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Reversible ON–OFF switching of single-molecule-magnetism associated with single-crystal-to-single-crystal structural transformation of a decanuclear dysprosium phosphonate†

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{Dy₅(EDDC)₂(μ₃-AcO)₂(μ₅-C₁₅H₁₁PO₃)(μ₄-C₁₅H₁₁PO₃)(μ₂-AcO)₂(AcO)₂(H₂O)(CH₃OH)₂}(μ₄-C₂O₄)·xH₂O (I), where H₂EDDC is *N',N',E,N',N',E-N',N'*-(ethane-1,2-diylidene)dipyrazine-2-carbohydrazide and C₁₅H₁₁PO₃H₂ is 9-anthrylmethylphosphonic acid, is found to undergo two consecutive single-crystal-to-single-crystal transformations. The first is under UV irradiation (λ = 365 nm for 3 d in air) to {Dy₅(EDDC)₂(μ₃-AcO)₂(μ₅-C₁₅H₁₁PO₃)₂(μ₂-AcO)₂(AcO)₂(H₂O)₃}(μ₄-C₂O₄)·xH₂O (I-UV) where the two CH₃OH are replaced by two H₂O and the second by annealing under N₂ at 100 °C on a diffractometer or under Ar in a glovebox to {Dy₅(EDDC)₂(μ₃-AcO)₂(μ₅-C₁₅H₁₁PO₃)₂(μ₂-AcO)₄(H₂O)₂}(μ₄-C₂O₄) (I-A-N₂ or I-A-Ar) where it has lost two H₂O molecules. The second transformation is reversible by exposure to air at room temperature (I-A-N₂-cool). While the overall structures are the same (retaining the space group *P2₁/c*), there is a considerable expansion of the unit cell from I (8171 Å³) to I-UV (8609 Å³) and I-A-N₂ (8610 Å³) and the coordinations of the Dy atoms undergo major reconstructions. This is associated with switching the single-molecule-magnetism (SMM) from OFF for I to ON for I-UV and to OFF again for I-A-Ar in air. Such a switching mechanism associated with the retention of crystallinity is unique in the chemistry of dysprosium. The structure of the molecule is formed from two symmetry related pentamers joined by an oxalate. A related compound containing two isolated neutral pentamers {Dy₅(EDDC)₂(μ₃-AcO)₂(μ₅-C₁₅H₁₁PO₃)₂(μ₂-AcO)₃(AcO)₂(H₂O)₂}{Dy₅(EDDC)₂(μ₃-AcO)₂(μ₅-C₁₅H₁₁PO₃)(μ₄-C₁₅H₁₁PO₃)(μ₂-AcO)₃(AcO)₂(CH₃OH)₂}·2CHCl₃ (II) has also been isolated with closely related Dy coordination and it exhibits similar SMM behaviour in zero field.

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Introduction

Molecular compounds that can have their physical properties switched from one state to another reversibly and retain their crystallinity are quite rare.¹ And clusters that can switch their single-molecule-magnetism (SMM) ON and OFF are even fewer. In

general, numerous compounds are known where the parameters defining their SMM are altered but are not switched ON and OFF reversibly. Since the discovery of SMM in the early nineties,² there is a general wish to use these SMMs as ON–OFF switches which will endow them with a bi-stable magnetic ground state that has been coined as a great attribute in applications for information storage and quantum computing.^{3–5} Consequently, there is major interest in realising these properties by design using both d- and f-elements.^{6–8} Lanthanide complexes are becoming more promising candidates for SMMs because of their significant single-ion magnetic anisotropy arising from the large unquenched orbital angular momentum and strong spin–orbit coupling, and thus may lead to high energy barriers for spin reversal.^{9–12} Additionally, the very weak intermolecular interaction compared to those made up of d-elements makes them very appealing for single-molecule magnetism. However, the magnetization dynamics of lanthanide based SMMs (Ln-SMMs) is still far from being fully understood, because they are influenced by different factors such as the symmetry and the charge distribution of the coordination sphere around the lanthanide ions, the hyperfine couplings, dipolar

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Scheme 1 Molecular structures of $N',N'',E,N',N'',E-N',N''$ -(ethane-1,2-diylidene)dipyrazine-2-carbohydrazide (H_2EDDC , left) and 9-anthrylmethylphosphonic acid ($C_{15}H_{11}PO_3H_2$, right).

spin-spin interactions, *etc.*^{9–13} For polynuclear clusters the magnetic behaviour can be further complicated by the different moment vector directions.

While great efforts have been devoted to the design and syntheses of new Ln-SMMs with high energy barriers and blocking temperatures,^{14–18} the manipulation of the magnetic properties of Ln-SMMs through external stimuli has rarely been explored. Such manipulations can be realized by imposing chemical influences such as ligand redox^{19–21} and pH control with a base or acid.²² Guest solvent exchange was also found to induce a reversible structural transformation but only tunable magnetization relaxation.^{23,24} Reversible switching of SMM behaviors can also be achieved by using physical stimuli. Light-induced switching of magnetic relaxation has been realized in Dy compounds containing optically active diarylethene²⁵ or 1,2-bis(4-pyridyl)ethane linkages.²⁶ Thermal treatment can remove lattice or coordinated solvent molecules, switching the magnetic behavior of Ln-SMMs.^{27,28} But only in one case, as far as we are aware, ON–OFF SMM switching coupled with reversible single-crystal-to-single-crystal (SC–SC) structural transformation was achieved for Ln-SMMs.²⁹

Cyclic lanthanide rings belong to a specific subclass of Ln-based clusters. These rings not only have structural aesthetics but also are magnetically interesting. In particular, the observation of a toroidal arrangement of local magnetization vectors in a Dy_3 triangular cluster^{30,31} stimulates the enthusiasm for the exploration of new cyclic Ln-based clusters.³² Unfortunately, there are a very few examples of cyclic lanthanide rings reported so far, and most contain even-numbered lanthanide ions.^{32–37} The syntheses of odd-numbered lanthanide rings larger than three have therefore been a great challenge.³⁸ Recently we developed a new synthetic strategy by combining pre-designed trimeric and dimeric building blocks and a phosphonate ligand as an adhesive co-ligand.³⁹ The present cyclic heptanuclear Dy_7 clusters were successfully obtained, which turn out to be the first with seven Dy on the ring and none inside.

