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Five new β -diketonate Ce^{3+} mononuclear complexes, $[\text{Ce}(\text{Btfa})_3(\text{H}_2\text{O})_2]$ (**1**), $[\text{Ce}(\text{Btfa})_3(\text{phen})]$ (**2**), $[\text{Ce}(\text{Btfa})_3(\text{bipy})]$ (**3**), $[\text{Ce}(\text{Btfa})_3(\text{terpy})]$ (**4**) and $[\text{Ce}(\text{Btfa})_3(\text{bathophen})(\text{DMF})]$ (**5**), where $\text{Btfa}^- = 4,4,4$ -trifluoro-1-phenyl-1,3-butanedionate, $\text{phen} = 1,10$ -phenanthroline, $\text{bipy} = 2,2'$ -bipyridyl, $\text{terpy} = 2,2':6',2''$ -terpyridine and $\text{bathophen} = 4,7$ -diphenyl-1,10-phenanthroline, have been synthesized and structurally characterized through X-ray diffraction of single crystals. The central Ce^{3+} atom displays a coordination number of 8 for **1**, **2** and **3** and of 9 for **4** and **5**. Under a 0 T external magnetic field, none of the given compounds exhibits single molecule magnet (SMM) behaviour. However, a small magnetic field, between 0.02 and 0.1 T, is enough for all the compounds to exhibit slow relaxation of the magnetization. A comprehensive magnetic analysis, with experimental magnetic data and *ab initio* calculations, was undertaken for all the complexes, and the study highlights the significance of the different spin relaxation mechanisms that must be considered for a Ce^{3+} lanthanide ion.

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Introduction

Magnetically bistable molecules that exhibit slow relaxation of the magnetization below a critical temperature, T_B , are named single molecule magnets (SMMs). The energy barrier separating the two bistable magnetic states is named the effective energy barrier (U_{eff}) and, for transition metal compounds, it is described by the magnetic anisotropy (D) and the square of the overall ground spin state (S): $U_{\text{eff}} = |D| \cdot S^2$ or $U_{\text{eff}} = |D| \cdot (S^2 - 1/4)$ for integer and non-integer S values respectively. The first molecular system for which single-molecular magnet type behaviour was observed was the mixed valence dodecanuclear Mn_{12} cluster with the formula $[\text{Mn}_{12}\text{O}_{12}(\text{CH}_3\text{COO})_{16}(\text{H}_2\text{O})_4]$ in 1993,¹ previously reported in 1980.² At the end of the twentieth century, efforts to obtain new SMMs with higher T_B and U_{eff} centred on transition metal clusters.³ SMM systems show very interesting properties and could be used in many potential

applications such as memory storage.^{3,4} After a plethora of 3d metal clusters were studied, the field of single molecule magnets moved onto lanthanide(III) ions, starting with the mononuclear $\text{Tb}(\text{III})$ phthalocyaninato-based double-decker complex $[\text{TbPc}_2]^-$ ($\text{Pc} = \text{phthalocyanine}$). It showed SMM behaviour and it was isolated in 2003 by N. Ishikawa *et al.*⁵ Due to the shielding of the $4f^n$ electrons by the filled $5s^2$ and $5p^6$ orbitals, the interaction between the lanthanide ion and the donor atoms can be considered as electrostatic in nature. Consequently, lanthanide ions have unquenched orbital angular momentum (L) that leads to strong spin-orbit coupling and therefore, intrinsic magnetic anisotropy; this makes them good candidates for SMMs. Mononuclear SMMs, also called single ion magnets (SIMs), are mononuclear systems that show slow relaxation of the magnetization.^{6–13} In lanthanides, the intrinsic magnetic anisotropy that the ions show is a key factor determining the magnetic properties of lanthanide(III) compounds, but the crystal field effect has relevance in the modulation of such anisotropy to tune the magnetic properties depending on the electron density of the m_j ground state. Rinehart and Long¹⁴ revealed that the electron density of the m_j ground state could be oblate (equatorially extended) or prolate (axially extended). For instance, in the $[\text{TbPc}_2]^-$ compound, the donor atoms of the ligand are located in the axial positions leading to an axial stressed coordination environment that enhances magnetic anisotropy on the easy axis leading to a good SMM performance of this Tb^{3+} compound with an oblate m_j ground state.⁵

In the design of SMMs, heavy metal lanthanides such as Tb^{3+} , Dy^{3+} and Er^{3+} ions have been extensively used because of

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their large magnetic and angular momentum J , where $J = L + S$ for half-filled Ln^{3+} ions, and consequently large magnetic anisotropy. Indeed, a rapid overview of the literature on lanthanide-based SMMs shows that most of the scientific production on this subject is related to Dy^{3+} , Tb^{3+} and Er^{3+} based species, with Dy^{3+} being the most popular.^{3,6–13} For instance, the mononuclear dysprosium complex $[(\text{Cp}^{\text{IPr}_5})\text{Dy}(\text{Cp}^*)][\text{B}(\text{C}_6\text{F}_5)_4]$, where Cp^{IPr_5} = pentaisopropylcyclopentadienyl and Cp^* = pentamethylcyclopentadienyl, was shown to exhibit the highest reported blocking temperature of 80 K.^{10,11,15} Still, the most relevant permanent magnets that are employed in industry are the ones based on light lanthanide ions, including the NdFeB and SmCo_5 ferromagnets.¹⁶ In this sense, the number of reported lanthanide-based SMMs of light lanthanides such as Ce^{3+} or Nd^{3+} ions is much smaller than those mentioned above.¹⁷ Focusing on cerium(III), its electronic configuration is $4f^1$ with a $^2F_{5/2}$ ground state and in some Ce^{3+} compounds magnetic relaxation can be observed because the spin-orbit effect of the $4f^1$ electron can create non-negligible magnetic anisotropy.

When the Ce^{3+} ion ($^2F_{5/2}$ ground state) is set in a coordination complex, the spin-orbit coupling component (J) splits into $(2J + 1) \pm m_J$ states: $+J$, $+J - 1, \dots, -J + 1$, $-J$, due to the crystal field: $\pm 5/2$, $\pm 3/2$ and $\pm 1/2$. For Ce^{3+} compounds the goal would be to choose the ideal coordination environment around the lanthanide ion to yield an axial geometry that would stabilize the $m_J = \pm 5/2$ Kramers doublet (KD) with an oblate $4f$ -shell electron density distribution as the ground state and destabilize the prolate electron distribution, $m_J = \pm 1/2$ and $\pm 3/2$ states. Such systems would lead to a reduced quantum tunneling of magnetization (QTM) effect with anisotropic barriers higher in energy (reduced m_J mixing) and, in principle, improve the SMM performance.^{14,18,19} Regarding β -diketonate systems, two series of mononuclear Ce^{3+} β -diketonate complexes with N-donor auxiliary ligands are found in the literature, Table S1.† First, 1,1,1-trifluoro-5,5-dimethyl-2,4-hexadione (Hfdh) led to the $[\text{Ce}(\text{fdh})_3(\text{bpy})]$ compound with D_{4d} symmetry.²⁰ Based on the 4,4,4-trifluoruro-1-(naphthalene-2-yl)butane-1,3-dionato (ntfa) β -diketonate ligand, we recently published a series of compounds: $[\text{Ce}(\text{ntfa})_3(\text{MeOH})_2]$, $[\text{Ce}(\text{ntfa})_3(5,5'\text{-Me}_2\text{bipy})]$ and $[\text{Ce}(\text{ntfa})_3(\text{bipy})_2]$, with field induced SIM behavior and D_{2d} , D_{2d} and C_{2v} point group symmetries, respectively.²¹ For these mononuclear Ce^{3+} β -diketonate complexes, *ab initio* calculations at zero field on the electron gyromagnetic ratio (g tensor) for each m_J state were described. Easy axial anisotropy was found (g_{xx} and $g_{yy} < g_{zz}$) through the calculations, which demonstrated that the effective g tensor of the ground doublet presented considerable transverse contributions (g_{xx} and g_{yy}) that led to the ground state wavefunction $|\pm 5/2\rangle$ mixed with $|\pm 1/2\rangle$ and $|\pm 3/2\rangle$ from the other excited m_J doublets.

Otherwise, the only Ce^{3+} compound showing SMM behavior to date is the trinuclear Zn-Ce-Zn compound prepared from an *o*-vanillin Schiff base based ligand.²² For this heterometallic compound, g_{zz} yielded 4.06 from *ab initio* calculations; this is close to the theoretical value of 4.28 for a pure $m_J = \pm 5/2$

2 Kramers doublet, $g_{zz} = 2g_J m_J$.²³ From the measured high-field electron paramagnetic resonance (HF-EPR) spectrum, g_{zz} is 3.27, which is assigned to the ground state of $m_J = 5/2$ KD with slight mixing of the $m_J = 1/2$ pair. Nevertheless, two mononuclear Ce^{3+} metallocene compounds, $[\text{Ce}(\text{Cp}^{\text{ttt}})_2\{\text{C}_6\text{F}_5\text{-k}^1\text{-F}\}\text{B}(\text{C}_6\text{F}_5)_3]$ and $[\text{Ce}(\text{Cp}^{\text{ttt}})_2\text{Cl}][\text{B}(\text{C}_6\text{F}_5)_4]$, are found in the literature where the g_{zz} axial component obtained by multiconfigurational analysis shows even higher values than the trinuclear Zn-Ce-Zn SMM compound, with values up to 4.16 and 4.22, respectively, and a ground state wavefunction of 100% $|\pm 5/2\rangle$ for the first compound.²⁴ The two Cp^{ttt} groups placed in the axial positions of the lanthanide cation and the low equatorial interactions of the $[\text{B}(\text{C}_6\text{F}_5)_4]^-$ and Cl^- anions, led to Ce^{3+} centers in an axial stressed symmetry. Furthermore, for these organometallic compounds, the EPR measurements were recorded and the experimental g^{eff} values were close to the computed ones. However, despite the calculated g_{zz} values being near 4.28, these compounds only showed slow relaxation of the magnetization when applying an external magnetic field of 1000 Oe. The other mononuclear cerium compounds found in the literature showing slow relaxation of the magnetization under an external H_{dc} , according to *ab initio* calculations, resulted from axial symmetry but with a considerable rhombic contribution to the ground state. Their magnetic properties are compiled in Tables S1 and S2.†^{25–29} A similar result was found for the dinuclear Ce^{3+} compound $[\text{Ce}_2(\text{RR-L})_2(\mu\text{-Cl})_6]$, where L is 1,4,7,10-tetrakis(2-pyridylmethyl)-1,4,7,10-tetraaza-cyclododecane, Table S3.†³⁰ In addition, the natural isotopes of Ce^{3+} do not have nuclear spin; this results in clearer and easier to interpret EPR spectra. Interestingly, in the studies published where EPR was also measured for the Ce^{3+} compounds, the g_{xx} , g_{yy} and g_{zz} experimental values were in good agreement with the computational experiments.^{22,24,25,30} At the moment, as far as we know, pure Ising type axial anisotropy ($g_z = 4.28$) has not been found in the computational calculations nor in EPR measurements for any of the Ce^{3+} compounds found in the literature. It is worth noting that slow relaxation of the magnetization under relatively small H_{dc} fields was measured for some Ce^{3+} coordination complexes where multiconfigurational CASSCF calculations indicated easy plane magnetic anisotropy that led to larger contributions of the $|\pm 1/2\rangle$ and $|\pm 3/2\rangle$ wavefunctions in the ground state $\pm m_J$ KD.^{25–29} Even though there was a transverse contribution to the ground state, the compounds showed similar magnetic behavior to those compounds with more axially strained coordination geometries. We attempted to compile the magnetic characteristics of the Ce^{3+} compounds found in the literature in Tables S1–S5.†

More examples of Ce^{3+} SMM compounds are needed to further understand their spin relaxation and so that we can control and manipulate it to be able to design compounds with better SMM performance. With the aim of increasing this limited family of compounds and providing new insights into the understanding of cerium(III) spin relaxation, five new β -diketonate Ce^{3+} coordination compounds are presented. The presented complexes are derived from the β -diketonate ligand



Btfa[−] = 4,4,4-trifluoro-1-phenyl-1,3-butanedionate with polypyridyl molecules as ancillary ligands and they have the formulae [Ce(Btfa)₃(H₂O)₂] (1), [Ce(Btfa)₃(phen)] (2), [Ce(Btfa)₃(bipy)] (3), [Ce(Btfa)₃(terpy)] (4) and [Ce(Btfa)₃(bathophen)(DMF)] (5), where phen = 1,10-phenanthroline, bipy = 2,2'-bipyridyl, terpy = 2,2':6',2''-terpyridine and bathophen = bathophenanthroline. For this purpose, the nitrogen donor N,N-L and N,N,N-L auxiliary ligands could be placed in the equatorial positions of the Ce³⁺ metallic center while the β-diketonates could be fixed in the axial positions, thus, in this way, stressing an axial geometry. Moreover, the five compounds were structurally and magnetically characterized and a thorough investigation of the experimental magnetic data and *ab initio* calculations was conducted to gain deeper insights into the spin dynamics involved in the magnetization relaxation of these compounds.

