



Cite this: *Dalton Trans.*, 2015, **44**, 12365

Magnesium amino-bis(phenolato) complexes for the ring-opening polymerization of *rac*-lactide†

Katalin Devaine-Pressing,^a Joshua H. Lehr,^a Michelle E. Pratt,^a Louise N. Dawe,^{‡a,b} Amy A. Sarjeant^c and Christopher M. Kozak*^a

Magnesium compounds of tetradentate amino-bis(phenolato) ligands, Mg[L1] (**1**) and Mg[L2] (**2**) (where [L1] = 2-pyridyl-*N,N*-bis(2-methylene-4-methoxy-6-*tert*-butylphenolato), and [L2] = dimethylamino-ethylamino-*N,N*-bis(2-methylene-4-methyl-6-*tert*-butylphenolato)) were prepared. The proligands, H₂[L1] and H₂[L2] were reacted with di(*n*-butyl)magnesium in toluene to give the desired compounds in high yields. Compounds **1** and **2** exhibit dimeric structures in solutions of non-coordinating solvents as observed by NMR spectroscopy and in the solid state as shown by the single crystal X-ray structure of **2**. These compounds exhibit good activity for *rac*-lactide polymerization in solution and in molten lactide.

Received 18th January 2015,
Accepted 17th March 2015

DOI: 10.1039/c5dt00236b

www.rsc.org/dalton

Introduction

Poly(lactide) continues to be at the forefront of biodegradable, biosourced and biocompatible polymer synthesis. It is central to the development of sustainable polymers, an area of critical importance for the design of new materials that reduce environmental and health impacts resulting from their manufacture, use, and their end-of-life properties.^{1–4} The array of different metal complexes studied for the ring-opening polymerization (ROP) of lactide has been described in several reviews^{5–8} and examples of metal-containing initiators vary considerably in terms of the metal centre and ligands employed. Complexes of amino-bis(phenolates) exhibit interesting structural and electronic properties, as well as catalytic behaviour; the variety of metal complexes with this ligand class has also been reviewed.⁹ Derivatives of this ligand class have been used in main group and d-block metal chemistry, including lithium,^{10–18} magnesium,^{19–25} calcium,²⁶ rare-earths,^{27,28} zinc,^{20,29–37} aluminum,^{7,38,39} zirconium^{40,41} and titanium.⁴² Many of these complexes have been reported to be excellent initiators for the ROP of cyclic esters such as lactide and ϵ -caprolactone. For the ROP of cyclic esters, earth

abundant alkali and alkaline earth metal compounds have become particularly important in the search for efficient, stable, inexpensive and non-toxic catalysts. We have previously reported the structures and catalytic lactide ROP activity of lithium and potassium amino-bis(phenolato) compounds.^{43,44} Magnesium compounds also have great potential as ROP catalysts for the preparation of PLA particularly for biomedical applications, as Mg is a biocompatible and non-toxic metal and thus the residue of these metals in the resulting poly(lactide) is not of great concern.^{45,46} Furthermore, magnesium compounds have been shown to elicit high reaction rates for copolymerization of CO₂ and epoxides.⁴⁷

Magnesium complexes reported by Ma and co-workers exhibited some of the highest activities in ROP of cyclic esters by this metal.^{48,49} The high activity for the ROP of *rac*-lactide may be due to the monomeric nature of these magnesium bis(silyl)amido complexes supported by tetradentate monophenolate ligands and the presence of a reactive monodentate amido ligand. A TOF of 36 560 h^{−1} was achieved when 10 000 equiv. of *rac*-lactide was converted to poly(lactide) in 15 minutes at 25 °C in the absence of any co-catalyst. One of these complexes showed very high activity even under melt conditions at 110 °C giving 72% conversion with a TOF of 86 880 h^{−1}. Magnesium compounds supported by tridentate monophenolate Schiff-base ligands and benzyl alkoxide also proved active for *l*-lactide polymerization in CH₂Cl₂ solutions.²¹ In both these cases, it is likely the monodentate amido or alkoxide ligand that serves as the initiation site.

In light of these high activities, we investigated the activity of magnesium amino-bis(phenolato) complexes towards ROP of *rac*-lactide under melt conditions and in solution. Magnesium complexes of aminophenolate ligands have previously been reported by others but their activity for ROP of lactides

^aDepartment of Chemistry, Memorial University of Newfoundland, St. John's, Newfoundland, Canada A1B 3X7. E-mail: ckozak@mun.ca; Tel: +1-709-864-8082

^bC-CART X-ray Diffraction Laboratory, Memorial University of Newfoundland, St. John's, Newfoundland, Canada

^cDepartment of Chemistry, Northwestern University, 2145 Sheridan Rd, Evanston, IL 60208, USA

† Electronic supplementary information (ESI) available: Crystallographic experimental and spectroscopic data (PDF). CCDC 1043113. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c5dt00236b

‡ Current Address: Department of Chemistry and Biochemistry, Wilfrid Laurier University, Waterloo, Ontario, Canada



was not tested or they showed very low or no activity.^{50–53} In this report, we present the synthesis and characterization of two new magnesium amine-bis(phenolate) complexes that show good lactide ROP activity in both melt and solution conditions.

Results and discussion

Synthesis and characterization of proligands and magnesium complexes

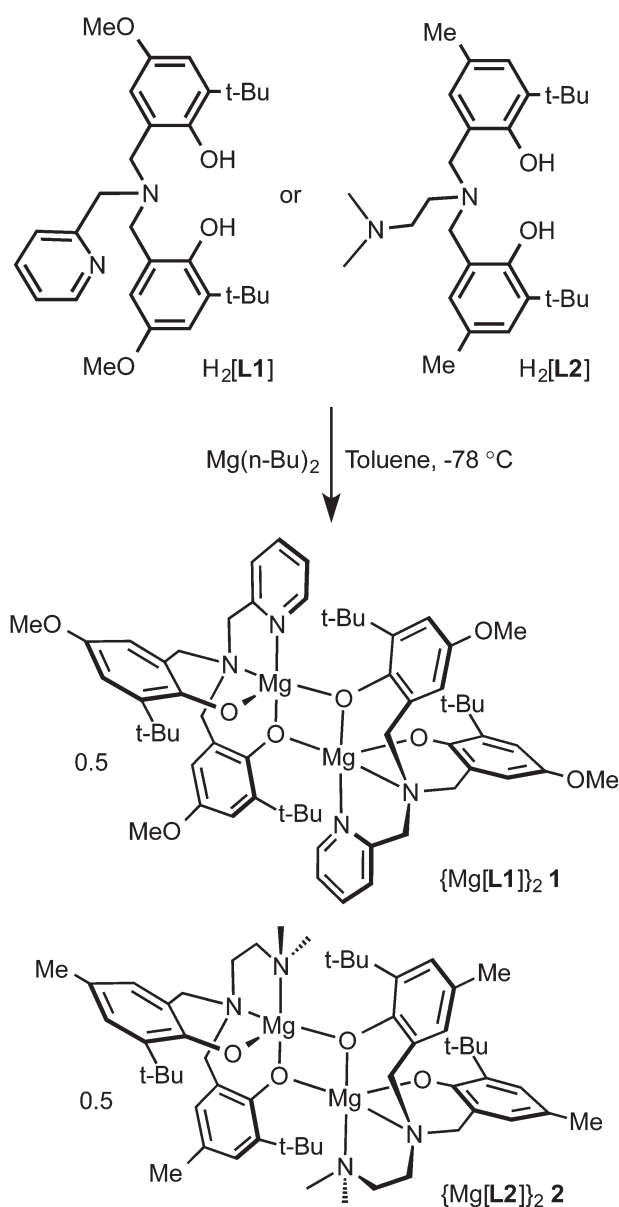
Tetradentate tripodal amine-bis(phenol) proligands (Scheme 1) were synthesized *via* a modified Mannich condensation reaction in water.⁵⁴ The substituted phenols contained *t*-butyl groups in the *ortho* positions and methoxy ($H_2[L1]$) or

methyl groups ($H_2[L2]$) *para* to the hydroxyl group. The proligands possessed pyridyl ($H_2[L1]$) or dimethylaminoethyl ($H_2[L2]$) groups as the neutral pendant N-donors. The synthesis of $H_2[L1]$ using a four step procedure⁵⁵ and without the addition of any further solvents⁵⁶ was previously reported. The synthesis of $H_2[L2]$ in methanol was reported by Bochmann and co-workers.⁵³

