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External stimuli modulate the magnetic relaxation of lanthanide single-molecule magnets

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Mutilfunctional molecular magnetic materials have promising potential for use as future molecular devices, such as for magneto-luminescence sensing, optical switching, information storage, *etc*. Lanthanide ions have been considered as excellent candidates to construct molecular switches as they are very sensitive to the subtle changes of structure including the change of molecular geometry, bond distance, even crystal packing, *etc*, while for transition metal ions, in most cases, only the change of electronic configuration can cause a switch of magnetic properties. This review aims to summarize the effects of stimuli-induced structural transformations on the magnetic relaxation involving light-, solvato-, proton- and redox-switching of Ln-SMM behaviour. Emphasis is placed on the role played by external stimuli in reversibly modulating the magnetic properties realized in the systems with single-crystal-to-single-crystal structural transformation.

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Introduction

Molecular magnetism, which started in the 1980s, is a multidisciplinary research field.[1] Distinguished from the traditional inorganic magnetic materials, molecular magnetic materials exhibit fascinating and valuable cooperative-electronic properties, *e.g.* combining magnetic properties with conductivity or optical phenomena.[2–4] In the past 27 years or so, single molecule magnets (SMMs) and multifunctional magnetic materials have been dominating the field for their practical applications in Quantum Information Technology (QIP) including molecule-based information storage, spintronic devices, quantum computing, *etc*.[5–10] In 2003, Ishikawa *et al.* reported the first example of lanthanide SMMs (Ln-SMMs),[11] *i.e.* double-decker lanthanide complexes $[Pc_2Ln][NBu_4]$ (Pc = phthalocynanine, Ln = Tb, Dy) containing only one magnetic ion, which motivated chemists to introduce lanthanides into SMMs due to their large magnetic anisotropy and strong spin–orbit coupling,[12,13] eventually leading to the discovery of magnetic hysteresis up to 80 K at a sweep rate of 25 Oe s^{-1} in a near-linear dysprosium metallocene compound

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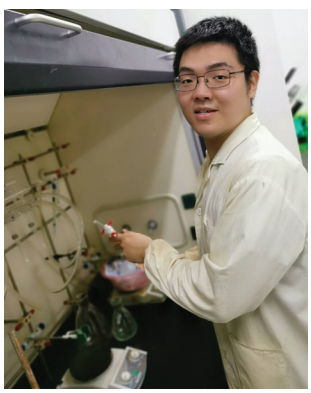
[(Cp^{iPr5})Dy(Cp*)][B(C$_6$F$_5$)$_4$] (Cp^{iPr5} = penta-iso-propylcyclopentadienyl, Cp* = pentamethylcyclopentadienyl).[14] As the 4f orbitals of lanthanides are well shielded by the outer 5s and 5p orbitals, a smaller crystal field (CF) splitting or weaker metal–ligand covalence is expected for lanthanides compared to d-block metals.[15] In most cases, the CF multiplet structure of lanthanides, which generally depends on the molecular geometry or the symmetry of the coordination environment, coordination numbers and the types of donor atoms, defines the magnetic properties (or magnetic relaxation) of the complexes.[12] The coordination geometries of lanthanides are determined by ligand steric factors therefore usually forming labile 'ionic' complexes that can undergo facile exchange of ligands, which provides more possibilities to modulate the magnetic properties. Taking Dy(III)-SMMs as an example, the perfect molecular geometries are C_n ($n > 7$), C_{5h}/D_{5h}, S_8/D_{4d} and S_{12}/D_{6d} where a pure axial CF was constructed based on the oblate spheroidal shape of the Dy(III) free-ion electron density.[16] Although it is difficult to make such ideal coordination environments for Dy(III) ions, building a *predominant bond* by lowering the coordination number or introducing anionic donor atoms with high electronegativity (*e.g.* F$^-$, O$^-$ or even carbanion) in complexes, which generally refers to the one with the shortest bond length, can also provide strong axial CF.[17] For a long time, we have been modeling magnetic relaxation of Dy(III)-SMMs on the general assumption of regarding the energy barrier as a sole criterion on the basis of an ideal condition where the temperature of the lattice is equal to that of the surrounding bath.[18] In recent years, consideration of other factors associated with molecular vibrations, such as coupling of anharmonic phonons and spin-phonons that addresses the presence of alternative relaxation processes, has been required.[19–21] Therefore, the design criteria for high-temperature Dy(III)-SMMs have been evolving, from the traditional requirements of high axial symmetry of CF based on the electrostatic repulsion interactions between lanthanides and ligand atoms to the new trend of considering the effects of molecular vibrational modes on magnetic relaxation.[20,22] In addition, the magnetic relaxation mechanism has also made great progress especially the quenching of the relaxation through the first excited state and the recognition of the vital importance of two-phonon Raman relaxation process.[23–25]

Apart from Ln-SMMs, in recent years, multifunctional molecular magnetic materials showing peculiar functionalities in addition to and/or interacting with the magnetic properties, such as microporous magnets, conducting magnets, chiral magnets, luminescent magnets *etc.*, have also attracted more and more attention in the molecular magnetism community due to their potential applications in sensors, optical switches, magnetic gas separations, *etc.*[26–32] So far, while numerous reviews or books have been published on Ln-SMMs and multifunctional molecular magnetic materials, however, to the best of our knowledge, the scope of stimuli-responsive Ln-SMMs has been far less highlighted.[33] In fact, the SMM behaviour is just a manifestation of the stimuli from alternating current (ac) magnetic field. Compared to the external stimuli modulated transition metal (TM) molecular magnets, the research of stimuli-responsive Ln-SMMs is still in its infancy. For example, the light-induced excited spin state trapping phenomenon or the LIESST effect has been a focus of research over 30 years; especially in the past ten years, the introduction of a photoisomerizable diarylethene-derived ligand into a spin-crossover iron(II) complex led to the construction of magneto-optical molecular devices with reversible photomagnetic conversions.[34–39] In 2009, the first example of photochemical control of SMMs was reported in a [Mn$_4$] one-dimensional (1D) chain structure and four years later, the light-actuated SMM combined with a spin-crossover phenomenon was also fabricated.[40–42] However, for Ln-SMMs, the efficient photomodulation of magnetic relaxation has not been realized until recently (*vide infra*).[43] Another typical example is that the first combination of high-pressure X-ray crystallography and magnetic measurements on TM-molecular magnets was reported