Experimental section

Starting materials

4,4,4-Trifluoro-1-phenyl-1,3-butanedione (HBtfa), 2,2'-bipyridyl (bipy), 1,10-phenanthroline (phen), 2,2':6',2''-terpyridine (terpy) and bathophenanthroline (bathophen) were purchased from Sigma-Aldrich. Cerium(III) chloride hexahydrate was obtained from Strem Chemicals and the other chemicals used were of analytical grade.

[Ce(Btfa)₃(H₂O)₂] (1). A methanol solution (5 mL) containing CeCl₃·6H₂O (2 mmol, 0.745 mg) was added to a previously mixed methanol solution (10 mL) composed of NaOH (6 mmol, 240 mg) and HBtfa (6 mmol, 1296.96 mg). The solution was stirred for 1 hour at room temperature. Then 80 mL of deionized water was added to the reaction mixture and it was allowed to stir overnight. The orange precipitate was filtered and dried under vacuum (yield: 75%). Anal. Calc. for C₃₀H₂₂CeF₉O₈ (821.59 g mol^{−1}): C, 43.9; H, 2.7%. Found: C, 43.2; H, 2.7%. Selected IR bands (ATR-IR, cm^{−1}): 3648(m), 3363(br), 1608(s), 1571(s), 1530(m), 1483(m), 1457(m), 1308(m), 1281(s), 1243(m), 1182(m), 1133(s), 1075(m), 943(m), 771(m), 700(m), 629(m) cm^{−1}.

[Ce(Btfa)₃(1,10-phen)] (2). An ethanol solution (15 mL) containing phen (1 mmol, 180.21 mg) was added to another ethanol solution (15 mL) containing [Ce(Btfa)₃(H₂O)₂] (1 mmol, 824.6 mg). The solution was stirred for 30 minutes and then left to stand undisturbed at room temperature. Orange single crystals suitable for X-ray diffraction were obtained within a week (yield: 56%). Anal. Calc. for C₄₂H₂₆CeF₉N₂O₆ (965.77 g mol^{−1}): C, 52.2; N, 2.9; H, 2.7%. Found: C, 52.0; N, 2.9; H, 2.6%. Selected IR bands (ATR-IR, cm^{−1}): 1609(s), 1576(s), 1521(s), 1486(m), 1470(m), 1320(m), 1287(s), 1241(m), 1180(s), 1133(s), 945(m), 844(m), 746(s), 699(s), 628(m), 578(m).

[Ce(Btfa)₃(bipy)(EtOH)]/[Ce(Btfa)₃(bipy)] (3a/3). An ethanol solution (15 mL) containing bipy (1 mmol, 156.2 mg) was added to another ethanol solution (15 mL) containing [Ce(Btfa)₃(H₂O)₂] (1 mmol, 824.6 mg). The solution was stirred for 30 minutes and then left to stand at room temperature.

Orange single crystals suitable for X-ray diffraction were obtained within a week (yield: 82%). X-ray diffraction analysis of several single crystals of the sample showed that the bulk compound was a mixture of two compounds: [Ce(Btfa)₃(bipy)(EtOH)] (3a) and [Ce(Btfa)₃(bipy)] (3). Compound 3 can be obtained in two different ways: after exposure of the single crystals from the bulk sample to the open air for three months or by heating the bulk sample in the oven for 1 h at 100 °C. Anal. Calc. for C₄₀H₂₆CeF₉N₂O₆ (941.7 g mol^{−1}) (3): C, 51.0; N, 3.0; H, 2.8%. Found: C, 51.2; N, 2.8; H, 2.8%. Selected IR bands (ATR-IR, cm^{−1}): for compound 3a 3054(m, broad), 1635(m), 1606(s, split), 1572(s), 1523(m), 1483(m), 1466(m), 1433(m), 1313(m, split), 1280(s), 1240(m), 1170(m), 1123(s, split), 1071(s), 1011(m), 941(m), 755(s, split), 699(s), 626(s), 573(s), for compound 3 1635(m), 1606(s, split), 1572(s), 1523(m), 1483(m), 1466(m), 1433(m), 1313(m, split), 1280(s), 1240(m), 1170(m), 1123(s, split), 1071(s), 1011(m), 941(m), 755(s, split), 699(s), 626(s), 573(s).

[Ce(Btfa)₃(terpy)] (4). An ethanol solution (15 mL) containing terpy (1 mmol, 233.3 mg) was added to another ethanol solution (15 mL) containing [Ce(Btfa)₃(H₂O)₂] (1 mmol, 824.6 mg). The solution was stirred for 30 minutes and then left to stand at room temperature. Orange single crystals suitable for X-ray diffraction were obtained within a week (yield: 21%). Anal. Calc. for C₄₅H₂₉CeF₉N₃O₆ (1018.8 g mol^{−1}): C, 53.0; N 4.1, H 2.9%. Found: C, 52.6; N, 3.4; H, 2.6%. Selected IR bands (ATR-IR, cm^{−1}): 1609(s, split), 1576(s), 1526(m), 1469(s), 1439(m), 1313(m), 1277(s), 1237(m), 1170(s), 1123(s), 1071(m), 1004(m), 938(m), 759(s), 716(s), 696(s), 652(m), 629(s), 573(m).

[Ce(Btfa)₃(bathophen)(DMF)] (5). An *N,N*-dimethylformamide (DMF) solution (10 mL) of bathophen (1 mmol, 332.40) was added to an ethanol solution (20 mL) containing [Ce(Btfa)₃(H₂O)₂] (1 mmol, 824.6 mg). The orange solution was stirred for 30 min at 80 °C, filtered and left to stand undisturbed at room temperature. Orange single crystals suitable for X-ray diffraction were obtained within a month (yield: 24%). Anal. Calc. for C₅₇H₄₁CeF₉N₃O₇ (1191.1 g mol^{−1}): C, 57.5; N, 3.5; H, 3.5%. Found: C, 57.1; N, 3.5; H, 3.4%. Selected IR bands (ATR-IR, cm^{−1}): 1662(w), 1612(s, split), 1572(m), 1526(m), 1466(m), 1433(w), 1310(m), 1277(s), 1234(m), 1180(s), 1123(s), 1067(m), 1021(m), 938(m), 762(s, split), 696(s), 669(m), 626(s), 573(s).

IR and elemental analyses

Infrared spectra of solid complexes were recorded on a Bruker Alpha P (platinum-ATR-cap) spectrometer. Elemental microanalyses were carried out with an Elementar Vario EN3 analyser.

Magnetic measurements

Magnetic measurements were performed on solid polycrystalline samples with a Quantum Design MPMS-XL SQUID magnetometer at the Magnetic Measurements Unit of the Scientific and Technological Center (CCiTUB) of the University of Barcelona. Data were collected for powdered microcrystal-



line samples or for crushed polycrystalline samples of compounds **1** to **5**, the purity and structural integrity of which were analyzed by powder X-ray diffraction (see Molecular structure section of the Results and discussion). The samples were mounted in a gelatin capsule. Direct current (dc) susceptibility measurements were made over the 2–300 K temperature range and under an external magnetic field of 0.3 T. Blank measurements were also recorded for the capsule and holder and their diamagnetic contributions were corrected. Pascal's constants were used to estimate the diamagnetic corrections, which were subtracted from the experimental susceptibilities to give the corrected molar magnetic susceptibilities. Alternating current (ac) susceptibility measurements were carried out by applying an oscillating ac field of 4 Oe with ac frequencies between 1 and 1500 Hz for **1**, **3** and **5** and between 10 and 1500 Hz for compounds **2** and **4** at different temperatures and different dc applied fields indicated in the text.

X-ray crystallography

Crystals of **1–5** were mounted in air on a D8VENTURE (Bruker) diffractometer with a CMOS detector. Crystallographic data, conditions required for intensity data collection and some features of the structure refinements are listed in Tables S6 and S7.† All the structures were refined by the least-squares method. Intensities were collected with multilayer monochromated Mo-K α radiation. Lorentz polarization and absorption corrections were made for all samples. The structures were solved by direct methods, using the SHELXS-97 computer program³¹ and refined by the full-matrix least-squares method, using the SHELXL-2018 computer program.³² The non-hydrogen atoms were located in successive difference Fourier transformations and refined with anisotropic thermal parameters on F^2 . For hydrogen atoms, isotropic temperature factors were assigned to be 1.2 or 1.5 times the respective parent.

Powder X-ray diffraction (PXRD) measurements (see Fig. S1–S5†) were used to check the bulk phase purity. The powder patterns were recorded on a Bruker D8 Advance diffractometer (Bruker AXS, Madison, WI, USA) with a LynxEye detector in Bragg–Brentano θ/θ geometry, with the sample dispersed thinly on a zero-background Si sample holder, $\lambda(\text{CuK}\alpha) = 1.54060 \text{ \AA}$, scanning from $2\theta = 5^\circ$ to 50° , with a step size of 0.02° .

Computational details

Ab initio calculations were performed using crystallographic structures and Orca software, version 5.0.3. The def2-TZVP basis set was used.³³ NEVPT2 calculations did not include dynamic correlations as it was previously demonstrated for Ce complexes that this was not essential due to the large ionic character of the Ln–O and Ln–N bonds.³⁴ The selected active space was (1,7), considering the seven doublets. The SINGLE_ANISO module, as implemented in Orca, was employed.³⁵

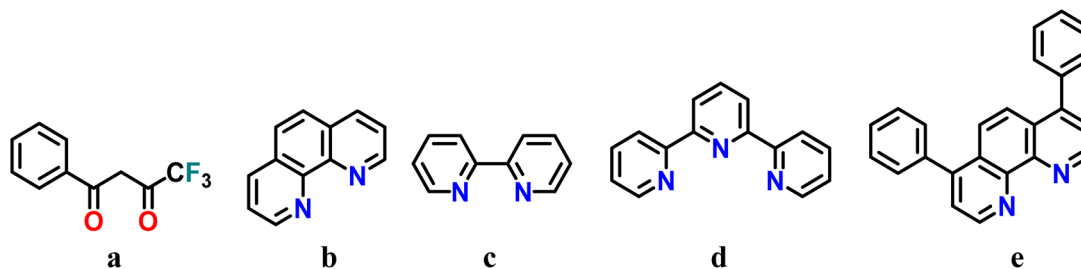
To evaluate the vibrational frequencies, the experimental geometries were optimized with Gaussian09,³⁶ the B3LYP functional³⁷ and using the Stuttgart pseudo/basis set³⁸ for Ce and a TZV basis set³⁹ for lighter atoms.

