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Coordination chemistry of [2 + 2] Schiff-base
macrocycles derived from the dianilines
[(2-NH₂C₆H₄)₂X] (X = CH₂CH₂, O): structural
studies and ROP capability towards cyclic esters†Kuiyuan Wang,^a Timothy J. Prior,^{ib} ^a David L. Hughes,^{ib} ^b Abdessamad Arbaoui^b
and Carl Redshaw^{ib} ^{*a}

Reaction of the [2 + 2] Schiff-base macrocycles {[2-(OH)-5-(R)-C₆H₂-1,3-(CH)₂][CH₂CH₂(2-C₆H₄N)₂]}₂ (R = Me, L¹H₂; tBu, L²H₂) with FeBr₂ afforded the complexes [FeBr(L¹H₂)₂][FeBr₃(O)]·2MeCN (**1**·2MeCN), [FeBr(L²H₂)]X (X = 0.5(FeBr₃)₂O, **2**·0.5MeCN, X = Br, **3**·5.5MeCN), respectively. Reaction of L²H₂ with [KFe(OtBu)₃(THF)] (formed *in situ* from FeBr₂ and KOtBu), following work-up, led to the isolation of the complex [Fe(L²)(L²H)]·3MeCN (**4**·3MeCN), whilst with [CuBr₂] afforded [CuBr(L²H₂)] [CuBr₂]·2MeCN (**5**·2MeCN). Attempts to form mixed Co/Ti species by reaction of [CoBrL²][CoBr₃(NCMe)] with TiCl₄ resulted in [L²H₄][CoBr₄]·2MeCN (**6**·2MeCN). Use of the related oxy-bridged Schiff-base macrocycles {[2-(OH)-5-(R)-C₆H₂-1,3-(CH)₂][O(2-C₆H₄N)₂]}₂ (R = Me, L³H₂; tBu, L⁴H₂) with CoBr₂ led to the isolation of the complexes [(CoBr)₂(L³)]·2C₃H₆O (**7**·2C₃H₆O), [Co(NCMe)₂(L⁴H₂)] [CoBr₄]·5MeCN (**8**·5MeCN), [Co(NCMe)₆][CoBr₃(MeCN)]₂·2MeCN (**9**·2MeCN). For comparative structural/polymerisation studies, the complexes {CoBr(NCMe)L⁵}₂·2MeCN (**10**·2MeCN) and [Co(NCMe)₂L⁵][CoBr₃(NCMe)]₂ (**11**), [FeBr(NCMe)L⁵]₂·2MeCN (**12**·2MeCN) where L⁵H = 2,6-(CHO)₂-4-tBu-C₆H₂OH, as well as the chelate-free salt [Fe(NCMe)₆][FeBr₃OFeBr₃] (**13**) have been isolated and structurally characterized. The ability of these complexes to act as catalysts for the ring opening polymerisation (ROP) of ε-caprolactone (ε-CL) and δ-valerolactone (δ-VL) was investigated, as well as co-polymerisation of ε-CL with *rac*-lactide (*r*-LA) and *vice versa*.

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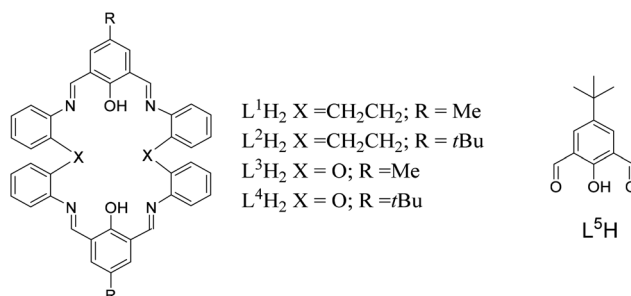
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Introduction

Global issues over plastic pollution continue to drive the search for alternative, more environmentally-friendly materials.¹ As part of our search for new catalysts capable of affording biodegradable polymers *via* the ring opening polymerisation (ROP) of cyclic esters, we have initiated investigations into the coordination chemistry of Schiff-base macrocycles derived from the [2 + 2] condensation of the dianilines [(X)(2-C₆H₄NH₂)₂] (X = CH₂CH₂, O) with the diformylphenols 2,6-(CHO)₂-4-R-C₆H₂OH (R = Me, tBu) (Chart 1).² To date, we have reported how remote alkylaluminium centres bound to the macrocycle (X = CH₂CH₂) exhibited beneficial cooperative effects in the ROP of ε-caprolactone (ε-CL), whereas aluminox-

ane type (Al–O–Al) bonding proved to be detrimental.³ Manganese complexes bearing these macrocycles were far less active (conversions <15%).⁴ However, studies of mixed cobalt/zinc systems revealed interesting catalytic properties, with homo-dinuclear systems exhibiting inactivity while mixed-metal systems proved to be efficient for the ROP of ε-CL and δ-valerolactone (δ-VL).⁵ It is also noteworthy that the structural chemistry of macrocycles of this type remains underexplored; a

Chart 1 Synthesis of ligands L¹H₂–L⁵H prepared herein.^aPlastics Collaboratory, Department of Chemistry, University of Hull, HU6 7RX, UK.
E-mail: c.redshaw@hull.ac.uk^bSchool of Chemistry, University of East Anglia, Norwich, NR4 7TJ, UK

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search of the CSD revealed no hits,⁶ other than our reported aluminium, manganese and cobalt systems.^{3–5} Given this, we have re-focused our efforts on such Schiff-base systems and have extended our studies to iron, cobalt and copper complexes bearing [2 + 2] macrocycles derived from the dianilines [(X)(2-C₆H₄NH₂)₂] (X = CH₂CH₂, O, Chart 2). Herein, we report the molecular structures of these complexes, and have screened a number of them for their capability in the ROP of ϵ -CL, δ -VL and *rac*-lactide (*r*-LA), and for the co-polymerisation of ϵ -CL with *r*-LA and *vice versa*. Poly(ϵ -caprolactone), PCL, and poly(lactide), PLA, are favoured polymers given their biodegradability properties, and their co-polymers are considered as potential environmentally-friendly commodity plastic.⁷

Results and discussion

–CH₂CH₂– bridged systems

Iron

Use $R = \text{Me}$ L^1H_2 . Iron is a cheap, earth abundant metal and its complexes have shown potential in the ROP of cyclic esters.⁸ Given this, we have initiated studies on the iron chemistry of our [2 + 2] macrocyclic systems. The reaction of the macrocycle {[2-(OH)-5-Me-C₆H₂-1,3-(CH₂)₂][CH₂CH₂(2-C₆H₄N)₂]}₂ (L^1H_2) with two equivalents of FeBr₂ in refluxing toluene afforded, following work-up (MeCN), a brown crystalline solid in moderate yield. Single crystals were grown from a saturated solution of acetonitrile on standing at ambient

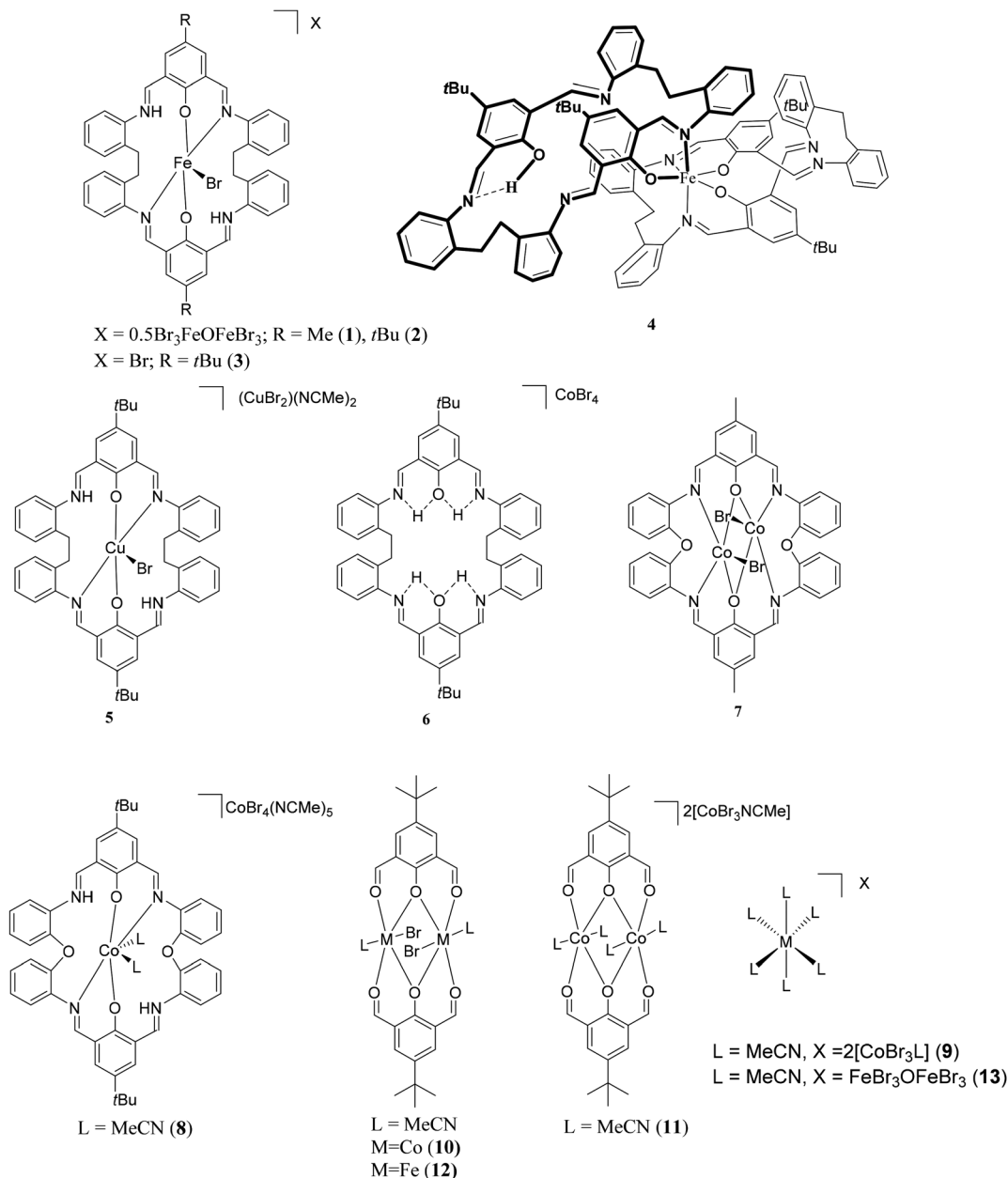


Chart 2 Structures of iron and cobalt complexes 1–13 prepared herein.



