

Cite this: *Dalton Trans.*, 2020, **49**, 223

# Heterodinuclear complexes featuring Zn(II) and M = Al(III), Ga(III) or In(III) for cyclohexene oxide and CO<sub>2</sub> copolymerisation†

Arron C. Deacy, Christopher B. Durr and Charlotte K. Williams \*

The ring opening copolymerisation of CO<sub>2</sub> and epoxides is a useful means to valorise waste emissions and to reduce pollution in polymer manufacturing. Heterodinuclear catalysts, particularly those of Zn(II)/Mg(II), have shown better performances than homodinuclear analogues in this reaction. As part of on-going efforts to better understand the catalytic synergy, this work describes a series of heterodinuclear complexes, combining Zn(II) with a metal from Group 13 (M = Al(III), Ga(III) or In(III)). The complexes are synthesised from a symmetrical macrocyclic ligand in high yields *via* sequential metalation steps and are the thermodynamic reaction products. The Zn(II)/Group 13 complexes are effective homogeneous catalysts for the ring opening copolymerisation (ROCOP) of cyclohexene oxide at 1 bar pressure of carbon dioxide, but all show inferior performances compared to the di-zinc analogue. The CO<sub>2</sub> uptake into the polymer increases in the order Al(III) < Ga(III) < In(III) which is attributed to lower Lewis acidity heavier Group 13 homologues showing a reduced tendency to form ether linkages. Concurrently, polycarbonate activity increases down the Group 13 series consistent with weaker metal–oxygen bonds which show enhanced lability to insertion reactions.

Received 15th July 2019,  
Accepted 22nd October 2019  
DOI: 10.1039/c9dt02918d

rsc.li/dalton

## Introduction

The transformation of CO<sub>2</sub> into valuable products is a cornerstone of sustainable chemistry,<sup>1</sup> driven by the necessity to reduce green-house gas emissions and fossil fuel consumption in chemical manufacturing.<sup>2</sup> In contrast to the plethora of esoteric stoichiometric CO<sub>2</sub> transformations, CO<sub>2</sub>/epoxide ring-opening copolymerisation (ROCOP) is a promising utilisation as it is truly catalytic and allows for CO<sub>2</sub> uptake of 30–50% by weight into the polymer.<sup>3,4</sup> The major immediate term opportunity for these materials is as low molecular mass, hydroxyl terminated polycarbonate or polyether-carbonate polyols.<sup>5</sup> These polyols are used to manufacture a range of coatings, urethanes and thermosets which show equivalent or better properties than incumbent materials that derive only from petrochemicals.<sup>6–9</sup> Importantly, the process of CO<sub>2</sub>/epoxide copolymerisation to yield polycarbonate polyols is amenable to large-scale manufacturing.<sup>5,10</sup> Furthermore, life cycle analysis has shown a ‘triple win’ in terms of CO<sub>2</sub> emissions, for every

CO<sub>2</sub> molecule incorporated into the polymer backbone, two more are saved by reducing epoxide consumption.<sup>2</sup> Catalysis is central to the success of CO<sub>2</sub>/epoxide ROCOP but overall the field suffers from a lack of diversity in catalyst structures with most studies focussing on either di-Zn(II) or bicomponent Co(III)/Cr(III)/Al(III) complex/co-catalyst systems.<sup>11,12</sup> These bicomponent systems can show high activity but are known to enchain by complex mechanisms and require the use of expensive and corrosive co-catalysts. Our research group have focused on single-component dinuclear catalysts which obviate co-catalyst requirements; di-Zn(II), Mg(II), Co(II/III) and Fe(III) macrocyclic complexes are all active and selective in the low pressure regime (1 bar CO<sub>2</sub> pressure).<sup>13–16</sup> In 2015, we reported the first Zn(II)/Mg(II) heterodinuclear catalyst which showed superior performances to either di-Zn(II) or di-Mg(II) counterparts, or any combinations of them.<sup>17</sup> Recently we investigated a series of these Zn(II)/Mg(II) heterocatalysts and proposed a chain shuttling mechanism whereby magnesium coordinates and activates the epoxide whilst zinc provides the nucleophilic carbonate group.<sup>18</sup> In 2018, Okuda and Mashima reported a series of hetero-tetranuclear catalysts of the form Zn<sub>3</sub>Ln which were also highly active.<sup>19</sup> Given the promise of these mixed metal catalysts it is important to understand which metal combinations are most effective and how synergy functions. Naturally, not all metal combinations are synergic as demonstrated by recent investigation of heterocomplexes

Department of Chemistry, Chemistry Research Laboratory, 12 Mansfield Rd, Oxford OX1 3TA, UK. E-mail: charlotte.williams@chem.ox.ac.uk

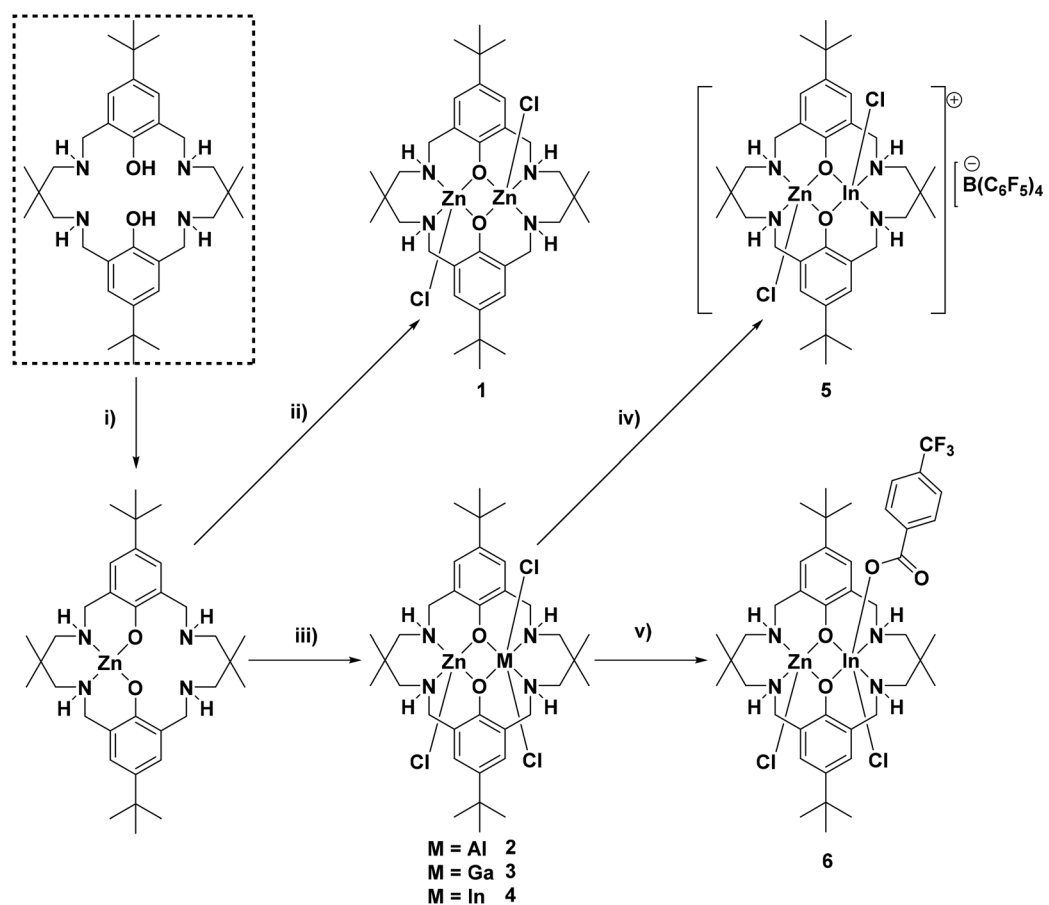
† Electronic supplementary information (ESI) available. CCDC 1939693–1939697. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9dt02918d



featuring Zn(II) with Group 1 elements [Li(I), Na(I) and K(I)] or Zn(II) with Group 2 elements [Mg(II) and Ca(II)]. Within the series of complexes, only the Zn(II)/Mg(II) combination showed synergy and all others were less active than the Zn(II)/Zn(II) homodinuclear complex.<sup>20</sup> In this work, we investigate a series of complexes featuring Zn(II) and Group 13 elements. The metal selection is motivated by some precedent for Al(III) complexes in this catalysis, for example Al(III) salen,<sup>21–23</sup> porphyrin<sup>24–28</sup> or trisphenolate<sup>29</sup> complexes are all active as part of bicomponent systems (*i.e.* with co-catalysts). Very recently, the first In(III) catalysts were reported and operate without co-catalyst, attributed to an unusual mononuclear mechanism.<sup>30</sup> Al-catalyst systems, with co-catalysts, are also widely investigated in the ring-opening copolymerisation of epoxides and cyclic anhydrides.<sup>31–36</sup> Group 13 complexes also show good activity for mechanistically related reactions such as in the ring-opening polymerisation of lactide, lactones or cyclic carbonates.<sup>37–42</sup> For the target macrocyclic catalysts, it was envisaged that combining the labile Zn(II) site with Group 13 elements could allow modification of metal Lewis acidity and alkoxide lability. When targeting new catalysts, it is proposed that both metals should have sufficiently small ionic radii to enable in-plane coordination in the macrocycle cavity.<sup>20</sup>

