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A Pseudo-Octahedral Cobalt(II) Complex with *bis*-Pyrazolylpyridine Ligands Acting as a Zero-Field Single-Molecule Magnet with Easy Axis Anisotropy

Luca Rigamonti,^{*,[a]} Nathalie Bridonneau,^[a,Δ] Giordano Poneti,^[b,∘] Lorenzo Tesi,^[b] Lorenzo Sorace,^[b] Dawid Pinkowicz,^[c] Jesus Jover,^[d] Eliseo Ruiz,^[d] Roberta Sessoli,^[b] and Andrea Cornia^[a]

- [a] *Dipartimento di Scienze Chimiche e Geologiche, Università degli Studi di Modena e Reggio Emilia and INSTM RU of Modena and Reggio Emilia, via G. Campi 103, 41125 Modena (Italy)*
E-mail: luca.rigamonti@unimore.it, luca.rigamonti@yahoo.com
- [b] *Laboratory of Molecular Magnetism (LAMM), Dipartimento di Chimica 'Ugo Schiff', Università degli Studi di Firenze and INSTM RU of Firenze, via della Lastruccia 3-13, 50019 Sesto Fiorentino, FI (Italy)*
- [c] *Faculty of Chemistry, Jagiellonian University, Gronostajowa 2, 30-387 Kraków (Poland)*
- [d] *Departament de Química Inorgànica i Orgànica and Institut de Química Teòrica i Computacional, Universitat de Barcelona, Diagonal 645, 08028 Barcelona (Spain)*
- [Δ] *current address: Laboratoire Interfaces Traitements Organisation et DYnamique des Systèmes (ITODYS), UMR 7086 CNRS, Université Paris 7 Diderot, Paris, Bât. Lavoisier, 15 rue Jean-Antoine de Baïf, 75205 Paris Cedex 13 (France)*
- [∘] *current address: Instituto de Química, Universidade Federal do Rio de Janeiro, 21941-909 Rio de Janeiro (Brazil)*

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Authors' ORCIDs: Luca Rigamonti: 0000-0002-9875-9765, Nathalie Bridonneau: 0000-0003-3789-9024, Giordano Poneti: 0000-0002-1712-4611, Lorenzo Tesi: 0000-0003-4001-8363, Lorenzo Sorace: 0000-0003-4785-1331, Dawid Pinkowicz: 0000-0002-9958-3116, Jesus Jover: 0000-0003-3383-4573, Eliseo Ruiz: 0000-0001-9097-8499, Roberta Sessoli: 0000-0003-3783-2700, Andrea Cornia: 0000-0001-9765-3128.

Abstract: The homoleptic mononuclear compound [Co(bpp-COOMe)₂](ClO₄)₂ (**1**) (bpp-COOMe = methyl 2,6-di(pyrazol-1-yl)pyridine-4-carboxylate) crystallizes in the monoclinic *C2/c* space group, and the cobalt(II) ion possesses a pseudo-octahedral environment given by the two *mer* coordinated tridentate ligands. Direct current magnetic data, together with single-crystal torque magnetometry and EPR measurements, disclosed the easy axis nature of this cobalt(II) complex, which shows

Single-Molecule Magnet behaviour when a static field is applied in alternating current susceptibility measurements. Diamagnetic dilution in the zinc(II) analogue, $[\text{Zn}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$ (**2**), afforded the derivative $[\text{Zn}_{0.95}\text{Co}_{0.05}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$ (**3**) that exhibits slow relaxation of magnetization even in zero field thanks to the reduction of dipolar interactions. Theoretical calculations confirmed the overall electronic structure and the magnetic scenario of the compound as drawn by experimental data, confirming the spin-phonon Raman relaxation mechanism, and a direct quantum tunnelling in the ground state as the most plausible relaxation pathway in zero field.

Introduction

Single-Molecule Magnets (SMMs) can in principle be used for encoding binary information and for data processing, exploiting their magnetic bistability and inherent quantum behaviour.^[1] Research in this field moves in different directions, such as finding better performing systems,^[2,3] developing reliable methods to deposit molecules on solid substrates,^[4,5] and finding ways to control molecular magnetism by external stimuli, including light and pressure.^[6–8] Complexes containing only one metal centre may represent the smallest chemically-tuneable SMMs for spin-based devices,^[9] and significant progresses have been made in this regard by using both lanthanides^{3,10} and, more recently, 3d metal ions.^[10,11] In the latter case, relaxation by quantum tunnelling is usually very fast in zero field, so that application of a static field is required to observe SMM behaviour by alternating current (ac) susceptibility measurements. However, few examples of zero-field SMMs based on mononuclear complexes of iron(I)^[12] iron(III),^[13] and cobalt(II) have been reported,^[14–25] which all feature half-integer spin states (3/2 or 5/2). In particular, individual cobalt(II) ions were found to exhibit zero-field SMM behaviour when embedded in tetrahedral,^[14–17,20,22,24] trigonal prismatic^[18,19,23] or linear^[21] coordination environments, or in a solid state lattice.^[25] In all these cases, the metal complexes possess a negative zero field splitting (*zfs*) parameter *D*, thus suggesting to look for other coordination environments suitable to enforce an easy axis magnetic anisotropy on the cobalt(II) ion.

The bpp (2,6-bis(pyrazol-1-yl)pyridine) class of ligands are well known to promote spin crossover (SCO) in their octahedral iron(II) metal complexes.^[26–28] These ligands can host both high spin (HS) and low spin (LS) iron(II) centres,^[27] thus enabling spin state modulation by temperature or application of light.^[29,30] In some cases, the distortion of the octahedral geometry is so pronounced as to block the metal centre in its HS state, as we recently found using bpp-COOMe (methyl 2,6-di(pyrazol-1-yl)pyridine-4-carboxylate).^[26] In particular, in the distorted molecular

structure of $[\text{Fe}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$, the pseudo-octahedral coordination environment leads to a stabilization of the d_{xy} orbital relative to d_{xz} and d_{yz} (where z represents the *trans*-N{pyridyl}–Fe–N{pyridyl} direction, and x and y the directions of the coordinated pyrazolyl nitrogen atoms).^[26]

Such orbital pattern is expected to produce an easy axis anisotropy for cobalt(II), as sought for.^[31] We have thus synthesized $[\text{Co}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$ (**1**) by reaction of cobalt(II) perchlorate with bpp-COOMe and found this compound to be isostructural to the iron(II) complex.^[26] From the study of its magnetic properties we positively discovered that **1** has an easy axis anisotropy and shows SMM behaviour under an applied external field. Magnetic dilution in its zinc(II) analogue, $[\text{Zn}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$ (**2**), afforded $[\text{Zn}_{0.95}\text{Co}_{0.05}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$ (**3**), where dipolar interactions are reduced and slow relaxation of the magnetization is detectable even in zero field. To the best of our knowledge, this discloses the first fully-characterized pseudo-octahedral cobalt(II) complex with easy-axis anisotropy acting as a zero-field SMM.^[32] Here we present the detailed and critical description of the experimental data collected by direct current (dc) and ac magnetic measurements, X-band Electron Paramagnetic Resonance (EPR) spectroscopy and single-crystal Cantilever Torque Magnetometry (CTM), together with *ab initio* theoretical calculations supporting the magnetic properties of **1**.

Results and Discussion

Synthesis, crystallization and molecular structures

Reaction of a suspension of bpp-COOMe in acetonitrile with $\text{M}(\text{ClO}_4)_2 \cdot 6\text{H}_2\text{O}$ ($\text{M} = \text{Co}, \text{Zn}$, either as pure salts or in mixture) at room temperature produced the immediate dissolution of the ligand and formation of a solution, which gave well-formed X-ray quality crystals of **1** (light red), **2** (colourless) or **3** (very light orange) in good yields by vapour diffusion with diethyl ether. The crystals of the three compounds are air stable, belong to monoclinic space group $C2/c$ and are isomorphous to the iron(II) derivative recently studied for its SCO properties (see Supporting Information, SI, Tables S1 and S2, for a comparison of the room temperature unit cell parameters).^[26] X-ray diffraction data were collected at 295 and 161 K for **1** and at 120 K for **2** in order to determine their molecular structures, and the main bond distances and angles so obtained are reported in Table 1.