In this paper, we report the syntheses, structures and magnetic properties of a novel series containing oxalate bridged pentamers, $\{Dy_5(EDDC)_2(\mu_3-AcO)_2(\mu_5-C_{15}H_{11}PO_3)(\mu_4-C_{15}H_{11}PO_3)(\mu_2-AcO)_2(AcO)_2(H_2O)_2(CH_3OH)_2\}_2(\mu_4-C_2O_4) \cdot xH_2O$ (**I**) and its SC–SC transformations to two other forms, $\{Dy_5(EDDC)_2(\mu_3-AcO)_2(\mu_5-C_{15}H_{11}PO_3)_2(\mu_2-AcO)_2(AcO)_2(H_2O)_3\}_2(\mu_4-C_2O_4) \cdot xH_2O$ (**I-UV**) and $\{Dy_5(EDDC)_2(\mu_3-AcO)_2(\mu_5-C_{15}H_{11}PO_3)_2(\mu_2-AcO)_4(H_2O)_2(\mu_4-C_2O_4)\}$ (**I-A-N₂**), where H_2EDDC is $N',N'',E,N',N'',E-N',N''$ -(ethane-1,2-diylidene)dipyrazine-2-carbohydrazide, and $C_{15}H_{11}PO_3H_2$ is 9-anthrylmethylphosphonic acid (Scheme 1), consisting of a fused pair of a cyclic pentamer by oxalate. **I** undergoes two consecutive SC–SC transformations accompanied by switching

ON and OFF of the single-molecule-magnetism reversibly. Another compound consisting of two independent structurally related pentamers, $\{Dy_5(EDDC)_2(\mu_3-AcO)_2(\mu_5-C_{15}H_{11}PO_3)_2(\mu_2-AcO)_3(AcO)_2(H_2O)_2\}_2\{Dy_5(EDDC)_2(\mu_3-AcO)_2(\mu_5-C_{15}H_{11}PO_3)(\mu_4-C_{15}H_{11}PO_3)(\mu_2-AcO)_3(AcO)_2(CH_3OH)_2\} \cdot 2CHCl_3$ (**II**), has also been isolated and characterised. It displays similar SMM behaviour. These phases are the first examples of odd-numbered rings with cyclic Ln-pentamers displaying the unique ON–OFF switching of SMM properties.

Results and discussion

Structures of **I** and **II**

I was synthesized as dark red crystals from a water–methanol (1 : 3) mixture containing H_2EDDC neutralised with triethylamine, $Dy(OAc)_3 \cdot 4H_2O$, sodium oxalate and $C_{15}H_{11}PO_3H_2$. As shown in Scheme 2, the crystals undergo two SC–SC transformations, the first under UV irradiation to form **I-UV** and the second by annealing at 100 °C under N_2 on a diffractometer (or under Ar in a glovebox) to form **I-A-N₂** (or **I-A-Ar**). Upon exposure to air, **I-A-N₂** (or **I-A-Ar**) transforms into **I-A-N₂-cool** (or **I-A-Ar-cool**). Therefore, we have determined the structure in each state (**I**, **I-UV**, **I-A-N₂** and **I-A-N₂-cool**). A summary of the crystallographic data is given in Tables 1 and S1.† We therefore describe the five structures with enough details for a thorough understanding. Further details can be seen in the ESI (Tables S2 and S3†) and the crystallographic information files. We note that there are three different molecular contents for the five structures where those of **I-UV**, **I-A-N₂-cool** and **I-A-Ar-cool** are the same. Therefore, only the structure of **I-UV** is described.

The crystals adopt the monoclinic space group $P2_1/c$ ($Z = 2$) in all the forms. The key feature of the structures is two symmetry-related cyclic $[Dy_5]$ units, looking like a distorted star, fused by a bis-chelating oxalate ligand (Fig. 1 and 2). Each Dy_5 unit contains a common fragment consisting of five independent Dy atoms, two $EDDC^{2-}$ and two μ_3 -acetate (shown in grey in Fig. 2). The oxygen atoms of one $EDDC^{2-}$ ligand form two bridges between three Dy atoms (Dy_1 , Dy_2 and Dy_3) in a pseudo-linear array. It also provides five chelate units (4 N–O and 1 N–N) to hold the three Dy atoms. The other $EDDC^{2-}$ is hinged at the apical Dy atom (Dy_1) and hold two other Dy atoms (Dy_4 and Dy_5)



Scheme 2 Protocol used for the transformations of **I**.



the star ring, in both independent molecules (Fig. 3 and Table S2†).

The remaining ligands adopt different coordination modes in all the five different pentamers of compounds **I** and **II**. Table 2 lists the modes of coordination of each ligand and the total coordination bonds per pentamer. We find that there are additionally a μ_2 -bridge (*syn-syn* mode), chelate and terminal acetate (shown in blue). The two phosphonates (shown in red) per pentamers can be only μ_5 - or mixed with μ_4 - $C_{15}H_{11}PO_3^{2-}$ as in **I** and **II-molecule B**. They sit on each face of the pentamers with a P...P distance of *ca.* 3.7 Å. Methanol or water molecules (shown in green) are also present in prepared **I** which contains two coordinated methanol and one water molecules. When it is exposed to UV light it is transformed to **I-UV** where the methanol molecules are replaced by two water molecules, presumably from the atmosphere. Upon heating **I-UV** at 100 °C under a flow of N_2 on a diffractometer it loses two of the three molecules of water to form **I-A-N₂**. During the two transformations major reconstructions have taken place as can be seen from Table 2 and in the ESI.† After the subsequent exposure of **I-A-N₂** in air, it reabsorbs two water molecules to **I-A-N₂-cool** which has almost the same molecular content and structure as **I-UV** (Tables 1 and S2†). When **I-UV** is heated at 100 °C under argon in a glovebox it loses its solvents forming **I-A-Ar**. Upon subsequent exposure to air it quickly absorbs water to give the same structure as **I-A-N₂-cool** and **I-UV**, confirmed by the crystal structure determination (Tables S1 and S3†).

