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Efficient synthesis of polycarbonate ether polyol via copolymerization of CO₂ and 1,2-butylene oxide over a layered Zn–Co double metal cyanide catalyst†

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Polycarbonate ether polyol synthesized by the copolymerization of carbon dioxide (CO₂) and epoxides is a promising technology for chemically fixing CO₂ and manufacturing degradable polymeric materials. However, research on the copolymerization of CO₂ and 1,2-butylene oxide (BO) to produce polycarbonate ether polyol is relatively scarce. Herein, we employed layered Zn–Co double metal cyanide (L-DMC) as a catalyst for the copolymerization of CO₂ and BO to obtain polycarbonate ether polyol. Under optimized conditions, the L-DMC-mediated copolymerization of CO₂ and BO displays excellent activity (catalytic productivity up to 1500 g polymer per g catalyst) and a high CO₂ incorporation fraction in polycarbonate ether polyol rate of 35.5%. *In situ* FTIR spectroscopy experiments showed a short induction period (10 min) in polymerization and competition between the synthesized polycarbonate ether polyol and butylene carbonate (BC). The properties of easy preparation, stability, and excellent catalytic performance of L-DMC suggest that it may be a promising candidate for large-scale production of CO₂–BO-based polymers.

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1. Introduction

The conversion of carbon dioxide (CO₂) into polycarbonate ether polyols is an effective strategy to achieve high value utilization of CO₂ and manufacture of degradable materials.^{1–7} Polycarbonate ether polyols are ideal soft segments for the preparation of high-performance polyurethane materials due to their unique degradability and inoxidizability.^{8–10} Currently, the synthesis of polycarbonate ether polyols is commonly achieved through the copolymerization of CO₂ and propylene oxide (PO).^{11–15} However, the deficiencies of CO₂–PO based polymers in terms of hydrophobicity and heat resistance prevent them from being used in hydrophobic coatings, waterproof film materials and high temperature-resistance adhesives.^{16–18} Compared with PO-based polymers, BO-based polymers synthesized from 1,2-epoxide butene (BO) with long alkyl side

chains have excellent hydrophobicity and heat resistance and have attracted extensive attention. Such polyethers synthesized by ring-opening polymerization of BO have been successfully used in oil-soluble and hydrophobic materials.^{19,20} To the best of our knowledge, research on the synthesis of CO₂–BO-based polymers has been relatively limited to date, resulting in the slow development of such materials.²¹ Therefore, synthesis of polycarbonate ether polyol by the development of efficient copolymerization methods for CO₂ and BO is highly desirable.²¹

Due to the properties of simple preparation and stability, double metal cyanide (DMC) catalysts can be used in the commercial production of polycarbonate ether polyols.²² Since 1966, intense research work have been performed to improve the catalytic efficiency of DMC catalysts.²³ Kim *et al.*²⁴ synthesized Zn–Co DMC catalysts using different dicarbonyl complexing agents which gave polypropylene carbonate polyols with low molecular weights (<4000 g mol^{−1}) and a high CO₂ incorporation fraction in polycarbonate ether rate of 46.7%. However, the productivity of the catalysts was lower than 160 g polymer per g DMC. In addition to altering the complexing agent of DMC, the development of novel DMC structures is an effective way to improve its performance. Zhang *et al.*²⁵ reported a nano-lamellar structure DMC catalyst for the copolymerization reaction between CO₂ with PO to obtain polycarbonates with number average molecular weights of >10 000 g mol^{−1}, but the conversion of PO was only 27.2% in 6 h. Trees *et al.*²⁶ had

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successfully synthesized a layered $\text{Zn}_2[\text{Co}(\text{CN})_6](\text{CH}_3\text{COO})$ double metal cyanide catalyst, which exhibited more excellent catalytic performance than conventional DMC for the ring-opening polymerization of epoxides. Experimental characterization results show that the two-dimensional layered structure of L-DMC with high specific surface area can expose more available active sites on the catalyst surface, which is essential for its high catalytic activity. Inspired by this work, we propose that the use of DMC catalysts with a layered structure for the copolymerization of CO_2 and BO will not only improve the polymerization efficiency but also increase the CO_2 incorporation fraction in the copolymer.

Herein, we synthesize a highly active L-DMC catalyst to achieve copolymerization of CO_2 and BO to prepare polycarbonate ether polyol (Fig. 1 and Scheme 1). The L-DMC mediated copolymerization can provide a catalytic productivity of 1553 g polymer per g catalyst and a CO_2 incorporation fraction in polycarbonate ether polyol rate of 35.5% at 110 °C and 3 MPa CO_2 . The high reactivity of L-DMC might be ascribed to its two-dimensional layered structure and high specific surface area, which was confirmed by BET, SEM, TEM, XRD and XPS studies. Moreover, the structure of the polycarbonate ether polyol was determined by FTIR, ^1H NMR, ^{13}C NMR and DOSY NMR characterization techniques. *In situ* FTIR spectroscopy monitored the copolymerization process of CO_2 and BO catalyzed, and the relationship between polycarbonate ether polyol and butylene carbonate (BC) during the formation process was revealed. The proposed approach successfully synthesized degradable CO_2 -BO-based polymer polyols creating better opportunities for preparing high-performance polyurethane materials.

2. Experimental part

2.1 Materials

1,2-Butylene oxide (BO), butane-1,4-diol (1,4-BDO), potassium hexacyanocobaltate(III) ($\text{K}_3[\text{Co}(\text{CN})_6]$), glacial acetic acid dihydrate ($\text{Zn}(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}$) and *tert*-butanol (*t*-BuOH) were purchased from Aladdin Industrial Corporation. Acetic acid (CH_3COOH) and zinc chloride (ZnCl_2) were provided by Sino-pharm Chemical Reagent Co., Ltd. BO was purified through refluxing calcium hydride for 48 h and subsequently kept in a nitrogen environment until needed. 1,4-BDO was dried on

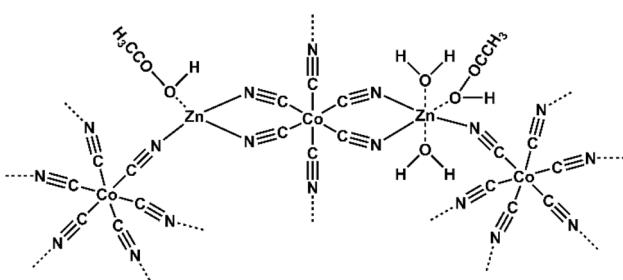


Fig. 1 Schematic structure of L-DMC catalyst with acetic acid as ligand.



