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16653Single ion magnets based on lanthanoid
polyoxomolybdate complexes†José J. Baldoví,^{‡a} Yan Duan,^{‡a} Carlos Bustos,^{b,c} Salvador Cardona-Serra,^d
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Polyoxometalate (POM) chemistry has recently offered excellent examples of single ion magnets (SIMs) and molecular spin qubits. Compared with conventional coordination compounds, POMs provide rigid and highly symmetric coordination sites. However, all POM-based SIMs reported to date exhibit a very limited range of possibilities for chemical processability. We present herein two new families of POM-based SIMs which are soluble in organic solvents: $[\text{Ln}(\beta\text{-Mo}_8\text{O}_{26})_2]^{5-}$ ($\text{Ln}^{\text{III}} = \text{Tb, Dy, Ho, Er, Tm}$ and Yb) and the functionalised POMs $[\text{Ln}(\text{Mo}_5\text{O}_{13}(\text{OME})_4\text{NNC}_6\text{H}_4\text{-}p\text{-NO}_2)_2]^{3-}$ ($\text{Ln}^{\text{III}} = \text{Tb, Dy, Ho, Er, Yb}$ and Nd). In addition, these two families represent the first SIMs based on polyoxomolybdates. A magneto-structural analysis of these families is presented, which is based on an effective crystal field model, and compared with the results reported in analogous lanthanoid SIMs based on polyoxotungstates.

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Introduction

The main goal of spintronics is the active manipulation of the electron spin degrees of freedom in solid-state systems for carrying information.^{1,2} Spintronic systems have experienced a rapid development and currently are used in a range of applications, including read-heads devices and non-volatile magnetic memories (MRAM).^{3,4} Extraordinary potential for these systems is expected for the fabrication of spin-transfer nano-oscillators (STNOs)⁵ and quantum computers.^{6–8} The progress of molecular electronics and molecular magnetism has led to the emergence of a new field known as molecular spintronics,^{9,10} which combines the ideas and concepts of spintronics with the singular possibilities offered by molecular electronics and molecular magnetism to develop a second generation of spintronic devices.^{11–14}

A particularly challenging area within this field is that of single-molecule spintronics, which intends to use individual molecules as main components of spintronic devices. In this context, single-molecule magnets (SMMs)¹⁵ have been proposed as promising candidates.^{16–18} These systems, which represent the limit of miniaturization of nanomagnets, are between the most complex magnetic entities, exhibiting slow relaxation of the magnetization¹⁹ and magnetic hysteresis²⁰ at liquid-helium temperatures. Moreover, they may also present quantum phenomena from purely molecular origin.^{21–24} In this context, SMMs based on mononuclear lanthanide complexes deserve a special attention.^{25,26} The first example of this class of molecular nanomagnets, also known as single-ion magnets (SIMs), was reported by Ishikawa and co-workers in the series with general formula $[\text{LnPc}_2]^-$, where lanthanoid ions are sandwiched between two phthalocyaninato moieties displaying a square-antiprismatic D_{4d} symmetry.²⁷ Derivatives closely related to this family include the oxidised terbium phthalocyaninato complex $[\text{TbPc}_2]^0$. This system, besides its “double-decker” structure which favours adsorption on surfaces, is electrically neutral. This feature facilitates its sublimation under UHV conditions. The processability of lanthanoid phthalocyaninato complexes has allowed a series of breakthroughs including the realization of molecular/supramolecular spin valves^{28,29} and the electrical control of nuclear spin qubits.^{30,31} Indeed, sublimable lanthanoid complexes have demonstrated a great potential in molecular spintronics. Among the few sublimable lanthanoid-based systems reported so far, we can highlight the trinuclear Tb^{3+} complex

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Tb₃(OQ)₉ (OQ = quinolinato), where spin-polarised hopping transport has been realised.³² It has also been proposed as a model system for quantum error correction in quantum computing.³³

A second series of SIMs is that provided by polyoxometalate (POM) chemistry,³⁴ a class of systems that has produced a number of conventional SMMs.^{35,36} In this case, mononuclear lanthanoid complexes encapsulated by POMs have produced key examples of spin qubits for quantum computing. Indeed, extended quantum coherence was recently achieved on concentrated samples of the holmium derivative of the series [Ln(W₅O₁₈)₂]⁹⁻ (in short LnW₁₀) through the use of atomic clock transitions.³⁷ Employing a different strategy, a large number of coherent manipulations was realised in the Gd³⁺ derivative³⁸ of the [LnP₅W₃₀O₁₁₀]¹²⁻ series (in short LnW₃₀).³⁹ Within the same POM family, current rectification was achieved in a single molecule diode of DyW₃₀.⁴⁰ Still, the incorporation of these molecular inorganic polyanions onto surfaces or its anchoring to electrodes have been limited by their poor solubility in organic solvents and by their difficult functionalization with organic ligands.

Here, we have prepared two families of POM-based mononuclear lanthanide complexes with a rigid square antiprism structure, analogous to that of LnW₁₀, which overcome these processing limitations. The first family is formulated as [Ln(β-Mo₈O₂₆)₂]⁵⁻ (in short, LnMo₁₆), {Ln^{III} = Tb, Dy, Ho, Er, Tm and Yb} and consists of a lanthanide ion trapped by two [β-Mo₈O₂₆]⁴⁻ moieties.⁴¹ The second family is formulated as [Ln{Mo₅O₁₃(OMe)₄NNC₆H₄-*p*-NO₂}₂]³⁻ (in short, LnMo₁₀), {Ln^{III} = Tb, Dy, Ho, Er, Yb and Nd}. In this cases the lanthanide ion is trapped by a functionalised POM based on a lacunary Lindqvist-type pentamolybdate.⁴²

Results and discussion

Solubility and stability

LnMo₁₆ series. The LnMo₁₆ series are very soluble in acetone, acetonitrile and benzonitrile. Their derivatives are insoluble in methanol, ethanol, dichloromethane or chloroform, giving rise to colorless solutions. The structural integrity of the POMs in solution is kept as long as the solvents are dried, as shown by electrospray ionization for acetonitrile solutions (see Fig. S6†). Wet solvents, however, give rise to yellow solutions, evidencing decomposition.