In all cases, metal coordination environments show pronounced distortion from perfect octahedral geometry (Figure 1 and SI, Figure S1). This is best outlined by the *trans*-N{pyridyl}–M–N{pyridyl} angle (ϕ) and by the dihedral angle between the least squares (l.s.) planes of the two

mer-coordinated ligands (θ), which lie far from the ideal 180° and 90° values, respectively.^[27] Although the cobalt(II) ion in **1** is Jahn-Teller active,^[33] the described distortion is most probably caused by crystal-packing effects, since it is present to almost the same extent in the zinc(II) derivative **2**. Notice that the most regular geometry would still exhibit departures from perfect octahedral symmetry due to the intrinsic shape of bpp ligands, which imposes a clamp angle $\psi < 180^\circ$. The distortion is also highlighted by the high values of the parameters Σ and Θ (Table 1). The first parameter is a general measure of the deviation of a metal ion from an ideal octahedral geometry, while Θ more specifically indicates its distortion from an octahedral towards a trigonal prismatic structure.^[28] In fact, a perfectly octahedral complex would give $\Sigma = \Theta = 0$.

The Co–N bond distances in **1** (Table 1) range from 2.08 Å with the pyridyl nitrogen atom (ring C) to about 2.15 Å with the pyrazolyl moieties (rings A and B), which are partially rotated by about $6\text{--}10^\circ$ with respect the l.s. plane of the whole ligand; these values remain the same within experimental error upon variation of the temperature. The ester group is coplanar with the pyridyl ring C and is involved in one of the two shortest intermolecular interactions with one pyrazolyl C–H group (at 161/295 K, $\text{C1(H1)}\cdots\text{O17} = 3.252/3.291$ Å, $\text{C1–H1}\cdots\text{O17} = 147.6/148.9^\circ$, which leads to a Co \cdots Co distance of 9.642 (9.664) Å. The second short intermolecular contact involves another C–H group and the pyrazolyl ring of the adjacent molecule (at 161/295 K: $\text{C12(H12)}\cdots\text{centroid}(\text{ring A}) = 3.47/3.52$ Å, $\text{C12–H12}\cdots\text{centroid}(\text{ring A}) = 150/149^\circ$) with a shorter intermolecular Co \cdots Co distance of 8.494/8.530 Å. Perchlorate anions are located in the voids between cations and all their oxygen atoms are involved in weak C–H \cdots O interactions (SI, Figure S1 and Tables S3 and S4).

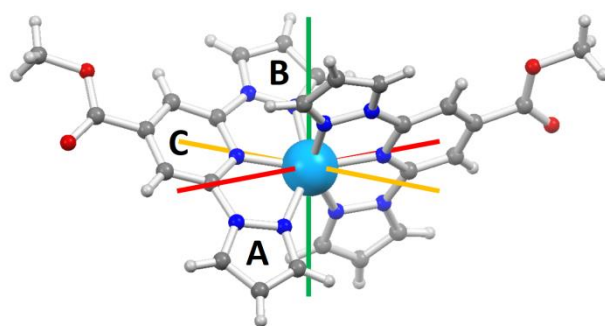


Figure 1. Molecular structure of the cation $[\text{Co}(\text{bpp-COOMe})_2]^{2+}$ of **1** at low T in a perspective view with ring labelling (ring A = N1N4C3C2C1, ring B = N14N10C11C12C13, ring C = N15C5C6C7C8C9, where N1, N14 and N15 are the donor atoms to cobalt) and magnetic anisotropy axes as obtained by single-crystal CTM and EPR measurements. Colour code: Co = azure, O = red, N = blue, C = light grey, H = white; easy axis = red, intermediate axis = green, hard axis = yellow.

The zinc derivative **2** shows similar structural features compared to **1**, with only slightly longer Zn–N distances with pyrazolyl rings (Table 1) but fully comparable crystal packing (SI, Table S5). This allowed to successfully obtain a diamagnetically-diluted crystalline sample **3** containing a Zn:Co = 18:1 ratio (i.e. $\text{Zn}_{0.9475}\text{Co}_{0.0525}$), as confirmed by ICP-OES measurements.

Table 1. Main bond lengths (Å) and angles (°) of **1** (Co) and **2** (Zn).

	1 ($T = 295$ K)	1 ($T = 161$ K)	2 ($T = 120$ K)
M1–N1	2.142(2)	2.141(2)	2.1428(16)
M1–N14	2.148(2)	2.152(2)	2.2056(16)
M1–N15 (py)	2.082(2)	2.080(2)	2.1216(16)
N1–M1–N15	74.56(9)	74.73(8)	74.43(6)
N15–M1–N14	74.91(9)	74.93(8)	73.49(6)
N1–M1–N14 (ψ)	148.76(9)	148.99(8)	147.72(6)
N15–M1–N15 ^a (ϕ)	161.12(13)	161.45(8)	160.36(6)
θ^b	81.21	81.59	82.92
Σ^c	140.1	139.1	145.6
Θ^c	445	444	457

^a primed atom is obtained from unprimed one through twofold rotation;
^b dihedral angle between the two ligands (the plane of each ligand was defined as the least-squares plane through its sixteen aromatic C/N atoms); ^c see ref. 28 for its definition.

Static magnetic properties

Dc magnetic susceptibility data of **1** were recorded in the 2–300 K temperature range in a low field regime (1 kOe up to 40 K and 10 kOe from 30 to 300 K), together with the field dependence of the isothermal molar magnetization, M_M , at low temperature (Figure 2). At 300 K, the product of the molar magnetic susceptibility, χ_M , with the temperature, T , is about $2.84 \text{ emu K mol}^{-1}$, against an expected spin-only value of $1.875 \text{ emu K mol}^{-1}$ for an $S = 3/2$ ion with $g = 2.00$. This difference indicates a relevant orbital contribution to the magnetism, which can be parametrized in first approximation by assuming $g = 2.46$. The $\chi_M T$ value remains almost unvaried upon cooling down to 90 K, where it begins to slowly decrease to $1.98 \text{ emu K mol}^{-1}$ at 2.0 K, consistent with the depopulation of the spin-orbit and low-symmetry split states of the $^4\text{T}_{1g}(\text{O}_h)$ electronic term.

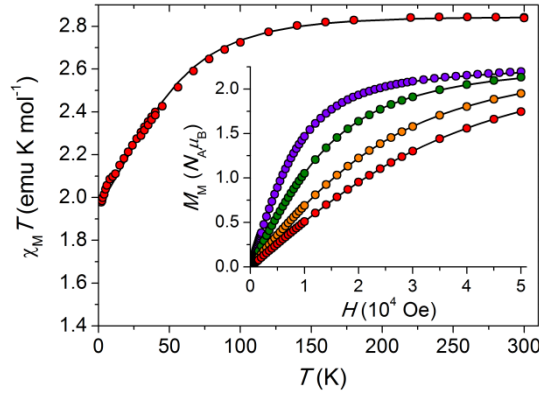


Figure 2. Temperature dependence of the $\chi_M T$ product for **1** (red circles in the main panel, overlapped points in the 30–40 K range are referred to measurements at both 1 and 10 kOe); inset: field dependence of the molar magnetization, M_M , at 1.8 (blue), 3.0 (green), 5.0 (yellow) and 7.0 (red) K. Solid lines are given by the best-fit parameters (see text).