Results and discussion

General syntheses

Deprotonation of the β -diketone 4,4,4-trifluoro-1-phenyl-1,3-butanedione (HBtfa; Scheme 1a) in a basic medium and further addition of cerium(III) chloride led to the precipitation of the neutral $[\text{Ce}(\text{Btfa})_3(\text{H}_2\text{O})_2]$ (**1**) aqua complex. Compound **1** was used as the precursor in subsequent reactions. Compounds **2–5** were obtained through the displacement of the H_2O molecules of **1** by different polypyridyl ligands. The nitrogen containing molecules used for this family of cerium(III) compounds were: 1,10-phenanthroline (phen; Scheme 1b), 2,2'-bipyridyl (bipy; Scheme 1c), 2,2';6',2''-terpyridine (terpy; Scheme 1d) and bathophenanthroline (bathophen; Scheme 1e). The reaction of precursor compound **1** with chelating N,N-L and N,N,N-L ligands in a 1 : 1 ratio and in an ethanolic solution (DMF/ethanol in the case of complex **5**) led to the mononuclear Ce^{3+} coordination complexes $[\text{Ce}(\text{Btfa})_3(\text{phen})]$ (**2**), $[\text{Ce}(\text{Btfa})_3(\text{bipy})]$ (**3**), $[\text{Ce}(\text{Btfa})_3(\text{terpy})]$ (**4**) and $[\text{Ce}(\text{Btfa})_3(\text{bathophen})(\text{DMF})]$ (**5**).

In the case of compounds **3a** and **3**, after the reaction of precursor **1** with the 2,2'-bipyridyl (bipy) ligand in ethanol as a solvent, structure determination by single crystal X-ray diffraction by using one chosen crystal from the bulk product indicated that the compound was $[\text{Ce}(\text{Btfa})_3(\text{bipy})(\text{EtOH})]$, **3a**. Powder X-ray diffraction (PXRD) of the whole powder sample, Fig. S3,† indicates the presence of two phases. After various recrystallization processes of the product, the PXRD spectra



Scheme 1 Molecular representations of (a) 4,4,4-trifluoro-1-phenyl-1,3-butanedione (HBtfa), (b) 1,10-phenanthroline (phen), (c) 2,2'-bipyridyl (bipy), (d) 2,2';6',2''-terpyridine (terpy) and (e) bathophenanthroline (bathophen).



were still the same. At that point, by choosing another single crystal of the bulk compound, SCXRD afforded a different structure with the formula $[\text{Ce}(\text{Btfa})_3(\text{bipy})]$ (**3**).

Complex **3** can be isolated by removing all the coordinated EtOH molecules present in the bulk product from $[\text{Ce}(\text{Btfa})_3(\text{bipy})(\text{EtOH})]$ (**3a**). We found two different ways to do so. First, by leaving the obtained product in the open air for 3 months. The second method was to keep sample **3a** in the oven for 1 h at 100 °C, Fig. S3e.† Moreover, TGA measurements were performed on a freshly prepared sample, which probably predominantly contained compound **3a**. As can be observed in Fig. S6,† at around 100 °C there is a weight decrease corresponding to the loss of 4.9% of the sample content. This percentage corresponds to one EtOH molecule; the decrease of the curve starts at 29.56 °C and finishes at 175.17 °C. Then complex **3** was isolated as can be seen by PXRD, Fig. S1.†

Molecular structures of the compounds

The structures of compounds **1–5** and **3a** are presented here. A partially labelled plot of the Ce^{3+} mononuclear complexes, coordination polyhedra and intermolecular interaction arrangements are presented in Fig. 1–8 and Fig. S7–S10.†

Crystallographic data are collected in Tables S6 and S7.† Selected bond lengths of compounds **1–5** are listed in Table 1.

$[\text{Ce}(\text{Btfa})_3(\text{H}_2\text{O})_2]$ (**1**). Complex **1**, with the formula $[\text{Ce}(\text{Btfa})_3(\text{H}_2\text{O})_2]$, crystallizes in an orthorhombic crystal system, $P2_12_12_1$ space group. The structure of **1** consists of mononuclear molecules in which each Ce^{3+} ion is octa-coordinated in a CeO_8 coordination environment (Fig. 1a). The central ion is coordinated to six oxygen atoms from three different Btfa[−] ligands, with Ce–O distances in the 2.390(3)–2.505(3) Å range. In addition, the other two coordination sites around the central atom are occupied by the O7 and O8 atoms of two water molecules, with Ce–O(water) distances of 2.535(4) and 2.520(3) Å, respectively. The estimation of the distortion coefficients of the CeO_8 coordination polyhedron for **1** with reference to the ideal eight vertex polyhedron was performed by employing the continuous shape measure theory and SHAPE software,⁴⁰ as for all the other compounds presented herein, showing that the O_8 disposition for **1** was intermediate between different coordination polyhedra. The best SHAPE estimation led to the triangular dodecahedron (TDD-8, D_{2d}) geometry with a continuous shape measurement (CSHM) value of 0.416, Fig. 1b.

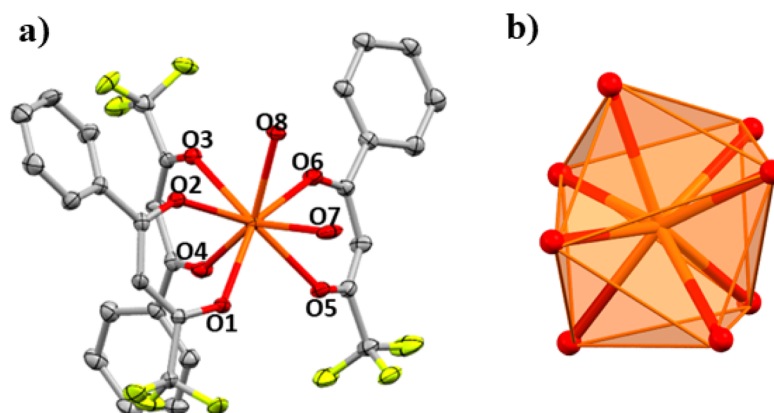


Fig. 1 (a) Partially labelled structure of **1** where hydrogen atoms have been omitted for clarity. (b) Idealized coordination polyhedron (triangular dodecahedron; TDD-8, D_{2d}) compared with the real positions of the coordinating atoms of compound **1**.

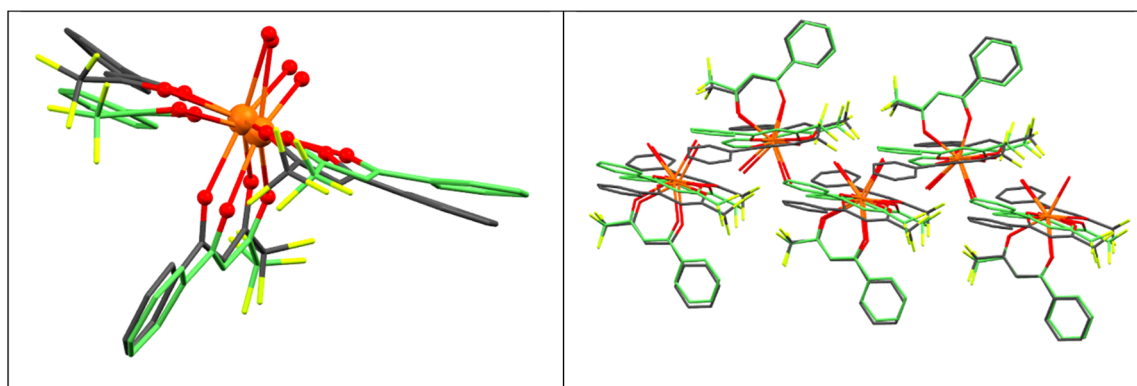


Fig. 2 Superimposed crystal structures of compound **1** obtained by single crystal XRD. The structure in default colours is the structure measured at 100 K and in green is the structure measured at 294 K.



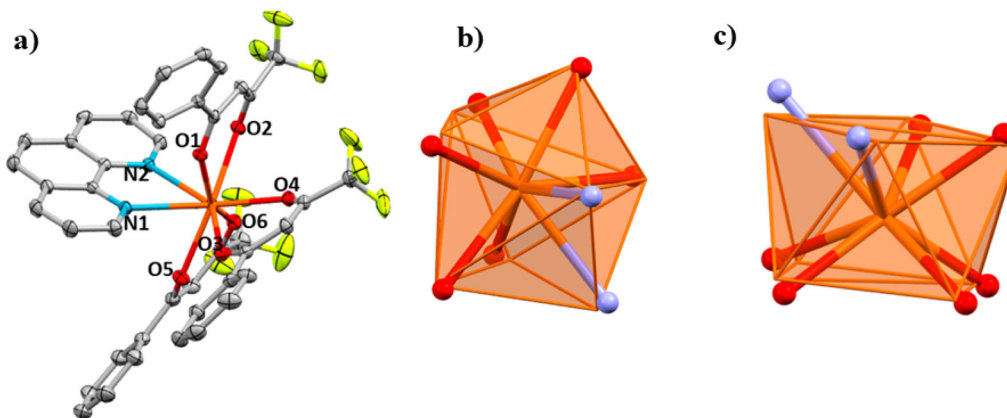


Fig. 3 (a) Partially labelled structure of **2**. The hydrogen atoms have been omitted for clarity. (b) Idealized coordination polyhedron (triangular dodecahedron; TDD-8, D_{2d}) compared with the real positions of the coordinating atoms of compound **2**. (c) Idealized coordination polyhedron (square antiprism; SAPR-8, D_{4d}) compared with the real positions of the coordinating atoms of compound **2**.

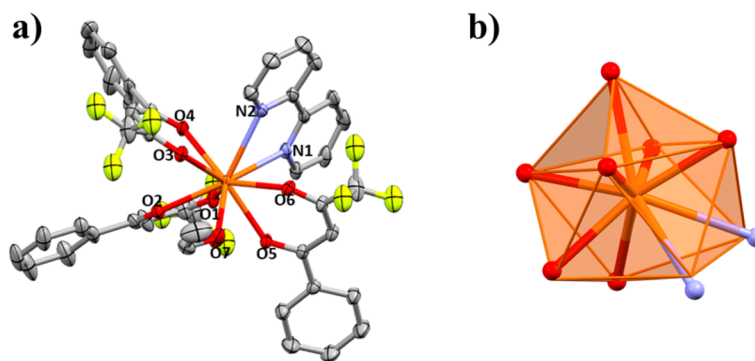


Fig. 4 (a) Partially labelled structure of **3a** where hydrogen atoms have been omitted for clarity. (b) Idealized coordination polyhedron (tricapped trigonal prism; TCTPR-9, D_{3h}) compared with the real positions of the coordinating atoms of compound **3a**.

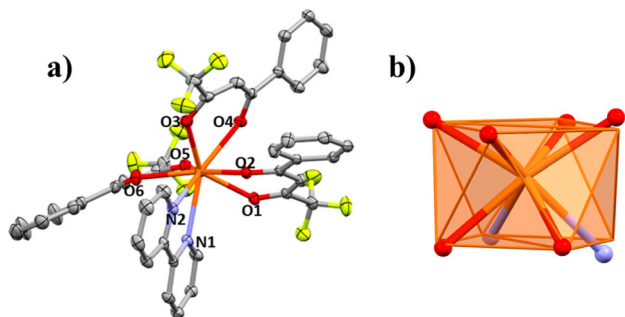


Fig. 5 (a) Partially labelled structure of **3** where hydrogen atoms have been omitted for clarity. (b) Idealized coordination polyhedron (square antiprism; SAPR-8, D_{4d}) compared with the real positions of the coordinating atoms of compound **3**.

