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Protic ionic liquids with low viscosity for efficient and reversible capture of carbon dioxide

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Abstract

Protic ionic liquids (PILs) are considered as potential solvents for CO₂ capture due to their simple synthetic routes and unique properties. In this work, three low viscous PILs, tetramethylguanidinium imidazole ([TMGH][Im]), tetramethylguanidinium pyrrole ([TMGH][Pyrr]) and tetramethylguanidinium phenol ([TMGH][PhO]) were synthesized and the effect of anions, temperature, CO₂ partial pressure and water content on CO₂ absorption performance of PILs was also systematically studied. It was found that the PILs with larger basicity show higher CO₂ absorption capacity, and [TMGH][Im] simultaneously shows relatively high absorption rate and CO₂ absorption capacity of 0.154 g CO₂/g IL at 40 °C, 1 bar. The addition of H₂O has a positive effect on gravimetric absorption capacity of CO₂ at the range of 0-20 wt% H₂O, and the highest capacity of 0.186 g CO₂/g IL was achieved as the water content was 7 wt%. *In-situ* FTIR, ¹³C NMR and theoretical calculations verified that more stable bicarbonate are produced during CO₂ absorption by [TMGH][Im]-H₂O system. However, neat [TMGH][Im] can react with CO₂ to form the reversible carbamate, leading to excellent recyclability after four absorption-desorption cycles. The results implied that neat [TMGH][Im] shows great potentials in CO₂ absorption applications.

Keywords: Ionic liquids, protic, low viscosity, CO₂ absorption, mechanisms

1. Introduction

Greenhouse effect caused by carbon dioxide (CO₂) emission has become one of the most serious environmental problems in the global area and awakened wide attention due to the excessive combustion of fossil fuels (Mondal et al., 2012). Therefore, developing new technologies to reduce CO₂ emission is the main concern (Guo et al., 2015; Ozturk, 2015; Lazarevic et al., 2017). One of the most promising methods to mitigate the impact of greenhouse gas for climate change and CO₂ utilization is developing carbon dioxide capture and storage technologies (CCS) (Deng, 2016). Alkanolamines (including monoethanolamine, diethanolamine and N-methyldiethanolamine) are the most commonly applied solvents for CO₂ capture in industries (Han et al., 2011). However, aqueous alkanolamines for CO₂ capture have some intrinsic drawbacks, such as corrosivity and degradation during the regeneration of solvents. Besides, large heat capacity of water and easy volatilization of alkanolamines result in high solvent loss and intensive energy consumption (Vaidya and Kenig, 2007). Accordingly, there is a pressing demand for seeking novel CO₂ capture solvents with favorable absorption performance, low solvent loss and energy consumption.

Ionic liquids (ILs) have emerged as potential solvents for gas separation (Zhang et al., 2013; Gao et al., 2015; Cao et al., 2017; Shang et al., 2017; Zhu et al., 2017) due to their outstanding properties, such as high thermal stability, low heat capacity, negligible vapor pressure and adjustable nature. ILs as CO₂ absorbents is one of the most effective alternatives to replace conventional aqueous alkanolamines solvents. Since Blanchard

et al. (1999) reported that CO₂ can dissolve in ILs, while ILs do not dissolve in CO₂, a large number of publications on CO₂ capture by ILs through physical interaction were reported (Palomar et al., 2011; Sistla and Khanna, 2011; Zhao et al., 2011; Ramdin et al., 2012). However, CO₂ absorption capacity in these ILs is only about 0.03 mol CO₂/mol IL under atmospheric pressure (Wang et al., 2011). In order to improve CO₂ absorption capacity in ILs, the chemisorption ILs were designed by introducing basic functional groups into ILs. Bates et al. (2002) proposed a new strategy for chemical absorption of CO₂ by amino-functionalized IL 1-propylamide-3-butylimidazolium tetrafluoroborate ([apbim][BF₄]), the absorption capacity of CO₂ in this IL is 0.074 g CO₂/g IL at room temperature and atmospheric pressure. Subsequently, there have been many reports on CO₂ absorption using amino-functionalized ILs, including amine-based and amino acid-based ILs (Hu et al., 2014; Lv et al., 2016; Huang et al., 2018). Riisager et al (2014). synthesized a series of amino acid-based ILs as CO₂ absorbents, and high CO₂ capacity of 0.13 g CO₂/g IL was measured for [N₆₆₆₁₄][Lys] at ambient condition. Recently, an extremely high CO₂ capacity up to 0.25 g CO₂/g IL have been achieved by using tri-*n*-butylethylphosphonium succinimide ([P₄₄₄₂][Suc]) (Huang et al., 2017). Nevertheless, the difficulty in chemisorbing ILs is the extremely high viscosity and complex synthetic routes of most task-specific ILs. For example, [P₄₄₄₂][Suc] with a viscosity as high as 998 mPa·s at 20 °C should be prepared through anion-exchange and neutralization reaction. Consequently, developing novel task-specific ILs with high CO₂ absorption performance, simple synthetic routes and low viscosity are the main tasks for CO₂ capture.

Protic ionic liquids (PILs) can be easily obtained through a neutralization reaction of Brønsted acid and base. Recently, PILs have caught considerable attention in CO₂ capture fields due to their simpler synthetic routes and lower costs compared to most of aprotic ionic liquids (AILs) (Mumford et al., 2015; Xu, 2017; Oncsik et al., 2018). For example, Alcantara et al. (2018) studied the CO₂ absorption performance in four potential PILs which were synthesized through one-step neutralization of diethylamine/ethanolamines and butanoic acids. The results indicated that the PILs 2-hydroxyethylammonium butanoate ([2HEA][Bu]) and N-methyl-2-hydroxyethylammonium butanoate ([m-2HEA][Bu]) exhibited significant higher CO₂ solubility than most ILs ones analyzed. Tetramethylguanidine (TMG), as a common kind of superbase, has been widely reacted with proton donors and the obtained TMG-based PILs were frequently used in different fields (Reddy et al., 2015; Porwal et al., 2016; Singh et al., 2017). In recent years, TMG-based PILs were applied in acid gases absorption due to low viscosity, simple synthetic routes and excellent absorption performance (Jin et al., 2011; Meng et al., 2018). However, current research of TMG-based PILs mainly focused on sulfur dioxide (SO₂) absorption, rarely on CO₂ absorption. The efficiency of SO₂ absorption mainly related to the reaction of SO₂ with N-H group in [TMGH]⁺ cation forming an N-S bond (Wu et al., 2004), while the reactivity of CO₂ and [TMGH]⁺ cation is not valid because of the low acidity of CO₂ (Jessop et al., 2012). Recently, a novel method for CO₂ capture by tunable anion-functionalized PILs based on the single-site interaction between the CO₂ and electronegative nitrogen or oxygen atom was reported, it was shown that CO₂

absorption capacity can be easily tuned by the IL basicity (Wang et al., 2011). Therefore, anion-functionalized PILs with TMG as the proton acceptor combining with reactive hydrogen in azole or phenol is suitable for CO₂ absorption because of the strong basicity of TMG.

In this work, considering the significant role of IL basicity on CO₂ absorption (Wang et al., 2011), three basic PILs were designed and synthesized by neutralizing TMG with different weak proton donors, including imidazole (Im), pyrrole (Pyrr) and phenol (PhO) with different dissociation constant (pK_a). CO₂ absorption performance of these ILs, including the effect of anions, temperature, CO₂ partial pressure and water content on CO₂ absorption, as well as recyclability in the absence and presence of H₂O were systematically studied. The CO₂ absorption mechanisms of these systems were also investigated by *in-situ* FTIR, ¹³C NMR analysis and theoretical calculations.

2. Experimental section

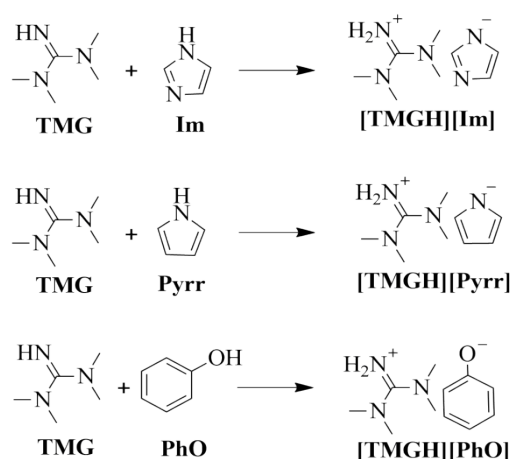
2.1. Materials

The materials TMG, Im and Pyrr were purchased from Aladdin Industrial Corporation. PhO was obtained from Xilong Scientific Limited. All the above chemical reagents were obtained in the analytical purity grade and used without further purification. CO₂ (>99.9% purity) and nitrogen (N₂, >99.9% purity) gases were supplied by Beijing Beiwen Gases Factory.