The corresponding ligand was reacted with one equivalent of di(*n*-butyl)magnesium in toluene at $-78\text{ }^\circ\text{C}$ to afford complexes **1** and **2** (Scheme 1). The NMR spectra of complexes **1** and **2** in non-coordinating solvents revealed that the compounds exist as dimers in solution. The $^1\text{H-NMR}$ spectrum of **2** in C_6D_6 showed two *t*-butyl environments implying inequivalent phenolates. This was further supported by the existence of four aromatic proton resonances arising from the phenolate rings occupying inequivalent coordination sites at the magnesium centres. The methylene resonances exist as four doublets. Two doublets are expected for a monometallic complex given the diastereotopic environments resulting from coordination to the metal centre. The existence of the four doublets arises from asymmetric phenolate sites where one phenolate of each ligand occupies a bridging position between two magnesium ions. The solid-state structure of compound **2** confirmed the proposed connectivity (see below). In coordinating solvents, such as pyridine or DMSO, the NMR spectra are consistent with monometallic compounds. The ^1H NMR spectra of **1** in C_6D_6 or toluene- d_8 showed the presence of several isomers suggesting the existence of dimeric species but also of more than one structural isomer (*e.g.* *syn* and *anti* oriented pendent donors) present in solution. Similar isomerism has been observed in zirconium amine-tris(phenolates).⁵⁷

MALDI-TOF mass spectrometry (see Fig. S1 and S2 in ESI†) showed complexes **1** and **2** exhibit dimeric structures in the gas phase as well as in solution and the solid state. Peaks at m/z 1028.50 (rel. intensity 100%) and m/z 514.26 (58%) corresponding to the dimer and monomer fragment ions, respectively, were observed for complex **1**. For complex **2**, the peak assigned to the dimer at m/z 924.52 was very weak (3%) and the peak at m/z 462.25 (100%) corresponds to the monomer.

Colourless crystals of **2** suitable for single crystal X-ray diffraction were obtained from saturated toluene solutions at $-35\text{ }^\circ\text{C}$. The solid-state molecular structure with selected bond lengths and angles is shown in Fig. 1. Crystallographic and structure refinement data can be found in Table S1 in ESI.† The dimeric structure with chemically distinct phenolate groups is consistent with the NMR studies discussed above. Each magnesium centre is five-coordinate and bound to two nitrogen donors and three oxygen donors. Four of the coordination sites are occupied by the chelating diamine-bis(phenolate) ligand, while the fifth coordination site is occupied by a bridging phenolate oxygen. The non-planar Mg_2O_2 core exhibits a $\text{O}(2)\text{-Mg}(1)\text{-O}(2)^*\text{-Mg}(1)^*$ torsion angle of $32.69(4)^\circ$. The magnesium complexes that show polymerization activity are believed to initiate cyclic ester ring-opening polymerizations through a coordination insertion mechanism;^{6,45} therefore, the creation of a vacant site by breaking the Mg_2O_2 ring (in the



Scheme 1 Synthesis of complexes **1** and **2**.



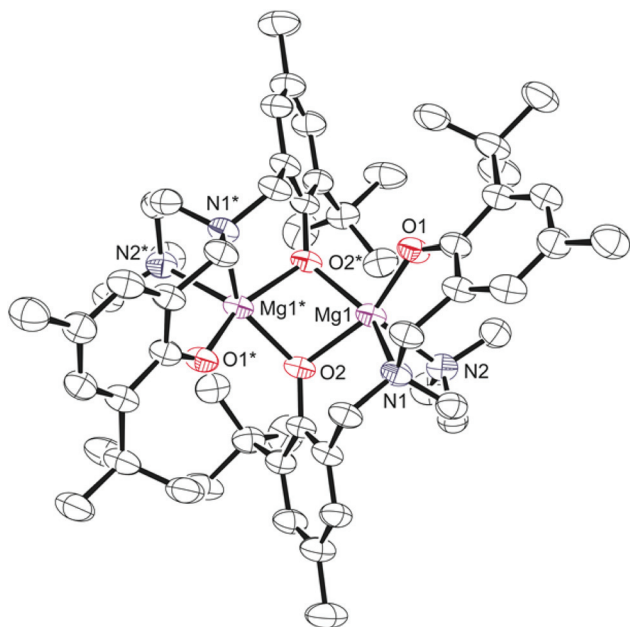


Fig. 1 Partially labelled molecular structure (ORTEP) of the dimer of complex **2**. Thermal ellipsoids are drawn at 50% probability and H atoms are excluded for clarity. Selected bond lengths (Å): Mg(1)–O(1), 1.950(2); Mg(1)–O(2), 2.049(2); Mg(1)–O(2)*, 2.049(2); Mg(1)–N(1), 2.243(3); Mg(1)–N(2), 2.265(3). Bond angles (°): O(1)–Mg(1)–O(2), 153.48(10); O(1)–Mg(1)–N(1), 88.25(9); O(2)–Mg(1)–O(2)*, 74.53(9); Mg(1)–O(2)–Mg(1)*, 95.71(8); O(2)*–Mg(1)–N(1), 140.38(9); O(2)–Mg(1)–N(2), 102.07(9); O(1)–Mg(1)–O(2)*, 91.64(9). *Symmetry operations used to generate equivalent atoms: $1 - x, y, \frac{1}{2} - z$.

presence of a polar monomer or a coordinating solvent, for example) may induce activity in these typically inactive bimetallic complexes. NMR studies of monomer formation in the presence of benzyl alcohol (a lactide ROP cocatalyst) are discussed below. The Mg(1)–O(1) bond (1.950(2) Å) in **2** is considerably shorter than the Mg(1)–O(2) bond (2.049(2) Å), because of the bridging bonding mode of the O(2) atom. The bond lengths for Mg(1)–N(1) and Mg(1)–N(2) were very similar at 2.243(3) Å and 2.265(3) Å, respectively, and are within the range observed in similar complexes.^{50–52}

A brief comparison of the structural properties of magnesium amine-bis(phenolate) complexes follows. The trigonality index,⁵⁸ τ , for **2** is 0.27 at both magnesium centres, which more closely approaches a distorted square pyramidal ($\tau = 0$) rather than a trigonal bipyramidal geometry ($\tau = 1$). The related magnesium complex reported by Jerzykiewicz and co-workers differs from **2** in that it contains one methyl substituent at the *para* position of the phenolate ring and no substituents at the *ortho* position resulting in a much less sterically encumbered metal site.⁵⁰ This is manifested in a rhombic Mg₂O₂ core that is nearly perfectly planar, possessing a sum of angles of 359.9°. The τ values for this compound are 0.35 and 0.61 for Mg(1) and Mg(2), respectively, indicating geometries around the metal centres that are intermediate to square pyramidal and trigonal bipyramidal.

A similar complex was reported by Sobota and co-workers possessing the tripodal amine-bis(phenolate) ligand, *N,N*-bis(3,5-di-*tert*-butylbenzyl-2-hydroxy)tetrahydrofurfurylamine.⁵² A dimeric complex was observed in C₆D₆ solutions and in the solid state, where it displayed a distorted rhombic Mg₂O₂ core whose sum of angles was 346.8°. The structure exhibited τ values for the two pentacoordinate magnesium centres that were almost identical at 0.41 and 0.39, intermediate to trigonal bipyramidal and square pyramidal arrangements, similar to complex **2**. This complex, however, also proved to be inactive in ROP of *L*-lactide at 25 °C in toluene.

A related phenolate-bridged magnesium dimer possessing *N,N'*-bis(2-hydroxido-3,5-di-*tert*-butyl)-*N,N'*-dimethylethane-1,2-diamine (salan) ligands was reported by Davidson, O'Hara and co-workers.⁵¹ As in the compound reported by Jerzykiewicz, each magnesium ion in this structure possesses different coordination environments with Mg(1) and Mg(2) centres having τ values of 0.32 and 0.25, respectively. A Mg₂O₂ core is also present in this structure with a sum of angles of 357.6°. ¹³C- and ¹H-NMR spectra for this compound in aromatic solvents suggested a dimeric structure in solution, similar to complex **2**. Two distinct phenolate environments and complicated methylene resonances indicate the dimer remains intact in non-coordinating aromatic solvents. This complex showed no activity in ring opening polymerization of lactide even at 110 °C in solution, which was attributed to the bulkiness of the ligand and the persistence of the dimeric nature of the complex in non-coordinating, aromatic solvents.

Magnesium *N,N*-bis(3,5-di-*tert*-butylbenzyl-2-hydroxy)dimethylethylenediamine was prepared by Bochmann and co-workers using magnesium bis[bis(trimethylsilyl)amide] (instead of the more commonly used di(*n*-butyl)magnesium) to give a complex that, although not structurally authenticated, is expected to be similar to those discussed above.⁵³ This complex showed poor activity (only 5% conversion) in the ROP of ϵ -caprolactone (CL) in toluene at 60 °C for a [CL]:[Mg] ratio of 200 : 1.

