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# The effect of additional methyl on the magnetic relaxation and toroidal moment of Dy<sub>6</sub> complex†

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Two new  $\mu_4$ -O bridged hexanuclear lanthanide complexes, [Dy<sub>6</sub>L'<sub>4</sub>( $\mu_4$ -O)(NO<sub>3</sub>)<sub>4</sub>]·4CH<sub>3</sub>OH (1) and [Sm<sub>6</sub>L'<sub>4</sub>( $\mu_4$ -O)(CH<sub>3</sub>COO)<sub>4</sub>]·2CH<sub>3</sub>OH (2), have been assembled by using methyl modified Schiff-base ligand H<sub>3</sub>L' (2,6-bis((2-hydroxypropylimino)methyl)-4-methylphenol). The complex 1 shares a similar  $\mu_4$ -O bridged Dy<sub>6</sub> core as the parent Dy<sub>6</sub>-2, while coordination geometries and magnetic interactions are slightly changed upon the modulation of the ligand, which results in distinct single-molecule magnetic (SMM) and single-molecule toroic (SMT) properties.

## Introduction

The design and construction of single-molecule magnets (SMMs) has attracted increasing attention since the discovery of SMM behaviour,<sup>1</sup> because they have potential applications in processing and storing magnetic information at a molecular level.<sup>2–7</sup> Thus a large number of SMMs, including 3d complexes, 3d/4f complexes and 4f complexes, have been reported. Thereinto, 4f-based complexes are particularly appealing due to the strong magnetic anisotropy of 4f ions, and a flurry of groundbreaking results have been produced.<sup>2,5,8–12</sup> It is noteworthy that some cyclic Dy<sub>3</sub>,<sup>13–15</sup> Dy<sub>4</sub>,<sup>16–18</sup> and cyclic/local cyclic Dy<sub>6</sub> complexes<sup>19–21</sup> have exhibited SMM behaviours and toroidal moments that arise from a vortex arrangement of local magnetic moments of the individual Dy(III) centers. These systems are defined as single-molecule toroics (SMTs), which are promising candidates for future applications in quantum computing and information storage.<sup>5,22</sup> Toroidal moment is generally influenced by the molecular symmetry, local magnetic moment as well as magnetic interactions. However, the study on magneto-structural relationships of SMTs is rare.<sup>16,21</sup> Thus, it is of primary importance to design and construct a comprehensive (quasi)isostructural series of complexes to improve our knowledge of magneto-structural relationships.

In reality, most lanthanide SMMs have been synthesized by serendipitous approaches. Modulating the structure of complexes by design and choice of ligands is still a challenge. In general, one type of ligand affords a particular type of

complexes possessing a certain nuclearity and topology.<sup>23–26</sup> In this regard, we have been interested to investigate the magneto-structural relationships of SMTs with one typical ligand. Recently, we have successfully assembled a Dy<sub>6</sub> SMTs with enhanced toroidal moment that was formed by linkage of two Dy<sub>3</sub> units (the complex is defined as Dy<sub>6</sub>-2).<sup>27</sup> To go a step further, we modified slightly the Schiff-base ligand by adding a methyl, and assembled two  $\mu_4$ -O bridged hexanuclear lanthanide clusters, [Dy<sub>6</sub>L'<sub>4</sub>( $\mu_4$ -O)(NO<sub>3</sub>)<sub>4</sub>]·4CH<sub>3</sub>OH (1) and [Sm<sub>6</sub>L'<sub>4</sub>( $\mu_4$ -O)(CH<sub>3</sub>-COO)<sub>4</sub>]·2CH<sub>3</sub>OH (2). Though Dy<sub>6</sub>-2 and complex 1 have similar structures, the coordination geometries and magnetic interactions are changed upon the slight modulation of the ligand. The coordination environment of Dy(III) ions will influence the local tensor of anisotropy and crystal-field splitting. Thus they exhibit dissimilar magnetic behaviour. Complex 1 shows SMM behaviour with obvious quantum tunnelling of the magnetization (QTM) and SMT property with large conventional magnetization.

## Experimental section

### General

All starting materials were of A.R. Grade and were used as commercially obtained without further purification. 2,6-Diformyl-4-methylphenol (DFMP) were prepared according to a previously published method.<sup>28</sup> The Schiff-base ligand H<sub>3</sub>L' (2,6-bis((2-hydroxypropylimino)methyl)-4-methylphenol) (shown in Scheme 1) was prepared by the *in situ* condensation of DFMP and (DL)-1-amino-2-propanol in a 1 : 2 ratio in methanol.