2.2. Synthesis and characterization of PILs

Three target basic PILs, tetramethylguanidinium imidazole [TMGH][Im],

tetramethylguanidinium pyrrole [TMGH][Pyrr] and tetramethylguanidinium phenol [TMGH][PhO] were prepared by direct neutralization of an appropriate TMG with different weak proton donors (Im, Pyrr and PhO) at room temperature and atmospheric pressure for 24 h (**Scheme 1**), and the synthetic procedures were similar to our previous work (Zhao et al., 2010, 2011). After that, all the ILs were dried in the vacuum oven at 60 °C for 40 h before use. The water contents in these ILs were measured by using Karl Fischer titration (Mettler Toledo Coulometric KF titrator C20), and all the water contents were below 320 ppm.



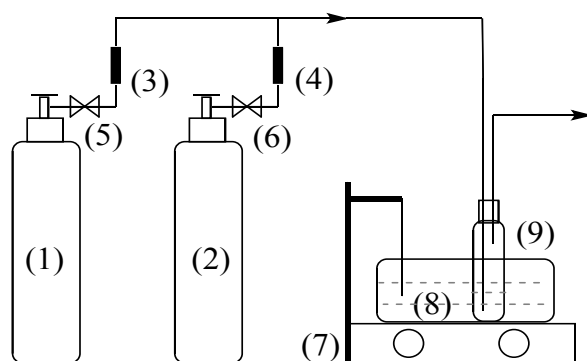
Scheme 1. Synthesis of [TMGH][Im], [TMGH][Pyrr] and [TMGH][PhO].

The obtained samples were characterized by ^1H and ^{13}C NMR using Bruker 600 spectrometer. FTIR spectra of these ILs were recorded in the range of 3600-400 cm^{-1} using Thermo Nicolet 380 spectrometer. Density and viscosity were measured using density meter (Anton Paar DMA 5000) and viscometer (Anton Paar AMVn) from 30 to 80 °C with 10 °C intervals under atmospheric pressure. The thermal stability of these ILs was measured using TGA Q5000 V3.15 with a heat rate of 10 °C/min in N_2 atmosphere at a flow rate of 20 ml/min. The ^1H NMR, ^{13}C NMR, FTIR spectra and

TGA data of all the studied PILs were concluded in the Supporting Information.

2.3. Absorption and desorption of CO₂

The experimental setup for CO₂ absorption is schematically represented in **Scheme 2**, and followed the procedures according to our previous work (Wang et al., 2014; Zeng et al., 2014, 2018). The absorption experiments were carried out in a glass container with an inner diameter of 2 cm. In a typical experiment, CO₂ gas (1) was bubbled through IL (about 5 g) in the glass container (9) at a flow rate of about 140 ml/min. The glass container was partly immersed in the water bath (8) at the desired temperature (the standard uncertainty of temperature is ± 0.1 °C). The amount of absorbed CO₂ was determined at regular intervals by an electronic balance with an accuracy of ± 0.1 mg, which could remain constant when absorption of CO₂ reached to equilibrium. To ensure precision during weight measurements of samples, the glass container was dried outside before weighted.



Scheme 2. Experimental diagram for CO₂ absorption. (1) CO₂ gas cylinder; (2) N₂ gas cylinder; (3) and (4) gas mass flowmeter; (5) and (6) valves; (7) magnetic stirrer; (8) water bath; (9) glass container with IL.

The effect of temperature on CO₂ absorption performance of ILs was carried out under atmospheric pressure by varying temperatures from 30 to 50 °C. During CO₂ absorption under reduced pressure, the absorption temperature was kept constant at 40 °C, the mixed gases with different CO₂ partial pressure were prepared by adjusting the flow rate of CO₂ and N₂ gases. To investigate the influence of water content on CO₂ absorption, CO₂ saturated with water vapor was bubbled through IL in the presence of different water content with 140 ml/min flow rate at 40 °C and atmospheric pressure.

For desorption process, CO₂-absorbed IL was transferred to a round-bottom flask and CO₂ was released by rotary evaporation at 60 °C for 2 h. The desorption of CO₂ from IL-H₂O system was similar to neat IL system, rotary evaporator was used to remove most of CO₂ and H₂O in CO₂ absorbed IL-H₂O system, the obtained product was dried in vacuum oven for 24 h to remove the residual moisture. Then fresh water was added to the obtained product at a certain ratio for next CO₂ absorption.

2.4 Computational methods

All calculations herein were performed with the Gaussian 09 program (Frisch et al., 2013) by using Density Functional Theory. All geometries of reactants, transition states and products were optimized at the M06-2X/def2-TZVP level of theory. The solvent effects of ILs were simulated by the SMD-GIL solvation model (Bernales et al., 2012). The frequency calculation was carried out at the same level to confirm the optimized structures to be energy minima without any imaginary frequency, and transition states has one and only one imaginary frequency. The interaction energies (or

energy barriers) were calculated as the energy difference between the products (or transition states) and the reactants.

3. Results and discussion

3.1. Physical properties of PILs

Physical properties, such as density and viscosity of ILs are important parameters for industrial application of CO₂ absorbents. Therefore, densities and viscosities of these PILs at various temperatures were measured and the results are shown in **Fig. 1** and **2**, respectively. The experimental density results for three ILs from 30 to 80 °C showed that temperature dependences of ILs densities have the linear behavior and decreased with the increasing of temperature. The density order of these ILs are: [TMGH][PhO] > [TMGH][Im] > [TMGH][Pyr].

The viscosity of absorbents has significant effect on mass transfer during CO₂ absorption. ILs with low viscosities can result in low mass transfer resistance between liquid and gas phases, and thereby increase CO₂ absorption rate. As seen in **Fig. 2**, the viscosities of all studied ILs decreased with an increase in temperature. The viscosities of [TMGH][Im], [TMGH][Pyr] and [TMGH][PhO] were 6.44, 2.10 and 26.77 mPa·s at 30 °C, respectively, which are far lower than most of conventional ILs, for example, the viscosity of [Bmim][BF₄] was 68.90 mPa·s at 30 °C (Zhao et al., 2010). The results indicated that all these ILs have relatively low viscosities, which facilitate CO₂ diffusion in ILs during absorption processes.

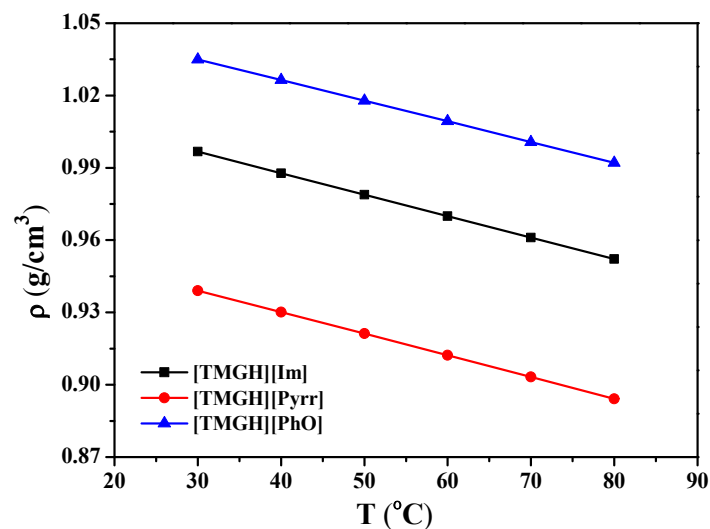


Fig. 1. Densities of [TMGH][Im], [TMGH][Pyrr] and [TMGH][PhO] at various temperatures.

