

Cite this: *Dalton Trans.*, 2022, **51**, 18118Slow magnetic relaxation in Fe(II) *m*-terphenyl complexes†Andrew J. Valentine,^a Ana M. Geer,^b Toby J. Blundell,^c Will Tovey,^a Matthew J. Cliffe,^a E. Stephen Davies,^a Stephen P. Argent,^a William Lewis,^d Jonathan McMaster,^a Laurence J. Taylor,^{a*} Daniel Reta^e and Deborah L. Kays^{a*}

Two-coordinate transition metal complexes are exciting candidates for single-molecule magnets (SMMs) because their highly axial coordination environments lead to sizeable magnetic anisotropy. We report a series of five structurally related two-coordinate Fe(II) *m*-terphenyl complexes (4-R-2,6-Xyl₂C₆H₂)₂Fe [R = *t*Bu (**1**), SiMe₃ (**2**), H (**3**), Cl (**4**), CF₃ (**5**)] where, by changing the functionalisation of the *para*-substituent (R), we alter their magnetic function. All five complexes are field-induced single-molecule magnets, with relaxation rates that are well-described by a combination of direct and Raman mechanisms. By using more electron donating R groups we were able to slow the rate of magnetic relaxation. Our *ab initio* calculations predict a large crystal field splitting (>850 cm⁻¹) and sizeable zero-field splitting parameters (ca. -60 cm⁻¹, |E| < 0.2 cm⁻¹) for **1–5**. These favourable magnetic properties suggest that *m*-terphenyl ligands have untapped potential as chemically versatile ligands able to impose highly axial crystal fields.

Received 1st November 2022,
Accepted 7th November 2022

DOI: 10.1039/d2dt03531f

rsc.li/dalton

Introduction

Single-Molecule Magnets (SMMs) are molecules that display slow magnetic relaxation without long-range magnetic ordering.^{1,2} As such, SMMs have attracted significant interest due to their potential applications in spintronics, quantum computing, and nanotechnology.^{3,4} SMM behaviour was first observed in the polynuclear cluster [Mn^{IV}₄Mn^{III}₈O₁₂(O₂CCH₃)₁₆(H₂O)₄],⁵ and later the mononuclear [nBu₄N][Pc₂Tb^{III}] (Pc = phthalocyaninato) demonstrated that SMM behaviour could occur in species containing a single lanthanide centre^{6,7} due to their inherently strong spin-orbit coupling and anisotropy. This facilitated the rational design of

mononuclear SMMs, converging upon highly axial dysprosium-based systems.^{8–11} Building upon this, mixed-valence dilanthanide complexes with strong magnetic exchange have recently shown record coercive fields at liquid nitrogen temperatures.¹²

While the first mononuclear SMMs incorporated lanthanide ions, the 3d metals have also received significant attention.¹³ The challenge for mononuclear 3d SMMs is that the crystal fields exerted on the metal centre by the ligands tend to quench the orbital angular momentum. This can be avoided through careful ligand design, and there have been a number of reports of Fe(I),^{14,15} Fe(II),^{16–27} Co(II),^{28–34} Ni(II),³⁵ and Mn(III)³⁶ complexes that display slow magnetic relaxation. Among these, two-coordinate metal centres in linear or near-linear geometries, most notably [Co{C(SiMe₂O-1-naphthyl)₃}₂],³⁴ have shown the best performance in terms of magnetic blocking temperature and open hysteresis^{15,24,28,34} as their large magnetic anisotropy inhibits quenching of the orbital angular momentum. Thus, to realise the potential of 3d SMMs, one promising pathway is to explore ligands which impose strong axial fields and are sufficiently bulky to achieve linear coordination.

To that end, *m*-terphenyl ligands stand out as potential candidates. These are a class of bulky, monoanionic ligands that have proven useful in the stabilisation of highly reactive or unusual d-block and main group species.^{37–39} Several two-coordinate 3d species have been reported,^{40–46} and have found applications in catalysis^{44–47} and small molecule activation^{41,43,48} due to the high reactivity of the coordinatively unsaturated metal centre. In addition, the *para*-substituent on

^aSchool of Chemistry, University Park, University of Nottingham, Nottingham, NG7 2RD, UK. E-mail: laurence.taylor@nottingham.ac.uk, deborah.kays@nottingham.ac.uk

^bDepartamento de Química Inorgánica, Instituto de Síntesis Química y Catálisis Homogénea (ISQCH), CSIC Universidad de Zaragoza, Pedro Cerbuna 12, 50009 Zaragoza, Spain

^cDepartment of Chemistry, Durham University, South Road, Durham, DH1 3LE, UK

^dSchool of Chemistry, The University of Sydney, F11, Eastern Ave, Sydney, NSW 2006, Australia

^eKimika Fakultatea, Euskal Herriko Unibertsitatea UPV/EHU & Donostia International Physics Center (DIPC), Euskadi, IKERBASQUE, Basque Foundation for Science, Bilbao, Spain. E-mail: daniel.reta@ehu.eus

† Electronic supplementary information (ESI) available: Full experimental details for the synthesis, characterization, and analysis. CCDC 2184650–2184653. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d2dt03531f>



the central aryl ring can be functionalised, making them amenable to tuning of their electronic properties.^{49–52}

Whilst the related *m*-terphenoxide ligands have been investigated for Dy(III) SMMs,⁵³ *m*-terphenyl carbanions have not been studied for SMM synthesis, despite imposing a near-linear coordination environment. Here, we report the synthesis and characterisation of five *m*-terphenyl Fe(II) complexes (1–5), four of which are novel, that all display field induced SMM behaviour. We show that, while 1–5 display similar molecular geometries in the solid state, their relaxation profiles are strongly influenced by the nature of the *para*-substituent R.