LnMo₁₀ series. Owing to the structural analogy between the [Mo₅O₁₃(OCH₃)₄NNC₆H₄-*p*-NO₂]³⁻ units with their nitrosyl [Mo₅O₁₃(OCH₃)₄(NO)]³⁻ analogues, one could *a priori* expect the transformation of the LnMo₁₀ family of compounds in other solvents than methanol, in which they are synthesised. However they proved to be unexpectedly stable in several organic solvents at least at room temperature. Indeed, they are soluble and can be recrystallised in acetonitrile and chlorinated solvents (chloroform or dichloromethane) without noticeable decomposition, at the difference of their nitrosyl analogues.⁴³ The combination of the stability and solubility

properties of these compounds (the solution we used for their recrystallization are about 10⁻² mol L⁻¹) make them particularly interesting in the frame of their processing and utilization in nanodevices.

Crystal structures

LnMo₁₆ series. This family of compounds crystallise in two different space groups: monoclinic (*P*2₁/*c*) for Tb³⁺ and orthorhombic (*Pbca*) for the rest of the series. Because of the rigidity of the POM, this difference in the crystallographic space group is in practice not noticeable in the vicinity of lanthanide ion. Thus, in all cases the structure exhibits the presence of a lanthanide ion sandwiched between two [β-Mo₈O₂₆]⁴⁻ units. The lanthanide ions are coordinated by the eight terminal oxygen atoms coming from the two octamolybdate units giving rise to an eight-fold square antiprismatic coordination (Fig. 1). The structural parameters concerning the lanthanide coordination sphere are listed in Table 1. Details of the crystal structure determination and crystallographic parameters (Table S1) are given in the ESI.†

LnMo₁₀ series. As in the case of the LnMo₁₆ series, these complexes display a weakly distorted square antiprism geometry, in which the central Ln³⁺ cation interacts with two [Mo₅O₁₃(OMe)₄NNC₆H₄-*p*-NO₂]³⁻ units playing the role of

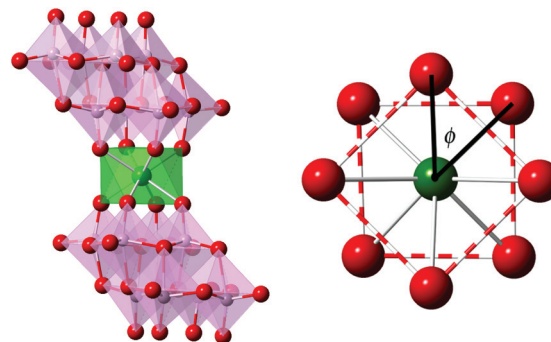


Fig. 1 (Left) Polyhedral and ball-and-stick representation of LnMo₁₆ series and (right) projection of the coordination sphere showing the square-antiprismatic coordination site.

Table 1 Structural parameters concerning the lanthanide coordination sphere

	LnMo ₁₆			LnMo ₁₀		
	<i>d</i> _{pp} ^a (Å)	<i>d</i> _{in} ^b (Å)	<i>φ</i> ^c (°)	<i>d</i> _{pp} ^a (Å)	<i>d</i> _{in} ^b (Å)	<i>φ</i> ^c (°)
Tb	2.604(13)	2.853(15)	40.2(5)	2.661(8)	2.816(8)	44.1(2)
Dy	2.552(18)	2.844(18)	44.1(5)	2.640(6)	2.797(6)	44.2(2)
Ho	2.553(10)	2.825(10)	44.3(3)	2.638(5)	2.786(5)	39.2(7)
Er	2.551(11)	2.819(11)	44.2(3)	2.614(6)	2.783(6)	40.0(6)
Tm	2.506(14)	2.818(14)	44.1(4)	—	—	—
Yb	2.506(14)	2.796(14)	44.3(4)	2.585(6)	2.759(6)	39.3(7)

^a *d*_{pp} defines the average distance between two oxygen based square planes. ^b *d*_{in} is the average O–O distance within the oxygen-based square planes. ^c *φ* is defined as the relative orientation between the two squares defined by the coordinating oxygen atoms.



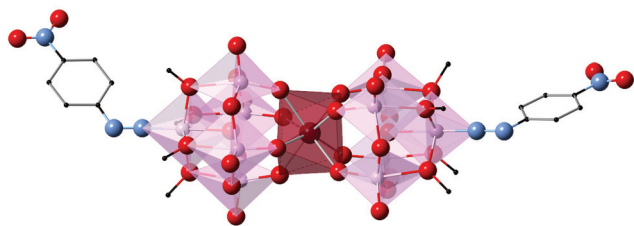


Fig. 2 Polyhedral and ball-and-stick representation of the LnMo₁₀ series.

planar tetradentate ligands, and providing an eight-coordination environment. The structural parameters concerning the lanthanide coordination sphere are listed in Table 1. The geometrical parameters of this series are comparable to those observed for the nitrosyl derivatives $[X\{Mo_5O_{13}(OMe)_4(NO)\}_2]^{n-}$ ($X = Ca^{2+}, Sr^{2+}, Ba^{2+}, Ce^{3+}, Eu^{3+}$ and Bi^{3+}),⁴⁴ thus indicating a slightly distorted D_{4d} geometry. The TbMo₁₀ crystal structure is shown in Fig. 2. Details of the crystal structure determination and crystallographic parameters (Table S2) are provided in the ESI.†

Magnetic properties

LnMo₁₆ series. A simultaneous fit was performed on the four magnetic susceptibility curves of the Tb³⁺, Dy³⁺, Ho³⁺ and Er³⁺ derivatives of this series using an effective crystal field approach (REC model; see section on experimental methods for details). The determined two parameters were tested by applying them to the real coordinates of the Tm³⁺ and Yb³⁺ derivatives. The predicted magnetic behaviour shows a very good agreement with the experimental data (Fig. 3).