Scheme 1 Copolymerization of CO_2 and BO to synthesize polycarbonate ether polyol by L-DMC catalyst.

a 4 Å molecular sieve for 48 h and also stored under nitrogen conditions for later use.

2.2 Preparation of L-DMC catalyst

The L-DMC catalyst was prepared according to the previous literature.²⁵ In order to avoid the influence of K^+ on catalyst activity, the required $\text{K}_3[\text{Co}(\text{CN})_6]$ in L-DMC synthesis was replaced with $\text{H}_3[\text{Co}(\text{CN})_6]$.²⁷ CH_3COOH (4.8 g, 0.08 mol) and $\text{Zn}(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}$ (660.0 mg, 3.0 mmol) were dissolved in distilled water (15.0 mL) to prepare solution A. $\text{H}_3[\text{Co}(\text{CN})_6]$ (496.0 mg, 1.5 mmol) was dissolved in distilled water (10.0 mL) to prepare solution B. Solution A was placed in an oil bath and heated to 90 °C with stirring, then solution B was added slowly and dropwise. The reaction solution took out from the oil bath right after the drop was finished. The solids that were suspended were gathered through centrifugation and rinsed three times with distilled water, and the resulting precipitate was dried under vacuum at 80 °C for 12 h.

2.3 Preparation of DMC catalyst

Preparation of DMC catalyst using methods reported in the literature.²⁸ Solution A was prepared by dissolving $\text{H}_3[\text{Co}(\text{CN})_6]$ (1.5 g, 4.8 mmol) in distilled water (20.0 mL). Solution B was prepared by dissolving ZnCl_2 (20.0 g, 0.15 mol) in a mixture of deionized water (50.0 mL) and *tert*-butanol (30.0 mL). Solution B was placed in an oil bath and heated to 50 °C with intense stirring, then solution A was added dropwise and continue stirring for 2 h. The white sediment was recovered by centrifugation and subsequently washed three times using a combined solution of deionized water and *tert*-butanol (the amount of *tert*-butanol in the washing solution is progressively increased), and the collected precipitate was dried under vacuum at 60 °C for 24 h.

2.4 Copolymerization of CO_2 and BO

The copolymerization of BO and CO_2 was conducted out in a pre-dehydrated 50 mL stainless steel autoclave reactor system. The catalyst was treated at 80 °C in a vacuum for 12 hours and was subsequently stored in a nitrogen atmosphere for further use. Taking the data from entry 2 in Table 1 as an example, the catalyst (8.0 mg), 1,4-BDO (63.0 mg, 0.7 mmol) and BO (8.0 g, 0.1 mol) were added into the 50 mL autoclave equipped with a magnetic stir. The autoclave was charged with CO_2 to reach specified pressure and heated to 110 °C. After cooling down to room temperature and releasing the pressure to obtain the crude product. The crude product was dissolved by methylene chloride, and the catalyst and solvent were removed by filtration and vacuum distillation, respectively. The pure product was

Table 1 Effect of reaction parameters on the copolymerization of CO_2 and BO^a

Entry	Catalyst (mg)	Temperature (°C)	Pressure (MPa)	$f_{\text{CO}_2}^d$ (%)	W_{BC}^d (wt%)	M_n^e (g mol $^{-1}$)	D^e	Productivity f (g polymer per g catalyst)
1 ^b	7.0	110	3	—	—	—	—	—
2	8.0	110	3	30.3	8.2	10 700	2.8	1239
3	10.0	110	3	35.0	22.1	6800	6.6	510
4	20.0	110	3	24.7	50.3	—	—	—
5	8.0	100	3	—	—	—	—	—
6	8.0	120	3	23.8	12.3	8000	3.0	925
7	8.0	140	3	4.0	1.0	1400	1.8	—
8	8.0	110	1	18.0	4.9	6300	3.1	1008
9	8.0	110	2	29.6	7.9	9700	3.5	1197
10	8.0	110	4	32.3	6.6	11 100	3.9	1401
11	8.0	110	5	35.5	10.2	12 600	3.2	1500
12 ^c	8.0	110	3	13.3	1.6	5000	2.5	861

^a All copolymerization reactions were carried out in 10 mL BO . ^b The copolymerization of CO_2 and BO was catalyzed by L-DMC. ^c The copolymerization of CO_2 and BO was catalyzed by DMC. ^d Determined by ^1H NMR and formula (1) and (2). ^e Determined by GPC. ^f Determined according to $W_{\text{product}}/W_{\text{cat}}$.

obtained by precipitation from methanol and dried to constant weight under vacuum condition at 80 °C.

The CO_2 incorporation fraction (f_{CO_2}) in polycarbonate ether polyol and the weight ratio of cyclic carbonate (W_{BC}) were determined by integrating the ^1H NMR signals at 4.7–4.9 ppm and 4.0–4.3 ppm (representing carbonate units), 3.2–3.7 ppm (indicating ether linkage chain) and 4.0 ppm, 4.5 ppm and 4.6 ppm (corresponding to 1,2-butylene carbonate), in accordance with the formulas (1) and (2) reported by Zhang *et al.*^{25,27}

$$f_{\text{CO}_2}\% = (A_{4.7-4.9} + A_{4.0-4.3} - 2A_{4.5})/[2(A_{4.7-4.9} + A_{4.0-4.3} - 2A_{4.5}) + A_{3.2-3.8}] \quad (1)$$

$$W_{\text{BC}}\% = (3A_{4.5})/(A_{4.7-4.9} + A_{4.5} + A_{4.0-4.3} + A_{3.2-3.8}) \quad (2)$$

3. Results and discussion

3.1 Characterization of L-DMC catalysts and DMC catalysts

Fig. S1† illustrated the powder XRD patterns of DMC and L-DMC catalysts. DMC exhibited broader diffraction peaks from 15° to 21°, whereas L-DMC displayed sharp diffraction peaks from 15° to 35°. This suggests that L-DMC possesses a more integrity crystal structure and exhibits higher crystallinity. Moreover, the layered phases of L-DMC were clearly characterized by reflections at $2\theta = 16.9^\circ$, 18.4° , 23.6° , 25.6° and 34.3° , which were consistent with the results of previous work.^{26,29}

