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Amplification of weak chiral inductions for excellent control over the helical orientation of discrete topologically chiral $(M_3L_2)_n$ polyhedra†

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Superb control over the helical chirality of discrete $(M_3L_2)_n$ polyhedra $(n=2,4,8, M=Cu^l)$ or Ag^l) created from the self-assembly of propeller-shaped ligands (L) equipped with chiral side chains is demonstrated here. Almost perfect chiral induction (>99:1) of the helical orientation of the framework was achieved for the largest $(M_3L_2)_8$ cube with 48 small chiral side chains (diameter: \sim 5 nm), while no or moderate chiral induction was observed for smaller polyhedra (n=2, 4). Thus, amplification of the weak chiral inductions of each ligand unit is an efficient way to control the chirality of large discrete nanostructures with high structural complexity.

Introduction

Amplification of chiral induction has previously been observed in double asymmetric synthesis, where high enantioselectivity or diastereoselectivity arises from two or more chiral sources (e.g., a chiral substrate and a chiral reagent). With increasing multiplication number (m), the selectivity should be exponentially enhanced because the small free energy differences ($\Delta E_{\rm m}$) that arise from each chiral source accumulate. This process roughly follows the Gibbs free energy equation, $\Delta G = \Sigma \Delta E_{\rm m} =$ $-RT \ln K$, where K is the reaction selectivity or the equilibrium constant of the two states. Exceptional amplification effects have often been observed in polymeric systems, particularly in the formation of helices in long-chain polymers, where the helical sense of the polymer main chain can be perfectly controlled, even when the chiral induction at every monomer unit is small.2-4 In discrete molecular systems, however, there are only a limited number of examples where large amplification effects have been observed,5,6 given that the incorporation of a large number of chiral sources into discrete molecular structures is usually not straightforward. In particular, chirality control over huge and complex discrete nanoarchitectures is believed to open up new concepts in the fields of material

We have recently reported a number of $(M_3L_2)_n$ polyhedra $(2a-4a; n = 2, 4, 8; M = Cu^{I} \text{ or } Ag^{I}; \text{ Fig. } 1)^{8} \text{ constructed from }$ ligand 1a through conventional metal-pyridyl coordination9 and secondary metal-acetylene π-coordination. The propellershaped tripodal ligand 1a can exist in two enantiomeric conformations (Fig. 1a) and, once coordinated to M, oligomerizes to form polyhedral frameworks with diameters of up to ca. 5 nm (Fig. 1b). Crystallographic and spectroscopic studies have shown that the helical sense of all the 1a ligands in the polyhedral frameworks are identical, although the polyhedra exist as racemates. We observed that the $(M_3L_2)_n$ polyhedra have many identical ligand arms directed to outer environments, and thus, substantial chiral induction in the polyhedral frameworks retaining cavity structures could be expected by simply introducing a chiral element to the ligand arm. In addition, note that a series of the $(M_3L_2)_n$ polyhedra contain highly entangled structures based on trifurcate motifs, 11-13 exceeding the well classified topologies of conventional knots and links.14 As these structures are inherently topologically chiral,15 it is expected to be interesting to demonstrate their precise regulation for invoking future applications of the newly exploited threedimensional molecular entanglements.

Therefore, in this study, we designed and synthesized ligands **1b-e** (Fig. 1a), which are derivatives of **1a** equipped with a chiral substituent on every arm. We found that, while chiral induction from the side chains is negligible or very small for polyhedra **2** (n = 2) and **3** (n = 4), perfect control over the helicity can be achieved in cubic octamer **4** (n = 8) via the remarkable amplification effect of, in total, 48 chiral substituents on the ligand arms. This result demonstrates that the chirality of

science including artificial reaction systems, sensors, and biochemical applications in higher degree of multicomponent manners.⁷