The most satisfactory agreement was obtained when the radial displacement (D_r) equals 0.72 Å and the effective charge (Z_i) is 0.253. If we compare the obtained parameters with the ones extracted in the families LnW₁₀ and $[Ln(\beta_2-SiW_{11}O_{39})_2]^{13-}$

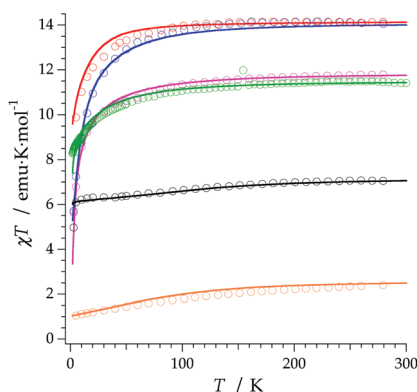


Fig. 3 Temperature-dependent magnetic susceptibility of the series LnMo₁₆ and theoretical calculations using the REC model: Dy³⁺ (red), Ho³⁺ (blue), Tb³⁺ (pink), Er³⁺ (green), Tm³⁺ (black), Yb³⁺ (orange). Markers give experimental data under a magnetic field of 1000 Oe, and solid lines theoretical results: fits for Tb³⁺–Er³⁺ and predictions for Tm³⁺ and Yb³⁺.

(in short: LnW₂₂),⁴⁵ the radial displacement is smaller in this case. This means that the effective point charge needs a smaller covalent correction to produce an adequate relation between the crystal-field parameters that describe the experimental data. This difference can be related to the different Pauling electronegativity of the Mo and W atoms (2.16 and 2.36 respectively).⁴⁶ The larger difference in electronegativity between Mo (2.16) and O (3.44) enhances the ionic character of the Mo–O bonding and also the coordination bond Ln–O.

The resulting energy levels for all the series are reported in the ESI (Fig. S10†). A general trend for LnMo₁₆ compared with LnW₁₀ and LnW₂₂ is a slightly larger crystal field splitting. In average, the Ln–O distance is practically identical in both series, e.g. in ErW₁₀ and ErMo₁₆ (2.367(7) Å and 2.367(3) Å). This would mean that polyoxomolybdates tend to produce a slightly stronger ligand field splitting compared with polyoxotungstates at a given metal–ligand distance. For a deeper analysis of the compounds that exhibit SMM properties in both families, the Stark sublevels of the HoMo₁₆ and ErMo₁₆ derivatives are plotted in Fig. 4. Regarding the lower states wave functions, one can observe that in the case of the HoMo₁₆ the ground state is defined by a mixture of 47% of $|+4\rangle$ and 47% of $|-4\rangle$. In this case, the crystal field operators enable a quantum tunnelling splitting between wave functions with a strong mixture of M_J components, and thus in absence of an external magnetic field no blocking of the magnetic moment is to be expected. Nevertheless when a longitudinal external field is applied, the purity of the $M_J = +4$ and $M_J = -4$ is recovered and thus, relaxation *via* tunnelling is cancelled and a blocking of the magnetization is possible. For ErMo₁₆ the wave function for the ground state appears to be $M_J = 0.78|\pm 1/2\rangle + 0.12|\pm 13/2\rangle$ with two states lying closely above it (at about 1.6 and 3.7 cm⁻¹). Those are described by $M_J = 0.78|\pm 15/2\rangle$ –marked in red in Fig. 4– and $M_J = 0.74|\pm 13/2\rangle + 0.15|\pm 15/2\rangle$ respectively. The low energy difference between the ground and the first excited state (and even the second) is definitively below the precision of the method. Also, thermal

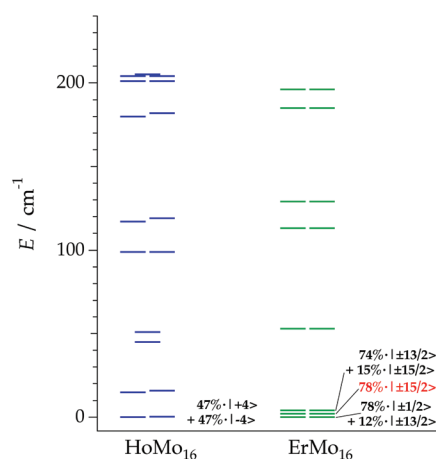


Fig. 4 Energy level scheme and main contributions to the ground state wave function for HoMo₁₆ and ErMo₁₆.



effects over the chemical structure, while only slightly affecting the energy level description,⁴⁷ are expected to become significant. This is expected to affect our modelling of the magnetization. As seen below, it is likely from the dynamic susceptibility behaviour that the Kramers doublet described by 78% of $\pm 15/2$ is the actual ground doublet. Within this assumption, the ground wave function would show a high M_J projection compatible with SMM behaviour.

The *ac* magnetic susceptibilities were collected in the range 2–12 K with an applied alternating field of 3.95 Oe at different frequencies in the range 1–10 000 Hz. ErMo₁₆ and HoMo₁₆ show frequency dependent out-of-phase signals at very low temperatures. In neither complex a clear peak is observed (Fig. S2 and S3†) even under an external applied field of 1000 Oe. (Fig. 5 and 6) This feature is a characteristic of the fast relaxation process that takes place at low temperatures. In none of the other derivatives an increase in the out-of-phase signal down to 2 K was observed.