Although the solution-state ROP experiments of the previously reported magnesium-bis(phenolate) complexes were not encouraging, we believed that under melt conditions compounds **1** and **2** may exhibit activity. Indeed, *rac*-lactide can be polymerized by **1** and **2** in the bulk at temperatures above 125 °C and, to our surprise, in toluene solutions at 90 °C.

Ring opening polymerization in the melt

Melt phase ROP has attracted much interest recently as it offers a viable technique for industry.⁵⁹ There are examples for lactide ROP in the melt catalysed by a variety of metal complexes, such as sulphonamide-supported aluminum complexes,⁶⁰ Ni(II) and Ni(II)–Sm(III) salen complexes,⁶¹ aluminum-salen⁶² and salen-like⁶³ complexes, zirconium and hafnium amine tris(phenolate) alkoxides,⁶⁴ but solvent-free lactide ROP with an earth-abundant, non-toxic metal initiator at low catalyst loading still remains a challenge. Magnesium complexes **1** and **2** were assessed for their capabilities toward the ROP of *rac*-lactide under industrially relevant melt (solvent-free) conditions above 125 °C. The polymerization results are summar-



Table 1 Solvent-free (melt) *rac*-lactide polymerization by complexes **1** and **2**

Entry ^a	Complex	Time (min)	<i>T</i> (°C)	[Mg]:[<i>rac</i> -LA]:[BnOH]	Conv. ^b (%)	Activity ^c (kg mol ⁻¹ h ⁻¹)	<i>M</i> _n ^d (calcd)	<i>M</i> _n ^e (GPC)	<i>D</i> (<i>M</i> _w / <i>M</i> _n)
1	1	100	125	1:100:0	99	8.56	14 400	20 300	2.74
2	1	100	125	1:100:1	99	8.56	14 400	11 300	1.77
3	1	100	125	1:500:0	99	42.8	72 000	36 200	3.01
4	1	100	125	1:500:1	98	42.4	70 600	9700	2.01
5	2	100	125	1:100:0	100	8.65	14 400	16 500	2.40
6	2	100	125	1:100:1	82	6.23	14 400	7200	1.54
7	2	100	125	1:500:0	72	31.1	51 800	30 600	2.72
8	2	100	125	1:500:1	98	42.4	70 600	2400	1.25
9	1	100	125	1:1000:0	93	80.4	133 900	44 900	1.25
10	1	100	125	1:1000:1	93	80.4	133 900	42 900	1.24
11	1	180	125	1:1000:0	93	44.7	133 900	74 300	1.05
12	1	180	125	1:1000:1	89	42.8	128 200	50 100	1.24
13	1	100	150	1:1000:0	92	79.6	132 500	35 200	1.22
14	1	100	150	1:1000:1	96	83.0	138 200	70 290	1.08
15	1	180	150	1:2500:0	16	19.2	57 600	ND ^f	ND

^a All polymerization reactions were carried out in neat *rac*-lactide (0.5–0.6 g). ^b Conversion determined by ¹H-NMR. ^c Calculated as mass of lactide converted in grams/(mol of Mg × time in h). ^d Calculated from $([LA]/[Mg]) \times \% \text{ conv.} \times 144.13 \text{ g mol}^{-1}$. ^e Molecular weights (g mol⁻¹) of entries 1 to 8 determined by gel permeation chromatography (GPC) in CHCl₃ calibrated against polystyrene standards using the Mark-Houwink correction of 0.58.⁶⁵ GPC of entries 9 to 15 were conducted in THF by triple detection. See Experimental for full details. ^f ND = Not Determined, yield too low.

ized in Table 1. Polymerization occurred with or without a co-initiator (benzyl alcohol, BnOH). In general, the polymer molecular weights observed by gel permeation chromatography were lower than expected, likely due to mass transfer issues in the melt. Also, lower molecular weight polymers were obtained when BnOH was used, but the dispersity values were lower indicating a more controlled polymerization. Higher molecular weight and monodisperse polymers were obtained when catalyst loading is decreased. Interestingly, the physical appearance of the polymers was found to depend on whether BnOH was used (which gave thin films) or not (which gave crystalline white solids). The mechanical properties of the polymers are discussed below. Complex **1** showed slightly better conversions than complex **2**, so **1** was chosen to carry out further reactions.

When the catalyst loading was lowered to 0.1 mol% (entries 9–14), lactide conversion remained high translating to an increase in activity with good control as shown by the narrow dispersities of the polymers obtained. Polymerization remained controlled even in the absence of BnOH co-initiator. Highest molecular weights were obtained when the polymerization was run for 180 min (entry 11) or when the temperature was raised to 150 °C (entry 14). Activity was still observed upon further decreasing the catalyst loading to 0.04 mol% at this temperature, but with low conversion (16%) (entry 15).

Kinetic studies performed with **1** under conditions given in Table 1, entry 3 revealed that the polymerization reached completion in 60 minutes (see Table 2). Molecular weights increase with increasing time and conversion and the dispersity values are generally low (1.14–1.40). The difference in conversion between entry 5 and 6 is not significant, but the difference in molecular weight is in accordance with the phenomenon observed earlier in Table 1, entry 11 that it is favourable to leave the polymerization to occur for longer time to achieve higher molecular weight polymers.

Table 2 Time dependence of melt-phase *rac*-lactide polymerization by **1**

Entry ^a	Time (min)	Conv. ^b (%)	<i>M</i> _n ^c (calcd)	<i>M</i> _n ^d (GPC)	<i>D</i> (<i>M</i> _w / <i>M</i> _n)
1	10	7	5000	5400	1.40
2	20	36	25 900	16 600	1.22
3	30	78	56 000	18 100	1.34
4	40	91	65 500	25 100	1.37
5	50	93	67 000	31 400	1.32
6	60	96	69 100	55 100	1.14

^a All polymerization reactions were carried out in neat *rac*-lactide (0.5 g), 125 °C with $[1]:[LA] = 1:500$. ^b Determined by ¹H-NMR. ^c Calculated from $([LA]/[Mg]) \times \% \text{ conv.} \times 144.13 \text{ g mol}^{-1}$. ^d Molecular weights (g mol⁻¹) determined by triple detection gel permeation chromatography (GPC) in tetrahydrofuran using a *dn/dc* value of 0.049 mL g⁻¹.

The conversion vs. time plot (Fig. S3 in ESI†) shows that polymerization starts once a homogenous melt is achieved (after 5 minutes at 125 °C), the rate of conversion remains almost linear for the next 10 min and then accelerates for 5 min before returning to a constant rate. The discontinuity in conversion rate implies an induction period, which could be a result of dimer dissociation in order to form the active species. Solution polymerization kinetics studies (see below) are consistent with this theory. The MALDI-TOF mass spectra of the polymers obtained in the melt in the absence of BnOH showed the presence of both cyclic poly(*rac*-lactide) and linear chains terminated with –OH and carboxylic acid end groups (Fig. S4 and S5 in ESI†).

Mechanical properties of the polymers obtained under melt conditions

Polymerizations were performed on a larger scale using 1.0 g (instead of 0.5 g) of *rac*-lactide according to the conditions in



Table 3 Polymerization of *rac*-lactide by **2** in toluene

Entry ^a	[Mg] : [LA] : [ROH]	Time (min)	Conv. ^b (%)	M_n ^c (calcd)	M_n ^d (GPC)	$D (M_w/M_n)$
1	1 : 100 : 1 (iPrOH)	30	99	14 300	5700	1.70
2	1 : 100 : 10 (iPrOH)	3	99	1400	480	1.42
3	1 : 100 : 0	210	100	14 400	46 000	1.26
4	1 : 100 : 1 (BnOH)	12	98	14 100	13 900	1.07
5	1 : 100 : 2 (BnOH)	7	97	7200	7100	1.02

^a All polymerization reactions were carried out in toluene (10 mL) at 90 °C. ^b Determined by ¹H-NMR. ^c Calculated from $([LA]/[Mg]/[ROH]) \times \% \text{ conv.} \times 144.13 \text{ g mol}^{-1}$. ^d Molecular weights (g mol^{-1}) determined by gel permeation chromatography (GPC) in CHCl_3 calibrated against polystyrene standards using the Mark-Houwink correction of 0.58.⁶⁵

entries 3 and 4 of Table 1 to investigate the correlation between conversion and isolated yield. The scaled-up reactions improved the recovery of the polylactide after washing as determined by ¹H NMR. The physical appearance of the polymers obtained with and without added benzyl alcohol is significantly different as a result of the difference in polymer molecular weights obtained (Fig. S6 in ESI†). Without BnOH, the polymer is obtained as a thin film whereas in the presence of alcohol a crystalline polymer results. Differential Scanning Calorimetry (DSC) and Thermogravimetric Analysis (TGA) were performed on these samples and revealed very similar thermal properties. The midpoint of the glass transition temperature taken from the third heating curve was 52.59 °C for the polymer obtained from entry 3 and 50.86 °C for entry 4, which is expected given the polymer's much lower observed M_n . TGA conducted with a ramp rate of 10 °C min^{-1} showed the decomposition temperature of the polymer obtained in the absence of BnOH was 354.5 °C and slightly higher (356.7 °C) for the polymer obtained in presence of BnOH. Slower rates of 5 °C min^{-1} resulted in slightly lower decomposition temperatures for both the polymers obtained with BnOH (onset at 340 °C, end at 375 °C) and without BnOH (onset at 344 °C, end at 367 °C, see Fig. S7 and S8 in ESI†). These decomposition temperatures are higher than expected,⁶⁶ but they are similar to carboxylic acid terminated *branched* polymers reported by S. H. Lee *et al.*⁶⁷ It should be noted that the polymers used for thermal analysis were highly purified by precipitation, centrifugation, decanting of solvents, washing with cold methanol and drying by vacuum oven overnight at 40 °C.