## Results and discussion

Firstly, the di-zinc complex (**1**) was prepared to allow for comparison with the target heterodinuclear complexes (**2–4**); all complexes were synthesised using a sequential metalation procedure (Fig. 1). The synthesis involved ligand deprotonation with diethyl zinc, in THF at 25 °C, to yield *in situ* a mono-zinc complex (not isolated). Next, ZnCl<sub>2</sub> (for **1**) or the appropriate MCl<sub>3</sub> {for (**2–4**) where M = Al(III), Ga(III) or In(III)} precursor was added and the reaction heated to 100 °C for 16 h. The complexes were isolated, without further purification, as white powders in reasonable isolated yields (>70%). To understand the importance of providing a free coordination site for monomers, cationic complex **5** was synthesised by chloride abstraction from complex **4**, by reaction with K[B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>] in THF at 25 °C, and was isolated as an off-white powder in 90% yield. To understand the influence of the co-ligand on initiation rates complex **6**, featuring a bridging carboxylate co-ligand, was synthesised *via* a metathesis reaction between complex **4** and KOBz<sub>p</sub>CF<sub>3</sub>. The reaction was conducted in MeOH, at 25 °C, and enabled isolation of the product as a white powder in 95% yield. All complexes were characterised by solution state NMR spectroscopy (Fig. S1–S25<sup>†</sup>), mass spectrometry



**Fig. 1** Synthesis of complexes **1–6**. (i) ZnEt<sub>2</sub>, THF, 25 °C, 16 h. (ii) ZnCl<sub>2</sub>, THF, 100 °C, 16 h, 81%. (iii) MCl<sub>3</sub> (M = Al, Ga or In), THF, 100 °C, 16 h, >70%. (iv) K[B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>], THF, 25 °C, 16 h, 90%. (v) KOBz<sub>p</sub>CF<sub>3</sub>, MeOH, 25 °C, 16 h, 95%.



(Fig. S26–S29†), elemental analysis and complexes **1–4** were also characterised by single crystal X-ray diffraction experiments (Fig. S30–S33†).

The  $^1\text{H}$  NMR spectrum of **1** shows multiple, complex signals at room temperature in all solvents tested (THF, methanol, TCE) but was successfully determined in tetrachloroethane (TCE) at 403 K, under these conditions the signals coalesce to give an assignable spectrum. The spectrum shows five resonances between 2.2 ppm and 5.0 ppm, representative of the benzylic, methylene and amine proton environments. There is a single resonance at 7.00 ppm corresponding to the phenyl protons and this peak is indicative of a homodinuclear complex coordinated by a symmetrical ligand. Its  $^{13}\text{C}$  NMR spectrum displays two signals, at 58.7 and 64.5 ppm, corresponding to the benzylic and methylene carbon atoms, along with four phenyl carbon signals, at 160.5 (*ipso*, phenolic carbon), 140.3 (*para*), 125.5 (*ortho*) and 128.4 ppm (*meta*), due to the complex's symmetry. The  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra are in-line with related macrocycle di-zinc complexes with coligands such as acetate, phenyl, benzoate and iodide.<sup>13,20,43,44</sup> The MALDI-ToF mass spectrum displays a peak at 717 *m/z*, corresponding to the molecular cation,  $[\text{LZn}_2(\text{Cl})]^+$ .

NMR spectroscopy is a useful means to determine the success of the heterodinuclear complex synthesis and comparisons between the signals for the heterodinuclear complexes and homodinuclear (**1**) are useful to determine purity. The homodinuclear complex is  $C_2$  symmetric but heterodinuclear complexes **2–4** lack this symmetry and as a result show different environments for all the protons in the  $^1\text{H}$  NMR spectra (Fig. 2). Another indicator of heterodinuclear complex formation is the presence of coupling between the two phenyl proton resonances, at  $\sim 7.00$  ppm, as confirmed by 2D COSY NMR spectroscopy (Fig. S5, S9 and S14†). If a mixture of homodinuclear complexes were formed, it would be expected to display two phenyl resonances but these would not couple

with one another. The presence of coupled aromatic signals is a strong indicator of heterodinuclear complex formation; that said, in some circumstances this coupling is more difficult to unambiguously assign as it depends upon the magnitude of the  $^4J_{\text{H-H}}$  coupling constant. The  $^1\text{H}$  NMR spectra for the heterodinuclear complexes can be used to infer the approximate purity, by comparing the integrals of signals at 6.79 (if any complex **1** is present) and  $\sim 6.97$  ppm (for heterocomplexes **2–4**). This analysis indicates that the heterodinuclear complexes are all formed as the major product, with purity values, within the detection limits of NMR spectroscopy, of 90% (**2**), 95% (**3**) and  $>99\%$  (**4**), respectively.

The  $^{13}\text{C}$  NMR spectra also show the loss of  $C_2$  symmetry compared to **1**, with the heterodinuclear complexes displaying double the number of signals for nearly all the carbon atoms (Fig. 2). Another diagnostic feature of heterodinuclear complex formation is the retention of a single resonance for the *ipso*- and *para*-phenyl carbons, at approx. 160 and 140 ppm, respectively (Fig. 2). A mixture of homodinuclear complexes would be expected to display two signals each for these carbons. The  $^{13}\text{C}$  NMR can be used in conjunction with  $^1\text{H}$  and COSY NMR to confirm heterodinuclear complex formation.

2D DOSY NMR spectra for complexes **2–4** all show a single diffusion coefficient, consistent with a discrete monomeric heterodinuclear complex in THF solution (as a representative solvent considering that polymerizations are conducted in neat epoxide) (Fig. S11, S16 and S25†). MALDI-ToF mass spectra showed molecular ions corresponding to the cation  $[\text{LZnM}(\text{Cl}_2)]^+$  at 714 *m/z*, 755 *m/z* and 800 *m/z*, where  $M = \text{Al}$ ,  $\text{Ga}$  and  $\text{In}$ , for complexes **2**, **3** and **4**, respectively (Fig. S27–S29†).

The  $^1\text{H}$  NMR spectrum of cationic complex **5** displays significant shifts to higher chemical shift values for all NH resonances compared to complex **4** (Fig. S17†), consistent with the increased positive charge on the complex. The NMR data clearly confirm that the heterodinuclear complex is main-