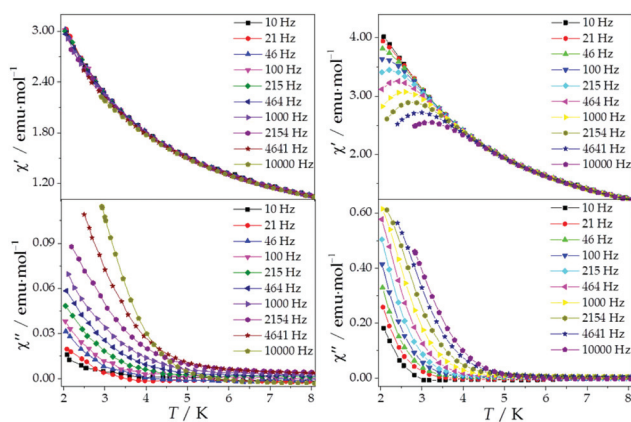


Fig. 5 In-phase (up) and out-of-phase (down) dynamic susceptibility of HoMo₁₆ (left) and ErMo₁₆ (right) with an applied field of 1000 Oe. The frequencies are shown in the legend. Solid lines are eye-guides.

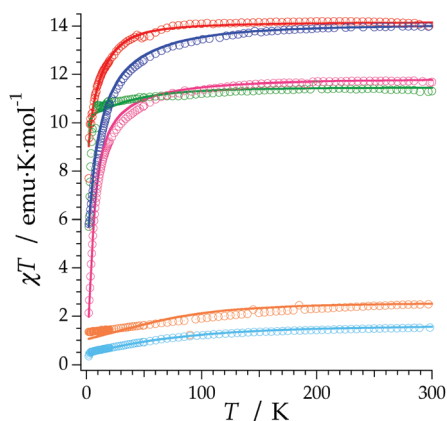


Fig. 6 Experimental (circles) and predicted (solid lines) χT product of the series LnMo₁₀ under a magnetic field of 1000 Oe: Dy³⁺ (red), Ho³⁺ (blue), Tb³⁺ (pink), Er³⁺ (green), Yb³⁺ (orange), Nd³⁺ (clear blue) from 2 to 300 K.

LnMo₁₀ series. For this series our initial prediction, based on the coordination geometry around the lanthanoid, pointed towards the possibility of obtaining SMM behaviour in the Dy derivative (97% of $M_J = \pm 11/2$), in Yb (97% of $M_J = \pm 5/2$) and in Er (99% of $M_J = \pm 15/2$) (see Experimental methods section). This was confirmed by the *ac* susceptibility data that showed slow relaxation of the magnetization in all three derivatives (Fig. 7, S4 and S5†).

The REC model permits to characterize the magnetic and spectroscopic properties of a family of complexes using a range of approaches. An extreme limit is the rough prediction with no free parameters,⁴⁸ which allows the discovery of new SIMs. On the opposite extreme, one can perform an individual fit for each metal, which describes the observables more accurately but has a more reduced ability to extrapolate the results to related derivatives. In this case, we have decided to employ an intermediate path. Owing to the chemical similarity of the two families of polyoxomolybdates presented in this work, especially in the coordination environment, the product of the parameters $D_r = 0.72 \text{ \AA}$ and $Z_i = 0.253$ obtained in the study of LnMo₁₆ was used to reduce the number of independent REC parameters for each metal when modelling the magnetic properties of LnMo₁₀. The magnetic susceptibility curves of this series were fitted individually keeping the relation $f = D_r \cdot Z_r$ constant, allowing just one free parameter per compound. The results are plotted in Fig. 6, where we can observe the excellent agreement between the model and the experiment. The sharp decrease that appears in ErMo₁₀ below 10 K is reminiscent of the behavior previously observed at the same temperature range in other complexes,^{49,50} such as [Er(COT)₂]⁻, [Er(COT'')₂]⁻ and Er(Cp*₂COT), where COT = cyclooctatetraenyl dianion, COT'' = 1,4-bis-(trimethylsilyl) cyclooctatetraenyl dianion and Cp* = pentamethylcyclopentadienide. As in the present case, in those examples the ground doublet can be characterized as an Ising state ($M_J = \pm 15/2$), and this may favour a stronger dipolar interaction at low temperature. The

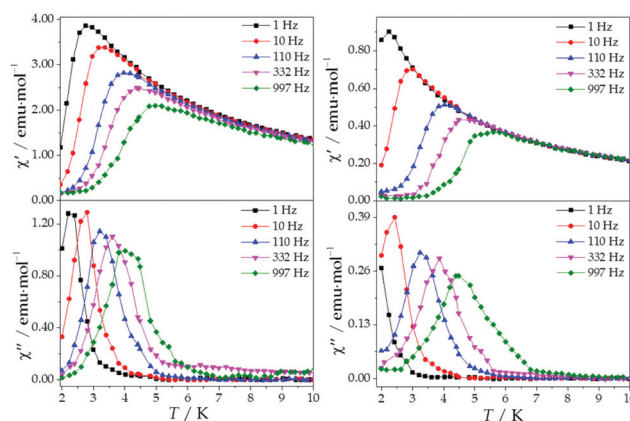


Fig. 7 In-phase (up) and out-of-phase (down) dynamic susceptibility of DyMo₁₀ (left) and YbMo₁₀ (right) under an applied field of 1000 Oe. The frequencies are shown in the legend. Solid lines are eye-guides.



estimation of the energy levels (Fig. S8†) and wave functions of this new family of polyoxomolybdates are reported in the ESI.†

