1 Introduction

SO_{2} emission, which is mainly produced by vehicles, not only pollutes the environment, such as acid rain\(^1\), but also harms the health of humans, such as cancer. In order to decrease the pollution caused by SO_{2}, many countries have laid down a series of rules for restricting the sulfur content of fuel oil. It is an aim to achieve little sulfur in oil (sulfur content < 10 mg g\(^{-1}\)).\(^2\) The traditional hydrodesulfurization (HDS) process, removing aliphatic and acyclic sulfur compounds, has been considered as a mature technology. However, HDS finds it difficult to remove aromatic sulfur compounds\(^3,4\) such as thiophene (TH), benzo-thiophene (BT) dibenzothiophene (DBT) and their derivatives because of their large steric hindrance. Therefore, some non-hydrodesulfurization processes, such as oxidative desulfurization (ODS),\(^5\) biodesulfurization (BDS),\(^6\) extractive desulfurization (EDS)\(^7\) and adsorption desulfurization (ADS),\(^8\) etc., have attracted wide attention.

Among these non-hydrodesulfurization processes, oxidative desulfurization (ODS) was considered as one of the most promising technologies due to its advantages,\(^9\) such as the high desulfurization rate and the mild reaction conditions. Various oxidants such as H\(_2\)O\(_2\), NO\(_2\), O\(_2\), O\(_3\), K\(_2\)FeO\(_4\), and organic peroxides have been investigated in the oxidative desulfurization system. Among these oxidants, H\(_2\)O\(_2\) is widely employed due to its high activity and low price.\(^10\)

A series of deep eutectic solvents (DESs) of ChCl/XCF\(_3\)SO\(_3\)H (X from 1.0 to 2.0) were synthesized by stirring a mixture of choline chloride (ChCl) and trifluoromethanesulfonic acid (CF\(_3\)SO\(_3\)H) at room temperature. The DESs were characterized by Fourier transform infrared (FT-IR) and \(^1\)H nuclear magnetic resonance (\(^1\)H NMR). The oxidative desulfurization of model oil was investigated using ChCl/1.5CF\(_3\)SO\(_3\)H as a catalyst and extraction agent, and H\(_2\)O\(_2\) as the oxidant. Some reaction parameters such as type of DES, molar ratio of CF\(_3\)SO\(_3\)H and ChCl in DESs, H\(_2\)O\(_2\) dose, reaction temperature, DES dose and type of sulfur compound were investigated. Under the optimum conditions, the removal rate of dibenzothiophene (DBT) and 4,6-dimethyldibenzothiophene (4,6-DMDBT) can reach up to 98.65% and 96.8%, respectively. After five recycling runs, the removal rate of DBT can still reach 97.16%.

In recent years, a Brønsted acid based ionic liquid was used in the oxidative desulfurization process. Fang \textit{et al.}\(^11\) reported the synthesis of 1-butyl-3-methyl-imidazolium trifluoroacetate acid ([C\(_4\)mim][TFA]) ionic liquids and their application in oxidative desulfurization and found that the anion of the ionic liquid has an important influence on the desulfurization rate. Lu \textit{et al.}\(^12\) synthesized 1-methylimidazolium tetrafluoroborate ([HMIm]BF\(_4\)) ionic liquids and the sulfur removal of DBT in model oil can reach 93% under optimum conditions. Gao \textit{et al.}\(^13\) synthesized an acidic ionic liquid N-butyl-N-methyl-imidazolium hydrogen sulfate ([BMIm]HSO\(_4\)), its oxidative desulfurization rate can reach 100% for DBT in model oil. The [BMIm]HSO\(_4\) ionic liquid can be recycled 5 times with only a slight reduction in activity. Zhao \textit{et al.}\(^14\) synthesized the Brønsted acid ionic liquid N-methyl-pyrrolidinium tetrafluoroborate ([Hnmp]BF\(_4\)) and used as the extractant and catalyst for desulfurization (ODS) of model oil. The experimental results indicated that 99.4% sulfur compounds in diesel fuel can be removed. Although these ionic liquids have better desulfurization activity in the oxidative desulfurization system, the raw materials of ionic liquids are expensive and the preparation process is complex.

