Synthesis of polyoxymethylene dimethyl ethers catalyzed by sulfonic acid-functionalized mesoporous SBA-15

Zhenzhen Xue, Hongyan Shang, Chunhua Xiong, Changbo Lu, Gaojun An, Zailong Zhang, Chuntao Cui and Mingjie Xu

Sulfonic acid-functionalized mesoporous SBA-15 (SO$_3$H-SBA-15) with different sulfur loading was synthesized via a post-synthesis method with 3-mercaptopropyltrimethoxysilane (MPTMS) and was used to investigate the catalytic performance for the synthesis of polyoxymethylene dimethyl ethers (PODE$_n$) from methyal (DMM) and trioxymethylene (TOX). X-ray diffraction, N$_2$ adsorption–desorption, Fourier transform infrared spectra, $^{13}$C-NMR and $^{29}$Si-NMR, elemental analyses, and X-ray fluorescence were used to characterize the structures and sulfur loading of the obtained catalysts, and chemistry titration was carried out to investigate the acid amounts of the catalysts. Through the comparison of the catalysts with different sulfur loading, it was found that the acidity of the catalysts has a decisive effect on the product distribution and the chain length of the products: under the catalysis of the SO$_3$H-SBA-15(0.8) catalyst, which has an acid amount of 0.166 mmol g$^{-1}$, the highest DMM and TOX conversion and PODE$_n$ yield and selectivity were achieved and the desired chain length of PODE$_{2,8}$ was obtained. The optimum reaction conditions such as reaction time, temperature, and molar ratio of DMM and TOX were investigated, and by comparison of different reaction temperatures, it was demonstrated that temperature played an important role in the TOX depolymerization rate and selectivity of the main products and by-products. By adjusting the molar ratio of DMM and TOX, it was observed that an increasing amount of TOX was beneficial for promoting the reaction to generate the products with a higher degree of polymerization. The comparison test of the reaction time confirmed that the reaction reached chemical equilibrium within 60 min and could not be promoted by further extending the time.

1. Introduction

With the increasing depletion of oil resources and environmental pressure, researchers have been developing a new diesel engine fuel or additive that could replace diesel and reduce engine emissions. According to the combustion characteristics of the diesel engine fuel, it is considered that the oxygenated compounds are very suitable to be used as diesel fuel additives. As a representative of an oxygenated compound, polyoxymethylene dimethyl ethers (PODE$_n$) have the characteristics of high cetane number,$^1$ high oxygen content,$^2$ and are free of sulfur and aromatics. They can achieve highly effective and clean combustion$^{1,4}$ that effectively reduces soot emission, PM emission, and particle number without changing the engine infrastructure.$^{3,5,6}$ On the other hand, the feedstock for PODE$_n$ synthesis primarily consists of methanol and methanol derivatives; therefore, the development of PODE$_n$ is of great significance in promoting the development of methanol downstream products of high added value and the healthy development of the coal chemical industry.

It is well known that the synthesis of PODE$_n$ is a type of acid catalytic process; at present, there are relatively few reports on the PODE synthesis catalysts. Several acid catalysts such as ion-exchanged resin,$^7$–$^9$ molecular sieve,$^{10,11}$ ionic liquids,$^{12,13}$–$^{15}$ heteropoly acids,$^{15}$ Zr-Alumina,$^{16}$ graphene oxide,$^{17}$ solid super acid,$^{18}$ etc. have been studied for the synthesis of PODE$_n$. However, only few catalysts are suitable for industrial application because of the unsatisfactory conversion rate and selectivity of PODE$_n$. Wu et al.$^{11}$ evaluated the catalytic performance of ZSM-5 molecular sieves with different Si/Al ratios for the synthesis of PODE$_n$, and they indicated that under the catalysis of ZSM-5 with the Si/Al ratio of 580, the conversion rate of TOX and selectivity of PODE$_n$ were 85.3% and 88.5%, respectively. Li et al.$^{18}$ prepared SO$_2$/Fe$_2$O$_3$/SiO$_2$ catalysts with varied acid strength and acid sites, by which the conversion rate of TOX and selectivity of PODE$_n$ were only 81.9% and 34.4%, respectively. Zhao and coworkers$^{10}$ compared the catalytic performance of HY, HZSM-5, Hf, and HMCM-22, and they indicated that HMCM-22 showed the best catalytic performance; however, the selectivity of PODE$_{3−8}$ was only 29%. Therefore, an effective
catalyst with high activity is urgently required for PODEₙ synthesis from methyal and trioxyxymethylene.