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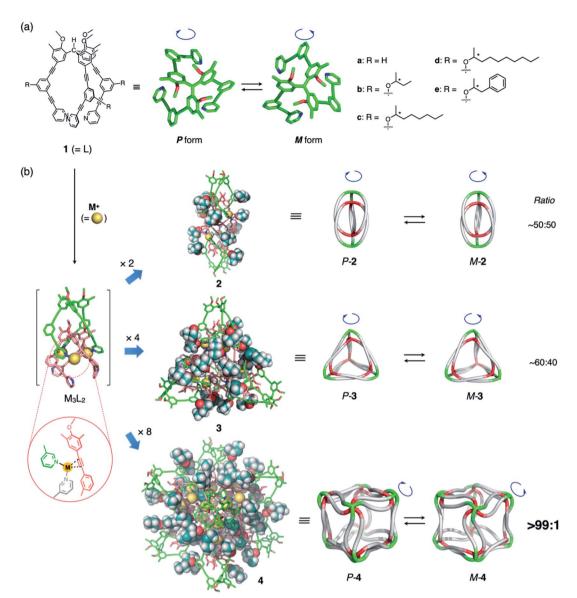


Fig. 1 (a) Chemical structure of propeller-shaped ligand 1 and its P- and M-forms. Unless otherwise noted, the (S)-2-alkoxy isomers of 1b-e were used in the main study. (b) Schematic representation of the self-assembly of the $(M_3L_2)_n$ coordination polyhedra 2-4 (n=2,4,8) with chiral side chains. Each assembly is formed by the oligomerization of n units of M_3L_2 , a hypothetical intermediate stabilized by metal-acetylene π -coordination. Illustrations in the second column from the left were generated by combining 2-butoxy side chains (molecular-mechanics (MM) models) with polyhedral frameworks based on the reported crystal structures. Base Cartoon illustrations in the third column exhibit the entangled structures of the $(M_3L_2)_n$ polyhedra, while metal-acetylene π -coordination is omitted for clarity.

discrete nanostructures assembled from numerous components can be easily controlled by the amplification effect of weakly chiral elements present on every component.

Results and discussion

Propeller-shaped tripodal ligands **1b–e**, which bear chiral (S)-alkoxy side chains on each arm, were designed and prepared according to literature procedures (for details, see the ESI†).^{8 α ,1 ϵ} Firstly, dimeric (M_3L_2)₂ cages 2 were synthesized by mixing [Cu(MeCN)₄]BF₄ (1.5 equiv., 3.75 mM) with ligand **1b** in CD₃NO₂. The ¹H NMR spectra showed two sets of proton signals attributable to dimer **2b** (Fig. 2b and S1†), suggesting the

formation of two diastereomeric cages with the ligands in either the M- or P-configuration. However, the diastereomeric ratio was $\sim 1:1$ in both cases even though a chiral substituent is present on each of the 12 side chains. A lack of chiral induction in the $(M_3L_2)_2$ cage frameworks was also confirmed by the circular dichroism (CD) spectra, which showed no chiral amplification in comparison with the CD spectra of the ligand solutions (Fig. S2 \dagger). This absence of chiral induction is presumably due to insufficient steric interactions between the chiral side chains and the cage framework of 2b. In fact, the single-crystal X-ray structure of the cage framework of 2b (Fig. 2d) revealed almost no steric crowding between the side chains and the cage framework. We thus examined ligand 1e

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Fig. 2 Self-assembly of complex 2b and tetrahedron 3b. Partial 1 H NMR spectra of (a) ligand 1b (500 MHz, 300 K, CDCl₃), (b) complex 2b (700 MHz, 300 K, CD $_3$ NO $_2$), and (c) tetrahedron 3b (500 MHz, 300 K, toluene- d_8). Red circles and blue triangles indicate two diastereomeric complexes, respectively. A 1 H DOSY spectrum is shown in addition (for details, see Fig. S5†). (d) X-ray crystal structure of 2b; 2-butoxy chains are shown as space-filling models.

with a sterically more crowded (*S*)-1-phenyl-2-propanoxy group as the chiral substituent (for details, see the ESI†). When cage **2e** was assembled from ligand **1e**, a small amount of diastereoselectivity was observed (62:38 by ¹H NMR analysis; Fig. S3†), which can be interpreted in terms of increased steric crowding of the bulky groups on the cage (MM model based on partially refined X-ray data; *cf.* Fig. S4†).