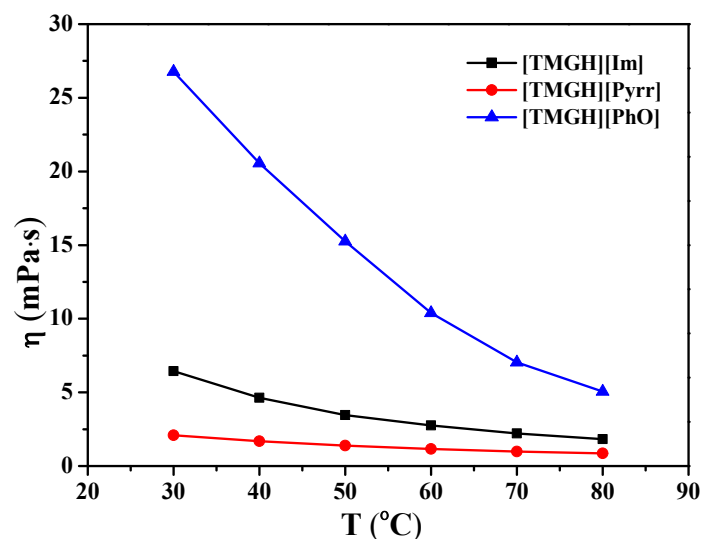


Fig. 2. Viscosities of [TMGH][Im], [TMGH][Pyrr] and [TMGH][PhO] at various temperatures.

3.2. Effect of anions on CO₂ absorption

In order to investigate the CO₂ absorption performance in these PILs, CO₂ capture experiments were firstly carried out in three PILs with different anions, [TMGH][Im], [TMGH][Pyrr] and [TMGH][PhO] at 40 °C under atmospheric pressure in **Fig. 3**. It was

found that CO₂ absorption in all the ILs is very quick at the beginning, then decreased gradually and finally achieved equilibrium. Among the three ILs, [TMGH][Im] and [TMGH][Pyrr] exhibited higher CO₂ absorption capacity of 0.154 and 0.159 g CO₂/g IL, respectively, while CO₂ absorption capacity of [TMGH][PhO] was only 0.010 g CO₂/g IL at the same condition. In fact, CO₂ absorption capacity is significantly affected by the basicity of ILs. In general, the anion of ILs with larger p*K*_a has higher CO₂ absorption capacity due to its stronger reactivity with CO₂ (Wang et al., 2011). As shown in **Table 1**, when the p*K*_a of anions in DMSO decreased from 23.0 to 16.4, CO₂ molar absorption capacity decreased remarkably from 0.66 to 0.05 mol CO₂/mol IL.

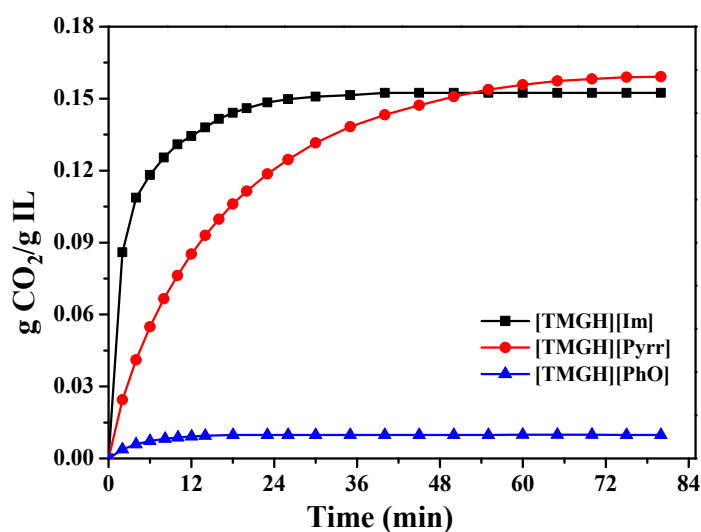


Fig. 3. CO₂ absorption in [TMGH][Im], [TMGH][Pyrr] and [TMGH][PhO] at 40 °C and 1 bar.

Table 1: The effect of anions pK_a on the CO₂ absorption capacity.

IL	M (g/mol)	T (°C)	Capacity (mol CO ₂ /mol IL)	pK_a in DMSO
[TMGH][Pyrr]	182.27	40	0.66	23.0 (Bordwell et al., 1981)
[TMGH][Im]	183.26	40	0.64	18.6 (Bordwell, 1988)
[TMGH][PhO]	209.29	40	0.05	16.4 (Ritchie, 1969)

Meanwhile, [TMGH][Im] also showed the faster rate of CO₂ absorption than other PILs, and the absorption can achieve equilibrium in 20 min. Although [TMGH][Pyrr] has the lowest viscosity, the absorption of CO₂ was completed in nearly 60 min, which is substantially three times that of [TMGH][Im]. The possible reason is that the solid state product was observed in [TMGH][Pyrr] after CO₂ absorption, which caused an increase in the apparent slurry viscosity, and reduction in effective diffusivity of CO₂ in the liquid (Kumar et al., 2003). As described above, [TMGH][Im] has a relative high CO₂ absorption capacity and the highest absorption rate compared with the other two ILs, which can be considered as a candidate solvent for CO₂ capture. Therefore, the effect of temperature, CO₂ partial pressure and water content on CO₂ absorption performance in [TMGH][Im] were further investigated and discussed.

3.3. Effect of temperature on CO₂ absorption

Temperature plays a key role in CO₂ absorption process, hence the effect of

temperature on CO₂ absorption performance of [TMGH][Im] was investigated from 30 to 50 °C under atmospheric pressure as shown in **Fig. 4**. The results indicated that CO₂ absorption capacity decreased obviously with temperature increasing. The absorption capacity of [TMGH][Im] was 0.177 g CO₂/g IL at 30 °C, but reduced to 0.124 g CO₂/g IL at 50 °C. This is because CO₂ absorption is an exothermic process, and the increasing of temperature is not favorable for CO₂ absorption. On the other hand, increasing temperature has a positive influence on mass transfer during the absorption of CO₂ due to lower viscosity of absorbents at higher temperature, and finally decreases saturation time. The saturation time decreased from 40 to 15 min when temperature changed from 30 to 50 °C.

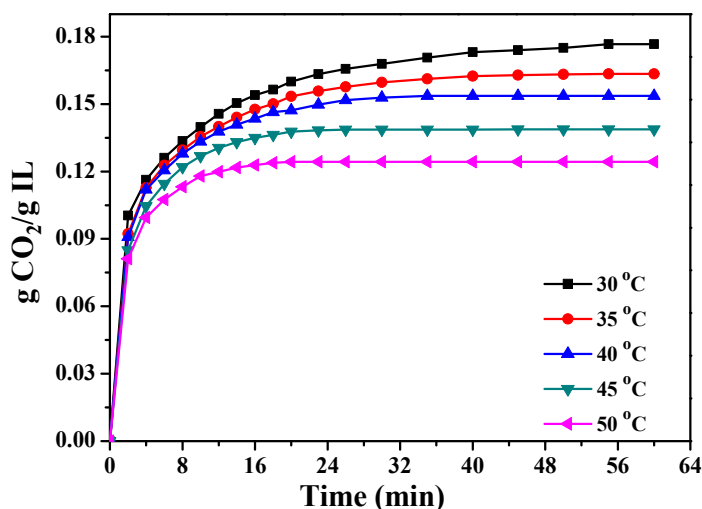


Fig. 4. CO₂ absorption in [TMGH][Im] at various temperatures.

3.4. Effect of partial pressure on CO₂ absorption

In industrial processes, CO₂ frequently coexists with other gases and CO₂ partial pressure almost lower than 1.0 bar. As illustrated in **Fig. 5**, CO₂ absorption performance of [TMGH][Im] was carried out at 40 °C with a CO₂ partial pressure of 0.1, 0.3, 0.5,

0.7, 0.9 and 1.0 bar to study the effect of CO₂ partial pressure on CO₂ absorption. With the decreasing of CO₂ partial pressure, CO₂ absorption capacity of [TMGH][Im] reduced gradually. The absorption capacity of [TMGH][Im] was as low as 0.05 g CO₂/g IL at 0.1 bar. From the investigation of the effect of temperature and CO₂ partial pressure on CO₂ absorption performance, both increasing CO₂ partial pressure and reducing temperature are benefit for enhancing CO₂ absorption capacity.