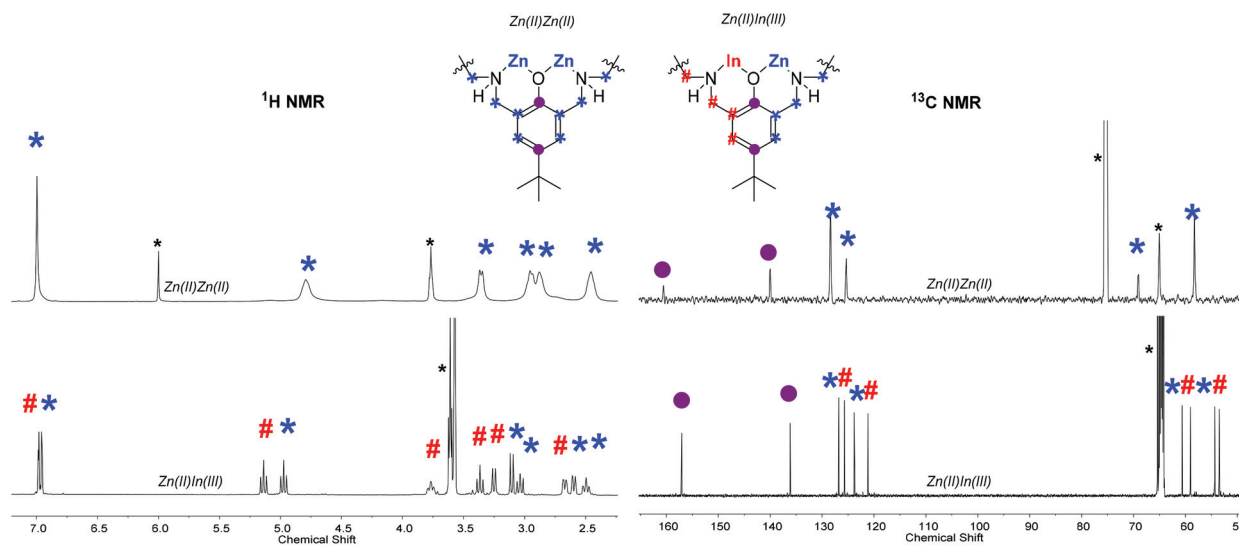


Fig. 2 Overlaid selected regions of the  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra of di-zinc complex (**1**) and heterodinuclear  $\text{Zn}(\text{II})\text{In}(\text{III})$  complex (**4**). Note the full spectra and assignment is available in the ESI (Fig. S1, S2, S12 and S15†). \*Residual TCE and THF.



tained and that there are not any metal re-distribution reactions.

The  $^{11}\text{B}$ ,  $^{13}\text{C}$  and  $^{19}\text{F}$  NMR spectra also all indicate the formation of a single complex (Fig. S19–S21†). The  $^1\text{H}$  NMR spectrum of complex **6** indicates the success of the metathesis reaction and is consistent with the solid state structure where the  $p^{\text{CF}_3}$ benzoate is located at the bridging position (Fig. S22 and S34;† single crystal analysis by X-ray diffraction gave insufficient data to fully resolve the structure, though its connectivity could still be obtained). The  $^1\text{H}$  NMR data confirms that the heterodinuclear complex is retained without any detectable metal redistribution. The NH resonances are not observed due to H–D exchange processes occurring between the complex and the solvent ( $d_4$ -MeOD). A major and minor set of resonances are observed for the  $p^{\text{CF}_3}$ benzoate protons which is attributed to both  $\kappa_1$  vs.  $\kappa_2$  coordination modes, as was previously observed for an analogous Mg(II)/Zn(II) complex.<sup>18</sup>

Single crystals, suitable for X-ray crystallography, were isolated by diffusion of pentane into a saturated solution of the complexes in acetonitrile (**1**, **2** and **3**) or THF (**4** and **6**), at  $-40\text{ }^\circ\text{C}$  in a glovebox (Fig. S30–S34 and Tables S1–S5†). The structures of **2–4** confirm the formation of heterodinuclear complexes, which are monomeric in the solid state, and which feature three chloride co-ligands. The structures show the

macrocycle adopts a ‘bowl’ conformation in each case. For complexes **1–4**, the zinc atom(s) adopts a pentacoordinate, distorted square based pyramidal geometry ( $\tau_5 = 0.35$  Zn1 (**1**),  $\tau_5 = 0.15$  Zn2 (**1**),  $\tau_5 = 0.07$  Zn1 (**2**),  $\tau_5 = 0.02$  Zn1 (**3**),  $\tau_5 = 0.01$  Zn1 (**4**)) and is coordinated by a chloride co-ligand. The Group 13 metals show octahedral coordination geometries and are coordinated by two chloride co-ligands in line with the higher oxidation state of these metals. For **1**, the Zn–Zn separation is  $3.04(1)\text{ \AA}$ , consistent with previous di-zinc complexes coordinated by this ligand and with the proposed distance for effective catalysis.<sup>20,44–46</sup> For **2–4**, Zn–M(III) separations increase with the increasing ionic radius of the Group 13 element and are  $3.02(3)\text{ \AA}$ ,  $3.12(3)\text{ \AA}$  and  $3.15(8)\text{ \AA}$  for complexes **2**, **3** and **4**, respectively (Fig. 3).

The heterodinuclear complexes, **2–6**, were tested for  $\text{CO}_2/\text{CHO}$  ROCOP and compared against the di-zinc catalyst (**1**). The polymerisations were all conducted at  $80\text{ }^\circ\text{C}$ ,  $0.1\text{ mol\%}$  catalyst loading and using  $1\text{ bar}$   $\text{CO}_2$  pressure. The conditions were selected as they have previously been shown to result in reasonable activity for related catalysts and to enable comparison against the existing literature.<sup>13,20</sup> Within the series of compounds, the di-zinc catalyst (**1**) shows the highest activity ( $9\text{ h}^{-1}$ ) and yields the highest carbonate linkage content ( $\text{CO}_2$  uptake) and polymer selectivity (minor quantities of

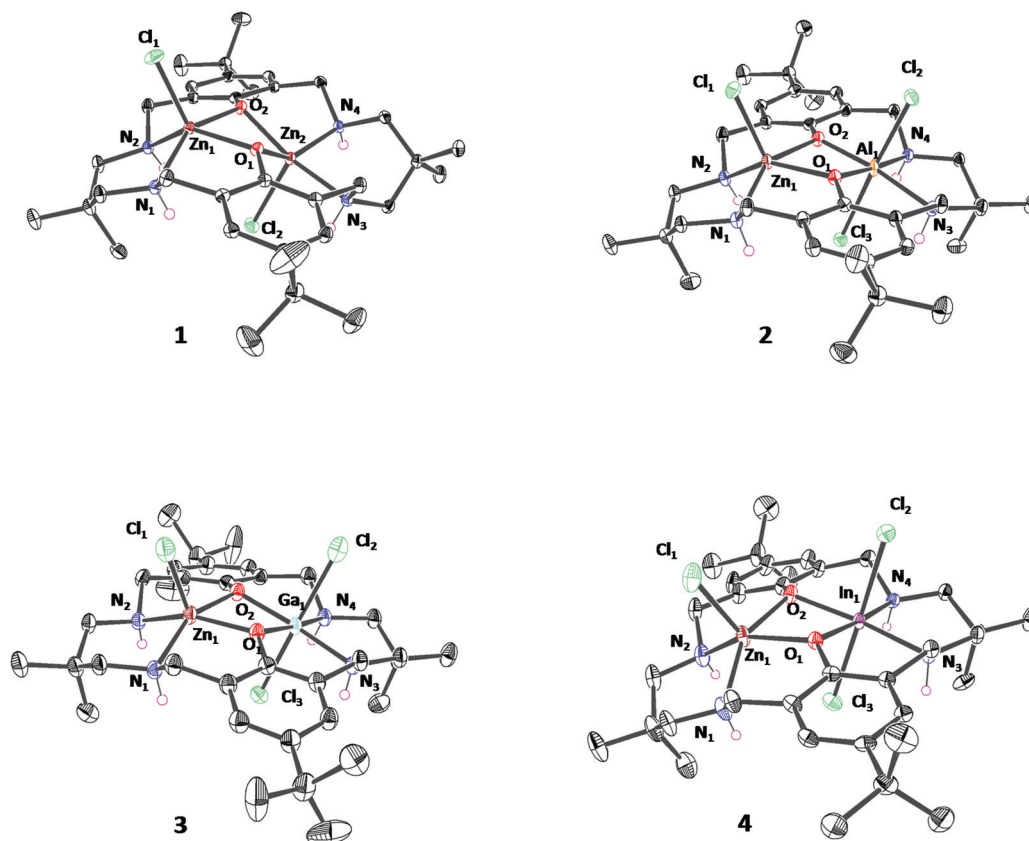


Fig. 3 ORTEP representation of the crystallographic structure of complex **1–4** and **6** obtained by single crystal X-ray diffraction. Disorder and H-atoms (excluding NH) have been omitted for clarity, with thermal ellipsoids represented at 40% probability. Selected bond lengths ( $\text{\AA}$ ) and angles ( $^\circ$ ) see Tables S1–S5.†



*trans*-cyclohexene carbonate are observed by  $^1\text{H}$  NMR spectroscopy). Indeed, its performance is equivalent to the previously reported di-zinc acetate analogue.<sup>13</sup> Heterodinuclear tri-chloride complexes (2–4) all show low activity and yield polymers with significant ether linkage contents and/or polyether contamination (up to 32%). These polyethers form as a result of the metal-alkoxide catalytic intermediate under-going sequential epoxide ring opening of an epoxide molecule (see Fig. S36,† for illustrations of the reactions occurring in the catalytic cycle and side-reactions). All heterodinuclear complexes show high polymer selectivity and there is very little cyclic carbonate by-product formation.