The small chiral induction effect observed for cage 2e encouraged us to examine higher-order oligomeric cages, in which a larger number of chiral side chains are present. The self-assembly of tetrahedral $(M_3L_2)_4$ 3 was thus examined with ligands 1b. After complexation with CuBF₄ (1.5 equiv., 3.75 mM; 50 °C for 30 min) in toluene- d_8 , the formation of 3b was confirmed by 1 H and 13 C NMR (Fig. 2c and S27–S28†), in addition to 1 H diffusion ordered spectroscopy (DOSY) showing a hydrodynamic diameter of \sim 4.7 nm (log D=-9.8 at 300 K) comparable to that was determined for 3a by X-ray analysis (4.5 nm). $^{8\alpha}$ Here, however, only small diastereomeric ratio (52 : 48) was observed by 1 H NMR (Fig. 2c), which is consistent with the moderately enhanced absorption observed in the CD spectra (Fig. S6†). 17

The helical orientation of ligand 1 forces the generation of a chiral trefoil knot topology¹⁴ on every face of the tetrahedral cage 3 (Fig. 2c, right). In this situation, the two enantiomeric

forms cannot be interconverted unless the trefoil knot topology is dissociated. Thus, the P- and M-helical orientations of the ligand do not interconvert, except under very slow equilibration via bond dissociation. Therefore, as shown by variable-temperature (VT) 1 H NMR experiments (Fig. S7†), the diastereomeric ratio of 3 is not affected by a change in temperature between 273 and 323 K. 18

We expected to see a further enhancement in the chiral induction in the $(M_3L_2)_n$ assembly of cube-structured 4 (n=8). As previously reported, ^{8c} cube 4 can only be constructed via the anion-templated conversion of pre-synthesized dimeric cages of 2. Thus, cage 2d was prepared by mixing ligand 1d (with an (S)-2-decanoxy side chain, 2.5 mM) and AgPF₆ (1.9 equiv.) in CDCl₃/CD₃OD (97/3, v/v) at 50 °C for 30 min. Finally, the solution was treated with tetramethylammonium nitrate (TMANO₃, 1.0 equiv. in CD₃OD at room temperature over 21 h; CDCl₃/CD₃OD ratio after the template addition was 94/6, v/v) (Fig. 3a). ¹⁹

To our surprise, we observed the exclusive formation of only one diastereomer of cube **4d** (>99:1) *via* ¹H NMR spectroscopy (Fig. 3b and S11a†). Time-dependent NMR measurements

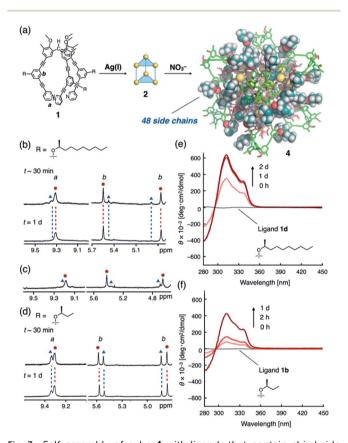


Fig. 3 Self-assembly of cube 4 with ligands that contain chiral side chains. (a) Schematic representation of the nitrate-templated self-assembly of 4 (depicted with R=2-butoxy for simplicity); partial $^1\mathrm{H}$ NMR spectra (500 MHz, 287 K, CDCl $_3$ /CD $_3$ OD (94/6, v/v)) of ligand 1d and cube 4 showing (b) time-dependent redistribution of the diastereomers, (c) cube 4 with ligand 1d in CDCl $_3$ /CD $_3$ OD (97/3, v/v, t=18 h), and (d) cube 4 with ligand 1b in CDCl $_3$ /CD $_3$ OD (97/3, v/v). (e) Changes in the CD (293 K, CHCl $_3$ /MeOH (94/6, v/v)) spectrum upon formation of cube 4d. (f) Changes in the CD (293 K, CHCl $_3$ /MeOH (97/3, v/v)) spectrum upon formation of cube 4b.