Because of the changes in the coordination of the ligands the coordination number (CN) per cluster is also varied from 41 to 44. For example, it is 42 for **I**, meaning that $3 \times CN = 8$ and $2 \times CN = 9$, 43 for **I-UV** ($2 \times CN = 8$ and $3 \times CN = 9$), 44 for **I-A-N₂** ($1 \times CN = 8$ and $4 \times CN = 9$), 42 for **II-molecule A** ($3 \times CN = 8$ and $2 \times CN = 9$), and 41 for **II-molecule B** ($4 \times CN = 8$ and $1 \times CN = 9$). The different coordination geometries suggested by the Shape analysis program (SchM)⁴⁰ are listed in Tables 2 and S5–S8.† While the coordination number of the Dy sharing the two EDDC²⁻ is always eight, their geometries are close to either a triangular dodecahedron (D_{2d}) or square antiprism (D_{4d}). The coordination numbers of the other atoms are eight or nine and are random with a wide range of geometries.

The oxalate anion serves as a bis-chelate bridge through its four oxygen atoms between the two pentamers in the different structures of compound **I**. The Dy...Dy distances over the oxalate bridges are 6.32 Å for **I**, 6.30 Å for **I-UV** and 6.21 Å for **I-A-N₂**, respectively. It is planar in each case.

The Dy–O and Dy–N bond lengths for the five pentamers are in the range 2.1–2.5 Å and 2.4–2.7 Å, respectively. One of the two anthracene of the coupled pentamers in the structure of **I** is oriented nearly face-to-face to the aromatic rings of the EDDC²⁻ and the other protrudes out of the long axis of the molecules in such a way to form π - π interaction (plane-to-plane distance of 3.39 Å and centroid-to-centroid distance of 3.68 Å) with neighbouring molecules to form chains (Fig. 1 and S7†). But for **II** the situation is completely different. The two anthracenes are oriented towards the EDDC²⁻ ligand and no π - π interaction is possible. However, the structure may be viewed as a 2D supra-molecular plane with a network arrangement of weakly

intermolecular H-bonds between the pentamers (Fig. S15†). The shortest intermolecular Dy...Dy distance is 9.1 Å.

Given the short distance of less than 4.2 Å between the peripheral anthracene moieties in **I**, it may be favourable for solid-state $[4\pi + 4\pi]$ photodimerization reaction. With this in mind, UV-irradiation with 365 nm of compound **I** was performed for 3 days at room temperature. It experiences SC–SC transformation to **I-UV** (Fig. 2). However, the structure of the UV irradiated crystal suggests that photodimerization does not occur but the methanol molecules of the original compound **I** were replaced by two water molecules. The replacement is not at the same Dy atom hosting the methanol. There are several concerted steps: (i) the departure of methanol on Dy₄ is compensated by the bridging of the terminal acetate of Dy₃, (ii) the methanol of Dy₅ is replaced by water and (iii) the bridging acetate (Dy₁–Dy₂) breaks one bond with Dy₁ and a water molecule takes its place and (iv) the terminal acetate (Dy₃) then becomes chelated to Dy₃. The π - π interaction between the anthracene groups in **I-UV** becomes weaker with a plane-to-plane distance of 3.42 Å and a centroid-to-centroid distance of 4.40 Å. The transformation of **I-UV** to **I-A-N₂** has two modifications: (i) the departing water of Dy₁ promotes the bridging of the terminal acetate on Dy₂ – a reverse of the changes from **I** to **I-UV**, and (ii) the chelate acetate of Dy₃ then forms a bridge between Dy₃ and Dy₂.

The lattice solvent molecules in **I** and **II** are heavily disordered and thus treated with the PLATON/SQUEEZE program. The solvent accessible volumes per unit cell are 665 Å³ for **I**, 1420 Å³ for **I-UV**, 1538 Å³ for **I-A-N₂** and 8741 Å³ for **II**, respectively. It is surprising that the volume of **I** is significantly smaller than that of the other related compounds. After a careful analysis of the structures of **I** and **I-UV**, we found that the methyl groups from the coordinated methanol molecules (C72 and C73) and the terminal acetate (C68) in **I** occupy partial voids which is different from the case of **I-UV** (Fig. S16 and S17†). This may explain the fact that the solvent accessible void in **I** is largely reduced compared with that in **I-UV**. The number of lattice solvent molecules is determined by thermal analyses.