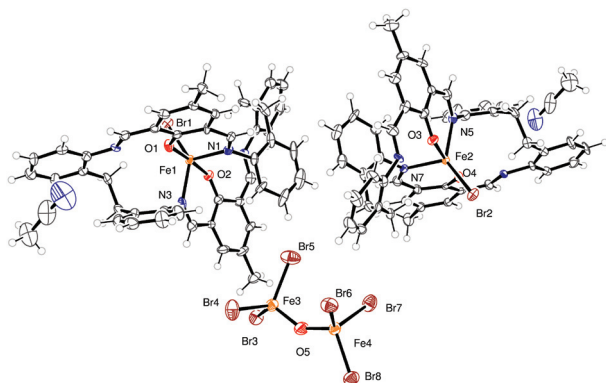


Fig. 1 Asymmetric unit for the structure of $[\text{FeBr}(\text{L}^1)]_2[\text{Br}_3\text{FeOFeBr}_3] \cdot 2(\text{MeCN})$ ($1 \cdot 2\text{MeCN}$). Thermal ellipsoids are drawn at the 50% probability level. Selected bond lengths (Å) and angles (°): $\text{Fe}(1) - \text{O}(1)$ 2.002(6), $\text{Fe}(1) - \text{O}(2)$ 2.020(6), $\text{Fe}(1) - \text{N}(1)$ 2.129(7), $\text{Fe}(1) - \text{N}(3)$ 2.138(7), $\text{Fe}(1) - \text{Br}(1)$ 2.5070(16), $\text{Fe}(2) - \text{O}(3)$ 2.031(6), $\text{Fe}(2) - \text{N}(5)$ 2.150(7), $\text{Fe}(2) - \text{N}(7)$ 2.135(7), $\text{Fe}(2) - \text{Br}(2)$ 2.5104(15); $\text{O}(1) - \text{Fe}(1) - \text{O}(2)$ 177.0(2), $\text{N}(1) - \text{Fe}(1) - \text{N}(3)$ 103.8(3), $\text{Br}(1) - \text{Fe}(1) - \text{N}(1)$ 120.26(19), $\text{N}(5) - \text{Fe}(2) - \text{N}(7)$ 102.9(3).

temperature for 3 days. The molecular structure is shown in Fig. 1, with selected bond lengths and angles given in the caption. There are two macrocyclic iron complexes in the asymmetric unit, related by a pseudosymmetric translation of $c/2$ (see ESI†), plus an anion of $[\text{Br}_3\text{FeOFeBr}_3]$ and two molecules of acetonitrile. Of the macrocyclic bound iron centres, both Fe1 and Fe2 adopt a distorted trigonal bipyramidal geometry ($\tau = 0.69$),⁹ bound by a bromide and two nitrogen atoms and two oxygen atoms of the macrocycle, apical sites are occupied by O atoms, with the bromide and N atoms in the equatorial sites. The composition is thus $[\text{FeBr}(\text{L}^1)]_2[\text{Br}_3\text{FeOFeBr}_3] \cdot 2(\text{MeCN})$ ($1 \cdot 2\text{MeCN}$). In terms of charge, the 2+ available from the anion which contains two Fe(III) centres, is balanced by the two cations ($2 \times +1$), each of which contains an Fe(II) centre.

Use $R = t\text{Bu}$ L^2H_2 . Similar reaction of $\{[2-(\text{OH})-5-(t\text{Bu})-\text{C}_6\text{H}_2-1,3-(\text{CH}_3)_2][\text{CH}_2\text{CH}_2(2-\text{C}_6\text{H}_4\text{N})_2]\}_2$ (L^2H_2) with 2.1 equivalents of FeBr_2 afforded, following work-up, red needles in good yield. Crystals suitable for X-ray diffraction were grown from a saturated solution of acetonitrile at ambient temperature. A view of the molecular structure is shown in Fig. 2, and bond lengths and angles are given in the caption. The crystal comprises $\text{FeBr}(\text{L}^2\text{H}_2)$ cations, $(\text{FeBr}_3)_2\text{O}$ anions and MeCN solvent molecules. The anions lie about a centre of symmetry and the solvent molecule sites refine best with half-occupancy; the molecular formula is therefore $[\text{FeBr}(\text{L}^2\text{H}_2)]$, $0.5[(\text{FeBr}_3)_2\text{O}] \cdot 0.5(\text{MeCN})$ ($2 \cdot 0.5\text{MeCN}$), and like $1 \cdot 2\text{MeCN}$, is a mixed oxidation state Fe(II)/Fe(III) system. The iron atom in the cation is five-coordinate with a trigonal bipyramidal pattern ($\tau = 0.82$),⁹ the apical sites are occupied by O atoms, with the bromide and N atoms in the equatorial sites. There is a pseudo two-fold symmetry axis along the Fe–Br bond. In the macrocyclic ligand, there are four N atoms, each involved in a double bond, *viz.* C11=N2 1.298(5) C52=N1 1.287(6) C26=N3 1.296(6) C37=N4 1.308(7), N1 and N3 are coordinated to the Fe atom,

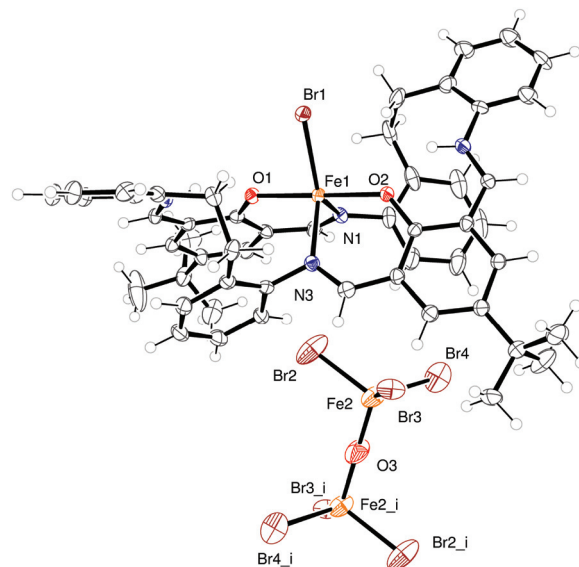


Fig. 2 View of the $[\text{FeBr}(\text{L}^2)][(\text{FeBr}_3)_2\text{O}]_{0.5} \cdot 0.5(\text{MeCN})$ ($2 \cdot 0.5\text{MeCN}$) complex ion, indicating the atom numbering scheme. Thermal ellipsoids are drawn at the 50% probability level. Symmetry equivalent atoms are generated by $i = 1 - x, 1 - y, 1 - z$. Selected bond lengths (Å) and angles (°): $\text{Fe}(1) - \text{O}(1)$ 2.052(3), $\text{Fe}(1) - \text{O}(2)$ 2.059(3), $\text{Fe}(1) - \text{N}(1)$ 2.131(4), $\text{Fe}(1) - \text{N}(3)$ 2.124(4), $\text{Fe}(1) - \text{Br}(1)$ 2.4967(8); $\text{Br}(1) - \text{Fe}(1) - \text{O}(1)$ 89.71(9), $\text{O}(1) - \text{Fe}(1) - \text{N}(3)$ 94.83(14).

while N2 and N4 are bonded to two C atoms (one with a double bond) and a hydrogen atom. The hydrogen atoms were included in the structure factor calculations in a planar, trigonal fashion, and their U_{iso} values were refined freely and satisfactorily; it is believed that both these groups are charged $\text{C}^--\text{N}^+\text{H}=\text{C}$ groups. Both these hydrogen atoms are involved in intramolecular hydrogen bonds.

The $(\text{FeBr}_3)_2\text{O}$ anion lies about a centre of symmetry. One of the bromide ligands is disordered over two sites, in an 82 : 18 occupancy ratio. The nearest neighbours of the bromide atoms are atoms of the disordered *t*-butyl group.

Similar treatment of $\{[2-(\text{OH})-5-(t\text{Bu})-\text{C}_6\text{H}_2-1,3-(\text{CH}_3)_2][\text{O}(2-\text{C}_6\text{H}_4\text{N})_2]\}_2$, but with limited FeBr_2 (1.1 equivalents), led to the isolation of the Fe(II) salt $[\text{FeBr}(\text{L}^2\text{H}_2)]\text{Br} \cdot 3\text{MeCN}$ (5.5MeCN). The molecular structure is provided in the ESI (Fig. S2†).