As a new type of solvent, deep eutectic solvents have attracted great attention because of its excellent physical and chemical properties, which is friendly for environment. DESs are composed of two or more components that interact via intermolecular hydrogen bonds.\(^15\) Compared with the traditional ionic liquids, DESs have many advantages such as cheap and easily accessible raw material, a simple preparation process, and wide range of applications.\(^16\) In developmental history of DESs, ChCl-based DESs are first reported by Abbott and his co-
workers, its physical properties have been researched. Thereafter, DESs are used in various fields such as electrochemistry, organic synthesis and separation processes. DESs also are used in the desulfurization process of fuel. For example, Zhu et al. reported the synthesis of ChCl-2CH3COOH and it was used as the extractant of oxidation desulfurization process. The study found that the removal rate of DBT in model oil can up to 98.6% under UV light irradiation. Gano Z. S. et al. reported that the FeCl3-based deep eutectic solvents for the extractive desulfurization process. In this work, the novel CF3SO3H-based DESs were synthesized by stirring a mixture of ChCl and CF3SO3H at the room temperature. The removal rate of DBT from model oil was 97.25%, which are much higher than those obtained with traditional and functional ionic liquids. Our research group has synthesized phenylpropanoic acid-based DESs and was applied to the oxidation desulfurization process. In the synthesis, the acid-based DESs was synthesized by heating method. As is known to all, simpler preparation process and higher acidity of DESs are favorable for the desulfurization. Therefore, it is necessary to develop a new DES using a simple preparation method and apply it to the oxidative desulfurization process.

Trifluoromethanesulfonic acid (CF3SO3H) is one of the strongest organic acids, it is widely used in the synthesis reaction. In this work, the novel CF3SO3H-based DESs were synthesized by stirring a mixture of ChCl and CF3SO3H at the room temperature. The removal rate of DBT from model oil was investigated using CF3SO3H-based DESs as an extraction agent and catalyst, H2O2 as the oxidant. The influence of different type DESs, the molar ratio of ChCl to CF3SO3H, the oxygen to sulfur (O/S) molar ratio, the reaction temperature and the amount of DESs on the desulfurization rate was investigated. Under the optimum conditions, the removal rate of dibenzothiophene (DBT) and 4,6-dimethyldibenzothiophene (4,6-DMDDBT) can reach up to 98.65% and 96.8%, respectively. After five recycling runs, the removal rate of DBT can still reach 97.16%.

2 Experimental

2.1 Chemical regent and instrument

Dibenzothiophene (DBT, 98%), benzo thiophene (BT, 97%), thiophene (TH, 99.8%) and 4,6-dimethyldibenzothiophene (4,6-DMDDBT, 97%) were purchased from Aladdin Chemistry Co. Ltd. The choline chloride (ChCl, AR), tetrabutylammonium chloride (TBAC, AR), tetaethylammonium chloride (TEAC, AR), n-octane, carbon tetrachloride (CCl4, 99.5%), trifluoromethanesulfonic acid (CF3SO3H, 98%), and hydrogen peroxide (H2O2, 30 wt%) were purchased from Sinopharm Chemical Reagent Co. Ltd. Gas chromatography was carried out on an Agilent 7890A GC with an FID detector using a 30 m packed HP5 column. Infrared spectra of DBT, oxidation of DBT, DESs and raw materials were determined on Fourier-transform infrared spectrometer (WQF-520; Beijing Beifen Ruiyi Instrument Company, China). 1H NMR of DESs were determined by using the Mercury Plus 400 MHz spectrometer (Varian Co., Ltd. American).

2.2 Synthesis of ChCl/CF3SO3H

ChCl/CF3SO3H was synthesized by stirring a mixture of ChCl and CF3SO3H at room temperature. The specific process is as follows: ChCl was added to a 100 mL round-bottomed flask. Then, the CF3SO3H was carefully added according to a certain molar ratio under stirring conditions. The two raw materials were stirred vigorously for 1 h in order to fully release the reaction heat. After the reaction, the ChCl/CF3SO3H (X from 1.0 to 2.0) DESs homogeneous liquid was obtained. The reaction was shown in Fig. 1. TBAC/CF3SO3H and TEAC/CF3SO3H were synthesized according to the same method.