The polymers' molecular masses were analysed using GPC and, in most cases, bimodal molecular mass distributions were apparent. Multi-modal molecular mass distributions are very common in ROCOP reactions and are attributed to two series of chains resulting from different initiating groups.<sup>49</sup> It is proposed that the lower molecular mass series are initiated from the catalyst M–Cl, whereas the higher molecular mass series are from 1,2-cyclohexanediol. The 1,2-cyclohexanediol is proposed to form by reaction between cyclohexene oxide and residual water (e.g. in the  $\text{CO}_2$ ). Darensbourg and co-workers have established the rates of such hydrolyses typically exceed polymerization initiation and propagation rates.<sup>50–52</sup> In the case of catalyst 4 at 1 bar  $\text{CO}_2$  pressure a trimodal molecular mass distribution was observed, with peaks being tentatively attributed to polyether and two polycarbonate series, respectively (Table 1, entry 4).<sup>30</sup> At 20 bar pressure, catalyst 4 does not produce any polyether and the expected bimodal molecular mass distribution is observed (Table 1, entry 5).

In terms of polymerization selectivity, the catalysts show decreasing amounts of ether linkages on descending the series and the complexes of Ga(III) and In(III) show reasonable/good selectivity. This change in selectivity is proposed to result from the lower Lewis acidity of heavier Group 13 congeners compared to Al(III) and highlights the importance of appropriate balance of Lewis acidity in controlling side-reactions. The high ether content may also signal that  $\text{CO}_2$  insertion is implicated in the rate limiting step and thus complex 4 was tested with 20 bar  $\text{CO}_2$  pressure. Under these conditions a perfectly alternating copolymer formed, with no ether linkages. In the higher pressure regime, there is also a significant increase in activity ( $28\text{ h}^{-1}$ ) which may be a preliminary indication of a rate law dependent on  $\text{CO}_2$  pressure or from the improved reaction stirring in the high pressure autoclaves.

All polymerisation catalysts show induction periods prior to the start of the catalysis. To rule out that metal redistribution reactions occur during these induction periods, catalysts 2–4 were heated at 100 °C for 3 days in THF (Fig. S37–39†). In all cases, the  $^1\text{H}$  NMR spectra were identical before and after prolonged heating, i.e. there was no evidence for the formation of dizinc 1. The low polymerisation activity and significant induction periods of these ZnM(III) complexes could be due to coordinative saturation retarding epoxide coordination and activation. To investigate whether this could be overcome, cationic complex 5 showing a 'free' coordination site was prepared. Unfortunately, it rapidly forms a polymer which is almost entirely polyether, probably due to the increased Lewis acidity of the In(III) site.<sup>53</sup>

A recent investigation into analogous ZnMg catalysts showed that by applying carboxylate ligands, in place of

**Table 1** Data for the ROCOP of cyclohexene oxide and 1 bar  $\text{CO}_2$  using Catalysts 1–6<sup>a</sup>

Entry	Catalyst	Time (d)	TON <sup>b</sup>	TOF <sup>c</sup> ( $\text{h}^{-1}$ )	$\text{CO}_2$ <sup>d</sup> (%)	Polym. <sup>e</sup> (%)	$M_n$ [D] <sup>f</sup> ( $\text{g mol}^{-1}$ )
1	1	2	417	9	>99	98	20 500 [1.04] 8800 [1.13]
2	2	3	24	0.3	68	88	n.d.
3	3	3	69	1	85	95	n.d.
4	4	3	399	6	85	91	16 800 [1.07] 6600 [1.07] 2700 [1.12] 12 400 [1.03] 5100 [1.18]
5	4 <sup>g</sup>	0.33	221	28	>99	>99	n.d.
6	5	0.25	696	116	<1	>99	n.d.
7	6	3	227	3	>99	90	11 200 [1.04] 4900 [1.12]
8	ZnMgBr <sub>2</sub> <sup>18</sup>	0.25	247	78	>99	>99	3000 [1.18]
9	PSalan-InCl <sup>h,30</sup>	2	350	15	>99	95	19 700 [2.00] 3400 [1.32]
10	Salan-AlCl + Bu <sub>4</sub> NCl <sup>i,21</sup>	0.33	404	35	98	3	n.d.
11	$\beta$ -Keto-Al <sub>2</sub> Me + Bu <sub>4</sub> NOAc <sup>j,47</sup>	0.25	36	15	>99	97	4200 [1.21]
12	Amino tris(phenolate)AlCl + PPNCI <sup>k,48</sup>	2	154	3	>99	>99	11 900 [1.49]

<sup>a</sup> Catalysis conditions: catalyst : CHO 1 : 1000, 80 °C, 1 bar pressure  $\text{CO}_2$  and in neat epoxide. <sup>b</sup> Turnover number (TON) = number of moles of cyclohexene oxide consumed/number of moles of catalyst. <sup>c</sup> Turnover frequency (TOF) = TON/time (h). <sup>d</sup> Expressed as a percentage of  $\text{CO}_2$  uptake vs. the theoretical maximum (100%), determined by comparison of the relative integrals of the  $^1\text{H}$  NMR proton resonances due to carbonate ( $\delta$  4.65 ppm) and ether ( $\delta$  3.45 ppm) linkages in the polymer backbone. <sup>e</sup> Expressed as a percentage of polymer formation vs. the theoretical maximum (100%), determined by comparison of the relative integrals of the  $^1\text{H}$  NMR proton resonances due to polymer (4.65 ppm) and *trans*-cyclic carbonate (4.00 ppm). <sup>f</sup> Determined by SEC, in THF, calibrated using narrow- $M_n$  polystyrene standards. Values for Zn/Al and Zn/Ga were not determined as the conversion was too low. <sup>g</sup> 20 bar  $\text{CO}_2$ . <sup>h</sup> 60 °C. <sup>i</sup> 35 bar  $\text{CO}_2$ . <sup>j</sup> 50 bar  $\text{CO}_2$ . <sup>k</sup> 15 bar  $\text{CO}_2$ , 40 °C, 0.5 mol% cat loading (Fig. S35† for molecular structures).



halides, it was possible to significantly enhance turn-over-frequency values.<sup>18</sup> Nonetheless, in this series of catalysts complex **6**, featuring a *p*-CF<sub>3</sub>benzoate ligand, showed no discernible difference in rate or selectivity, but a slight decrease in ether linkages, compared with **4**. Attempts to further substitute the two remaining chloride ligands in **6** with *p*-CF<sub>3</sub>benzoate groups were unsuccessful and resulted in broad, complex NMR spectra.

In terms of the catalytic activity of the series of ZnM(III) catalysts, there is a marginal increase on descending the series of Group 13 elements from ~1 h<sup>-1</sup> (Al(III)) to 9 h<sup>-1</sup> (In(III)). This increasing activity may correlate with the decreasing M–O bond strengths and the increasing lability of the metal alkoxide intermediates. Overall, it is quite clear that the activity values for these complexes are very low and particularly compared to leading catalysts in this field.<sup>18,19,54,55</sup> The Zn(II)Mg(II) catalyst shows a TOF of ~78 h<sup>-1</sup> and quantitative selectivity for carbonate linkages, under the same conditions (Table 1). Nonetheless, it is worth noting that the activities are in-line with other (previously reported) Group 13 catalysts (Table 1).<sup>56</sup> For example, the high pressure CO<sub>2</sub> activity of **4** (ZnIn(III)) is approximately equivalent to that of a salanAl/Bu<sub>4</sub>NCl system and significantly greater than a di-Al(III) catalyst system (β-ketoAl<sub>2</sub>Et/Bu<sub>4</sub>NOAc) or the amino tris(phenolate)Al/PPNCl systems (Table 1).<sup>21,47,48</sup> Overall, these heterodinuclear ZnM(III) catalysts show some promise in terms of activity and selectivity, particularly compared to other Group 13 catalysts. Future investigations should target ancillary ligands capable of reducing the Group 13 element Lewis acidity, and associated M–O bond strength, so as to increase activity and selectivity for carbonate linkage formation.