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revealed that the selectivity is thermodynamically controlled. Immediately following the addition of TMANO₃ (\sim 30 min), we observed a relatively low selectivity (82:18) that increased to >99:1 after stirring the solution for 21 h. This kinetic-tothermodynamic conversion process was also monitored using CD spectroscopy (Fig. 3e). As precursor 2 is a $\sim 1:1$ mixture of diastereomers, the relatively rapid interconversion of 2 to 4 by addition of nitrate initially results in the formation of a kinetic mixture of diastereomers of 4d, which is then converted into the thermodynamic product. Cage 4 was characterized by comparing the ¹H and ¹³C NMR spectra of 4 (Fig. S12†)²⁰ with those of the previously reported 4a.8c 1H DOSY measurements indicated the formation of a single species with a diffusion coefficient of $D = 1.6 \times 10^{-10} \text{ m}^2 \text{ s}^{-1} (\log D = -9.8 \text{ at } 287 \text{ K};$

The use of high polarity solvents (CDCl₃/CD₃OD (94/6, v/v)) is particularly important to achieve high selectivity. In fact, when the same experiment was conducted in a slightly less polar solvent (CDCl₃/CD₃OD (97/3, v/v)), the selectivity decreased significantly (max.: 85: 15; Fig. 3c and S11b†). Presumably, the steric contact of long hydrophobic alkyl chains with the framework of cube 4d becomes more efficient in polar solvents, thus enhancing the chiral induction, as observed in helical polymers.4

To clarify the secondary steric effects, ligands 1c and 1b with shorter (S)-2-alkoxy side chains were used in the self-assembly of cube 4. In CDCl₃/CD₃OD (97/3, v/v), a ligand with a 2-octoxy chain (1c) offered comparable selectivity (84:16; Fig. S15-S16†) to that of 1d (85:15). In the case of 1b, *i.e.*, a ligand with a much shorter 2-butoxy chain, much lower selectivity (70:30) was observed even after 18 h of treatment with tetrabutylammonium nitrate (TBANO₃) (Fig. 3d and f).

The P and M equilibrium constant (K) can be described by the Gibbs free energy equation, $K = \exp(-\Delta G/RT)$, where ΔG may roughly be described as $m \cdot E_h$ (E_h : energy difference per chiral unit). 4a,b Even though E_h is very small, the accumulation of 48 chiral units (m = 48) in 4d is enough to induce a relatively large ΔG that accounts for the perfect chiral induction. In other words, the diastereomeric ratio of 99 : 1 for cube 4 (n = 8, m =48) can be ascribed to an energy difference of only -0.23 kJ mol⁻¹ per chiral group. Whilst the absolute value is small, the accumulation of 48 chiral groups on the surface of the polyhedron can generate a large ΔG that perfectly controls the chirality of the huge polyhedral framework.

Conclusions

In conclusion, we have successfully controlled the helicity of discrete $(M_3L_2)_n$ polyhedra by adding small chiral side chains to their highly entangled higher-order frameworks. The results shown here are inspiring as they allow the precise structural regulation of large, complex nanoarchitectures, that are usually difficult to synthesize via the self-assembly of numerous components. This strategy can be expected to stimulate new approaches toward advanced functional nanomaterials¹⁵ based on discrete molecular designs.

Data availability

All the data are shown in the ESI.†

Author contributions

Y. D. and M. F. conceived and designed the study. Y. D. and M. F. co-wrote the paper. Y. D., K. Y., S. H., and Z. Y. performed the chemical experiments and analysed the data. All authors discussed the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

‡ Crystallographic data for 2b: M = 20583.18, monoclinic, space group $P2_1/n$, a =18.838(4) Å, b = 32.458(6) Å, c = 64.439(13) Å, $\alpha = \gamma = 90^{\circ}$, $\beta = 94.55(3)^{\circ}$, $V = 94.55(3)^{\circ}$ $39277(14) \text{ Å}^3$, Z = 1, $D_c = 0.870 \text{ g cm}^{-3}$, T = 100(2) K, $1.784 < \theta < 47.56^{\circ}$, 20892unique reflections out of 69369 with $I > 2\sigma(I)$, GoF = 1.290, final R factors R_1 = $0.1440 (I > 2\sigma(I)), wR_2 = 0.3746, CCDC$ deposition number 2108704.