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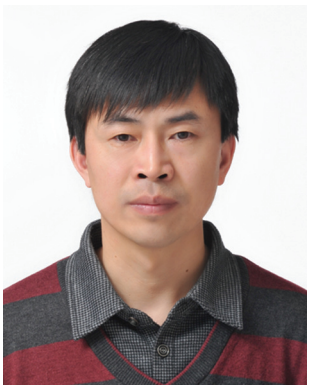

Jinkui Tang

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in 2008, while a similar investigation on Ln-SMMs was published in 2020.[44,45]

Although light-actuated TM-molecular magnets have become a fascinating topic in recent years, most Fe(II) complexes hardly exhibited good SMM properties due to the non-Kramers systems and the quenching of the fist-order spin–orbit coupling (SOC) unless they possess a strict symmetry of CF.[46] In contrast, Co(II) complexes usually displayed impressive SMM properties with a record effective energy barrier (U_{eff}) of 413 cm^{-1}; however, the fact that Co(II) ions can quench the luminescence of organic ligands prevents the design of photo-modulated SMMs.[47,48] In this connection, some significant advances in stimuli-responsive Ln-SMMs have been made owing to the unique 4f orbitals of lanthanides. In this review, we focus on the manipulations of the magnetic relaxation of Ln-SMMs through external stimuli generally classified as physical stimuli (such as light, temperature, pressure, electric potential) and chemical influences (such as redox, pH-responsive, guest-exchange), shedding light on the utilization of SMMs in molecular devices.

Photo-switching of Ln-SMM behaviour

Light as an extremely attractive tool in the field of molecular materials due to its easy availability, environmental friendliness, sustainability, *etc*. has garnered substantial attention. Photomagnets refer to a very special class of multifunctional molecular magnetic materials where the magnetization can be controlled by photons.[50] Until now, the most widely used strategy for designing and synthesizing photo-switchable Ln-SMMs is to introduce a photochromic ligand on account of its light-driven reversible transformation between two isomeric forms. The most formidable challenge to this approach is that the light-driven structural changes hardly alter the first coordination environment of lanthanide ions, which is closely related to the magnetic anisotropy and slow relaxation of magnetization of Ln-SMMs, leading to only a few examples being known. However, these structural changes often lead to modulation of electronic properties, polarity, conjugation, molecular geometry, physical dimensions, intermolecular interactions, *etc*.,[51] which provide other ways to externally manipulate the SMM behaviour.

Since 2011, the Yamashita group has used a diarylethene photochromic ligand (DTE) to explore the photo-control of magnetic interactions and slow relaxation of magnetization in Dy(III)/Ho(III)-complexes. Although there are some differences in the sub-level of the ground state and quantum-tunneling relaxation between the open form and the closed form of DTE, the unclear photo-control mechanism and the unsatisfactory SMM properties attributed to the lack of single-crystal-to-single-crystal (SC–SC) structural transformation and the reasonable CF environment prevent further investigations.[52–54]

Recently, the Long group reported a photochromic chain of Dy(III)-SMM based on a highly magnetic anisotropic compound, $[Dy(Tp^{py})(pyridine)_2]PF_6$ (Tp^{py} = tris(3-(2-pyridyl)pyrazolyl)hydroborate), using a bispyridyl dithienylethene photochromic ligand as the photoactive unit (Fig. 1a).[49,55] Compared to previous efforts, the main breakthroughs in this contribution are (1) the magnetic unit has a significantly improved SMM property due to the presence of a strong axial magnetic anisotropy provided by a relatively short Dy–F bond; (2) the SC–SC isomerization was achieved upon visible irradiation, which is the essential prerequisite for *ab initio* calculations, providing possibilities to explore the mechanism of photo-modulated SMM behaviour (Fig. 1b). Photocrystallography investigations showed that continuous irradiation of a single crystal of compound **1c** (closed form) of 1 h with green light resulted in obvious variations of the unit cell parameters with a small increase of 1.3% and 2.75% in *b* and *c* lattice parameters, respectively, and a decrease of 1.5% in the *a* lattice parameter (Fig. 1c). However, only partial conversions of **1o** (open form) toward **1c** with 365 nm light were observed, suggesting a more preferred packing rearrangement in **1o**. In order to investigate the influences of photoisomerization on the magnetic properties, static (dc) and dynamic (ac) magnetic measurements were performed for compounds **1c** and **1o**, and the results revealed that the peak in out-of-phase susceptibility of **1c** was observed as high as 44 K under a zero dc field giving the U_{eff} of 157 cm^{-1}, which displayed almost identical magnetic behaviour to **1o** due to the intact first coordination environment. The most striking difference between **1c** and **1o** was the quantum-tunneling relaxation (τ_{tunnel}), 0.0337 s for **1c** while 0.0015 s for **1o**. The significantly decreased τ_{tunnel} value of **1o** was also reflected by the narrower hysteresis hoop, for which a slightly stronger nearest-neighbor dipolar coupling in **1o** than **1c** induced by the subtle changes in crystal packing should be responsible, evident by *ab initio* calculations.