Regarding polymer tacticity, the solvent-free (melt) ROP of *rac*-lactide using **1** gives an atactic microstructure of poly(*rac*-lactide). This is not surprising given the achiral ligands used in compounds **1** and **2** and the poor polymer M_n control. A representative ¹H{¹H}-NMR spectrum of the methine resonances of poly(*rac*-lactide) is given in Fig. S9 in ESI†. Highly isotactic polylactide stereopolymers can be obtained solvent-free at 130 °C using salen aluminum alkoxides at a monomer/initiator molar ratio of 200, but the reaction rates were very slow (95% conversion after 2 days).⁶⁸

Ring opening polymerization in solution

Polymerization at room temperature did not occur in either CH_2Cl_2 or THF with complexes **1** and **2**. However, upon

heating to 90 °C with 1 equivalent of isopropyl alcohol (iPrOH) as co-initiator complex **2** exhibited fast propagation in toluene achieving 98% conversion in 12 minutes (Table 3, entry 1 and Fig. 2). Use of iPrOH accelerates the polymerization activity compared to using **2** alone, but gives lower than expected polymer molecular weights likely resulting from transesterification. Increasing the amount of iPrOH as co-initiator to 10 equivalents with respect to **2** resulted in faster reaction rate but much lower molecular weight and narrower dispersity (entry 2). The plot of conversion vs. time (Fig. 2) indicates an induction period of approximately 7 minutes before the onset of the expected conversion curve. This non-linearity is apparent when attempting to model the data using a first order kinetics plot over the whole reaction time (Fig. S10 in ESI†). These observations can be explained if one considers the dimeric form of **2** in toluene as being inactive to ROP of *rac*-lactide. Hence, activation of the complex is achieved in the presence of co-initiator, possibly *via* formation of monomeric alcohol adducts. Once an active species is obtained, ROP proceeds quickly at 90 °C, thus explaining the rapid increase in conversion after 7 minutes and effectively complete consumption of monomer after 12 minutes. This is consistent with the activity vs. *t* plot (Fig. S11 in ESI†), which shows the highest activity was observed between 9 and 15 min. NMR spectra following the reaction profile are given in Fig. S12 in ESI†.

The influence of the co-initiator on polymerizations in toluene was investigated. The polymerization without a co-initiator (Table 3, entry 3) proceeded very slowly and any conversion of lactide was only observed after 30 min in toluene at

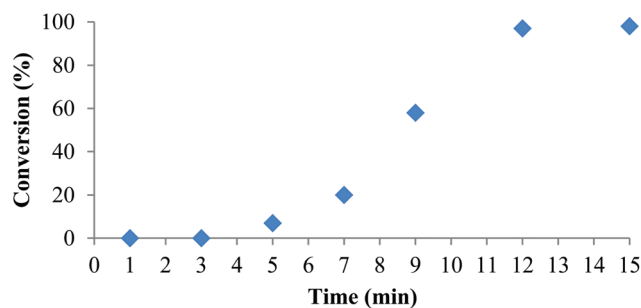


Fig. 2 Conversion (%) vs. time for the ROP of *rac*-LA initiated by **2** under the conditions in Table 3, entry 1.



90 °C, which is consistent with a slow initiation period. In the presence of BnOH, however, the reaction proceeded very quickly and achieved 98% conversion within 6 minutes (Table 3, entry 4). The polymerization demonstrates first order reaction profiles with respect to [LA] in either the presence or absence of BnOH (Fig. 3). The observed rate constant, k_{obs} , in the presence of BnOH was 0.828 min^{-1} whereas in the absence of BnOH was $9.87 \times 10^{-2} \text{ min}^{-1}$. When the concentration of BnOH was doubled (Table 3, entry 5), the molecular weight of the polymer decreased by half, which suggests a well-behaved *immortal* ROP with rapid, reversible chain transfer between growing PLA fragments and dormant hydroxyl-terminated polylactide chains.^{69,70}

Mechanistic proposal

Without addition of exogenous alcohol, the ROP of LA is slower in solution than when the alcohol co-initiator is

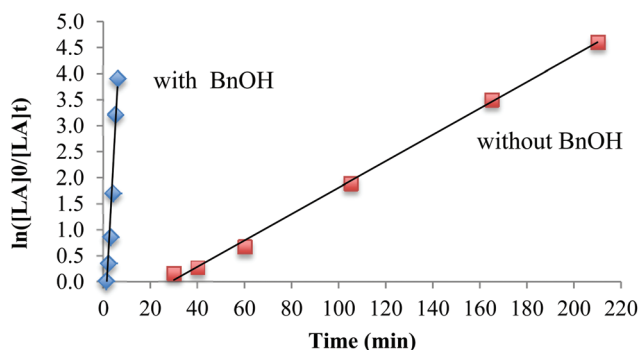
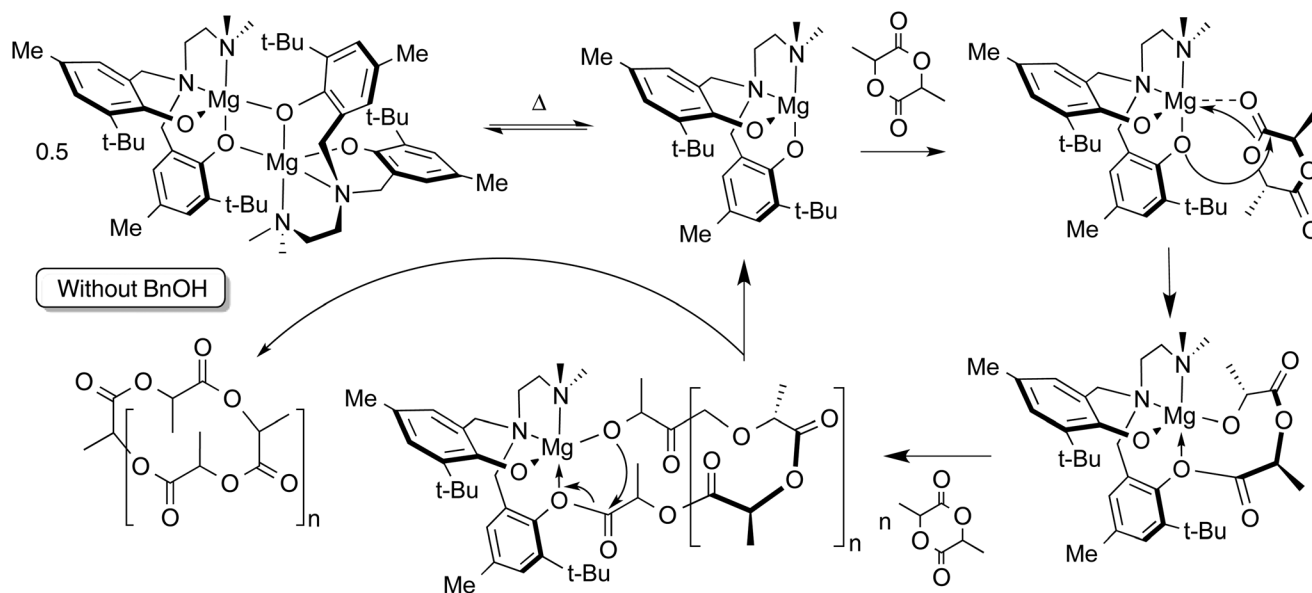


Fig. 3 Plot of $\ln([LA]_0/[LA]_t)$ vs. t , $[LA]_0/[Mg]_0 = 100$, in toluene at 90 °C according to the conditions in Table 3, entries 3 and 4.

present. Initiation by nucleophilic attack of the phenolate ligand on the monomer has been observed in related lithium-phenolate complexes.^{10,44,69} In melt polymerization, however, there is no significant loss of control of the polymerization when conducted in the absence of alcohol co-initiator. The observation in the MALDI-TOF mass spectrum of cyclic polylactide alongside acyclic polymer possessing carboxylic acid and hydroxyl end groups (resulting from hydrolytic quenching of polymerization with acidified methanol) is consistent with phenolate-initiated ROP (Scheme 2).