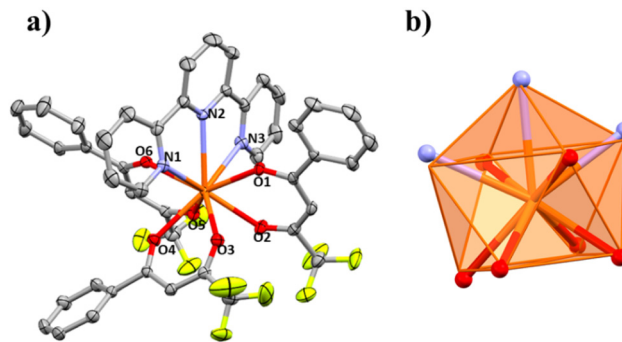


Fig. 6 (a) Partially labelled structure of **4** where hydrogen atoms have been omitted for clarity. (b) Idealized coordination polyhedron (capped square antiprism; CSAPR-9, C_{4v}) compared with the real positions of the coordinating atoms of compound **4**.

Molecules of **1** are arranged in space through hydrogen bonds forming an infinite 1D chain in the $[1\ 0\ 0]$ direction, Fig. S7.† The atoms contributing to these intermolecular interactions are the hydrogens from the two water molecules coordinated to the cerium(III) atoms. On one hand, the H atoms

from O8 interact with O1 and O5 from two different β -diketonates of the neighbouring coordination complex with donor (D)⋯acceptor (A) distances of 2.777 and 2.761 Å. On the other hand, the H atoms from the other O7 water molecule are connected through H-bonding with the O3 and F34 atoms



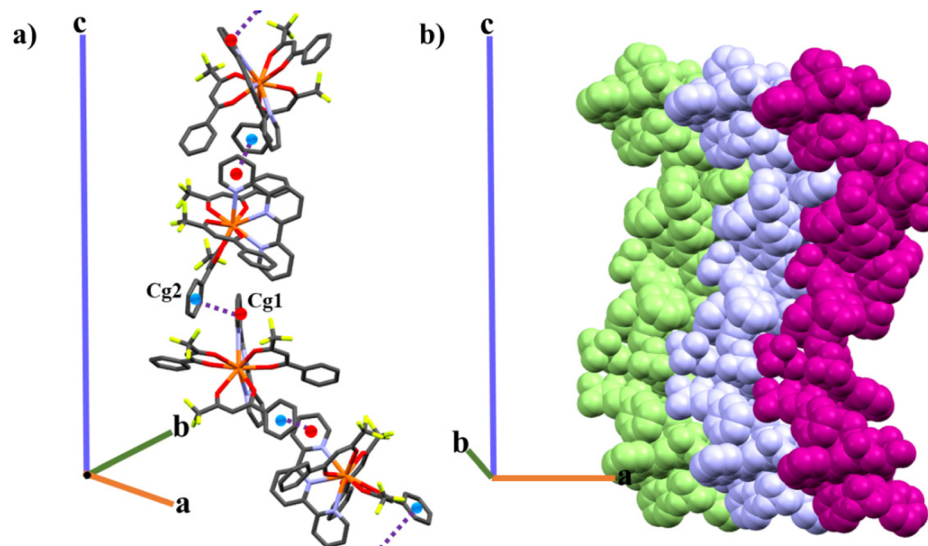


Fig. 7 (a) Helical chain formed by the intermolecular interactions of compound 4. The π - π stacking interactions are represented by the purple dotted lines and (b) helical chains along the 4_3 screw axis.

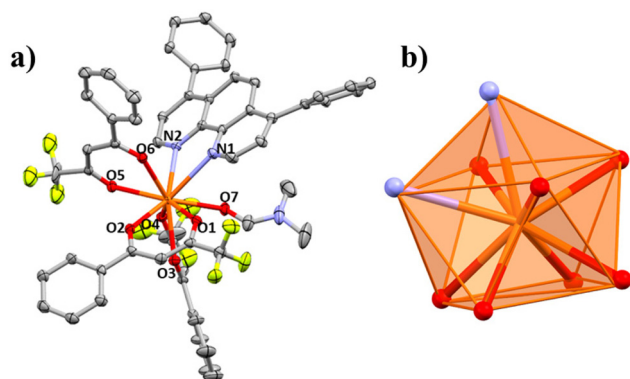


Fig. 8 (a) Partially labelled structure of 5 where hydrogen atoms have been omitted for clarity. (b) Idealized coordination polyhedron (capped square antiprism; CSAPR-9, C_{4v}) compared with the real positions of the coordinating atoms of compound 5.

from the nearby complex with D-A distances of 2.868 and 3.026 Å respectively. The shortest Ce-Ce intermolecular distance is 6.105 Å.

PXRD for **1** was measured at 294 K, and the resulting spectrum was compared with the calculated one for the single crystal measured at 100 K to verify that the entire polycrystalline product was the same as the structure obtained by single crystal X-ray diffraction, Fig. S1.† However, the spectra do not match very well. Not even after several recrystallizations of the product. Then, one single crystal was measured by XRD at room temperature (300 K). The calculated spectrum for this new structure determination matches the spectrum of the powder one measured at 294 K, Fig. S1.† Comparing the crystallographic data for compound **1** at 100 K and at room temperature (Table S6†), we can see that, on decreasing the temperature, there is a slight structural change. The crystal system and space group are still the same (orthorhombic, $\alpha = \beta = \gamma = 90^\circ$ and $P2_12_12_1$), but the cell parameters change from $a = 10.854$, $b = 13.219$ and $c = 23.317$ Å at room temperature to $a = 10.809$, $b = 14.77$ and $c = 21.079$ Å at 100 K. The structure at room temperature (294 K) shows Ce-O distances and angles that are slightly different from those measured at 100 K, Table 1. At room temperature, the coordination polyhedral geometry is still close to an ideal triangular dodecahedron

Table 1 Selected bond lengths (Å) of compounds 1–5

	1	1 r.t.	2	3a	3b	5	6
Ce–O1	2.455(3)	2.438(7)	2.453(2)	2.443(3)	2.421(2)	2.450(2)	2.448(3)
Ce–O2	2.427(4)	2.474(6)	2.467(2)	2.481(2)	2.467(1)	2.454(2)	2.461(3)
Ce–O3	2.505(3)	2.522(6)	2.425(2)	2.460(3)	2.405(2)	2.531(2)	2.511(4)
Ce–O4	2.390(3)	2.379(7)	2.419(1)	2.496(3)	2.459(2)	2.490(2)	2.470(3)
Ce–O5	2.471(3)	2.422(7)	2.434(2)	2.464(3)	2.417(2)	2.428(2)	2.476(3)
Ce–O6	2.402(4)	2.473(7)	2.416(2)	2.490(2)	2.446(2)	2.461(2)	2.479(3)
Ce–O7	2.535(4)	2.509(7)	—	2.570(3)	—	—	2.541(3)
Ce–O8	2.520(3)	2.553(7)	—	—	—	—	—
Ce–N1	—	—	2.659(2)	2.713(3)	2.660(2)	2.688(3)	2.701(4)
Ce–N2	—	—	2.697(2)	2.730(3)	2.668(2)	2.717(2)	2.701(4)
Ce–N3	—	—	—	—	—	2.680(3)	—



(TDD-8, D_{2d}) with a CShM value of 0.327, whereas the CShM value for **1** at 100 K is 0.416.

In Fig. 2 the structures obtained at 300 K and 100 K are superimposed. The slight difference between the distances and angles is enough to change the cell parameters when cooling the single crystal resulting in different powder XRD patterns.

[Ce(Btfa)₃(phen)] (2). Complex **2** with the formula [Ce(Btfa)₃(phen)] crystallizes in the monoclinic crystal system, $P2_1/c$ space group. In **2** each Ce^{3+} ion is octacoordinated with a CeN_2O_6 coordination environment formed by one phen and three Btfa[−] ligands, Fig. 3a. Two of the coordination sites around the central Ce^{3+} ion are occupied by the N1 and N2 atoms of the phen ligand, with Ce–N distances of 2.659(2) and 2.697(2) Å, respectively. In addition, the Ce(III) ion is also coordinated to six oxygen atoms from three different Btfa[−] chelating ligands, with Ce–O distances in the 2.416(2)–2.467(2) Å range.

The CeN_2O_6 disposition for **2** is intermediate between different coordination polyhedra. The lowest CShM values for **2** correspond to the triangular dodecahedron (TDD-8, D_{2d}) and square antiprism (SAPR-8, D_{4d}) geometries with CShM values of 1.700 and 1.835, respectively, Fig. 3b and c. The PXRD spectrum of the powder sample of **2** compared to the one calculated for the single crystal structure matches perfectly indicating the purity of the bulk product, Fig. S2.†

The [Ce(Btfa)₃(phen)] molecules are arranged in the crystal lattice through π – π stacking interactions between the ring formed by C5–C10 from one Btfa[−] ligand (Cg3) and the three rings with delocalized electron density from the phen ligand (Cg9). The intermolecular distance between the two centroids (Cg3...Cg9) is 3.921 Å and the interaction grows along the [0 0 1] base vector as an infinite 1D chain Fig. S8.† The shortest Ce–Ce intermolecular distance is 8.806 Å.

[Ce(Btfa)₃(bpy)(EtOH)] (3a). Complex **3a**, [Ce(Btfa)₃(bpy)(EtOH)], crystallizes in the triclinic crystal system, with the $P\bar{1}$ space group, Fig. 4a. The structure of **3a** consists of mononuclear molecules in which each Ce^{3+} ion is nonacoordinated with a CeN_2O_7 coordination environment formed by six O atoms from the three deprotonated β -diketonate ligands with Ce–O distances in the 2.496–2.443 Å range, two N atoms from the bpy ancillary ligand with Ce–N distances of 2.713 and 2.730 Å and the O atom from the ethanol molecule with a Ce–O distance of 2.570 Å. The fluorine atoms from the –CF₃ group of the substituted β -diketone molecule show disorder. The CeN_2O_7 nine coordination polyhedron has a coordination geometry close to that of a tricapped trigonal prism (TCTPR-9, D_{3h}), Fig. 4b, with a CShM value of 0.562.

Powder X-ray diffraction was performed and the result was compared with the calculated PXRD spectrum of the single crystal structure of **3a**, but the spectra did not properly match, Fig. S3b.† As mentioned before, this is because, after the reaction of the [Ce(Btfa)₃(H₂O)₂] precursor with bpy, a mixture of structures [Ce(Btfa)₃(bpy)(EtOH)] (**3a**) and [Ce(Btfa)₃(bpy)] (**3**) crystallizes. Unlike **3**, compound **3a** cannot be isolated.

[Ce(Btfa)₃(bpy)] (3). Complex [Ce(Btfa)₃(bpy)] (**3**) crystallizes in the monoclinic crystal system, with the $P2_1/n$ space group.

The structure of **3** consists of molecules in which each Ce^{3+} ion is octacoordinated with a CeO_6N_2 coordination environment formed by six O atoms from the three Btfa[−] molecules with Ce–O distances in the range of 2.467–2.405 Å and by two N atoms from the bpy ligand with Ce–N distances of 2.660 and 2.668 Å, Fig. 5a, Table 1. In this mononuclear complex the Ce–O and Ce–N distances are slightly shorter compared to the ones found in [Ce(Btfa)₃(bpy)(EtOH)] (**3a**). The CeO_6N_2 octacoordinated polyhedron has a coordination geometry that is close to an ideal square antiprism (SAPR-8, D_{4d}) with a CShM value of 1.058, Fig. 5b. As explained in the General syntheses section, compound **3** can be isolated from the mixture of **3a** and **3** that is obtained after the reaction by heating the sample in an oven for 1 h at 100 °C.

The intermolecular interactions to consider for both structures, **3a** and **3**, are the π – π stacking interactions between the rings from the bpy ligands with Cg1–Cg2 distances of 3.687 Å for **3a** and of 3.999 Å for **3**. Moreover, other weak intermolecular interactions such as π –H build up a three-dimensional supramolecular structure, see Fig. S9a and S9b.† The shortest Ce–Ce intermolecular distance is 8.647 Å.