While a spin Hamiltonian formalism might be inadequate to account for the electronic structure of the cobalt(II) ion,^[34] we first used the following Hamiltonian to fit simultaneously the experimental $\chi_M T(T)$ and $M_M(H)$ data with PHI program.^[35]

$$\begin{aligned} \mathcal{H} &= \mathcal{H}_{Zeeman} + \mathcal{H}_{fs} = \mu_B \hat{\mathbf{B}} \cdot \bar{\mathbf{g}} \cdot \hat{\mathbf{S}} + \hat{\mathbf{S}} \cdot \bar{\mathbf{D}} \cdot \hat{\mathbf{S}} \\ &= g_x \mu_B B_x S_x + g_y \mu_B B_y S_y + g_z \mu_B B_z S_z + D \left[S_z^2 - \frac{1}{3} S(S+1) \right] + E(S_x^2 - S_y^2) \end{aligned} \quad (1)$$

where D and E are the axial and transverse second-order anisotropy parameters (defined from the principal components of the $\bar{\mathbf{D}}$ matrix as $D = 3D_{zz}/2$ and $E = (D_{xx} - D_{yy})/2$), and g_x , g_y and g_z are the principal components of the $\bar{\mathbf{g}}$ matrix for the $S = 3/2$ state ($\bar{\mathbf{D}}$ and $\bar{\mathbf{g}}$ were assumed to be collinear). Taking an axial $\bar{\mathbf{g}}$ matrix with $g_x = g_y$ for simplicity, a systematic survey was performed with g_α ($\alpha = x, y, z$) values from 1.8 to 3.2 and diagonal components of $\bar{\mathbf{D}}$ from -90 to 90 cm^{-1} . An accurate fit was only possible for negative D values and a unique solution fulfilled the condition expected for easy axis cobalt(II) ions, namely $g_z > g_x = g_y$: $D = -57.5(7) \text{ cm}^{-1}$, $|E| = 15.7(3) \text{ cm}^{-1}$, $g_x = g_y = 2.332(4)$, $g_z = 2.6687(15)$. As can be noticed, the cobalt(II) ion has an easy-axis anisotropy but with a relevant rhombic distortion as given by $|E/D| = 0.273$ (the sign of E is irrelevant for powder measurements when $g_x = g_y$). The resulting splitting of the $S = 3/2$ manifold is $2(D^2 + 3E^2)^{1/2} = 127 \text{ cm}^{-1}$.

Torque magnetometry

To gain independent experimental evidences of the magnetic anisotropy, CTM experiments were performed on a face-indexed single crystal of **1** mounted on a rotating capacitive cantilever

torquemeter. The device measures the torque component (τ_Y) along the rotation axis (Y) whereas the magnetic field is applied normal to it in the XZ plane at an angle θ from Z (θ increases going from Z towards $-X$).^[36,37] The measured torque is thus given by:

$$\tau_Y = m_Z B_X - m_X B_Z \quad (2)$$

where $\mathbf{m} = (m_X, m_Y, m_Z)$ is the magnetic moment of the sample and $\mathbf{B} = (B_X, 0, B_Z)$ is the applied magnetic field. Two rotations (Rot1 and Rot2) were performed with the XYZ frame corresponding to the orthogonalized crystallographic frame a^*bc (Rot1) or $-a^*cb$ (Rot2), see ESI for further details. Thanks to the fact that the cobalt(II) ion in **1** is located on a binary axis of monoclinic $C2/c$ space group, only one molecule is magnetically independent. Moreover, one of the principal directions of the magnetic anisotropy tensor (say, y) must coincide with this binary axis, which is parallel to b , while the others (x and z) must lie in the a^*c plane. When scanning this plane, as in Rot1, and working in the low-field limit eq 2 becomes:

$$\tau_Y(\phi) = B^2(\chi_{zz} - \chi_{xx})\sin\phi\cos\phi \quad (3)$$

where $\bar{\chi}$ is the susceptibility tensor and $\phi = \theta_z - \theta$ is the angle between the principal anisotropy axis z and \mathbf{B} (i.e. at $\theta = \theta_z$ the magnetic field is applied along z). The principal directions of the anisotropy tensor, separated by $\pi/2$ but arbitrarily located in the a^*c plane are easily detected as zero-torque points. Furthermore their easy or hard character becomes evident by increasing the field at low temperature: when the low-field limit is no longer valid the curves become steeper around the hard-axis zero-torque point.^[37] The measurements, performed at the temperature of 10 K under a static magnetic field of 2 and 4 T, and at 100 K with a magnetic field of 10 and 12 T, are reported in Figure 3, while data at intermediate temperatures are available in SI, Figure S2.

In Rot1 ($XYZ = a^*bc$) at 10 K we find the first torque zero at $\theta = 43^\circ$ going from c toward $-a^*$; thus the first in-plane principal direction (x) deviates by only ca. 10° from the a crystallographic axis (SI, Figure S3). The second torque zero occurs at $\theta = 133^\circ$ and corresponds to the second in-plane principal direction z ($\theta_z = 133^\circ$). The sign of the measured torque signal directly indicates that z (x) is the easy (hard) in-plane direction. The same conclusion can be reached looking at the 4 T data. At this field the torque curve deviates from the simple angular dependence of eq 3, showing a steeper angular variation $\theta = 43^\circ$, which is identified as the hard direction in the scanned plane.

For Rot2 ($XYZ = -a^*cb$) the two torque zeros are found at $\theta = 0^\circ$ and 90° as expected, i.e. when the field is applied along b and a^* crystallographic axes, respectively. Unlike the former, the latter field orientation does not individuate a principal anisotropy axis but only its projection on the a^*b plane. Both the sign and the shape of the torque signal in this rotation indicate that the crystal is more easily magnetized along a^* than along b . However, from a qualitative analysis it is not

possible to establish whether b is the overall intermediate or hard anisotropy axis. To solve this ambiguity, the data at $T = 10$ K were numerically simulated with the $S = 3/2$ spin Hamiltonian in eq 1. As a first step, \bar{g} and \bar{D} matrices were constrained to have the principal values determined by dc magnetic measurements, but their common principal directions were left free to reorient in space and an overall scale factor was refined to account for the incertitude on the mass of the crystal (ca. 80 μ g). In a second step, the principal values were also left free to vary, but this did not provide any improvement in the agreement.

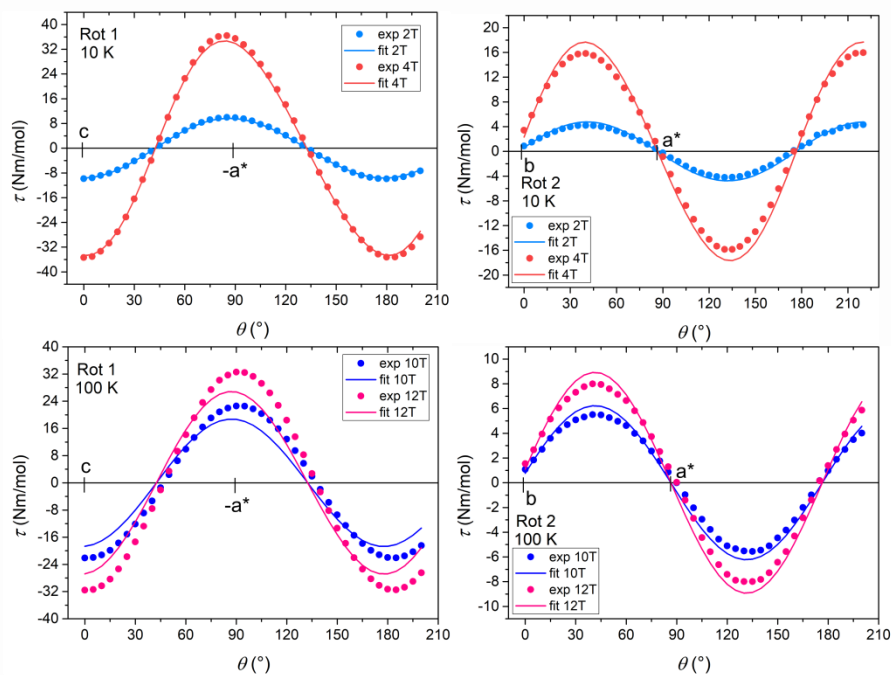


Figure 3. Experimental (dots) data and simulated curves with $S = 3/2$ spin Hamiltonian (solid lines, N m mol⁻¹) of **1** obtained with CTM for two rotations at 10 and 100 K under different static magnetic fields. The common orientation of \bar{g} and \bar{D} matrices was optimized on the 10 K data using the principal values obtained from magnetization and susceptibility data.