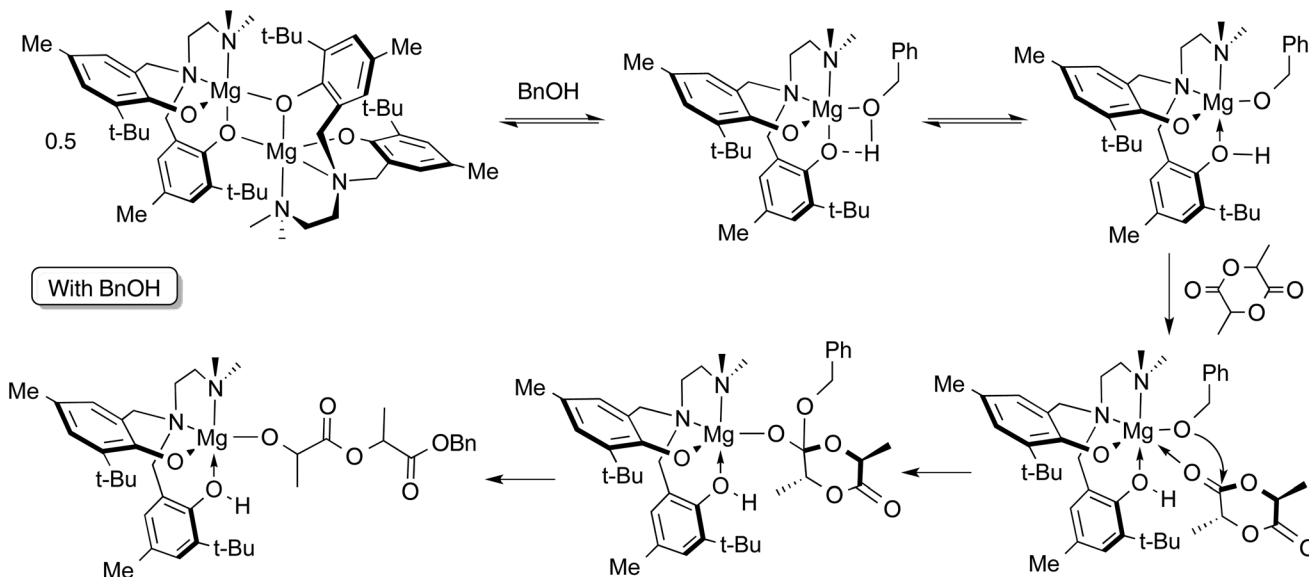
The mechanism of polymerization and the role of the alcohol co-initiator require closer examination, particularly since the majority of binary systems (employing a Lewis acidic metal site and an added alcohol as initiators) hypothesize coordination-insertion mechanisms. Carpentier, Sarazin and co-workers suggest that activated monomer mechanisms should not be excluded and have performed elegant studies of alkali aminoether-phenolate complexes that provide strong evidence that this mechanism is at play under living or immortal ROP conditions of L-LA.⁶⁹

Preliminary stoichiometric studies by ¹H NMR spectroscopy of reactions involving complex 2, *rac*-LA and BnOH were conducted in toluene-*d*₈ at 363 K, *i.e.* the conditions employed in solution ROP catalysis shown in Table 3. The reaction of a 1 : 1 ratio (per Mg centre) of 2 and BnOH was followed by ¹H NMR at 363 K. After 10 minutes at this temperature, in addition to those of complex 2 alone, new resonances were observed consistent with monomeric Mg compounds (see Fig. S14 ESI† for ¹H NMR spectra). It is not unequivocal whether the BnOH is coordinated to the metal centre since the benzyl methylene peak is slightly broadened at this temperature, possibly from an exchange process as suggested by the continued presence



Scheme 2 Proposed mechanism for ROP of LA by 2 without added alcohol co-initiator. Generation of cyclic polymer is shown, but hydroxyl or carboxylic acid end groups may be obtained by protonolysis.





Scheme 3 Proposed mechanism for ROP of LA by 2 with benzyl alcohol co-initiator.

of dimer in the spectrum. Also, a very small peak at 9.19 ppm could be observed upon expansion of the region of the spectrum, suggesting that the presence of protonated ligand, $H_2[L2]$ is negligible.

The reaction of a 1 : 1 : 1 mixture of *rac*-LA, 2 and BnOH was monitored under the same conditions. Within 10 minutes at 363 K the product of lactide ring-opening, benzyl-2-((2-hydroxypropanoyl)oxy)propanoate or its deprotonated alkoxide form, was observed. It is possible that the ring-opened lactide remains coordinated to the Mg site, since a phenolic -OH resonance is observable at 9.22 ppm indicating protonation of a phenolate oxygen. The proposed mechanism is shown in Scheme 3. Based on these observations, an activated monomer mechanism may be occurring, but further investigation is needed.

Conclusions

Magnesium amine-bis(phenolate) compounds 1 and 2 showed catalytic activity for the ROP of *rac*-lactide both in toluene solutions and solvent-free (melt) to give atactic poly(*rac*-lactide). Kinetic studies showed that conversion of *rac*-lactide is essentially complete within 60 minutes in the melt at 125 °C, although the resulting polymers exhibit broad dispersities. Solution polymerization studies in toluene at 90 °C indicate long initiation periods in the absence of an alcohol co-initiator, but rapid initiation when a suitable co-initiator is used. Complex 2 proved to be an efficient initiator in solution at 90 °C with a [LA] : [Mg] ratio of 100 : 1 regardless of whether a co-initiator was used or not. Without a co-initiator an initiation period of ~30 min was observed, after which time the propagation showed a slow first order reaction profile with respect to *rac*-lactide concentration and effectively complete

conversion within 210 minutes. The polymerization was accelerated significantly by the addition of a co-initiator, either *i*PrOH or BnOH. The highest activity was observed for [LA] : [Mg] : [BnOH] ratios of 100 : 1 : 1 at 90 °C in toluene giving 177.3 kg of *rac*-PLA per mol Mg per h ($k_{obs} = 1.38 \times 10^{-2} s^{-1}$). Preliminary mechanistic studies support an activated monomer pathway, but more detailed studies are ongoing.

Experimental

General experimental conditions

Unless otherwise stated, all manipulations were performed under an atmosphere of dry, oxygen-free nitrogen by means of standard Schlenk techniques or using an MBraun Labmaster DP glove box. Anhydrous THF was distilled from sodium benzophenone ketyl under nitrogen. Isopropyl alcohol was dried over calcium oxide, then distilled under nitrogen. Toluene was purified by an MBraun Manual Solvent Purification System. Reagents were purchased either from Aldrich or Alfa Aesar and used without further purification. *Rac*-lactide was purchased from Aldrich or Alfa Aesar and dried over sodium sulphate in THF, recrystallized and stored under an inert atmosphere prior to use. Benzyl alcohol was purchased from Alfa Aesar and dried over activated 4 Å molecular sieves, distilled under reduced pressure and stored under nitrogen in an ampule prior to use.

Instrumentation

MALDI-TOF MS was performed using an Applied Biosystems 4800 MALDI TOF/TOF Analyzer equipped with a reflectron, delayed ion extraction and high performance nitrogen laser (200 Hz operating at 355 nm). Samples were prepared in the glove box and sealed under nitrogen in a Ziploc® bag for trans-



port to the instrument. Anthracene was used as the matrix for compounds **1** and **2** and 2,5-dihydroxybenzoic acid (DHBA) was used as the matrix for the polymers. The matrix was dissolved in THF at a concentration of 10 mg mL⁻¹. Polymer was dissolved in THF at approximately 1 mg mL⁻¹. The matrix and polymer solutions were mixed together at ratios of 5 to 1, 4 to 1 or 3 to 1; 1 μL of this was spotted on the MALDI plate and left to dry. Images of mass spectra were prepared using mMass™ software (<http://www.mmass.org>).