Magnetic properties

The direct-current (dc) magnetic susceptibility was measured as a function of temperature (1.8–300 K) in a field of 1 kOe on polycrystalline samples of **I**, **I-UV**, **I-A-Ar** and **II** (Fig. S18†). At 300 K, the $\chi_M T$ ($cm^3 K mol^{-1}$) values per Dy₁₀ are 139.9 for **I**, 137.4 for **I-UV**, 135.6 for **I-A-Ar** and 140.4 for **II**, close to the expected values of 141.7 for ten non-interacting Dy^{III} ($S = 5/2$, $L = 5$, ${}^6H_{15/2}$, $g = 4/3$, and $C = 14.17 cm^3 K mol^{-1}$). The $\chi_M T$ of all the compounds behaves similarly showing a decrease on lowering the temperature from 300 to 30 K, which is mainly due to the progressive depopulation of the excited Stark sublevels of the m_J states of the Dy^{III}.^{41–44} Below 30 K, weak intra-/inter-molecular antiferromagnetic coupling becomes operative and the $\chi_M T$ shows a sharp decline to 88.1 (**I**), 79.2 (**I-UV**), 81.5 (**I-A-Ar**) and 86.7 $cm^3 K mol^{-1}$ (**II**). This behaviour is typical of Dy^{III}.^{10,11}

The isothermal magnetizations up to 70 kOe were measured for each sample at different temperatures (Fig. S19†). The





Table 2 Summary of coordination geometries of the dysprosium atoms and bonding modes of the components in I, I-UV, I-A-N₂ and II

	I	I-UV	I-A-N ₂	II-molecule A	II-molecule B
Formula	{Dy ₅ (EDDDC) ₂ (μ ₃ -AcO) ₂ (μ ₅ -C ₁₅ H ₁₁ PO ₃)(μ ₄ -C ₁₅ H ₁₁ PO ₃)(μ ₂ -AcO) ₂ (AcO) ₂ (H ₂ O)(CH ₃ OH) ₂ (μ ⁴ -C ₂ O ₄)	{Dy ₅ (EDDDC) ₂ (μ ₃ -AcO) ₂ (μ ₅ -C ₁₅ H ₁₁ PO ₃)(μ ₂ -AcO) ₂ (c-AcO) ₂ (H ₂ O) ₂ (μ ⁴ -C ₂ O ₄)	{Dy ₅ (EDDDC) ₂ (μ ₃ -AcO) ₂ (μ ₅ -C ₁₅ H ₁₁ PO ₃)(μ ₂ -AcO) ₂ (H ₂ O) ₂ (μ ⁴ -C ₂ O ₄)	{Dy ₅ (EDDDC) ₂ (μ ₃ -AcO) ₂ (μ ₅ -C ₁₅ H ₁₁ PO ₃)(μ ₂ -AcO) ₂ (c-AcO) ₂ (H ₂ O) ₂	{Dy ₅ (EDDDC) ₂ (μ ₃ -AcO) ₂ (μ ₅ -C ₁₅ H ₁₁ PO ₃)(μ ₄ -C ₁₅ H ₁₁ PO ₃)(μ ₂ -AcO) ₂ (c-AcO) ₂ (CH ₃ OH) ₂
Acetate	2 μ ₃ <i>syn-syn-anti</i> (Dy ₃ -Dy ₄), 2 μ ₂ <i>syn-syn</i> (Dy ₁ -Dy ₂ , Dy ₁ -Dy ₅), 2 terminal (Dy ₃)	2 μ ₃ <i>syn-syn-anti</i> (Dy ₃ -Dy ₄), 2 μ ₂ <i>syn-syn</i> (Dy ₁ -Dy ₅ , Dy ₃ -Dy ₄), 1 chelate (Dy ₃), 1 terminal (Dy ₂)	2 μ ₃ <i>syn-syn-anti</i> (Dy ₃ -Dy ₄), 4 μ ₂ <i>syn-syn</i> (Dy ₁ -Dy ₂ , Dy ₁ -Dy ₅ , Dy ₂ -Dy ₃ , Dy ₃ -Dy ₄)	2 μ ₃ <i>syn-syn-anti</i> (Dy ₄ -Dy ₅), 3 μ ₂ <i>syn-syn</i> (Dy ₂ -Dy ₃ , Dy ₂ -Dy ₄ , Dy ₄ -Dy ₅), 2 chelate (Dy ₄ , Dy ₅)	2 μ ₃ <i>syn-syn-anti</i> (Dy ₆ -Dy ₇), 3 μ ₂ <i>syn-syn</i> (Dy ₈ -Dy ₉ , Dy ₉ -Dy ₁₀ , Dy ₆ -Dy ₇), 2 chelate (Dy ₆ , Dy ₇)
Oxalate	Chelate (Dy ₄)	Chelate (Dy ₄)	Chelate (Dy ₄)	—	—
H ₂ O	1 terminal (Dy ₂)	3 terminal (Dy ₁ , Dy ₂ , Dy ₅)	1 terminal (Dy ₅)	2 terminal (Dy ₃ , Dy ₅)	2 terminal (Dy ₈ , Dy ₁₀)
CH ₃ OH	2 terminal (Dy ₄ , Dy ₅)	None	None	None	None
PO ₃	P1 - 2 μ ₂ (Dy ₁ -Dy ₂ , Dy ₁ -Dy ₅), 1 terminal (Dy ₄), P2 - 1 μ ₂ (Dy ₁ -Dy ₂), 2 terminal (Dy ₄ , Dy ₅)	P1 - 2 μ ₂ (Dy ₁ -Dy ₅ , Dy ₄ -Dy ₅), 1 terminal (Dy ₂), P2 - 2 μ ₂ (Dy ₁ -Dy ₂ , Dy ₂ -Dy ₃), 1 terminal (Dy ₅)	P1 - 2 μ ₂ (Dy ₁ -Dy ₅ , Dy ₄ -Dy ₅), 1 terminal (Dy ₂), P2 - 2 μ ₂ (Dy ₁ -Dy ₂ , Dy ₂ -Dy ₃), 1 terminal (Dy ₅)	P1 - 2 μ ₂ (Dy ₂ -Dy ₃ , Dy ₃ -Dy ₄), 1 terminal (Dy ₁)	P3 - 2 μ ₂ (Dy ₈ -Dy ₉ , Dy ₇ -Dy ₈), 1 terminal (Dy ₁₀), P4 - 1 μ ₂ (Dy ₉ -Dy ₁₀), 2 terminal (Dy ₆ , Dy ₈)
EDDC	2 N-O chelate (Dy ₁ , Dy ₃), 1 N-N chelate (Dy ₂), 2 N-O chelate (Dy ₁ , Dy ₄), 1 N-N chelate (Dy ₅), carbonyl bridges (Dy ₁ -Dy ₂ , Dy ₁ -Dy ₅ , Dy ₄ -Dy ₅)	2 N-O chelate (Dy ₁ , Dy ₃), 1 N-N chelate (Dy ₂), 2 N-O chelate (Dy ₁ , Dy ₄), 1 N-N chelate (Dy ₅), carbonyl bridges (Dy ₁ -Dy ₂ , Dy ₁ -Dy ₅ , Dy ₂ -Dy ₃ , Dy ₄ -Dy ₅)	2 N-O chelate (Dy ₁ , Dy ₃), 1 N-N chelate (Dy ₂), 2 N-O chelate (Dy ₁ , Dy ₄), 1 N-N chelate (Dy ₅), carbonyl bridges (Dy ₁ -Dy ₂ , Dy ₁ -Dy ₅ , Dy ₂ -Dy ₃ , Dy ₄ -Dy ₅)	2 N-O chelate (Dy ₂ , Dy ₄), 1 N-N chelate (Dy ₃), 2 N-O chelate (Dy ₂ , Dy ₅), 1 N-N chelate (Dy ₁), carbonyl bridges (Dy ₂ -Dy ₃ , Dy ₂ -Dy ₅ , Dy ₃ -Dy ₅ , Dy ₅ -Dy ₁)	2 N-O chelate (Dy ₉ , Dy ₇), 1 N-N chelate (Dy ₈), 2 N-O chelate (Dy ₉ , Dy ₆), 1 N-N chelate (Dy ₁₀), carbonyl bridges (Dy ₉ -Dy ₁₀ , Dy ₁₀ -Dy ₆ , Dy ₉ -Dy ₈ , Dy ₉ -Dy ₇)
Coordination bonds	42 (3 × 8 + 2 × 9)	43 (2 × 8 + 3 × 9)	44 (1 × 8 + 4 × 9)	42 (3 × 8 + 2 × 9)	41 (4 × 8 + 1 × 9)
Coordination geometries	Dy ₁ - triangular dodecahedron (D _{2d}), Dy ₂ - Hula-hoop (C _{2v}), Dy ₃ - Snub diphenooid J84 (D _{2d}), Dy ₄ - spherical capped square antiprism (C _{4v}), Dy ₅ - Hula-hoop (C _{2v})	Dy ₁ - triangular dodecahedron (D _{2d}), Dy ₂ - Hula-hoop (C _{2v}), Dy ₃ - spherical capped square antiprism (C _{4v}), Dy ₄ - Hula-hoop (C _{2v})	Dy ₁ - square antiprism (D _{4d}), Dy ₂ - Hula-hoop (C _{2v}), Dy ₃ - biaugmented trigonal prism (C _{2v}), Dy ₄ - spherical capped square antiprism (C _{4v}), Dy ₅ - Hula-hoop (C _{2v})	Dy ₁ - spherical tricapped trigonal prism (D _{3h}), Dy ₂ - Hula-hoop (C _{2v}), Dy ₃ - square antiprism (D _{4d})	Dy ₄ - spherical capped square antiprism (C _{4v}), Dy ₅ - Hula-hoop (C _{2v}), Dy ₆ - square antiprism (D _{4d})