In the end, the fitting procedure indicated that b is the overall intermediate axis (i.e. y in eq 1 with $E > 0$), while the a^*c plane contains the easy (z) and hard (x) molecular magnetic axes, with the latter at ca. 10° from a . In Figure 1 the orientation of the anisotropy axes is superimposed to the molecular structure, where the principal axes are not pointing towards the ligand atoms but between them. The good agreement between simulated and experimental torque data confirms the spin Hamiltonian parameters extracted from static magnetic measurements, in particular the negative sign of D with a relatively large rhombicity. On the other hand, Figure 3 shows that the agreement is less satisfactory at 100 K. The inclusion of rhombicity in the \bar{g} matrix (see EPR section below),

i.e. $g_x \neq g_y$, does not improve the quality of the fit, as indeed expected because torque measurements are mainly sensitive to the $\bar{\mathbf{D}}$ tensor. Keeping in mind that the orbital angular momentum of the cobalt(II) ion may be not completely quenched, on increasing temperature the magnetic properties are affected by spin-orbit coupled states at higher energies and the description of the metal ion as a simple $S = 3/2$ spin may be no longer valid. This would result in a failure of the spin Hamiltonian approach leaving *ab initio* methods and explicit inclusion of orbital contribution^[34] as the only way to appropriately describe the system (see below).

EPR spectroscopy

Since it can easily provide detailed information on the $\bar{\mathbf{g}}$ matrix, EPR spectroscopy can be regarded as a complementary technique to CTM. Low-temperature continuous-wave X-band EPR spectra were recorded on both powder and single crystal samples. The experimental spectrum of a powder sample of **3**, shown in Figure 4, can be easily interpreted as arising from an effective spin doublet with strongly anisotropic g factor and hyperfine coupling with the nuclear spin $I = 7/2$ of ^{59}Co (natural abundance = 100%). Similar spectra were obtained for **1**, although dominant dipolar broadening precludes resolution of the hyperfine structure (SI, Figure S4). In this frame, it is evident that the anisotropy of the ground doublet is of easy axis type, with “parallel” transition at low fields, and a non-negligible rhombicity evidenced by the separation of the perpendicular transitions in the 400–500 mT region. The experimental spectrum was simulated using the following spin Hamiltonian:^[38]

$$\mathcal{H}' = \mathcal{H}'_{\text{Zeeman}} + \mathcal{H}'_{\text{hyperfine}} = \mu_B \hat{\mathbf{B}} \cdot \bar{\mathbf{g}}' \cdot \hat{\mathbf{S}}' + \hat{\mathbf{S}}' \cdot \bar{\mathbf{A}}' \cdot \hat{\mathbf{I}}' \quad (4)$$

where the primed parameters are associated with the effective spin $S' = 1/2$. The best simulation was obtained by assuming an axial hyperfine coupling tensor ($\mathcal{A}'_{1,2} = 2 \pm 1 \cdot 10^{-3}$ and $\mathcal{A}'_3 = 3.20 \pm 0.05 \cdot 10^{-2} \text{ cm}^{-1}$) and a moderately rhombic effective $\bar{\mathbf{g}}'$ matrix ($g'_1 = 1.37 \pm 0.02$, $g'_2 = 1.62 \pm 0.02$, and $g'_3 = 7.930 \pm 0.002$).

The principal values of the effective $\bar{\mathbf{g}}'$ matrix can be related to the spin Hamiltonian parameters of an $S = 3/2$ system through the following equations, valid only for easy axis systems, *i.e.* for $D < 0$:^[39]

$$g'_1 = g_x \left(1 - \frac{1-3\eta}{\sqrt{1+3\eta^2}} \right), \quad g'_2 = g_y \left(1 - \frac{1+3\eta}{\sqrt{1+3\eta^2}} \right), \quad g'_3 = g_z \left(\frac{2}{\sqrt{1+3\eta^2}} + 1 \right) \quad (5)$$

where $\eta = E/D$. For purely axial anisotropy ($\eta = 0$) eq 5 affords $g'_1 = g'_2 = 0$ and $g'_3 = 3g_z$. The non-zero experimental values of g'_1 and g'_2 are then consistent with a significant rhombic anisotropy. It is evident that, depending on the value of η , different solutions for g_x , g_y and g_z can be found, since the problem is over-parametrized. However, sets of solutions in reasonable accordance

with those extrapolated from magnetic measurements and confirmed by CTM can be found by assuming the rhombicity to lie in the range $-0.28 < \eta < -0.22$. In particular, if one fixes $\eta = -0.273$, as obtained from the magnetic analysis, application of eq 5 provides the following estimates of the principal g values for the $S = 3/2$ system: $g_x = 2.125$, $g_y = 1.935$ and $g_z = 2.825$ (alternatively, assuming $\eta = 0.273$, $g_x = 1.935$, $g_y = 2.125$ and $g_z = 2.825$). We note that no reproduction of the effective g' pattern can be obtained for $\eta = 0.273$ and $g_x > g_y$ or for $\eta = -0.273$ and $g_y > g_x$. The outcome of this analysis is that also the \bar{g} matrix has a non-negligible rhombicity, which can be clearly detected only by EPR, whereas the magnetic characterization techniques are mainly sensitive to \bar{D} anisotropy. One should also notice that the only way to reconcile the description of Hamiltonian in eq 4 with that of Hamiltonian in eq 1 is assuming that \bar{g} and \bar{D} for the $S = 3/2$ state have opposite rhombicities. This means that the smallest \bar{g} principal value is collinear with the intermediate direction of the \bar{D} tensor, while the intermediate \bar{g} component is along the hard direction of the \bar{D} tensor. The resulting partial compensation of the two rhombicities is consistent with the relatively small separation of the perpendicular transitions at high fields. Such behaviour is at first sight counter-intuitive because both anisotropies arise from the same phenomenon, i.e. spin-orbit coupling (SOC).

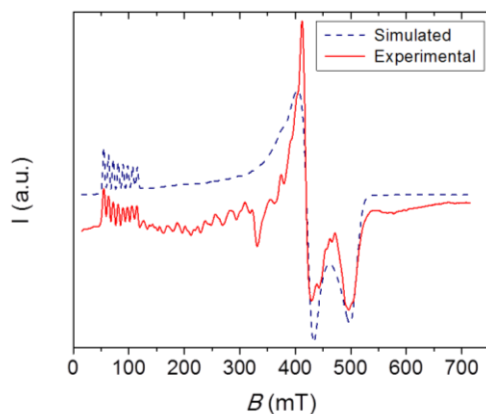


Figure 4. X-band ($\nu = 9.43$ GHz) continuous wave EPR spectrum recorded at 5 K on a powder sample of **3** (solid line), along with its simulation (dash line).

Finally we measured the angular dependence of the X-band EPR spectrum of a single crystal of **1** at low temperature to assign the easy, hard and intermediate directions of the magnetic anisotropy of the ground doublet. The sample was set to rotate around b axis from $\theta = 0^\circ$, where \mathbf{B} is parallel to c , to $\theta = 90^\circ$, where \mathbf{B} is parallel to $-a^*$, as in Rot1 of CTM measurements. The angular dependence of the spectra is shown in Figure 5: the maximum resonant field ($B_{\text{res}} = 491.5$ mT, corresponding to

$g_{\text{eff}} = 1.37$) is observed at $\theta = 40^\circ$, while the minimum one is found at $\theta = 130^\circ$ ($B_{\text{res}} = 85.7$ mT, $g_{\text{eff}} = 7.93$). Within experimental error ($\pm 5^\circ$), these two extremes occur along the principal directions of the $\bar{\mathbf{D}}$ tensor, as determined by CTM (Figure 3). Furthermore, since $g_{\text{eff}} = 1.37$ and 7.93 are also the highest and the lowest resonance fields observed in the powder spectrum (Figure 4), both the easy and the hard direction of g_{eff} lie in the a^*c plane, confirming our assignment of b as the intermediate anisotropy axis, though it corresponds to the smallest component of the true $\bar{\mathbf{g}}$ tensor.

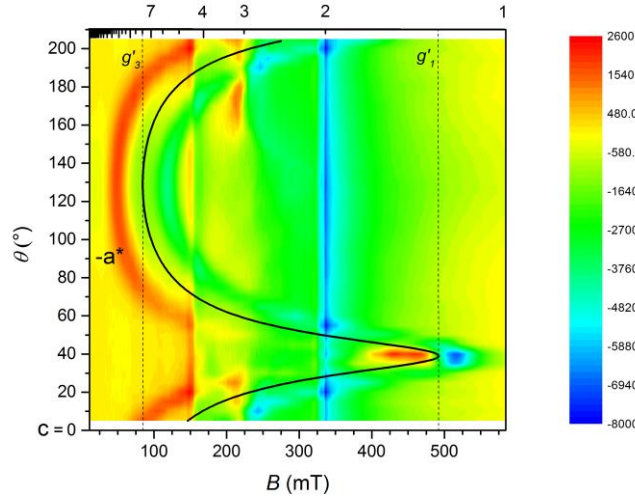


Figure 5. Angular dependence of the resonant fields of X-band ($\nu = 9.41$ GHz) EPR of a single crystal of **1** at 5 K. The black solid line is the simulated angular dependence of the resonance field using the parameters reported in the text. The blue line marks a $g = 2.00$ reference impurity in the cavity.