The surface compositions of DMC and L-DMC were analyzed using XPS. The results indicates that Zn, Co, C, N, and O were common components of both catalysts (Fig. S2†). According to previous reports,³⁰ the main role of Co in Zn-Co DMC catalysts is to act as a stabilizer for the cyanide-bridged complexes to ensure that the catalysts remain heterogeneous, while Zn is the main key catalytic active site. Fig. 2 presented the expanded Zn 2p XPS spectra for the DMC and L-DMC catalysts. The binding energies of Zn atoms in both DMC and L-DMC catalyst systems were significantly reduced compared to the binding energy of Zn atoms in ZnCl_2 ($E_a = 1023.7$ eV), which suggests a change in

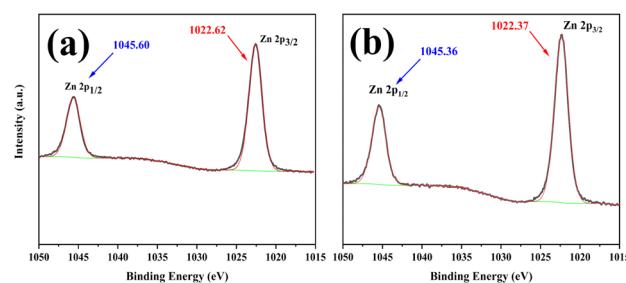


Fig. 2 Zn 2p XPS spectra of (a) DMC, (b) L-DMC.

the chemical coordination environment around the active site of Zn(II), thereby influencing its electronic structure properties.³¹ The DMC exhibited two singlet peaks of the Zn 2p spin-orbits with binding energies at 1022.62/1045.60 eV, and similarly the L-DMC also exhibited two singlet peaks of the Zn 2p spin-orbits with binding energies at 1022.37/1045.36 eV. This result suggests that the coordination between Zn atoms and different complexing agents can affect the binding energy of Zn 2p. CH_3COOH as a complexing agent tends to promote the transfer of electrons to the Zn center, which is contributing to the increase of catalytic performance.²⁴

Transmission electron microscopy (TEM) was used to characterize the structural morphology of DMC and L-DMC. As illustrated in Fig. 3(a) and (b), the particle shape of DMC predominantly exhibits a hexagonal configuration with a variation in particle size. In contrast, the particle size of L-DMC, as depicted in Fig. 3(c) and (d), is significantly smaller and exhibits a lamellar structure. These lamellar particles are primarily arranged in a random overlapping manner, leading to a layered structure. Fig. S3† presents the characterization results of the surface morphology as observed through SEM images of the two catalyst samples. As illustrated Fig. S3(a) and (b),† the DMC particles exhibited uneven particle sizes, irregular shapes, and low dispersion, with many particles adhering to one another and not easily separable. In contrast, the surface particles of L-



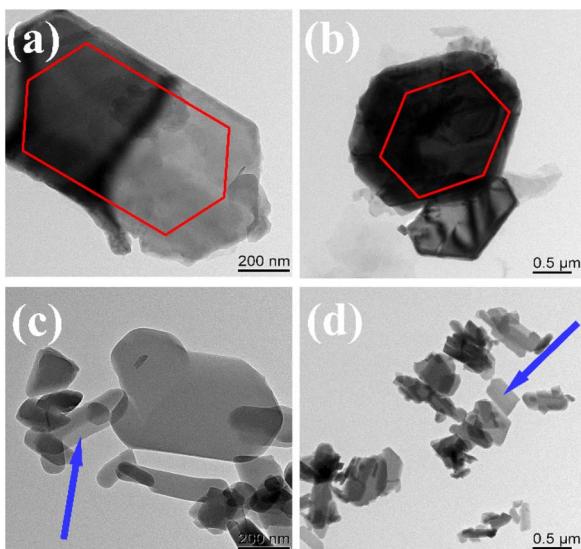


Fig. 3 TEM images (a and b) DMC, (c and d) L-DMC.

DMC demonstrate greater dispersion, as displayed in Fig. S3(c) and (d).[†] Furthermore, it is observed that the L-DMC particles are aggregated to form a layered structure, as highlighted in the enlarged image of Fig. S3(d),[†] which aligns with the findings from TEM characterization.

The N_2 adsorption isotherm curves for DMC and L-DMC were presented in Fig. S4.[†] The N_2 adsorption isotherm of the DMC catalyst exhibits a typical type V isotherm with a type H3 hysteresis loop,³² indicating a mesoporous structure. On the contrary, the adsorption isotherm of L-DMC catalyst only shows a small hysteresis loop.³³ The characteristics of the pore structure properties for both catalysts are presented in Table S1,[†] where the specific surface area ($S_{BET} = 668 \text{ m}^2 \text{ g}^{-1}$) and external specific surface area ($S_{ext} = 32 \text{ m}^2 \text{ g}^{-1}$) of L-DMC are significantly higher than those of DMC ($S_{BET} = 26 \text{ m}^2 \text{ g}^{-1}$ and $S_{ext} = 21 \text{ m}^2 \text{ g}^{-1}$). It has been reported that specific surface area is a crucial factor influencing the activity of the L-DMC catalyst, as the larger specific surface area typically increases the number of active sites exposed on the catalyst surface, thereby improving catalytic activity.^{34,35} Furthermore, based on the SEM analysis of the samples, it can be inferred that the high specific surface area of L-DMC may be attributed to its smaller particle size and highly dispersed particles without aggregation.

