a coupled oxidation extraction desulfurization mechanism is proposed according to these studies. As seen in Scheme 1, take DBT as an example, firstly, some kinds of radicals (mainly, \( \text{HO}_2^- \)) could be produced by \( \text{H}_2\text{O}_2 \) with the acidic [PrSO_3HMIm]HSO_4 and the Bronsted acid sites in MIL-100(Fe) as the catalysts. DBT was then oxidized to the corresponding sulfones by these radicals. Due to their high polarity, the sulfones could be easily removed from the model oil phase by a polar extractant, such as acetonitrile in this study. The sulphur-free fuel was then obtained, and the heterogeneous catalyst ([PrSO_3HMIm]HSO_4@MIL-100(Fe)) could be recycled by static settlement or centrifugation.

4 Conclusions

In summary, [PrSO_3HMIm]HSO_4 immobilized in MIL-100(Fe) was successfully synthesized through wet-impregnation. The XRD, FT-IR, EA and \( \text{N}_2 \) adsorption–desorption indicated that the ILs@MIL-100(Fe) samples remained in the mesoporous structure of MIL-100(Fe). The [PrSO_3HMIm]HSO_4@MIL-100(Fe) catalysts displayed significantly higher catalytic performance than [PrSO_3HMIm]HSO_4 for the oxidative desulfurization of DBT. In the experimental system, the effects of different ILs loading, catalyst dosage, O/S ratio, and different sulphur substrates on the desulfurization rate were investigated, and the optimum reaction conditions were determined. The 1.1-[PrSO_3HMIm]HSO_4@MIL-100(Fe) sample had the highest oxidative desulfurization efficiency, by which, the sulphur removal of DBT reached 99.3%.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

1. J. Wu, Y. Gao, W. Zhang, Y. Tan, A. Tang, Y. Men and B. Tang, Appl. Organomet. Chem., 2015, 29, 96–100.
2. D. Bokare and W. Choi, J. Hazard. Mater., 2016, 304, 313–319.
3. A. Stanislaus, A. Marafi and M. S. Rana, Catal. Today, 2010, 153, 1–68.
4. W. Jiang, W. Zhu, Y. Chang, Y. Chao, S. Yin, H. Liu, F. Zhu and H. Li, Chem. Eng. J., 2014, 250, 48–54.
5. C. Li, D. Li, S. Zou, Z. Li, J. Yin, A. Wang, Y. Cui, Z. Yao and Q. Zhao, Green Chem., 2013, 15, 2793–2799.
6. W. Ding, W. Zhu, J. Xiong, L. Yang, A. Wei, M. Zhang and H. Li, Chem. Eng. J., 2015, 266, 213–221.
7. U. Domanska and M. Wlazlo, Fuel, 2014, 134, 114–125.
8. P. S. Kulkarni and C. A. M. Afonso, Green Chem., 2010, 12, 1139–1149.
9. H. Liu, S. Bao, Z. Cai, T. Xu, N. Li, L. Wang, H. Chen, W. Lu and W. Chen, Chem. Eng. J., 2017, 317, 1092–1098.
10. Z. Klimont, S. J. Smith and J. Cofala, Environ. Res. Lett., 2013, 8, 014003.
11. B. Jiang, H. Yang, L. Zhang, R. Zhang, Y. Sun and Y. Huang, Chem. Eng. J., 2016, 283, 89–96.
12. B. Rodriguez-Cabo, H. Rodriguez, E. Rodil, A. Arce and A. Soto, Fuel, 2014, 117, 882–889.