Use of $[\text{KFe}(\text{OtBu})_3(\text{THF})]$. Given alkoxide species play a central role in metal-catalysed ROP, we attempted to generate an iron alkoxide species. Reaction of L^2H_2 with *in situ* generated $[\text{KFe}(\text{OtBu})_3(\text{THF})]$ in refluxing toluene resulted, after work-up, in the isolation of a brown crystalline material. Crystals grown from a saturated solution of acetonitrile were found to be a bis-chelate structure $[\text{Fe}(\text{L}^2)(\text{L}^2\text{H}_2)] \cdot 3\text{MeCN}$ ($4 \cdot 3\text{MeCN}$) (see Fig. 3), in which a distorted octahedral iron(III) centre is bound to two of the macrocyclic ligands. The asymmetric unit contains one iron complex and 3 molecules of acetonitrile. The coordination at the iron is such that one macrocycle is bound only in chelate fashion *via* N,O-type ligation, whilst the second macrocycle utilizes four atoms to bind in $2 \times \text{N,O}$ -type fashion. We have observed similar binding modes recently for aluminium.^{3c}



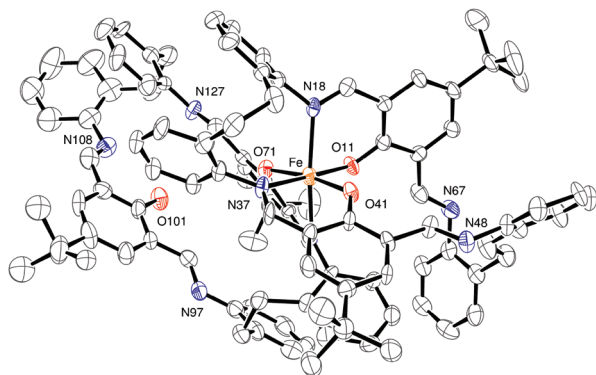


Fig. 3 View of $[\text{Fe}(\text{L}^2)(\text{L}^2\text{H})]\cdot 3\text{MeCN}$ (**4**·**3MeCN**), indicating the atom numbering scheme. Thermal ellipsoids are drawn at the 50% probability level. For clarity hydrogen atoms are not shown. Selected bond lengths (Å) and angles (°): Fe(1)–O(11) 1.936(4), Fe–O(41) 1.931(4), Fe–O(71) 1.905(4), Fe–N(18) 2.234(5), Fe–N(37) 2.189(5), Fe–N(78) 2.240(5); O(11)–Fe–O(41) 92.1(2), O(11)–Fe–N(37) 176.6(2), N(18)–Fe–N(78) 164.0(2).

Use of copper bromide. Copper is also an earth abundant metal, though it has had only limited success in the ROP of cyclic esters.¹⁰ For the successful systems, Schiff-base ligation appears beneficial, suggesting that interaction with the macrocyclic systems of the type herein could lead to a ROP active complex. Reaction of L^2H_2 with two equivalents of CuBr_2 afforded, following work-up, the orange/brown Cu salt complex $[\text{CuBr}(\text{L}^2\text{H}_2)]\cdot 2\text{MeCN}$ (**5**·**2MeCN**) in good yield. In the cation, the copper centre is distorted trigonal bipyramidal with oxygens at the apex and two nitrogens and a bromide in the equatorial plane ($\tau = 0.90$).⁹ The anion $\text{Cu}(\text{I})\text{Br}_2^-$ resides close to a pocket formed by the macrocycle, see Fig. 4.

Attempted mixed-metal systems

Having successfully isolated mixed cobalt/zinc systems, which were active for ring opening polymerization whereas the analo-

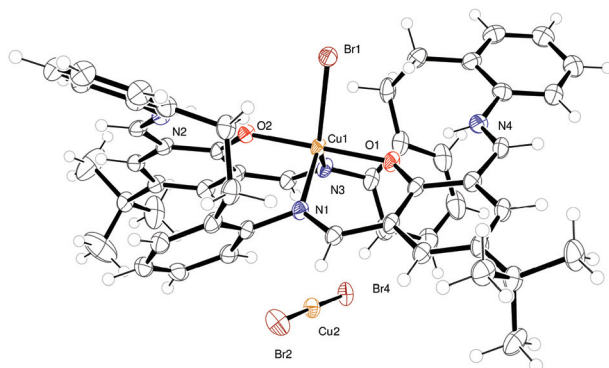


Fig. 4 Molecular structure of $[\text{CuBr}(\text{L}^2\text{H}_2)]\cdot 2\text{MeCN}$ (**5**·**2MeCN**) with atoms drawn as 50% probability ellipsoids. Solvent molecules are not shown. Selected bond lengths (Å) and angles (°): Cu(1)–O(1) 1.955(3), Cu(1)–O(2) 1.948(3), Cu(1)–N(1) 2.106(3), Cu(1)–N(3) 2.074(3), Cu(1)–Br(1) 2.4790(6); O(1)–Cu(1)–O(2) 179.8(1), N(1)–Cu(1)–Br(1) 119.58(9), N(3)–Cu(1)–Br(1) 125.8(1).

gous homodinuclear species were not,⁵ we attempted to prepare mixed cobalt/titanium complexes. Our entry point was again the cobalt complex $[\text{CoBrL}^2\text{H}_2][\text{CoBr}_3(\text{NCMe})]$, and reaction with $[\text{TiCl}_4]$ resulted in $[\text{L}^2\text{H}_4][\text{CoBr}_4]\cdot 2\text{MeCN}$ (**6**·**2MeCN**) as the only crystalline product. The molecular structure of the salt **6** comprises a protonated macrocycle L^2H_2 and the $\text{Co}(\text{II})$ containing anion $[\text{CoBr}_4]^{2-}$; see ESI (Fig. S4†) for details.

–O– bridged systems

We have also initiated a programme to investigate the coordination chemistry of the oxy-bridged macrocycles $\{[2-(\text{OH})-5-(\text{R})-\text{C}_6\text{H}_2-1,3-(\text{CH}_2)_2][\text{O}(2-\text{C}_6\text{H}_4\text{N})_2]\}_2$ ($\text{R} = \text{Me}$ L^3H_2 , $t\text{Bu}$ L^4H_2). Reaction of L^3H_2 with two equivalents of CoBr_2 afforded, following work-up (acetone), the complex $[(\text{CoBr})_2\text{L}^3]\cdot 2\text{C}_3\text{H}_6\text{O}$ (**7**·**2C₃H₆O**), containing two $\text{Co}(\text{II})$ centres, as black crystals in *ca.* 40% isolated yield. Single crystals can be grown from a saturated solution of acetone at ambient temperature, and the molecular structure is shown in Fig. 5; selected bond lengths and angles are given in the caption. Both cobalt centres adopt distorted trigonal bipyramidal with a bromide at each apex ($\tau = 0.59$ and 0.53 for Co1 and Co2 , respectively).⁹

In the case of L^4H_2 , reaction with CoBr_2 afforded, following work-up (MeCN), two sets of crystals ($\sim 90 : 10$). Both sets were subjected to single crystal X-ray diffraction, and the major product, as shown in Fig. 6, was found to be the $\text{Co}(\text{II})$ containing salt $[\text{Co}(\text{NCMe})_2(\text{L}^4\text{H}_2)][\text{CoBr}_4]\cdot 5\text{MeCN}$ (**8**·**5MeCN**). In the cation, the cobalt centre is distorted octahedral, and is bound by two phenoxide oxygens and the two nitrogens N(1) and N(3) of the macrocycle plus two bound acetonitrile ligands. In the solid-state, there are $\text{C}-\text{H}\cdots\text{Br}$ interactions present. The minor product was found to be $\text{Co}(\text{II})$ containing salt $[\text{Co}$

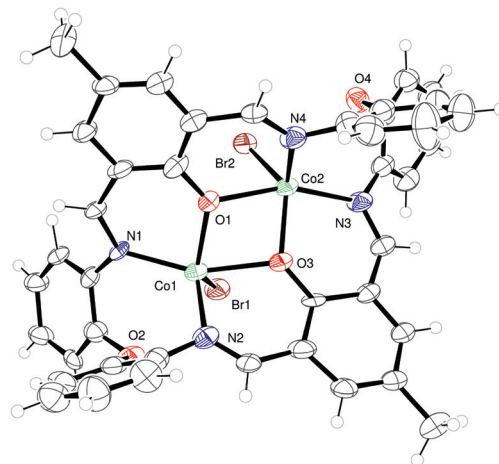


Fig. 5 Molecular structure of $[(\text{CoBr})_2(\text{L}^3)]\cdot 2\text{C}_3\text{H}_6\text{O}$ (**7**·**2C₃H₆O**) with atoms drawn as 50% probability ellipsoids. Selected bond lengths (Å) and angles (°): Co(1)–O(1) 1.982(11), Co(1)–O(3) 2.216(11), Co(1)–N(1) 2.067(12), Co(1)–N(2) 2.091(13), Co(1)–Br(1) 2.421(3), Co(2)–O(1) 2.179(11), Co(2)–O(3) 1.977(11), Co(2)–N(3) 2.102(14), Co(2)–N(4) 2.079(13), Co(2)–Br(2) 2.421(3); O(1)–Co(1)–O(3) 70.0(4), N(1)–Co(1)–N(2) 112.3(5), O(1)–Co(1)–Br(1) 122.2(3), O(1)–Co(2)–N(3) 157.8(5), N(4)–Co(2)–Br(2) 111.2(4).



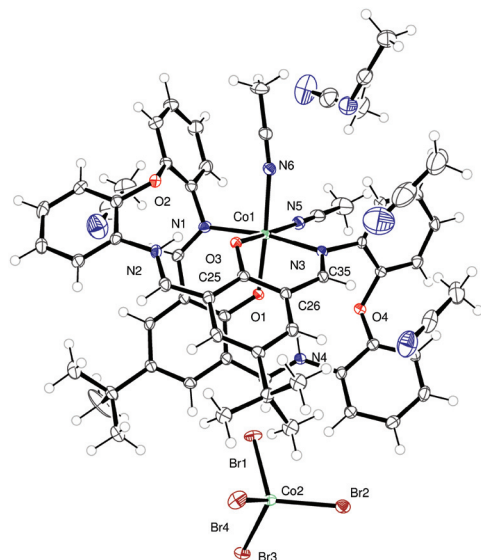


Fig. 6 Molecular structure of $[\text{Co}(\text{NCMe})_2(\text{L}^4)][\text{CoBr}_4] \cdot 5\text{MeCN}$ (**8·5MeCN**) with atoms drawn as 50% probability ellipsoids. Selected bond lengths (Å) and angles (°): $\text{Co}(1) - \text{O}(1)$ 2.016(3), $\text{Co}(1) - \text{O}(3)$ 2.056(3), $\text{Co}(1) - \text{N}(1)$ 2.156(3), $\text{Co}(1) - \text{N}(3)$ 2.156(3), $\text{Co}(1) - \text{N}(5)$ 2.121(3), $\text{Co}(1) - \text{N}(6)$ 2.114(3); $\text{O}(1) - \text{Co}(1) - \text{O}(3)$ 91.98(11), $\text{N}(1) - \text{Co}(1) - \text{N}(3)$ 169.15(12), $\text{O}(1) - \text{Co}(1) - \text{N}(6)$ 174.71(13), $\text{O}(3) - \text{Co}(1) - \text{N}(5)$ 174.77(13)°.