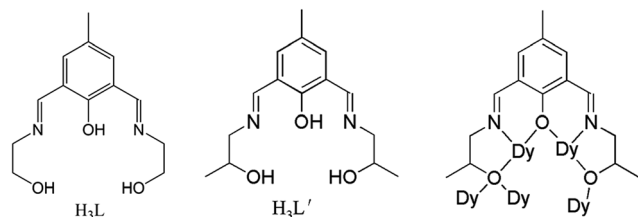
Elemental analyses for C, H, and N were carried out on a Perkin-Elmer 2400 analyzer. Fourier transform IR spectra (4000–300 cm<sup>-1</sup>) were measured using KBr pellets by a Fourier transform infrared spectrometer Nicolet 6700. All magnetization data were recorded on a Quantum Design MPMS-XL7 SQUID magnetometer equipped with a 7 T magnet. The

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Scheme 1 Ligands  $H_3L$  and  $H_3L'$  and the coordination formation of  $H_3L'$ .

variable-temperature magnetization was measured with an external magnetic field of 1000 Oe in the temperature range of 1.9–300 K. The experimental magnetic susceptibility data are corrected for the diamagnetism estimated from Pascal's tables and sample holder calibration.

### X-ray crystallography

Suitable single crystals of **1** and **2** were selected for single-crystal X-ray diffraction analysis. Crystallographic data were collected at 185(2) K on a Bruker ApexII CCD diffractometer with graphite monochromated Mo-K $\alpha$  radiation ( $\lambda = 0.71073$  Å). The structure was solved by direct methods and refined on  $F^2$  with full-matrix least-squares techniques using SHELXS-97 and SHELXL-97 programs.<sup>29,30</sup> The locations of Dy(III) ion were easily determined, and O, N, and C atoms were subsequently determined from the difference Fourier maps. Anisotropic thermal parameters were assigned to all non-hydrogen atoms. The H atoms were introduced in calculated positions and refined with a fixed geometry with respect to their carrier atoms. CCDC 1566095 and 1566096 contain the supplementary crystallographic data for this paper.†

### Synthesis of $[Dy_6L'_4(\mu_4-O)(NO_3)_4] \cdot 4CH_3OH$ (**1**)

A solution of  $Dy(NO_3)_3 \cdot 6H_2O$  (137.8 mg, 0.30 mmol) in 5 ml methanol was added in a solution of DFMP (0.15 mmol) and (DL)-1-amino-2-propanol (0.3 mmol) in 5 ml methanol in presence of triethylamine. The mixture was continually stirred at room temperature for 20 min. The resulting reaction mixture was sealed in an autoclave and maintained at 90 °C for 3 days, and then cooled slowly to room temperature to yield yellow single crystals suitable for X-ray diffraction. Yield: 33 mg, (27%, based on the metal). Elemental analysis (%) calcd for  $Dy_6C_{64}H_{92}N_{12}O_{29}$ : C, 31.14, H, 3.76, N, 6.81; found C, 30.96, H, 3.67, N, 6.50. IR (KBr,  $cm^{-1}$ ): 2856 (w), 1646 (vs), 1549 (m), 1497 (m), 1452 (s), 1396 (m), 1368 (w), 1323 (s), 1281 (m), 1231 (m), 1082 (m), 1042 (s), 902 (w), 873 (w), 813 (m), 775 (m), 739 (w).

### Synthesis of $[Sm_6L'_4(\mu_4-O)(CH_3COO)_4] \cdot 2CH_3OH$ (**2**)

A procedure similar to that for **1** was followed except that  $Dy(NO_3)_3 \cdot 6H_2O$  was replaced by  $Sm(CH_3COO)_3 \cdot H_2O$  (0.3 mmol, 105.2 mg). Yield: 27 mg (23%, based on the metal salt). Elemental analysis (%) calcd for  $Sm_6C_{70}H_{94}N_8O_{23}$ : C, 36.27; H, 4.09; N, 4.83. Found: C, 36.16; H, 3.97; N, 4.53. IR (KBr,  $cm^{-1}$ ): 2910 (w), 1644 (s), 1548 (vs), 1444 (s), 1395 (m), 1323 (m), 1231

(w), 1231 (w), 1138 (m), 1033 (s), 928 (w), 854 (w), 816 (w), 678 (w).

## Results and discussion

To explore the effect of additional methyl groups on the lanthanide compound, the symmetrical Schiff-base  $H_3L'$  (Scheme 1) was formed by the *in situ* condensation of DFMP and (DL)-1-amino-2-propanol (1 : 2) in methanol. Similar to the procedure for  $Dy_6-2$ , the reactions of *in situ* formed  $H_3L'$  with  $Dy(NO_3)_3 \cdot 6H_2O/Sm(CH_3COO)_3 \cdot H_2O$  in the presence of triethylamine under solvothermal condition produce block-shaped crystals of  $[Dy_6L'_4(\mu_4-O)(NO_3)_4] \cdot 4CH_3OH$  (**1**) and  $[Sm_6L'_4(\mu_4-O)(CH_3COO)_4] \cdot 2CH_3OH$  (**2**). The structures of **1** and **2** were established by single crystal X-ray diffraction. The crystal data and structure refinement are summarised in Table 1.