1H NMR spectra of CF3SO3H, ChCl and ChCl/CF3SO3H were obtained on hydrogen spectrum detector. Coupling constants (J1) in Hz and chemical shifts (d) are given in ppm. The multiplicities of signals in 1H NMR are given with chemical shifts (s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet), and the data are listed as follows:

- CF3SO3H for 1H NMR (500 MHz, D2O) δ 4.86 (s, 1H).
- ChCl for 1H NMR (500 MHz, D2O) δ 4.71 (d, 26H), 4.08 (m, 59H), 3.56-3.46 (m, 39H), 3.25 (s, J = 68.7 Hz, 30H), 3.18 (d, 176H), 3.03 (s, 2H).
- ChCl/CF3SO3H for 1H NMR (500 MHz, D2O) δ 4.77 (s, 2H), 3.90 (dd, J = 7.7, 5.3, 2.8 Hz, 1H), 3.35 (dd, J = 5.8, 4.2 Hz, 1H), 3.04 (s, 5H).

2.3 Desulfurization experiment

The 500 μg g⁻¹ model oil was prepared by dissolving 1.437 g DBT in 500 mL n-octane. The model oil, DESs, and 30 wt% H2O2 were added into a three-necked flask. The mixture was stirred in a water bath at a certain temperature. A small amount of the upper oil phase was removed every 20 min and determined by gas chromatography. The desulfurization rate was calculated using the following formula:

$$\text{ODS rate} = \frac{S_{\text{tot}} - S_{\text{res}}}{S_{\text{tot}}} \times 100\%$$

where $S_{\text{tot}}$ (500 μg g⁻¹) is the initial concentration of the sulfur compound in the model oil and $S_{\text{res}}$ is the residual concentration of the sulfur compound after the ODS process.

![Fig. 1 Synthetic mechanism of DESs.](source_url)
3 Results and discussion

3.1 Characterization

3.1.1 Infrared spectra analysis. The infrared spectra of ChCl, CF$_3$SO$_3$H and ChCl/1.5CF$_3$SO$_3$H were shown in Fig. 2. Compared with the peaks of ChCl and CF$_3$SO$_3$H, the stretching vibration peak of O–H at 3272 cm$^{-1}$, stretching vibration peak of C–H at 3018 cm$^{-1}$ and stretching vibration peak of S=O at 1406 cm$^{-1}$ in ChCl, and stretching vibration peak of F–C at 947 cm$^{-1}$ and bending vibration peak of F–C at 621 cm$^{-1}$ in CF$_3$SO$_3$H shifted to 3466, 3048, 1416, 960 and 644 cm$^{-1}$ in ChCl/1.5CF$_3$SO$_3$H were clearly seen (Fig. 2). The peaks happening to compare with ChCl in Fig. 3(b).

IR-FT (a) ChCl; (b) CF$_3$SO$_3$H; (c) ChCl/1.5CF$_3$SO$_3$H.

Fig. 2

1H NMR characterization. In order to further confirm the interactions between ChCl and CF$_3$SO$_3$H, 1H NMR spectra of CF$_3$SO$_3$H, ChCl and DESs were obtained in Fig. 3. The 1H NMR spectra of ChCl/1.5CF$_3$SO$_3$H in Fig. 3(c), the peak of hydrogen bond obviously became stronger and slightly shift to right compare with ChCl in Fig. 3(b). 1H NMR spectra of CF$_3$SO$_3$H demonstrate there is a peak of hydrogen bond in Fig. 3(a). The peak of ChCl labeled “4” and the peak of hydrogen bond in CF$_3$SO$_3$H disappeared and there is a new peak of hydrogen bond appeared in ChCl/1.5CF$_3$SO$_3$H. These results demonstrate that there are formation of hydrogen bond between ChCl and CF$_3$SO$_3$H.