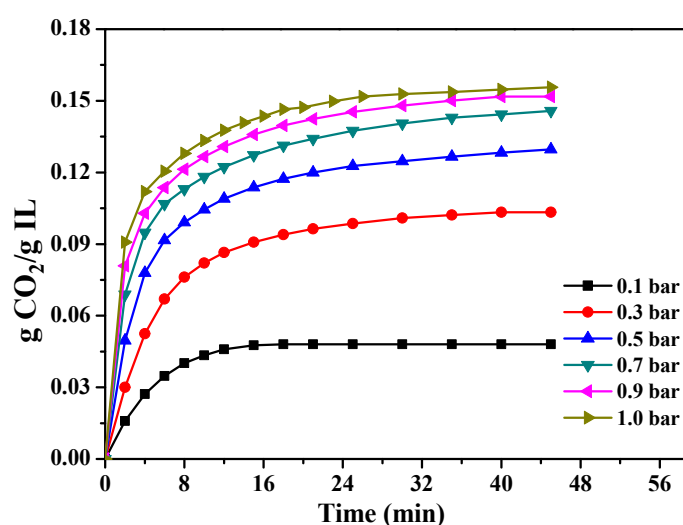


Fig. 5. CO₂ absorption in [TMGH][Im] at various CO₂ partial pressures.

3.5. Effect of water content on CO₂ absorption

The effect of water content on CO₂ absorption performance of [TMGH][Im] was also investigated under 40 °C and atmospheric pressure. As seen in **Fig. 6** and **7**, water contents can remarkably affect the CO₂ capacity of [TMGH][Im]. At the beginning, both CO₂ gravimetric and molar absorption capacity increased greatly with the increasing of water content, the highest absorption capacity up to 0.186 g CO₂/g absorbent (0.83 mol CO₂/mol IL) when 7 wt% H₂O was added to [TMGH][Im]. CO₂ gravimetric absorption capacity of absorbents reduced with an increase of water content

but still higher than that in neat [TMGH][Im] in the range from 7 to 20 wt% H₂O, which was 0.161 g CO₂/g absorbent in 80 wt% [TMGH][Im]-20 wt% H₂O system, but CO₂ molar absorption capacity was almost maintain at 0.83 mol CO₂/mol IL. The possible reason is that CO₂ can directly react with the imidazole anion to form the carbamate product in neat [TMGH][Im], while in the presence of H₂O, the imidazole anion interacts more strongly with the H₂O than CO₂, and a more stable bicarbonate and neutral Im may generate after absorption of CO₂, resulting in an increase in CO₂ absorption capacity (Thompson et al., 2014). On the contrary, excessive quantities of H₂O have a negative influence on CO₂ gravimetric absorption capacity of [TMGH][Im]-H₂O systems due to the reducing of [TMGH][Im] content, it is also shown that CO₂ gravimetric absorption capacity becomes lower when 25 wt% H₂O was existed in [TMGH][Im], which was 0.149 g CO₂/g absorbent. The detail reason will be discussed in the section of absorption mechanism.

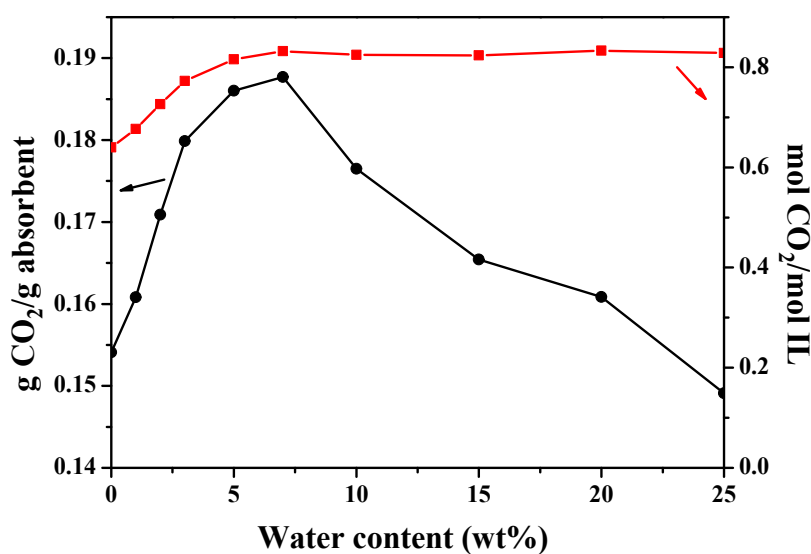


Fig. 6. CO₂ absorption capacity of [TMGH][Im]-H₂O systems at 40 °C and 1 bar.

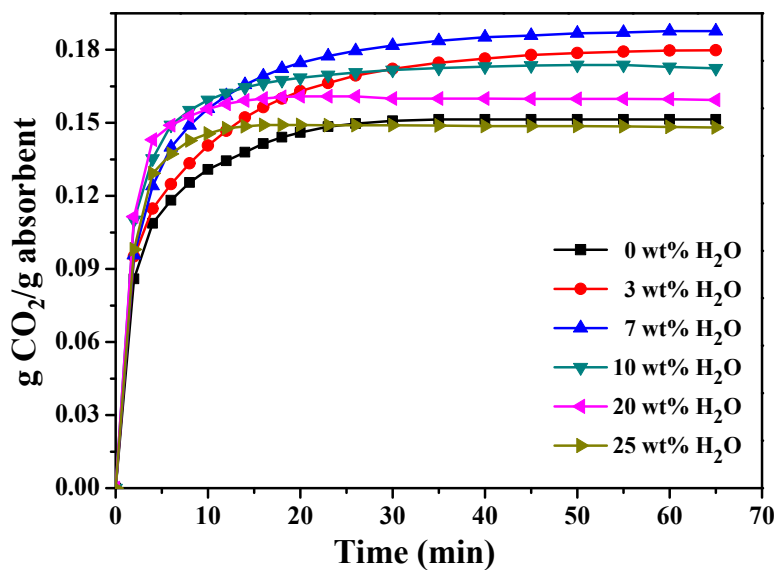


Fig. 7. Effect of water content on CO₂ absorption performance of [TMGH][Im] at 40 °C and 1 bar.

3.6. CO₂ absorption mechanisms in PILs

As aforementioned, the addition of H₂O changed the CO₂ absorption performance of [TMGH][Im]. In order to understand the absorption mechanism of [TMGH][Im] in the absence and presence of H₂O, *in-situ* FTIR, ¹³C NMR and theoretical calculations were used to analyze the interaction between ILs or IL-H₂O system and CO₂.

3.6.1 In-situ FTIR and NMR analysis

The [TMGH][Im]-H₂O system containing 80 wt% [TMGH][Im] and 20 wt% H₂O showed higher absorption capacity and faster absorption rate. Therefore, neat [TMGH][Im] and 80 wt% [TMGH][Im]-20 wt% H₂O system were selected to investigate the absorption mechanism of [TMGH][Im] in the absence and presence of H₂O via *in-situ* FTIR and ¹³C NMR spectra. As shown in **Fig. 8**, the FTIR spectra of

the CO₂-absorbed [TMGH][Im] contained three new peaks compared to neat [TMGH][Im]. The peaks at 1700 and 1293.7 cm⁻¹ can be attributed to the stretching vibrations of C=O and C-O of carbamate, respectively. The peak at 1176.3 cm⁻¹ was appeared due to the formation of C-N. The results indicated that the carbamate was formed between the basic nitrogen of imidazole anion and CO₂ (Wang et al., 2011; Zhu et al., 2017). For [TMGH][Im]-H₂O system, two new characteristic peaks at 1252.7 and 1040.2 cm⁻¹ appeared as shown in **Fig. 9** verified the formation of the bicarbonate during the absorption of CO₂, which means that new product was generated in the presence of H₂O. Furthermore, By analyzing ¹³C NMR of [TMGH][Im]-based solvents before and after absorption of CO₂ listed in **Fig. 10**, a new carbon signal at 161.16 was observed in the CO₂-absorbed [TMGH][Im] as compared to neat [TMGH][Im], which was attributed to carbonate carbonyl carbon, while the peak at 160.26 ppm appeared after CO₂ absorption in [TMGH][Im]-H₂O was due to the formation of bicarbonate. (Andrews et al., 2011; Simon et al., 2017; Chen et al., 2018). The results were in agreement with relevant researches reported in literature that the basicity of anion can activate the reaction between CO₂ and H₂O to form conjugate acid of the basic anion and bicarbonate (Huang et al., 2019). Based on the above analysis, the possible CO₂ absorption mechanism of [TMGH][Im] in the absence and presence of H₂O was proposed in **Scheme 3**.