[Ce(Btfa)₃(terpy)] (4). Complex **4** crystallizes in a tetragonal crystal system in the Sohncke $P4_3$ space group. The Sohncke space groups are formed of 65 space groups, which contain only rotation and translation symmetry operations (operations of the first kind) and where the reflection, glide reflection, rotoinversion and inversion mirror operations are restricted. Some of the molecular structures containing these non-enantiogenic space groups, 22 out of 65, result in chiral crystal structures (which are different from chiral molecules). For example, proteins generally crystallize in Sohncke space groups.⁴¹ The structure of **4** consists of molecules in which each Ce^{3+} ion is nonacoordinated with a CeO_6N_3 coordination environment, Fig. 6a. The CeO_6N_3 coordination sphere is generated by six O atoms from the three Btfa[−] ligands with Ce–O distances ranging from 2.428 to 2.531 Å and the three N atoms from the terpy ligand with Ce–N distances ranging from 2.680 to 2.717 Å. The CeN_3O_6 nine coordination polyhedron has a coordination geometry close to an ideal capped square antiprism (CSAPR-9, C_{4v}) with a CShM value of 0.751, Fig. 6b.

Each asymmetric unit is formed by one mononuclear entity. For **4**, the asymmetric unit grows in space forming a laevorotatory 4_3 -helical chain through the c axis (left-handed chain) and, therefore, homo-chiral packing of helices in the crystal is induced. The helix is formed around the crystallographic 4_3 screw axis *via* the π – π stacking interaction between two different rings with delocalized electron density, one from the terpy ligand (Cg1) and the other from one Btfa[−] β -diketone molecule (Cg2) with an intermolecular distance of 3.767 Å, Fig. 7a. The helices align themselves along the a and b axis through weak Van der Waals interactions, resulting in the formation of a chiral crystal structure, Fig. 7b. The shortest Ce...Ce intermolecular distance is 10.563 Å.

The enantiomeric pair for this Sohncke space group is $P4_1$ in which the helix would be dextrorotatory. To find a crystal with the enantiomeric $P4_1$ Sohncke space group, a second



single crystal was measured. For this second measurement, the same $P4_3$ space group was found. Though both crystals presented the same chirality, we cannot assert that the final product is purely chiral and neglect the idea that it could be a racemic mixture. The powder XRD results presented in Fig. S4† support one single phase for the whole powder product of 4.

[Ce(Btfa)₃(bathophen)(DMF)] (5). Complex [Ce(Btfa)₃(bathophen)(DMF)] (5) crystallizes in the monoclinic crystal system, $P2_1/c$ space group. The structure of 5 consists of mononuclear molecules in which each Ce³⁺ ion is nonacoordinated with a CeN₂O₇ coordination environment, Fig. 8a. The coordination sphere encompasses six O atoms from three Btfa[−] ligands with Ce–O distances in the 2.448 to 2.511 Å range, two N atoms from the bathophen ligand with Ce–N distances of 2.724 and 2.701 Å and a DMF molecule through the O atom with a Ce–O distance of 2.541 Å. The N₂O₇ vertexes are established around the Ce³⁺ metal in a polyhedral geometry that is close to an ideal capped square antiprism (CSAPR-9, C_{4v}) with a CShM value of 0.316, Fig. 8b. The mononuclear complexes are arranged in space through π – π stacking interactions between one of the aromatic rings from the bathophen ligand (Cg1) and one phenyl ring from a Btfa[−] ligand. The Cg(1)–Cg(2) intermolecular distance is 3.818 Å and it grows along the c axis as shown in Fig. S4†. The shortest Ce–Ce intermolecular distance is 10.770 Å. Moreover, PXRD measurements were performed on the powder product. The calculated PXRD spectrum from the single crystal structure was compared with the experimental one, which affirmed a single phase throughout the sample, Fig. S5†.

Magnetic properties

Static magnetic properties

Direct current (dc) magnetic susceptibility and magnetization measurements were performed on polycrystalline samples of compounds 1–5. Since compound 3a was not isolated as a pure compound, the magnetic study was not performed for this sample. The $\chi_M T$ vs. T plots in the 2–300 K temperature

range under a dc magnetic field of 0.3 T are shown in Fig. 9 (left). At room temperature (300 K) the $\chi_M T$ values for compounds 1–5 are 0.67, 0.73, 0.68, 0.73 and 0.76 cm³ mol^{−1} K respectively. These $\chi_M T$ values are close to the calculated value for an isolated Ce³⁺ cation in the ground state, ²F_{5/2} ($S = 1/2$, $L = 3$, $J = 5/2$, $g_J = 6/7$), which is 0.8 cm³ mol^{−1} K.⁸ On cooling the samples, the $\chi_M T$ values decrease gradually due to the progressive thermal depopulation of the m_J states to finite values at 2 K of 0.23, 0.40, 0.44, 0.32 and 0.38 cm³ mol^{−1} K for compounds 1–5 respectively. Other mononuclear Ce³⁺ coordination compounds show similar $\chi_M T$ vs. T curves.^{21,42,43} The field dependence of the magnetization (M) recorded at 2 K is depicted in Fig. 9 (right). On increasing the external magnetic field up to 5 T, the magnetization values increase to 0.62, 0.84, 0.89, 0.83 and 0.89 N_Bμ_B for 1–5 respectively without stabilization of the M values, showing no evidence of saturation of the magnetization ($g_J J$). Similar behavior is found for other Ce³⁺ compounds showing significant magnetic anisotropy.^{17,21,30}

Dynamic magnetic properties

No frequency dependence of the out-of-phase magnetic susceptibility component (χ_M'') was found under a 0 T external magnetic dc field for complexes 1–5 (Fig. S11†), suggesting the lack of a pure axial Ising type geometry, which sets $m_J = \pm 5/2$ as the ground doublet, and makes the reorientation of the magnetization occur through the fast quantum tunneling of magnetization (QTM) mechanism preventing the SMM behavior.²³ Nevertheless, by applying a small direct current (dc) magnetic field (H_{dc}), the QTM was suppressed enough so that all the above compounds showed slow relaxation of the magnetization.

To decide which external dc magnetic field was optimal for each compound, ac measurements were performed at a constant temperature of 2.5 K for 1, 3 and 5, of 3.5 K for 2, and of 2 K for 4 at oscillating frequencies between 1 and 1488 Hz and under different H_{dc} fields. The measurements of χ_M' and χ_M'' with frequency (ν) plots, at a constant temperature, are depicted in Fig. S12† for samples 1–5. When applying low H_{dc} (0.002–0.01 T), χ_M'' shows maximum values at higher frequencies. On increasing H_{dc} , the peaks increase in intensity, reach-

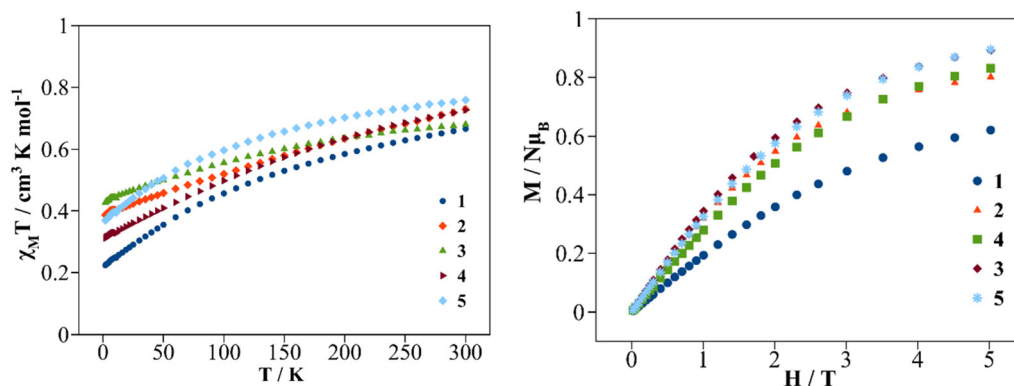


Fig. 9 $\chi_M T$ vs. T plot (left) and M vs. magnetic field plot (right) for compounds 1–5.



ing their maximum values and shifting to the 96–177 Hz frequency zone, then staying constant until higher H_{dc} is applied. Then χ_M'' shifts again to higher frequencies, diminishing its intensity. The optimal H_{dc} was 0.1 T for **1** and **3**, 0.05 T for **2** and 0.02 T for **4** and **5**, Fig. S13.† Given the shorter Ce...Ce intermolecular distance of 6.105 Å, compound **1** might require a greater external H_{dc} . The shortest Ce...Ce intermolecular distances found for structures **2–5** are 8.806, 8.647, 10.563 and 10.770 Å respectively.

This behavior is represented as τ^{-1} based on the H_{dc} plots, Fig. 10. At low magnetic fields, τ^{-1} values decrease, corresponding to suppression of the QTM mechanism by the external magnetic field. After 0.3 T for **1–3** and **5**, the τ^{-1} values remain almost constant, accounting for the predominance of Raman and other thermally dependent relaxation processes. For compound **4**, the constant trend at intermediate H_{dc} is rather poor and it is not as well differentiated as for the other compounds. When surpassing the specific H_{dc} , this constant tendency is broken, and τ^{-1} values increase exponentially following the H^4 power law, showing that the direct process is taking place at higher magnetic fields. ($\tau^{-1} = AH^4T$ for relaxation of the magnetization through a direct process.)

Under the chosen dc external magnetic field, dynamic measurements were performed under an oscillating ac field of 4×10^{-4} T in the 1–1488 Hz frequency range for **1**, **3** and **5**, and of 10–1488 Hz for **2** and **4**. Maximum values of the out of phase magnetic susceptibility component can be seen below 5 K for **1–5**, Fig. S14.† All the samples show slow relaxation of the magnetization over a similar temperature range; compound **2** is the one for which the χ_M'' dependence extends to higher temperature (5.7 K).

The χ_M'' vs. oscillating frequency plots at different temperatures are depicted in Fig. 11 and χ_M' vs. ν plots at different temperatures are found in Fig. S15.† For complexes **1–5**, the χ_M'' peaks shift progressively to higher frequencies as the

temperature increases, demonstrating the thermal dependence of magnetization relaxation. Therefore, we could assume that the QTM relaxation process would be considerably reduced after the application of an external dc field.³ When comparing compounds **1–5**, **2** shows χ_M'' dependence on T at lower frequencies, whereas **3** shows a dependence at higher ones.

The Cole–Cole plots for **1–5** (Fig. 12) are not perfectly symmetrical semicircles. The curves could be fitted using the generalized Debye model described with the Casimir–Du Pré function (eqn (S1)†)⁴⁴ from which the relaxation times for magnetization can be extracted. The resulting parameters obtained from the fitting are presented in Tables S8–S12.† The distribution of relaxation times (α) is narrow for the presented compounds mostly for **1**, **4** and **5** in which α values are 0.077–0.0041, 0.0529–0.0031 and 0.05–0.0118 respectively while for **2** and **3** α values are over a slightly wider range of 0.1804–0.0018 and 0.1335–0.0146 respectively.