As shown in Fig. 1, complex **1** crystallizes in the triclinic space group  $P\bar{1}$  and consists of six Dy(III) ions, four  $(L')^{3-}$  ligands, one  $\mu_4-O$  anion and four  $NO_3^-$  anions. The core of **1**, similar to the  $Dy_6-2$ , consists of two  $[Dy_3(\mu_3-O)_2(\mu_2-O)_2]$  triangular units linked by one  $\mu_4-O^{2-}$  ion in an edge-to-edge arrangement. The triangular  $Dy_3$  units in **1** are more equilateral than that in  $Dy_6-2$  with Dy...Dy distances in range 3.4933(8)–3.4017(6) Å. The dihedral angle between the two  $Dy_3$  planes in **1** is 30.101(12)° that is slightly larger than that in  $Dy_6-2$  with 29.6568°.

It is notable that all donors in ligand  $H_3L'$  are coordinated, and each of the four  $(L')^{3-}$  binds five Dy(III) ions utilizing their tridentate pocket ( $O_2N$ ) along with bridging alkoxido oxygen (Scheme 1). This is different from some Schiff-base ligands that only coordinated partially.<sup>31</sup> Two arms of alkoxido oxygens in ligands bridge distinctly: while one arm bridges two Dy(III) ions of one triangular edge, the other arm links three Dy(III) ions upon the other triangular unit. Additionally, the phenoxido oxygen bridges two Dy(III) ions from two triangular units. Therefore, alkoxido oxygens from four ligands fasten  $Dy_3$  triangles, one  $\mu_4-O$  anion and four phenoxido oxygens of ligands further strengthen the “ $Dy_3 + Dy_3$ ” construction from center and periphery, respectively. This is the same with the reported  $Dy_6-2$ .

Table 1 Crystal data and structure refinement for complexes **1** and **2**

Compound	<b>1</b>	<b>2</b>
Empirical formula	$Dy_6C_{64}H_{92}N_{12}O_{29}$	$Sm_6C_{70}H_{94}N_8O_{23}$
Fw ( $g\ mol^{-1}$ )	2468.50	2317.63
Crystal system	Triclinic	Monoclinic
Space group	$P\bar{1}$	$C2/c$
$a$ (Å)	12.4459(5)	18.6094(11)
$b$ (Å)	13.8185(5)	17.218(1)
$c$ (Å)	25.2852(10)	25.3390(14)
$\alpha$ (°)	82.073(1)	90
$\beta$ (°)	81.870(1)	92.634(1)
$\gamma$ (°)	75.400(1)	90
$V$ (Å <sup>3</sup> )	4142.1(3)	8110.5(8)
$Z, \rho_{calcd}$ ( $Mg\ m^{-3}$ )	2, 1.979	4, 1.898
$F(000), R_{int}$	2376, 0.0306	4504, 0.0744
$R_1, wR_2 [I > 2\sigma(I)]$	0.0455, 0.1085	0.0495, 0.1146
$R_1, wR_2$ (all data)	0.0750, 0.1271	0.0887, 0.1394
GO F	1.006	1.008