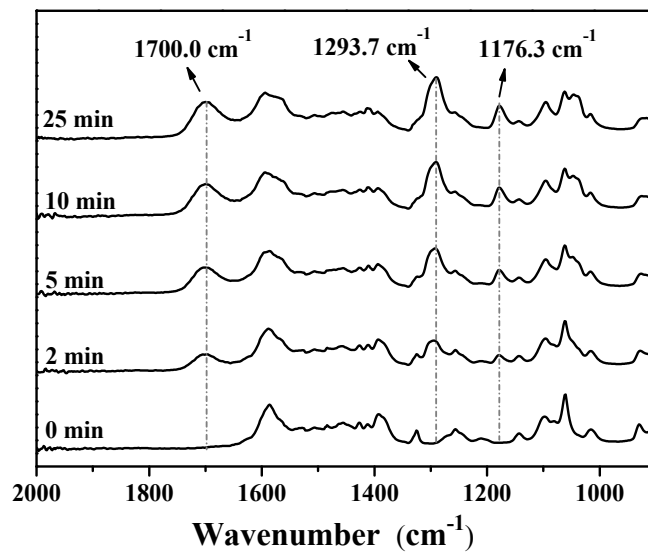


Fig. 8. FTIR spectra of [TMGH][Im] before and after absorption of CO₂.

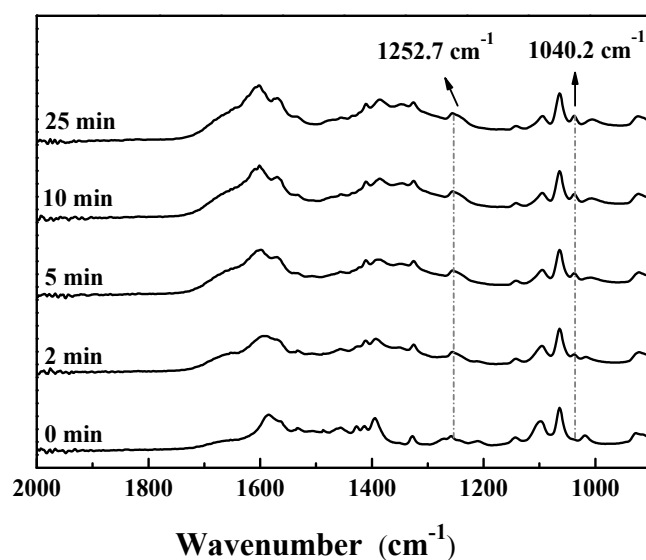


Fig. 9. FTIR spectra of [TMGH][Im]-H₂O system before and after absorption of CO₂.

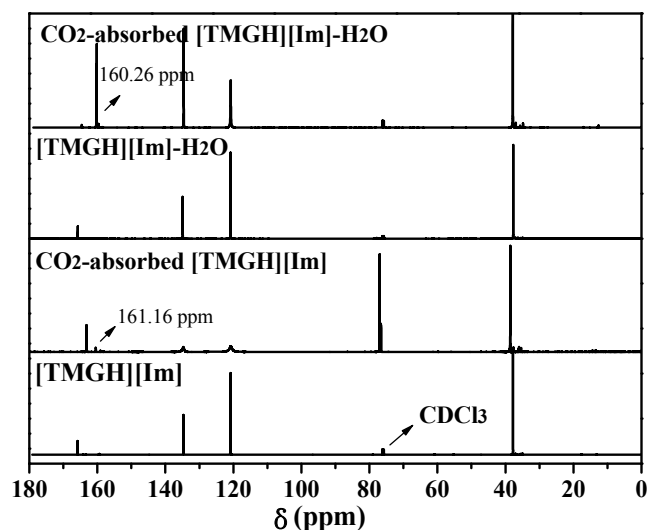
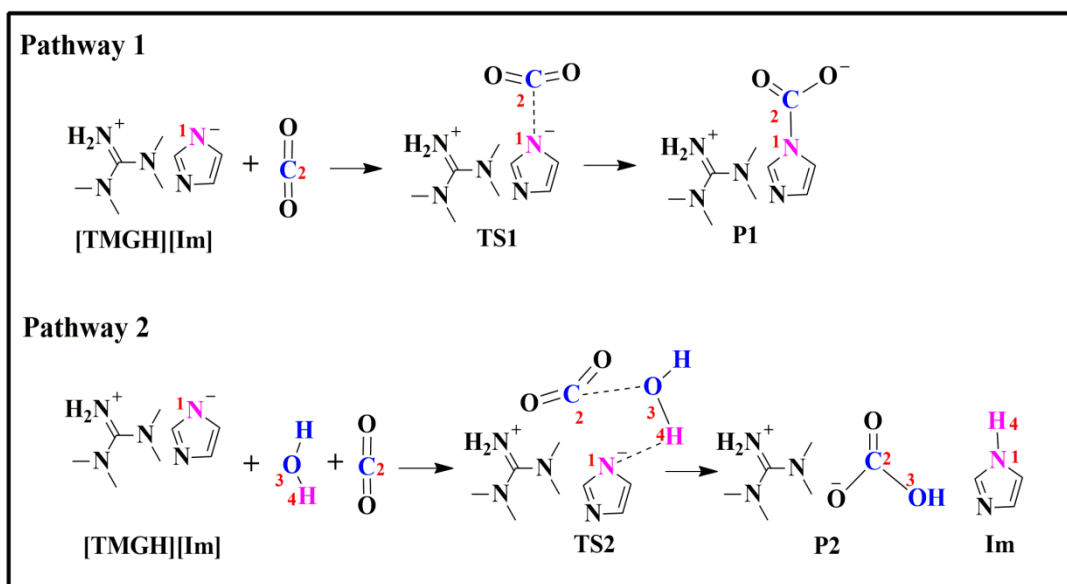


Fig. 10. ^{13}C NMR spectra of $[\text{TMGH}][\text{Im}]$ and $[\text{TMGH}][\text{Im}]\text{-H}_2\text{O}$ systems before and after absorption of CO_2 .

3.6.2 Quantum chemistry calculations

Based on the above experimental results and spectral characterizations, we proposed the possible CO_2 absorption pathways of $[\text{TMGH}][\text{Im}]$ in the absence (pathway 1) and presence (pathway 2) of H_2O in **Scheme 3**. During the absorption of CO_2 in $[\text{TMGH}][\text{Im}]$, carbamate (P1) is generated from the combination of nitrogen atom of N1 on the imidazole anion and CO_2 through transition states (TS1). When H_2O is existed in $[\text{TMGH}][\text{Im}]$, the formation of N1-H4 and C2-O3 bonds generates the bicarbonate (P2) and Im via the transition states (TS2).



Scheme 3. CO₂ absorption pathways of [TMGH][Im] and [TMGH][Im]-H₂O systems

The CO₂ reaction potential energies for two systems are shown in **Fig. 11**. It can be seen that the reaction energy barriers of CO₂ with [TMGH][Im] and [TMGH][Im]-H₂O were 4.45 and 5.61 kcal/mol, respectively. The low energy barriers make CO₂ absorption easy to perform both in neat [TMGH][Im] and [TMGH][Im]-H₂O systems. The interaction energies of products P1 and P2 + Im were -12.26 and -14.86 kcal/mol, respectively. These data suggested that more stable bicarbonate was generated in the presence of H₂O, which is in good agreement with the experimental results.

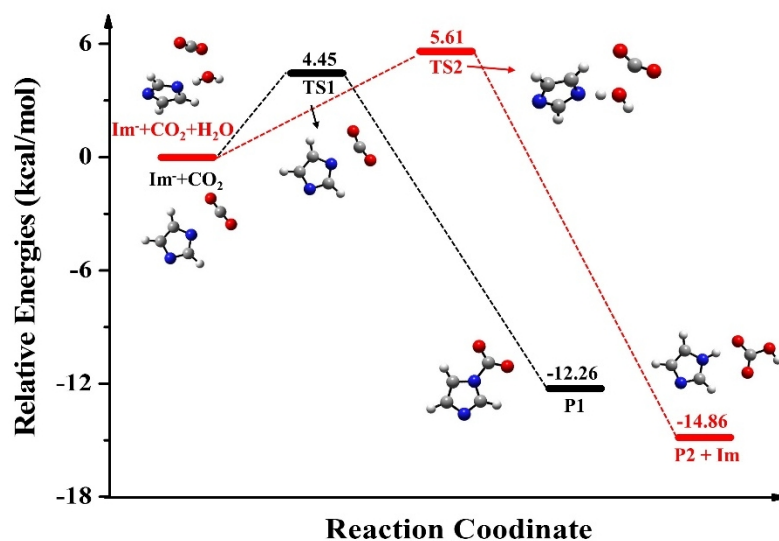


Fig. 11. Optimized structures and potential energy profiles of CO₂ absorption in [TMGH][Im] and [TMGH][Im]-H₂O systems

3.7. Recycle of [TMGH][Im] systems

In order to examine the recyclability of [TMGH][Im] systems, CO₂-absorbed IL systems were regenerated and reused for CO₂ absorption. As illustrated in **Fig. 12**, four absorption-desorption experiments by [TMGH][Im] showed that [TMGH][Im] can be repeatedly recycled without obvious loss of CO₂ absorption capacity. The results indicated that [TMGH][Im] as a novel CO₂ absorbent has an excellent performance for the reutilization and effective CO₂ absorption-desorption ability.