For a spin system, magnetization relaxation can occur through different relaxation mechanisms. The dependence of τ on temperature (T) and the applied external magnetic field (H) is expressed in eqn (1).

$$\tau^{-1} = aH^4T + d\left(\frac{1 + eH^2}{1 + fH^2}\right)T^n + \frac{B_1}{1 + B_2H^2} + \tau_0^{-1} \exp\left(\frac{-\Delta E}{k_B T}\right) \quad (1)$$

The first term is the field dependent direct process, which involves one phonon, the second term refers to the Raman process that occurs through a virtual state involving two phonons. Here it is represented as a field dependent term using the Brons–Van Vleck equation, which has a coefficient field dependence and exponential dependence on temperature. The d parameter represents zero-field relaxation, e is related to the paramagnetic center concentration and the f parameter reports the effect of the external field on suppres-

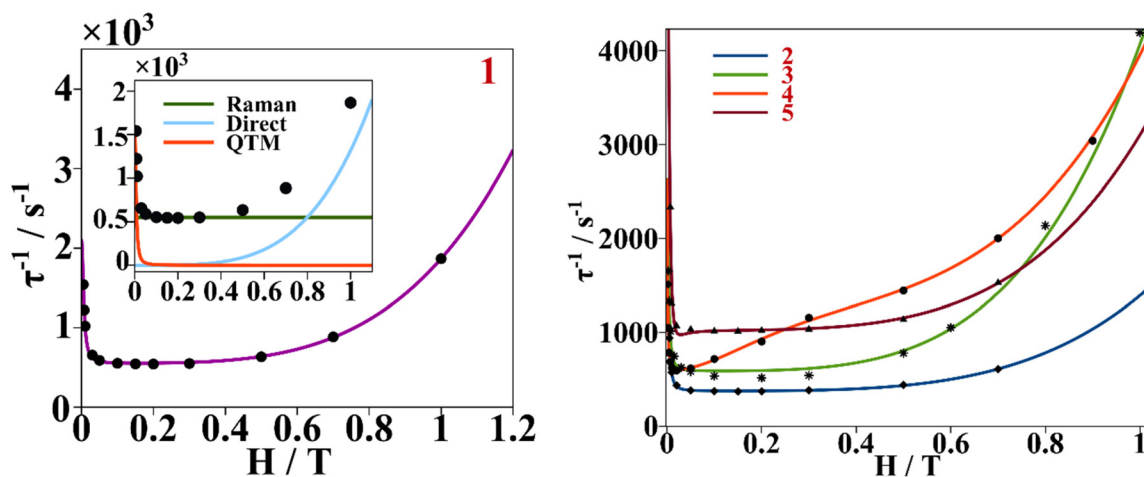


Fig. 10 Left, dependence of τ^{-1} on a static magnetic field (black dots) for compound **1**, continuous purple line accounts for the fitting using field dependent eqn (3). Inset shows the contribution of Raman, direct and QTM mechanisms to τ^{-1} vs. H for compound **1**. Right, dependence of τ^{-1} on a static magnetic field for compounds **2–5** respectively with eqn (3). The continuous blue, green, orange and red lines correspond to the fitting of the experimental magnetic data (discrete dots) of compounds **2** to **5** respectively with eqn (3).



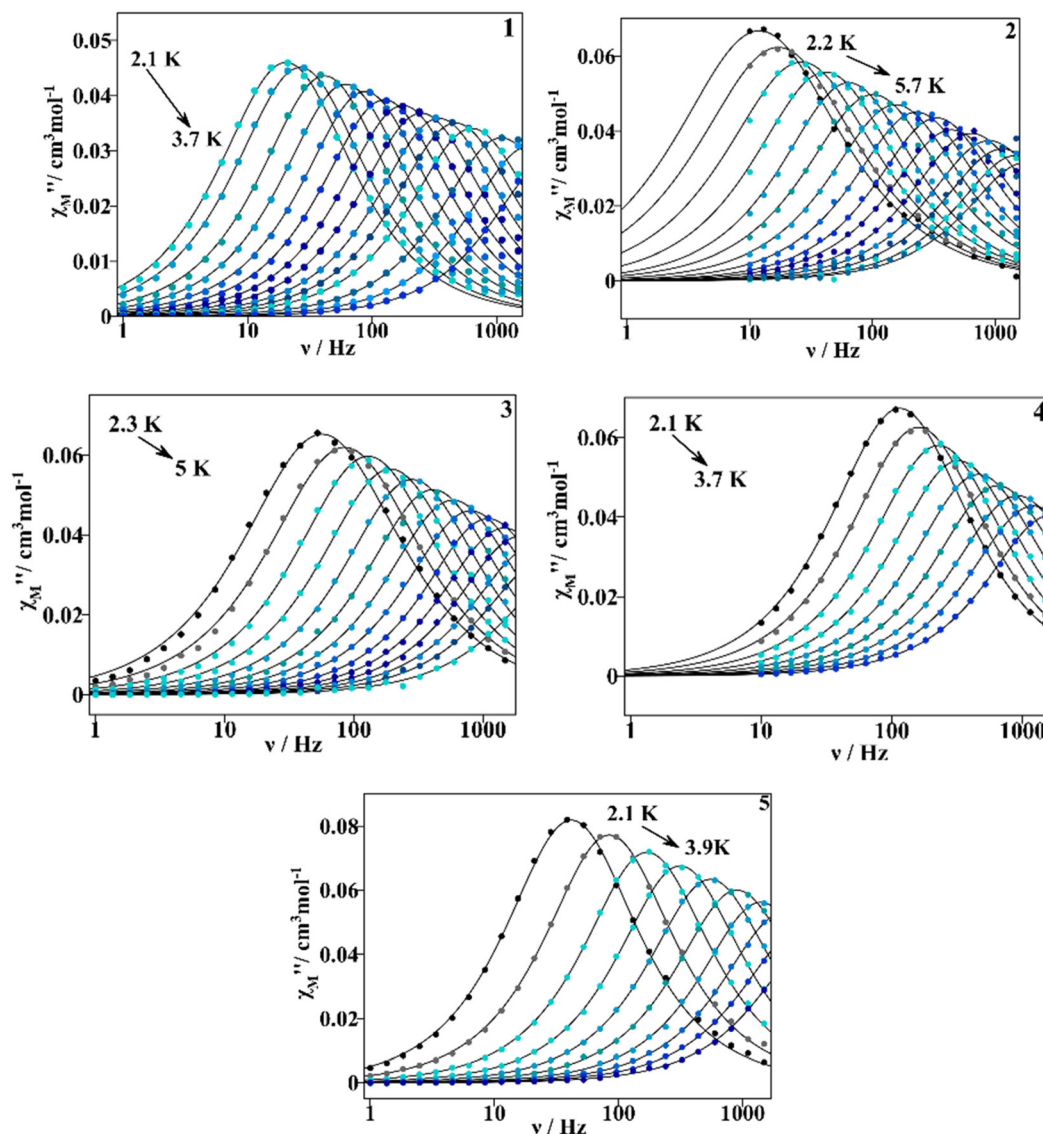


Fig. 11 χ_M'' vs. frequency plots for compounds 1–5 measured at an oscillating ac field of 4×10^{-4} T, and at a H_{dc} of 0.1 T for 1–3, and of 0.02 T for 4 and 5. The continuous lines correspond to the fitting with eqn (S1).†

sing spin relaxation. The third term of eqn (1) describes the fast QTM and, finally, the last term relates to the so called thermally assisted relaxation process, the Orbach process. It follows the Arrhenius law and experimentally, when the Orbach process takes place, a linear trend of $\log(\tau)$ with temperature is expected. This spin–lattice mechanism involves two phonons, and relaxation occurs through an effective energy barrier from the ground state $\pm m_j$ through a real excited state. Normally, the Orbach process takes place at a higher temperature range since the system needs enough energy to surpass the whole energy barrier between the $\pm m_j$ ground states.^{45,46}

For the title compounds a linear slope appears to be followed in the higher temperature range. We fit the experimental τ values of 1–5 using the Arrhenius law, and the values obtained for the activation energy barrier are in the range of

12.2–19.8 cm^{-1} with τ_0 values between 10^{-7} and 10^{-8} s, Table S13 and Fig. S16.† This linear trend could indicate that the relaxation of magnetization is taking place through an over-barrier Orbach mechanism for 1–5 or a thermally assisted tunneling process through the first $\pm m_j$ excited state. Nevertheless, for all the cerium(III) compounds found in the literature where *ab initio* calculations were performed, the energy differences between the $\pm m_j$ ground Kramers doublet (KD) and the first excited $\pm m_j$ KD are in the range of 258–339 cm^{-1} for mononuclear Ce^{3+} – β -diketonate complexes,^{20,21} 220–787 cm^{-1} for other Ce^{3+} –mononuclear compounds,^{23–28,47,48} 303–348 cm^{-1} for Ce^{3+} –polynuclear compounds^{30,49} and 180–303 cm^{-1} for Ce^{3+} –heterometallic compounds.^{29,34,50,51} Interestingly, the activation energy barrier calculated from experimental data (ΔE_{exp}), with the



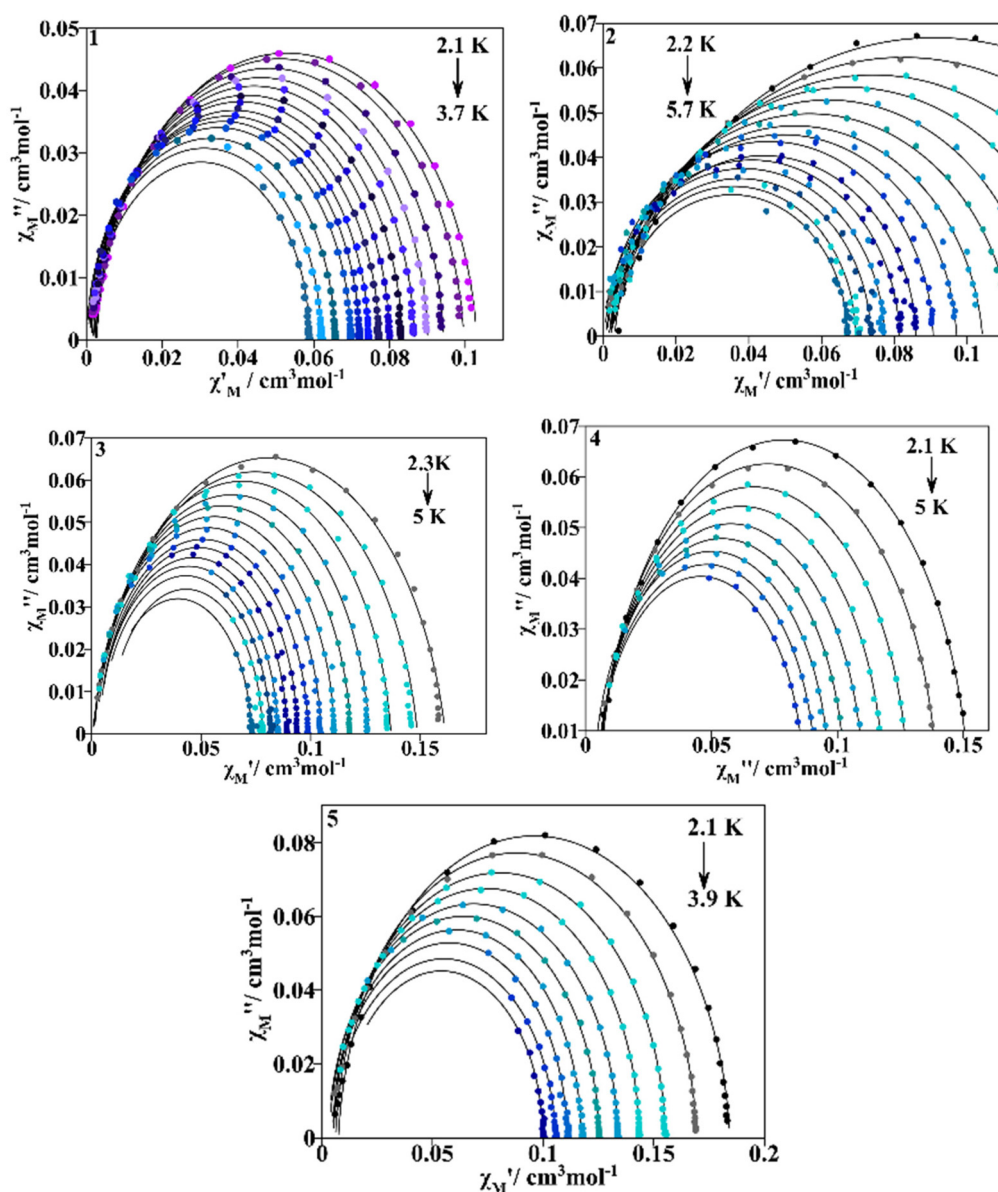


Fig. 12 Cole–Cole plots for compounds 1–5. The continuous black lines refer to the fitting with the generalized Debye model.