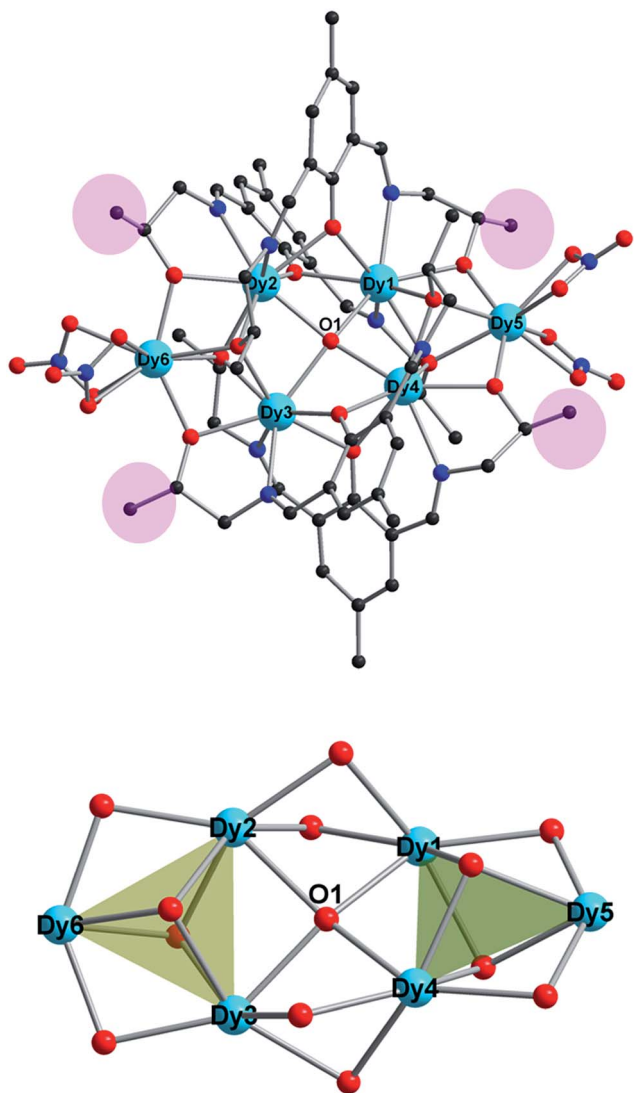


Fig. 1 (Top) Structure of 1 with hydrogen atoms omitted for clarity. (Bottom)  $\mu_4$ -O bridged hexanuclear core.

Such a robust edge-to-edge arrangement will enhance the toroidal magnetic moment of the molecule.

The coordination spheres of Dy5 and Dy6 ions are completed respectively by two nitrate ions, and all Dy(III) ions are eight-coordinate (Fig. 2). This is different from Dy<sub>6</sub>-2, where five Dy(III) centers are eight-coordinate and one Dy(III) ion is nine-coordinate. The exact geometries of all Dy(III) centers were determined by using the SHAPE 2.1<sup>32,33</sup> and the results are listed in Table S1.† For complex 1, Dy1/Dy3/Dy5/Dy6 centers are situated in distorted square antiprismatic geometries, and Dy2/Dy4 centers are situated in distorted triangular dodecahedron. For Dy<sub>6</sub>-2, Dy1/Dy3/Dy5 centers are situated in distorted square antiprismatic geometries, Dy2/Dy4 centers are situated in distorted triangular dodecahedron, and Dy6 ion is in mono-capped square antiprism. The changes probably have a great impact on the magnetic anisotropy and further the magnetic properties.<sup>34–37</sup>

In addition, the Dy–O bond lengths fall in the range of 2.235(6)–2.478(7) Å with the average bond of 2.37 Å, the Dy–N

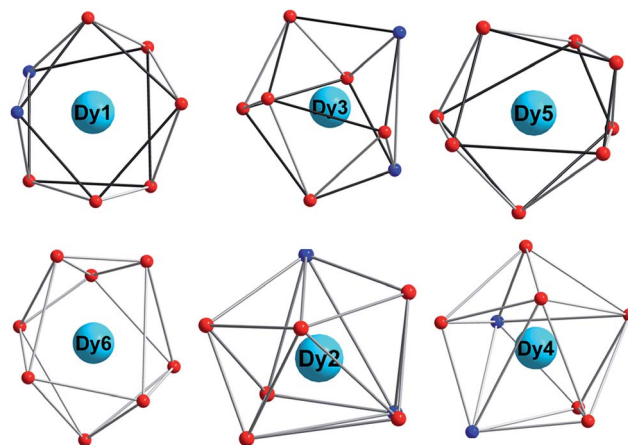


Fig. 2 Coordination polyhedra (distorted square antiprism for Dy1, Dy3, Dy5 and Dy6; distorted triangular dodecahedron for Dy2 and Dy4) for six Dy<sup>3+</sup> ions in 1.

bond lengths fall in the range of 2.457(8)–2.490(8) Å with the average bond of 2.47 Å. Those average bonds in complex 1 are similar to that of Dy<sub>6</sub>-2 (average bonds are 2.48 and 2.38 Å for Dy–O and Dy–N, respectively). This indicates the metal–ligand interactions of complex 1 and Dy<sub>6</sub>-2 are similar.

In [Dy<sub>6</sub>L<sub>4</sub>( $\mu_4$ -O)(NO<sub>3</sub>)<sub>4</sub>] core, the four ligands are twisted along the C–N bonds, and each ligand straddles two triangles to coordinate five Dy(III) ions, resulting in formation of quadruple-stranded helicates (Fig. 3).<sup>38–40</sup> Significantly, two stereoisomers of right- ( $\Delta$ ) and left-hand ( $\Lambda$ ) configurations are in crystal, forming a racemic mixture. The packing arrangement of 1 reveals that  $\Delta$ - and  $\Lambda$ -configuration helicates are alternate along the crystallographic *a* axis (Fig. 4).