magnetization rises abruptly in a low field then slowly reach the maximum values of 53.1 (I), 51.4 (I-UV) 49.9 (I-A-Ar) and 59.2 μ_B (II) at 1.8 K. The magnetizations do not saturate at 70 kOe and reach values which are lower than the expected 100 μ_B for ten Dy^{III}, and are associated with the considerable crystal-field effects.^{45,46} Furthermore, the non-saturation of the magnetization together with the non-superimposed M vs. H/T curves (Fig. S20†) suggests the presence of significant magnetic anisotropy and/or low-lying excited states that might be populated when a dc field is used.^{47–49}

Alternating-current (ac) susceptibility measurements were carried out on polycrystalline samples in the temperature range 1.8 to 26 K without a dc field in the frequency range 1 to 1000 Hz to investigate the dynamics of the magnetization reversal.²¹

The behaviours of I, I-UV and I-A-Ar are drastically different from one another. I exhibits a lack of slow dynamics with clear frequency independence of both χ' (in-phase) and χ'' (out-of-phase) (Fig. 4a). This is speculatively associated with facile tunnelling allowing demagnetization without overcoming the barrier.

However, upon transforming I to I-UV by exposure to 365 nm light for 3 days strong frequency dependent ac-susceptibilities are now observed indicating that slow relaxation is completely integrated (Fig. 4b, S21 and S22†). The Cole–Cole plots (χ'' vs. χ')⁵⁰ are quasi-semicircles fitting well to a generalized Debye model between 3.0 and 7.6 K (Fig. 4c, inset and Table S9†). To

quantify the effective energy barrier to magnetization reversal, the relaxation time was obtained from the fit of the frequency-dependent data between 1.8 and 7.6 K. The plot of $\ln \tau$ versus $1/T$ shows three distinct regions based on an Orbach process of high temperatures, a Raman process for intermediate temperatures and a QTM process at lower temperatures.^{51–54} These data follow the function including the three above-mentioned relaxation processes, $\tau^{-1} = AT + B + CT^n + \tau_0^{-1} \exp(-U_{\text{eff}}/kT)$,⁵⁵ with the following parameters: $U_{\text{eff}} = 47.3$ K (32.8 cm^{-1}), $\tau_0 = 1.26 \times 10^{-7}$ s, $A = 3.58$ $\text{s}^{-1} \text{K}^{-1}$, $B = 0.0036$ s^{-1} , $C = 0.0011$ $\text{s}^{-1} \text{K}^{-5.21}$, $n = 5.21$ and $\alpha \approx 0.15$ (Fig. 4c). The α parameter indicates a narrow distribution of slow relaxation times (τ).^{56,57} This drastic change of behaviour may be brought about by the different geometrical changes of the coordination spheres of the five Dy atoms of the asymmetric unit.