Sulfonic acid mesoporous SBA-15 catalyst as a type of excellent and efficient Brønsted acid catalyst has been applied in different fields such as ethylene polymerization, biodiesel synthesis, bio-oil upgrading, fatty acid esterification, glycerol acetylation, and dimethyl ether synthesis. In our previous research, we synthesized a series of mesoporous Al-SBA-15 molecular sieves with different Si/Al ratios and investigated the influences of acid strength, acid amount, acid type, and pore sizes on the PODEₙ synthesis. In this study, as a continuation of our previous study, a series of sulfonic acid-functionalized mesoporous SBA-15 with different sulfur loading were synthesized by a post-synthesis method. X-ray diffraction (XRD), N₂ adsorption-desorption, X-ray fluorescence (XRF), nuclear magnetic resonance (NMR), Fourier transform infrared (FT-IR) spectroscopy, elemental analysis, and acid–base titration were used to characterize the structure and acidity of the as-synthesized catalysts. Moreover, reusability performance of sulfonic acid-functionalized mesoporous SBA-15 was also studied herein.

2. Experimental

2.1. Catalyst preparation

Preparation of SBA-15. Pure silica SBA-15 materials were synthesized according to the procedure described by Zhao et al. Typically, 4 g P123 (Aldrich, EO₂₀₋₅₋PO₇₀₋₅₋EO₂₀) was dispersed in 30 ml of deionized water and 120 ml of 2 M HCl solution under sufficient stirring, followed by the drop-wise addition of 9 g TEOS (Aldrich, Si(OC₂H₅)₄) to the above-mentioned solution. The mixtures were continuously stirred at room temperature for 24 h and crystallized in a Teflon-lined autoclave at 100 °C. The mixtures were filtered, washed, dried at 100 °C for 12 h, and then calcined at 550 °C for 6 h.

Preparation of sulfonic acid-functionalized SBA-15. The procedure for the synthesis of organosilane-sulfonated SBA-15 was carried out as reported in a previous study. Typically, 1 g SBA-15 was dispersed in 30 ml of toluene and stirred for 10 min under a N₂ atmosphere, and specific amounts of 3-mercaptopropyltrimethoxysilane (0.2, 0.5, 0.8, 1, and 2 g per gram SBA-15) were added dropwise using an injector. The mixture solution was refluxed at 110 °C under a N₂ atmosphere for 24 h. Then, the mixture was filtered and washed with toluene and ethanol and dried at 60 °C for 12 h. The obtained solid was oxidized with 30 ml of aqueous H₂O₂ (30 wt%) at room temperature for 24 h under a N₂ atmosphere. The catalysts were filtered and washed with deionized water and ethanol and dried at 60 °C under vacuum for 12 h. The sulfonated SBA-15 was denoted as SO₃H-SBA-15(n), where n refers to the sulfur loading on the SBA-15 surface.

2.2. Catalyst characterization

X-ray diffraction (XRD) was carried out using an XPert Pro MPD equipment produced by Panalytical with CuKα radiation. The scan range was set between 0.5° and 5° for small angle measurements. N₂ adsorption-desorption experiments were carried out via a TriStar II instrument (micromeritics, USA) using nitrogen as the adsorption agent at −196 °C. The specific surface areas were calculated from the Brunauer–Emmett–Teller (BET) equation, and the total pore volumes were calculated by the BJH method. The BJH pore sizes were calculated from the desorption branch of the isotherms. The S, C, and H elemental analyses were carried out by a Thermo EA1112 Element Analyzer. X-ray fluorescence (XRF) was performed via a PANalytical Axios instrument. Fourier transform infrared spectra (FT-IR) were obtained using a Thermo Fisher Nicolet-6700 instrument. The ¹³C-NMR and ²⁹Si-NMR spectra were obtained using a Bruker Avance III HD 400 MHz spectrometer with the resonance frequencies of 75.5 MHz and 208.3 MHz, respectively.