Complex 2 crystallizes in the monoclinic space group *C*2/*c*, the asymmetric unit contains four Sm(III) ions, two (L)<sup>3-</sup> ligands and two CH<sub>3</sub>COO<sup>-</sup> ions. The structure of 2 is essentially isomorphous to 1 (Fig. S1†), consists of a  $\mu_4$ -O<sup>2-</sup> ion linked two triangular units in an edge-to-edge arrangement, where Sm...Sm distances are in range of 3.4439(10)–3.6234(96) Å, the dihedral angle between the two Sm<sub>3</sub> planes in 2 is 28.176° that is smaller than that in 1. In the asymmetric unit, Sm2 and Sm3 ions are eight-coordinate with 6O<sub>2</sub>N, and both display distorted triangular dodecahedral geometries based on SHAPE 2.1

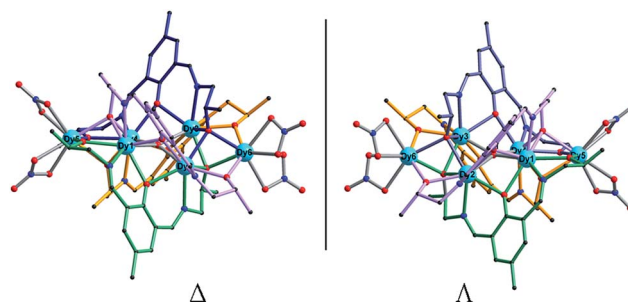


Fig. 3 Two stereoisomers of right- ( $\Delta$ ) and left-hand ( $\Lambda$ ) configurations in 1.



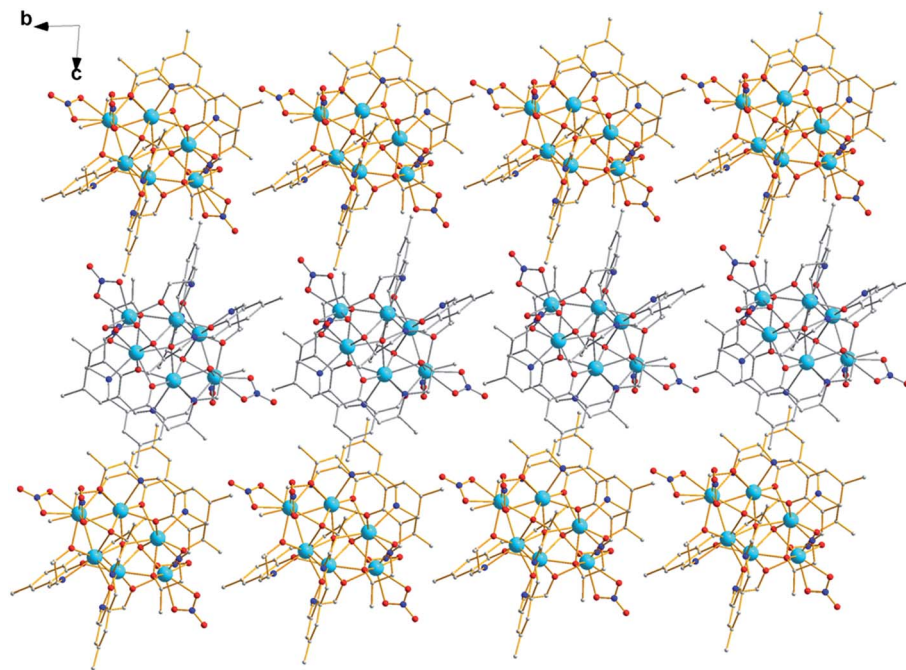


Fig. 4 Crystal packing of **1** along the crystallographic *a* axis showing the stereoisomers.

(Fig. S2 and Table S2†). The coordination spheres of Sm1 and Sm4 ions are filled respectively by two  $\text{CH}_3\text{COO}^-$  ions, forming eight-coordinate Sm1 and Sm4 ions with 8O. The exact geometries of Sm1 and Sm4 ions are both distorted square antiprisms based on SHAPE 2.1. In addition, the Sm–O bond lengths fall in the range of 2.331(6)–2.506(8) Å, the Sm–N bond lengths fall in the range of 2.507(9)–2.552(9) Å.

In  $[\text{Sm}_6\text{L}'_4(\mu_4\text{-O})(\text{CH}_3\text{COO})_4]$  core, each of four ligands coordinates five Sm(III) ions and straddles two triangular units, producing formation of quadruple-stranded helicates (Fig. S3†). As shown in Fig. S4,† the two stereoisomers of  $\Delta$  and  $\Lambda$  configurations form a racemic mixture, which can be seen in molecular structures and crystal packing.