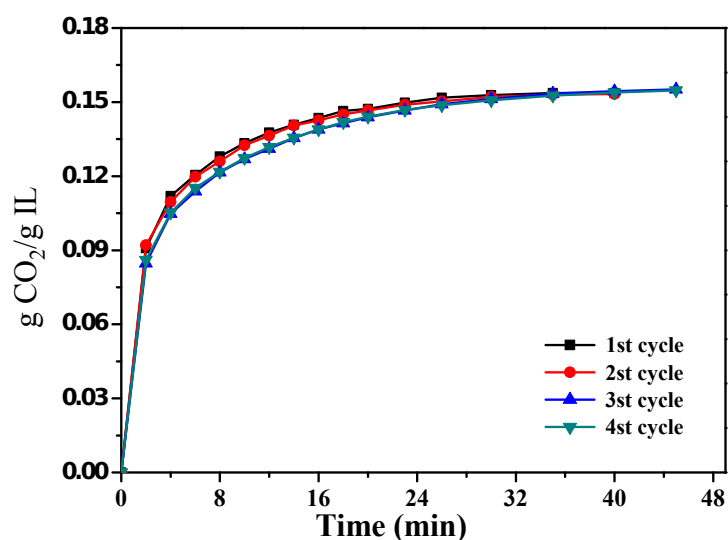


Fig. 12. Recycling of [TMGH][Im] for CO₂ absorption at 40 °C and 1 bar.

For the system of 80 wt% [TMGH][Im]-20 wt% H₂O, CO₂ absorption capacity obviously decreased with the increasing of recycle times as demonstrated in **Fig. 13**. Such solvent is not suitable for industrial CO₂ capture even it has high CO₂ capacity in the first cycle. As seen in **Fig. 11**, energy barriers for the desorption of CO₂ from carbamate and bicarbonate + Im systems were 16.71 and 20.47 kcal/mol, separately. The higher energy barrier makes it difficult to strip CO₂ from bicarbonate and Im.

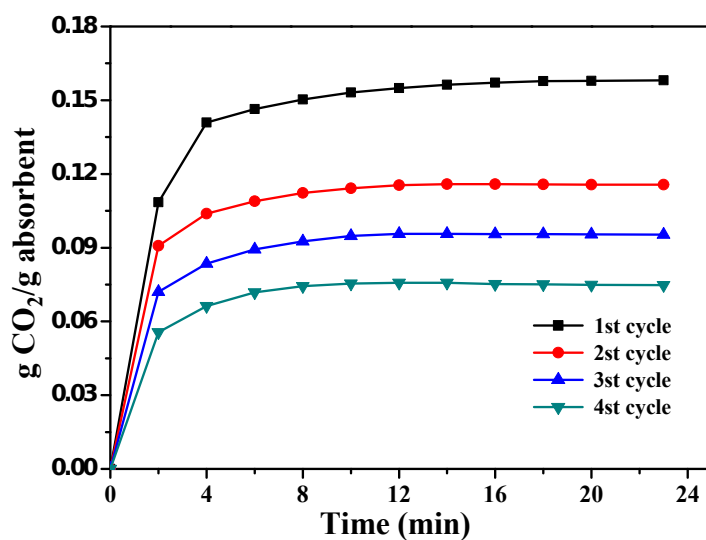


Fig. 13. Recycling of [TMGH][Im]-H₂O system for CO₂ absorption at 40 °C and 1

bar.

4. Conclusions

In summary, three low viscous PILs with different basicity were designed and synthesized by one-step route for efficient absorption of CO₂. The viscosity of [TMGH][Im], [TMGH][Pyrr] and [TMGH][PhO] were 6.44, 2.10 and 26.77 mPa·s at 30 °C, separately, which is quite low compared with conventional ILs. The relationships between the basicity of ILs and their absorption performance of CO₂ indicated that ILs with larger p*K*_a of anions has higher CO₂ absorption capacity. The addition of H₂O changed CO₂ absorption performance of [TMGH][Im], gravimetric absorption capacity of CO₂ firstly increased and then decreased gradually with water content, the highest absorption capacity of 0.186 g CO₂/g absorbent (0.83 mol CO₂/mol IL) at 40 °C and 1 bar was obtained in 93 wt% [TMGH][Im]-7 wt% H₂O system. *In-situ* FTIR, ¹³C NMR and theoretical calculations confirmed that carbamate can be formed reversibly during the absorption of CO₂ in [TMGH][Im], the good recyclability after four absorption-desorption cycles indicating that [TMGH][Im] as a potential alternative for CO₂ capture has an excellent performance for the effective CO₂ absorption-desorption ability. The existence of H₂O result in larger energy barrier during the desorption of CO₂, so it is necessary to make sure that water content in [TMGH][Im] is quite low to ensure excellent recyclability of [TMGH][Im] in CO₂ capture application.

Acknowledgment

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Supporting Information

Protic ionic liquids with low viscosity for efficient and reversible capture of carbon dioxide

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FTIR data of all studied ionic liquids

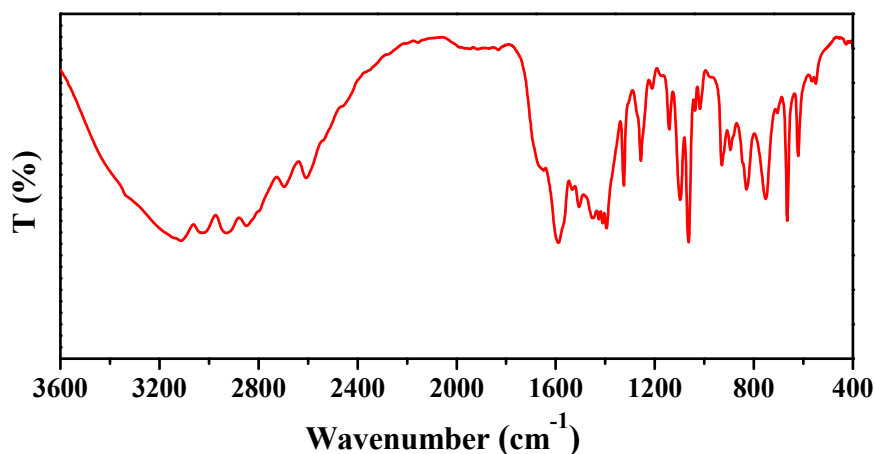


Fig. S1. FTIR spectra of [TMGH][Im]

$\nu = 2925.51$ (the stretching vibration of CH_3), 2696.10 (the characteristic peak of N-CH_3), 1589.07 (the stretching vibration of C=N), 1506.18 and 1410.72 (the stretching vibration of imidazole ring skeleton), 1063.47 cm^{-1} (the stretching vibration of C-N).

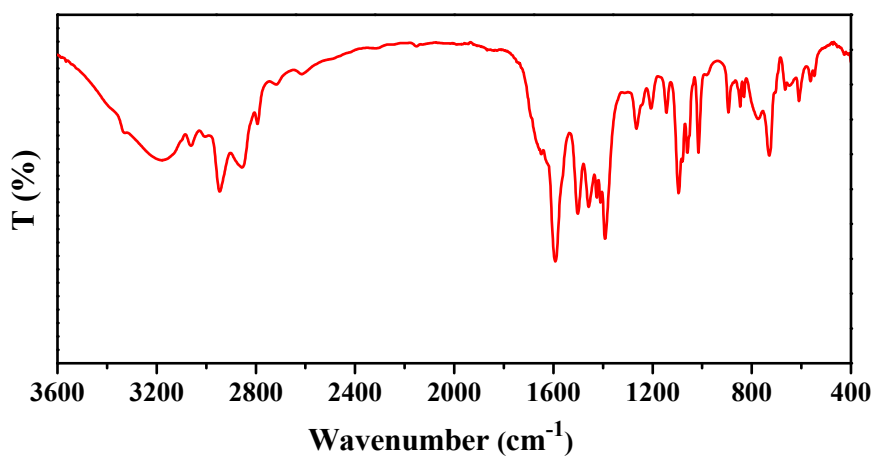


Fig. S2. FTIR spectra of [TMGH][Pyrr]

$\nu = 2945.69$ (the stretching vibration of CH_3), 1592.28 (the stretching vibration of C=N), 1501.53 and 1457.70 (the stretching vibration of pyrrole ring skeleton), 1095.02 cm^{-1} (the stretching vibration of C-N).