Results and discussion

Synthesis and structural characterisation

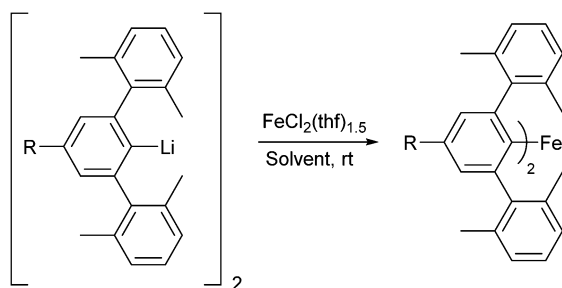
Reaction of the organolithium complexes [4-*R*-2,6-Xyl₂C₆H₂Li]₂ (R = *t*Bu, SiMe₃, H, Cl, CF₃)⁵¹ in solution with FeCl₂·(thf)_{1.5} resulted in the formation of the corresponding Fe(II) *m*-terphenyl complexes (4-*R*-2,6-Xyl₂C₆H₂)₂Fe [R = *t*Bu (1), SiMe₃ (2), H (3), Cl (4), CF₃ (5); Scheme 1]. Note that these complexes are numbered in order of the electron donating/withdrawing power of the R group (as quantified by their Hammett constant, σ_{para})⁵⁴ with 1 (*t*Bu) being the most electron donating, and 5 (CF₃) the most electron withdrawing. For 1–4, the reaction was carried out in a mixture of toluene and THF (*ca.* 10 : 1). For complex 5 (R = CF₃), carrying the reaction out in diethyl ether gave significantly better yields (57% *vs.* 36%). The complexes were purified by recrystallisation from hexane (1, 2, 5) or toluene (3, 4) in moderate to good yields (42–58%). Complexes 1–5 are highly air- and moisture-sensitive, but stable over long periods (>1 year) at room temperature under inert atmosphere. Complex 5 is noted to be particularly sensitive, with crystalline samples undergoing rapid surface decomposition (from yellow to orange/dark brown) upon exposure to trace amounts of air.

Complex 3 (R = H) has been reported previously, and its characterisation matched that of the literature.⁴⁸ The novel complexes 1, 2, 4 and 5 have been characterised by X-ray crystallography, mass spectrometry, elemental analysis; and IR, UV/Vis, and ¹H NMR spectroscopy. These complexes crystallise as yellow-green or yellow solids, dissolving to give bright yellow

solutions with broad UV/Vis absorptions (λ_{max} *ca.* 370 nm; ϵ = 830–1550 dm³ mol⁻¹ cm⁻¹; Fig. S9†).

The crystal structures for 1–5 determined by X-ray diffraction (Fig. 1, Fig. S10–S13†) show the iron is two-coordinate and the complexes are all monomeric in the solid state, with bent C–Fe–C bond angles over a range of 167.71(9)–174.9(2)° and a narrow range of Fe–C bond lengths [2.019(4)–2.041(4) Å] (Table 1). All structures feature large Fe...Fe separations of >9 Å and, despite displaying similar molecular geometries, they crystallise in different space groups (Table S2†) with different numbers of symmetry independent molecules in the unit cell (*Z'* = 1 for 1, 3, 5; 1.5 for 2; 3 for 4). Additionally, the crystal structure of 5 contains 0.5 co-crystallised *iso*-hexane molecules per formula unit (ESI, section S4.1†), which is also observed in the ¹H NMR spectrum of 5 (Fig. S6†).

Notably, the structures with *Z'* > 1 (2 and 4) show relatively broad variation in C–Fe–C bond angles and C–Fe bond lengths (Table 1). This suggests that changing the electronic character of the R group has little effect on the solid-state structures, with the variations in bond length and angle likely dominated by crystal packing effects. This is in line with our previous work on these ligands with lithium and Group 12 metal ions.^{51,52} The bond lengths and angles about the Fe centres are similar to those observed for the related *m*-terphenyl carbanion complex, (2,6-Mes₂C₆H₃)₂Fe.⁴⁰ However, the C–Fe–C bond angles are less acute than those seen in the more sterically demanding *m*-terphenyl carbanion complex, (2,6-Dipp₂C₆H₃)₂Fe [C–Fe–C = 159.34(6) Å].⁴¹ These non-linear structures can be contrasted with the field-induced SMMs Fe[N(H)Ar]₂ [Ar = C₆H₃-2,6-(C₆H₃-2,6-*i*Pr₂)₂; C₆H₃-2,6-(C₆H₂-2,4,6-



Scheme 1 Synthesis of diaryl complexes (4-*R*-2,6-Xyl₂C₆H₂)₂Fe 1–5 [R = *t*Bu (1), SiMe₃ (2), H (3), Cl (4), CF₃ (5)]. Solvent = toluene/THF (10 : 1) for 1–4; diethyl ether for 5.