### Magnetic studies

The direct current (dc) susceptibility studies of complexes **1** and **2** were investigated under a 1000 Oe field in the temperature range 2–300 K (Fig. 5). At room temperature, the  $\chi_{\text{M}}T$  value of complex **1** is  $86.37 \text{ cm}^3 \text{ K mol}^{-1}$ . The value is in agreement with the expected theoretical value ( $85.02 \text{ cm}^3 \text{ K mol}^{-1}$ ) for six free Dy(III) ions ( ${}^6\text{H}^{15/2}$ ,  $S = 5/2$ ,  $L = 5$ ,  $g = 4/3$ ,  $C = 14.17 \text{ cm}^3 \text{ K mol}^{-1}$ ). On cooling,  $\chi_{\text{M}}T$  decreases throughout the whole temperature range: firstly,  $\chi_{\text{M}}T$  decreases monotonously down to 50 K, then decreases abruptly to  $21.03 \text{ cm}^3 \text{ K mol}^{-1}$  at 2.0 K. The thermal variation of the  $\chi_{\text{M}}T$  indicates the progressive depopulation of the excited Stark sublevels of Dy(III) ions and/or weak molecular magnetic interactions.<sup>41,42</sup> Actually, this profile of  $\chi_{\text{M}}T$  is similar but higher than that of Dy<sub>6</sub>-2, and the minimum is larger than that of Dy<sub>6</sub>-2 ( $6.59 \text{ cm}^3 \text{ K mol}^{-1}$  at 2.0 K), which indicates the weaker magnetic interaction in complex **1**. The large minimum ( $21.03 \text{ cm}^3 \text{ K mol}^{-1}$ ) also indicates the magnetic ground state of complex **1** is different from the net toroidal moment.<sup>16,20,43</sup>

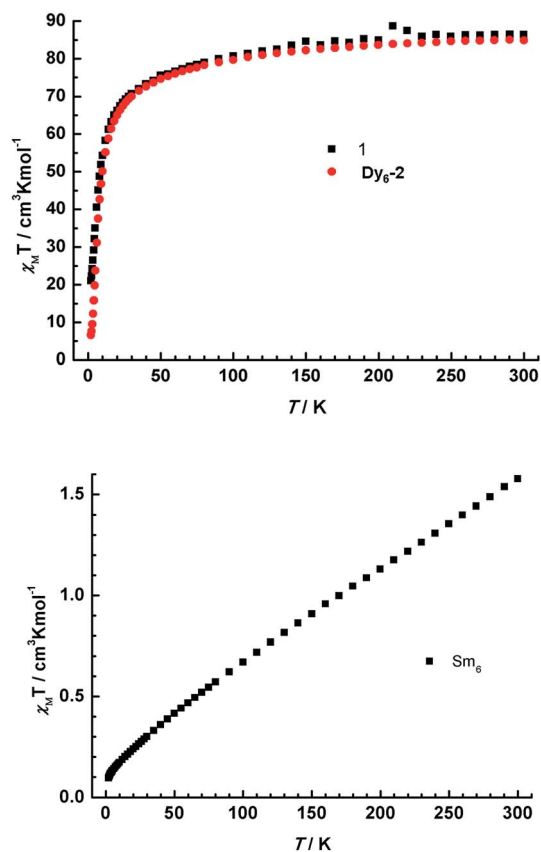


Fig. 5 (Top) Plots of  $\chi_{\text{M}}T$  vs.  $T$  for **1** (black square) and Dy<sub>6</sub>-2 (red circle) in a dc field of 1000 Oe (2–300 K); (bottom) plots of  $\chi_{\text{M}}T$  vs.  $T$  for **2** in a dc field of 1000 Oe.



For complex **2**, the  $\chi_M T$  value (Fig. 5) observed at room temperature is  $1.58 \text{ cm}^3 \text{ K mol}^{-1}$ , which is significantly larger than the theoretical one. This is attributed to the first or even the higher excited states ( ${}^6\text{H}^{7/2}$ ,  ${}^6\text{H}^{9/2}$ , ...,  ${}^6\text{H}^{15/2}$ ) of the  $\text{Sm}(\text{III})$  ion that can be populated obviously at room temperature.<sup>44</sup> On cooling, the  $\chi_M T$  value decreases continuously to a minimum of  $0.10 \text{ cm}^3 \text{ K mol}^{-1}$  at 2.0 K.

The field dependence of magnetization ( $M$ ) was measured between 1.9 and 5 K. As shown in Fig. 6, it is obviously that  $M$  under variable field at 1.9 K shows a "S" shape, which is consistent with SMTs.<sup>22</sup> The peak ( $d(M/M_s)/dH$ ) vs. field suggests inflection around 12 kOe. The inflection is less evident than that in  $\text{Dy}_6\text{-2}$ .  $M$  reaches a maximum value  $33.83 \mu_B$  at about 45 kOe, which is far lower than the expected saturation value of  $60 \mu_B$  for six noninteracting  $\text{Dy}(\text{III})$  ions. This can be mostly attributed to crystal-field effects and the possible weak antiferromagnetic interactions that make the low-lying excited states accessible. Meanwhile, the non-superposition of the  $M$  vs.  $H/T$  data implies the presence of significant magnetic anisotropy. The static magnetic behaviours, including the dc susceptibility and the field dependence of  $M$ , reveal that the ground state of **1** could be

toroidal magnetic moment and the conventional magnetic moment should be larger than that of  $\text{Dy}_6\text{-2}$ .<sup>22</sup>