Although the photo-modulated magnetic hysteresis was observed in the above compounds, the On/Off switching of SMM behaviour has not been realized. In 2020, the Wang group reported a series of chain compounds, $[Ln_3(HEDP^{3-})_3(HEDP^{2-})_3]\cdot 2TPT^{3+}\cdot HEDP\cdot 10H_2O$ (QDU-1(Ln)) (Ln = Dy, Gd, Y; HEDP = hydroxyethylidene diphosphonate, TPT = 2,4,6-tri(4-pyridyl)-1,3,5-triazine), prepared *via* solvothermal reactions, among which, the radical-actuated On/Off SMM behaviour *via* room temperature (RT) light irradiation was achieved for the first time.[43] Note that the TPT ligand is in the form of pyridine salt with three positive charges. As shown in Fig. 2a, the oxygen atoms from HEDP serve as electron donors, while the nitrogen atoms in TPT units play the role of electron acceptors showing obvious H-bonding interactions between them. When illuminated by UV light in air, the crystal, QDU-1Dy, quickly turned blue, named QDU-1a(Dy), from colorless. QDU-1a(Dy) displayed a very stable state when stored below −20 °C and can return to the initial colorless state (QDU-1b(Dy)) by heating to 100 °C for 0.5 h (Fig. 2b). UV-vis, fluorescence, and ESR spectra confirmed the presence of photoinduced electron transfer (PET) from HEDP-Dy(III) chains to TPT ligands, during which the stable photogenerated anionic $O^{\bullet}$ and cationic $TPT^{\bullet}$ radicals were observed upon

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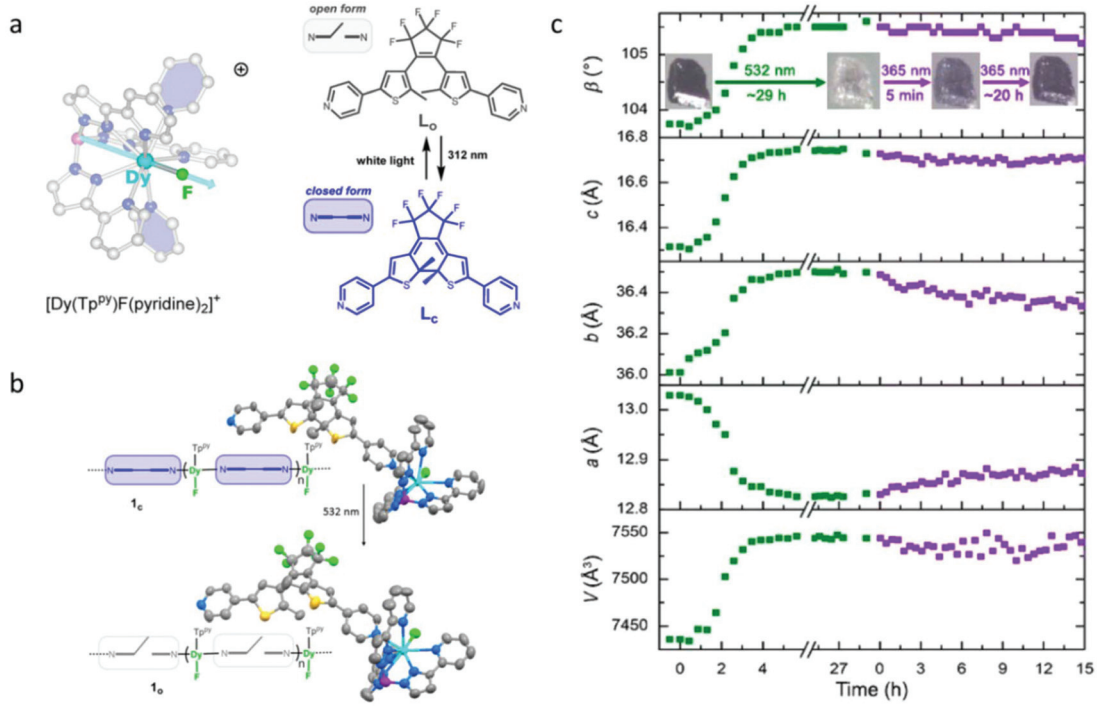

Fig. 1 (a) Molecular structure of $[Dy(Tp^{py})(pyridine)_2]PF_6$ in the cationic form and the open and closed form of the photoactive bispyridyl dithienylethene unit used in building **1c**; (b) single crystal structures of **1c** and **1o**; (c) evolution of the unit cell parameters of **1c** upon continuous irradiation with 532 nm light (green dots) followed by 365 nm light (purple dots). Reproduced with permission from ref. 49. Copyright 2020 American Chemical Society.[49]

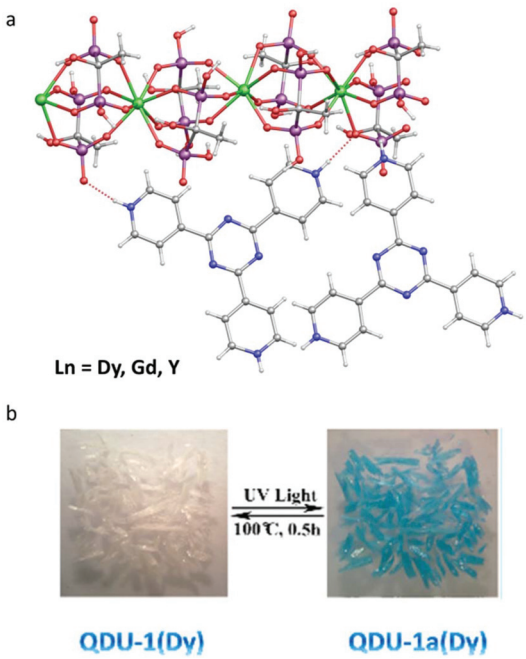

Fig. 2 (a) Crystal structures of QDU-1(Ln); (b) reversible photochromic behaviour between QDU-1(Dy) and QDU-1a(Dy). Reproduced with permission from ref. 43. Copyright 2020 American Chemical Society.[43]

irradiation of QDU-1(Dy). More importantly, only electron transfer process took place before and after irradiation, namely the main structure was maintained without photo-induced photolysis or isomerization, proved by the unchanged XRD, IR spectra and elemental analysis. The single crystal structures of QDU-1(Dy) and QDU-1a(Dy) confirmed this PET process with an obvious volume reduction of the unit cell and a closer N···O distance between the HEDP-Ln chain and TPT ligand.