3.2 Influence of chain transfer agent type on the copolymerization of CO_2 and BO

Utilizing a chain transfer agent is essential for the synthesis of polycarbonate ether polyol on the copolymerization of CO_2 and BO. Research shows that low number average molecular weight PPGs ($200\text{--}600 \text{ g mol}^{-1}$) alcohols, non-complexing alcohols and carboxylic acid as chain transfer agents have different effects on the polymerization.^{13,36\text{--}38} This study investigated the effects of polypropylene glycol (PPG-400), butane-1,4-diol (1,4-BDO), and sebacic acid (SA) chain transfer agents on the copolymerization of CO_2 and BO. When SA as a chain transfer agent added to the

polymerization system (Table S2 (entry 3)[†]), targeted polymer was not obtained, which is attributed to the deactivation of the catalyst resulted from the strong coordination of SA with the Zn sites of L-DMC.³⁹ Furthermore, as shown in Table S2 (entries 1 and 2),[†] the employment of PPG-400 as the chain transfer agent, the f_{CO_2} in the product was only 10.9%, which is significantly lower than the desired ($f_{\text{CO}_2} > 30\%$). In contrast, the use of 1,4-BDO as a chain transfer agent in the copolymerization of CO_2 and BO demonstrated superior performance, achieving an f_{CO_2} of 30.3%, W_{BC} less than 10%, and a catalytic productivity of 1239 g polymer per g L-DMC. This suggested that the coordination of 1,4-BDO activates L-DMC, followed by initiation and propagation with BO. Therefore, 1,4-BDO was selected as the chain transfer agent for further investigations into the copolymerization of CO_2 and BO catalyzed by L-DMC. In addition, Table S2 (entry 4)[†] showed that the copolymerization of CO_2 and BO catalyzed by L-DMC can synthesize polycarbonate ether polyol with higher number average molecular weight ($M_n = 18\,300 \text{ g mol}^{-1}$) without the addition of chain transfer agent, which may be the result of adventitious traces of water (such as water present in the monomer) as the chain transfer agent.

3.3 Effect of reaction parameters on the copolymerization of CO_2 and BO

The effects of reaction parameters on the copolymerization of CO_2 and BO catalyzed by L-DMC are summarized from entries 1–11 in Table 1. Firstly, the effect of the dosage of L-DMC on polymerization was investigated. When the dosage of L-DMC was 7.0 mg (entry 1) under 110 °C and 3 MPa CO_2 conditions, no polymer was obtained. It is attributed to the total inactivation of the effective Zn active site of L-DMC after coordination with the chain transfer agent. When the dosage of L-DMC was increased to 8.0 mg (entry 2), copolymerization between CO_2 and BO was achieved, yielding 30.3% f_{CO_2} and a productivity of 1239 g polymer per g L-DMC. As the dosage of L-DMC was further increased to 10.0 mg (entry 3), f_{CO_2} rose to 35.0%; however, the weight ratio of BC increased significantly to 22.1% and the dispersity (D) value rose to 6.6. This is due to the fact that the number of available zinc active sites increases with the amount of L-DMC, which enhances the monomer conversion. It not only promotes the copolymerization of CO_2 with BO, but also facilitates the cycloaddition reaction of BO with CO_2 . Furthermore, the increased catalyst dosage could lead to polycarbonate ether polyol with high D , because the rise in the number of active centers contributes to a higher quantity of initiated chains.⁴⁰ When the dosage of L-DMC was increased to 20.0 mg (entry 4), W_{BC} further escalated to 50.3%, while f_{CO_2} decreased to 24.7%. This result may be due to the fact that excessive use of the L-DMC accelerated the activation of CO_2 in the reaction, leading to the inhibition of copolymerization by a huge amount of activated CO_2 . While excessive dosage of catalyst increased the number of Zn active sites, improved the conversion of monomer and promoted the formation of BC through the cycloaddition reaction of BO and CO_2 .

The effects of temperature on the copolymerization of CO_2 and BO catalyzed by L-DMC were summarized in Table 1



(entries 2 and 5–7). The copolymerization does not take place at 100 °C (entry 5). Data presented in Table 1 (entry 2) indicated that the lowest reaction temperature at which the copolymerization of BO and CO₂ taken place catalyzed by L-DMC was 110 °C. The result indicates that the ligands site composed of 1,4-BDO and active sites are more easily replaced by BO to activate L-DMC at higher temperatures. As the reaction temperature increased to 120 °C, the f_{CO_2} in polycarbonate ether polyol decreased from 30.3% to 23.8% while the content of the W_{BC} rose from 8.2% to 12.3% (entry 6). Because higher temperatures promoted the cycloaddition reaction of CO₂ and BO to replace part of the copolymerization of CO₂ and BO, which is consistent with previous research.^{41,42} When the reaction temperature was raised to 140 °C (entry 7), f_{CO_2} quickly decreased to 4.0% with only 1.0% of the byproduct BC formed, indicating that the majority of BO underwent homopolymerization into polyether products in a short time. Therefore, temperatures exceeding 120 °C are not conducive to the copolymerization of CO₂ and BO. This finding is consistent with the result of previous research, which suggests that lower reaction temperatures are beneficial for the copolymerization catalyzed by L-DMC.⁴³

Furthermore, the influence of CO₂ pressure factors on the reaction was investigated. As shown in Table 1 (entries 2 and 8–11), the copolymerization of CO₂ and BO catalyzed by L-DMC under different CO₂ pressures can synthesize polycarbonate ether polyol with M_n in the range of 6300 g mol⁻¹ to 12 600 g mol⁻¹ and D in the range of 3.1–3.9. The f_{CO_2} in polycarbonate ether polyol increases from 18.0% to 35.5% with the CO₂ pressure rising from 1 to 5 MPa. According to Fig. S5,† the glass transition temperatures (T_g) of the polycarbonate ether polyol are –26.1 °C (f_{CO_2} : 18.0%) and –13.5 °C (f_{CO_2} : 30.3%), suggesting that T_g increases with the addition of carbonate units. The low glass transition temperature of polycarbonate ether polyol highlights its potential as suitable parts for the synthesis of polyurethane soft segments. This finding indicates that increase of CO₂ pressure is an effective way to increase the CO₂ incorporation fraction in polycarbonate ether polyols⁴⁴ In

addition, the content of BC also increased with the increase of pressure, indicating that the synthesis of BC was facilitated by the increase in pressure.