An analysis of the frequency dependence of the χ'' peaks through Arrhenius plots (Fig. S6†) has permitted to estimate the magnetization-relaxation parameters. Compared with analogous polyoxotungstates, polyoxomolybdates display poorer behaviour as SIMs. Out of the whole series, Dy^{3+} , Er^{3+} and Yb^{3+} complexes can be characterised as SIMs (displaying clear peaks above 2 K) and only under an external dc field $H = 1000$ G. Under these conditions, a naïve fitting of the ac data of the Dy^{3+} , Yb^{3+} derivatives assuming an Orbach process would result in a barrier height ($U_{\text{eff}}/k_{\text{B}}$) of 38.5 K with a pre-exponential factor (τ_0) of 6.6×10^{-8} s for the Dy^{3+} case and 23.3 K with $\tau_0 = 6.7 \times 10^{-6}$ s for the Yb^{3+} case, compared with the archetypical ErW_{10} ($U_{\text{eff}}/k_{\text{B}}$) = 55.2 K and $\tau_0 = 1.6 \times 10^{-8}$ s. These plots do not contain enough information to independently fit a Raman process, but this should not be taken as a proof of an Orbach mechanism, especially lacking spectroscopic indication of levels at these energies. This is even more marked for the Er^{3+} derivative, which displays only two clear χ'' peaks, precluding an Arrhenius fit.

The calculated energy level schemes for DyMo_{10} , ErMo_{10} and YbMo_{10} are represented in Fig. 8. According to the individual magnetic susceptibility data fits based on the predictions of the REC model, DyMo_{10} possesses a ground state dominated by $M_J = \pm 11/2$ (86%) with an excited state of $M_J = 0.92|\pm 9/2\rangle$, which is placed at about 11 cm^{-1} above it. Such scheme is compatible with the observed SMM behaviour. One can notice that in this example all the M_J states are bunched below 100 cm^{-1} with the exception of the highest $M_J = 15/2$ which is located at 260 cm^{-1} . In ErMo_{10} , the ground state is characterised by 99% of $M_J = 15/2$ and the first excited Kramers doublet is placed at about 19 cm^{-1} . In the case of YbMo_{10} , the level scheme also supports the relationship between the ground-state wave function (described by $M_J = 97\%$ of $|\pm 5/2\rangle$) and the SMM behaviour. The rest of the energy levels appear at about 167 cm^{-1} above it, starting with the $M_J = 0.69|\pm 3/2\rangle$. The absence of mixing provides a quasi-pure

M_J function in the ground state without a clear path to invert its moment, thus the magnetic moment may be blocked at low temperatures. Here one again needs to highlight the risk of an Orbach-only interpretation in absence of richer experimental information.

Magneto-structural correlations

The optimization of f-element single-molecule magnets is still an open problem.²⁶ Thus, at this point, it is worth to briefly compare the results of a given lanthanide ion as a function of the different ligand fields created in the different families of POMs showing an approximate D_{4d} symmetry. For that, we will focus in the erbium analogue because its crystal structure has been determined for ErW_{10} , ErW_{22} , ErMo_{16} and ErMo_{10} . Regarding the lowest energy levels, we found a ground state characterised by $\pm 13/2$ in both polyoxotungstates, and a first excited state dominated by $\pm 1/2$ (placed at about 16 cm^{-1} in ErW_{10} and 11 cm^{-1} in ErW_{22}). In the case of ErMo_{10} , the ground doublet is apparently dominated by $\pm 15/2$ (99%) and the first excited state (located at about 19 cm^{-1}) presents 85% of $\pm 1/2$, according to our theoretical calculations based on available susceptibility data. The explanation of such a different energy scheme can be attributed to differences in the first coordination sphere, as well as differences in the effective charges and distances of the oxygen donor atoms. Thus, while in the two polyoxotungstate series the interplanar distances, d_{pp} , are equal to 2.46 \AA in ErW_{10} and 2.48 \AA in ErW_{22} , in the ErMo_{10} this distance increases to a value of 2.61 \AA . ErMo_{16} represents an intermediate situation, having $d_{\text{pp}} = 2.55 \text{ \AA}$ and a skew angle comparable to that of ErW_{10} (44.2°). In this case, these inexpensive calculations indicate that there are three Kramers doublets very close in energy, two of them dominated by high M_J values and one dominated by $\pm 1/2$, being difficult to determine the true nature of the ground Kramers doublet.

Conclusions

In the last years, a sufficiently large set of lanthanoid-based SIMs has already been obtained. This has provided a good basis for relating their magnetic properties with the nature and structure of the coordination site around the lanthanide ion. A subsequent development that has been recently initiated is that related with the improvement in the chemical processability of these magnetic molecules in order to incorporate them into functional devices.

The present work contributes towards these two goals. Thus, we have used a robust family of ligands based on POMs to prepare new SIMs exhibiting SMM properties, which can be modelled using an effective crystal field approach previously developed for this kind of ligands. The key novelty has been the unprecedented use of polyoxomolybdates instead of polyoxotungstates for obtaining SIMs. While the chemistry of polyoxomolybdates is more challenging, this result has opened the possibility of making these magnetic molecules soluble in organic solvents, thus facilitating their processing and

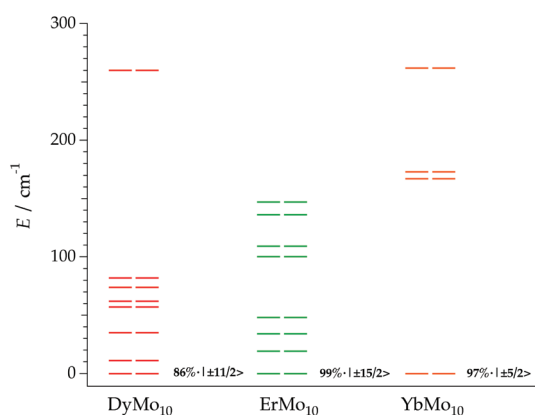


Fig. 8 Energy level scheme and main contributions to the ground state wave function for DyMo_{10} , ErMo_{10} and YbMo_{10} .