## Conclusions

The syntheses and characterisations of a series of heterodinuclear Zn(II)/Al(III), Ga(III) or In(III) complexes, coordinated by a symmetrical macrocycle are reported. The compounds are synthesised using a sequential metalation route, operating under thermodynamic control, which yields the heterodinuclear complexes as the major reaction products. The complexes are applied as homogeneous catalysts for cyclohexene oxide/CO<sub>2</sub> ring opening copolymerisation but all show worse performance than the di-zinc analogue. It seems that the enhanced Lewis acidity and significant M–O bond strengths of these Group 13 elements, compared to Zn(II), are less desirable in this catalysis. The ability to use inexpensive, colourless and abundant elements, such as Al(III) and Mg(II), remains an important goal for the field. Future work needs to address optimising the ligand design so as to improve catalytic activity and selectivity.

## Experimental section

All experimental manipulations were performed using a dual-manifold nitrogen-vacuum Schlenk line or in a nitrogen filled

glovebox. All solvents and reagents were obtained from commercial sources and used as received unless stated otherwise. THF, acetonitrile and pentane were obtained from an SPS system, degassed by several freeze–pump–thaw cycles, further dried with 3 Å molecular sieves and stored under nitrogen. Cyclohexene oxide was dried over calcium hydride (2 days) and purified *via* fractional distillation prior to use and stored under a nitrogen atmosphere.

Low pressure copolymerisation studies were performed using a triple manifold youngs tap CO<sub>2</sub>/N<sub>2</sub>/vacuum Schlenk line where research-grade carbon dioxide was dried through a drierite column, and two addition drying columns (Micro Torr, Model number: MC1-804FV) in series, before use in copolymerisation studies. High pressure copolymerisation studies were performed in a 25 mL Parr 5500 HP Compact Reactor using research-grade carbon dioxide that was dried by passing through two drying columns in series (VICI, Thames Restek).

<sup>1</sup>H, <sup>11</sup>B, <sup>13</sup>C, <sup>19</sup>F and 2D NMR spectra were obtained using a Bruker AV 400 MHz spectrometer at 298 K, unless stated otherwise. MALDI-ToF analysis was performed on a Micromass MALDI micro MX spectrometer. The matrix used was *trans*-2-[3-(4-*tert*-butylphenyl)-2-methyl-2-propenylidene]-malonitrile.

Gel permeation chromatography analysis was carried out on a two mixed bed PSS SDV linear S column in series, with THF as the eluent, at a flow rate of 1 mL min<sup>-1</sup>, on a Shimadzu LC-20AD instrument, at 40 °C. Polymer molecular mass values (*M<sub>n</sub>*) were obtained by calibration of the instrument using a series of narrow molecular mass polystyrene standards and are reported without correction.

Elemental Analysis was determined by Mr Stephen Boyer at London Metropolitan University.

### General procedure of complexes 1–4 synthesis

ZnEt<sub>2</sub> (0.11 g, 0.91 mmol) was added dropwise to a solution of H<sub>2</sub>L (0.50 g, 0.91 mmol) in THF (10 mL) and stirred for 16 h at 25 °C. ZnCl<sub>2</sub> or MCl<sub>3</sub> (M = Al, Ga or In) (0.91 mmol) in THF (5 mL) was added to the stirring solution (LZn) and the reaction heated to 100 °C for 16 h. The solvent was removed *in vacuo* to afford the products as white powders.

**Complex 1.** (0.55 g, 0.73 mmol, 81%). <sup>1</sup>H NMR (d<sub>2</sub>-TCE, 400.20 MHz, 403 K) 6.99 (s, 4H, PhH) 4.77 (s, 4H, Zn–HN–CH<sub>2</sub>–Ph) 3.38 (s, 4H, Zn–HN–CH<sub>2</sub>–Ph) 3.03 (s, 4H, Zn–NH) 2.82 (s, 4H, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.50 (s, 4H, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 1.40–1.30 (s, 24 H, C(CH<sub>3</sub>)<sub>2</sub>, C(CH<sub>3</sub>)<sub>3</sub>) 1.12 (s, 6H, C(CH<sub>3</sub>)<sub>2</sub>). <sup>13</sup>C NMR (d<sub>2</sub>-TCE, 125.81 MHz, 403 K) 160.5 (*ipso*-Ph) 140.3 (*p*-Ph) 128.3 (*m*-Ph) 124.9 (*o*-Ph) 64.5 (Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 58.7 (Zn–HN–CH<sub>2</sub>–Ph) 35.2 (C(CH<sub>3</sub>)<sub>2</sub>) 34.9 (C(CH<sub>3</sub>)<sub>3</sub>) 32.9 (C(CH<sub>3</sub>)<sub>3</sub>) 29.4 (C(CH<sub>3</sub>)<sub>2</sub>) 25.3 (C(CH<sub>3</sub>)<sub>2</sub>). Elemental analysis for C<sub>34</sub>H<sub>54</sub>Cl<sub>2</sub>N<sub>4</sub>O<sub>2</sub>Zn<sub>2</sub> (748.22 g mol<sup>-1</sup>): calculated; C, 54.3; H, 7.2; N, 7.5%. Found; C, 54.5; H, 7.4; N, 7.6%. MS (MALDI-ToF): *m/z* 717 [LZnZnCl]<sup>+</sup> (90%).

**Complex 2.** (0.63 g, 0.84 mmol, 95%). <sup>1</sup>H NMR (d<sub>3</sub>-MeCN, 400.20 MHz, 298 K): 7.06 (d, 4H, Al–PhH, Zn–PhH) 5.04–4.89 (m, 4H, Al–HN–CH<sub>2</sub>–Ph, Zn–HN–CH<sub>2</sub>–Ph) 3.65 (t, 2H, <sup>3</sup>*J* = 11.45 Hz, Al–NH) 3.35 (d, 2H, <sup>3</sup>*J* = 13.17 Hz, Zn–HN–CH<sub>2</sub>–Ph) 3.29 (s, 2H, <sup>3</sup>*J* = 12.60 Hz, Al–HN–CH<sub>2</sub>–Ph) 2.93 (t, 2H,



$^3J = 11.45$  Hz, Al–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.83 (t, 2H,  $^3J = 13.17$  Hz, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.78–2.64 (m, 4H, Al–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.54 (t, 2H,  $^3J = 11.99$  Hz, Zn–NH) 1.36 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 1.27 (s, 18H, C(CH<sub>3</sub>)<sub>3</sub>) 1.07 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 0.93 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>). <sup>13</sup>C NMR (d<sub>3</sub>-MeCN, 125.81 MHz, 298 K) 155.0 (*ipso*-Ph) 141.5 (*p*-Ph) 129.1 (*m*-PhZn) 127.7 (*m*-PhAl) 124.7 (*o*-PhZn) 64.8 (Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 63.2 (Al–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 57.1 (Zn–HN–CH<sub>2</sub>–Ph) 56.7 (Al–HN–CH<sub>2</sub>–Ph) 34.2 (Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 34.0 (Al–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 33.3 (C(CH<sub>3</sub>)<sub>3</sub>) 31.6 (C(CH<sub>3</sub>)<sub>3</sub>) 27.9 (C(CH<sub>3</sub>)<sub>2</sub>) 27.7 (C(CH<sub>3</sub>)<sub>2</sub>) 22.1 (C(CH<sub>3</sub>)<sub>2</sub>) 20.4 (C(CH<sub>3</sub>)<sub>2</sub>). Elemental analysis for C<sub>34</sub>H<sub>54</sub>AlCl<sub>3</sub>N<sub>4</sub>O<sub>2</sub>Zn (746.24 g mol<sup>-1</sup>): calculated; C, 54.5; H, 7.3; N, 7.5%. Found; C, 54.0; H, 6.9; N, 7.2%. MS (MALDI-ToF): *m/z* 714 [LZnAlCl<sub>2</sub>]<sup>+</sup> (90%).

