Unsymmetrical difunctionalization of cyclooctadiene under continuous flow conditions: expanding the scope of ring opening metathesis polymerization†

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Functionalized cyclooctenes (FCOEs) are important monomers in ring-opening metathesis polymerization (ROMP). Herein, a new library of disubstituted FCOEs bearing adjacent heteroatoms were synthesized and applied in ROMP. To address the issues associated with the handling of the reactive thienyl chloride intermediate, a two-step continuous flow method has been developed to prepare 5-thio-6-chlorocyclooctene compounds from abundant cyclooctadiene starting materials. These newly synthesized FCOE monomers were subsequently polymerized through ROMP, giving rise to a range of functionalized polymers with high molecular weights. Furthermore, we demonstrated that the thermal properties of these polymers could be fine-tuned by changing the functional groups in the FCOE monomers. We expect that this functionalization-polymerization strategy will enable the preparation of a range of polymeric materials with complex structures.

Introduction

The development of synthetic methods to access functionalized polymers is of considerable interest due to the interesting physical and chemical properties associated with these materials. As a result, extensive efforts have been made to accomplish this task by designing well-tailored monomers for different synthetic methods, such as controlled radical polymerization1 and ring-opening metathesis polymerization (ROMP).2 Alternatively, a number of methods for the postsynthetic modification of polymers have also been developed.3 Due to the robustness and functional group tolerance of ROMP, it has become one of the most powerful methods for accessing polymers bearing a wide range of functionalities,4 thus enabling the development of materials for drug delivery,5 the manipulation of liquids,6 ion exchange7 and other uses.8 While this method is widely utilized, the most frequently used monomers are norbornene, cyclobutene and cyclooctadiene.9 A simple method that could provide cyclic olefins with various substituents is important for expanding the scope of functionalized polymers.

FCOE derivatives are a class of the most widely used monomers for ROMP.5,8a–c,9 Among the many applications of poly (FCOE)s,5,8a,9 ROMP of FCOEs followed by hydrogenation yields linear polyolefins with well-defined chemical structures possessing a wide range of side chains.5,8a–c This represents a useful approach to high-precision functionalized polyolefins,5,8a–c which are otherwise difficult to synthesize.10 To further explore the utility of ROMP, it is necessary to expand the scope of the FCOEs. Thanks to the efforts devoted to catalyst development and monomer scope exploration, a variety of FCOEs have shown high reactivity in ROMP.5,8b–c,11 Among these, most examples are of mono-substituted compounds (Fig. 1A) prepared via C=C bond addition of cyclooctadienes (CODs).5,8b,c–f,11 allyl C–H bond functionalization of cyclooctenes (COEs).5,8f or other methods.9,6

In contrast to monosubstituted FCOEs, polysubstituted FCOEs are much less investigated for ROMP reactions.5,12 Grubbs and coworkers reported the synthesis and ROMP of symmetrically disubstituted COEs connected with two adjacent hydroxyl groups and their derivatives (Fig. 1B).12a,12c Hillmyer and coworkers reported the preparation and ROMP of ester and methyl/phenyl disubstituted COEs (Fig. 1C).12d Nuyken found that the polymerization of diacyano COEs is sluggish, while the monocyano COE polymerizes efficiently.12d However, the ROMP
of FCOEs possessing different vicinal heteroatoms (Fig. 1D) has not been reported so far. The incorporation of these functional side chains could not only allow for the fine tuning of polymer properties, but also open up new opportunities to introduce orthogonal reactive sites, and is thus highly desirable.

In this regard, we have designed a two-step sequence of thienyl chloride formation/C≡C bond addition to prepare FCOEs from cis,cis-1,5-COD (Fig. 1D: X = SR, Y = Cl). Since the chloride group is easily cleavable through the assistance of the adjacent thioether via neighboring group participation,14 we envisioned that the 5-Cl,6-SR-COE would be a versatile intermediate to prepare FCOEs with different functionalities (Fig. 1D: X = SR, Y = OR2/NR2). Although the thienyl chloride (RSCI) species has been known for over half a century, the explosive nature14 and unpleasant smell of these compounds somewhat limits their application. Flow processes are useful alternatives to traditional batch procedures.15 Many examples have shown the possibility to safely handle hazardous intermediates under flow conditions.16 Given our experience with this technique,17 we anticipated that a flow approach would significantly enhance the practicality of olefin chlorothiolation processes using thienyl chloride by allowing for the safe and convenient handling of these reactive intermediates.