Table 1 (entries 2 and 12) presented the catalytic effects of L-DMC and DMC catalysts on the copolymerization of CO₂ and BO. Notably, the f_{CO_2} in the polycarbonate ether polyol increased from 13.3% (entry 2) for the DMC catalyst to 30.3% (entry 12) for the synthesized L-DMC catalyst under identical reaction conditions. Furthermore, the catalytic efficiency comparison revealed that L-DMC achieved 1239 g polymer per g L-DMC, surpassing the 861 g polymer per g DMC, thereby indicating the superior catalytic activity of L-DMC. It has been reported that reactions catalyzed by DMC typically occur on the catalyst's surface.³⁴ Compared with DMC catalyst, L-DMC catalyst has a unique layered structure and smaller particle size, while the particles are also better dispersed. These characteristics are beneficial to the increase of the specific surface area of the catalyst, resulting in increased exposure to the available Zn active sites. The characterization results of BET (L-DMC: $S_{BET} = 668 \text{ m}^2 \text{ g}^{-1}$, $S_{ext} = 32 \text{ m}^2 \text{ g}^{-1}$, DMC: ($S_{BET} = 26 \text{ m}^2 \text{ g}^{-1}$, $S_{ext} = 21 \text{ m}^2 \text{ g}^{-1}$) also prove the possibility of the above analysis. Therefore, L-DMC showed excellent catalytic activity than DMC on the copolymerization of CO₂ and BO.³⁰

3.4 *In situ* FTIR spectroscopy for the copolymerization of CO₂ and BO catalyzed by L-DMC

In situ FTIR three-dimensional stack plot of L-DMC-catalyzed copolymerization of CO₂ and BO is presented in Fig. 4(a). The $\nu(C=O)$ and $\nu(C-O-C)$ characteristic absorption peaks at 1752 cm⁻¹ and 1101 cm⁻¹ were attributed to the carbonate units and the ether linkages of the polycarbonate ether polyol, respectively. Additionally, the characteristic absorption peak at 1819 cm⁻¹ was assigned to cyclic carbonate. In Fig. 4(b), the existence of an induction period for polymerization can be clearly observed, in which no change in the characteristic absorption peaks was observed within 10 min (stage 1) of the polymerization, indicating that the polymerization did not start

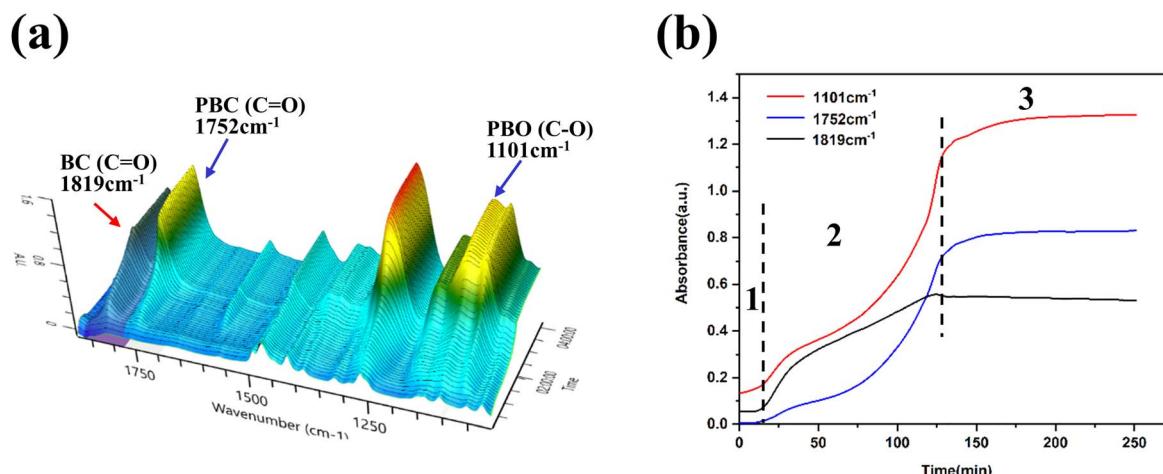


Fig. 4 (a) Three-dimensional stack plots of *in situ* FTIR spectroscopy. (b) Reaction profile monitored by *in situ* FTIR. Reaction was carried out at 110 °C, 1.5 MPa CO₂ pressure.



during this period. This phenomenon indicates that the L-DMC catalyst complies with the coordination polymerization mechanism of conventional DMC catalysts. As the reaction time increases from 10 min to 130 min (stage 2), the characteristic absorption peaks (C—O—C stretching vibration and C=O stretching vibration) of the ether linkages and carbonate units at 1101 cm^{-1} and 1752 cm^{-1} were gradually intensified. Meanwhile, the characteristic absorption peak (C=O stretching vibration) of cyclic carbonate was also gradually intensified. This result suggests that BO and CO_2 were rapidly converted to polycarbonate ether polyol and BC after ending the induction period. Notably, the growth rate of the characteristic absorption peaks of ether linkages and cyclic carbonate at the very beginning of polymerization is significantly faster than that of the characteristic absorption peak of carbonate units, indicating that the main formation of ether linkage chain segments and BC. As the reaction proceeds the reaction rate of BC is gradually decreased and the rate of synthesis of carbonate units and ether linkage in the polymer is nearly equal. The production of BC ceased in the reaction after 130 minutes (stage 3), and the synthesis rate of carbonate units and ether linkages in the polymer decreased significantly until BO was nearly fully converted. These findings indicate that the competitive formation between the target product polycarbonate ether polyol and the by-product BC during the copolymerization of CO_2 and BO using 1,4-BDO as chain transfer agent and L-DMC as catalyst.