GPC analysis was performed either in CHCl₃ on a Viscotek VE 2001 GPCMax at 35 °C equipped with a Viscotek VE 3580 Refractive Index Detector or in THF at 25 °C on a Wyatt Triple Detection (triple angle light scattering, viscometry and refractive index) system with Agilent 2600 series sample and solvent handling. The Viscotek system used two Phenogel 5μ Linear Mixed Bed 300 × 4.60 mm columns whereas the Wyatt system used two Phenogel 10³ Å 300 × 4.60 mm columns. Samples were prepared at a concentration of 2 mg mL⁻¹ in CHCl₃ and left to equilibrate for ~2 h. The samples were filtered through syringe filters before analysis and eluted with HPLC grade solvents at flow rates of 0.30 mL min⁻¹ with 100 μL injection volumes. For conventional calibration, six polystyrene standards (Viscotek) were used in making the calibration curve, bracketing molecular ranges from 1050 to 400 000 Da. A correction factor of 0.58 was used to calculate the M_n value.⁶⁵

NMR spectra were recorded at 300 MHz for ¹H and 75.5 MHz for ¹³C or at 500 MHz for ¹H NMR kinetic studies. CDCl₃, C₆D₆, DMSO-d₆ were purchased from Cambridge Isotope Laboratories and pyridine-d₅ from Aldrich. All deuterated solvents except for DMSO-d₆ used in the analysis of the magnesium complexes were dried over calcium hydride (CDCl₃ and pyridine-d₅) or sodium/potassium alloy (C₆D₆), vacuum transferred and stored under nitrogen in ampules fitted with Teflon valves. Elemental analyses were performed at Guelph Chemical Laboratories, Guelph, Ontario, Canada.

Thermogravimetric analysis (TGA) was performed with a TA Instruments Q500. Samples (4–15 mg) were loaded onto a platinum pan and subjected to a dynamic high-resolution scan, with an initial heating rate of 5 or 10 °C min⁻¹. Each sample was heated from room temperature to 600 °C. Glass transition (*T_g*) temperatures were measured using a Mettler Toledo DSC 1 STAR^e System equipped with a Julabo FT 100 immersion cooling system, using R1150 refrigerant in an EtOH bath with a working range of -100 to +20 °C. Samples (~5 mg) were weighed into 40 μL aluminum pans and subjected to three heating cycles. The first heating cycle consisted of heating from 0 to 100 °C at a rate of 10 °C min⁻¹, held for 2 min at 100 °C and then cooled back to 0 °C at 10 °C min⁻¹. The sample was held at this temperature for 2 min and subjected to a two heating cycles from 0 to 190 °C at a rate of 10 °C min⁻¹.

Synthesis of compounds

H₂[L1]. Aminomethylpyridine (6.65 g, 0.0615 mol) was slowly added to a vigorously stirred mixture of 3-*tert*-butyl-5-hydroxyanisole (22.17 g, 0.123 mol) and formaldehyde

(9.16 mL of a 37% aqueous solution, 0.123 mol) in deionized water (100 mL) in a round bottom flask equipped with a magnetic stir bar. This mixture was heated to reflux and stirred overnight. Upon cooling, a large quantity of white solid formed. The solvent was decanted and the solid was washed with cold methanol then recrystallized from a chloroform-methanol solvent mixture to give a white crystalline powder (25.06 g, 82% yield). Anal. Calcd For C₃₀H₄₀N₂O₄: C 73.14, H 8.18, N 5.69. Found: C 72.88, H 7.90, N 5.53. ¹H NMR (300 MHz, 298 K, CDCl₃, δ): 10.30 (br s, OH, 2H), 8.69 (ddd, ³J_{HH} = 5.03 Hz, ⁴J_{HH} = 1.79, ⁵J_{HH} = 0.93 Hz, Py: NCH, 1H), 7.70 (ddd, ³J_{HH} = 7.69 Hz, ³J_{HH} = 7.69 Hz, ⁴J_{HH} = 1.79 Hz, Py: NCHCH, 1H), 7.29 (ddd, ³J_{HH} = 7.69 Hz, ⁴J_{HH} = 5.01 Hz, ⁵J_{HH} = 1.05 Hz, Py: NCHCHCH, 1H), 7.12 (dt, ³J_{HH} = 7.80 Hz, ⁴J_{HH} = 1.05 Hz, Py: NCCH, 1H), 6.83 (d, ⁴J_{HH} = 3.08 Hz, Ar: CHCCH₂N, 2H), 6.51 (d, ⁴J_{HH} = 3.06 Hz, Ar: CHCCMe₃, 2H), 3.80 (s, NCH₂Py, 2H), 3.78 (s, NCH₂Ar, 4H), 3.75 (s, OMe, 6H), 1.40 (s, CMe₃, 18H). ¹³C{¹H} NMR (75.5 MHz, CDCl₃, 298 K, δ): 155.99 (COH); 151.96 (Ar: COMe) 150.38 (Ar: CCH₂N), 148.37 (Py: NCCH), 139.06 (Py: NC), 137.68 (Py: NCCHCH), 124.18 (Py: NCH), 122.85 (Ar: CCMe₃); 122.81 (Py: NCHCH), 113.72 (Ar: CHCCH₂N); 112.54 (Ar: CHCCMe₃); 56.44 (CH₂Py); 55.91 (OMe); 55.12(NCH₂Ar), 35.28 (CMe₃), 29.66 (CMe₃).

H₂[L2]. *N,N*-Dimethylethylenediamine (5.45 g, 61.8 mmol) was slowly added to a vigorously stirred mixture of 2-*tert*-butyl-4-methylphenol (20.23 g, 123 mmol) and formaldehyde (10.0 mL of a 37% aqueous solution, 0.123 mol) in deionized water (100 mL) in a round bottom flask equipped with a magnetic stir bar. The mixture was heated to reflux and stirred for approximately 24 hours, at which point a large amount of orange-brown solid had formed. The mixture was decanted and the remaining solid residue was washed with ice-cold methanol to yield a white powder. Recrystallization from chloroform produced analytically pure product (yield: 23.8 g, 88%). Spectroscopic characterization was consistent with that previously reported.⁵³

1: 2.46 g (5.00 mmol) of H₂[L1] was dissolved in 100 mL of toluene and cooled to -78 °C. To this solution was added 5.0 mmol of MgBu₂ (5.0 mL of a 1.0 M solution in heptane) *via* syringe. The solution was warmed to room temperature and stirred overnight. Volatiles were removed *in vacuo* to yield a pale yellow powder. Yield = 2.42 g (94%). Anal. calc'd for C₃₀H₃₈MgN₂O₄: C 69.97, H 7.44, N 5.44. Found: C 69.26, H 7.17, N 5.13. The lower than expected %C is attributed to a 5–10% contamination by MgO or Mg(OH)₂ (*e.g.* anal. calc'd for C₃₀H₃₈MgN₂O₄(MgO₂H₂)_{0.05}: C 69.58, H 7.42, N 5.41. ¹H NMR (300 MHz, 298 K, DMSO-d₆, δ): 8.49 (d, ³J_{HH} = 4.43 Hz, Py: NCH, 1H); 7.68 (td, ³J_{HH} = 7.62 Hz, ⁴J_{HH} = 1.51 Hz, Py: NCHCH, 1H); 7.21 (dd, coupling constants could not be determined because of the overlapping residual toluene peaks, Py: NCHCHCH, 1H); 7.06 (d, ³J_{HH} = 7.8 Hz, Py: NCCH, 1H); 6.44 (d, ⁴J_{HH} = 3.23 Hz, Ar: CHCCH₂N, 2H); 6.38 (d, ⁴J_{HH} = 3.23 Hz, Ar: CHCCMe₃, 2H); 3.76 (d, ³J_{HH} = 11.65 Hz, NCH₂Ar, 2H); 3.58 (s, NCH₂Py, 2H); 3.55 (s, OCH₃, 6H); 3.15 (d, ³J_{HH} = 11.72 Hz, NCH₂Ar, 2H); 1.29 (s, C(CH₃)₃, 18H). ¹³C{¹H} NMR (75.5 MHz, 298 K, DMSO-d₆, δ): 160.71 (COMg); 158.04 (COMe); 148.60



(Ar: CCH₂N); 145.35 (Py: NCCH); 138.46 (Py: NC); 136.66 (Py: NCCHCH); 130.38 (Py: NCHCH); 122.69 (Py: NCH); 122.03 (Ar: CCM₃); 113.46 (Ar: CHCCH₂N); 112.94 (Ar: CHCCM₃); 60.68 (CH₂Py); 56.10 (CH₂Ar); 55.69 (OMe); 34.54 (CMe₃); 29.34 (CMe₃). MS (MALDI-TOF) *m/z* (% ion): 514.26 (58, [M]⁺); 1028.50 (100, [M]₂⁺).