Static and dynamic magnetic measurements were carried out to explore the difference in magnetic properties before and after irradiation. Expectedly, QDU-1a(Dy) showed a χT value of 43.67 $cm^3\ mol^{-1}$ K at 300 K, slightly larger (3%) than the value of QDU-1(Dy) indicating the presence of photogenerated radicals. As shown in Fig. 3a, QDU-1(Dy) exhibited a gradual decrease of the χT value suggesting weak interactions; however, a steady increase of the χT value for QDU-1a(Dy) was observed, reaching up to 52.67 $cm^3\ mol^{-1}$ K at 2 K, demonstrating the presence of strong ferromagnetic interactions between Dy(III) ions and anionic $O^{\bullet}$ radicals, which were also confirmed by the isostructural Y(III) and Gd(III) complexes. Once heated, QDU-1b(Dy) showed a similar trend of χT values toQDU-1(Dy), that is to say, the manipulation of On/Off magnetic interactions was realized. In addition, a more exciting result is that the out-of-phase susceptibility of QDU-1a(Dy) exhibited strong temperature dependency with clear χ'' peaks under a zero-dc field corresponding to the typical SMM behav-

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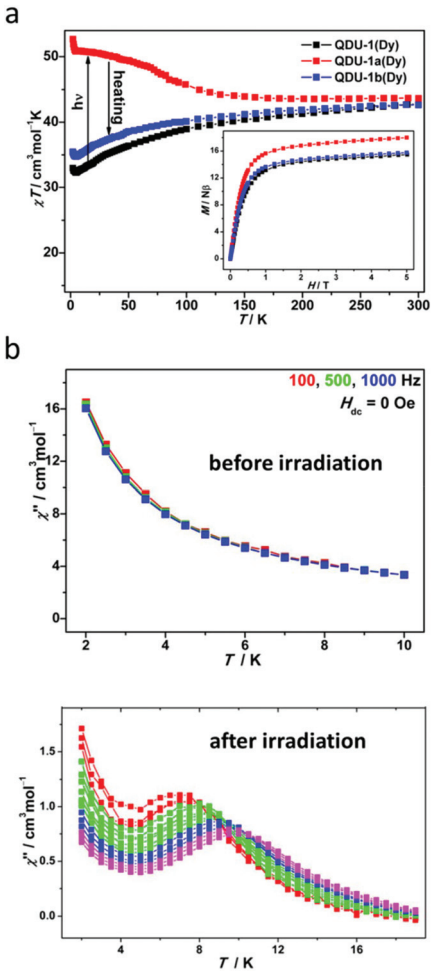

Fig. 3 (a) Temperature dependence of $\chi_m T$ values for QDU-1(Dy), QDU-1a(Dy) and QDU-1b(Dy); (b) plots of $\chi''(T)$ in the absence of a dc field at various ac frequencies for QDU-1(Dy) (up) and QDU-1a(Dy) (bottom). Reproduced with permission from ref. 43. Copyright 2020 American Chemical Society.[43]

iour, which can be attributed to the combination of large anisotropy of Dy(III) ions and strong ferromagnetic coupling between photogenerated anionic O$^{\bullet}$ radicals and Dy(III) ions induced by the PET process, while only slow relaxation of magnetization was observed in QDU-1(Dy) (Fig. 3b).

In addition, photoactive ligands including 1,2-bis(4-pyridyl) ethene (bpe) and anthracene (or its derivatives) were also used to construct photo-responsive Ln-SMMs. Although the SC–SC transformation was achieved in bde-based Ln-SMMs (Ln = Dy, Er) by the photodimerized [2 + 2] cycloaddition reaction of two bde ligands upon UV irradiation, this physical process was not reversible.[56–58] In 2018, the Zheng group reported a mononuclear Dy(III)-compound, Dy(depma)$(NO_3)_3$(hmpa)$_2$ (hmpa = hexamethylphosphoramide, depma = 9-diethylphosphonomethylanthracene), exhibiting a reversible SC–SC transformation in anthracene-based lanthanide complexes.[59] When the yellow crystal of this compound was irradiated at 365 nm for 0.5 h, a colorless crystal of $Dy_2(depma_2)(NO_3)_6(hmpa)_4$ without loss of crystallinity was obtained involving the [4 + 4] cycloaddition reaction of the neighboring anthracene moieties and meanwhile accompanied by a profound change in magneto-optic properties, *i.e.* from a single-ion magnet with yellow-green emission to a single-molecule magnet with blue-white emission. A completely reversed transformation occurred if this dinuclear compound was heated at 100 °C in air, opening a new route for designing photo-switchable smart molecular materials.