Results and discussion

We began our studies on the thienyl chloride intermediate formation/difunctionalization sequence with the setup depicted in Scheme 1A with p-toluenethiol 1a as the model substrate. In the flow setup, a solution of 1a in anhydrous dichloromethane (DCM) was mixed with SO2Cl2 in anhydrous DCM and introduced into a tubing reactor (R1) immersed in a cooling bath. After the arylthiol was completely converted, as monitored by thin layer chromatography (TLC) analysis, R1 was assembled with the following setup of step II via a T-mixer, allowing the solution from R1 to combine with the COD (3) solution in-line. The resultant mixture was further delivered into the second tubing reactor (R2), which was submerged in another cooling bath, to perform the direct difunctionalization of the C≡C double bond. After the reaction, the mixture was collected and directly analyzed without the isolation of 4a. Upon investigating a variety of reaction parameters, we determined that the synthesis of 4a proceeded in good yield with a 1/1.05/4 ratio of 1a/2/3, and two reactors cooled at 0 °C and −20 °C respectively (Scheme 1B, entry 1). Notably, this two-step flow method only needed a residence time (tR) of less than 4 min.18 As shown in entries 2 to 7, changing the temperature of either reactor or the molar ratio of the three components resulted in a lower yield of the target product 4a (see Section II in the ESI†). In contrast, when this reaction sequence was performed under batch conditions only 50% yield of 4a was obtained in 2.5 h of reaction time, as detected by 1H NMR analysis.

Following the two-step flow synthesis (Table 1, step I & II), the solution of compound 4a was directly added into a vial with anhydrous methanol at room temperature (Table 1, step III). The Cl group on 4a was efficiently replaced by a OMe group under mild conditions within several hours, as monitored by TLC analysis. The resultant mixture was purified by silica gel column chromatography to afford FCOE 5a in 64% yield over three steps. In comparison, when 5-Cl-1-cyclooctene19 was reacted with MeOH at room temperature for 48 h instead of 4a, no substitution product was detected by LC-MS, supporting our hypothesis of a vicinal SR group assisted substitution process.14

With the method established for the preparation of 5a, we turned our attention to the synthesis of FCOEs with different
SR1 substituents. To our delight, all the R'SH substrates (1) investigated in Table 1 underwent complete conversion to 5b–5g in about 4 h of reaction time (Table 1, step I to III). After the consecutive three-step transformations, the resultant mixtures were purified by silica gel column chromatography to afford the FCOEs 5b–5g in satisfactory yields (55–70%). Notably, since aryl halides (e.g. Cl and Br) are versatile functional groups in metal-catalysed cross-coupling reactions, the incorporation of such groups (5c and 5d) would bring in reactivity orthogonal to the substituent on the COE backbone. All FCOE monomers were characterized by nuclear magnetic resonance (NMR), infrared radiation (IR), and high-resolution mass spectroscopy (HRMS) analysis (Section III and Fig. S3–S23†), demonstrating the successful introduction of the two adjacent heteroatom substituents SR1 and OMe into the COEs.

Moreover, to streamline the synthesis of the FCOEs 5, a three-step continuous-flow setup has been developed (Fig. S2†) using a pressurised heating system at 80 °C for step III. As exemplified with 5a, the reaction time was reduced to 20 min, facilitated by the efficient heat transfer under the flow conditions, affording 5a in 66% isolated yield.

The 5-SR1,6-OMe-COE monomers 5a–5g were polymerized with the second-generation Grubbs carbene complex (G2) in DCM at room temperature (step IV). As illustrated in Table 1, full conversions of all monomers upon G2-catalyzed ROMP was achieved when the arylthio group was substituted with an electron-donating group (Me, entry 1, 5a), an electron-withdrawing group (F, entry 2, 5b; Cl, entry 3, 5c; Br, entry 4, 5d), or a phenyl group (entry 5, 5e), affording a variety of functionalized polymers in high yields (6a–6f: 90–96% yields) following isolation via a three-time precipitation from methanol. Similar to the Ru-promoted ROMP of alkylthio monosubstituted COEs reported by Noels and coworkers, when 5-S1012H12S-6-OMe-COE (5g, entry 7) was used a decreased polymerizing reactivity was observed, providing 6g ([Mn/Mn] = 311 kg mol⁻¹, Mw/Mn = 1.67) with 36% monomer conversion in 48 h of reaction time. This is probably due to the increased coordinating effect of an alkylthio group to the metal center compared
Table 2  Synthesis and ROMP of 7a–7h