(NCMe)₆][CoBr₃NCMe]₂·2MeCN (**9·2MeCN**), details of this structure can be found in the ESI (Fig. S4†).

'Dialdehyde' systems

To probe the role played by the presence of the macrocycle during catalysis (see ROP section), we have also prepared cobalt complexes bearing chelate ligands derived from the 4-*tert*-butyl-2,6-diformylphenol 2,6-(CHO)₂-4-*t*Bu-C₆H₂OH (L^5H). Reaction of L^5H with CoBr_2 in the presence of excess Et_3N afforded, after work-up (MeCN), the $\text{Co}(\text{II})$ complex $\{[\text{CoBr}(\text{NCMe})] \text{L}^5\}_2 \cdot 2\text{MeCN}$ (**10·2MeCN**) as orange/brown prisms in good yield. The molecular structure is shown in Fig. 7, with selected bond lengths and angles given in the caption. Each cobalt centre is distorted octahedral, and are bound by oxygens from two L ligands and a bromide and acetonitrile ligand. The ligand arrangement is such that one bromide and an MeCN ligand reside above the Co_2O_6 plane and one bromide and an MeCN ligand are below this plane.

If the same reaction is conducted in the absence of Et_3N , then, following work-up (MeCN), green prisms are isolated in good yield. The molecular structure (see Fig. 8 for the cation) revealed the structure of the complex to be $\text{Co}(\text{II})$ containing salt $[\text{Co}(\text{NCMe})_2 \text{L}^5]_2 [\text{CoBr}_3(\text{NCMe})]_2$ (**11**). The dimer is centrosymmetric – the second half of the dimer is generated by the symmetry operation $-x, 1 - y, 1 - z$. Each cobalt centre in the cation is distorted octahedral, but unlike **10**, there is no bound bromide.

If a similar reaction is conducted with FeBr_2 in the presence of Et_3N , then, following work-up (MeCN), black prisms are iso-

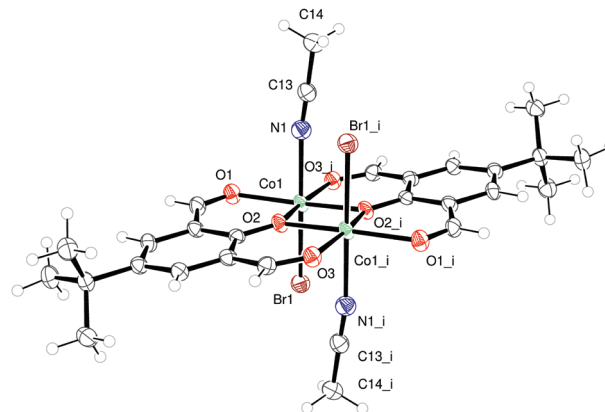


Fig. 7 Molecular structure of $\{[\text{CoBr}(\text{NCMe})] \text{L}^5\}_2 \cdot 2\text{MeCN}$ (**10·2MeCN**) with atoms drawn as 50% probability ellipsoids (unbound solvent not shown). Selected bond lengths (Å) and angles (°): $\text{Co}(1) - \text{O}(1)$ 2.052(2), $\text{Co}(1) - \text{O}(2)$ 2.079(2), $\text{Co}(1) - \text{Br}(1)$ 2.5751(6), $\text{Co}(1) - \text{N}(1)$ 2.163(3); $\text{O}(1) - \text{Co}(1) - \text{O}(2)$ 87.17(8), $\text{Co}(1) - \text{O}(2) - \text{Co}(1_i)$ 100.59(9), $\text{Br}(1) - \text{Co}(1) - \text{N}(1)$ 169.51(8).

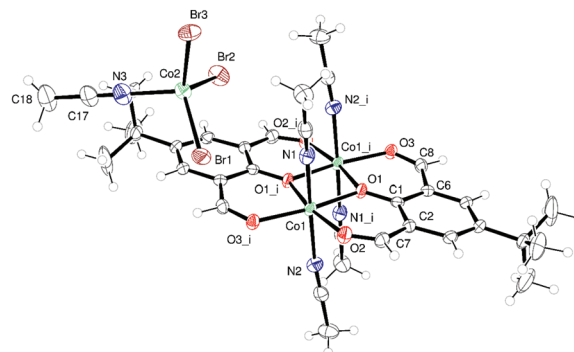


Fig. 8 Molecular structure of $[\text{Co}(\text{NCMe})_2 \text{L}^5]_2 [\text{CoBr}_3(\text{NCMe})]_2$ (**11**) with atoms drawn as 50% probability ellipsoids. Symmetry operation used to generate equivalent atoms: $i = -x, 1 - y, 1 - z$. Selected bond lengths (Å) and angles (°): $\text{Co}(1) - \text{O}(1)$ 2.046(2), $\text{Co}(1) - \text{O}(2)$ 2.050(2), $\text{Co}(1) - \text{N}(1)$ 2.108(3), $\text{Co}(1) - \text{N}(2)$ 2.143(3); $\text{O}(1) - \text{Co}(1) - \text{O}(2)$ 88.56(9), $\text{Co}(1) - \text{O}(2) - \text{Co}(1_i)$ 99.59(9), $\text{N}(1) - \text{Co}(1) - \text{N}(2)$ 172.07(11).

lated in good yield. The molecular structure (Fig. S5†) revealed the structure of the $\text{Fe}(\text{II})$ complex to be $[\text{FeBr}(\text{NCMe}) \text{L}^5]_2 \cdot 2\text{MeCN}$ (**12·2MeCN**). Each iron centre in the cation is distorted octahedral, bound by oxygens from two L ligands, a bromide and an acetonitrile ligand.

Finally, for comparative catalytic studies, the salt $[\text{Fe}(\text{NCMe})_6][\text{FeBr}_3\text{OFeBr}_3]$ (**13**) was prepared from L^1H_2 and FeBr_3 .

Ring opening polymerisation (ROP) of ϵ -CL and δ -VL

The Fe, Cu and Co complexes **1**, **2**, **4**, **5**, **9–13** have been screened for their ability to ring open polymerise ϵ -caprolactone (ϵ -CL) and δ -valerolactone (δ -VL); runs were conducted in the presence of benzyl alcohol (BnOH). For complexes **1**, **2** and **4**, a variety of conditions were used in the attempted ROP of ϵ -



Table 1 Synthesis of polycaprolactone using catalysts **1**, **2**, **4**, **9–13**

Run	Cat.	[Monomer]:[cat]:BnOH	<i>T</i> (°C)	<i>t</i> /h	Conv. ^a (%)	<i>M_n</i> × 10 ³ ^b	<i>M_n</i> _{calcd} × 10 ⁴ ^c	PDI ^d
1	1	500:1:1	130	24	63.2	3.70	3.61	1.95
2	1	250:1:1	130	24	88.7	2.95	2.54	3.28
3	1	500:1:1	130	12	79.4	0.97	4.53	1.11
4	1	500:1:1	130	12	66.3	0.70	3.79	1.17
5	2	500:1:1	130	24	99.3	5.92	5.66	1.56
6	2	250:1:1	130	24	55.0	2.60	1.58	1.06
7	4	500:1:1	130	24	68.9	8.67	3.94	1.75
8	4	250:1:1	130	24	87.6	7.52	2.50	1.58
9	9	500:1:1	130	24	—	—	—	—
10	10	500:1:1	130	24	—	—	—	—
11	11	500:1:1	130	24	—	—	—	—
12	12	500:1:1	130	24	8.4	—	0.45	—
13	13	500:1:1	130	24	15.3	—	0.87	—

^a Conversion was confirmed by ¹H NMR spectroscopy. ^b Determined by GPC analysis calibrated with polystyrene standards and multiplied by correction factor of 0.56. ^c F.W. ([M]/[BnOH])(conversion) + BnOH. ^d Polydispersity index (*M_w*/*M_n*) were determined by GPC.

CL including differing ratios of [ε-CL]:[Cat]:[BnOH] and run times, and it was found that the Fe compounds were far more active than the other metal complexes screened herein. ROP systems based on Fe,⁸ Cu¹⁰ and Co^{5,11} have been reported with mixed success.

The polymerisation data for the ε-CL runs is given in Table 1. Highest conversion was achieved using **2** in run 5 using the ratio 500:1:1 at 130 °C over 24 h, albeit with less control than observed in run 6 using 250:1:1. Systems employing **1** and **4** were also less controlled, whilst the non-macrocylic cobalt systems **9–11** proved to be inactive under the conditions employed herein. Interestingly, the non-macrocylic iron complexes **12** and **13** proved to be poorly active with low conversions for ε-CL. However, for δ-VL, a 32% conversion (affording small molecular weight oligomers) was observed for **13**, whereas **12** was inactive. The MALDI-TOF spectra can be interpreted using the formula C₆H₆–CH₂O[O(CH₂)₅CO]_{*n*}OH, for example see run 5 (Fig. S7, ESI†). The ¹H NMR spectrum of the PCL indicated the presence of benzyloxy and hydroxyl end

groups (Fig. S8, ESI†). There was evidence of significant transesterification, whilst all observed *M_n* values were significantly lower than the calculated values.