Thermogravimetric analysis was carried out under a high-purity N₂ atmosphere at the heating rate of 10 °C min⁻¹. The acid amounts of the catalysts were detected via a titration method as follows: 0.1 g of SO₃H-SBA-15 catalyst was added to 20 ml of NaCl (2 M) aqueous solution; the mixture was stirred at room temperature for 12 h and then titrated with 0.05 M NaOH solution. 3.2. Catalytic performance test

Experiments for the catalytic performance tests were carried out using a 100 ml stirred autoclave reactor. DMM, TOX, and SO₃H-SBA-15 catalysts were quantitatively loaded into the reactor. The amounts of DMM and TOX were basically added according to the molar ratio from 2.5 : 1 to 1 : 2 and the amount of SO₃H-SBA-15 catalysts were 2 wt% of the total amount of the feedstock loading. The reaction temperature was set at 70–130 °C and controlled with ±1 °C of the set value via a modular controller; the reaction times were set from 20 to 120 min. The reaction pressure was fixed at 1 MPa and the agitation speed was set at 300 rpm.

Synthetic products were analyzed by an Agilent 7820 gas chromatograph with an FID detector. An Agilent HP-5 capillary column (30 m × 0.32 mm × 0.25 μm) was used. Chromatographic conditions were set as follows: injection port temperature, 280 °C; detector temperature, 280 °C; oven temperature program: initially the temperature was maintained at 32 °C for 5 min, then increased to 45 °C at the rate of 3 °C min⁻¹, and finally to 245 °C at the rate of 50 °C min⁻¹ and maintained for 5 min. The internal standard method was adopted as a quantitative method and octane was used as the internal standard.

To estimate the conversion of feedstocks and the yield and selectivity of PODEₙ, the calculation formulas were employed as follows: DMM conversion:

$$X_{\text{DMM}} = \frac{m_{\text{DMM, feed}} - m_{\text{DMM, product}}}{m_{\text{DMM, feed}}} \times 100\%$$

TOX conversion:

$$X_{\text{CH}_2\text{O}} = \frac{m_{\text{CH}_2\text{O, feed}} - m_{\text{CH}_2\text{O, product}}}{m_{\text{CH}_2\text{O, feed}}} \times 100\%$$
PODE$_n$ yield:

$$Y_{\text{PODE}_n} = \frac{m_{\text{PODE}_n, \text{product}}}{m_{\text{CH}_2\text{O}, \text{feed}}} \times 100\%$$

PODE$_n$ selectivity:

$$S_{\text{PODE}_n} = \frac{m_{\text{PODE}_n}}{\sum m_{\text{PODE}_n}} \times 100\%$$

where $m_{\text{DMM,feed}}$ and $m_{\text{CH}_2\text{O,feed}}$ is the feeding quality of methylyl and trioxymethylene, respectively; $m_{\text{DMM,product}}$ and $m_{\text{CH}_2\text{O,product}}$ is the quality of methylyl and trioxymethylene in the product, respectively.

3. Results and discussion

3.1. Structural features of the catalysts

The small angle XRD patterns of SBA-15 and SO$_3$H-SBA-15 with different sulfur loading are shown in Fig. 1. All the samples have three well-resolved diffraction peaks at about 0.92°, 1.55°, and 1.77°, corresponding to the (100), (110), and (200) reflections, respectively, which can be identified as typical two-dimensional hexagonal mesostructures (space group $p6m/m$). XRD patterns for SO$_3$H-SBA-15 catalysts did not change after the grafting process, evidencing that the two-dimensional hexagonal mesostructure was stable during the functionalization process.\(^{21}\)

The N$_2$ adsorption–desorption isotherms and pore size distribution of SBA-15 and SO$_3$H-SBA-15 are shown in Fig. 2. It can be seen that both the SBA-15 and SO$_3$H-SBA-15 catalysts show type IV isotherm with H1 hysteresis capillary at 0.65 < $P/P_0$ < 0.80 due to the presence of mesoporous structure and have narrow pore size distribution, which centralizes at about 6.5 nm.

The physical properties of SO$_3$H-SBA-15 catalysts are shown in Table 1. The acid amounts are proportional to the surface area, pore volume, and pore sizes of the as-synthesized materials gradually decreased with the increase of sulfur loading, indicating that the grafting reactions not only proceeded on the surface of the catalyst, but also occurred inside of the pore and caused a certain degree of plugging phenomenon. The sulfur loadings of SO$_3$H-SBA-15 catalysts, which are shown in Table 1, were determined by elemental analysis and further confirmed via XRF detection. It can be seen that the sulfur content on the surface of SBA-15 supports gradually increased with the increase of silane concentration, and the actual measured sulfur contents are always lower than that of the feeding amount used for the grafting, which is consistent with a previous report.\(^{31}\)

Acid amounts of SO$_3$H-SBA-15 catalysts with different sulfur loading are shown in Table 1. The acid amounts are proportional to the sulfur content of the catalyst and gradually increased from 0.056 to 0.443 mmol g$^{-1}$.