<table>
<thead>
<tr>
<th>Entry</th>
<th>R1</th>
<th>R3</th>
<th>7 yield b (%)</th>
<th>7 conv c (%)</th>
<th>8 yield d (%)</th>
<th>(M_n,\text{calc} e) (kg mol(^{-1}))</th>
<th>(M_n,\text{GPC} f) (kg mol(^{-1}))</th>
<th>(\bar{D}^g)</th>
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<tr>
<td>1</td>
<td>2</td>
<td></td>
<td>67 (7a)</td>
<td>&gt;99</td>
<td>41</td>
<td>20 (8a)</td>
<td>51</td>
<td>80</td>
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<tr>
<td>2</td>
<td>67 (7a)</td>
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<td>3 i</td>
<td></td>
<td></td>
<td>63 (7b)</td>
<td>&gt;99</td>
<td>81 (8b)</td>
<td>6.2</td>
<td>6.8</td>
<td>1.71</td>
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<tr>
<td>4 j</td>
<td>64 (7c)</td>
<td></td>
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<td>5</td>
<td>65 (7c)</td>
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<td>6</td>
<td>62 (7d)</td>
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<td>7</td>
<td>54 (7e)</td>
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<td>8</td>
<td>59 (7f)</td>
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\(R_1SH\) was synthesized using the optimized conditions shown in Scheme 1B, entry 1; (III) silica gel; (IV) 7b and 7c: TBSCI, imidazole, DMAP, DCM, 0 °C to rt; 7d: BnBr, NaH, 0 °C to rt; 7e: AcOH, DCC, DMAP, DCM, 0 °C to rt; (V) 7f: morpholine, rt; 7g \(\text{N}_(n\text{Bu})_4\text{N}^+\), rt; 7h: imidazole, rt; (VI) G2 was used to initiate the ROMP, \([M]/[G2] = 500/1\), room temperature. \(a\) Isolated yields of three steps (7a and 7f) or four steps (7b–7e), calculated based on the recovered monomer by column chromatography. \(b\) Isolated yields were calculated based on the monomers added in the ROMP. \(c\) Calculated based on the conversions of 7. \(d\) Analyzed by GPC. \(e\) X-ray structure of 7f. \(f\) \([M]/[G2] \geq 20/1.\) \(g\) \([M]/[G2] = 200/1.\) \h Reaction temperature = 45 °C.
connected to a eluent. During the column chromatography process, di were successfully polymerized. We further focused on expand-

to each other. This is consistent with the vicinal SR group con

The newly synthesized FCOE monomers (7a–7h) were next polymerized in the presence of G2 at room temperature (Table 2, step VI). When FCOE 7a with an unprotected hydroxy group was employed in a [7a]/[G2] ratio of 500/1, less than 50% conversion was achieved in 48 h of reaction time, providing 8a in 20% isolated yield (M_{n,GPC} = 80 kDa mol^{-1}, entry 1). Although decreasing the monomer/G2 ratio to 20/1 led to complete monomer conversion within 24 h, 8a′ with a much lower M_{n,GPC} of 6.8 kDa was provided (entry 2), with a D value similar to 8a (for 8a, D = 1.66, for 8a′, D = 1.65). We hypothesized that the improved monomer conversion was due to less of the transition-metal being poisoned by increasing the G2/monomer ratio. When the reaction temperature was increased from room temperature to 45 °C, poly(FCOE)s were generated with a similar M\n and slightly improved control over the molecular weight distributions (entry 3, M_n = 109 kDa and D = 1.71 vs. entry 4, M_n = 106 kDa and D = 1.62). When the third-generation of Grubbs carbene complex (G3) was used to initiate the ROMP of 7b ([7b]/[G3] = 200/1) at room temperature, the corresponding polymer was produced with D = 1.65 and M_n = 94 kDa at ~99% conversion.