In the case of δ-VL (Table 2), complexes **1** and **2** exhibited better conversions *versus* **4** and **12**, whilst **5**, **9**, **10** and **11** proved to be inactive. In the MALDI-TOF mass spectra of the PVL, a number of families of peaks were observed separated by 100 mass units (Fig. S22, ESI†) can be assigned by the formula C₆H₆–CH₂O[O(CH₂)₄CO]_{*n*}OH; the highest were typically about 6000 (run 4, Fig. S21†). The ¹H NMR spectra of the PVL (*e.g.* Fig. S20†) revealed the presence of benzyloxy and OH end groups. The observed molecular weights were lower than the calculated values, suggesting the presence of a chain transfer agent (H₂O or BnOH).

From a kinetic study (Fig. S11 and S18†), it was observed that the polymerisation rate exhibited near first order dependence on the ε-CL or δ-VL concentration at 130 °C, and complex **2** displayed the best rate in both δ-VL [*K_{obs}* = 7.29 × 10^{−3} h^{−1}] and ε-CL polymerisations [*K_{obs}* = 4.70 × 10^{−3} h^{−1}].

Table 2 ROP of δ-VL using complexes **1**, **2**, **4**, **5**, **9–13**

Run	Cat.	[VL]:[Cat]:BnOH	<i>T</i> (°C)	<i>t</i> /h	Conv. ^a (%)	<i>M_n</i> × 10 ³ ^b	<i>M_n</i> _{calcd} × 10 ⁴ ^c	PDI ^d
1	1	500:1:1	130	24	98.6	2.52	4.94	1.14
2	1	250:1:1	130	24	94.3	3.10	2.36	1.28
3	1	500:1:1	130	12	39.7	—	2.00	—
4	2	500:1:1	130	24	96.1	6.00	4.92	1.38
5	2	250:1:1	130	24	93.2	4.53	2.41	1.23
6	2	500:1:1	130	12	83.8	—	4.20	—
7	4	500:1:1	130	24	34.5	2.59	1.84	1.25
8	4	250:1:1	130	24	23.5	2.51	0.60	1.27
9	5	500:1:1	130	24	—	—	—	—
10	9	500:1:1	130	24	—	—	—	—
11	10	500:1:1	130	24	—	—	—	—
12	11	500:1:1	130	24	—	—	—	—
13	12	500:1:1	130	24	—	—	—	—
14	13	500:1:1	130	24	32.0	0.76	2.18	1.26

^a Conversion was confirmed by ¹H NMR spectroscopy. ^b Determined by GPC analysis calibrated with polystyrene standards and multiplied by correction factor of 0.58. ^c F.W. ([M]/[BnOH])(conversion) + BnOH. ^d Polydispersity index (*M_w*/*M_n*) were determined by GPC.



Table 3 Synthesis of block copolymers from cyclic ester monomers using the Fe catalysts **1**, **2** and **4**

Run	Composition	Catalyst	<i>t</i> (h)	Incorporated amount ^a	<i>M_n</i> × 10 ³ ^b	PDI
1	Poly (ε-CL + <i>r</i> -LA) [ε-CL] : [<i>r</i> -LA] : [cat] : [BnOH] = 500 : 500 : 1 : 1	1	24 + 24 :	ε-CL : <i>r</i> -LA = 47 : 53	3.75	2.22
2	Poly (δ-VL + <i>r</i> -LA) [δ-VL] : [<i>r</i> -LA] : [cat] : [BnOH] = 500 : 500 : 1 : 1	1	24 + 24	δ-VL : <i>r</i> -LA = 14 : 86	5.51	1.73
3	Poly (<i>r</i> -LA + ε-CL) [<i>r</i> -LA] : [ε-CL] : [cat] : [BnOH] = 500 : 500 : 1 : 1	2	24 + 24	<i>r</i> -LA : ε-CL = 49 : 51	10.55	1.51
4	Poly (<i>r</i> -LA + ε-CL) [ε-CL] : [<i>r</i> -LA] : [cat] : [BnOH] = 250 : 250 : 1 : 1	2	24 + 24	<i>r</i> -LA : ε-CL = 93 : 7	6.20	1.82
5	Poly (ε-CL + <i>r</i> -LA) [ε-CL] : [<i>r</i> -LA] : [cat] : [BnOH] = 500 : 500 : 1 : 1	2	24 + 24	ε-CL : <i>r</i> -LA = 60 : 40	9.31	1.64
6	Poly (ε-CL + <i>r</i> -LA) [ε-CL] : [<i>r</i> -LA] : [cat] : [BnOH] = 250 : 250 : 1 : 1	2	24 + 24	ε-CL : <i>r</i> -LA = 63 : 37	12.63	3.02
7	Poly (δ-VL + <i>r</i> -LA) [δ-VL] : [<i>r</i> -LA] : [cat] : [BnOH] = 500 : 500 : 1 : 1	2	24 + 24	δ-VL : <i>r</i> -LA = 55 : 45	3.40	1.26
8	Poly (δ-VL + <i>r</i> -LA) [δ-VL] : [<i>r</i> -LA] : [cat] : [BnOH] = 250 : 250 : 1 : 1	2	24 + 24	δ-VL : <i>r</i> -LA = 88 : 12	2.57	2.56
9	Poly (ε-CL + <i>r</i> -LA) [ε-CL] : [<i>r</i> -LA] : [cat] : [BnOH] = 500 : 500 : 1 : 1	4	24 + 24	ε-CL : <i>r</i> -LA = 100 : 0	4.61	1.23
10	Poly (ε-CL + <i>r</i> -LA) [ε-CL] : [<i>r</i> -LA] : [cat] : [BnOH] = 250 : 250 : 1 : 1	4	24 + 24	ε-CL : <i>r</i> -LA = 90 : 10	3.61	1.18

^a Conversion was confirmed by ¹H NMR spectroscopy. ^b Determined by GPC analysis calibrated with polystyrene standards and multiplied by correction factor of 0.56.

Synthesis of block co-polymers

The co-polymerisation of ε-CL with *r*-LA and of δ-VL with *r*-LA was also examined (Table 3). Noteworthy, the co-polymerisation of ε-CL with *r*-LA is much easier than δ-VL with *r*-LA when using the Fe complexes. In the presence of complex **2**, the *M_n* of the block co-poly(ε-CL + *r*-LA) reached around 12 000. Complexes **1** and **4** were less active in these co-polymerisations. As observed by ¹H NMR spectroscopy (Fig. S14 and S24†), the co-polymers were also capped by benzyloxy and hydroxyl end groups. The composition of the copolymer was further illustrated by ¹³C NMR spectroscopy (Fig. S17 and S25†). From the MALDI-ToF mass spectrum (positive mode) of poly(ε-CL + *r*-LA) a gap of 114 corresponding to the molecular weight of ε-CL was evident, whilst running the spectra in negative method revealed a gap of 144 corresponding to the molecular weight of *r*-LA (Fig. S12 and 13†). The MALDI-ToF mass spectrum for co-poly (δ-VL + *r*-LA) was also recorded, see Fig. S26.† The 2D J-resolved ¹H NMR spectrum for the copolymer was recorded and the peaks were assigned to the corresponding tetrads (see Fig. S17 and S22, ESI†) according to literature reports,¹² which revealed an atactic LA-LA-LA chain.

Conclusion

In conclusion, we have successfully synthesised iron, cobalt and copper complexes bearing [2 + 2] Schiff-base macrocycles derived from dianilines containing CH₂CH₂ bridges. Cobalt complexes were prepared from a related dianiline containing an oxy bridge. A number on non-macrocylic Fe and Co complexes were also prepared for comparative catalytic studies. The iron complexes outperformed the other metal systems

herein for the ring opening polymerisation of both ε-caprolactone and δ-valerolactone. Best results were obtained using the iron salt **2**, however the non-macrocylic system **12** also performed well. Complex **2** was also capable of the copolymerisation of ε-CL (or δ-VL) with *rac*-lactide, affording copolymers appreciable amounts of each monomer incorporated.

Experimental

General

All manipulations were carried out under an atmosphere of nitrogen using standard Schlenk and cannula techniques or in a conventional nitrogen-filled glove-box. Toluene was refluxed over sodium, whilst acetonitrile was refluxed over calcium hydride. IR spectra (nujol mulls, KBr windows) were recorded on a Nicolet Avatar 360 FT-IR spectrometer; ¹H and ¹³C NMR spectra were recorded at room temperature on a Varian VXR 400 S spectrometer at 400 MHz or a Gemini 300 NMR spectrometer or a Bruker Advance DPX-300 spectrometer. The ¹H NMR spectra were calibrated against the residual protio impurity of the deuterated solvent. Elemental analyses were performed by the elemental analysis service at the London Metropolitan University, the Chemistry Department at the University of Hull or Nanjing University. The precursors 2,6-(CHO)₂-4-R-C₆H₂OH (R = Me, *t*Bu) and 2,2'-ethylenedianiline (or 2,2'-oxydianiline) and the Schiff-base pro-ligands were prepared by the literature.^{3a,c,13–15} For the iron and cobalt complexes, all manipulations were carried out under an atmosphere of dry nitrogen using conventional Schlenk and cannula techniques or in a conventional nitrogen-filled glove box. All solvents were distilled and degassed prior to use (Table 2).