2: 2.21 g (5.0 mmol) of H₂[L₂] was dissolved in 70 mL of toluene and cooled to -78 °C. To this solution was added 5.0 mmol of MgBu₂ (5.0 mL of a 1.0 M solution in heptane) *via* syringe, forming a bright yellow solution. The solution was warmed to room temperature and stirred overnight, upon which a large amount of white precipitate formed within a colourless solution. Volatiles were removed *in vacuo*, yielding an analytically pure white powder. Yield = 2.29 g (99%). Recrystallization from toluene at -35 °C afforded colourless crystals. Anal. calc'd for C₃₄H₅₄MgN₂O₂: C 72.64, H 9.14, N 6.05. Found: C 72.38, H 9.09, N 6.22. ¹H NMR (300 MHz, 298 K, C₆D₆, δ): 7.29 (m, Ar, 4H); 6.80 (d, ⁴J_{HH} = 2.35 Hz, Ar, 2H); 6.69 (d, ⁴J_{HH} = 2.30, Ar, 2H); 4.84 (d, ²J_{HH} = 12.01 Hz, Ar-CH(H)-N, 2H); 4.63 (d, ²J_{HH} = 12.41 Hz, Ar-CH(H)-N, 2H); 2.97 (d, ²J_{HH} = 12.38 Hz, Ar-CH(H)-N, 2H); 2.63 (d, ²J_{HH} = 12.06 Hz, Ar-CH(H)-N, 2H); 2.00–2.70 (br, CH₂NMe₂, 4H); 2.35 (s, NMe₂, 6H); 2.31 (s, NMe₂, 6H); 1.05–1.70 (br, CH₂CH₂NMe₂, 4H); 1.69 (s, CMe₃, 18H); 1.65 (s, ArMe, 6H); 1.59 (s, ArMe, 6H); 1.29 (s, CMe₃, 18H). ¹³C{¹H} NMR (75.5 MHz, 298 K, C₆D₆, δ): 164.59 (C-O); 160.09 (C-O); 137.80, 136.95, 130.13, 129.61, 128.92, 128.06, 128.74, 125.55, 123.01, 119.89 (All Ar); 65.49 (ArCH₂); 64.48 (ArCH₂); 60.54 (CH₂CH₂NMe₂); 48.69 (CH₂NMe₂); 47.55 (NMe₂); 47.42 (NMe₂); 35.12 (CMe₃); 34.77 (CMe₃); 32.82 (CMe₃); 30.34 (CMe₃); 20.98 (ArMe); 20.90 (ArMe). ¹H NMR (300 MHz, 298 K, pyridine-d₅, δ): 7.35 (d, ⁴J_{HH} = 2.45 Hz, Ar, 2H); 6.92 (d, ⁴J_{HH} = 2.51 Hz, Ar, 2H); 4.04 (d, ²J_{HH} = 12.49, Ar-CH(H)-N, 2H); 3.11 (d, ²J_{HH} = 12.51 Hz, Ar-CH(H)-N, 2H); 2.75 (br, CH₂NMe₂, 2H); 2.40 (s, NMe₂, 6H); 2.23 (br, CH₂CH₂NMe₂, 2H); 1.97 (s, ArMe, 6H); 1.75 (s, CMe₃, 18H). ¹³C{¹H} NMR (75.5 MHz, 298 K, pyridine-d₅, δ): 167.80 (C-O); 138.71, 132.41, 130.65, 130.08, 126.64 (All Ar); 66.88 (ArCH₂); 61.54 (CH₂CH₂NMe₂); 51.27 (CH₂NMe₂); 49.51 (NMe₂); 37.24 (CMe₃); 32.43 (CMe₃); 23.05 (CMe₃). MS (MALDI-TOF) *m/z* (% ion): 462.25 (100, [M]⁺); 924.52 (3, [M]₂⁺).

Polymerization procedure

Melt polymerization. A representative polymerization with a 500:1 [*rac*-lactide]:[**1**] ratio was carried out with 500 mg (3.47 mmol) *rac*-lactide and 3.57 mg (6.93 × 10⁻³ mmol) of complex **1** weighed into a 10 mL scintillation vial equipped with a small stir bar in the glove box. The closed vial was taken out of the glove box and placed into an aluminum block vial holder, which was pre-heated on a hotplate to 125 °C. A typical polymerization was run for 100 minutes. The vial was then placed in an ice bath to halt the reaction and solidify the polymer. The solids were dissolved in dichloromethane or chloroform and the polymer was precipitated with acidified methanol. Centrifugation was applied where needed for better separation of the solids. Solvents were decanted and the white

solids (either crystalline or thin film-like) were dried *in vacuo* followed by drying in a vacuum oven at 40 °C overnight.

Solution polymerization. The appropriate amount of complex **1** or **2** and alcohol (iPrOH or BnOH) were placed in an ampule equipped with a PTFE valve. A second ampule was prepared containing a stir bar and the required amount of monomer and toluene (7.50 mL). Both ampules were heated to 90 °C for ~30 min, after which time the lactide solution was transferred by cannula to the complex mixture. Timing of the reaction began when all the lactide was transferred. An aliquot of the reaction mixture was taken for ¹H NMR analysis at certain time intervals. When the presence of monomer was no longer apparent by ¹H NMR, volatiles were removed *in vacuo* and the solid residue was dissolved in CH₂Cl₂ and precipitated using cold methanol. Solvents were decanted and the white solids were dried *in vacuo* followed by drying in a vacuum oven at 40 °C overnight.

Acknowledgements

Financial support was provided by the Canada Foundation for Innovation (CFI), Newfoundland and Labrador Research Development Corporation, the Natural Sciences and Engineering Research Council (NSERC) of Canada and Memorial University of Newfoundland. J. H. L. and M. E. P. thank NSERC for Undergraduate Student Research Awards.

References

- 1 R. T. Martin, L. P. Camargo and S. A. Miller, *Green Chem.*, 2014, **16**, 1768–1773.
- 2 J. M. Becker and A. P. Dove, in *Green Polymerization Methods: Renewable Starting Materials, Catalysis and Waste Reduction*, eds. R. T. Mathers and M. A. R. Meier, Wiley-VCH, Weinheim, 1st edn, 2011, pp. 201–220.
- 3 R. E. Drumright, P. R. Gruber and D. E. Henton, *Adv. Mater.*, 2000, **12**, 1841–1846.
- 4 O. Dechy-Cabaret, B. Martin-Vaca and D. Bourissou, *Chem. Rev.*, 2004, **104**, 6147–6176.
- 5 M. J. Stanford and A. P. Dove, *Chem. Soc. Rev.*, 2010, **39**, 486–494.
- 6 C. A. Wheaton, P. G. Hayes and B. J. Ireland, *Dalton Trans.*, 2009, 4832–4846.
- 7 M. H. Chisholm and Z. P. Zhou, *J. Mater. Chem.*, 2004, **14**, 3081–3092.
- 8 B. J. O'Keefe, M. A. Hillmyer and W. B. Tolman, *J. Chem. Soc., Dalton Trans.*, 2001, 2215–2224.
- 9 O. Wichmann, R. Sillanpää and A. Lehtonen, *Coord. Chem. Rev.*, 2012, **256**, 371–392.
- 10 N. Ikpo, C. Hoffmann, L. N. Dawe and F. M. Kerton, *Dalton Trans.*, 2012, **41**, 6651–6660.
- 11 Y. Huang, Y.-H. Tsai, W.-C. Hung, C.-S. Lin, W. Wang, J.-H. Huang, S. Dutta and C.-C. Lin, *Inorg. Chem.*, 2010, **49**, 9416–9425.