Arrhenius law, considering an Orbach mechanism, led to very similar values for all the previous Ce^{3+} compounds. The maximum value reported is 38 cm^{-1} ,²⁸ but the ΔE_{exp} values for the vast majority of published Ce^{3+} complexes lie in the $20\text{--}30 \text{ cm}^{-1}$ range and it appears that there is no difference between compounds presenting easy axial or easy plane symmetry.^{20,21,26,27,29,43,47,50,52,53} Besides, *ab initio* calculations (see the Computational results section) of the former cerium (iii) systems result in an energy difference between the ground and first excited m_j states much higher in energy ($151.9\text{--}372.6 \text{ cm}^{-1}$, see Table S14†) than the energy barrier obtained from experimental data using the Arrhenius law that describes the Orbach process.

The lack of a linear trend in the $\log(\tau)$ vs. T plots throughout the temperature range could be explained by considering the

presence of other magnetization relaxation mechanisms occurring at the same time as the Orbach process or taking into consideration that the Orbach process is not taking place at all, as a comparison of experimental data with *ab initio* calculations appears to indicate. Furthermore, hyperfine interactions between the electronic spin and the nuclear spin of a lanthanide ion can split the electronic doublet into different manifolds making new pathways for magnetization that could explain the linear trend with temperature. But the natural isotopes of cerium do not possess nuclear spin. Another proposed spin–lattice relaxation mechanism is the local mode, and it is referred to as a thermally dependent process, which considers that spin relaxation goes through a higher energy state that is a local vibration from the same m_j ground state following eqn (2). Then Δ_{loc} can be extracted and is



referred to as the local mode energy (cm^{-1}).⁵⁴ Δ_{loc} could be experimentally assigned from vibrational spectroscopic techniques.^{55,56}

$$\tau^{-1} = A_{\text{loc}} \left(\frac{e^{-\frac{\Delta_{\text{loc}}}{k_B T}}}{\left(e^{-\frac{\Delta_{\text{loc}}}{k_B T}} - 1 \right)^2} \right) \quad (2)$$

With the above exposed considerations, we tried to fit the magnetic data with the thermally dependent components of eqn (1) but by adding the local mode function instead of the Orbach one. Given that the temperature ranges over which there is a magnetic response for 1–5 are small and the risk of over-parametrization when many parameters are used in the equation, we first proceed to fit the dependence of τ^{-1} on the magnetic field so in this way we can extract the parameters from the equations that are field dependent: dT^n , e and f for Raman, B_1 and B_2 for the QTM and a for the direct process (eqn (3)) and then the parameters corresponding to these equations can be fixed in the temperature dependent function.

The Cole–Cole plots of the field dependent experimental magnetic data measured at a constant temperature were fitted using the generalized Debye model, Fig. S17 and Tables S15–S19.† The field dependent magnetic data could be fitted following eqn (3), for all the compounds, Fig. 10. In the τ^{-1} vs H plot, the different tendencies from the predominant QTM, Raman and direct mechanisms are well differentiated, first,

each part is fitted independently with the corresponding mechanism and the fitted parameters are used as the starting point for the whole set. The obtained parameters from the field dependent fitting of compounds 1–5 are compiled in Table 2. The inset in Fig. 10 and Fig. S18† show the contributions of each relaxation mechanism to magnetization relaxation along the magnetic fields of compounds 1–5.

$$\tau^{-1} = aH^4T + \frac{B_1}{1+B_2H^2} + d \left(\frac{1+eH^2}{1+fH^2} \right) T^n + K(T) \quad (3)$$

Then the temperature dependent experimental magnetic data were fitted with eqn (4) and the parameters n from the Raman process and c and ω from the local mode spin relaxation mechanism were obtained for 1–5. The τ^{-1} dependence on T plots are depicted in Fig. 13 and the obtained parameters from the best fit are compiled in Table 3. The contribution of each relaxation mechanism to τ^{-1} with temperature is shown in the inset of Fig. 13 and in Fig. S19.†

$$\tau^{-1} = A(H)T + K(H) + D(H)T^n + A_{\text{loc}} \left(\frac{e^{-\frac{\Delta_{\text{loc}}}{k_B T}}}{\left(e^{-\frac{\Delta_{\text{loc}}}{k_B T}} - 1 \right)^2} \right) \quad (4)$$

When looking at the dependence of τ^{-1} on the field, for all the compounds under low external bias dc magnetic fields, the QTM mechanism is the one dominating magnetization relaxation. Under high external dc fields for 1 and 3 the direct

Table 2 Values obtained from the fitting of τ^{-1} vs. H curves with the field dependent curve function: $\tau^{-1} = AH^4T + \frac{B_1}{1+B_2H^2} + d \left(\frac{1+eH^2}{1+fH^2} \right) T^n + K(T)$

Compound	Raman		Direct		QTM	
	dT^n (s^{-1})	e (T^{-2})	f (T^{-2})	a ($\text{s}^{-1} \text{T}^{-4}$)	B_1 (s^{-1})	B_2 (T^{-2})
1	549.21	82.78	22 760.23	526.71	1549.17	22 760.21
2	370.07	214.49	77 778.39	291.57	1691.25	77 778.40
3	575.89	1279.27	87 007.58	1394.18	2054.14	87 010.40
4	585.82	23.25	15.40	1270.65	2055.59	556 254.6
5	197.48	14 891.10	3561.34	832.12	5754.71	42 402.08

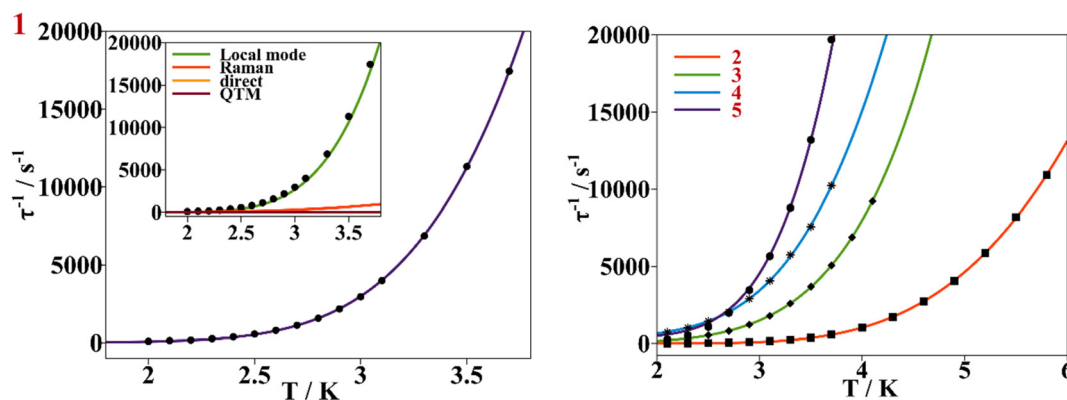


Fig. 13 Dependence of τ^{-1} on temperature obtained from the experimental magnetic data measured at an external magnetic field of 0.1 T for 1 and 3, of 0.05 for 2 and of 0.02 T for 4 and 5. Left, the plot of compound 1. Inset shows the contributions of each magnetization relaxation mechanism separately. Right, the plots for compounds 2–5.



Table 3 Values obtained from the fitting of τ^{-1} vs. H curves with the field dependent curve function:

$$\tau^{-1} = C \left(\frac{e^{-\frac{H}{T}}}{(e^{-\frac{H}{T}} - 1)^2} \right) T^n + D(H)T^n + A(H)T + K(H)$$

Compound	Raman n	Local mode	
		C (s ⁻¹)	ω (cm ⁻¹)/(K)
1	4.82	4.03×10^7	19.95/28.71
2	7.20	1.7×10^7	20.89/30.07
3	8.27	3.16×10^5	11.55/16.61
4	2.40	3.05×10^6	15.42/22.19
5	6.16	1.20×10^7	17.70/25.47

process is the dominant one while for 2, 4 and 5 the Raman process is still relevant, Fig. 10 and Fig. S18.†

From the global fit the obtained local mode energy values for the different compounds are in the range of 11.55–20.90 cm⁻¹ (16.61–30.07 K). With this approach, the spin relaxation of compound 1 appears to be governed by the local mode mechanism, Fig. 13 inset. While for 2, 4 and 5 spin relaxation over the temperature range is best described by both the Raman and local mode mechanisms, Fig. S19.† In fact for compound 4 it is well discerned that the Raman mechanism prevails in the lower temperature range while the local mode mechanism plays an important role in the higher temperature range, Fig. S19(4).† Moreover, compound 3 shows a smaller contribution of the local mode (ω), which is the smallest of the presented compounds with a value of 11.55 cm⁻¹ (16.61 K), but there is a higher contribution from the n parameter from the Raman process ($n = 8.2$) and the direct mechanism (it has the highest $A(H)$ value). This suggests that for 3, spin relaxation is mostly taking place through the Raman and direct processes.

Additionally, the temperature dependence of τ could also be fitted with only the Raman term as the $\ln(\tau)$ vs. $\ln(T)$ plots lead to straight lines for the magnetic data for all compounds, Fig. S20.† A fitting of these plots taking into account just Raman relaxation of the magnetization mechanism described by the $\ln(\tau) = -n \cdot \ln(T) + \ln(C)$ equation leads to good values of n and C , which are $n = 8.9$ and $C = 5.9$ for 1, $n = 7.0$ and $C = 17.8$ for 2, $n = 5.7$ and $C = 0.3$ for 3, $n = 4.7$ and $C = 0.0519$ for 4 and $n = 7.5$ and $C = 0.9$ for 5. This leads us to consider that the Raman process may actually be the relevant mechanism for the magnetic relaxation of these cerium(III) compounds. However, a fitting of temperature dependent data with eqn (3) without considering the local mode but taking into account

the contribution of all the field dependence terms, leads to unsuccessful fitting of the data, Fig. S21.† Therefore, two different approaches are presented to explain the spin dynamics of 1–4.

The first is that, when considering the contributions of QTM and direct processes to magnetic relaxation, this would lead to overparametrization, so the parameters for those mechanisms are first obtained from the field dependent magnetic data and fixed in the temperature dependent fitting. In this approach, the local mode has to be considered to obtain a good fitting of magnetization relaxation times with temperature since only considering the Raman process leads to unsuccessful fitting. The second approach is just to consider the Raman relaxation mechanism and neglect the contribution that direct and QTM mechanisms may have on spin relaxation; this also leads to reasonable results. Still the contributions of the QTM and direct mechanisms to spin relaxation should not be overlooked, especially the direct mechanism, if an external magnetic field is applied to measure the alternating current magnetic susceptibility.

The existence of multiple potential explanations to describe the spin dynamics phenomenon of compounds 1–5, once again, highlights the need for continued investigation into the magnetic behavior, at the molecular level, of Ce(III) coordination compounds.

Geometry optimization and calculation of the DFT vibrational frequencies were performed for 1, 2 and 4. The analysis of the frequencies shows a large amount of low energy vibrational frequencies (Table S20†) that can contribute to spin relaxation through the local-mode process; however, a spin-phonon calculation of these frequencies to verify their contribution is out of the scope of this paper.