Synthesis of $[\text{FeBr}(\text{L}^1\text{H}_2)]_2[(\text{FeBr}_3)_2\text{O}] \cdot 2(\text{MeCN})$ (1·2MeCN)

L^1H_2 (0.52 g, 0.77 mmol) and FeBr_2 (0.34 g, 1.58 mmol) were combined in a Schlenk and toluene (20 mL) was added. After refluxing for 12 h, the volatiles were removed *in-vacuo*, and the residue was extracted into MeCN (20 mL). Prolonged standing at room temperature afforded orange/brown prisms. Yield: 0.72 g, 78%. Elemental analysis calculated for $\text{C}_{96}\text{H}_{82}\text{Br}_8\text{Fe}_4\text{N}_{10}\text{O}_5$: required: C 49.32% H 3.60% N 5.00% Found: 49.88% H 3.58% N 5.09%. IR (KBr, cm^{-1}): 3378 (s), 2923 (s), 2726 (w), 2671 (w), 2360 (w), 1633 (m), 1585 (m), 1538 (m), 1462 (s), 1377 (s), 1300 (w), 1280 (w), 1238 (m), 1102 (m), 977 (w), 873 (w), 801 (m), 754 (m), 722 (m), 688 (w), 622 (w), 574 (w), 534 (w), 502 (m), 486 (w), 455 (w). M.S. (MALDI-ToF): 817 (M^+). Magnetic moment: 6.35 B.M.¹⁶

Synthesis of $[\text{FeBr}(\text{L}^2\text{H}_2)]_2[(\text{FeBr}_3)_2\text{O}] \cdot 0.5\text{MeCN}$ (2·0.5MeCN)

To the pro-ligand L^2H_2 (0.50 g, 0.65 mmol) in toluene was added 2.1 equivalents of FeBr_2 (0.30 g, 1.39 mmol), and the system was refluxed for 12 h. On cooling, the volatiles were removed under vacuum, and the residue was extracted into acetonitrile (30 mL) to afford 2·MeCN as red needles (0.48 g, 56%). Elemental analysis calculated for $\text{C}_{106}\text{H}_{107}\text{Br}_8\text{Fe}_4\text{N}_9\text{O}_5$: C 51.97, H 4.40, N 5.15%; found: C 51.21, H 4.43, N 5.55%. IR (KBr, cm^{-1}): 3168 (m), 2925 (s), 2854 (s), 2726 (w), 1704 (m), 1620 (m), 1587 (m), 1542 (s), 1462 (s), 1377 (s), 1317 (w), 1260 (w), 1233 (m), 1214 (m), 1180 (m), 1130 (w), 933 (w), 890 (w), 839 (w), 799 (m), 744 (m), 722 (m), 526 (m), 449 (m). M.S. (MALDI-ToF): 901 ($\text{M} - \text{anion}$). Magnetic moment: 5.86 B.M.¹⁶

Synthesis of $[\text{FeBr}(\text{L}^2\text{H}_2)]\text{Br} \cdot 3\text{MeCN}$ (5.5MeCN)

As for 2, but using L^2H_2 (0.52 g, 0.68 mmol) and FeBr_2 (0.16 g, 0.74 mmol) affording 3·5.5MeCN as brown prisms. Yield: 0.52 g, 65%. IR (KBr, cm^{-1}): 3168 (m), 2925 (s), 2854 (s), 2726 (w), 1636 (s), 1619 (s), 1599 (m), 1587 (m), 1537 (s), 1463 (s), 1377 (s), 1333 (m), 1283 (w), 1241 (m), 1183 (m), 1101 (m), 1063 (s), 1007 (m), 976 (m), 880 (s), 794 (m), 753 (s), 722 (m), 689 (m), 622 (w), 595 (w), 575 (w), 535 (w), 513 (m), 494 (m), 476 (w). M.S. (MALDI-ToF): 901 ($\text{M} - 5.5\text{MeCN} - \text{anion}$).

Synthesis of $[\text{Fe}(\text{L}^2)(\text{L}^2\text{H})] \cdot 3\text{MeCN}$ (4·3MeCN)

To FeBr_2 (1.00 g, 4.64 mmol) was added $\text{KO}t\text{Bu}$ (1.04 g, 9.27 mmol) in THF (30 mL) at 0 °C and the system was stirred for 5 h. Following removal of the volatiles, L^2H_2 (3.57 g, 4.64 mmol) and toluene (20 mL) was added and the system was refluxed for 12 h. On cooling, the volatiles were removed and the residue was extracted into cold acetonitrile (30 mL). 4·3MeCN formed. Yield 1.62 g, 41%. Elemental analysis calculated for $\text{C}_{104}\text{H}_{101}\text{FeN}_8\text{O}_4$ (sample dried *in-vacuo* for 2 h): C 78.92, H 6.43, N 7.08%. Found C 79.09, H, 6.49, N On prolonged standing at ambient dark brown block. Yield: 1.84 g, 46.6%. IR (KBr, cm^{-1}): 3377 (w), 2958 (s), 2924 (s), 2854 (s), 2726 (w), 1630 (m), 1587 (m), 1461 (s), 1415 (w), 1377 (s), 1260 (s), 1202 (w), 1093 (s), 1019 (s), 863 (m), 800 (s), 755 (w), 740 (w), 723 (m), 705 (w), 662 (w), 566 (w), 530 (w), 504 (w), 465 (w). MS (Maldi): 1584 ($\text{M} + \text{H}$)⁺. Magnetic moment: 5.07 B.M.¹⁶

Synthesis of $[\text{CuBr}(\text{L}^2\text{H}_2)][\text{CuBr}_2] \cdot 2\text{MeCN}$ (5·2MeCN)

As for 1, but using L^2H_2 (0.52 g, 0.68 mmol) and CuBr_2 (0.30 g, 1.34 mmol), affording 5 as brown prisms. Yield 0.69 g, 84%. Elemental analysis calculated for $\text{C}_{52}\text{H}_{52}\text{Br}_3\text{Cu}_2\text{N}_4\text{O}_2$: C 55.18, H 4.63, N 4.95%. Found C 54.59, H, 4.48, N 4.93%. IR (KBr, cm^{-1}): 3171 (m), 2922 (s), 2853 (s), 2727 (s), 2671 (w), 2350 (w), 1633 (s), 1618 (s), 1594 (s), 1538 (s), 1463 (s), 1377 (s), 1336 (w), 1285 (m), 1260 (w), 1249 (m), 1239 (w), 1183 (m), 1156 (w), 1103 (m), 1062 (m), 1022 (w), 977 (w), 950 (w), 938 (w), 876 (s), 846 (m), 830 (m), 794 (m), 754 (s), 738 (w), 722 (s), 688 (m), 623 (m), 594 (w), 575 (m), 558 (m), 550 (m). M.S. (ESI): 908 ($\text{M} - \text{anion}$), 827 ($\text{M} - \text{anion} - \text{Br}$). Magnetic moment: 1.19 B.M.¹⁷

Synthesis of $[\text{L}^2\text{H}_4][\text{CoBr}_4] \cdot 2\text{MeCN}$ (6·2MeCN)

To $[\text{CoBrL}^2][\text{CoBr}_3(\text{NCMe})]$ (1.00 g, 0.80 mmol) in toluene (30 mL) was added $[\text{TiCl}_4]$ (0.80 mL, 1.0 M, 0.80 mmol) and the system was refluxed for 12 h. On cooling, the volatiles were removed and the residue was extracted into acetonitrile (30 mL). On prolonged standing at 0 °C small red blocks of 6·2MeCN formed. Yield: 0.38 g, 83%. Elemental analysis calculated for $\text{C}_{52}\text{H}_{54}\text{Br}_4\text{CoN}_4\text{O}_2$: C 54.52, H 4.75, N 4.89%. Found C 53.34, H, 5.19, N 4.95%. IR (KBr, cm^{-1}): 3381 (m), 2925 (s), 2924 (s), 2854 (s), 2726 (w), 2359 (w), 1636 (w), 1618 (s), 1594 (m), 1574 (w), 1538 (s), 1462 (s), 1377 (s), 1333 (s), 1285 (m), 1260 (m), 1183 (s), 1102 (m), 1063 (m), 1021 (m), 873 (m), 799 (s), 756 (m), 722 (m), 688 (w), 668 (w), 624 (w), 576 (w), 558 (w), 522 (w). M.S. (ESI): 847 ($\text{M} - \text{CoBr}_3$). 829 ($\text{M} - \text{Br}_4$). 766.97 ($\text{M} - \text{CoBr}_4$). Magnetic moment: 5.05 B.M.¹⁸

Synthesis of $[(\text{CoBr})_2(\text{L}^3)] \cdot 2\text{C}_3\text{H}_6\text{O}$ (7·2C₃H₆O)

To the pro-ligand L^3H_2 (1.00 g, 1.52 mmol) in toluene was added 2.1 equivalents of CoBr_2 (0.70 g, 3.20 mmol), and the system was refluxed for 12 h. On cooling, the volatile were removed under vacuum, and the residue was extracted into acetone (30 mL) to afford 7·2C₃H₆O as black crystals (0.98 g, 62%). Single crystals can also be grown from a saturated acetone solution of 7. Elemental analysis calculated for $\text{C}_{42}\text{H}_{30}\text{Br}_2\text{Co}_2\text{N}_4\text{O}_4$: C 54.10, H 3.24, N 6.01%. Found C 53.88, H 3.19, N 6.25%. IR (KBr, cm^{-1}): 2924 (s), 2853 (s), 2727 (w), 1746 (w), 1693 (w), 1620 (m), 1587 (m), 1537 (m), 1463 (s), 1377 (s), 1261 (m), 1235 (m), 1215 (m), 1151 (w), 1106 (w), 1072 (m), 1030 (w), 891 (w), 867 (w), 842 (w), 800 (m), 757 (w), 722 (m), 534 (w), 478 (w), 449 (w). M.S. (ESI): 851 ($\text{M} - 2\text{C}_3\text{H}_6\text{O} - \text{Br}$), 714 ($\text{M} - 2\text{C}_3\text{H}_6\text{O} - \text{CoBr}$). Magnetic moment: 6.10 B.M.¹⁸

Synthesis of $[\text{Co}(\text{NCMe})_2(\text{L}^4\text{H}_2)][\text{CoBr}_4] \cdot 5\text{MeCN}$ (8·5MeCN)