To produce poly(FOE)s with high molecular weights, a monomer/G2 ratio of 500/1 was used during the ROMP reaction of the other FCOEs. When 7c–7e were applied in the ROMP for 24 h, isolated yields of 45–78% were obtained for the polymers 8c–8e with M_{n,GPC} values of 71–160 kDa (entries 5–7). When the SR\(^1\) group was adjacent to a morpholine group instead, polymer 8f was isolated in 82% yield (M_{n,GPC} = 193 k g mol^{-1}, entry 8). Both NMR and IR analyses clearly demonstrate that both types of functional group have been successfully incorporated in polymers 8a–8f (Section VI and Fig. S83–S112†). Replacing the morpholine group with an azide or an imidazole group provided less than 5% monomer conversion, which is probably caused by the irreversible coordin-

ation of the functional group to the Ru-center, as observed by Noels and coworkers.\(^{14}\) Notably, these represent the first ROMPs of FCOEs possessing adjacent substituents of SR\(^1\) and OR\(^2/\)NR\(^2\) functionalities.

To investigate the ROMP of difunctionalized FCOEs at different monomer/G2 ratios, 5a and 7f were employed. As shown in Fig. 2, when the [M]/[G2] ratios were increased from 20/1 to 500/1 for both monomers, poly(FCOE)s were produced with different M_{n,GPC} values, while the D values stayed at a similar level (D = 1.47–1.71 in Fig. 2A, D = 1.52–1.78 in Fig. 2B). Notably, a linear increase of M_{n,GPC} vs. [M]/[G2] was observed for both examples, which demonstrated that these poly(FCOE)s can be generated at the desired M_n by choosing a proper [M]/[G2] ratio within the investigated range.\(^{22}\)

The thermal properties for the polymers 6a–6g and 8a–8f were analyzed by differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA). The summarized results of their glass-transition temperature (T_g) and decomposition temperature (T_d) are shown in Fig. 3\(^{23,24}\) (for the DSC and TGA
profiles, see Section IV and VI of the ESI†). From 6a to 6g, while keeping the MeO group constant, changing the alkylthio side chains to arylthio chains resulted in polymers possessing increased \( T_g \) values (6g: \(-56^\circ\)C vs. 6a–6f: \(-12^\circ\)C to 35°C). Among 6b–6g, an increased functional group size on the aryl ring (from 6b to 6e: \(-12^\circ\)C, \(-3^\circ\)C, 13°C and 35°C respectively) or an increased degree of conjugation (e.g., 6f: 26°C vs. 6a: 4°C) led to increased \( T_g \) values. These results are in agreement with the sidechain influence on the glass transition temperature, as observed by others.9,24,26 For the polymers 8b–8e, when the hydroxy side groups were protected with groups larger than methyl, the resultant \( T_g \) values were higher than 6b (8b–8e: 0–18°C vs. 6b: \(-12^\circ\)C). Replacement of the MeO group with a morpholine group also led to an increased glass-transition temperature (8f: 45°C vs. 6a: 4°C). The thermogravimetric analysis in Fig. 3 shows that the thermal stabilities of these polymers are also dictated by the connection of different functional groups. Polymers 6a–6g and 8a–8f possess \( T_d \) values ranging from 225°C to 350°C at 5% weight loss.

Finally, the hydrogenation of polymer 6e was conducted to demonstrate the preparation of linear polyolefins possessing two different side chains on every seventh and eighth backbone carbon, from the corresponding poly(FCOE)s. The hydrogenation reaction was performed using p-toluene sulfonohydrazide as the reductant and tri-\( n \)-hexylamine as the base with a catalytic amount of 2,6-di-\( t \)-butyl-4-methylphenol (BHT) in \( o \)-xylene solvent.\(^{14,25}\) The reduced product 9 was obtained in 88% isolated yield via precipitation from methanol. As shown in the \(^1\)H NMR spectra (Fig. 4A1 and A2; Section VIII and Fig. S104–S108†), during the hydrogenation process, the signals found between 5.3–5.5 ppm corresponding to H\(_8\) and H\(_9\) of polymer 6e are completely absent in the spectrum of polymer 9. As a result, an increase in the signal region corresponding to alkyl protons is clearly observed for polymer 9 (Fig. 4A1 vs. 4A2 in the 1.0–2.5 ppm region), indicating the successful hydrogenation transformation. The GPC analyses of 6e and 9 (Fig. 4B) show: (1) similar \( M_n,GPC \) and \( M_w/M_n \) values, and (2) no new shoulder peaks in the GPC traces, suggesting that the polymer backbone remains intact during the reduction process. Moreover, the hydrogenated polymer 9 has a lower \( T_g \) value than 6e (Fig. 4C), indicating that the formation of a saturated backbone results in a higher molecular mobility. Hillmyer\(^*\) and Tanaka\(^*\) have also reported a decrease in the \( T_g \) values upon hydrogenating the corresponding poly(FCOE)s.