Furthermore, the anisotropy axes of complex **1** were calculated using the Magellan program<sup>45–47</sup> based on an electrostatic model (Fig. 7 and Table S3†). The results indicate that complex **1** shows toroidal moment in both  $\text{Dy}_3$  units. This agrees with the magnetic properties. The *ab initio* calculations<sup>48–50</sup> are planned to provide quantitative evaluation of the anisotropy axes and Dy–Dy coupling interactions.

To compare the magnetic relaxation between  $\text{Dy}_6\text{-2}$  and **1**, the dynamics of the magnetization of **1** were investigated using alternating current (ac) susceptibility measurements (Fig. 8). Under zero dc field, the variable-temperature ac susceptibility of **1** shows the temperature dependence maximum, which signals the freezing of the spins by the anisotropy barrier and is typical for SMM behaviour. Both **1** and  $\text{Dy}_6\text{-2}$  show broad peak in the out-of-phase susceptibility, but the former appears at 7 K that is lower than that of  $\text{Dy}_6\text{-2}$  (at about 9 K). This indicates lower effective energy barrier for complex **1**. Obviously, unlike the  $\text{Dy}_6\text{-2}$ , the rise in the out-of-phase and in-phase susceptibility on cooling at low temperatures indicates the presence of QTM for **1**, which may be due to the weak magnetic interaction that is too weak to suppress the QTM.<sup>51–53</sup> This agrees well with dc magnetic susceptibility that indicates weaker magnetic interaction in complex **1**.

The reported isomorphous  $\text{Dy}_6\text{-2}$  displayed impressive dynamic magnetic properties with effective suppression of QTM and remarkable toroidal moment. However, obvious QTM is present and toroidal moment is less distinct in **1**. Comparing the structure data of complex **1** and  $\text{Dy}_6\text{-2}$ , they have the identical metal–ligand interaction, but distinctive coordination geometries mainly on  $\text{Dy}_6$  ion. For  $\text{Dy}_6\text{-2}$ ,  $\text{Dy}_6$  ion is nine-coordinate with a mono-capped square antiprism. While  $\text{Dy}_6$

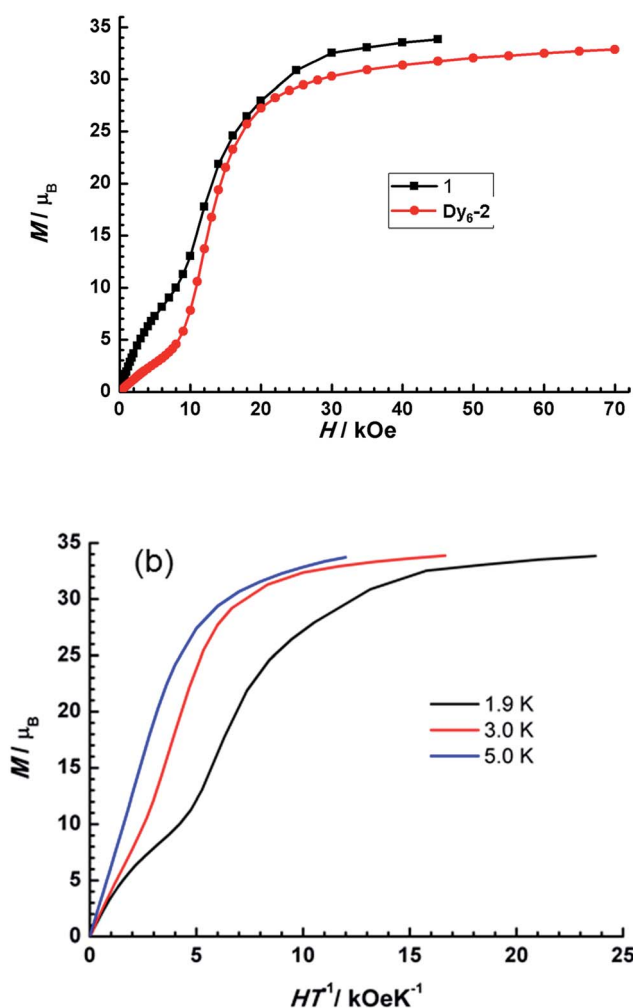


Fig. 6 (Top) Field dependence of magnetization at 1.9 K for **1** and  $\text{Dy}_6\text{-2}$ . (Bottom)  $M$  versus  $H/T$  plots for **1** at 1.9 K, 3 K and 5 K.

