Efficient catalyst removal and recycling in copolymerization of epoxides with carbon dioxide *via* simple liquid–liquid phase separation†

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A simple and efficient catalyst removal system has been developed in the cobalt–salen-catalyzed copolymerization of propylene oxide with carbon dioxide. The present system requires no prior modification of the catalyst, and the removal is achieved by simple addition of myristic acid, followed by organic liquid–liquid phase separation.

The alternating copolymerization of epoxides with carbon dioxide to give aliphatic polycarbonates, pioneered by Inoue and coworkers in the 1960s,¹ has attracted much attention as one of the most promising processes for CO₂ utilization.² In the last decade, a variety of catalyst systems based on metal–salen complexes have been developed,^{2c,3–5} many of which show much higher catalytic activities than other conventional systems. Unfortunately, however, because metal–salen complexes are highly coloured, even a small amount of residual metal–salen complexes in the product polycarbonates leads to undesired coloration of these inherently colourless copolymers. In addition, highly active catalyst systems usually employ cobalt^{4,6} or chromium^{5,7} complexes, and contamination of these metals in the product polycarbonates could also be problematic due to their toxicity. It is therefore highly desirable to develop a method that allows for easy and efficient removal of these metal complexes from the polymers in this alternating copolymerization.

The conventional and most typical method for purification of the polycarbonates from metal catalysts is achieved by precipitation of polycarbonates with methanol, but repeated precipitation is required to obtain pure polycarbonates. To overcome this cumbersome purification procedure, several catalyst systems have been designed for easy separation of the metal catalysts.

For example, insoluble polymer-supported chromium/porphyrin⁸ and zinc/diiminato⁹ catalysts were employed in the copolymerization of cyclohexene oxide (CHO) with CO₂, and these catalysts could be removed by filtration after the polymerization reaction. In addition to these catalyst immobilization approaches, modification of homogeneous metal catalysts has also been explored. Lee and coworkers reported a highly active cobalt–salen complex bearing four quaternary ammonium salt units for copolymerization of propylene oxide (PO) with CO₂.^{4j} The product polymers could be purified by passing through a pad of silica gel, and the catalyst remaining on the silica gel could also be recovered and reused after sequential anion exchange processes. Darensbourg, Bergbreiter, and coworkers introduced a chromium–salen complex having polyisobutylene tethers for copolymerization of CHO with CO₂.^{5d} In this system, due to the highly non-polar nature of the polyisobutylene tethers, the chromium–salen complex could be efficiently removed from the polycarbonates through HCl treatment in MeCN followed by extraction with heptane, and the recovered catalyst could also be reused for the copolymerization.

Although significant progress has been made for efficient catalyst removal as mentioned above, the major drawback for both immobilized and homogeneous methods is the requirement of design and synthesis of elaborate metal–ligand complexes. In addition, due to the modification of the catalyst, the polymerization activity can sometimes become lower than those of the corresponding parent metal complexes. To address this issue, Jessop and coworkers demonstrated an efficient purification of poly(cyclohexene carbonate) from a simple chromium–salen catalyst by way of switching the polarity of the amine solvent through introduction of CO₂.^{5f}

In this context, herein we describe the development of a simple catalyst removal system *via* liquid–liquid phase separation in the copolymerization of PO with CO₂. The present system employs a readily available, standard cobalt–salen complex with no prior modification, and the separation is achieved by simple addition of a commercially available, long-chain alkylcarboxylic acid after the copolymerization reaction.

A schematic view of the catalyst separation system in our present study is illustrated in Fig. 1. The copolymerization of

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Fig. 1 Schematic representation of the catalyst separation in the copolymerization of propylene oxide with CO_2 .

PO with CO_2 is conducted with readily available cobalt-salen complex **1**^{4a,b,10} as the catalyst in the presence of bis(triphenylphosphine)iminium chloride ([PPN]Cl), and the reaction mixture is treated with a long-chain alkylcarboxylic acid (RCOOH) after the copolymerization. The anionic propagated polymer chain on the cobalt centre is replaced by the carboxylate and released as neutral poly(propylene carbonate) (PPC). Because of the long-chain alkyl carboxylate on cobalt, the resulting cobalt-salen complex becomes highly soluble in non-polar hydrocarbon solvents, while the product PPC stays selectively soluble in polar solvents such as MeCN and DMF. Accordingly, the cobalt-salen complex can be easily separated from the product PPC *via* simple liquid-liquid phase separation by using immiscible non-polar and polar organic solvents.