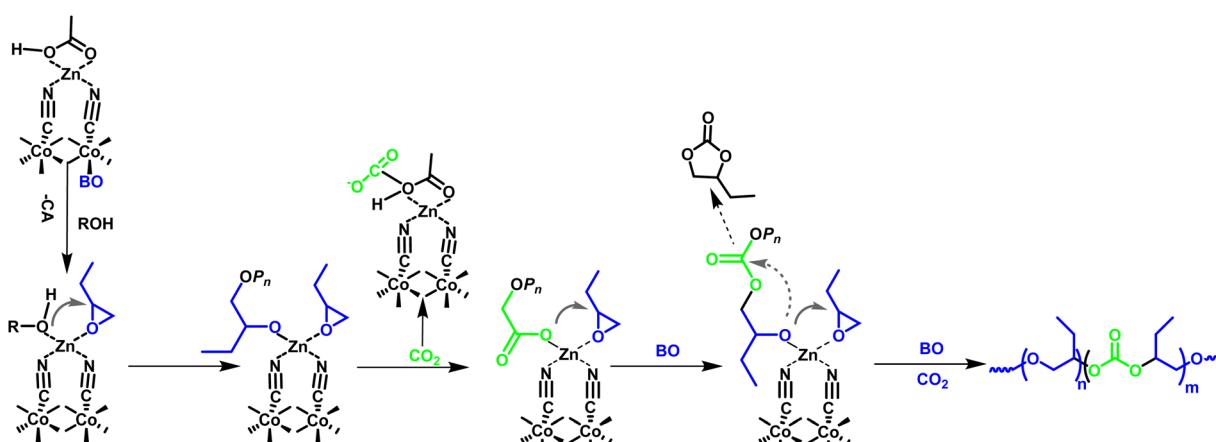
3.5 Plausible reaction mechanism

Given that the L-DMC catalyst demonstrates high catalytic activity in the copolymerization of CO_2 and BO, it is essential to propose a reaction mechanism that elucidates the interactions of its active sites with CO_2 and BO. The potential reaction mechanism has been formulated based on the characterization analysis detailed above, as well as previously reported studies on the polymerization processes involving analogous DMC catalysts.^{24,40} As illustrated in Scheme 2, the process begins with the oxygen atom (O) of the BO monomer and the hydroxyl group

of the chain transfer agent replacing the complexing agent of L-DMC, thereby coordinating with the Zn active sites. Subsequently, the hydroxyl group on the chain transfer agent nucleophilically attacks the BO coordinated with the Zn active sites, leading to the opening of its ring and the generation of propagating species. The process of propagation is carried out by the nucleophilic attack of the growing polymer chain on activated BO, while the absorbed CO_2 is activated by the Zn active sites and incorporated into the chain until the monomer is fully consumed. The byproduct BC may be generated by the cycloaddition reaction of CO_2 and BO.^{23,43,45} In this context, the acetic acid complexing agent in L-DMC not only interacts with the Zn atoms to influence the activity of catalyst but may also facilitates the absorption of CO_2 and enhances the CO_2 incorporation fraction in polycarbonate ether polyol.⁴⁶ This proposed mechanism aligns with the previously mentioned competitive relationship between BC and polycarbonate ether polyol.

3.6 Characterization of polymers

The composition of the polycarbonate ether polyol can be analyzed using FTIR and ^1H NMR spectroscopy. It can be observed in the FTIR spectra (Fig. 5(a)), polycarbonate ether polyol exhibits the characteristic $\nu(\text{C=O})$ of the carbonate units and the $\nu(\text{C—O})$ vibrations associated with the ether units at approximately 1752 cm^{-1} and 1255 cm^{-1} , respectively. Meanwhile, the stretching vibration of the C—O—C bond in the polyether segment is detected at 1101 cm^{-1} . ^1H NMR spectra of copolymer (Fig. 5(c)) exhibited characteristic signals of the methyl to illustrates the composition of the polycarbonate ether polyol. The signals detected in the range of 4.7–4.9 ppm and 4.0–4.3 ppm corresponding to the CH and CH_2 groups in the carbonate units, while those at 3.2–3.8 ppm was assigned to the CH and CH_2 groups in the ether linkages. Additionally, characteristic signals at 0.9–1.0 ppm and 1.4–1.7 ppm are assigned to the ethyl side chains in the polycarbonate ether polyol. Signals at 4.6 ppm, 4.5 ppm, and 4.0 ppm were ascribed to the aromatic protons of BC. The structure of the polycarbonate ether polyol



Scheme 2 Proposed mechanism for the copolymerization of CO_2 and BO using L-DMC catalyst.