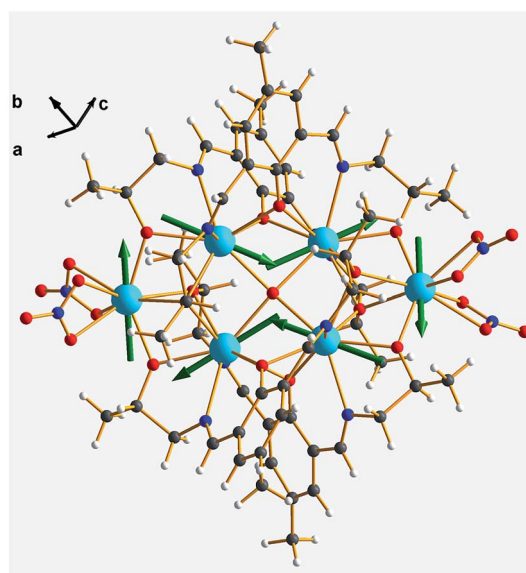


Fig. 7 Ground-state magnetic anisotropy of complex **1**. The green arrows represent the orientations of the anisotropic axes for each  $\text{Dy}(\text{III})$  ion, as calculated by the electrostatic model.



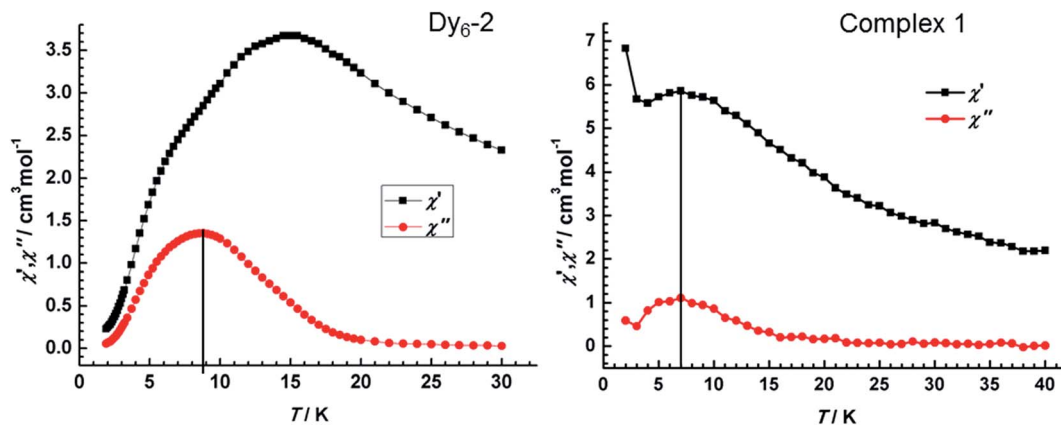


Fig. 8 Temperature dependence of the out-of phase and in-phase ac susceptibility for  $\text{Dy}_6\text{-2}$  (left) and **1** (right) at 1000 Hz under zero dc field.

center in complex **1** is eight-coordinate situated in distorted square antiprismatic geometries, which is due to the steric hindrance from additional methyl groups of ligands (Fig. 1). The changes in coordination geometries have a great impact on the orientation of the easy axes. Consequently, the toroidal moment retains in complex **1** but with a large conventional magnetization, SMM behaviour also is observed but with a non-ignorable QTM. It should be note that the weaker magnetic interaction in complex **1** could not suppress the QTM.

## Conclusions

In conclusion, methyl modified Schiff-base ligand has been successfully used to synthesize two new hexanuclear lanthanide complexes,  $[\text{Dy}_6\text{L}'_4(\mu_4\text{-O})(\text{NO}_3)_4]\cdot 4\text{CH}_3\text{OH}$  (**1**) and  $[\text{Sm}_6\text{L}'_4(\mu_4\text{-O})(\text{CH}_3\text{COO})_4]\cdot 2\text{CH}_3\text{OH}$  (**2**), under solvothermal condition. Single crystal X-ray diffraction studies revealed that complex **1** has a  $\mu_4\text{-O}$  bridged  $\text{Dy}_6$  core,  $\mu_4\text{-O}$  bridged two  $[\text{Dy}_3(\mu_3\text{-O})_2(\mu_2\text{-O})_2]$  triangular units in an edge-to-edge arrangement, that is the same as the parent  $\text{Dy}_6\text{-2}$ . While the coordination geometries especially on  $\text{Dy}_6$  site and magnetic interactions change, which result in distinct magnetic properties. Complex **1** shows SMM behaviour with a non-ignorable QTM and SMT behaviour with a large conventional magnetization. The work provides an efficient strategy to probe the magneto-structural relationships of SMTs, further to design the special SMTs and SMMs.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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