**Complex 3.** (0.51 g, 0.65 mmol, 72%). <sup>1</sup>H NMR (d<sub>3</sub>-MeCN, 400.20 MHz, 298 K): 7.06 (d, 2H,  $^4J = 2.63$  Hz, Ga–PhH) 7.03 (d, 2H,  $^4J = 2.63$  Hz, Zn–PhH) 4.99 (t, 2H,  $^3J = 12.92$  Hz, Ga–HN–CH<sub>2</sub>–Ph) 4.87 (t, 2H,  $^3J = 11.90$  Hz, Zn–HN–CH<sub>2</sub>–Ph) 3.71 (t, 2H,  $^3J = 11.22$  Hz, Ga–NH) 3.26 (d, 2H,  $^3J = 12.41$  Hz, Ga–HN–CH<sub>2</sub>–Ph) 3.25 (d, 2H,  $^3J = 12.66$  Hz, Zn–HN–CH<sub>2</sub>–Ph) 3.03 (t, 2H,  $^3J = 12.29$  Hz, Ga–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.86 (t, 2H,  $^3J = 12.78$  Hz, Zn–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.73 (t, 2H,  $^3J = 11.55$  Hz, Ga–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.66 (t, 2H,  $^3J = 10.20$  Hz, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.43 (t, 2H,  $^3J = 10.08$  Hz, Zn–NH) 1.35 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 1.28 (s, 18H, C(CH<sub>3</sub>)<sub>3</sub>) 1.05 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 0.96 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 0.93 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>). <sup>13</sup>C NMR (d<sub>3</sub>-MeCN, 125.81 MHz, 298 K): 157.2 (*ipso*-Ph) 140.9 (*p*-Ph) 129.4 (Zn–*m*-Ph) 128.7 (Ga–*m*-Ph) 125.2 (Ga–*o*-Ph) 124.2 (Zn–*o*-Ph) 64.1 (Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 63.3 (Ga–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 56.9 (Zn–HN–CH<sub>2</sub>–Ph) 56.7 (Ga–HN–CH<sub>2</sub>–Ph) 33.6 (C(CH<sub>3</sub>)<sub>3</sub>) 34.2 (C(CH<sub>3</sub>)<sub>2</sub>) 34.0 (C(CH<sub>3</sub>)<sub>2</sub>) 31.7 (C(CH<sub>3</sub>)<sub>3</sub>) 27.9 (C(CH<sub>3</sub>)<sub>2</sub>) 26.3 (C(CH<sub>3</sub>)<sub>2</sub>) 22.0 (C(CH<sub>3</sub>)<sub>2</sub>) 20.5 (C(CH<sub>3</sub>)<sub>2</sub>). Elemental analysis for C<sub>34</sub>H<sub>54</sub>Cl<sub>3</sub>GaN<sub>4</sub>O<sub>2</sub>Zn (788.19 g mol<sup>-1</sup>): calculated; C, 51.5; H, 6.9; N, 7.1%. Found; C, 51.7; H, 6.9; N, 7.1%. MS (MALDI-ToF): *m/z* 755 [LZnGaCl<sub>2</sub>]<sup>+</sup> (100%).

**Complex 4.** (0.65 g, 0.78 mmol, 86%). <sup>1</sup>H NMR (d<sub>8</sub>-THF, 400.20 MHz, 298 K) 6.99 (d, 2H,  $^4J = 2.10$  Hz, In–PhH) 6.96 (d, 2H,  $^4J = 1.96$  Hz, Zn–PhH) 5.15 (t, 2H,  $^3J = 11.87$  Hz, In–HN–CH<sub>2</sub>–Ph) 4.98 (t, 2H,  $^3J = 11.65$  Hz, Zn–HN–CH<sub>2</sub>–Ph) 3.77 (t, 2H,  $^3J = 10.68$  Hz, In–NH) 3.37 (t, 2H,  $^3J = 12.04$  Hz, In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 3.26 (d, 2H,  $^3J = 12.95$  Hz, In–HN–CH<sub>2</sub>–Ph) 3.11 (d, 2H,  $^3J = 12.27$  Hz, Zn–HN–CH<sub>2</sub>–Ph) 3.05 (t, 2H,  $^3J = 12.04$  Hz, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.68 (d, 2H,  $^3J = 12.04$  Hz, In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.60 (d, 2H,  $^3J = 11.36$  Hz, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.50 (t, 2H,  $^3J = 11.59$  Hz, Zn–NH) 1.31 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>) 1.25 (s, 18H, C(CH<sub>3</sub>)<sub>3</sub>) 1.09 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>) 0.99 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>) 0.95 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>). <sup>13</sup>C NMR (d<sub>8</sub>-THF, 125.81 MHz, 298 K) 159.7 (*ipso*-Ph) 138.8 (*p*-Ph) 129.4 (Zn–*m*-Ph) 128.4 (In–*m*-Ph) 126.5 (In–*o*-Ph) 123.8 (Zn–*o*-Ph) 63.3 (Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 61.7 (In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 57.0 (Zn–HN–CH<sub>2</sub>–Ph) 56.1 (In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 35.0 (C(CH<sub>3</sub>)<sub>2</sub>) 34.2 (C(CH<sub>3</sub>)<sub>2</sub>) 34.0 (–C(CH<sub>3</sub>)<sub>3</sub>) 31.7 (–C(CH<sub>3</sub>)<sub>3</sub>) 28.9 (–C(CH<sub>3</sub>)<sub>2</sub>) 28.2 (–C(CH<sub>3</sub>)<sub>2</sub>) 21.7 (–C(CH<sub>3</sub>)<sub>2</sub>) 20.7 (C(CH<sub>3</sub>)<sub>2</sub>). Elemental analysis for C<sub>34</sub>H<sub>54</sub>Cl<sub>3</sub>InN<sub>4</sub>O<sub>2</sub>Zn (834.16 g mol<sup>-1</sup>): calculated; C, 48.8; H, 6.5; N, 6.7%. Found; C, 48.7; H, 6.6; N, 6.6%. MS (MALDI-ToF): *m/z* 799 [LZnInCl<sub>2</sub>]<sup>+</sup> (90%).

**Complex 5.** K[B(C<sub>6</sub>F<sub>5</sub>)<sub>4</sub>] (0.33 g, 0.46 mmol) was added to a solution of complex 4 (0.38 g, 0.46 mmol) in THF (5 mL) and

stirred for 16 h at 25 °C. The solution was filtered and reduced *in vacuo* to dryness. The product was extracted into toluene and washed with pentane (3 × 5 mL) to afford a white solid (0.61 g, 0.41 mmol, 90%). <sup>1</sup>H NMR (d<sub>8</sub>-THF, 400.20 MHz, 298 K) 7.06 (s, 2H, In–PhH) 7.03 (s, 2H, Zn–PhH) 5.05 (t, 2H,  $^3J = 11.87$  Hz, In–HN–CH<sub>2</sub>–Ph) 4.84 (t, 2H,  $^3J = 11.65$  Hz, Zn–HN–CH<sub>2</sub>–Ph) 4.30 (t, 2H,  $^3J = 10.68$  Hz, In–NH) 3.36 (t, 2H,  $^3J = 12.04$  Hz, In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 3.28 (d, 2H,  $^3J = 12.95$  Hz, In–HN–CH<sub>2</sub>–Ph) 3.23 (d, 2H,  $^3J = 12.27$  Hz, Zn–HN–CH<sub>2</sub>–Ph) 2.96 (t, 2H,  $^3J = 12.04$  Hz, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.90 (d, 2H,  $^3J = 12.04$  Hz, In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.75 (d, 2H,  $^3J = 11.36$  Hz, Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.69 (t, 2H,  $^3J = 11.59$  Hz, Zn–NH) 1.27 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>) 1.25 (s, 18H, C(CH<sub>3</sub>)<sub>3</sub>) 1.09 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>) 1.00 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>) 0.97 (s, 3H, –C(CH<sub>3</sub>)<sub>2</sub>). <sup>11</sup>B NMR (d<sub>8</sub>-THF, 128.39 MHz, 298 K) –16.6 (s, B(ArF<sub>5</sub>)<sub>4</sub>). <sup>13</sup>C<sup>57</sup> NMR (d<sub>8</sub>-THF, 125.81 MHz, 298 K) 158.2 (*ipso*-Ph) 149.2 (br. *ipso*-BArF) 147.2 (br. *p*-BArF) 139.6 (*p*-Ph) 137.2 (br. *m*-BArF) 135.1 (br. *m*-BArF) 129.1 (Zn–*m*-Ph) 128.3 (In–*m*-Ph) 125.5 (In–*o*-Ph) 123.1 (Zn–*o*-Ph) 63.6 (Zn–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 61.3 (In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 56.0 (Zn–HN–CH<sub>2</sub>–Ph) 55.3 (In–HN–CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 34.2 (C(CH<sub>3</sub>)<sub>2</sub>) 33.3 (C(CH<sub>3</sub>)<sub>3</sub>) 30.8 (C(CH<sub>3</sub>)<sub>3</sub>) 27.8 (C(CH<sub>3</sub>)<sub>2</sub>) 27.2 (C(CH<sub>3</sub>)<sub>2</sub>) 20.7 (C(CH<sub>3</sub>)<sub>2</sub>) 19.8 (C(CH<sub>3</sub>)<sub>2</sub>). <sup>19</sup>F NMR (d<sub>8</sub>-THF, 470 MHz, 298 K) –132.7 (s, 8H, *o*-PhF) –165.1 (t, 4H,  $^3J = 24.4$  Hz, *p*-PhF) –168.5 (t, 8H, 22.3 Hz, *m*-PhF). Elemental analysis for C<sub>58</sub>H<sub>54</sub>BCl<sub>2</sub>F<sub>20</sub>InN<sub>4</sub>O<sub>2</sub>Zn (1478.17 g mol<sup>-1</sup>): calculated; C, 47.0; H, 3.7; N, 3.8%. Found; C, 46.7; H, 4.0; N, 4.1%.