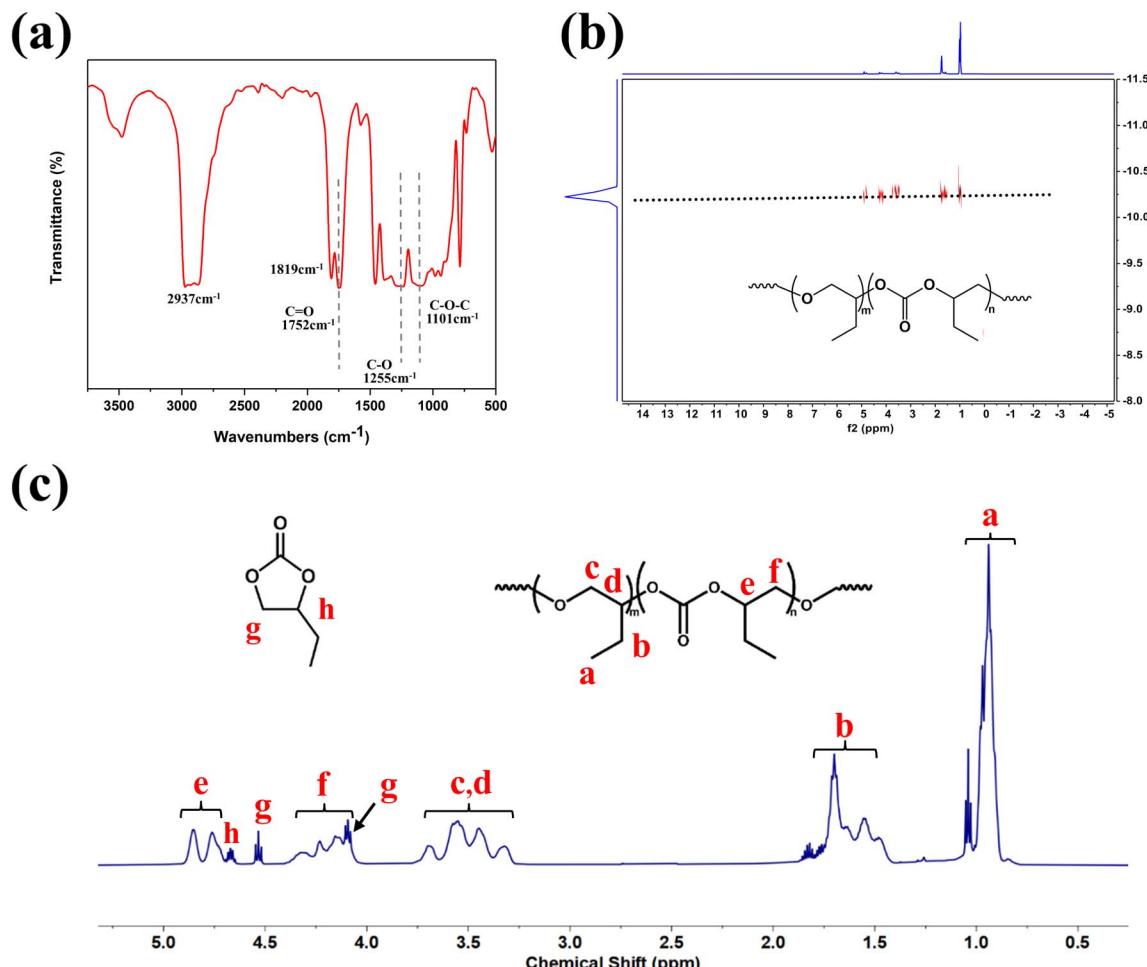


Fig. 5 Preparation of the polycarbonate ether polyol (Table 1, entry 2) by the copolymerization of CO_2 and BO using L-DMC. (a) FTIR spectrum. (b) DOSY NMR spectrum. (c) ^1H -NMR spectrum.

was obtained through FTIR spectra and ^1H NMR spectra, which also provided a basis for calculating the f_{CO_2} and W_{BC} . The ^{13}C NMR spectra of the polycarbonate ether polyol (Fig. S11†) reveals tail-to-tail (T-T) and head-to-tail (H-T) linkages, as evidenced by the typical signals observed at 154.84 ppm, 154.63 ppm, and 154.57 ppm. Consequently, it can be concluded that the BO units in the polycarbonate ether polyol are predominantly connected in the T-T configuration.^{15,47}

There is only one diffusion coefficient in the DOSY spectra of the CO_2 -BO copolymer in Fig. 5(b), and this result indicates that the product is a copolymer with mixed carbonate and ether linkages instead of mixture of polycarbonate and polyether. In addition, the MALDI-TOF results (Fig. S17†) showed that the molar masses of the polycarbonate ether polyol can be determined using the formula $m/z = 72.1 \times (m+n) + 44 \times n + 90.1 + 23$, where 72.1, 44, 90.1 and 23 represent the respective molar masses of the BO unit, the CO_2 unit, 1,4-BDO and sodium ion. For example, the peak at $m/z = 11197.3$ corresponds to the theoretical mass of the desired 1,4-BDO-initiated polycarbonate ether polyol (calculated for $[\text{M} + \text{Na}]^+ = 72.1 \times (70 + 52) + 44 \times 52 + 90.1 + 23 = 11197.3$).

4. Conclusions

In summary, a series of CO_2 -BO-based polymers were synthesized by high efficiency copolymerization of CO_2 and BO. Using L-DMC catalyst, the copolymerization of CO_2 and BO can be conducted under 110 °C and 3 MPa CO_2 conditions enables the synthesis of polycarbonate ether polyol with number average molecular weights of 6300–12600 g mol⁻¹ with moderate distributions ($D = 3.1\text{--}3.9$). The catalytic productivity of L-DMC up to 1500 g polymer per g catalyst and CO_2 incorporation fraction (f_{CO_2}) up to 35.5%. Compared with the conventional DMC catalysts, the two-dimensional layered structure and high specific surface area of L-DMC resulted in a significant increase in the number of effective Zn active sites facilitating the enhancement of CO_2 and BO copolymerization activities. Mechanistic studies revealed that the CH_3COOH complexing agent in L-DMC is essential for increasing the CO_2 incorporation fraction in polycarbonate ether polyols. ^{13}C NMR spectrum of the polycarbonate ether polyol revealed that BO units are primarily linked in a T-T configuration. *In situ* FTIR monitoring of the copolymerization process revealed the competitive



formation between the target product polycarbonate ether polylol and the byproduct BC and the induction period of the reaction was only 10 min with the use of 1,4-BDO chain transfer agent. Based on the advantages of simple preparation, good stability, and excellent catalytic performance of L-DMC, our next step is to promote its application to the large-scale production of CO₂-BO-based polymer materials.

Data availability

Data are available on request to the authors.