Infrared spectrum detection was used as a qualitative method to identify whether silane was successfully grafted on the silica surface.\(^{31,34}\) The strong and wide absorption band at 1093 cm$^{-1}$ can be attributed to the Si–O–Si antisymmetric stretching vibration (Fig. 3). The bands at 809 cm$^{-1}$ and 460 cm$^{-1}$ are assigned to the Si–O stretching vibration and bending vibration, respectively. The peak at 958 cm$^{-1}$ belongs to the bending vibration absorption peaks of the Si–OH bond. The attachment of silane on the silica surface was identified by the bands at 2940 cm$^{-1}$ and 2854 cm$^{-1}$, which are assigned to the asymmetric and symmetric stretching peaks of methylene, and the band at 1460 cm$^{-1}$ can be attributed to the methylene symmetric bending vibration.

The incorporation of silane on the silica surface was further confirmed by $^{29}$Si MAS NMR and $^{13}$C-MAS NMR.\(^{31,35-37}\) The $^{13}$C
MAS NMR spectra of SO$_3$H-SBA-15(0.8) (Fig. 4(A)) exhibit the signals at 12.77, 17.52, and 55.39 ppm, which are assigned to the C1, C2, and C3 carbon peaks, respectively. No peak can be found at about 23 and 41 ppm, indicating that no disulphide was formed during the preparation. Fig. 4(B) shows the 29Si MAS NMR spectra of SO$_3$H-SBA-15(0.8). The two distinct resonances at $\gamma$/C0 112 ppm and $\gamma$/C0 103 ppm refer to Q4 [Si(OSi)$_3$] and Q3 [HOSi(OSi)$_3$], respectively. The spectrum of SO$_3$H-SBA-15(0.8) shows another two resonances at $\gamma$/C0 67 ppm and $\gamma$/C0 59 ppm, which illustrate that the Si–C bonds were successfully generated, and the organic functional groups were incorporated into the frameworks.

3.2. Catalytic performance

The catalytic performance of SO$_3$H-SBA-15 catalysts with different sulfur loading for the synthesis of PODE$_n$ is presented in Fig. 5. There is a significant influence of the sulfur loading on the conversion rate of feedstock DMM and TOX and the selectivity and yield of main product PODE$_2$–8. It can be seen from Fig. 5a that the conversion of DMM and TOX and the yield of PODE$_2$–8 first increased and then decreased when the sulfur loading increased from 0.2 to 2 wt% and reached maximum values of 51.84, 95.57, and 61.86%, respectively, under the catalysis of SO$_3$H-SBA-15(0.8). In addition, as shown in Fig. 5b, with the increase of sulfur loading, the selectivity of by-product methyl formate gradually increased and reached its highest value under the catalysis of SO$_3$H-SBA-15(2.0), which could be attributed to the highest acid amount of SO$_3$H-SBA-15(2.0), as illustrated in Table 1. Therefore, the SO$_3$H-SBA-15(0.8) catalyst has the best catalytic performance for PODE$_n$ synthesis.

It is well known that the production process of PODE$_n$ can be roughly summarized as two steps: depolymerization of trioxymethylene into formaldehyde monomer and condensation reaction of formaldehyde monomer with DMM and PODE$_n$. Moreover, the excessive formaldehyde monomer in the acidic environment will cause the generation of the by-product methyl formate in a Tischenko reaction. It was reported that the trioxymethylene dissociation process was a crucial step and depolymerization and transformation situation of trioxymethylene was related with the catalyst acidity, which directly