**Complex 6.** KOBzPCF<sub>3</sub> (0.01 g, 0.05 mmol) was added to a solution of complex 4 (0.04 g, 0.05 mmol) in THF (5 mL) and stirred for 16 h at 25 °C. The solution was filtered and reduced *in vacuo* to dryness. The product was extracted into toluene (0.5 mL) and further dried *in vacuo*. The crude product was washed with pentane (3 × 5 mL) to afford a white solid (0.05 g, 0.048 mmol, 95%) <sup>1</sup>H NMR (d<sub>4</sub>-MeOD, 500.20 MHz, 298 K) 7.96 (d, 2H,  $^3J = 8.1$  Hz, *ortho*-OBzPCF<sub>3</sub>) 7.53 (d, 2H,  $^3J = 8.4$  Hz, *meta*-OBzPCF<sub>3</sub>) 7.10 (d, 2H,  $^4J = 2.5$  Hz, Zn–PhH) 7.07 (d, 2H,  $^4J = 2.60$  Hz, In–PhH) 4.64 (d, 2H,  $^3J = 12.2$  Hz, Ph–CH<sub>2</sub>–NH) 4.25 (d, 2H,  $^3J = 12.2$  Hz, Ph–CH<sub>2</sub>–NH) 3.41 (d, 4H,  $^3J = 12.4$  Hz, Ph–CH<sub>2</sub>–NH) 3.33 (d, 2H,  $^3J = 12.1$  Hz, CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.95 (d, 2H,  $^3J = 12.1$  Hz, CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.78 (d, 2H,  $^3J = 12.1$  Hz, CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 2.75 (d, 2H,  $^3J = 12.1$  Hz, CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 1.50 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 1.43 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 1.19 (s, 18 H, C(CH<sub>3</sub>)<sub>3</sub>) 1.08 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>) 1.05 (s, 3H, C(CH<sub>3</sub>)<sub>2</sub>). <sup>13</sup>C NMR (d<sub>4</sub>-MeOD, 125.81 MHz, 298 K) 170.3 (CO<sub>2</sub>Ph) 159.3 (*ipso*-OPh) 139.4 (*ipso*-OBz) 138.0 (*ortho*-OPh) 130.0 (*ortho*-OBz) 129.5 (*meta*-OPh) 129.0 (*meta*-OBz) 124.2 (*para*-OBz) 124.1 (*para*-OBz) 61.8 (CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 60.3 (CH<sub>2</sub>–C(CH<sub>3</sub>)<sub>2</sub>) 54.8 (Ph–CH<sub>2</sub>–NH) 54.3 (Ph–CH<sub>2</sub>–NH) 34.3 (C(CH<sub>3</sub>)<sub>2</sub>) 33.1 (C(CH<sub>3</sub>)<sub>3</sub>) 30.5 (C(CH<sub>3</sub>)<sub>3</sub>) 27.5 (C(CH<sub>3</sub>)<sub>2</sub>) 27.3 (C(CH<sub>3</sub>)<sub>2</sub>) 20.9 (C(CH<sub>3</sub>)<sub>2</sub>) 20.3 (C(CH<sub>3</sub>)<sub>2</sub>). <sup>19</sup>F NMR (d<sub>4</sub>-MeOD, 376.64 MHz, 298 K): –64.4 (s, CF<sub>3</sub>). Elemental analysis for C<sub>42</sub>H<sub>58</sub>Cl<sub>2</sub>F<sub>3</sub>InN<sub>4</sub>O<sub>2</sub>Zn (988.21 g mol<sup>-1</sup>): calculated; C, 50.9; H, 5.9; N, 5.7%. Found; C, 50.3; H, 6.1; N, 6.0%.

#### General procedure for the synthesis of polycarbonates from CO<sub>2</sub>/epoxide, at 1 bar CO<sub>2</sub> pressure

The catalyst (15 μmol) was dissolved in CHO (15 mmol) and placed under 1 bar CO<sub>2</sub> in a Schlenk tube and was heated to



80 °C, at a stirring rate of 350 RPM. Aliquots were taken at regular intervals and quenched by addition, in air, into chloroform.

### General procedure for the synthesis of polycarbonates from CO<sub>2</sub>/epoxide, at 20 bar CO<sub>2</sub> pressure

The catalyst (60 μmol) was dissolved in CHO (60 mmol) and placed into a 25 mL Parr Reactor inside a nitrogen filled glove-box. The reactor was removed from the glove box and charged with CO<sub>2</sub>, to a pressure of 20 bar, and heated to 80 °C.

### X-ray crystallography

Crystalline samples were isolated in a glovebox, under a pool of fluorinated oil, and mounted on MiTeGen MicroMounts. The crystals were cooled to 150 K, using an Oxford Cryostream nitrogen cooling device. Data collection was carried out with an Oxford Diffraction supernova diffractometer using Cu K $\alpha$  ( $\lambda = 1.5417 \text{ \AA}$ ) radiation. The resulting raw data was processed using CrysAlis Pro.<sup>58</sup> Structures were solved using SHELXT and Full-matrix least-squares refinements based on  $F^2$  were performed in SHELXL-14,<sup>59</sup> as incorporated in the WinGX package.<sup>60</sup> For each methyl group, the hydrogen atoms were added at calculated positions, using a riding model with  $U(\text{H}) = 1.5U_{\text{eq}}$  (bonded carbon atom). The rest of the hydrogen atoms were included in the model at calculated positions, using a riding model with  $U(\text{H}) = 1.2U_{\text{eq}}$  (bonded atom). Neutral atom scattering factors were used and included terms for anomalous dispersion.<sup>61</sup> All crystal structures have been registered with the Cambridge structural database: CCDC 1939693–1939697.†

### Conflicts of interest

CKW is a director of Eonic Technologies.

### Acknowledgements

The EPSRC (EP/L017393/1, EP/K014668/1) and eonic technologies (CASE award to AD) are acknowledged for research funding.