incorporation into devices. This contrasts with the SIMs based on polyoxotungstate ligands, which are commonly only soluble in water. Furthermore, we have also shown that polyoxomolybdate-based SIMs can be functionalised with organic groups, facilitating their processability -e.g. grafting onto surfaces/electrodes – and the incorporation of a second property coming from the organic ligand. Last but not least, the enhanced reduction potential of polyoxomolybdates may also be of interest to incorporate delocalised electrons into the mixed-valence POM framework, thus increasing the interest of these magnetic molecules in spintronics.^{51–54}

Experimental methods

All reagents and solvents used were commercially available. Acetonitrile was distilled under argon before use. The precursor for the LnMo₁₆ series: [TBA]₄[β-Mo₈O₂₆], where TBA = (*n*-C₄H₉)₄N was prepared according to a previously described literature procedure.⁵⁵

The precursor [TBA]₃[Mo₆O₁₈NN-C₆H₄-*p*-NO₂] of the LnMo₁₀ series was prepared as described elsewhere.⁵⁶ The subsequent synthesis of the ligand [TBA]₂[Mo₅O₁₃(MeO)₄(NNC₆H₄-*p*-NO₂){Na(MeOH)}] was slightly modified from the initial report (see below).⁴²

Synthesis of LnMo₁₆ series

The salts [TBA]₅[Ln(β-Mo₈O₂₆)₂] (Ln^{III} = Tb, Dy, Ho, Er, Tm, and Yb) were prepared following a slight modification of a previously described procedure.⁴³ A solution of lanthanide salt Ln(NO₃)₃ (0.200 mmol) in 5 mL of acetonitrile was added drop-wise to a solution of [TBA]₄[β-Mo₈O₂₆] (0.376 mmol) in 25 mL of acetonitrile under stirring. Then, 50 mL of diethyl ether were added carefully, in such a way that two layers were formed. After three days of slow diffusion, block-shaped crystals formed and were collected by filtration, washed with a small amount of acetonitrile, and dried under vacuum overnight (see ESI† for details of the elemental analysis and yield IR spectra).

Synthesis of LnMo₁₀ series

Synthesis of the ligand [TBA]₂[Mo₅O₁₃(MeO)₄(NNC₆H₄-*p*-NO₂){Na(MeOH)}]. [TBA]₃[Mo₆O₁₈NN-C₆H₄-*p*-NO₂] (1.740 g, 1.000 mmol) was introduced in 15 mL of methanol and 5 mL of 0.2 mol L⁻¹ solution of NaOH in methanol (1 mmol) was then added. The mixture was heated in refluxing methanol (65 °C) for 1 hour, leading to a red limpid solution. This solution was then allowed to cool to room temperature and filtered on a glass frit (por. 4). Diethyl-ether (20 mL) was added to the resulting filtrate, which was then placed at -18 °C for one day. The red crystals thus obtained were filtered, rinsed with 3 portions of 5 mL of diethyl-ether and dried in air, yielding 1.200 g of [TBA]₂[Mo₅O₁₃(MeO)₄(NNC₆H₄-*p*-NO₂){Na(MeOH)}].

Synthesis and crystallization of LnMo₁₀ series. [TBA]₂[Mo₅O₁₃(MeO)₄(NNC₆H₄-*p*-NO₂){Na(MeOH)}] (0.376 g, 0.250 mmol) was dissolved in 15 mL of methanol. 0.125 mmol

of the lanthanide salt {TbCl₃·6H₂O: 0.048 g; ErCl₃·6H₂O: 0.0342 g; YbCl₃·6H₂O: 0.048 g; Dy(NO₃)₃·5H₂O: 0.048 g; Nd(NO₃)₃·6H₂O: 0.055 g; Ho(NO₃)₃·5H₂O: 0.055 g} was then added. The resulting mixture was refluxed in methanol for 30 min. The red/orange precipitate formed after cooling the solution to room temperature was filtered on a glass frit (por. 4) and washed twice with diethyl-ether. All compounds were recrystallised by slow diffusion of diethyl-ether in a solution in chloroform (*ca.* 0.300 g in 15 mL of CHCl₃), leading to the formation of dark red crystalline materials with general formula [TBA]₃[Ln{Mo₅O₁₃(MeO)₄NNC₆H₄-*p*-NO₂}₂].1.5CHCl₃.

Magnetic measurements

Magnetic static and dynamic measurements of samples of LnMo₁₆ and LnMo₁₀ were performed by compacted powder molded from ground crystalline samples. Variable-temperature susceptibility measurements were carried out under a magnetic field of 1000 Oe in the temperature range 2–300 K on a magnetometer equipped with a SQUID sensor (Quantum Design MPMS-XL-5). The data were corrected for diamagnetic contribution from eicosane and for the diamagnetic contributions of the polyanions as deduced by using the Pascal's constant tables. Isothermal magnetization measurements at low temperature (2 K) were performed up to a field of 5 T in the same apparatus (see ESI†).

Ligand field calculations

LnMo₁₆ series. The four magnetic susceptibility data of the Tb³⁺, Dy³⁺, Ho³⁺ and Er³⁺ polyanions were used to carry out a simultaneous fit of an effective ligand field hamiltonian. This was based on the Radial Effective Charge (REC) model, which uses effective point charges to model the ligand effects and thus estimate the crystal field parameters in mononuclear lanthanoid complexes (see details in ESI,† section 7).⁵⁷ For this purpose we employed the SIMPRE package, a freely available *fortran77* code.^{58,59} This procedure provided the two parameters, namely the radial displacement and the effective charge, that can be used to describe the ligand field effects of the eight chemically equivalent oxygens of the polyoxomolybdate ligands. As a verification, the magnetic susceptibility of the two remaining members of the series (Tm³⁺ and Yb³⁺) was predicted using the same parameters.