Fig. 3

3.2 Oxidative desulfurization reaction conditions

3.2.1 Desulfurization rate of different desulfurization system. In this study, the removal rate of DBT in desulfurization system of ChCl/CF$_3$SO$_3$H, TBAC/CF$_3$SO$_3$H and TEAC/CF$_3$SO$_3$H was investigated. The results are shown in Fig. 4(a). It can be seen from Fig. 4(a), the desulfurization rate of DBT reached 90.58% for ChCl/CF$_3$SO$_3$H, 41.54% for TBAC/CF$_3$SO$_3$H and 85.38% for TEAC/CF$_3$SO$_3$H after the 180 min, respectively. The difference of desulfurization rate may be due to the difference in steric hindrance of DESs. It is well known that the steric hindrance of DESs follow the order TBAC/CF$_3$SO$_3$H > TEAC/CF$_3$SO$_3$H > ChCl/CF$_3$SO$_3$H. The order of desulfurization rate with these DESs is ChCl/CF$_3$SO$_3$H > TEAC/CF$_3$SO$_3$H > TBAC/CF$_3$SO$_3$H. The research shows that the bigger steric hindrance of long cation chain is not conducive to the oxidative desulfurization. Four cation chains of TBAC/CF$_3$SO$_3$H are the longest, therefore its steric hindrance is the biggest result in lowest desulfurization rate. Four cation chains of ChCl/CF$_3$SO$_3$H was the shortest, so the its steric hindrance is the smallest result in the highest desulfurization rate. Therefore, ChCl/CF$_3$SO$_3$H was chosen as the catalyst and extractant for the system of oxidation desulfurization.

In this study, the removal rate of DBT in desulfurization system of ChCl/CF$_3$SO$_3$H, TBAC/CF$_3$SO$_3$H and TEAC/CF$_3$SO$_3$H were synthesized using CF$_3$SO$_3$H and ChCl as raw materials. A series of ChCl/CF$_3$SO$_3$H (X from 1.0 to 2.0) with different acidity were investigated by changing the molar ratio of ChCl to CF$_3$SO$_3$H. As shown in Fig. 4(b), the desulfurization rate obviously increased from 90.58% to 98.85% when the molar ratio of CF$_3$SO$_3$H to ChCl were increased from 1 to 1.5. However, the desulfurization rate dropped from 98.85% to 87.3% when the molar ratio of CF$_3$SO$_3$H to ChCl was increased to 2. It is because that acidity of DESs is too strong when the molar ratio of CF$_3$SO$_3$H to ChCl were increased to 2. It could be attributed to the higher acidity rapidly decomposed H$_2$O$_2$ into oxygen, thereby decrease the oxidative ability of the system. The highest desulfurization rate of 98.85% can be attached when the molar ratio of ChCl to CF$_3$SO$_3$H is 1.5. Moreover, the viscosity of DESs have important influence on desulfurization, the high viscosity of DESs decrease extractive ability of DES. The high conductivity show there are a lot of ions in DESs. Therefore, high conductivity is benefit for desulfurization. As shown in Table 1, the conductivity and viscosity of DESs decrease with increasing molar ratio of CF$_3$SO$_3$H to ChCl. The consideration for influence of acidity, the conductivity and the viscosity of DESs on desulfurization, ChCl/1.5CF$_3$SO$_3$H can be selected as extractant and catalyst in the desulfurization system.