- 1 S. Masamune, W. Choy, J. S. Petersen and L. R. Sita, Angew. Chem., Int. Ed., 1985, 24, 1-30.
- 2 Y. Okamoto, H. Mohri, T. Nakano and K. Hatada, J. Am. Chem. Soc., 1989, 111, 5952-5954.
- 3 (a) M. M. Green, C. Andreola, B. Muñoz and M. P. Reidy, J. Am. Chem. Soc., 1988, 110, 4063-4065; (b) M. M. Green, N. C. Peterson, T. Sato, A. Teramoto, R. Cook and S. Lifson, Science, 1995, 268, 1860-1866; (c) H. Nakako, R. Nomura and T. Masuda, Macromolecules, 2001, 34, 1496-1502.
- 4 (a) T. Yamada, Y. Nagata and M. Suginome, Chem. Commun., 2010, **46**, 4914–4916; (b) Y. Nagata, T. Yamada, T. Adachi, Y. Akai, T. Yamamoto and M. Suginome, J. Am. Chem. Soc., 2013, **135**, 10104–10113; (c) Y. Nagata, T. Nishikawa, M. Suginome, S. Sato, M. Sugiyama, L. Porcar, A. Martel, R. Inoue and N. Sato, J. Am. Chem. Soc., 2018, 140, 2722-2726.
- 5 (a) N. Ousaka, J. K. Clegg and J. R. Nitschke, Angew. Chem., Int. Ed., 2012, 51, 1464-1468; (b) F. Huang, L. Ma, Y. Che,

- H. Jiang, X. Chen and Y. Wang, *J. Org. Chem.*, 2018, **83**, 733–739; (*c*) J. Zhang, D. Luo, C. Ma, L. Huang and Q. Gan, *Nat. Commun.*, 2021, **12**, 2659.
- 6 (a) K. S. Jeong, Y. S. Kim, Y. J. Kim, E. Lee, J. H. Yoon, W. H. Park, Y. W. Park, S.-J. Jeon, Z. H. Kim, J. Kim and N. Jeong, Angew. Chem., Int. Ed., 2006, 45, 8134–8138; (b) C. Gütz, R. Hovorka, C. Klein, Q.-Q. Jiang, C. Bannwarth, M. Engeser, C. Schmuck, W. Assenmacher, W. Mader, F. Topić, K. Rissanen, S. Grimme and A. Lützen, Angew. Chem., Int. Ed., 2014, 53, 1693–1698; (c) K.-R. Wang, D.-S. Guo, B.-P. Jiang and Y. Liu, Chem. Commun., 2012, 48, 3644–3646; (d) M. Gaeta, G. Sortino, R. Randazzo, I. Pisagatti, A. Notti, M. E. Fragalà, M. F. Parisi, A. D'Urso and R. Purrello, Chem.-Eur. J., 2020, 26, 3515–3518.
- 7 (a) L.-J. Chen, H.-B. Yang and M. Shionoya, *Chem. Soc. Rev.*, 2017, 46, 2555–2576; (b) M. Pan, K. Wu, J.-H. Zhang and C.-Y. Su, *Coord. Chem. Rev.*, 2019, 378, 333–349.
- 8 (a) Y. Domoto, M. Abe, T. Kikuchi and M. Fujita, Angew. Chem., Int. Ed., 2020, 59, 3450-3454; (b) Y. Domoto, M. Abe, K. Yamamoto, T. Kikuchi and M. Fujita, Chem. Sci., 2020, 11, 10457-10460; (c) Y. Domoto, M. Abe and M. Fujita, J. Am. Chem. Soc., 2021, 143, 8578-8582.
- (a) M. M. Conn and J. Rebek, Chem. Rev., 1997, 97, 1647–1668; (b) D. L. Caulder and K. N. Raymond, Acc. Chem. Res., 1999, 32, 975–982; (c) R. W. Saalfrank, H. Maid and A. Schuerer, Angew. Chem., Int. Ed., 2008, 47, 8794–8824; (d) M. M. Smulders, I. A. Riddell, C. Brown and J. R. Nitschke, Chem. Soc. Rev., 2013, 42, 1728–1754; (e) T. R. Cook and J. P. Stang, Chem. Rev., 2015, 115, 7001–7045; (f) H. Hou, K. Zhou, F. Jiang, Q. Chen and M. Hong, Isr. J. Chem., 2019, 59, 140–150; (g) M. Han, D. M. Engelhard and G. H. Clever, Chem. Soc. Rev., 2014, 43, 1848–1860.
- 10 (a) H. Lang, A. Jakob and B. Milde, *Organometallics*, 2012, 31, 7661–7693; (b) H. Schmidbaur and A. Schier, *Organometallics*, 2010, 29, 2–23.
- 11 (a) M. O'Keeffe and M. M. J. Treacy, *Acta Cryst. A*, 2020, 76, 611–621; (b) T. Castle, M. E. Evans and S. T. Hyde, *New J. Chem.*, 2008, 32, 1484–1492; (c) S. T. Hyde and M. E. Evans, *Proc. Natl. Acad. Sci. U. S. A.*, 2022, 119, e2110345118.