Upon annealing I-UV under Ar at 100 °C the crystal is transformed to I-A-Ar and its ac-susceptibilities are reverted back to being non-SMM. When I-A-Ar is allowed to stand in air under ambient conditions, it regains its SMM characteristics, albeit with slight modification of its metrics, $U_{\text{eff}}/k_B = 37.6$ K (26.1 cm^{-1}), $\tau_0 = 1.04 \times 10^{-7}$ s, $A = 2.23$ $\text{s}^{-1} \text{K}^{-1}$, $B = 0.0031$ s^{-1} , $C = 0.0009$ $\text{s}^{-1} \text{K}^{-4.79}$, $n = 4.79$ and $\alpha \approx 0.19$ (Fig. 4e and f). This means that annealing at 100 °C can be reverted by standing under ambient conditions but I-UV cannot be reverted back to I. This corroborates well with the crystal structures which indicates that annealing by heating at 100 °C removes

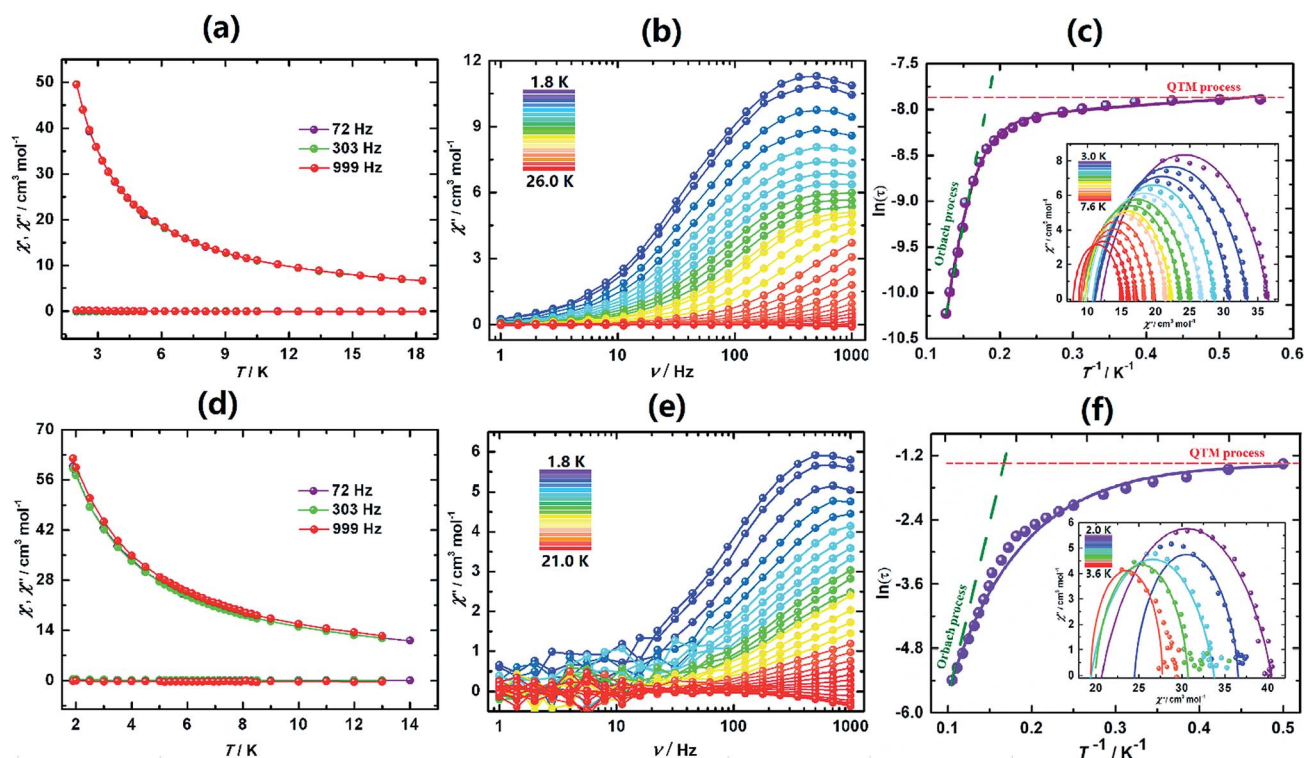


Fig. 4 Magnetic properties of I in its different forms. (a) Temperature dependence of the ac-susceptibilities of I for three different frequencies. (b) Frequency dependence of the out-of-phase ac susceptibility of I-UV and (c) Arrhenius plots of the relaxation rate and its fitting (see the text) for I-UV; inset: Cole–Cole plot and fits at different temperatures. (d) Temperature dependence of the ac-susceptibilities of I-A-Ar for three different frequencies. (e) Frequency dependence of the out-of-phase ac susceptibility of I-A-Ar exposed to air and (f) Arrhenius plots of the relaxation rate and its fitting (see the text) for I-A-Ar exposed to air; inset: Cole–Cole plot and fits at different temperatures.



two water molecules and it takes up the water reversibly upon standing in air.

The ac susceptibilities of **II** (Fig. 5, S25 and S26†) exhibit progressive frequency dependence characteristics of SMM. Analyses of the data led to characteristic parameters, $U_{\text{eff}} = 138 \text{ K}$ (95.9 cm^{-1}), $\tau_0 = 2.79 \times 10^{-7} \text{ s}$, $A = 2.62(3) \text{ s}^{-1} \text{ K}^{-1}$, $B = 0.19(6) \text{ s}$, $C = 0.0014 \text{ s}^{-1} \text{ K}^{-1}$ and $n = 4.83(5)$. The energy barrier is much higher than that of **I-UV**.