An alternative description of the electronic structure of distorted octahedral cobalt(II), which explicitly takes into account the unquenched orbital angular momentum, should involve the use of Griffith Hamiltonian:^[40]

$$\mathcal{H} = -\frac{3}{2}\kappa\lambda\hat{L} \cdot \hat{S} + \Delta_{ax} \left[\hat{L}_z^2 - \frac{1}{3}L(L+1) \right] + \Delta_{rh} [\hat{L}_x^2 - \hat{L}_y^2] + \mu_B \mathbf{B} \cdot \left(g_e \hat{S} - \frac{3}{2}\kappa\hat{L} \right) \quad (6)$$

which operates on the ground $^4T_{1g}$ term of the octahedral cobalt(II) ion. Using the T-P isomorphism,^[41] the orbital ($L = 1$) and spin ($S = 3/2$) angular momenta interact via SOC as described by the first term of the Hamiltonian. Here, the orbital reduction parameter (κ) takes into account both the covalence effect and the mixing of $^4T_{1g}(^4F)$ with $^4T_{1g}(^4P)$ due to the crystal field, while $-3/2$ is a constant required by T-P isomorphism. The second term of this Hamiltonian represents the effect of an axial crystal field, resulting in a splitting of the orbital triplet $^4T_{1g}$ in octahedral symmetry into $^4A_{2g}$ ($M_L = 0$) and 4E_g ($M_L = \pm 1$) in tetragonal symmetry. The third term models the rhombic component of the crystal field, which removes the degeneracy of the orbital doublet as symmetry is further lowered to C_2 . In this framework, a negative (positive) value of Δ_{ax}

results in a ground 4E_g (${}^4A_{2g}$) term and corresponds to easy axis (easy plane) anisotropy.^[34,42,43] Finally, the fourth term is the Zeeman interaction, which comprises both spin and orbital contributions. Despite the large number of parameters, by fixing the SOC constant to the free ion value ($\lambda = -180 \text{ cm}^{-1}$) the pattern of effective g factors for the ground doublet could be reproduced^[38] using parameters lying in a relatively narrow range: $-2100 < \Delta_{ax} < -1650 \text{ cm}^{-1}$, $100 < \Delta_{rh} < 160 \text{ cm}^{-1}$, $0.8 < \kappa < 1.0$ (SI, Figure S5). The best agreement was found for $\Delta_{ax} = -1850 \text{ cm}^{-1}$, $\Delta_{rh} = 130 \text{ cm}^{-1}$, $\kappa = 0.9$, $\lambda = -180 \text{ cm}^{-1}$, corresponding to $|\Delta_{rh}/\Delta_{ax}| \sim 0.07$. With this set of parameters, the first- and second-excited spin-orbit doublets lie at 210 and 558 cm^{-1} from the ground state (SI, Table S6), respectively. This suggests that a spin Hamiltonian formalism might not be entirely appropriate for modelling the high temperature magnetic behaviour, since the thermal energy is not completely negligible compared to that of second excited doublet. Interestingly, when the same set of parameters is used to simulate the powder dc magnetic data, the resulting curves are in reasonable agreement with the experimental ones (SI, Figure S6). On the other hand, if applied to torque measurements, the same set well reproduces the experimental data of Rot1 but it is less accurate in simulating those of Rot2 at 100 K (SI, Figure S7). This indicates that even this approach is too rough to capture the finest details of the electronic structure of this system and in particular those related to its rhombicity.

Dynamic magnetic properties

ac magnetic susceptibility measurements of **1** in zero static field did not display any temperature-dependent peak in the out-of-phase susceptibility, χ_M'' . Given the Kramers nature of cobalt(II), this suggests that quantum tunnelling (QT) relaxation paths, induced by hyperfine interactions or intermolecular dipolar couplings, accelerate the relaxation of the magnetization beyond detection. In a 1 kOe static field, however, clear maxima appeared in the 10–10000 Hz frequency range, whose position is temperature dependent (Figure 6a and SI, Figure S8). Treatment of the isothermal $\chi_M''(\nu)$ plots with a generalized Debye model allowed to extract the relaxation time τ at each temperature.^[44] These data are shown in Figure 6b as a $\ln \tau(1/T)$ plot. In the highest temperature interval, between 6.5 and 9.5 K, the plot appears to be linear and can be suitably fitted with the Arrhenius law ($\ln \tau = \ln \tau_0 + U_{\text{eff}}/k_B T$) with best-fit parameters $U_{\text{eff}}/k_B = 43.6(2) \text{ K}$ and $\tau_0 = 1.2(2) \times 10^{-7} \text{ s}$ (green line in Figure 6b). These values are in the range previously observed for field-induced pseudo-octahedral mononuclear SMMs.^[45] The observed activation barrier to magnetic relaxation, however, is well below the splitting between the two lowest-lying doublets (127 cm^{-1} or 165 K, as estimated from dc magnetic data of **1**, see above). On lowering temperature, a distinct curvature

appears in the plot; this deviation from the simple Arrhenius-like behaviour can be due to processes induced by either non-resonant spin-phonon interactions in the solid state^[46,47] or to tunnel mechanisms. In order to reproduce the temperature dependence of the magnetic relaxation in the whole temperature range, QT (A), direct (BT) and Raman (CT^n) relaxation processes were taken into account in addition to the previous Orbach term, as outlined by the following equation:

$$\tau^{-1} = \tau_0^{-1} \exp(-U_{\text{eff}}/k_B T) + A + BT + CT^n \quad (7)$$

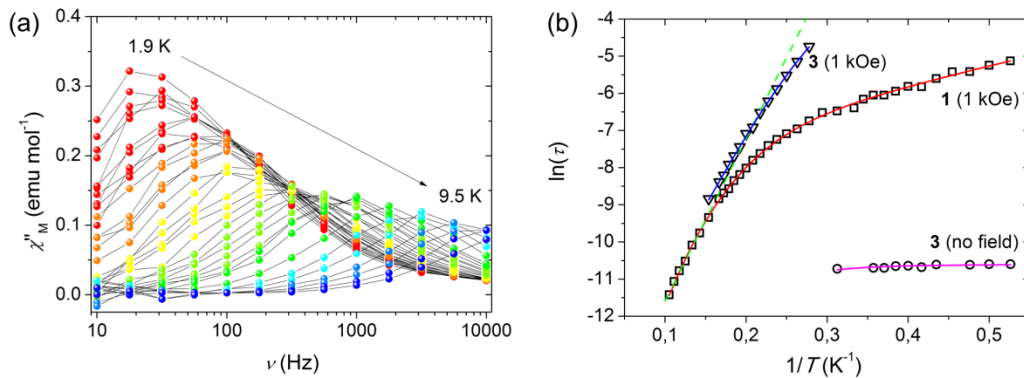


Figure 6. (a) Imaginary component of the ac susceptibility, χ''_M , of **1** measured at 1 kOe applied static field in the 10–10000 Hz frequency range and in the 1.9(red)–9.5(blue) K temperature range; grey lines are a guide for the eye. (b) Arrhenius plot for **1** at 1 kOe applied static field (\square) and **3** in zero (∇) and 1 kOe static field (\circ). The green dotted line is given by the fitting of **1** in the 6.5–9.5 K range with an Orbach model, while the red curve employs a mixed Raman and direct one. The magenta and blue lines display the data fitting of **3** at zero and 1-kOe applied field, respectively; see text for further discussion.