- 12 L. Wang, X. B. Pan, L. H. Yao, N. Tang and J. C. Wu, *Eur. J. Inorg. Chem.*, 2011, 632–636.
- 13 C.-A. Huang, C.-L. Ho and C.-T. Chen, *Dalton Trans.*, 2008, 3502–3510.
- 14 C. A. Huang and C. T. Chen, *Dalton Trans.*, 2007, 5561–5566.
- 15 F. M. Kerton, C. M. Kozak, K. Lüttgen, C. E. Willans, R. J. Webster and A. C. Whitwood, *Inorg. Chim. Acta*, 2006, **359**, 2819–2825.
- 16 B.-T. Ko and C.-C. Lin, *J. Am. Chem. Soc.*, 2001, **123**, 7973–7977.
- 17 W. Clegg, M. G. Davidson, D. V. Graham, G. Griffen, M. D. Jones, A. R. Kennedy, C. T. O'Hara, L. Russo and C. M. Thomson, *Dalton Trans.*, 2008, 1295–1301.
- 18 Z. Janas, T. Nerkowski, E. Kober, L. B. Jerzykiewicz and T. Lis, *Dalton Trans.*, 2012, **41**, 442–447.
- 19 L. E. Breyfogle, C. K. Williams, V. G. Young, M. A. Hillmyer and W. B. Tolman, *Dalton Trans.*, 2006, 928–936.
- 20 J. Ejfler, S. Szafert, K. Mierzwicki, L. B. Jerzykiewicz and P. Sobota, *Dalton Trans.*, 2008, 6556–6562.
- 21 W. C. Hung and C. C. Lin, *Inorg. Chem.*, 2009, **48**, 728–734.
- 22 E. L. Marshall, V. C. Gibson and H. S. Rzepa, *J. Am. Chem. Soc.*, 2005, **127**, 6048–6051.
- 23 Y. Sarazin, V. Poirier, T. Roisnel and J.-F. Carpentier, *Eur. J. Inorg. Chem.*, 2010, 3423–3428.
- 24 A. D. Schofield, M. L. Barros, M. G. Cushion, A. D. Schwarz and P. Mountford, *Dalton Trans.*, 2009, 85–96.
- 25 X. Zhang, T. J. Emge and K. C. Hultsch, *Organometallics*, 2010, **29**, 5871–5877.
- 26 D. J. Darensbourg, W. Choi, O. Karroonnirun and N. Bhuvanesh, *Macromolecules*, 2008, **41**, 3493–3502.
- 27 H. E. Dyer, S. Huijser, A. D. Schwarz, C. Wang, R. Duchateau and P. Mountford, *Dalton Trans.*, 2008, 32–35.
- 28 A. Amgoune, C. M. Thomas and J.-F. Carpentier, *Macromol. Rapid Commun.*, 2007, **28**, 693–697.
- 29 M. H. Chisholm, J. C. Gallucci, H. H. Zhen and J. C. Huffman, *Inorg. Chem.*, 2001, **40**, 5051–5054.
- 30 C. K. Williams, N. R. Brooks, M. A. Hillmyer and W. B. Tolman, *Chem. Commun.*, 2002, 2132–2133.
- 31 C. K. Williams, L. E. Breyfogle, S. K. Choi, W. Nam, V. G. Young, M. A. Hillmyer and W. B. Tolman, *J. Am. Chem. Soc.*, 2003, **125**, 11350–11359.
- 32 D. J. Doyle, V. C. Gibson and A. J. P. White, *Dalton Trans.*, 2007, 358–363.
- 33 J. D. Farwell, P. B. Hitchcock, M. F. Lappert, G. A. Luinstra, A. V. Protchenko and X.-H. Wei, *J. Organomet. Chem.*, 2008, **693**, 1861–1869.
- 34 G. Labourdette, D. J. Lee, B. O. Patrick, M. B. Ezhova and P. Mehrkhodavandi, *Organometallics*, 2009, **28**, 1309–1319.
- 35 L. Wang and H. Ma, *Dalton Trans.*, 2010, **39**, 7897–7910.
- 36 N. Ikpo, L. N. Saunders, J. L. Walsh, J. M. B. Smith, L. N. Dawe and F. M. Kerton, *Eur. J. Inorg. Chem.*, 2011, 5347–5359.
- 37 V. Poirier, T. Roisnel, J.-F. Carpentier and Y. Sarazin, *Dalton Trans.*, 2011, **40**, 523–534.
- 38 J. Lewinski, P. Horeglad, M. Dranka and I. Justyniak, *Inorg. Chem.*, 2004, **43**, 5789–5791.
- 39 J.-T. Issenhuth, J. Pluvinage, R. Welter, S. Bellemin-Laponnaz and S. Dagorne, *Eur. J. Inorg. Chem.*, 2009, 4701–4709.
- 40 S. Gendler, S. Segal, I. Goldberg, Z. Goldschmidt and M. Kol, *Inorg. Chem.*, 2006, **45**, 4783–4790.
- 41 S. Groysman, E. Sergeeva, I. Goldberg and M. Kol, *Inorg. Chem.*, 2005, **44**, 8188–8190.
- 42 L. M. Broomfield, Y. Sarazin, J. A. Wright, D. L. Hughes, W. Clegg, R. W. Harrington and M. Bochmann, *J. Organomet. Chem.*, 2007, **692**, 4603–4611.
- 43 L. N. Saunders, L. N. Dawe and C. M. Kozak, *J. Organomet. Chem.*, 2014, **749**, 34–40.
- 44 R. K. Dean, A. M. Reckling, H. Chen, L. N. Dawe, C. M. Schneider and C. M. Kozak, *Dalton Trans.*, 2013, 3504–3530.
- 45 R. H. Platel, L. M. Hodgson and C. K. Williams, *Polym. Rev.*, 2008, **48**, 11–63.
- 46 J. H. Khan, F. Schue and G. A. George, *Polym. Int.*, 2009, **58**, 296–301.
- 47 M. R. Kember and C. K. Williams, *J. Am. Chem. Soc.*, 2012, **134**, 15676–15679.
- 48 L. Wang and H. Ma, *Macromolecules*, 2010, **43**, 6535–6537.
- 49 S. Song, H. Ma and Y. Yang, *Dalton Trans.*, 2013, **42**, 14200–14211.
- 50 E. Kober, Z. Janas, T. Nerkowski and L. B. Jerzykiewicz, *Dalton Trans.*, 2013, **42**, 10847–10854.
- 51 M. G. Davidson, M. D. Jones, D. Meng and C. T. O'Hara, *Main Group Chem.*, 2006, **5**, 3–12.
- 52 J. Ejfler, K. Krauzy-Dziedzic, S. Szafert, L. B. Jerzykiewicz and P. Sobota, *Eur. J. Inorg. Chem.*, 2010, 3602–3609.
- 53 Y. Sarazin, R. H. Howard, D. L. Hughes, S. M. Humphrey and M. Bochmann, *Dalton Trans.*, 2006, 340–350.
- 54 F. M. Kerton, S. Holloway, A. Power, R. G. Soper, K. Sheridan, J. M. Lynam, A. C. Whitwood and C. E. Willans, *Can. J. Chem.*, 2008, **86**, 435–443.
- 55 S. Menage, G. Gellon, J.-L. Pierre, D. Zurita and E. Saint-Aman, *Bull. Soc. Chim. Fr.*, 1997, **134**, 785–791.
- 56 S. Heidari, E. Safaei, A. Wojtczak, P. Cotic and A. Kozakiewicz, *Polyhedron*, 2013, **55**, 109–116.
- 57 T. R. Forder, M. F. Mahon, M. G. Davidson, T. Woodman and M. D. Jones, *Dalton Trans.*, 2014, **43**, 12095–12099.
- 58 A. W. Addison, T. N. Rao, J. Reedijk, J. van Rijn and G. C. Verschoor, *J. Chem. Soc., Dalton Trans.*, 1984, 1349.
- 59 A. C. Silvino, D. Bonilha de Abreu Talina Martins, A. da Costa Rodrigues and M. L. Dias, *J. Polym. Environ.*, 2013, **21**, 1002–1008.
- 60 A. D. Schwarz, Z. Chu and P. Mountford, *Organometallics*, 2010, **29**, 1246–1260.
- 61 G. Xiao, B. Yan, R. Ma, W. J. Jin, X. Q. Lu, L. Q. Ding, C. Zeng, L. L. Chen and F. Bao, *Polym. Chem.*, 2011, **2**, 659–664.
- 62 C. Paetz and R. Hagen, *Chem. Ing. Tech.*, 2014, **86**, 519–523.
- 63 M. Bouyahyi, E. Grunova, N. Marquet, E. Kirillov, C. M. Thomas, T. Roisnel and J.-F. Carpentier, *Organometallics*, 2008, **27**, 5815–5825.



- 64 A. J. Chmura, M. G. Davidson, C. J. Frankis, M. D. Jones and M. D. Lunn, *Chem. Commun.*, 2008, 1293–1295.
- 65 J. Baran, A. Duda, A. Kowalski, R. Szymanski and S. Penczek, *Macromol. Rapid Commun.*, 1997, **18**, 325–333.
- 66 K. Jamshidi, S. H. Hyon and Y. Ikada, *Polymer*, 1988, **29**, 2229–2234.
- 67 S.-H. Lee, S. Hyun Kim, Y.-K. Han and Y. H. Kim, *J. Polym. Sci., Part A: Polym. Chem.*, 2001, **39**, 973–985.
- 68 Z. Zhong, P. J. Dijkstra and J. Feijen, *Angew. Chem., Int. Ed.*, 2002, **41**, 4510–4513.
- 69 Y. Sarazin, S.-C. Rosca, D.-A. Rosca, V. Dorcet, J.-F. Carpentier, F. M. Kerton and C. M. Kozak, *Dalton Trans.*, 2013, 9361–9375.
- 70 N. Ajellal, J.-F. Carpentier, C. Guillaume, S. M. Guillaume, M. Helou, V. Poirier, Y. Sarazin and A. Trifonov, *Dalton Trans.*, 2010, **39**, 8363–8376.