The contrasting relaxation processes of the different structures of compound **I** are quite astonishing and have not been seen previously. Although **I** is a non-SMM, its UV irradiated form **I-UV** behaves as a SMM and its heat annealed form under N_2 or argon **I-A** is reverted back to a non-SMM and **II** behaving as a SMM, and they all contain pentameric units which are structurally related. Therefore, the fusion of two pentamers with an oxalate bridge in **I**, **I-UV** and **I-A** cannot be the cause of the switching compared to the isolated pentamers in **II** which is a SMM.

The fortuitous advantage of having the above process with complete single-crystal transformation to single-crystals allows for a thorough structure–property correlation. The differences in the magnetic dynamics may be correlated with the different crystal field environments with the distinct coordination modes of the peripheral assistant ligands for crystallographically independent Dy^{III} ions. Careful analysis of the three cyclic arrangements reveals important disparities (Fig. 2). One piece of evidence is that the cyclic pentanuclear unit of **I** is surrounded by six acetate groups adopting three different coordination modes (Fig. 2a), while three of the six acetate groups in **I-UV** show one kind of crossfade based on the terminal and chelate modes (Fig. 2b). Only one coordination mode (bidentate fashion) can be observed for the four peripheral acetate groups of **I-A-N}_2** (Fig. 2c). On the other hand, the coordination spheres of the central metal ions of **I** are completed by one water and two methanol molecules, while it has three terminal water molecules in **I-UV** and only one water molecule in **I-A-N}_2**. Obviously, these structural modifications are most likely to influence the emergence of magnetic anisotropy through alteration of some crucial parameters of the structure (such as Dy–O/N bond lengths and Dy–O–Dy angles), and the coordination geometries of Dy^{III} ions (see Table S1† for details), hence

causing the drastic difference of magnetization dynamic behaviours.

In order to get an insight into the magneto-structure relationship, theoretical investigation was conducted for all these complexes. Complete-active-space self-consistent field (CASSCF) calculations on individual Dy^{III} fragments for complexes **I**, **I-UV**, **I-A-N}_2**, **II-molecule A** and **II-molecule B** on the basis of X-ray determined geometries have been carried out by the CASSCF/RASSI method with the MOLCAS 8.2 program package^{58–60} (see the ESI† for details). The Single_Aniso^{61–63} program was used to obtain the g tensors, energy levels, magnetic axes, *etc.*, based on the above CASSCF/RASSI calculations. The lowest eight spin-orbit energies, the corresponding g tensors and the m_j values of individual Dy^{III} fragments for complexes **I**, **I-UV**, **I-A-N}_2**, **II-molecule A** and **II-molecule B** are shown in Table S12,† where the calculated energy gaps of the lowest two Kramers doublets (KDs) and the ground g tensors of five or three types of individual Dy^{III} fragments for each complex are much different from each other. The tunnelling gaps of individual Dy^{III} fragments for each complex are also different from each other (Fig. S28–S32†). Thus, it is difficult to differentiate them only from the energy gaps of the lowest two KDs and the ground g tensors of individual Dy^{III} fragments for the five complexes. From the energy states and g tensors of the five complexes in Table S12,† however, we can find some differences between them. For **I**, the energy gap of the lowest two KDs of the Dy_5 fragment is only 15.1 cm^{-1} , much smaller than that of the others, and its ground g_z value is far from 20. For **I-UV**, both of the energy gaps of the lowest two KDs of five individual Dy^{III} fragments and the ground g_z values are larger than those of **I**, which can be one of the reasons for its showing SMM behaviour.

As usual, the energy barriers of the polynuclear lanthanide complexes are mainly from the single-ion anisotropy of the Dy^{III} centres. But the Dy–Dy interactions have some influence on the relaxation rate of incoherent quantum tunnelling to give drastically different effective relaxation barriers. For the current systems, it is difficult to fit the Dy–Dy interactions using the Lines model⁶⁴ since each complex has a large pair of interactions. However, the dipole–dipole interactions between Dy^{III} ions can be calculated exactly according to the ground g_z values and the magnetic axes shown in Fig. S33.† Simply, we only gave



Fig. 5 Magnetic properties of **II**. (a) Frequency and (b) temperature dependence of the out-of-phase ac-susceptibilities. (c) Arrhenius plots of the relaxation rate and its fitting (see the text); inset: Cole–Cole plot and fits at different temperatures.