3.2.2 Influence of acidity of DESs on desulfurization rate. ChCl/XCF$_3$SO$_3$H were synthesized using CF$_3$SO$_3$H and ChCl as raw materials. A series of ChCl/XCF$_3$SO$_3$H (X from 1.0 to 2.0) with different acidity were investigated by changing the molar ratio of ChCl to CF$_3$SO$_3$H. As shown in Fig. 4(b), the desulfurization rate obviously increased from 90.58% to 98.85% when the molar ratio of CF$_3$SO$_3$H to ChCl were increased from 1 to 1.5. However, the desulfurization rate dropped from 98.85% to 87.3% when the molar ratio of CF$_3$SO$_3$H to ChCl was increased to 2. It is because that acidity of DESs is too strong when the molar ratio of CF$_3$SO$_3$H to ChCl were increased to 2. It could be attributed to the higher acidity rapidly decomposed H$_2$O$_2$ into oxygen, thereby decrease the oxidative ability of the system. The highest desulfurization rate of 98.85% can be attached when the molar ratio of ChCl to CF$_3$SO$_3$H is 1.5. Moreover, the viscosity of DESs have important influence on desulfurization, the high viscosity of DESs decrease extractive ability of DES. The high conductivity show there are a lot of ions in DESs. Therefore, high conductivity is benefit for desulfurization. As shown in Table 1, the conductivity and viscosity of DESs decrease with increasing molar ratio of CF$_3$SO$_3$H to ChCl. The consideration for influence of acidity, the conductivity and the viscosity of DESs on desulfurization, ChCl/1.5CF$_3$SO$_3$H–H$_2$O$_2$ system with various O/S molar ratio was
Fig. 4  (a) Influence of different type DESs on desulfurization rate (5 mL model oil, O/S molar ratio of 8, 1 mL DESs, 40 °C); (b) influence of different acidity DESs on desulfurization rate (5 mL model oil, O/S molar ratio of 8, 1 mL DESs, 40 °C); (c) influence of amount of H$_2$O$_2$ on desulfurization rate (5 mL model oil, 1 mL DESs, 40 °C); (d) influence of reaction temperature on desulfurization rate (5 mL model oil, O/S molar ratio of 6, 1 mL DESs); (e) influence of amount of DESs on desulfurization rate (5 mL model oil, O/S molar ratio of 6, 40 °C).
influencing oxidative desulfurization. As shown in Fig. 4(d), the reaction rate of desulfurization was obviously increased and removal rate of DBT increase from 98.27% to 98.65% when the reaction temperature increased from 30 °C to 40 °C. However, the desulfurization rate no longer increased when the reaction temperature was beyond 40 °C. It could be attributed to high temperature accelerate the decomposition of H2O2, which decrease oxidative ability in the oxidative desulfurization process. So, the optimal temperature in the oxidative desulfurization system is 40 °C.

3.2.5 Influence of volume ratio of DESs/oil on desulfurization rate. Fig. 4(e) illustrates the DESs dosage has an important influence on removal of DBT. The removal rate for DBT was increased from 98.46% to 98.65 when the volume ratio of DESs/oil increased from 0.15 to 0.2. Continue to increase the DESs/oil ratio, removal rate of DBT no longer increase. So, it can be concluded that the optimal volume ratio of DESs/oil is 0.2 in the desulfurization system.

3.2.6 Influence of different sulfur compounds on desulfurization rate. In this study, DBT in model oil could be effectively removed using ChCl/1.5CF3SO3H as the catalyst and extractant, with H2O2 as the oxidant. In order to determine the desulfurization efficiency of the ChCl/1.5CF3SO3H for other organic sulfides in fuels, such as 4,6-DMDBT, BT and TH were researched in oxidation desulfurization under identical conditions. The results are shown in Fig. 5. The removal rate of BT, DBT, 4,6-DMDBT and TH were 39.53%, 98.65%, 98.6% and 24.35% after 180 min, respectively. The removal rate of four sulfur compound in the desulfurization system followed the order DBT > 4,6-DMDBT > BT > TH. The desulfurization rate is related to the electron density of the sulfur atoms in organic sulfide. The electron cloud densities of these compounds are 5.758 for DBT, 5.760 for 4,6-DMDBT, 5.739 for BT, and 5.696 for TH, respectively. The electron cloud densities of 4,6-DMDBT and DBT are very approximation. However, removal rate for DBT was higher than 4,6-DMDBT due to the steric hindrance of the methyl substituent in 4,6-DMDBT. Therefore, the removal rate of 4,6-DMDBT is a little lower than DBT.