To avoid over-parametrization, the number of fitting terms was kept as low as possible, and the quantum tunnelling mechanism was considered to be quenched by the applied static field ($A = 0$). Upon either including an Orbach mechanism (with fixed τ_0 and U_{eff} values taken from the high- T region) or excluding the Raman term, the model was unable to account for the low temperature relaxation behaviour. A Raman process along with direct phonon-induced relaxation at low temperature ($C = 0.45(8) \text{ s}^{-1} \text{ K}^{-n}$, $n = 5.38(9)$, $B = 103(4) \text{ K}^{-1}$) gave the best results (red line in Figure 6b). The exponent n of the Raman relaxation process for a Kramer ion should be equal to 9,^[20,48] but several factors can contribute to lower its value,^[20] like the presence of low-lying excited electronic states, for which $n = 5$ is postulated.^[48]

Dipolar interactions in the solid state can provide efficient relaxation pathways in zero field and hinder the detection of magnetization dynamics. Dilution of SMMs in a diamagnetic matrix has been identified as a key tool for reducing such long-range couplings, affording a measurable zero-

field relaxation.^[14,49] The effect of dipolar interactions in **1** was then ascertained by analysing the ac response of diamagnetically-diluted sample **3**. Unlike in **1**, a set of clearly visible peaks appeared in the $\chi_M''(\nu)$ isothermal profiles of **3** in zero field in the 1.9–3.2 K range (SI, Figure S9). The maxima of these peaks display a feeble dependence on temperature that could be reproduced considering a Raman relaxation process with QT. A best-fit procedure, whose results are reported as a magenta line in Figure 6b, yielded $C = 11(2) \text{ s}^{-1} \text{ K}^{-n}$ and a tunnelling frequency $A = 40.3(5) \text{ kHz}$, being the n exponent kept fixed at 5.38 as in **1** to avoid over-parametrization. The onset of a QT relaxation path in zero field is in line with the rhombicity of **1** and the unavoidable presence of hyperfine interactions which are necessary to promote tunnelling. These data show, to the best of our knowledge for the first time, that slow magnetic relaxation in zero field can be achieved in a pseudo-octahedral cobalt(II) complex, provided that intermolecular magnetic interactions are efficiently reduced. A similar effect given by magnetic dilution on easy axis systems was previously reported only in a tetrahedral cobalt(II) compound.^[14] It is also most probably the reason for zero-field slow magnetic relaxation in the $\text{Co}^{\text{II}}\text{Co}^{\text{III}}_3$ SMM family, where the magnetically-active trigonal prismatic cobalt(II) centre is partly isolated from the neighbours by three diamagnetic cobalt(III) ions.^[18]

Upon application of a 1 kOe static field, the magnetization dynamics of **3** slows down significantly due to the further suppression of relaxation pathways induced by QT (SI, Figure S10). The high-temperature data almost overlap with those detected for **1** suggesting that, when a linear regime in the Arrhenius plot is observed, the effective barrier is much smaller than the separation between the ground and the first excited doublets. This indicates the key role played by non-resonant phonons in the high-temperature range.^[46] The data can be phenomenologically reproduced including Raman and direct relaxation processes (blue line in Figure 6b), as for data fitting of **1**. The best-fit parameters ($C = 0.015(5) \text{ s}^{-1} \text{ K}^{-n}$, $n = 7.0(2)$, $B = 0(4) \text{ K}^{-1}$) indicate that application of a 1 kOe dc field suppresses the direct pathway and leads to an increased n exponent, in line with a previous report on trigonal antiprismatic cobalt(II) complexes with uniaxial anisotropy.^[50]

Theoretical calculations

To better understand and quantify the overall electronic structure of cobalt(II) and its magnetic anisotropy in **1**, *ab initio* multi-reference calculations using the ORCA^[51] and MOLCAS^[52] software packages were carried out. ORCA produces two sets of results: CASSCF and NEVPT2, both including spin-orbit contributions introduced by the quasi-degenerate perturbation theory (QDPT). MOLCAS has been employed to provide CASSCF and CASPT2 results, including spin-

orbit effects that have been introduced with SO-RASSI method. SI, Tables S7–S10 report the spin-free (δ_E) and spin-orbit (Δ_E) state energies, while Table 2 and SI, Tables S11 and S12 gather the second-order anisotropy parameters, as described by eq 1, the diagonalized \bar{g} and \bar{D} matrices and the energy difference between the ground and the first excited Kramers’ doublet (KD) computed with different methods and the standard basis set starting from the structures of **1** at 161 and 295 K, respectively.

Before including the spin-orbit effects, all calculations agree to indicate an orbitally non-degenerate $S = 3/2$ ground state. In fact, the crystal field term of the octahedrally-coordinated high-spin cobalt(II) ion ($^4T_{1g}$) splits into three orbital singlets as the symmetry is lowered to C_2 . Each of these orbitally nondegenerate spin quartets is further split by SOC into two KDs (SI, Figure S11). In all calculations, the ground and first excited KDs are energetically well separated from the higher KDs, suggesting that a spin Hamiltonian formalism may be justified in the present case. Moreover, the energy of the first, second and third excited doublets (*ca.* 191, 648 and 929 cm^{-1} by MOLCAS/CASPT2) resemble those found when the Griffith Hamiltonian is employed to reproduce EPR data (210, 558, 820 cm^{-1} , see SI, Table S6).

Table 2. D , E , diagonalized \bar{g} and \bar{D} matrices and energy difference between the ground and the first excited KD (Δ_E) computed using the structure of **1** at 161 K with different methods and the standard basis set (see SI). For comparison, the first excitation energy (δ_E) in the absence of SOC for the ground state is also reported.

Method	g_{xx}, g_{yy}, g_{zz}	D_{xx}, D_{yy}, D_{zz} (cm^{-1})	D (cm^{-1})	E (cm^{-1})	Δ_E (cm^{-1})	δ_E (cm^{-1})
ORCA/CASSCF	2.156, 1.945, 3.069	48.68, 13.42, -62.11	-93.2	17.6	196.1	364.7
ORCA/NEVPT2	2.152, 1.960, 3.004	43.11, 14.34, -57.45	-86.2	14.4	179.4	467.4
MOLCAS/CASSCF	2.169, 1.963, 3.070	49.22, 12.19, -61.41	-92.1	18.5	195.1	368.9
MOLCAS/CASPT2	2.149, 1.960, 3.058	47.30, 13.25, -60.55	-90.8	17.0	191.0	364.6

Calculations also produce similar and negative values of D , with a negligible effect due to the use of the molecular structures at the two different temperatures (161 and 295 K, see SI, Table S11). However, a small but systematic trend can be observed, consisting in a decrease of $|D|$ and an increase of E upon cooling. The calculated $|D|$ values are *ca.* 50% larger than obtained from the fitting of the experimental magnetic data, while the computed values of E are in good agreement with experiments. In order to evaluate the possible effect of the employed basis set on the computed zfs parameters, calculations were repeated with a larger basis set using the structure of **1** at 161 K (SI, Experimental Section and Table S12). The new results show just small differences as compared

with the standard basis, with slightly larger $|D|$ and E . The effect is more pronounced in MOLCAS than in ORCA.

The sign and value of D can be rationalized using the spin-orbit operator, which is responsible for the coupling between ground and excited states.^[10,53] When the excitation occurs between orbitals with the same $|m_l|$ values, the $M_S = \pm 3/2$ components become more stable, and thus a negative contribution to D is obtained. On the other hand, an excitation between orbitals involving a $|\Delta m_l| = 1$ change, which produce stabilized $M_S = \pm 1/2$ components, leads to a positive contribution to the D value. Compound **1** displays a distorted octahedral core that produces the d -orbital splitting shown in Figure 7, which is obtained from the ORCA/NEVPT2 calculation by using the *ab initio* ligand field theory (AILF) method.^[10,54] As may be observed, the degeneracy of the t_{2g} and e_g orbitals of the regular octahedron is lifted. This indicates that the initial octahedral $^4T_{1g}$ ground state should evolve into a lower-energy term in C_2 symmetry (SI, Figure S11). The AILF method allows the identification of the d orbitals in **1** after the splitting and the lowest-energy doubly-occupied orbital is d_{xy} , which is followed by the doubly occupied orbital d_{xz} (or d_{yz} , because these cannot be distinguished), while the first semi-occupied orbital is d_{yz} (or d_{xz}) (where z here represents the *trans*-N{pyridyl}-Co-{pyridyl} direction, and x and y the directions of the pyrazolyl nitrogen atoms). The other two orbitals ($d_{x^2-y^2}$ and d_{z^2}) are found at higher energies. Thus, the first excitation should occur within the d_{xz}/d_{yz} orbital pair and, since these orbitals have the same $|m_l|$ value, the larger contribution to D should be negative, and consequently the magnetic easy axis nature of **1** is confirmed. This is in opposition with the most common orbital splitting for pseudo-octahedral cobalt(II) complexes, which usually brings the d_{xy} orbital as first semi-occupied orbital,^[31] resulting then in easy plane systems.