- 34 V. Chandrasekhar, P. Bag and E. Colacio, *Inorg. Chem.*, 2013, **52**, 4562–4570.
- 35 L. G. Westin, M. Kritikos and A. Caneschi, *Chem. Commun.*, 2003, 1012–1013.
- 36 H. Tian, S.-S. Bao and L.-M. Zheng, *Dalton Trans.*, 2015, **44**, 14208–14212.
- 37 L. Zhao, S. Xue and J. Tang, *Inorg. Chem.*, 2012, **51**, 5994–5996.
- 38 (a) S. Biswas, S. Das, J. Acharya, V. Kumar, J. van Leusen, P. Kögerler, J. M. Herrera, E. Colacio and V. Chandrasekhar, *Chem.–Eur. J.*, 2017, **23**, 5154–5170; (b) X.-Y. Zheng, J.-B. Peng, X.-J. Kong, L.-S. Long and L.-S. Zheng, *Inorg. Chem. Front.*, 2016, **3**, 320–325; (c) X.-Y. Zheng, J. Xie, X.-J. Kong, L.-S. Long and L.-S. Zheng, *Coord. Chem. Rev.*, 2017, DOI: 10.1016/j.ccr.2017.10.023.
- 39 (a) H. Tian, S.-S. Bao and L.-M. Zheng, *Chem. Commun.*, 2016, **52**, 2314–2317; (b) H. Tian, S.-S. Bao and L.-M. Zheng, *Eur. J. Inorg. Chem.*, 2016, 3184–3190.
- 40 M. Pinsky and D. Avnir, *Inorg. Chem.*, 1998, **37**, 5575–5582.
- 41 O. Kahn, *Molecular Magnetism*, Wiley-VCH, New York, 1993.
- 42 P. H. Lin, T. J. Burchell, R. Clerac and M. Murugesu, *Angew. Chem., Int. Ed.*, 2008, **47**, 8848–8851.
- 43 L. Ungur, S. K. Langley, T. N. Hooper, B. Moubaraki, E. K. Brechin, K. S. Murray and L. F. Chibotaru, *J. Am. Chem. Soc.*, 2012, **134**, 18554–18557.
- 44 M. L. Kahn, R. Ballou, P. Porcher, O. Kahn and J. P. Sutter, *Chem.–Eur. J.*, 2002, **8**, 525–531.
- 45 S. Osa, T. Kido, N. Matsumoto, N. Re, A. Pochaba and J. Mrozinski, *J. Am. Chem. Soc.*, 2004, **126**, 420–421.
- 46 R. J. Blagg, L. Ungur, F. Tuna, J. Speak, P. Comar, D. Collison, W. Wernsdorfer, E. J. L. McInnes, L. F. Chibotaru and R. E. P. Winpenny, *Nat. Chem.*, 2013, **5**, 673–678.
- 47 P. Zhang, M. Perfetti, M. Kern, P. P. Hallmen, L. Ungur, S. Lenz, M. R. Ringenberg, W. Frey, H. Stoll, G. Rauhut and J. van Slageren, *Chem. Sci.*, 2018, **9**, 1221–1230.
- 48 E. Moreno Pineda, N. F. Chilton, R. Marx, M. Dörfel, D. O. Sells, P. Neugebauer, S.-D. Jiang, D. Collison, J. van Slageren, E. J. L. McInnes and R. E. P. Winpenny, *Nat. Commun.*, 2014, **5**, 5243–5251.
- 49 S. G. McAdams, A. Ariciu, A. K. Kostopoulos, J. P. S. Walsh and F. Tuna, *Coord. Chem. Rev.*, 2017, **346**, 216–239.
- 50 K. S. Cole and R. H. Cole, *J. Chem. Phys.*, 1941, **9**, 341–351.
- 51 I. J. Hewitt, J. Tang, N. T. Madhu, C. E. Anson, Y. Lan, J. Luzon, M. Etienne, R. Sessoli and A. K. Powell, *Angew. Chem., Int. Ed.*, 2010, **49**, 6352–6356.
- 52 J. J. Le Roy, M. Jeletic, S. I. Gorelsky, I. Korobkov, L. Ungur, L. F. Chibotaru and M. Murugesu, *J. Am. Chem. Soc.*, 2013, **135**, 3502–3510.
- 53 K. L. M. Harriman, J. L. Brosmer, L. Ungur, P. L. Diaconescu and M. Murugesu, *J. Am. Chem. Soc.*, 2017, **39**, 1420–1423.
- 54 B. S. Dolinar, D. I. Alexandropoulos, K. R. Vignesh, T. James and K. R. Dunbar, *J. Am. Chem. Soc.*, 2018, **140**, 908–911.
- 55 A. J. Brown, D. Pinkowicz, M. R. Saber and K. R. Dunbar, *Angew. Chem., Int. Ed.*, 2015, **54**, 5864–5868.
- 56 G. Abbas, Y. Lan, G. E. Kostakis, W. Wernsdorfer, C. E. Anson and A. K. Powell, *Inorg. Chem.*, 2010, **49**, 8067–8072.
- 57 Y. Bi, Y.-N. Guo, L. Zhao, Y. Guo, S. -Y. Lin, S. -D. Jiang, J. Tang, B. -W. Wang and S. Gao, *Chem.–Eur. J.*, 2011, **17**, 12476–12481.
- 58 F. Aquilante, L. De Vico, N. Ferré, G. Ghigo, P.-Å. Malmqvist, P. Neogrády, T. B. Pedersen, M. Pitonak, M. Reiher, B. O. Roos, L. Serrano-Andrés, M. Urban, V. Veryazov and R. Lindh, *J. Comput. Chem.*, 2010, **31**, 224.
- 59 V. Veryazov, P. -O. Widmark, L. Serrano-Andres, R. Lindh and B. O. Roos, *Int. J. Quantum Chem.*, 2004, **100**, 626.
- 60 G. Karlström, R. Lindh, P.-Å. Malmqvist, B. O. Roos, U. Ryde, V. Veryazov, P.-O. Widmark, M. Cossi, B. Schimmelpfennig, P. Neogrády and L. Seijo, *Comput. Mater. Sci.*, 2003, **28**, 222.
- 61 L. F. Chibotaru, L. Ungur and A. Soncini, *Angew. Chem., Int. Ed.*, 2008, **47**, 4126.
- 62 L. Ungur, W. Van den Heuvel and L. F. Chibotaru, *New J. Chem.*, 2009, **33**, 1224.
- 63 L. F. Chibotaru, L. Ungur, C. Aronica, H. Elmoll, G. Pilet and D. Luneau, *J. Am. Chem. Soc.*, 2008, **130**, 12445.
- 64 M. E. Lines, *J. Chem. Phys.*, 1971, **55**, 2977.
- 65 I. P. Beletskaya and M. A. Kazankova, *Russ. J. Org. Chem.*, 2002, **38**, 1391–1430.
- 66 *SHELXTL (version 5.0), Reference Manual*, Siemens Industrial Automation, Analytical Instruments, Madison, WI, 1995.
- 67 A. L. Spek, *J. Appl. Crystallogr.*, 2003, **36**, 7–13.