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>Specific surface area m$^2$ g$^{-1}$</th>
<th>Pore volume cm$^3$ g$^{-1}$</th>
<th>Pore diameter nm</th>
<th>S content$^a$ wt%</th>
<th>S content$^b$ wt%</th>
<th>Amount of acid sites mmol g$^{-1}$</th>
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<tr>
<td>SBA-15</td>
<td>747</td>
<td>1.04</td>
<td>5.92</td>
<td>—</td>
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<tr>
<td>SO$_3$H-SBA-15(0.2)</td>
<td>658</td>
<td>1.02</td>
<td>5.89</td>
<td>0.18</td>
<td>0.16</td>
<td>0.056</td>
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<tr>
<td>SO$_3$H-SBA-15(0.5)</td>
<td>639</td>
<td>0.98</td>
<td>5.86</td>
<td>0.42</td>
<td>0.43</td>
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<tr>
<td>SO$_3$H-SBA-15(0.8)</td>
<td>625</td>
<td>0.95</td>
<td>5.85</td>
<td>0.53</td>
<td>0.54</td>
<td>0.166</td>
</tr>
<tr>
<td>SO$_3$H-SBA-15(1.0)</td>
<td>618</td>
<td>0.92</td>
<td>5.81</td>
<td>0.84</td>
<td>0.89</td>
<td>0.262</td>
</tr>
<tr>
<td>SO$_3$H-SBA-15(2.0)</td>
<td>609</td>
<td>0.92</td>
<td>5.79</td>
<td>1.42</td>
<td>1.45</td>
<td>0.443</td>
</tr>
</tbody>
</table>

$^a$ S content detected from the elemental analysis. $^b$ Calculated by XRF.

Fig. 3 The Fourier transform infrared spectra of SBA-15 and SO$_3$H-SBA-15 catalysts.

Fig. 4 (A) 13C-NMR and (B) 29Si-NMR spectra.
affected the further condensation reaction and side reaction. Fig. 6 shows the reaction and depolymerization tendency of TOX in the presence of catalysts with different acid amounts, which can clearly explain the reaction rules. It can be seen that under the catalysis of SO₃H-SBA-15(0.2) and SO₃H-SBA-15(0.5) catalysts, which have relatively weak acidity, trioxymethylene was insufficiently depolymerized; therefore, the corresponding concentration of trioxymethylene, which was converted to the main product PODE₂₈ and byproduct MF, was relatively low. Under the catalysis of SO₃H-SBA-15(0.8), which has moderate acidity, the undecomposed trioxymethylene in the product reached a minimum value and at the same time the depolymerized trioxymethylene, which was converted to PODE₉, reached a maximum value. With the further increase of catalyst acidity, the depolymerized trioxymethylene, which was converted to the byproduct methyl formate, obviously increased, indicating that the depolymerization rate of trioxymethylene is far greater than its condensation rate with methylal in the presence of the catalyst with large amounts of acid, and accordingly, the excessive formaldehyde monomer generates methyl formate. Moreover, the trioxymethylene content in the product also increased under the catalysis of SO₃H-SBA-15(1.0) and SO₃H-SBA-15(2.0), illustrating that the excessive amount of acid will not only promote the occurrence of the side reaction, but also inhibits the depolymerization reaction of trioxymethylene, or accelerates the repolymerization of formaldehyde monomers into trioxymethylene. It can be concluded that the depolymerization of TOX requires a relatively moderate acid amount as an insufficient or excessive amount of acid will lead to incomplete depolymerization.

PODEₙ with the structure of CH₃O–(CH₂O)ₙ–CH₃ is a mixture of different degrees of polymerization, and with the increase of the degree of polymerization, the cetane number, which is one of the most important indexes for diesel oil, gradually improve. However, a high degree of polymerization (n > 8) will cause the increase of pour point and the cold filter plugging point of the mixture, which is not suitable to be used as a diesel fuel blending component and will need to be returned to the synthetic unit. Moreover, the PODE₂ component does not meet the security criterion of the diesel fuel due to its low boiling point and low flash point; therefore, it cannot be used as a diesel blending component either. Thus, among PODEₙ products, relatively more PODE₃ₘ–₈ should be generated, and PODE₂ and PODEₙₘ–₈ components should be largely suppressed. The effects of sulfur loading on the product distribution and chain length are shown in Fig. 7. It can be clearly seen that the selectivity of PODE₂₈ with different polymerization and chain length obviously varied with the change of sulfur loading. With the increase of sulfur loading, the selectivity of PODE₂₈ increased first and then decreased, and on the SO₃H-SBA-15(0.8) catalyst, the selectivity of PODE₅ₘ–₈ reached the highest value among all the catalysts. In addition, the maximum chain length of PODEₙ, obviously increased with the increase of sulfur.
loading. Under the catalysis of SO\textsubscript{3}H-SBA-15(0.2) catalyst, only PODE\textsubscript{2–6} were generated and the product with the polymerization degree of \( n > 7 \) was not formed. On the SO\textsubscript{3}H-SBA-15(0.5) catalyst, PODE\textsubscript{2} can be detected, and furthermore, PODE\textsubscript{10} was generated under the catalysis of SO\textsubscript{3}H-SBA-15(2.0). Fortunately, only the desired components PODE\textsubscript{3–8} were formed under the catalysis of SO\textsubscript{3}H-SBA-15(0.8). This result fully explains that the acidity of the catalysts also has a decisive effect on the product distribution and the chain length of the products. Thus, it is also an effective way to obtain the desired product distribution by adjusting the acidity of the catalyst.