The calculated \bar{D} matrix has its intermediate axis (i.e. y in eq 1 with $E > 0$) along the twofold crystallographic axis b and the hard (x) and easy (z) directions in the a^*c plane (SI, Figure S12), with the hard axis x at 10° from a . Thus, the computed orientation of the zfs tensor compares very well with that obtained by both single-crystal CTM and EPR spectroscopy. Turning now to g factors, their *ab initio* values (Table 2) are remarkably close to those obtained using eq 5. For instance the g factors resulting from ORCA/NEVPT2 calculations are $g_x = 2.152$, $g_y = 1.960$ and $g_z = 3.004$ to be compared with those estimated by analysis of the EPR spectra, $g_x = 2.125$, $g_y = 1.935$ and $g_z = 2.825$. The effective \bar{g}' factors of the ground KD found in this same calculation, which can be directly compared to those obtained from EPR experiments, are slightly more anisotropic: $g'_1 = 0.999$, $g'_2 = 1.172$ and $g'_3 = 8.566$ (from EPR: $g'_1 = 1.37 \pm 0.02$, $g'_2 = 1.62 \pm 0.02$, and $g'_3 = 7.930 \pm 0.002$). The results obtained in the MOLCAS/CASPT2 calculation are also quite similar to both

these sets of values: $g'_1 = 1.108$, $g'_2 = 1.287$ and $g'_3 = 8.570$. The principal directions of the \bar{g} matrix are collinear with those of \bar{D} , the largest g factor being along z . However, as shown in SI, Figure S12, the smallest g value is computed along the twofold crystallographic axis (y) rather than along the hard direction of \bar{D} (x). As observed in single-crystal EPR experiments, the theoretical calculations indicate an inversion of the intermediate and hard axes between \bar{g} and \bar{D} , i.e. opposite rhombicities.

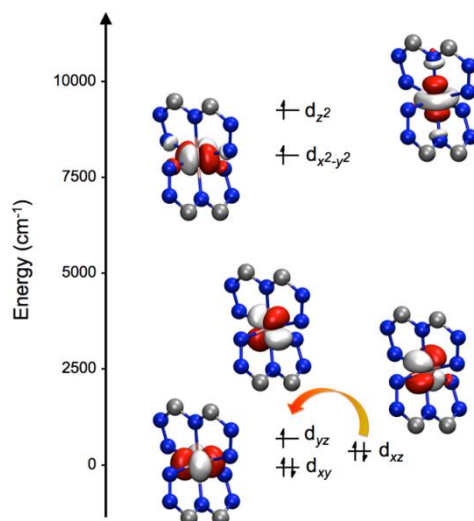


Figure 7. AILF computed d -orbital splitting for **1** (orbital energies: d_{xy} 0.0 cm^{-1} , d_{xz} 444.6 cm^{-1} , d_{yz} 697.9 cm^{-1} , $d_{x^2-y^2}$ 8121.0 cm^{-1} , d_{z^2} 9257.3 cm^{-1}). Colour code: Co = pink, C = grey, N = blue; H atoms and part of the ligands omitted for clarity. The orange arrow indicates the lowest-energy transition.

The CTM experiments can be nicely reproduced in the MOLCAS calculations by re-computing the magnetization of the system along the different axes upon application of a rotating magnetic field (see Experimental Section in ESI for further details). The agreement between experimental and theoretical torque values for Rot1 in the 10–100 K temperature range is very satisfactory, while some discrepancy is observed in Rot2 when increasing the temperature (SI, Figures S13 and S14), as observed by applying Griffith model. By the way, it is important to underline that Rot2, probing mainly the intermediate and hard magnetic directions, is mostly affected by the inversion of the corresponding components of \bar{g} and \bar{D} matrices, a phenomenon whose origin remains unclear.

The spin relaxation pathways, which can be related to the dynamic magnetic properties of transition metal complexes, can be obtained directly through the SINGLE_ANISO^[55] code as implemented in MOLCAS. The computed relative energies of the lowest-lying KDs and the magnetization reversal pathway for **1**, as obtained with the MOLCAS/CASPT2 calculation, are

shown in Figure 8. The calculations indicate a plausible relaxation pathway *via* a direct QT in the ground state; the matrix element of the transition within the 1- / 1+ ground doublet takes a value of 0.40, higher than the required threshold of 0.1 for an efficient relaxation mechanism.^[10] In addition, an Orbach process could be also plausible; for instance, the first excited KD is quite low in energy (191.0 cm⁻¹) and the matrix elements related to the vertical and diagonal (Orbach) excitations are high enough (1.09 and 1.11, respectively) to allow the spin relaxation through these pathways. However, the experimental effective barrier for the thermally activated process is significantly smaller than the calculated one. Obviously, the horizontal transition between the 2- and 2+ states is allowed, enabling the possibility of QT between them. The second and third excited KDs are much higher in energy and therefore they are not expected to participate in the relaxation mechanism. These results, combined with the ac magnetic susceptibility measurements of **1**, confirm the spin-phonon Raman relaxation mechanism.

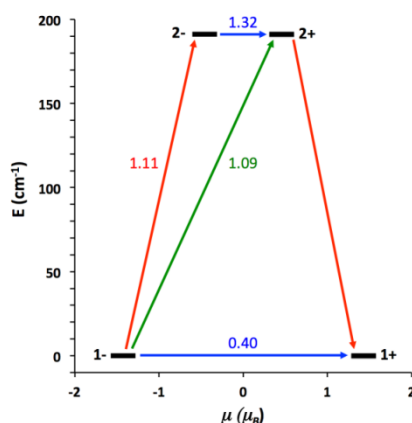


Figure 8. Lowest two KDs and *ab initio* MOLCAS/CASPT2 computed relaxation mechanism for **1**. The thick black lines imply KDs as a function of their magnetic moment along the main anisotropy axis. Red lines indicate the magnetization reversal mechanism. The blue lines correspond to ground state QT and thermally-assisted QT via the first excited KD, and green lines show possible Orbach relaxation processes. The values close to the arrows indicate the matrix elements of the transition magnetic moments (above 0.1 an efficient spin relaxation mechanism is expected).

If **1** were to follow the Orbach mechanism depicted by the calculations, the U_{eff} value would take a higher value than the one obtained in the fitting (43.6(2) K) and this is clearly not the case, indicating that the spin relaxation pathway does not proceed by overcoming a thermal activation barrier. This behaviour is quite general for systems with large D and an explanation can be based on the fact that the Arrhenius law is only expected if the Debye description of the phonons is correct.

Much smaller energy barrier and a deviation from Arrhenius at low temperature can be ascribed to the anharmonicity of the vibrational modes (a source of Raman-like relaxation).

Conclusions

We have here presented a detailed magnetic and spectroscopic characterization, both on powder and single crystals, of a pseudo-octahedral cobalt(II) complex, which we used as a test bench for different theoretical approaches, ranging from *ab initio* calculations to Ligand Field model and spin Hamiltonian. We can conclude that the main features of the magnetic behaviour of this system are accounted for by using an $S = 3/2$ spin Hamiltonian approach. However, by exploiting the variety of the experimental and theoretical techniques employed, we were also able to evidence the hitherto unreported inversion of the hard and intermediate axes of \bar{g} and \bar{D} matrices, a phenomenon which is nicely reproduced by *ab initio* calculations.

The present results further highlight the possibility to obtain slow magnetic relaxation in a mononuclear pseudo-octahedral HS cobalt(II) complex with dominant easy axis anisotropy in the absence of an applied dc field. In particular, the distortion of the coordination environment from perfect octahedral as created by the two *mer* bpp-COOMe ligands in **1** induces the easy axis nature of the compound.