Author contributions

Chen Liu: investigation, methodology, writing – original draft. Yan Cao: writing – review & editing. Xianqiang Zeng: investigation. Zheng Zheng: investigation. Ziqiang Han: investigation. Peng He: investigation. Liguo Wang: conceptualization, project administration, supervision, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 G. S. Sulley, G. L. Gregory, T. T. D. Chen, L. Pena Carrodeguas, G. Trott, A. Santmarti, K. Y. Lee, N. J. Terrill and C. K. Williams, *J. Am. Chem. Soc.*, 2020, **142**, 4367–4378.
- 2 C. Chen, Y. Gnanou and X. Feng, *Macromolecules*, 2023, **56**, 892–898.
- 3 X. Zhang, J. Dong, Y. Su, E. G. Lee, Z. Duan, I. Kim and B. Liu, *Polym. Chem.*, 2023, **14**, 1263–1274.
- 4 M. Scharfenberg, S. Hofmann, J. Preis, J. Hilf and H. Frey, *Macromolecules*, 2017, **50**, 6088–6097.
- 5 Y. Li, X. Kou, X. Wang, L. Xia and Z. Li, *Polym. Chem.*, 2024, **15**, 2476–2481.
- 6 Z.-Q. Huang, J.-Q. Chen, X.-Y. Zhang, C.-K. Yuan, P. Wang, Y. Zhao, B.-C. Zhao, W.-X. Qi and W.-Y. Sun, *CrystEngComm*, 2024, **26**, 1701–1709.
- 7 M. Martínez de Sarasa Buchaca, F. de la Cruz-Martínez, L. F. Sánchez-Barba, J. Tejeda, A. M. Rodríguez, J. A. Castro-Osma and A. Lara-Sánchez, *Dalton Trans.*, 2023, **52**, 3482–3492.
- 8 S. Tang, H. Suo, R. Qu, H. Tang, M. Sun and Y. Qin, *Polymers*, 2023, **15**, 748.
- 9 J. E. Seong, S. J. Na, A. Cyriac, B.-W. Kim and B. Y. Lee, *Macromolecules*, 2009, **43**, 903–908.
- 10 H. I. Mao, C. W. Chen, L. Y. Guo and S. P. Rwei, *J. Appl. Polym. Sci.*, 2022, **139**, e52986.
- 11 M. Pohl, E. Danieli, M. Leven, W. Leitner, B. Blümich and T. E. Müller, *Macromolecules*, 2016, **49**, 8995–9003.
- 12 M. Scharfenberg, J. Hilf and H. Frey, *Adv. Funct. Mater.*, 2018, **28**, 1704302.
- 13 S. Liu, Y. Miao, L. Qiao, Y. Qin, X. Wang, X. Chen and F. Wang, *Polym. Chem.*, 2015, **6**, 7580–7585.
- 14 J. Wang, H. Zhang, Y. Miao, L. Qiao and X. Wang, *Green Chem.*, 2017, **19**, 2194–2200.
- 15 G.-W. Yang, C.-K. Xu, R. Xie, Y.-Y. Zhang, C. Lu, H. Qi, L. Yang, Y. Wang and G.-P. Wu, *Nat. Synth.*, 2022, **1**, 892–901.
- 16 S. Liu, K. Yan, P. Ren, X. Shang and X. Zhang, *Case Stud. Constr. Mater.*, 2024, **20**, e03136.
- 17 F. Yu, X. Xu, N. Lin and X. Y. Liu, *RSC Adv.*, 2015, **5**, 72544–72552.
- 18 D. Wang, Z. Huang, S. Shi, J. Ren and X. Dong, *J. Appl. Polym. Sci.*, 2022, **139**, e52437.
- 19 X. Zeng, L. Wang, Y. Cao, C. Liu, Z. Han, P. He and H. Li, *Eur. Polym. J.*, 2022, **179**, 111483.
- 20 X. Zeng, L. Wang, Y. Cao, C. Liu, P. He and H. Li, *Ind. Eng. Chem. Res.*, 2024, **63**, 1215–1227.
- 21 L. Kunze, S.-Y. Tseng, R. Schweins, T. Sottmann and H. Frey, *Langmuir*, 2019, **35**, 5221–5231.
- 22 D. A. Pyatakov and I. E. Nifant'ev, *Pet. Chem.*, 2024, **63**, 1170–1193.
- 23 S. Inoue, K. Matsumoto and Y. Yoshida, *Makromol. Chem.*, 2003, **181**, 2287–2292.
- 24 C. H. Tran, S. A. Kim, Y. Moon, Y. Lee, H. M. Ryu, J. H. Baik, S. C. Hong and I. Kim, *Catal. Today*, 2021, **375**, 335–342.
- 25 X.-H. Zhang, R.-J. Wei, X.-K. Sun, J.-F. Zhang, B.-Y. Du, Z.-Q. Fan and G.-R. Qi, *Polymer*, 2011, **52**, 5494–5502.
- 26 C. Marquez, A. Simonov, M. Wharmby, C. Goethem, I. Vankelecom, B. Bueken, A. Krajnc, G. Mali, D. Vos and T. De Baerdemaeker, *Chem. Sci.*, 2019, **10**, 4869–4875.
- 27 Y. H. Seo, Y. B. Hyun, H. J. Lee, J. W. Baek, H. C. Lee, J. H. Lee, J. Lee and B. Y. Lee, *J. CO₂ Util.*, 2021, **53**, 101755.
- 28 Z. Li, Y. Qin, X. Zhao, F. Wang, S. Zhang and X. Wang, *Eur. Polym. J.*, 2011, **47**, 2152–2157.
- 29 J. Sebastian and D. Srinivas, *Appl. Catal., A*, 2015, **506**, 163–172.
- 30 Z. Song, B. Subramaniam and R. V. Chaudhari, *ACS Sustainable Chem. Eng.*, 2019, **7**, 5698–5710.
- 31 I. Kim, J.-T. Ahn, C. S. Ha, C. S. Yang and I. Park, *Polymer*, 2003, **44**, 3417–3428.
- 32 H. Yang, L. Wang, S. Xu, Y. Cao, P. He, J. Chen, Z. Zheng and H. Li, *Green Energy Environ.*, 2022, **7**, 1361–1376.
- 33 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 34 A. Peeters, P. Valvekens, R. Ameloot, G. Sankar, C. E. A. Kirschhock and D. E. De Vos, *ACS Catal.*, 2013, **3**, 597–607.
- 35 N. J. Robertson, Z. Qin, G. C. Dallinger, E. B. Lobkovsky, S. Lee and G. W. Coates, *Dalton Trans.*, 2006, **45**, 5390–5395.
- 36 J. K. Varghese, D. S. Park, J. Y. Jeon and B. Y. Lee, *J. Polym. Sci., Part A: Polym. Chem.*, 2013, **51**, 4811–4818.



37 S. Liu, Y. Qin, L. Qiao, Y. Miao, X. Wang and F. Wang, *Polym. Chem.*, 2016, **7**, 146–152.

38 Y. J. Huang, G. R. Qi and Y. H. Wang, *J. Polym. Sci., Part A: Polym. Chem.*, 2002, **40**, 1142–1150.

39 R.-J. Wei, Y.-Y. Zhang, X.-H. Zhang, B.-Y. Du and Z.-Q. Fan, *RSC Adv.*, 2014, **4**, 21765–21771.

40 J. Zhao, B. G. Li and H. Fan, *Macromol. Theory Simul.*, 2021, **30**, 2000101.

41 S. Chen, Z. J. Hua, Z. Fang and G. R. Qi, *Polymer*, 2004, **45**, 6519–6524.

42 K. Ma, Q. Bai, L. Zhang and B. Liu, *RSC Adv.*, 2016, **6**, 48405–48410.

43 Y. Gao, Y. Qin, X. Zhao, F. Wang and X. Wang, *J. Polym. Res.*, 2012, **19**, 1–9.

44 S. Liu, Y. Qin, X. Chen, X. Wang and F. Wang, *Polym. Chem.*, 2014, **5**, 6171–6179.

45 P. Góarecki and W. Kuran, *J. Polym. Sci., Polym. Lett. Ed.*, 2003, **23**, 299–304.

46 F. Liu, Y. Gu, P. Zhao, J. Gao and M. Liu, *ACS Sustainable Chem. Eng.*, 2019, **7**, 5940–5945.

47 W. Zhang, Q. Lin, Y. Cheng, L. Lu, B. Lin, L. Pan and N. Xu, *J. Appl. Polym. Sci.*, 2012, **123**, 977–985.