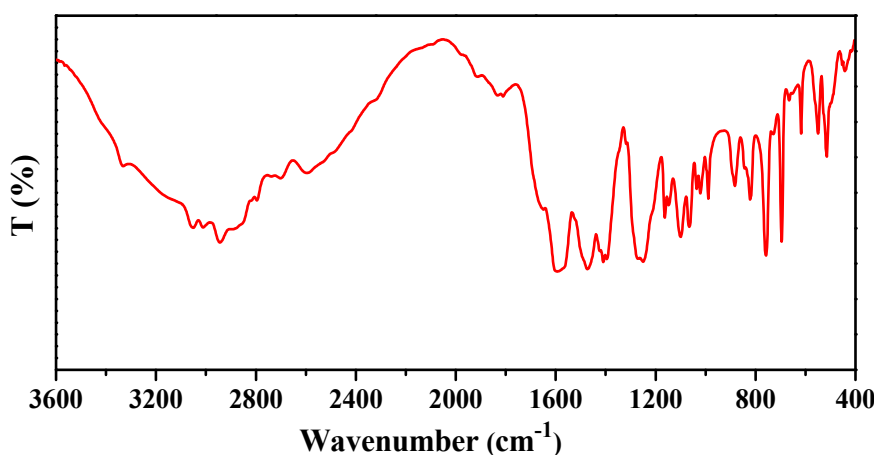


Fig. S3. FTIR spectra of [TMGH][PhO]

$\nu = 2943.95$ (the stretching vibration of CH_3), 2595.64 (the characteristic peak of N-CH_3), 1593.44 (the stretching vibration of C=N), 1472.91 and 1409.27 (the stretching vibration of imidazole ring skeleton), 1250.10 (the stretching vibration of C-O), 1099.71 cm^{-1} (the stretching vibration of C-N).

^1H NMR and ^{13}C NMR data of all studied ionic liquids

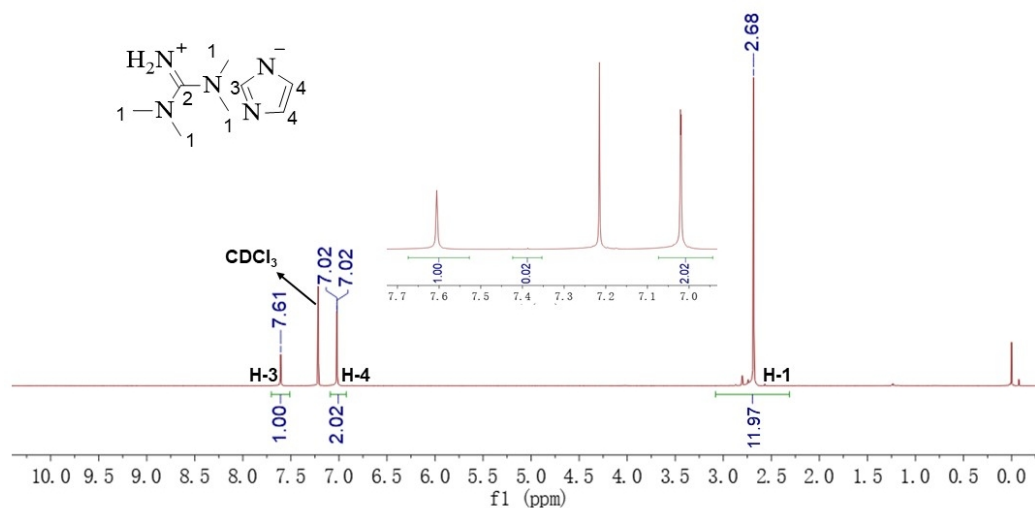


Fig. S4. ^1H NMR spectra of [TMGH][Im]

^1H NMR (600 MHz; CDCl_3 ; TMS): 2.68 (12H, s, CH_3), 7.02 (2H, s, CH), 7.61 ppm (1H, s, CH).

The purity of [TMGH][Im] was estimated by comparing area of the CH-3 signal with its signal area between δ 7.35 and 7.38, the obtained purity is 0.98.

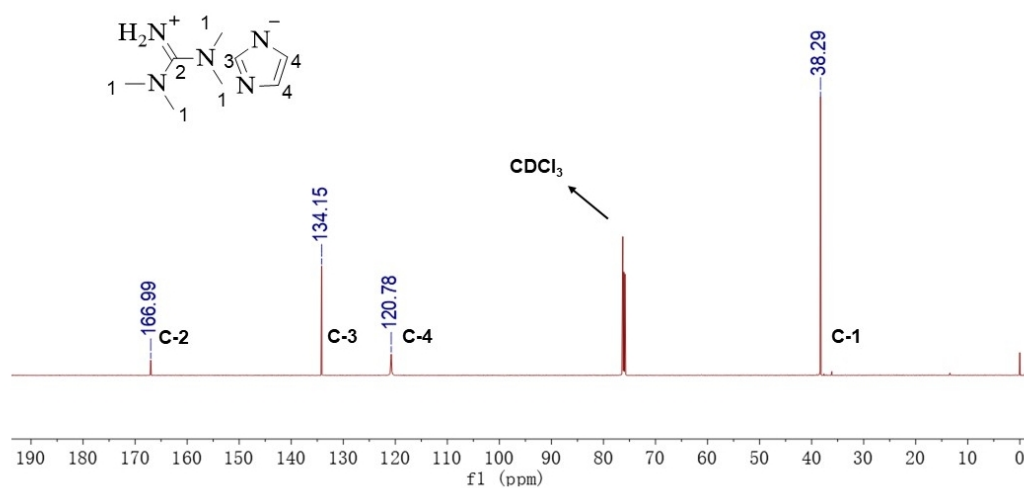


Fig. S5. ^{13}C NMR spectra of [TMGH][Im]

^{13}C NMR (600 MHz; CDCl_3 ; TMS): 38.29, 120.78, 134.15, 166.99 ppm.

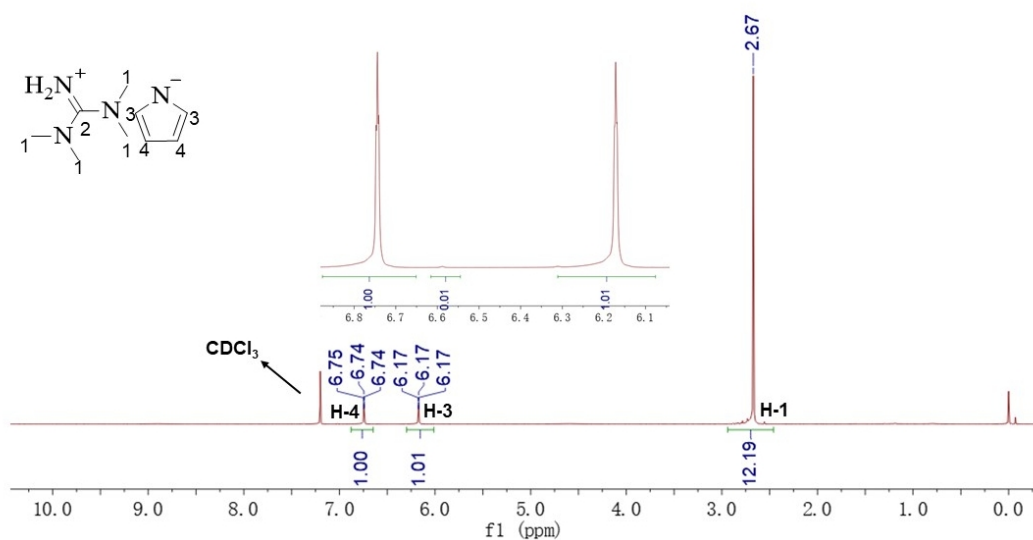


Fig. S6. ^1H NMR spectra of [TMGH][Pyrr]

^1H NMR (600 MHz; CDCl_3 ; TMS): 2.67 (12H, s, CH_3), 6.17 (2H, d, CH), 6.74 ppm (2H, d, CH).