**Conclusions**

In conclusion, the synthesis and ROMP of FCOEs bearing adjacent heteroatom groups have been successfully realized. Notably, the unstable thienyl chloride species has been generated and used under flow conditions for the first time, allowing for an efficient synthesis of 5-SR,6-Cl-COE compounds, which were employed as versatile intermediates for the preparation of a library of FCOEs. Moreover, the ROMP of these new cyclic monomers has produced a library of polyolefins with different substituents connected by S, O or N heteroatoms in high molecular weights. This represents a useful avenue to synthesize polymers with a high level of complexity. The investigation of the thermal properties of these functionalized polymers has shown the effect of the side chains on their glass-transition temperatures and thermal stabilities. Finally, this approach complements the useful strategy of producing high precision model polyolefins via ROMP, allowing the preparation of terpolymers of ethylene, vinyl thioether, and a variety of polar...
olefins including vinyl ethers, vinyl esters and vinyl amines, which are inaccessible via other methods.

**Experimental**

The experimental procedure for the preparation of 5a with the optimized reaction conditions: a syringe was loaded with a solution of p-toluenethiol 1a (1.0 M, flow rate = 250 µL min⁻¹) in anhydrous DCM, and fitted to the syringe pump. Another syringe was loaded with a solution of 2 (1.05 M, flow rate = 250 µL min⁻¹) in anhydrous DCM, and fitted to a same syringe pump. The third syringe was loaded with a solution of COD (0.5 M, flow rate = 2.0 mL min⁻¹) in anhydrous DCM, and fitted to the second syringe pump. Following the setup as shown in Scheme 1, the solutions of 1a and 2 were mixed and reacted in the tubing reactor R1 (volume = 1.0 mL, t₁ = 2.0 min) submerged in a cooling bath. When the reaction was complete, the resultant solution was mixed with the solution of COD and reacted in the tubing reactor R2 (volume = 5 mL, t₂ = 2.0 min) submerged in another cooling bath. After the reaction, the resultant mixture was passed through a back-pressure regulator (BPR, 20 psi) before collection. After reaching steady state (waiting for 12 min), 1.0 mmol samples (10 mL reaction solution) were collected into an oven-dried vial equipped with a stir bar.

Anhydrous MeOH (10 mmol) was added into the vial via syringe at room temperature. When the reaction was completed, as monitored by TLC analysis, the mixture was treated with DCM (150 mL) and NaHCO₃ saturated aqueous solution (20 mL). The separated organic layer was washed with brine two times (2 × 10 mL), dried over Na₂SO₄ and then concentrated under vacuum. The residue was purified by column chromatography (eluting with 0-2% EtOAc in petroleum ether) to afford 5a in 64% isolated yield.

An oven-dried vial equipped with a stir bar was charged with a 1.0 mL solution of 5a (0.5 M) in anhydrous DCM under N₂. The G2 compound solution (100 µL, 8.5 mg mL⁻¹ in degassed DCM) was added via micro syringe into the vial at room temperature. After stirring for 24 h, the mixture was concentrated and then added dropwise into MeOH with vigorous stirring. The solid compound was collected and re-dissolved in a minimal amount of DCM. The precipitation procedure was repeated three times in total to afford the target product. The produced polymer was characterized by ¹H NMR, ¹³C NMR, FT-IR, GPC, DSC and TGA analysis.

**Conflicts of interest**

There are no conflicts to declare.

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**Notes and references**


When the reaction time was extended, oligomerization of COD was observed.


20 The asymmetry of the substituted FCOEs allowed for the preparation of polymers with regiorandom placement of the functional groups.

21 Increasing the [M]/[G2] ratio to 1000/1 (e.g., M = 7e) led to 52% conversion (Mn = 208 kDa and D = 1.79).

22 Although the molar mass of a polymer influences its glass-transition temperature, it only undergoes a very slight change of the Tg value when a high molecular weight range is reached as illustrated by the Flory-Fox equation.