Solvato-switching of Ln-MOF-SMM behaviour

Compared to photomodulation, thermal treatment may be a more effective physical stimulus for SMM systems, during which, the desolvation or solvation process induced phase transformation usually took place resulting in the changes of the coordination environment of lanthanides therefor affecting the magnetic properties. Herein, we only discuss Ln-MOF-SMMs due to the generally higher structural stability of MOFs than low-dimensional coordination compounds providing greater possibilities to achieve reversible SC–SC structural transformation. In 2013, the Shi group investigated the magnetic behaviour of a very classic MOF system with relatively high thermal stability, Dy(BTC)(H_2O)·DMF and Dy(BTC) (BTC = 1,3,5-benzenetricarboxylate), which was first reported by Qiu and co-workers in 2006.[60,61] As shown in Fig. 4a, when Dy(BTC)(H_2O)·DMF was heated under vacuum at 240 °C for 12 h, the terminal coordinated water molecules and DMF were removed producing Dy(BTC), accompanied by the change of coordination geometry of Dy(III) ions from seven-coordinate pentagonal-bipyramidal to six-coordinate distorted trigonal

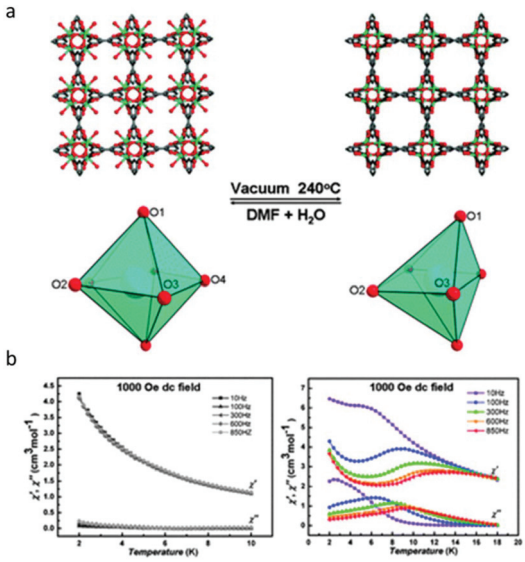

Fig. 4 (a) The structural or coordination geometry changes of the framework and Dy(III) ions; (b) temperature dependence of χ' and χ'' for Dy(BTC)(H_2O)·DMF (left) and Dy(BTC) (right). Reproduced with permission from ref. 61. Copyright 2013 Royal Society of Chemistry.[61]

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prism. If Dy(BTC) was immersed in the mixture of DMF and H_2O, it returned to Dy(BTC)(H_2O)·DMF as confirmed by single crystal X-ray diffraction, XRD, IR and elemental analysis. Ac susceptibility measurements showed that Dy(BTC) is a field-induced SMM while no χ'' signals were observed for Dy(BTC)(H_2O)·DMF under a 1000 Oe dc field (Fig. 4b), so the reversible switching of SMM behaviour by the desolvation and solvation process in a Ln-MOF system was achieved for the first time.

In the same year, the Maji group reported a Dy(III)–K(I) oxalate framework, $\{KDy(ox)_2(H_2O)_4\}_n$ (ox = oxalate dianion), where each Dy(III) ion is octacoordinated possessing a distorted square antiprismatic coordination geometry.[62] Static magnetic measurements suggested the presence of ferromagnetic interactions operating between the ox-connected Dy(III) centers in the system. When it was heated at 130 °C for 6 h under vacuum, the H_2O molecules bonded to K(I) ions were removed producing a dehydrated framework, $\{KDy(ox)_2\}_n$, in which the coordination environment of Dy(III) ions was unchanged. However, the antiferromagnetic interaction was observed in $\{KDy(ox)_2\}_n$ confirmed by the magnetically diluted sample $\{KDy_{0.25}Y_{0.75}(ox)_2\}_n$ and if it was exposed to H_2O vapor for three days, the rehydrated framework was obtained showing the ferromagnetic interactions again. The result indicated that in this Ln-MOF system, the reversible ferromagnetic to antiferromagnetic phase transition can be completed by the dehydration and rehydration process. However, the SMM properties have hardly changed when the magnetic interactions were switched to antiferromagnetic from ferromagnetic attributed to the almost unchanged D_{4d} coordination geometry of Dy(III) ions (U_{eff}: 417 K for $\{KDy(C_2O_4)_2(H_2O)_4\}_n$; 418 K for $\{KDy(C_2O_4)_2\}_n$).

The above two examples inspired us that the change of the first coordination environment of Dy(III) ions can induce the switching of SMM behaviour more clearly. In 2019, the Ohkoshi group reported a bimetal cyanido-bridged Dy–Co MOF system, $\{[Dy(III)(H_2O)_2][Co(III)(CN)_6]\}\cdot 2.2H_2O$, where each Dy(III) ion is eight-coordinate (N_6O_2) with square antiprism geometry (D_{4d}) (Fig. 5).[63] The single crystal of its dehydrated compound, $\{Dy(III)[Co(III)(CN)_6]\}$, was obtained by heating $\{[Dy(III)(H_2O)_2][Co(III)(CN)_6]\}\cdot 2.2H_2O$ to *ca.* 470 K under the nitrogen flow in which the coordination environment of Dy(III) ions was changed to trigonal prism geometry (D_{3h}). In addition, during this phase transformation, water of crystallization that occupied the interstitial space in the hydrated compound was also removed completely; meanwhile the space group changed to *P*63/*mmc* of a hexagonal crystal system from *Cmcm* of an orthorhombic crystal system. When keeping a crystal of $\{Dy(III)[Co(III)(CN)_6]\}$ for several days under a humid atmosphere, a rehydrated compound with excellent crystallinity was obtained showing the isostructural characteristics to the hydrated compound. For magnetic properties, no obvious $\chi''(\nu)$ signals were observed in the hydrated compound under a zero dc field or even after the application of the external dc magnetic field. In contrast, the dehydrated compound exhibited the typical SMM behaviour with clear maximum values of $\chi''(\nu)$ under a zero or 2000 Oe dc field giving the corresponding U_{eff} of 52 K and 58 K, respectively. The result proved that dehydration driven phase transformation between $\{[Dy(III)(H_2O)_2][Co(III)(CN)_6]\}\cdot 2.2H_2O$ and $\{Dy(III)[Co(III)(CN)_6]\}$ leads to the reversible ON/OFF switching of SMM behaviour.