According to the experimental results, ChCl/1.5CF3SO3H show excellent oxidative desulfurization activity. Table 2 shows a comparison of the desulfurization activity of ChCl/1.5CF3SO3H and other acidic ionic liquids. It can be seen that ChCl/1.5CF3SO3H has a high desulfurization rate under mild experimental conditions. Furthermore, ChCl/1.5CF3SO3H can be prepared by a simple method and the raw material are easily obtained.

| Catalysts       | S content (μg g⁻¹) | Reaction conditions | Removal of DBT | Ref. |
|-----------------|-------------------|---------------------|----------------|-----|
| [C₄mim]TFEA     | 202               | IL : oil = 1 : 3, H2O2 = 1 mL, 70 °C, 30 min | 100%           | 20  |
| [HMim]BF4       | 1000              | IL : oil = 5 : 3, O/S = 10 : 90 °C, 360 min | 93%            | 21  |
| BMIM[HSO4]      | 1000              | IL : oil = 1 : 2, O/S = 5, 25 °C, 90 min | 99.6%          | 22  |
| [Hnmp]BF4       | 1550              | IL : oil = 1 : 1, O/S = 3, 60 °C, 60 min | 100%           | 23  |
| ChCl/1.5CF3SO3H | 500               | IL : oil = 1 : 5, O/S = 6, 40 °C, 60 min | 98.65%         | This work |
3.2.7 Recovery and regeneration of DESs. After the oxidative desulfurization reaction, the upper oil phase was removed by using a separating funnel. The water in the DESs was removed by rotary evaporation. The DESs were extracted three times using CCl₄ of equal volume. Fresh H₂O₂, recovered DESs were added into model oil under the optimal conditions. As shown in Fig. 6, the removal rate of DBT decreased from 98.65% to 97.16% after five recycles. Two reasons result in the decrease of desulfurization rate. On the one hand, it could be attributed to minimal loss in DESs during the recovery process; on the other hand, it might be attributed to a little of residual oxidation products of sulfide in the recovered DESs.

3.2.8 FT-IR characterization of oxidative production. As shown in Fig. 7(a), the mixture of oil phase and DESs phase is transparent liquid before the desulfurization reaction. However, there is white precipitate between oil and DESs after the desulfurization reaction in Fig. 7(a). In order to identify the white precipitate, a reverse extraction experiment was carried out using carbon tetrachloride (CCl₄) as the extraction agent and the DESs as the extraction phase at room temperature. It can be found that the white substance is quickly dissolved into carbon tetrachloride. After the carbon tetrachloride were remove by distillation, the white powder was obtained. The infrared analysis of the white powder is shown in Fig. 7(b). Three infrared absorption peaks are observed at 1166, 1047, and 1288 cm⁻¹ corresponding to the three characterization peaks of DBTO₂. 36,37 This shows that the DBT was oxidized to DBTO₂ in the oxidative desulfurization system.

3.2.9 Mechanism of catalytic oxidative desulfurization. As seen in Fig. 8, H₂O₂ is a strong oxidant in the acidic medium. DBT was extracted into the DESs phase and was oxidized to DBTO₂ under the action of ChCl/1.5CF₃SO₃H and H₂O₂. During oxidative desulfurization process, DESs acted as not only extractant but also catalyst, so a continuous decrease of the content of DBT in n-octane was observed until H₂O₂ was completely decomposed.

3.3 Catalytic oxidative desulfurization of diesel oil
The S-removal activity of ChCl/1.5CF₃SO₃H for the real diesel oil with a total S-content of 375 mg L⁻¹ was also investigated. Under the optimal condition, the desulfurization rate of 68.5% was obtained. Sulfur removal efficiency was found to be lower in diesel oil than that obtained in model oil (98.65%), which may be attributed to the more complex chemical composition of diesel. 38
4 Conclusion

A series of ChCl/CF_{3}SO_{3}H (X from 1.0 to 2.0) were synthesized by stirring a mixture of ChCl and CF_{3}SO_{3}H. The DESs were used as extractant and catalyst in the oxidative desulfurization process. Compared to other acidic ionic liquids, the synthesis method of ChCl/1.5CF_{3}SO_{3}H is simple. The ChCl/1.5CF_{3}SO_{3}H exhibits a high desulfurization activity for DBT in model oil and can be recycled 5 times without a significant decrease in activity. The results demonstrate the DESs exhibit the high catalytic activity and stability for the desulfurization system.

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