- 12 T. Sawada and M. Fujita, *Bull. Chem. Soc. Jpn.*, 2021, **94**, 2342–2350.
- 13 X. Zhao, H. Wang, B. Li, W. Zhang, X. Li, W. Zhao, C. Janiak, A. W. Heard, X.-J. Yang and B. Wu, *Angew. Chem., Int. Ed.*, 2022, **61**, e202115042.
- 14 (a) E. E. Fenlon, Eur. J. Org. Chem., 2008, 5023-5035; (b)
 K. E. Horner, M. A. Miller, J. W. Steed and P. M. Sutcliffe, Chem. Soc. Rev., 2016, 45, 6432-6448; (c) S. D. P. Fielden, D. A. Leigh and S. L. Woltering, Angew. Chem., Int. Ed., 2017, 56, 11166-11194; (d) W.-X. Gao, H.-J. Feng, B.-B. Guo, Y. Lu and G.-X. Jin, Chem. Rev., 2020, 120, 6288-6325.
- 15 (a) N. Pairault and J. Niemeyer, Synlett, 2018, 29, 689–698; (b)
 A. H. G. David and J. F. Stoddart, Isr. J. Chem., 2021, 61, 608–621; (c) J. R. J. Maynard and S. M. Goldup, Chem, 2020, 6, 1914–1932; (d) Y. Sang and M. Liu, Chem. Sci., 2022, 13, 633–656.
- 16 J. Kobayashi, Y. Domoto and T. Kawashima, *Chem. Commun.*, 2009, 6186–6188.
- 17 Even the use of the ligand with longer side chain (1d) did not show the improvement of the diastereoselectivity (60:40) in the case of assembly 3.
- 18 Significant solvent effects on diastereoselectivity were observed also for the self-assembly of the tetrahedral cages 3 (Fig. S8–S10†).
- 19 The sources and equivalents of the silver (AgPF₆ or AgSbF₆, 1.5–2.0 equiv.) and nitrate ions (TMANO₃ or TBANO₃, 0.5–1.5 equiv.) were adjusted to achieve complete convergence of the self-assembly process, which depends on the ligand used.
- 20 Treatment of the solution with a chiral-shift reagent anion (Δ-tris(tetrachlorobenzenediolate)phosphate(v) (Δ-TRISPHAT), 0-0.35 equiv., ref. 21) induced significant shifts of every ¹H NMR signal at 287 K (Fig. S14†), indirectly supporting the assignment as a pair of diastereomers.
- 21 (a) J. J. Jodry and J. Lacour, *Chem.-Eur. J.*, 2000, 6, 4297–4304;
 (b) S. Hiraoka, K. Harano, T. Tanaka, M. Shiro and M. Shionoya, *Angew. Chem., Int. Ed.*, 2003, 42, 5182–5185.