Proton-switching of Ln-SMM behaviour

In this section, we introduce two typical examples of proton switching of Ln-SMM behaviour. The first one is a tetraphenylporphyrin (TPP)-based Tb(III) double-decker complex reported by Ishikawa and collaborators in 2012 (Fig. 6).[64] Single crystal structure analysis showed that the Tb(III) ion is hepta-coordinate with seven Tb–N bond lengths in the range of 2.430(7)–2.627(7) Å in the protonated complex, [Tb(III)H(TPP)$_2$], where a proton is located on a nitrogen atom of the TPP pyrrole ring leading to a significantly longer distance between the Tb(III)

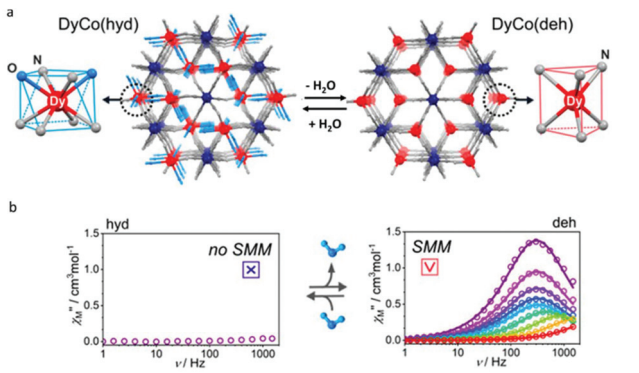

Fig. 5 (a) Crystal structures and (b) plots of $\chi''(\nu)$ in the absence of a dc field for $\{[Dy(III)(H_2O)_2][Co(III)(CN)_6]\}\cdot 2.2H_2O$ (left) and $\{Dy(III)[Co(III)(CN)_6]\}$ (right). Reproduced with permission from ref. 63. Copyright 2019 American Chemical Society.[63]

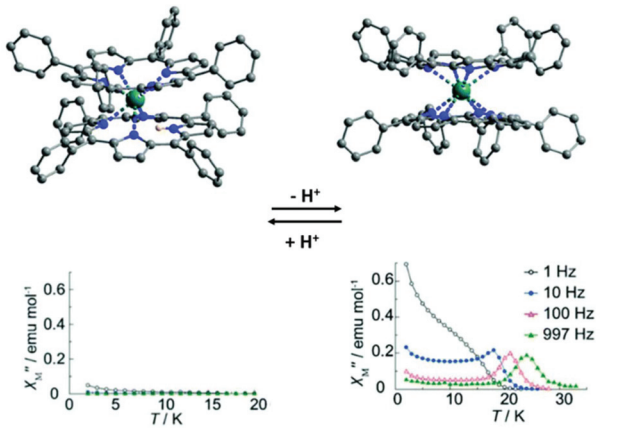

Fig. 6 Crystal structures (up) and plots of $\chi''(T)$ in the absence of a dc field (bottom) for the protonated (left) and the deprotonated (right) compounds. Reproduced with permission from ref. 64. Copyright 2012 Royal Society of Chemistry.[64]

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ion and N atom of 2.840(6) Å. When DBU (a base as an organic proton acceptor) is added to the solution of this protonated complex, deprotonation occurs and the complex, [Tb(III)(TPP)$_2$(H-DBU)], was obtained in which the Tb(III) ion possesses a square antiprism coordination geometry with eight identical Tb–N bond lengths of 2.513(4) Å. On the other hand, if the acetic acid is added to the solution of [Tb(III)(TPP)$_2$(H-DBU)], the protonation occurs, giving the protonated complex again. The protonation/deprotonation process is reversible as corroborated by UV-vis and IR spectra as well as cyclic voltammetry (CV) investigations. Ac magnetic susceptibility measurements were performed to study the effects of the protonation/deprotonation process on the magnetic relaxation. The results indicated that the protonated complex is not a SMM under a zero dc field without obvious $\chi''(\nu)$ signals at a frequency of 1000 Hz and temperature of 2 K; however, the deprotonated complex exhibited a typical SMM behaviour under a zero dc field with the U_{eff} of 269 cm^{-1} due to the high anisotropy of CF experienced by Tb(III) ions.

The other example of proton-controlled Ln-SMM behaviour is a polyoxometalate (POM)-based lanthanide complex containing a dinuclear dysprosium core reported by the Mizuno group in 2013.[65] POM 1 ($[\{Dy(H_2O)_2(CH_3COCH_3)\}_2(\gamma\text{-}SiW_{10}O_{36})_2]^{10-}$) synthesized by the reaction of $TBA_4H_4[\gamma\text{-}SiW_{10}O_{36}]$ (TBA = tetra-*n*-butylammonium) with Dy(acac)$_3$ (acac = acetylacetonato) contains two isolated Dy(III) ions and each one is seven-coordinate with four Dy–O bonds from the $[\gamma\text{-}SiW_{10}O_{36}]^{8-}$ unit and the remaining three one from two inner water molecules and one outer acetone molecule possessing a distorted monocapped trigonal prism geometry, as shown in Fig. 7. When TBAOH (2 equiv.) was added to the dichloromethane solution containing POM 1 and molecular sieve 3A and this system was kept at 313 K for 1 day, colorless crystals of POM 2 ($[Dy_2(\mu_2\text{-}OH)_2(\gamma\text{-}SiW_{10}O_{36})_2]^{12-}$) were obtained due to the successive dehydration condensation between two Dy(III) ions.