To the pro-ligand L^4H_2 (1.00 g, 1.36 mmol) in toluene was added 2.1 equivalents of CoBr_2 (0.62 g, 2.83 mmol), and the system was refluxed for 12 h. On cooling, the volatile were removed under vacuum, and the residue was extracted into acetonitrile (30 mL) to afford 8·3MeCN as red crystals, yield 0.86 g, 44%, and a smaller amount of green crystals of 9, yield *ca.* 5%. Elemental analysis calculated for $\text{C}_{62}\text{H}_{65}\text{Br}_4\text{Co}_2\text{N}_{11}\text{O}_4$: C 49.55, H 4.00, N 6.67% Found for 8 C, 49.35, H 3.96, N 6.61%. IR (KBr, cm^{-1}): 3172 (m), 2956 (s), 2923 (s), 2853 (s),



2726 (w), 1619 (m), 1595 (m), 1536 (m), 1461 (s), 1377 (s), 1260 (s), 1155 (w), 1096 (s), 1060 (s), 1022 (s), 892 (w), 874 (w), 800 (s), 722 (s). M.S. (ESI): 933 (M – 7MeCN – 3Br), 817 (M – 7MeCN – 3Br – 2Co). Magnetic moment: 6.94 B.M.¹⁸

IR (KBr, cm⁻¹) for Complex **9**: 2922 (s), 2854 (s), 2727 (w), 2308 (m), 2282 (m), 2247 (m), 1658 (s), 1635 (s), 1580 (w), 1531 (s), 1464 (s), 1416 (s), 1377 (s), 1367 (s), 1354 (s), 1327 (w), 1287 (m), 1255 (m), 1244 (w), 1222 (m), 1149 (w), 1130 (w), 1098 (w), 1040 (m), 1014 (s), 930 (w), 917 (w), 846 (s), 801 (m), 772 (m), 757 (m), 727 (s), 627 (w), 550 (s), 531 (m), 430 (m).

Synthesis of {CoBr(NCMe)L⁵}₂·2MeCN (10·2MeCN)

To L⁵H (1.00 g, 4.85 mmol) and [CoBr₂] (1.05 g, 4.85 mmol) was added toluene (20 mL) and Et₃N (0.3 mL, 0.22 mmol) and the system was refluxed for 12 h. Following removal of volatiles *in-vacuo*, the residue was extracted into MeCN (20 mL), and on standing at ambient temperature large orange/brown prisms of **10**·3MeCN formed. Isolated yield: 1.41 g, 68%. Elemental analysis calculated for C₃₂H₃₈Br₂Co₂N₄O₆ (sample dried *in-vacuo* for 2 h): C 43.66, H 4.19, N 3.64. Found C 43.81, H 4.23, N 3.67%. IR (KBr, cm⁻¹): 2957 (s), 2924 (s), 2360 (s), 2342 (m), 2312 (m), 2284 (s), 1696 (m), 1641 (s), 1566 (m), 1535 (s), 1464 (s), 1402 (w), 1377 (s), 1367 (s), 1352 (w), 1259 (s), 1225 (m), 1093 (s), 1038 (s), 939 (m), 911 (m), 867 (w), 845 (m), 800 (s), 768 (s), 729 (s), 668 (m), 620 (m), 556 (s), 532 (m), 435 (m). M.S. (ESI): 733 (M – MeCN), 585 (M – 3MeCN – Co), 573 (M – MeCN – 2Br). Magnetic moment: 6.35 B.M.¹⁸

Synthesis of [Co(NCMe)₂L⁵]₂[CoBr₃(NCMe)]₂ (11)

To L⁵H (1.00 g, 4.85 mmol) and [CoBr₂] (1.05 g, 4.85 mmol) was added toluene (20 mL) and the system was refluxed for 12 h. Following removal of volatiles *in-vacuo*, the residue was extracted into MeCN (20 mL), and on standing at ambient temperature large green prisms of **11** formed. Isolated yield: 1.53 g, 46%. Elemental analysis calculated for C₃₆H₄₄Co₄N₆O₆ (sample dried *in-vacuo* for 2 h): C 47.43, H 4.73, N 6.91%. Found C 47.82, H 4.79, N 6.98%. IR (KBr, cm⁻¹): 2926 (s), 2854 (s), 2360 (m), 2342 (w), 2312 (s), 2284 (m), 1641 (s), 1621 (s), 1566 (m), 1535 (s), 1464 (s), 1402 (w), 1377 (s), 1367 (s), 1352 (w), 1259 (s), 1225 (m), 1093 (s), 1038 (s), 1019 (s), 845 (s), 800 (s), 768 (m), 758 (m), 729 (s), 668 (m), 620 (m), 556 (s), 532 (s), 435 (s). M.S. (ESI): 412 (M – 4MeCN – 2Co). Magnetic moment: 6.69 B.M.¹⁸

Synthesis of [FeBr(NCMe)L⁵]₂·2MeCN (12·2MeCN)

To L⁵H (1.00 g, 4.85 mmol) and [FeBr₂] (1.06 g, 4.85 mmol) was added toluene (20 mL) and Et₃N (0.7 mL, 0.485 mmol) and the system was refluxed for 12 h. Following removal of volatiles *in-vacuo*, the residue was extracted into MeCN (20 mL), and on standing at ambient temperature large orange/brown prisms of **10**·3MeCN formed. Isolated yield: 0.837 g, 40%. Elemental analysis calculated for C₂₈H₃₂Br₂Fe₂N₂O₆ (sample dried *in-vacuo* for 2 h): C 44.02, H 4.22, N 3.67. Found C 44.61, H 4.52, N 3.72%. IR (KBr, cm⁻¹): 3352 (w), 2954 (s), 2923 (s), 2854 (s), 2727 (w), 2360 (w), 2341 (w), 1643 (m), 1620 (m), 1528 (m), 1462 (s), 1377 (s), 1259 (m), 1160 (m), 1037 (s), 1016 (m), 842

(w), 800 (s), 769 (w), 755 (w), 722 (m), 668 (w), 618 (m), 539 (m). M.S. (ESI): 519 (M – 4MeCN – 2Br). Magnetic moment: 6.70 B.M.¹⁸

Synthesis of [Fe(NCMe)₆][FeOBr₃]₂ (13)

L¹H₂ (0.52 g, 0.77 mmol) and [FeBr₃] (0.47 g, 1.58 mmol) were combined in a Schlenk and toluene (20 mL) was added. After refluxing for 12 h, the volatiles were removed *in-vacuo*, and the residue was extracted into MeCN (20 mL). Prolonged standing at room temperature afforded brown prisms; isolated yield 0.39 g 81%. Elemental analysis calculated for C₁₂H₁₈Br₆Fe₃N₆O (sample dried *in-vacuo* for 2 h): C 15.85, H 2.00, N 9.24%. Found C, 15.61 H, 2.12 N 9.31%. IR (KBr, cm⁻¹): 2925 (s), 2854 (s), 2361 (s), 2339 (s), 1868 (s), 1844 (s), 1830 (m), 1792 (m), 1772 (w), 1750 (m), 1734 (m), 1717 (m), 1700 (m), 1684 (m), 1669 (m), 1653 (m), 1646 (m), 1635 (m), 1616 (s), 1576 (s), 1559 (s), 1540 (s), 1521 (s), 1506 (s), 1497 (m), 1489 (m), 1457 (s), 1419 (m), 1377 (s), 1260 (s), 1089 (s), 800 (s), 668 (s).

ROP procedure

ε-Caprolactone and δ-valerolactone. Typical polymerisation procedure in the presence of one equivalent of benzyl alcohol (Table 1, run 1) is as follows. A toluene solution of **1** (0.010 mmol, in 1.0 mL toluene) and BnOH (0.010 mmol) were added into a Schlenk tube in the glove-box at room temperature. The solution was stirred for 2 min, and then ε-caprolactone (2.5 mmol) or δ-valerolactone along with 1.5 mL toluene was added to the solution. The reaction mixture was then placed into an oil bath pre-heated to the required temperature, and the solution was stirred for the prescribed time. The polymerisation mixture was then quenched by addition of an excess of glacial acetic acid (0.2 mL) into the solution, and the resultant solution was then poured into cold methanol (200 mL). The resultant polymer was then collected on filter paper and was dried *in vacuo*.

Co-polymerisations

A toluene solution of catalyst (0.010 mmol, in 1.0 mL toluene) was added into a Schlenk tube in the glove-box at room temperature. The solution was stirred for 2 min, and then 1 equivalent of BnOH (from 1 mmol BnOH in 100 mL toluene) and monomer were added, after 24 h the other monomer was added to the solution. The reaction mixture was then placed into an oil bath pre-heated to the 130 °C, and the solution was stirred for another 24 h. The polymerisation mixture was then quenched by addition of an excess of glacial acetic acid (0.2 mL) into the solution, and the resultant solution was then poured into cold methanol (200 mL). The resultant polymer was then collected on filter paper and was dried *in vacuo*.