LnMo₁₀ series. At the time of our first contact with this family, we had not yet investigated the LnMo₁₆ derivatives with the REC model, and thus we performed an initial (predictive) estimation to evaluate the likelihood of obtaining SMM behaviour in some of the derivatives of this series. For this purpose, we introduced the crystal structures of the Tb³⁺, Ho³⁺ and Er³⁺ derivatives as input in SIMPRE, applying the two REC parameters (radial displacement: *D_r* = 0.895 Å and effective charge: *Z_i* = 0.105) that had been determined for LnW₁₀ and LnW₂₂ in a previous work.⁴⁵ On a second stage, we used the product of the radial displacement and the effective charge parameters obtained in the study of the series LnMo₁₆ (*f* = *D_r*·*Z_i* = 0.18216) in order to fit the magnetic data of the series LnMo₁₀ while



keeping constant f , allowing only one free parameter per compound.

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Notes and references

- S. A. Wolf, D. D. Awschalom, R. A. Buhrman, J. M. Daughton, S. Von Molnár, M. L. Roukes, A. Y. Chtchelkanova and D. M. Treger, *Science*, 2001, **294**, 1488–1495.
- M. E. Flatté, *Nature*, 2009, **462**, 419–420.
- J. Akerman, *Science*, 2005, **308**, 508–510.
- I. L. Prejbeanu, M. Kerekes, R. C. Sousa, H. Sibuet, O. Redon, B. Dieny and J. P. Nozières, *J. Phys.: Condens. Matter*, 2007, **19**, 165218.
- B. Georges, J. Grollier, M. Darques, V. Cros, C. Deranlot, B. Marcilhac, G. Faini and a. Fert, *Phys. Rev. Lett.*, 2008, **101**, 4–7.
- J. R. Petta, A. C. A. Johnson, J. Taylor, E. A. Laird, A. Yacoby, M. D. Lukin, C. M. Marcus, M. Hanson and A. Gossard, *Science*, 2005, **309**, 2180–2184.
- J. F. Gregg, *Nat. Mater.*, 2007, **6**, 798–799.
- K. L. Wang, I. Ovchinnikov, F. Xiu, A. Khitun and M. Bao, *J. Nanosci. Nanotechnol.*, 2011, **11**, 306–313.
- A. R. Rocha, V. M. García-Suárez, S. W. Bailey, C. J. Lambert, J. Ferrer and S. Sanvito, *Nat. Mater.*, 2005, **4**, 335–339.
- S. Sanvito, *Nat. Phys.*, 2010, **6**, 562–564.
- J. Camarero and E. Coronado, *J. Mater. Chem.*, 2009, **19**, 1678–1684.
- F. Prins, M. Monrabal-Capilla, E. A. Osorio, E. Coronado and H. S. J. Van Der Zant, *Adv. Mater.*, 2011, **23**, 1545–1549.
- J. Dugay, M. Giménez-Marqués, T. Kozlova, H. W. Zandbergen, E. Coronado and H. S. J. Van Der Zant, *Adv. Mater.*, 2015, **27**, 1288–1293.
- A. Holovchenko, J. Dugay, M. Giménez-Marqués, R. Torres-Cavanillas, E. Coronado and H. S. J. van der Zant, *Adv. Mater.*, 2016, DOI: 10.1002/adma.201600890.
- R. Sessoli, H. L. Tsai, A. R. Schake, S. Wang, J. B. Vincent, K. Folting, D. Gatteschi, G. Christou and D. N. Hendrickson, *J. Am. Chem. Soc.*, 1993, **115**, 1804–1816.
- L. Bogani and W. Wernsdorfer, *Nat. Mater.*, 2008, **7**, 179–186.
- W. Wernsdorfer, *Int. J. Nanotechnol.*, 2010, **7**, 497–522.
- E. Cremades, C. D. Pemmaraju, S. Sanvito and E. Ruiz, *Nanoscale*, 2013, **5**, 4751–4757.
- L. Thomas, F. Lioni, R. Ballou, D. Gatteschi, R. Sessoli and B. Barbara, *Nature*, 1996, **383**, 145–147.
- C. Paulsen, J.-G. Park, B. Barbara, R. Sessoli and A. Caneschi, *J. Magn. Magn. Mater.*, 1995, **140–144**, 1891–1892.
- A. Ardavan and S. J. Blundell, *J. Mater. Chem.*, 2009, **19**, 1754–1760.
- F. Troiani and M. Affronte, *Chem. Soc. Rev.*, 2011, **40**, 3119–3129.
- P. C. E. Stamp and A. Gaita-Ariño, *J. Mater. Chem.*, 2008, **19**, 1718–1730.
- J. Lehmann, A. Gaita-Arino, E. Coronado and D. Loss, *Nat. Nanotechnol.*, 2007, **2**, 312–317.
- D. N. Woodruff, R. E. P. Winpenny and R. A. Layfield, *Chem. Rev.*, 2013, **113**, 5110–5148.
- S. T. Liddle and J. van Slageren, *Chem. Soc. Rev.*, 2015, **44**, 6655–6669.
- N. Ishikawa, M. Sugita, T. Ishikawa, S. Y. Koshihara and Y. Kaizu, *J. Am. Chem. Soc.*, 2003, **125**, 8694–8695.
- M. Urdampilleta, S. Klyatskaya, J.-P. Cleuziou, M. Ruben and W. Wernsdorfer, *Nat. Mater.*, 2011, **10**, 502–506.
- M. Urdampilleta, S. Klyatskaya, M. Ruben and W. Wernsdorfer, *ACS Nano*, 2015, **9**, 4458–4464.
- S. Thiele, F. Balestro, R. Ballou, S. Klyatskaya, M. Ruben and W. Wernsdorfer, *Science*, 2014, **344**, 1135–1138.
- C. Wäckerlin, F. Donati, A. Singha, R. Baltic, S. Rusponi, K. Diller, F. Patthey, M. Pivetta, Y. Lan, S. Klyatskaya, M. Ruben, H. Brune and J. Dreiser, *Adv. Mater.*, 2016, **28**, 5195–5199.
- A. Bedoya-Pinto, H. Prima-García, F. Casanova, E. Coronado and L. E. Hueso, *Adv. Electron. Mater.*, 2015, 1500065.
- J. J. Baldoví, S. Cardona-Serra, J. M. Clemente Juan, L. Escalera-Moreno, A. Gaita-Ariño and G. Mínguez Espallargas, *Europhys. Lett.*, 2015, **110**, 33001.
- J. M. Clemente-Juan, E. Coronado and A. Gaita-Ariño, *Chem. Soc. Rev.*, 2012, **41**, 7464–7478.
- N. V. Izarova and P. Kögerler, *Polyoxometalate-based single-molecule magnets, Trends in Polyoxometalates Research*, 2015.
- M. Vonci and C. Boskovic, *Aust. J. Chem.*, 2014, **44**, 6655–6669.
- M. Shiddiq, D. Komijani, Y. Duan, A. Gaita-Ariño, E. Coronado and S. Hill, *Nature*, 2016, **531**, 348–351.
- J. J. Baldoví, S. Cardona-Serra, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño and H. Prima-García, *Chem. Commun.*, 2013, **49**, 8922–8924.
- S. Cardona-Serra, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño, A. Camón, M. Evangelisti, F. Luis,