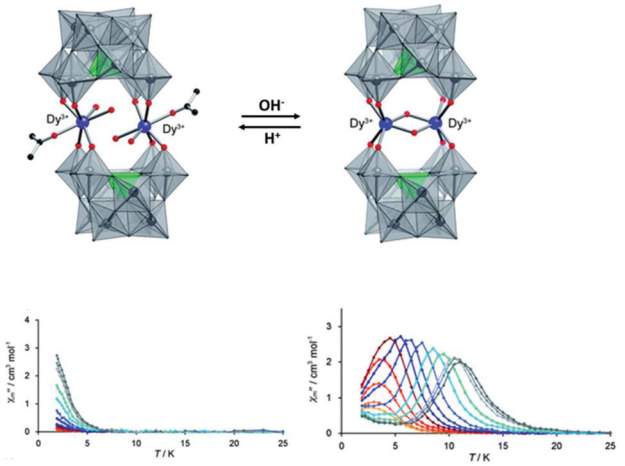

Fig. 7 (Up) Reversible SC–SC transformation between POM 1 and POM 2; (bottom) plots of $\chi''(T)$ in the absence of a dc field for POM 1 (left) and POM 2 (right). Reproduced with permission from ref. 65. Copyright 2013 Royal Society of Chemistry.[65]

Single crystal X-ray diffraction showed that these two Dy(III) ions in POM 2 are bridged by two μ_2-OH ligands and each one is six-coordinate with four Dy–O bonds from the $[\gamma\text{-}SiW_{10}O_{36}]^{8-}$ unit and two from the μ_2-OH ligand resulting in the change of coordination geometry from a distorted monocapped trigonal prism to distorted trigonal prism. More importantly, this chemical process is reversible benefitting from the excellent stability of the POM ligand, that is to say, if two equiv. HNO_3 and a small amount of water are added to a solution mixture of acetone and acetonitrile of POM 2, the bridging ligand μ_2-OH will be eliminated to form POM 1. For magnetic properties, only weak slow relaxation of magnetization was observed in POM 1 under a zero dc field due to the fast quantum-tunneling relaxation process while the typical SMM behaviour with clear frequency-dependent relaxation peaks was observed in POM 2 giving a U_{eff} of 65.7 K.

Redox-switching of Ln-SMM behaviour

In the field of molecular magnetism, the most known redox-active ligand is Pc derivatives due to the revolutionary discovery of the Pc-based Ln-SMMs in 2003. Four years later, Ishikawa and collaborators synthesized an anionic complex, $[\{Pc(OEt)_8\}_2Tb]^-(nBu_4N)^+$ and a cationic complex, $[\{Pc(OEt)_8\}_2Tb]^+(SbCl_6)^-$, based on the electrically neutral complex, $[\{Pc(OEt)_8\}_2Tb]^0$ through a redox reaction.[66] Note that the modification of Pc with an octaethoxy group ensures a moderate solubility. Although every compound has D_{4d} symmetry, compared to the anionic complex, a longitudinal contraction of the coordination space of Tb(III) ions in the cationic complex was observed, leading to a significant increase of U_{eff} (8%) accompanied by an extension of the χ'' peak from 40 K to 52 K at a frequency of 1000 Hz. In 2010, the Veciana group used magnetic circular dichroism (MCD) spectroscopy to study their SMM behaviour and the results showed that a greater coercive field for $[\{Pc(OEt)_8\}_2Tb]^+$ than $[\{Pc(OEt)_8\}_2Tb]^-$ was observed, which was consistent with the results of ac magnetic susceptibility measurements.[67]

In addition to Pc derivatives, other redox-active ligands such as tetraoxolene and tetrathiafulvalene were also used in the design of electro-active Ln-SMMs. However, as far as we are aware, most oxidized or reduced species based on these two ligands were not isolated in the solid state preventing possible simultaneous investigation of SMM behaviour and redox activity.[68–71] In 2018, J. van Slageren and collaborators succeeded in obtaining an anionic radical-bridged dinuclear Dy(III) complex $[(HBpz_3)_2Dy(\mu\text{-}CA^{\bullet})Dy(HBpz_3)_2]^-[CoCp_2]^+$ (1Dy) (HBpz = hydrotris(pyrazol-1-yl)borate, $CA^{3\bullet-}$ = chloroanilate) in the solid state with enough purity by chemical reduction of the neutral compound $[(HBpz_3)_2Dy(\mu\text{-}CA)Dy(HBpz_3)_2]\cdot 2CH_2Cl_2$ (2Dy) using cobaltocene as the reducing agent.[72] Although the low quality of crystal limited its crystal structure, its chemical formation was confirmed by ESI-MS (positive and negative mode), IR spectra and elemental analysis. Static magnetic

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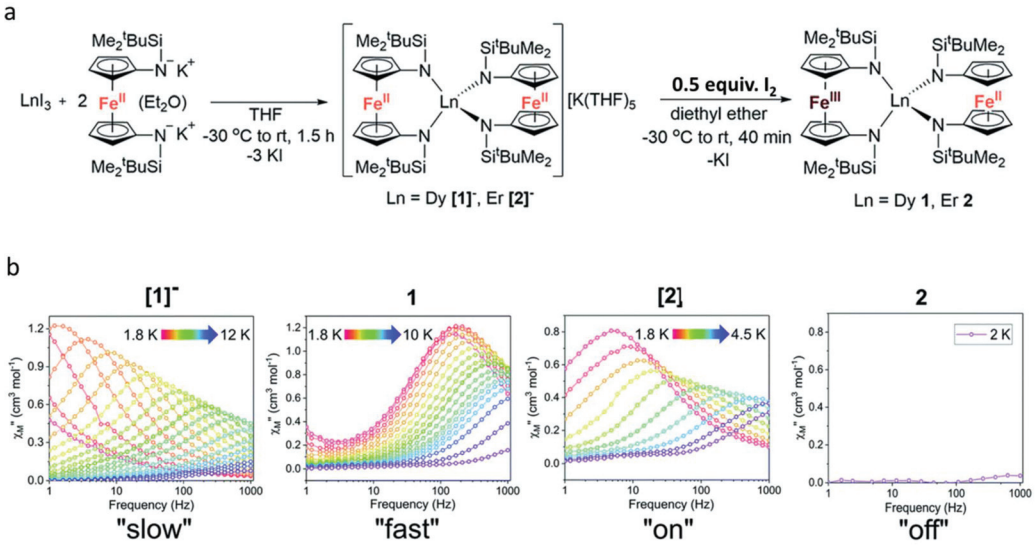