From a thermodynamic point of view, the ring opening depolymerization reaction of trioxymethylene is endothermic; thus, a relatively higher temperature is in favor of the depolymerization of trioxymethylene. However, the condensation reaction of formaldehyde and methylal is conversely a slightly exothermic reaction. Thus, it is very important to select an appropriate temperature to achieve the maximum efficiency of the reaction. The influence of temperature on the conversion rate of DMM and TOX and selectivity of PODE\textsubscript{n} and by-product is shown in Fig. 8. With the increase of temperature, the conversion rate of TOX gradually increased, indicating that a higher temperature will accelerate the depolymerization process of TOX. A relatively low temperature will lead to the incomplete depolymerization of TOX. However, the selectivity of the byproduct MF also increased with the increase of temperature, especially after a temperature greater than 100 °C. This is probably due to the fact that the depolymerization rate of trioxymethylene is far greater than its condensation rate with methylal at high temperature; thus, the competitive reaction of the formation of the byproduct methyl formate is greatly promoted. Therefore, considering the feedstock conversion rate and product selectivity, 100 °C is the most appropriate temperature for the PODE\textsubscript{n} synthesis.

The molar ratio of feedstocks has a great influence on the conversion rate of the feedstocks and the yield of the products. It can be seen that increasing the amount of trioxymethylene is beneficial for enhancing the yield of PODE\textsubscript{2–8} (Fig. 9a) and improving the conversion rate of DMM; however, the conversion rate of TOX was gradually reduced due to the limit of the equilibrium of the reaction. In addition, as shown in Fig. 9b, the product distribution and chain length obviously varied with the change of the molar ratio of DMM and TOX. With the increase of TOX proportion, the selectivity of PODE\textsubscript{2} decreased, whereas the selectivity of PODE\textsubscript{3–8} gradually increased, indicating that increasing the amount of TOX is in favor of promoting the reaction to generate products with a higher degree of polymerization. When the DMM/TOX molar ratio was 1 : 2, the product with the chain length of 10 was generated. With the increase of DMM proportion, the high degree of polymerization selectivity gradually decreased, and when the molar ratio of DMM/TOX reached 1 : 1, no product of \( n > 8 \) was formed. Therefore, to avoid the generation of products with a high degree of polymerization, a relatively large proportion of methylal was required. Although with the change of the DMM/TOX molar ratio from 2 : 1 to 1 : 2, the yield of PODE\textsubscript{2–8} gradually decreased, whereas the conversion rate of TOX significantly increased. Thus, a relatively large proportion of methylal will not only inhibit the generation of high degree of polymerization products, but also greatly improves the conversion rate of TOX. Therefore, to avoid the generation of a high degree of polymerization products, it is necessary to lose a certain yield of PODE\textsubscript{2–8}. Considering the yield of PODE\textsubscript{n} and conversion of TOX under the premise of no generation of PODE\textsubscript{n} in which \( n > 8 \), 1 : 1 is the optimum molar ratio of DMM and TOX in the reaction system.

The yield of PODE\textsubscript{n} and the conversion rate of DMM and TOX all have a significant increase in the reaction for the first 60
minutes, which is shown in Fig. 10. Because the depolymerization process of TOX is relatively slow, a certain period of time is required to achieve the balance of trioxymethylene depolymerization. However, when the reaction time is prolonged to 90 min and 120 min, the conversion rate of feedstocks and the yield of the products are essentially at compared with those at 60 min, which shows that the reaction system has basically reached chemical equilibrium and cannot be promoted by extending time. Thus, 60 min is the most appropriate time for PODe$_n$ synthesis.