In an initial investigation, we evaluated the efficiency of the catalyst separation by changing the chain length of the carboxylic acid after the copolymerization of PO with CO_2 . The copolymerization was carried out by using cobalt-salen complex **1** in the presence of [PPN]Cl (PO/1/[PPN]Cl = 4000/1/0.5, 2.0 MPa of CO_2). After the unreacted PO and CO_2 were removed, the polymerization mixture was dissolved in CH_2Cl_2 and treated with a carboxylic acid (2 equiv. to **1**). After removal of the solvent, the residue was dissolved in CH_3CN and the cobalt-salen complex was extracted with hexane. As a starting point, acetic acid ($\text{R} = \text{CH}_3$) was used and the amount of extracted cobalt-salen complex by this method was estimated to be 74% based on UV-VIS spectroscopy ($\lambda_{\text{max}} = 411 \text{ nm}$, $\epsilon = 6.67 \times 10^3 \text{ M}^{-1} \text{ cm}^{-1}$) (Table 1, entry 1). As expected, the use of longer alkylcarboxylic acids led to higher

Table 1 Effect of carboxylic acid for the efficiency of the catalyst separation

Co complex 1 (0.025 mol%) + [PPN]Cl (0.013 mol%) $\xrightarrow[22^\circ\text{C, 13 h}]{\text{CO}_2 (2.0 \text{ MPa})}$ RCOOH (0.050 mol%) $\xrightarrow[\text{CH}_2\text{Cl}_2, \text{ r.t., 2 h}]{\text{hexane/MeCN extraction}}$ $\text{PPC/PC} = 99/1$ carbonate linkage: >99%					
Entry	R	Catalyst removal ^a (%)	Yield of PPC + PC ^b (%)	M_n^c (g mol^{-1})	M_w/M_n^c
1	CH_3	74	36	26 700	1.1
2	C_5H_{11}	79	37	23 100	1.1
3	C_9H_{19}	90	39	35 800	1.1
4	$\text{C}_{13}\text{H}_{27}$	96	36	39 800	1.1

^a Determined using UV-VIS spectroscopy. ^b PC = propylene carbonate.

^c Determined using size-exclusion chromatography using a polystyrene standard.

efficiency of the catalyst separation. Thus, 79% of the cobalt complex was removed by using caproic acid ($\text{R} = \text{C}_5\text{H}_{11}$; entry 2), and the use of capric acid ($\text{R} = \text{C}_9\text{H}_{19}$) led to 90% removal under the same conditions (entry 3). Finally, as high as 96% of the cobalt-salen complex was extracted into the hexane phase by using myristic acid ($\text{R} = \text{C}_{13}\text{H}_{27}$) (entry 4). In this experiment, isolated PPC from the MeCN phase was also analyzed by ICP, and the amount of cobalt remaining in the polymer was found to be 15.6 ppm, which corresponded to 3.9% of the initially loaded cobalt-salen complex **1** (eqn (1)). It is worth noting that further purification of the polymer by repeating the same separation method once again gave the PPC with 1.9 ppm cobalt, showing that >99.5% of the cobalt could be removed by employing two cycles of this sequence. In comparison, ICP analysis was also carried out for the PPC obtained by the conventional precipitation method after the copolymerization reaction (eqn (2)). As much as 108 ppm of cobalt remained in the polymer after a single precipitation process, indicating that only 74% of the cobalt complex could be removed. 65.0 ppm of cobalt (16% of the used cobalt) was still detected even after the second precipitation, clearly demonstrating the superiority of our present purification method.

Having established an easy and efficient process for the catalyst removal in the present copolymerization, we turned our attention to recycling of the catalyst. Because the extracted cobalt-salen complex in the hexane phase is supposed to bear a carboxylate ligand (myristate instead of acetate), it should be ready to be used as the copolymerization catalyst after evaporation of the solvent. Indeed, the recovered catalyst displayed a

Table 2 Recycling experiments for the cobalt–salen-catalyzed copolymerization of PO with CO₂

Cycle	Yield of PPC + PC (%)	PPC/PC ^a	TOF for PPC (h ^{−1})	Carbonate linkage ^a (%)	M _n ^b (g mol ^{−1})	M _w /M _n ^b	Catalyst recovery ^c (%)
1	36	99/1	111	>99	39 800	1.1	96
2	29	99/1	89	>99	28 200	1.1	94
3	27	98/2	80	>99	22 700	1.1	94
4	28	99/1	84	>99	22 400	1.1	96
5	27	98/2	82	>99	23 900	1.1	94
6	23	96/4	69	>99	20 500	1.1	91

^a Determined using ¹H NMR. ^b Determined using size-exclusion chromatography using a polystyrene standard. ^c Determined using UV-VIS spectroscopy.

reasonable activity for the copolymerization of PO with CO₂ without further purification and the results of the recycling experiments of up to 5 times are summarized in Table 2. It is worth noting that the efficiency of catalyst removal (recovery) stays high for each cycle of the copolymerization (91–96%), although the catalyst activity gradually decreases.

In summary, we have developed a simple and efficient catalyst removal system in the cobalt–salen-catalyzed alternating copolymerization of propylene oxide with carbon dioxide. The present system requires no prior modification of the cobalt–salen complex, and the catalyst removal is achieved by simple addition of a long-chain alkylcarboxylic acid after the copolymerization, followed by organic liquid–liquid phase separation. The removal efficiency is much higher than the conventional purification method by precipitation, and the recovered cobalt catalyst can also be directly reused multiple times.

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