### References

- 1 J. Artz, T. E. Müller, K. Thenert, J. Kleinekorte, R. Meys, A. Sternberg, A. Bardow and W. Leitner, *Chem. Rev.*, 2018, **118**, 434–504.
- 2 N. von der Assen and A. Bardow, *Green Chem.*, 2014, **16**, 3272–3280.
- 3 B. Grignard, S. Gennen, C. Jérôme, A. W. Kleij and C. Detrembleur, *Chem. Soc. Rev.*, 2019, **48**, 4466–4514.
- 4 A. J. Kamphuis, F. Picchioni and P. P. Pescarmona, *Green Chem.*, 2019, **21**, 406–448.
- 5 J. Langanke, A. Wolf, J. Hofmann, K. Böhm, M. A. Subhani, T. E. Müller, W. Leitner and C. Gürtler, *Green Chem.*, 2014, **16**, 1865–1870.
- 6 W. Kuran, M. Sobczak, T. Listos, C. Debek and Z. Florjanczyk, *Polymer*, 2000, **41**, 8531–8541.
- 7 P. Alagi, R. Ghorpade, Y. J. Choi, U. Patil, I. Kim, J. H. Baik and S. C. Hong, *ACS Sustainable Chem. Eng.*, 2017, **5**, 3871–3881.
- 8 M. Scharfenberg, J. Seiwert, M. Scherger, J. Preis, M. Susewind and H. Frey, *Macromolecules*, 2017, **50**, 6577–6585.
- 9 M. DeBolt, A. Kiziltas, D. Mielewski, S. Waddington and M. J. Nagridge, *J. Appl. Polym. Sci.*, 2016, **133**, 9.
- 10 M. D. Burkart, N. Hazari, C. L. Tway and E. L. Zeitler, *ACS Catal.*, 2019, **9**, 7937–7956.
- 11 Y. Wang and D. J. Darensbourg, *Coord. Chem. Rev.*, 2018, **372**, 85–100.
- 12 C. M. Kozak, K. Ambrose and T. S. Anderson, *Coord. Chem. Rev.*, 2018, **376**, 565–587.
- 13 M. R. Kember, P. D. Knight, P. T. R. Reung and C. K. Williams, *Angew. Chem., Int. Ed.*, 2009, **48**, 931–933.
- 14 M. R. Kember, A. J. P. White and C. K. Williams, *Macromolecules*, 2010, **43**, 2291–2298.
- 15 A. Buchard, M. R. Kember, K. G. Sandeman and C. K. Williams, *Chem. Commun.*, 2011, **47**, 212–214.
- 16 M. R. Kember and C. K. Williams, *J. Am. Chem. Soc.*, 2012, **134**, 15676–15679.
- 17 P. K. Saini, C. Romain and C. K. Williams, *Chem. Commun.*, 2014, **50**, 4164–4167.
- 18 G. Trott, J. A. Garden and C. K. Williams, *Chem. Sci.*, 2019, **10**, 4618–4627.
- 19 H. Nagae, R. Aoki, S. Akutagawa, J. Kleemann, R. Tagawa, T. Schindler, G. Choi, T. P. Spaniol, H. Tsurugi, J. Okuda and K. Mashima, *Angew. Chem., Int. Ed.*, 2018, **57**, 2492–2496.
- 20 A. C. Deacy, C. B. Durr, J. A. Garden, A. J. P. White and C. K. Williams, *Inorg. Chem.*, 2018, **57**, 15575–15583.
- 21 D. J. Darensbourg and D. R. Billodeaux, *Inorg. Chem.*, 2005, **44**, 1433–1442.
- 22 N. Ikpo, S. M. Barbon, M. W. Drover, L. N. Dawe and F. M. Kerton, *Organometallics*, 2012, **31**, 8145–8158.
- 23 T. Ohkawara, K. Suzuki, K. Nakano, S. Mori and K. Nozaki, *J. Am. Chem. Soc.*, 2014, **136**, 10728–10735.
- 24 T. Aida and S. Inoue, *J. Am. Chem. Soc.*, 1983, **105**, 1304–1309.
- 25 T. Aida, M. Ishikawa and S. Inoue, *Macromolecules*, 1986, **19**, 8–13.
- 26 H. Sugimoto, H. Ohtsuka and S. Inoue, *J. Polym. Sci., Part A: Polym. Chem.*, 2005, **43**, 4172–4186.
- 27 C. Chatterjee and M. H. Chisholm, *Inorg. Chem.*, 2012, **51**, 12041–12052.
- 28 H. Sugimoto, H. Goto, S. Honda, R. Yamada, Y. Manabe and S. Handa, *Polym. Chem.*, 2016, **7**, 3906–3912.
- 29 L. Peña Carrodeguas, J. González-Fabra, F. Castro-Gómez, C. Bo and A. W. Kleij, *Chem. – Eur. J.*, 2015, **21**, 6115–6122.
- 30 A. Thevenon, A. Cyriac, D. Myers, A. J. P. White, C. B. Durr and C. K. Williams, *J. Am. Chem. Soc.*, 2018, **140**, 6893–6903.





- 31 E. H. Nejad, C. G. W. van Melis, T. J. Vermeer, C. E. Koning and R. Duchateau, *Macromolecules*, 2012, **45**, 1770–1776.
- 32 N. J. Van Zee and G. W. Coates, *Angew. Chem., Int. Ed.*, 2015, **54**, 2665–2668.
- 33 J. Li, Y. Liu, W. M. Ren and X. B. Lu, *J. Am. Chem. Soc.*, 2016, **138**, 11493–11496.
- 34 M. E. Fieser, M. J. Sanford, L. A. Mitchell, C. R. Dunbar, M. Mandal, N. J. Van Zee, D. M. Urness, C. J. Cramer, G. W. Coates and W. B. Tolman, *J. Am. Chem. Soc.*, 2017, **139**, 15222–15231.
- 35 F. Isnard, F. Santulli, M. Cozzolino, M. Lamberti, C. Pellicchia and M. Mazzeo, *Catal. Sci. Technol.*, 2019, **9**, 3090–3098.
- 36 J. Li, B. H. Ren, S. Y. Chen, G. H. He, Y. Liu, W. M. Ren, H. Zhou and X. B. Lu, *ACS Catal.*, 2019, **9**, 1915–1922.
- 37 C. Bakewell, A. J. P. White, N. J. Long and C. K. Williams, *Inorg. Chem.*, 2013, **52**, 12561–12567.
- 38 K. Press, I. Goldberg and M. Kol, *Angew. Chem.*, 2015, **127**, 15071–15074.
- 39 K. M. Osten and P. Mehrkhodavandi, *Acc. Chem. Res.*, 2017, **50**, 2861–2869.
- 40 M. Chwatko and N. A. Lynd, *Macromolecules*, 2017, **50**, 2714–2723.
- 41 D. Myers, A. J. P. White, C. M. Forsyth, M. Bown and C. K. Williams, *Angew. Chem.*, 2017, **129**, 5361–5366.
- 42 T. Stößer, D. Mulryan and C. K. Williams, *Angew. Chem., Int. Ed.*, 2018, **57**, 16893–16897.
- 43 Y. Zhu, C. Romain and C. K. Williams, *J. Am. Chem. Soc.*, 2015, **137**, 12179–12182.
- 44 C. Romain, J. A. Garden, G. Trott, A. Buchard, A. J. P. White and C. K. Williams, *Chem. – Eur. J.*, 2017, **23**, 7367–7376.
- 45 M. R. Kember, A. Buchard and C. K. Williams, *Chem. Commun.*, 2011, **47**, 141–163.
- 46 S. Klaus, M. W. Lehenmeier, C. E. Anderson and B. Rieger, *Coord. Chem. Rev.*, 2011, **255**, 1460–1479.
- 47 K. Nishioka, H. Goto and H. Sugimoto, *Macromolecules*, 2012, **45**, 8172–8192.
- 48 C. Martín and A. W. Kleij, *Macromolecules*, 2016, **49**, 6285–6295.
- 49 F. Jutz, A. Buchard, M. R. Kember, S. B. Fredriksen and C. K. Williams, *J. Am. Chem. Soc.*, 2011, **133**, 17395–17405.
- 50 S. Inoue, *J. Polym. Sci., Part A: Polym. Chem.*, 2000, **38**, 2861–2871.
- 51 G.-P. Wu and D. J. Darensbourg, *Macromolecules*, 2016, **49**, 807–814.
- 52 D. J. Darensbourg, *Green Chem.*, 2019, **21**, 2214–2223.
- 53 D. Franz and S. Inoue, *Chem. – Eur. J.*, 2019, **25**, 2898–2926.
- 54 S. Kissling, M. W. Lehenmeier, P. T. Altenbuchner, A. Kronast, M. Reiter, P. Deglmann, U. B. Seemann and B. Rieger, *Chem. Commun.*, 2015, **51**, 4579–4582.
- 55 J. A. Garden, P. K. Saini and C. K. Williams, *J. Am. Chem. Soc.*, 2015, **137**, 15078–15081.
- 56 N. Ikpo, J. C. Flogeras and F. M. Kerton, *Dalton Trans.*, 2013, **42**, 8998–9006.
- 57 P. C. Andrikopoulos, D. R. Armstrong, H. R. L. Barley, W. Clegg, S. H. Dale, E. Hevia, G. W. Honeyman, A. R. Kennedy and R. E. Mulvey, *J. Am. Chem. Soc.*, 2005, **127**, 6184–6185.
- 58 *CrysAlis CCD, CrysAlis RED and associated programs: Oxford Diffraction*, Oxford Diffraction Ltd, Abingdon, England, 2006.
- 59 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 60 L. J. Farrugia, *J. Appl. Crystallogr.*, 2012, **45**, 849–854.
- 61 A. J. C. Wilson, *International Tables for Crystallography*, Kluwer Academic Publishers, Dordrecht, 1st edn, 1992, vol. C.