The catalytic performance of SO$_3$H-SBA-15(0.8) catalyst was further evaluated by comparing with the reference catalysts such as Amberlyst-15 sulfonic acid resin, ZSM-5 zeolite with Si/Al ratio of 200, and HY zeolite with the Si/Al ratio of 5.3. It can be seen from Fig. 11 that the conversion rate of TOX and the PODe$_n$ yield and selectivity under the catalysis of SO$_3$H-SBA-15(0.8) are obviously higher than those of Amberlyst-15 sulfonic acid resin, and ZSM-5 and HY zeolite. One reason for the better performance of the SO$_3$H-SBA-15(0.8) catalyst is attributed to the enhanced mesoporosity, which would overcome the diffusion limitation and promote the transformation of the reactants and products in the two-dimensional hexagonal mesostructure. Another more important reason is probably ascribed to the moderate acidity of the SO$_3$H-SBA-15(0.8) catalyst. We know that the depolymerization of TOX requires a relatively moderate acid amount; insufficient or excessive amount of acid will promote the formation of the by-product methyl formate and lead to the incomplete depolymerization of TOX, thus affecting the conversion of TOX and yield of PODe$_n$. There is a relatively high acid amount in the pores of Amberlyst-15 sulfonic acid resin, ZSM-5 zeolite, and HY zeolite, which leads to the far greater depolymerization rate of TOX than its condensation rate with methylal; therefore, a large number of methyl formate is generated and the depolymerization of TOX is suppressed.

The catalytic stability of the as-synthesized catalyst, which is an important index for selecting the catalysts, was evaluated in the PODe$_n$ synthesis. SO$_3$H-SBA-15(0.8) catalyst was used to evaluate the reusability. After completing a catalytic reaction, SO$_3$H-SBA-15(0.8) was separated from the reaction system by filtration and was dried in vacuum at 60 °C for 12 h and then used for the next reaction. As illustrated in Fig. 12, the activity of the catalysts only slightly declined after being reused six times; the conversion rate of TOX decreased from 95.57% to 93.09%.

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and the yield of PODE$_{2-8}$ also decreased from 61.86% to 59.13%. Moreover, the as-synthesized and the 6th run recycled SO$_3$H-SBA-15(0.8) catalyst were characterized by elemental analysis and further verified by TGA (Fig. 13), showing that the sulfur content of the catalyst slightly reduced from 0.53% to 0.51%.

4. Conclusion

Herein, SO$_3$H-SBA-15 catalysts with different sulfur loading were successfully prepared for the synthesis of PODE$_n$ from DMM and TOX. In the study of the effect of different sulfur loading (acid property) on the catalytic performance, it was found that a moderate amount of acid was required for PODE$_n$ synthesis, as proven by SO$_3$H-SBA-15(0.8) catalyst, which showed the best catalytic performance, in that the conversion rate of DMM and TOX and the yield of PODE$_{2-8}$ reached high values of 51.84, 95.57, and 61.86%, respectively. Inadequate or excessive amounts of acid will lead to the incomplete depolymerization of trioxymethylene. Moreover, the excessive amount of acid can cause the formation of a large number of by-product, methyl formate. In addition, the acidity of the catalysts also had an effect on the product distribution and the chain length of the products. The optimum reaction conditions were investigated; through the comparison of reaction temperature, it can be concluded that a relatively low temperature ranging from 70 to 90 °C will lead to the incomplete depolymerization of trioxymethylene, and a higher temperature ranging from 110 to 130 °C will accelerate the depolymerization process of TOX and greatly promote the formation of the by-product methyl formate; thus, 100 °C was the most appropriate temperature. By adjusting the molar ratio of DMM and TOX, it was found that an increasing amount of TOX was in favor of improving the yield of PODE$_n$ and promoting the generation of the products with a higher degree of polymerization. Considering a higher product yield and reasonable product chain length, 1 : 1 was the optimum molar ratio of DMM and TOX, at which no PODE$_{2-8}$ was generated. The reaction time analysis demonstrated that the reaction system had basically reached chemical equilibrium within 60 min and the reaction could not be promoted by further extending time. Moreover, according to the comparison of SO$_3$H-SBA-15(0.8) catalyst and the reference catalysts, it can be concluded that the SO$_3$H-SBA-15(0.8) catalyst had better catalytic activity than Amberlyst-15 sulfonic acid resin, and ZSM-5 and HY zeolite for PODE$_n$ synthesis. The activity of SO$_3$H-SBA-15(0.8) catalyst just slightly declined after being reused six times, showing good application prospects.

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Notes and references