Computational results

Ab initio calculations were performed on the crystallographic structures, see the Computational details section. CASSCF calculations incorporating spin-orbit effects *via* the RASSI approach were conducted to delve into the magnetic characteristics of compounds 1–5. Cerium(III) complexes exhibit a $4f^1$ and a $^2F_{5/2}$ ground state. The simulated magnetic susceptibility and magnetization profiles are similar to the experimentally observed curves, see Fig. S22 and S23,† indicating good agreement between theoretical predictions and actual experimental results.

Except compound 1, the remaining systems studied show a large g_z component, see Table 4, indicating large axial anisotropy. The values are similar to those for other previously

Table 4 Calculated components of the g tensor and wavefunction analysis of the ground Kramers doublet state at the CASSCF-RASSI level

Compound	g_x	g_y	g_z	Wavefunction
1	0.563	1.391	2.392	$0.28 \pm 5/2 \rangle + 0.42 \pm 3/2 \rangle + 0.30 \pm 1/2 \rangle$
2	0.108	0.481	3.754	$0.90 \pm 5/2 \rangle + 0.005 \pm 3/2 \rangle + 0.09 \pm 1/2 \rangle$
3	0.460	0.940	3.600	$0.87 \pm 5/2 \rangle + 0.04 \pm 3/2 \rangle + 0.09 \pm 1/2 \rangle$
4	0.775	1.255	3.379	$0.89 \pm 5/2 \rangle + 0.08 \pm 3/2 \rangle + 0.03 \pm 1/2 \rangle$
5	0.501	1.500	3.350	$0.82 \pm 5/2 \rangle + 0.04 \pm 3/2 \rangle + 0.14 \pm 1/2 \rangle$



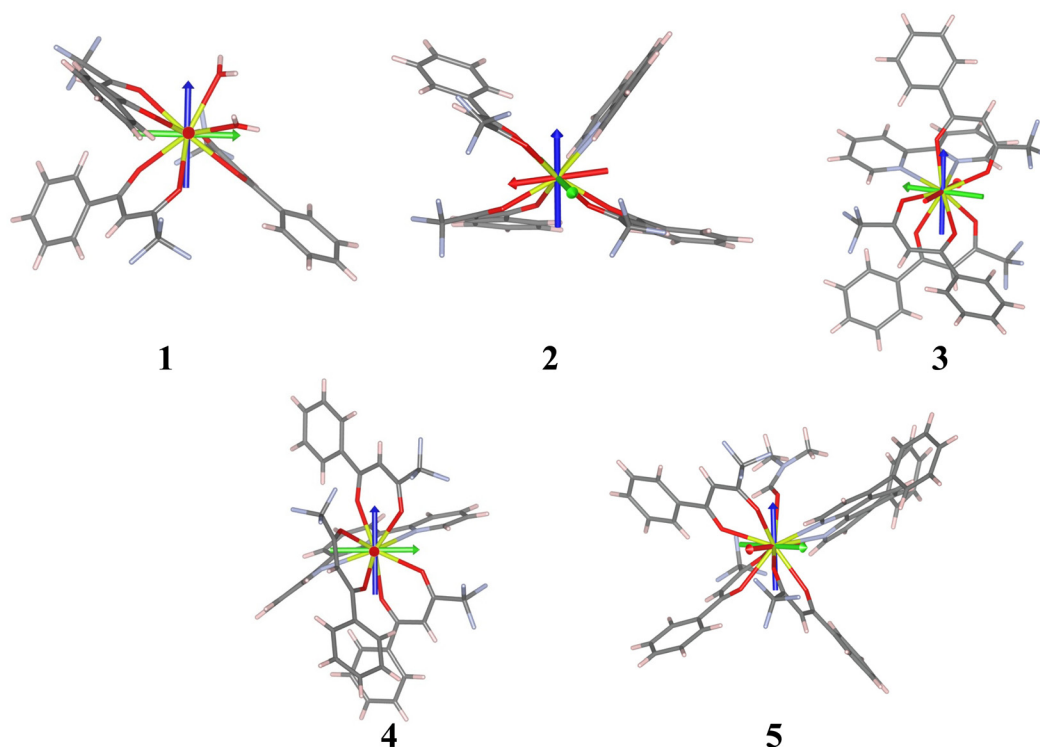


Fig. 14 Calculated orientations of the g tensor of the ground Kramers doublet, where g_z , g_y and g_x components are represented by blue, green and red arrows, respectively. Cerium, fluorine, oxygen, nitrogen, carbon and hydrogen are represented in yellow, light blue, red, blue, gray and pink.

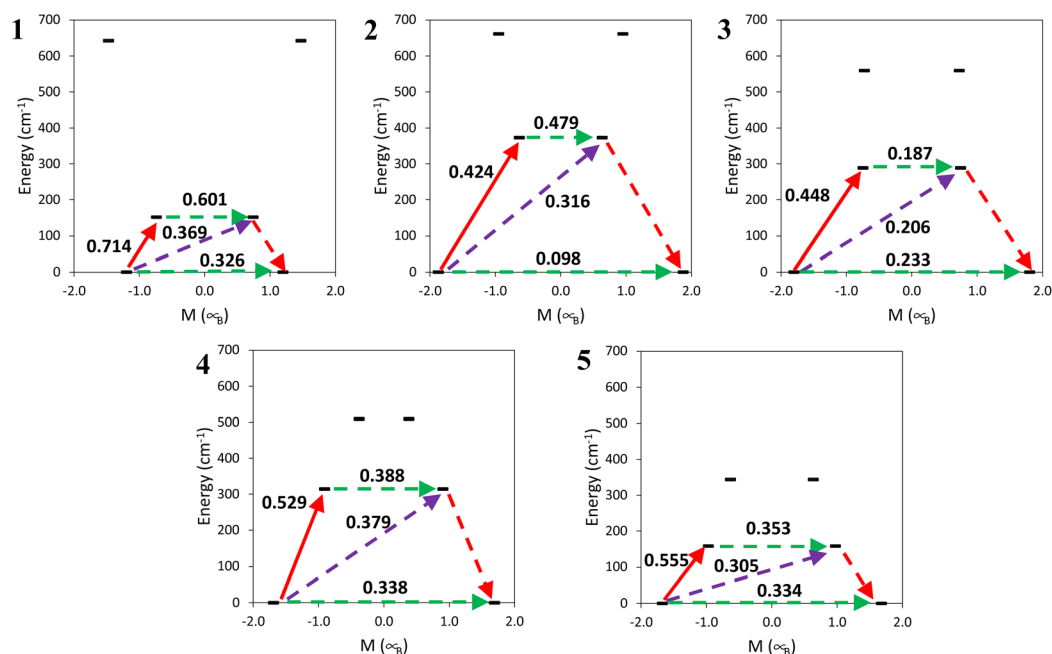


Fig. 15 Calculated state energies as a function of their magnetic moment, M_z , along the main anisotropy axis. The arrows correspond to the quantum tunneling mechanism of ground and first excited states (green), the hypothetical Orbach relaxation process (purple) and the transition between the ground and first excited states (red). The values close to the arrows indicate the matrix elements of the transition magnetic moments (efficient spin relaxation mechanism is expected above 0.1).



reported Ce systems (see Tables S1–S5†); however, for a perfectly axial Ce^{3+} complex the value should be 4.19 (30/7). Compounds **2–4** with values in the range of 3.3–3.8 are consistent with an $m_J = \pm 5/2$ ground state with a contribution from $m_J = \pm 3/2$ states. The orientation of the calculated g_z is quite different for the compounds (see Fig. 14). The coordination numbers are also different, eight for **1–3**, and nine for **4** and **5**. In the case of an $m_J = \pm 5/2$ ground state an oblate f electron density, with a perpendicular g_z orientation, is expected. The g_z tensor would be oriented in a way that the metal–ligand electrostatic repulsion would be reduced. It can be achieved by (i) the alignment of g_z with the shortest Ce–O distances, in a way that the f electron density disc stays perpendicular to that direction or (ii) the location of the f electron density disc in the plane containing the longest Ce–L distances. In this family of compounds in general g_z is located in a way that it is not pointing to any ligand and the electron density disc tries to avoid all the ligands. This happens for **1**, **2**, **4** and **5**. However, for **3** it is mainly oriented with one of the Ce–O bonds (Ce1–O5) and with the f electron density disc oriented close to the Ce–N bonds. In the case of **4** g_z is oriented in a way that the f electron density disc avoids most of the ligands but is in the same plane as some of the Ce–N bonds of the terpyridine ligand.

The analysis of the decomposition of the RASSI wavefunction of the ground Kramers doublet can be found in Table 4. It shows, as expected, a different contribution for **1**, where a large mixture of the three components is observed; this correlates with smaller g_z and larger g_x and g_y components. For the other compounds a larger contribution from the wave function $|\pm 5/2\rangle$ (80–90%) mixed with $|\pm 1/2\rangle$ and $|\pm 3/2\rangle$ from the other excited m_J doublets is found; this also explains the larger anisotropy of these Ce complexes.

The energy of the Kramers doublet and transition probabilities between the states were calculated, see Fig. 15. The first excited Kramers doublet is around 150 cm^{-1} in **1** and **5** and around 300 cm^{-1} or more for the other complexes, see also Table S14.† In all the cases, as commented on before, the energy difference is much larger than the energy barrier that can be obtained from the fit of the dependence of relaxation times on temperature with the Arrhenius equation, $12\text{--}20\text{ cm}^{-1}$. All the complexes show a large tunnelling probability in the ground state, with complex **2** having the smallest one, of 0.1. This agrees with the necessity to apply an external dc field to show an out-of-phase magnetic susceptibility signal (χ_M''). Besides **2** having the smallest tunnelling probability in this family of compounds the optimal field needed is not the smallest one. This is probably also related to the shorter Ce...Ce distance in **2** in comparison with **4** and **5**.

Conclusions

As a continuation of our interest in the less studied lanthanide (III) SIM compounds derived from Ce^{3+} ions,²¹ in this paper we have presented five new β -diketonate– Ce^{3+} coordination com-

pounds with the aim of providing new insights into understanding the cerium spin relaxation. The five structurally characterized new Ce^{3+} complexes are derived from the β -diketonate HBTfa = 4,4,4-trifluoro-1-phenyl-1,3-butanedione with polypyridyl molecules as ancillary ligands: $[\text{Ce}(\text{Btfa})_3(\text{H}_2\text{O})_2]$ (**1**), $[\text{Ce}(\text{Btfa})_3(\text{phen})]$ (**2**), $[\text{Ce}(\text{Btfa})_3(\text{bipy})]$ (**3**), $[\text{Ce}(\text{Btfa})_3(\text{terpy})]$ (**4**) and $[\text{Ce}(\text{Btfa})_3(\text{bathophen})(\text{DMF})]$ (**5**), where phen = 1,10-phenanthroline, bipy = 2,2'-bipyridine, terpy = 2,2':6',2''-terpyridine and bathophen = 4,7-diphenyl-1,10-phenanthroline. In the new reported compounds, Ce^{3+} displays a coordination number of 8 for **1**, **2** and **3** and a coordination number of 9 for **4** and **5**. Moreover, the five compounds have been magnetically characterized and a meticulous study has been carried out to further understand the spin dynamics in the magnetization relaxation of these new field induced SMM compounds. Furthermore, *ab initio* calculations were performed for the former compounds. For all the compounds, the Orbach mechanism was disregarded in light of the published literature and the obtained *ab initio* results. The local mode mechanism was used instead. The calculations also show the axially of the compounds and corroborate the presence of low energy vibrational modes that could contribute to spin relaxation through the local mode. Finally, a compilation of the cerium(III) coordination complexes found in the literature before 2023 and showing slow magnetization relaxation is given in Tables S1–S5.† From the published experimental data and *ab initio* calculations, we can infer that the stabilization of m_J with the highest value as the ground state does not imply the presence of SMM behavior for the previously magnetically studied Ce^{3+} coordination compounds.

Conflicts of interest

There are no conflicts to declare.

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