Fig. 8 (a) Synthesis of $[\mathbf{1}]^-$, **1**, $[\mathbf{2}]^-$, and **2**; (b) plots of $\chi''(\nu)$ for $[\mathbf{1}]^-$ and **1** under a 1000 Oe dc field and $[\mathbf{2}]^-$ and **2** under a 500 Oe dc field. Reproduced with permission from ref. 76. Copyright 2017 Royal Society of Chemistry.[76]

susceptibility measurements showed that an obvious increase of χT value was observed for 1Dy when the temperature was cooled to 50 K indicating a strong magnetic interactions between Dy(III) ions and π radicals, whereas 2Dy did not display similar behaviour due to the long Dy···Dy distance of 8.64(9) Å causing relatively weak magnetic interactions. Dynamic magnetic susceptibility measurements were also performed to study their magnetic relaxation and the results suggested that only 2Dy displayed SMM behaviour with clear χ'' peaks under a zero dc field.

Another method of constructing redox-switching of Ln-SMMs is to employ the redox activity of 3d or 4d transition metals.[73–75] Herein, a typical example reported by the Nippe group in 2017 was introduced.[76] As shown in Fig. 8a, the $FeCp_2$ ligand was used to associate with lanthanide ions aiming to modulate the dynamic magnetic properties by its redox properties. The trinuclear compounds $[\mathbf{1}]^-$ and $[\mathbf{2}]^-$ were synthesized by the reaction between two equivalents of $[K_2(OEt_2)]fc[NSi(t\text{-}Bu)Me_2]_2$ and one equivalent of LnI_3. Although the complete crystal structures of these two compounds were not obtained due to the presence of severe disorder, the important structural feature of equivalent distances between the lanthanide ion and the two Fe^{2+} ions can be determined. When half an equivalent of iodine was added, the oxidation reaction occurred to yield the corresponding neutral compounds, **1** and **2**, where the mixed valence of Fe^{3+}/Fe^{2+} was observed. Single crystal X-ray diffraction showed that these two neutral compounds are isostructural featuring an inequivalent Ln···Fe distance and the lanthanide ion was surrounded by a lower symmetry coordination environment. Ac magnetic susceptibility measurements confirmed the SMM behaviour of $[\mathbf{1}]^-$ under a zero dc field with a U_{eff} of 39.3 K; however, this behaviour was switched off in **1** owing to the appreciable loss of symmetry and the change in CF upon oxidation. When a 1000 Oe dc field was applied, both $[\mathbf{1}]^-$ and **1** exhibited a field-induced SMM behaviour but a faster magnetic relaxation was observed in **1**. For both Er(III) compounds, $[\mathbf{2}]^-$ and **2**, no reasonable χ'' signals were observed under a zero dc field at an ac frequency of 1000 Hz; however, a typical field-induced SMM behaviour was observed for $[\mathbf{2}]^-$ in the presence of a 500 Oe dc field while **2** exhibited no evidence of slow relaxation at any investigated temperature or field (Fig. 8b).

Conclusions and perspectives

In recent years, organometallic chemistry have helped researchers obtain many eye-catching results in the field of molecular magnetism, some of which represented the state-of-the-art achievements; however, it also hampered the addition of other physical properties to build multi-responsive molecular magnetic materials as most such Ln-SMMs were air-sensitive. In this review, we mainly discussed four external stimuli switching of SMM behaviour, *i.e.* light irradiation, thermal treatment, protonation/deprotonation and oxidation/reduction. No matter what kind of modulation, any slight change of the first coordination environment of lanthanide ions usually has remarkable impact on the magnetic relaxation making the manipulation of On/Off switching of Ln-SMM behaviour possible. On the other hand, for some structural changes which occurred far from the centre metal, such as the photoisomerization of DTE, they can also alter the magnetic properties through the change of intramolecular magnetic interactions, diplolar interactions, crystal packing, *etc.* as lanthanides are very sensitive to the subtle changes of structure. Most examples involved here have achieved reversible SC–SC transformation, which is very vital for determining the changes in the molecular structure and the further investi-

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gations of magneto-structural correlations by *ab initio* calculations. In fact, apart from these examples, other dynamic systems on the subject of this review have also been reported such as solvato-switching of SMM behaviour in low-dimensional lanthanide compounds,[77] the effects of guest-exchange on the magnetic relaxation,[78] multistimuli-responsive Ln-SMMs,[79] *etc*. Recently, the ultrafast transient absorption spectra were used to unveil the mechanism of the effects of ultrafast photoexcitation on the magnetic relaxation in a very simple Mn_3 SMM, providing new possibilities for optical control of the magnetization in SMMs on femtosecond timescales.[80]

From the perspective of the practical applications of Ln-SMMs, designing and building external stimuli modulated high-temperature Ln-SMMs with highly efficient reversible transformation and enough chemical stability in solid and solution will become the focus of future research.

Conflicts of interest

There are no conflicts to declare.

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