The observed magnetization dynamics is strongly affected by tunnel processes due to the rhombic magnetic anisotropy in zero field. In applied magnetic field, the behaviour deviated from the Arrhenius behaviour expected for an ideal SMM as the highest estimation of the energy barrier to be overcome is significantly smaller than the separation between the ground and first doublets. This behaviour has been observed in other highly anisotropic complexes^[56] and attributed to the efficiency of non-resonant optical phonons,^[46,47] which cannot be neglected if anharmonicity is taken into account. Even if the optimization of the magnetization dynamics of SMM requires therefore the control of several factors like tunnelling mechanisms and vibrational modes, the first step remains the rational design of the coordination environment able to generate magnetic bistability through the appropriate sign of the magnetic anisotropy. Octahedral cobalt(II), despite its large orbital contribution, has so far eluded to provide successful examples of genuine SMM behaviour. The combination of a detailed experimental investigation with theoretical modelling at different levels here described can provide valuable hints for a rational design of cobalt(II)-based SMM.

Experimental Section

General Information

Co(ClO₄)₂·6H₂O and Zn(ClO₄)₂·6H₂O were of reagent grade and used as received. Diethyl ether was pre-dried over CaCl₂ overnight and distilled from sodium/benzophenone under N₂ before use, CH₃CN was treated with CaH₂ and then distilled under N₂. Elemental analyses were recorded using a Carlo Erba EA1110 CHNS-O automatic analyser. Zn and Co contents in **3** were evaluated with a Perkin Elmer Optima 5300 DV Inductively Coupled Plasma Optical Emission Spectrometer (ICP-OES) equipped with a concentric nebulizer. ¹H NMR spectra were recorded on a 400 MHz Bruker FT-NMR Advance400 spectrometer at room temperature. Proton chemical shifts are given in parts per million (ppm) versus external TMS, and were determined by reference to the solvent residual signals (1.94 ppm for CHD₂CN); coupling constants are given in Hz. IR spectra were recorded as KBr discs using a Jasco FTIR-4700LE spectrophotometer with a 2 cm⁻¹ resolution. Ligand bpp-COOMe was synthesized as previously reported.^[26] Further details on the X-ray structure determination of **1** and **2** (SI, Table S1), dc and ac magnetic measurements, CTM and EPR experiments and theoretical calculations can be found in SI.

Synthesis of bis(methyl 2,6-di(pyrazol-1-yl)pyridine-4-carboxylate)cobalt(II) bis(perchlorate), [Co(bpp-COOCH₃)₂](ClO₄)₂ (**1**)

Ligand bpp-COOMe (36.0 mg, 0.134 mmol) was suspended in acetonitrile (3 mL). Addition of Co(ClO₄)₂·6H₂O (24.3 mg, 0.0664 mmol) caused dissolution of the ligand upon formation of the desired complex with an immediate colour change to purple-red. After 30 minutes stirring, the filtered solution was put in vapour diffusion with diethyl ether (5 mL). X-ray quality light red crystals appeared within a day, and were collected after one week at complete diffusion (39.4 mg, 74.5%). Elemental analysis (%) found: C, 39.90; H, 2.88; N, 17.99. Calcd for C₂₆H₂₂Cl₂CoN₁₀O₁₂ (796.35): C, 39.21; H, 2.78; N, 17.59. IR (KBr): $\nu_{\max}/\text{cm}^{-1}$ 1738s ($\nu\text{C=O}$ ester), 1632s, 1580s, 1523s and 1405s ($\nu\text{C=C}$, $\nu\text{C=N}$), 1460s and 1444m ($\delta\text{C-H}$), 1095s (νClO_4^-) and 1056s ($\nu\text{C-O}$).

Synthesis of bis(methyl 2,6-di(pyrazol-1-yl)pyridine-4-carboxylate)zinc(II) bis(perchlorate), [Zn(bpp-COOCH₃)₂](ClO₄)₂ (**2**)

The above-described procedure, starting from bpp-COOMe (39.0 mg, 0.145 mmol) and Zn(ClO₄)₂·6H₂O (27.6 mg, 0.0741 mmol), afforded colourless crystals of the title compound suitable for X-ray diffraction (40.0 mg, 68.8%). Elemental analysis (%) found: C, 39.05; H, 3.04; N,

17.48. Calcd for $C_{26}H_{22}Cl_2N_{10}O_{12}Zn$ (802.83): C, 38.90; H, 2.76; N, 17.45. IR (KBr): ν_{max}/cm^{-1} 1738s ($\nu C=O$ ester), 1632s, 1584s, 1522s and 1405s ($\nu C=C$, $\nu C=N$), 1458s and 1444m ($\delta C-H$), 1095s (νClO_4^-) and 1058s ($\nu C-O$). 1H NMR (400 MHz, CD_3CN , 293 K): δ = 8.73 (d, J_1 = 2.8 Hz, 2H, H^5 -pz), 8.45 (s, 2H, $H^{3/5}$ -py), 7.62 (d, J_2 = 1.6 Hz, 2H, H^3 -pz), 6.70 (dd, J_1 = 2.8 Hz, J_2 = 1.6 Hz, 2H, H^4 -pz), 4.11 (s, 3H, OCH_3) ppm.

Synthesis of $[Zn_{0.95}Co_{0.05}(bpp-COOCH_3)_2](ClO_4)_2$ (3)

The above-described procedure, starting from $bpp-COOMe$ (81.2 mg, 0.302 mmol), $Co(ClO_4)_2 \cdot 6H_2O$ (2.9 mg, 0.0079 mmol) and $Zn(ClO_4)_2 \cdot 6H_2O$ (53.9 mg, 0.145 mmol) using 6 mL of acetonitrile and 10 mL of diethyl ether, afforded very light orange crystals of the title compound suitable for X-ray diffraction (91.7 mg, 75.8%). Elemental analysis (%) found: C, 39.22; H, 2.83; N, 17.46; Co, 0.40; Zn, 8.04. Calcd for $C_{26}H_{22}Cl_2N_{10}O_{12}Zn_{0.95}Co_{0.05}$ (802.51): C, 38.91; H, 2.76; N, 17.45; Co, 0.37; Zn, 7.74. IR (KBr): ν_{max}/cm^{-1} 1737s ($\nu C=O$ ester), 1632s, 1584s, 1522s and 1405s ($\nu C=C$, $\nu C=N$), 1458s and 1444m ($\delta C-H$), 1095s (νClO_4^-) and 1058s ($\nu C-O$).

Acknowledgments

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Conflict of Interest

The authors declare no conflict of interest.

Contributor Roles

L.R. conceived and designed the experiments, L.R. and N.B. performed the synthesis and crystallization, G.P. performed dc and ac magnetic measurements, L.T. performed CTM and dc magnetic measurements and helped L.S. in single crystal EPR experiments, L.S. performed EPR experiments and simulations and wrote the code for analysis of CTM using Griffith Hamiltonian, D.P. performed the single-crystal X-ray diffraction studies, J.J. and E.R. performed theoretical calculations, all authors analyzed the data, R.S. and A.C. contributed reagents/materials/analysis tools; L.R. wrote the paper with the help of all co-authors.

Keywords: cobalt(II), single-molecule magnets, torque magnetometry, electron paramagnetic resonance spectroscopy, *ab initio* calculations

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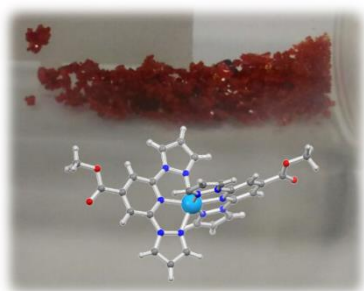
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Graphical abstract



Distorted relaxation: The homoleptic high-spin cobalt(II) compound $[\text{Co}(\text{bpp-COOMe})_2](\text{ClO}_4)_2$ with two *mer* tridentate nitrogen-rich ligands possesses the right distortion from octahedral coordination geometry to show dominant easy axis anisotropy and slow relaxation of the magnetization in the absence of applied field once magnetically diluted in its zinc analogue.

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Author/s:

Rigamonti, L; Bridonneau, N; Poneti, G; Tesi, L; Sorace, L; Pinkowicz, D; Jover, J; Ruiz, E; Sessoli, R; Cornia, A

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