The purity of [TMGH][Im] was estimated by comparing area of the CH-4 signal with its signal area between δ 6.55 and 6.61, the obtained purity is 0.99.

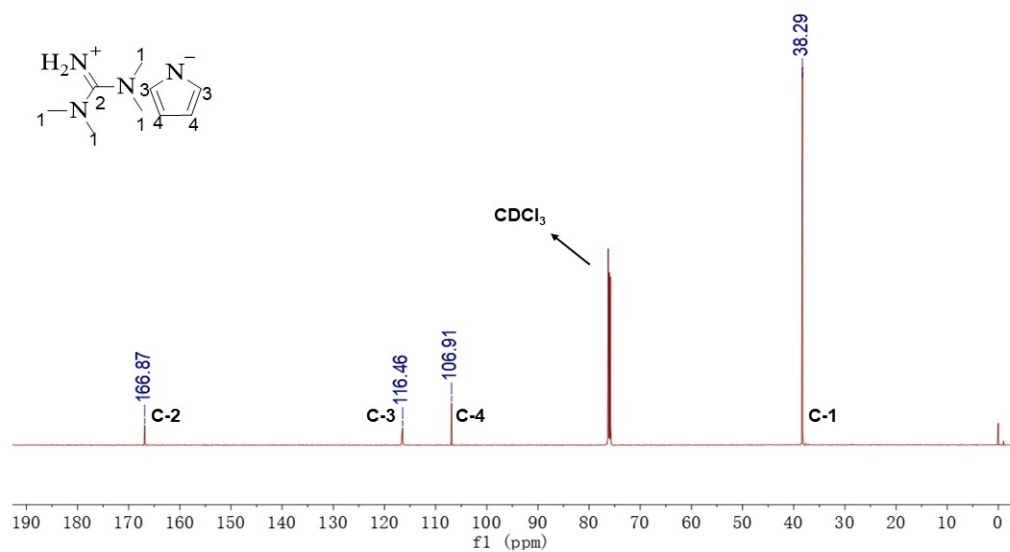


Fig. S7. ^{13}C NMR spectra of [TMGH][Pyrr]

^{13}C NMR (600 MHz; CDCl_3 ; TMS): 38.29, 106.91, 116.46, 166.87 ppm.

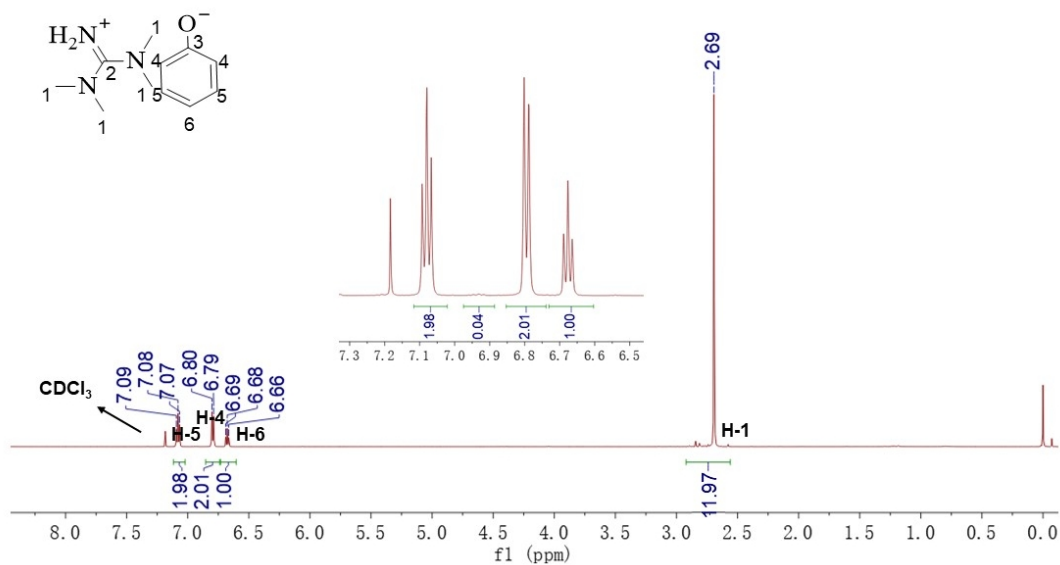


Fig. S8. ^1H NMR spectra of [TMGH][PhO]

^1H NMR (600 MHz; CDCl_3 ; TMS): 2.69 (12H, s, CH_3), 6.68 (H, t, CH), 6.79 (2H, d, CH), 7.08 ppm (2H, t, CH).

The purity of [TMGH][Im] was estimated by comparing area of the CH-5 signal with its signal area between δ 6.89 and 6.96, the obtained purity is 0.98.

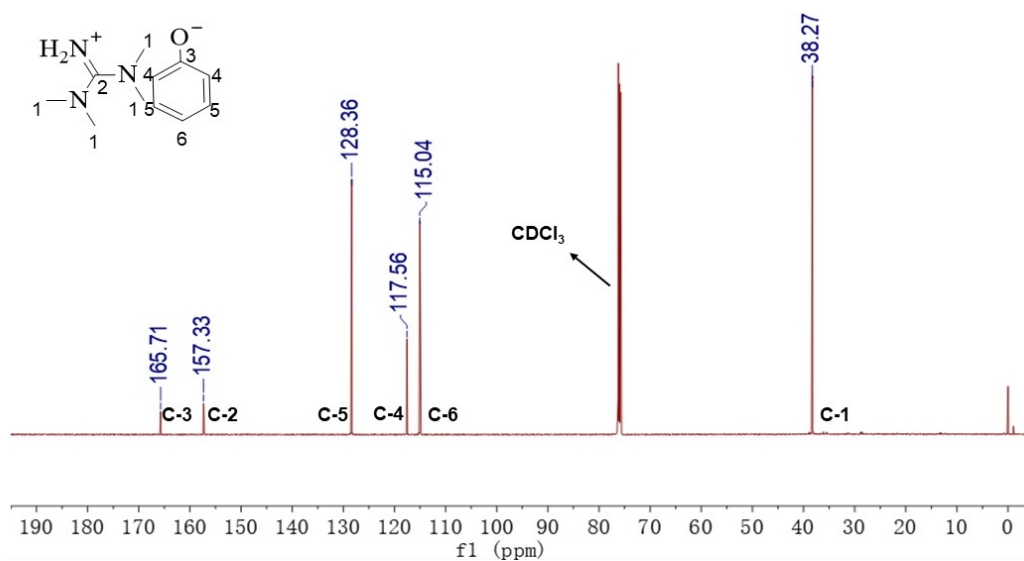


Fig. S9. ^{13}C NMR spectra of [TMGH][PhO]

^{13}C NMR (600 MHz; CDCl_3 ; TMS): 38.27, 115.04, 117.56, 128.36, 157.33, 165.71 ppm.

TGA of all studied ionic liquids

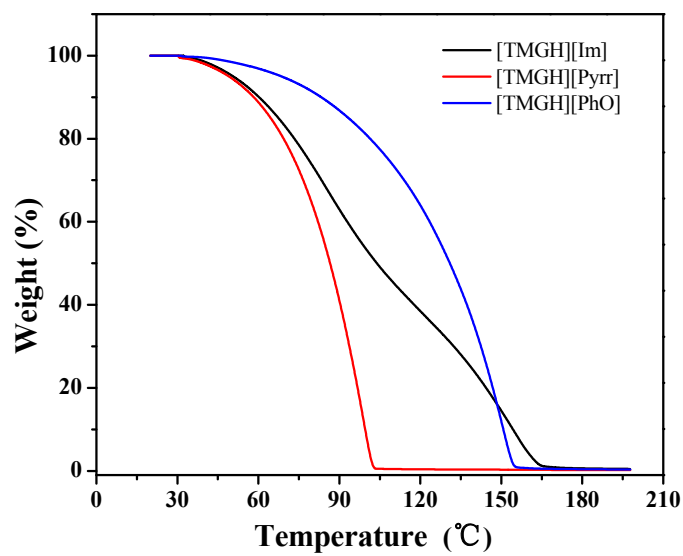


Fig. S10. TGA of [TMGH][Im]、[TMGH][Pyrr] and [TMGH][PhO]