Kinetic studies

The polymerisations were carried out at 130 °C in toluene (1 mL) using 0.010 mmol of complex. The molar ratio of monomer to initiator was fixed at 500:1, and at appropriate time intervals, 0.5 μL aliquots were removed (under N₂) and





Table 4 Crystallographic data

Compound	1:2MeCN	2:0.5MeCN	3:5.5MeCN	4:3MeCN
Formula	C ₉₆ H ₈₆ Br ₈ Fe ₄ N ₁₀ O ₅	C ₅₂ H ₅₂ BrFeN ₄ O ₂ , C ₂ H ₃ N, 0.5(Br ₆ Fe ₂ O)	C ₅₂ H ₅₂ BrFeN ₄ O ₂ , Br, 5.5(C ₃ H ₃ N)	C ₁₀₄ H ₁₀₁ FeN ₈ O ₄ , (3MeCN)
Formula weight	2322.42	1224.841185.89	1185.89	1705.9
Crystal system	Monoclinic	Triclinic	Monoclinic	Monoclinic
Space group	<i>P</i> ₂ ₁ / <i>n</i>	<i>P</i> $\bar{1}$	<i>P</i> ₂ ₁ / <i>n</i>	<i>P</i> ₂ ₁ / <i>c</i>
Unit cell dimensions				
<i>a</i> (Å)	15.6606(4)	13.6454(5)	14.2077(6)	12.6268(5)
<i>b</i> (Å)	16.0326(4)	13.9070(7)	27.6241(9)	30.3249(10)
<i>c</i> (Å)	37.4062(11)	15.6432(8)	15.1579(8)	24.6577(10)
α (°)	90	67.263(5)	90	90
β (°)	99.896(3)	88.642(4)	112.484(6)	100.817(3)
γ (°)	90	81.638(4)	90	90
<i>V</i> (Å ³)	9252.2(4)	2707.0(2)	5496.9(5)	9273.8(6)
<i>Z</i>	4	1	4	4
Temperature (K)	100(2)	100(2)	100(2)	140(1)
Wavelength (Å)	1.54184	0.71075	0.71073	0.71073
Calculated density (g cm ⁻³)	1.667	1.503	1.185	1.222
Absorption coefficient (mm ⁻¹)	9.439	3.531	1.766	0.223
<i>T</i> _{min} , <i>T</i> _{max}	0.755, 1.000	0.592, 1.000	0.555, 1.000	0.749, 1.185
Crystal size (mm ³)	0.08 × 0.05 × 0.02	0.30 × 0.015 × 0.01	0.26 × 0.045 × 0.01	0.74 × 0.18 × 0.16
θ (max) (°)	68.2	27.5	27.5	20.0
Reflections measured	68 070	12 337	47 911	51 064
Unique reflections	16 805	12 337	12 466	8583
<i>R</i> _{int}	0.107	0.067	0.056	0.143
Number of parameters	1127	604	556	1129
<i>R</i> ₁ [<i>F</i> ² > 2σ(<i>F</i> ²)]	0.070	0.061	0.088	0.057
w <i>R</i> ₂ (all data)	0.178	0.164	0.168	0.123
GOOF, <i>S</i>	1.011	1.049	1.050	0.913
Largest difference peak and hole (e Å ⁻³)	1.42 and −0.86	1.53 and −0.80	0.90 and −1.61	0.30 and −0.25

Compound	5:2MeCN	6:2MeCN	7:2C ₃ H ₆ O	8:3MeCN
Formula	C ₅₆ H ₅₈ Br ₃ Cu ₂ N ₆ O ₂	C ₅₂ H ₅₄ Br ₄ CoN ₄ O ₂ ·2(C ₃ H ₃ N)	C ₄₂ H ₃₀ Br ₂ Co ₂ N ₄ O ₄ ·2(C ₃ H ₆ O)	C ₆₂ H ₆₅ Br ₄ Co ₂ N ₁₁ O ₄
Formula weight	1213.89	1145.56	1048.56	1465.8
Crystal system	Triclinic	Triclinic	Monoclinic	Monoclinic
Space group	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$	<i>P</i> ₂ ₁ / <i>c</i>	<i>Ia</i>
Unit cell dimensions				
<i>a</i> (Å)	10.0808(2)	12.2629(5)	11.4774(8)	15.77810(10)
<i>b</i> (Å)	15.0015(4)	13.2489(6)	14.5787(10)	25.0018(2)
<i>c</i> (Å)	18.6604(5)	17.0384(7)	26.021(3)	16.28380(10)
α (°)	71.442(2)	96.850(3)	90	90
β (°)	82.738(2)	91.115(3)	100.235(8)	91.911(10)
γ (°)	76.600(2)	99.672(4)	90	90
<i>V</i> (Å ³)	2598.2(1)	2707.2(2)	4284.6(6)	6420.08(8)
<i>Z</i>	2	2	4	4
Temperature (K)	100(2)	100(2)	100(2)	100(2)
Wavelength (Å)	1.54184	0.71075	1.54178	0.71073
Calculated density (g cm ⁻³)	1.552	1.405	1.535	1.516
Absorption coefficient (mm ⁻¹)	4.572	3.309	8.652	3.060
<i>T</i> _{min} , <i>T</i> _{max}	0.149, 1.000	0.652, 1.000	0.404, 1.000	0.698, 1.000
Crystal size (mm ³)	0.70 × 0.60 × 0.15	0.10 × 0.04 × 0.02	0.14 × 0.03 × 0.01	0.25 × 0.18 × 0.05
θ (max) (°)	68.3	21.7	68.2	28.7



Table 4 (Contd.)

Compound	5-2MeCN	6-2MeCN	7-2C ₃ H ₆ O	8-3MeCN
Reflections measured				
Unique reflections				
R_{int}	44 092	10 619	6241	147 825
Number of parameters	9337	10 619	3471	16 480
$R_1 [F^2 > 2\sigma(F^2)]$	0.083	0.179	0.054	0.047
wR_2 (all data)	622	564	526	762
GOOF, S	0.066	0.114	0.128	0.032
Largest difference peak and hole (e Å ⁻³)	0.184	0.297	0.371	0.064
	1.062	1.098	1.226	1.023
	3.45 and -1.24	1.02 and -1.54	2.52 and -2.02	0.52 and -0.46
Compound	9	10-2MeCN	11-2MeCN	12-2MeCN
Formula	C ₂₀ H ₃₀ Br ₆ Co ₃ N ₁₀	C ₂₈ H ₃₂ Br ₂ Co ₂ N ₂ O ₆ ·2(C ₂ H ₃ N)	C ₃₆ H ₄₄ Br ₆ Co ₄ N ₆ O ₆	C ₃₂ H ₃₈ Br ₂ Fe ₂ N ₄ O ₆
Formula weight	1066.79	852.34	1371.95	846.18
Crystal system	Monoclinic	Triclinic	Triclinic	Triclinic
Space group	$P2_1/n$	$P\bar{1}$	$P\bar{1}$	$P\bar{1}$
Unit cell dimensions				
a (Å)	8.34720(10)	8.4718(5)	10.416(2)	8.51990(10)
b (Å)	18.8457(3)	9.2109(3)	11.156(4)	9.20980(10)
c (Å)	12.5811(2)	12.2495(5)	11.738(4)	12.1881(2)
α (°)	90	101.186(3)	78.04(3)	101.2110(10)
β (°)	108.941(2)	102.715(4)	70.59(2)	102.4660(10)
γ (°)	90	97.208(4)	87.78(2)	97.4090(10)
V (Å ³)	1871.95(5)	900.47(7)	1257.8(7)	901.29(2)
Z	2	1	1	6
Temperature (K)	100(2)	150(2)	100(2)	100(2)
Wavelength (Å)	0.71075	1.54184	0.71073	1.54184
Calculated density (g cm ⁻³)	1.893	1.572	1.811	2.230
Absorption coefficient (mm ⁻¹)	7.743	10.197	6.105	23.225
$T_{\text{min}}, T_{\text{max}}$	0.154, 0.733	0.423, 1.000	0.725, 0.820	0.646, 1.000
Crystal size (mm ³)	0.216 × 0.125 × 0.10	0.32 × 0.16 × 0.10	0.31 × 0.13 × 0.09	0.30 × 0.20 × 0.10
θ (max) (°)	27.49	68.2	29.2	70.3
Reflections measured	48 302	15 481	11 602	32 046
Unique reflections	4301	3240	6632	3387
R_{int}	0.0235	0.066	0.044	0.0342
Number of parameters	183	217	268	217
$R_1 [F^2 > 2\sigma(F^2)]$	0.017	0.042	0.035	0.0299
wR_2 (all data)	0.0037	0.119	0.072	0.0822
GOOF, S	1.170	1.072	0.823	1.091
Largest difference peak and hole (e Å ⁻³)	0.79 and -0.35	0.85 and -0.83	0.84 and -0.63	0.413 and -0.69
				0.38 and -0.51

were quenched with wet CDCl_3 . The percent conversion of monomer to polymer was determined by ^1H NMR spectroscopy.

Mass spectrometry

Polymer samples were run at the University of Hull using MALDI-TOF MS analysis. Samples were dissolved in THF, and the matrix, 2-(4-hydroxyphenylazo) benzoic acid (HPABA) with added NaOAc was employed, which was dissolved in THF to give a saturated solution. 50 μL of the sample solution was then mixed with 50 μL matrix solution, and 1 μL of the mixed solution applied to the sample target. The sample was allowed to dry in air before analysis. The co-poly (δ -VL + r -LA) (run 7) sample was analysed by MALDI in positive-linear and reflection modes, with DCTB matrix and NaOAc additive. The sample was fully soluble in THF at 10 mg mL^{-1} .

X-ray crystallography

Single-crystal X-ray diffraction data for the structures (except for **4** and **11**) were collected at the UK National Crystallography service on a range of Rigaku Oxford Diffraction ultra-high intensity instruments employing modern areas detectors. Samples were held at 100 K for data collection. For **4**, data were from a Rigaku Oxford Diffraction Xcalibur-3 CCD diffractometer at 140 K. For **11**, data were from a Stoe IPDS2 image plate diffractometer utilising monochromated Mo radiation ($\lambda = 0.71073 \text{ \AA}$). In all cases standard procedures were employed for integration and processing of data. Crystallographic data for all samples are collated in Table 4.

Crystal structures were solved using direct methods (in SHELXS^{19a}) or dual space methods implemented within SHELXT.^{19b} Completion of structures was achieved by performing least squares refinement against all unique F^2 values using SHELXL-2018.²⁰ All non-H atoms were refined with anisotropic displacement parameters. Hydrogen atoms were placed using a riding model. Where the location of hydrogen atoms was obvious from difference Fourier maps, C–H and O–H bond lengths were refined subject to chemically sensible restraints. Minor disorder was treated using standard methods.

The crystal of **5** examined was the best of those available. The crystal was split and intensity data from the major component were used for refinement. The final refinement has good quality of fit ($R_1 = 6.66\%$) but it is not perfect because of the imperfect nature of the crystals available.

Conflicts of interest

There are no conflicts of interest to declare.

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