- M. J. Martínez-Pérez and J. Sesé, *J. Am. Chem. Soc.*, 2012, **134**, 14982–14990.
- 40 S. Sherif, G. Rubio-Bollinger, E. Pinilla-Cienfuegos, E. Coronado, J. C. Cuevas and N. Agraït, *Nanotechnology*, 2015, **26**, 291001.
- 41 A. Kitamura, T. Ozeki and A. Yagasaki, *Inorg. Chem.*, 1997, **36**, 4275–4279.
- 42 C. Bustos, D. M.-L. Carey, K. Boubekeur, R. Thouvenot, A. Proust and P. Gouzerh, *Inorg. Chim. Acta*, 2010, **363**, 4262–4268.
- 43 R. Villanneau, A. Proust, F. Robert and P. Gouzerh, *Chem. – Eur. J.*, 2003, **9**, 1982–1990.
- 44 R. Villanneau, A. Proust, F. Robert and P. Gouzerh, *J. Chem. Soc., Dalton Trans.*, 1999, 421–426.
- 45 J. J. Baldoví, J. M. Clemente Juan, E. Coronado, Y. Duan, A. Gaita-Ariño and C. Giménez-Saiz, *Inorg. Chem.*, 2014, **53**, 9976–9980.
- 46 J. J. Baldoví, A. Gaita-Ariño and E. Coronado, *Dalton Trans.*, 2015, **44**, 12535–12538.
- 47 K. Qian, J. J. Baldoví, S.-D. Jiang, A. Gaita-Ariño, Y.-Q. Zhang, J. Overgaard, B.-W. Wang, E. Coronado and S. Gao, *Chem. Sci.*, 2015, **6**, 4587–4593.
- 48 J. J. Baldoví, Y. Duan, R. Morales, A. Gaita-Ariño, E. Ruiz and E. Coronado, *Chem. – Eur. J.*, 2016, DOI: 10.1002/chem.201601741.
- 49 K. R. Meihaus and J. R. Long, *J. Am. Chem. Soc.*, 2013, **135**, 17952–17957.
- 50 S. K. Singh, T. Gupta and G. Rajaraman, *Inorg. Chem.*, 2014, **53**, 10835–10845.
- 51 S. Cardona-Serra, J. M. Clemente-Juan, A. Gaita-Ariño, N. Suaud, O. Svoboda and E. Coronado, *Chem. Commun.*, 2013, **49**, 9621–9623.
- 52 S. Cardona-Serra, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño, N. Suaud, O. Svoboda, R. Bastardis, N. Guihéry and J. J. Palacios, *Chem. – Eur. J.*, 2015, **21**, 763–769.
- 53 C. Bosch-Serrano, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño, A. Palií and B. Tsukerblat, *ChemPhysChem*, 2012, **13**, 2662–2665.
- 54 C. Bosch-Serrano, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño, A. Palií and B. Tsukerblat, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **86**, 024432.
- 55 W. Klemperer, *Inorg. Synth.*, 2007, **27**, 74–85.
- 56 C. Bustos, B. Hasenknopf, R. Thouvenot, J. Vaissermann, A. Proust and P. Gouzerh, *Eur. J. Inorg. Chem.*, 2003, **15**, 2757–2766.
- 57 J. J. Baldoví, J. J. Borrás-Almenar, J. M. Clemente-Juan, E. Coronado and A. Gaita-Ariño, *Dalton Trans.*, 2012, **41**, 13705.
- 58 J. J. Baldoví, S. Cardona-Serra, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño and A. Palií, *J. Comput. Chem.*, 2013, **34**, 1961–1967.
- 59 J. J. Baldoví, J. M. Clemente-Juan, E. Coronado, A. Gaita-Ariño and A. Palií, *J. Comput. Chem.*, 2014, **35**, 1930–1934.