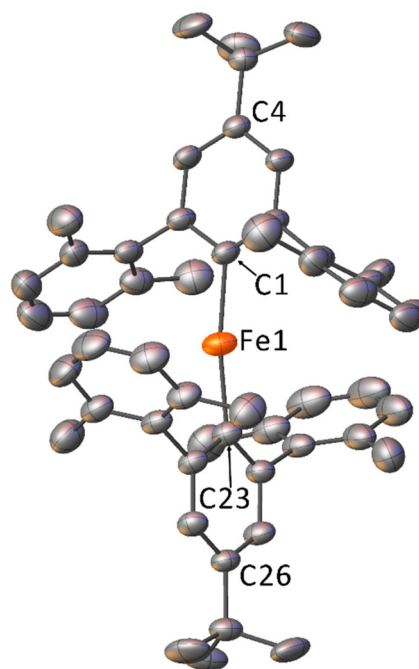


Fig. 1 Crystal structure of 1 with displacement ellipsoids set to 30%. Hydrogen atoms and minor disorder components omitted for clarity.



Table 1 Selected bond lengths (Å) and angles (°) for 1–5. Values for 3 taken from the literature.⁴⁸ Multiple entries refer to the symmetry independent molecules present in the asymmetric unit

Compound	Fe1–C1	Fe1–C23	C1–Fe1–C23	Fe1–C1...C4	Fe1–C23...C26
1	2.034(4)	2.022(4)	170.0(1)	174.6(2)	167.8(2)
2 ₁	2.026(4)	2.018(4)	172.2(2)	169.4(2)	170.4(2)
2 ₂	2.019(4)	2.019(4)	174.9(2)	170.7(2)	170.7(2)
3	2.027(2)	2.029(2)	167.71(9)	167.4(1)	174.9(1)
4 ₁	2.031(2)	2.030(2)	170.50(8)	169.0(1)	171.5(1)
4 ₂	2.023(2)	2.034(2)	170.74(8)	175.7(1)	165.1(1)
4 ₃	2.041(2)	2.033(2)	168.30(8)	165.7(1)	170.99(9)
5	2.039(1)	2.034(1)	170.04(5)	168.77(7)	172.99(7)

$iPr_2)_2]$, which also feature *m*-terphenyl moieties in their amide ligands, but which display a linear coordination environment around iron.²⁴

The ¹H NMR spectroscopic signals of these complexes are broad and occur over a wide range of chemical shifts (+181.6 to –59.5 ppm) owing to their paramagnetism (Fig. S3–S6†). Despite this, it is possible to assign protons based on integration of the signals. For 1–5, a linear correlation is observed between σ_{para} ⁵⁴ of the R group and some of the ¹H NMR signals; with the C₆H₂ *m*-H signals shifting upfield and the methyl signals for the xyllyl substituents shifting downfield with increasing σ_{para} (Table S1, Fig. S1†). Both trends bring the signals closer to 0 ppm (*i.e.* reduce the magnitude of the pseudocontact shift) as the R group becomes more electron withdrawing (Fig. S2†). These data suggest that the degree of spin delocalisation onto the ligand can be controlled by varying the electronics of R.⁵⁵ It should be noted, however, that no correlation is observed between σ_{para} and the computed zero-field splitting (ZFS) parameters of 1–5 (see below).

Electrochemistry

Cyclic voltammetry (CV) studies on THF solutions of 1–5 revealed that all complexes undergo a reduction at negative potentials (Fig. 2). Previous CV studies on related Group 12 *m*-terphenyl complexes [(4-R-2,6-Xyl₂C₆H₃)₂Hg, R = *t*Bu, CF₃]

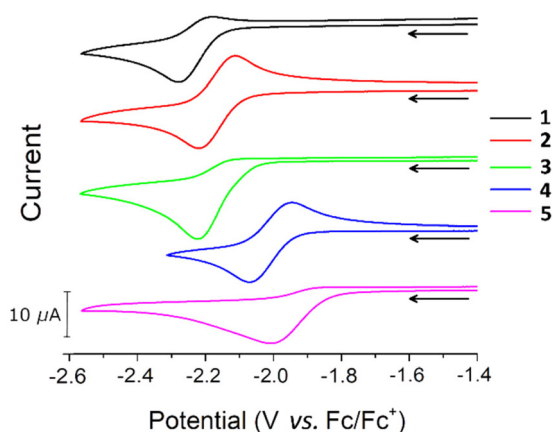


Fig. 2 Cyclic voltammograms for 1–5. All voltammograms recorded in 0.5 M [*n*Bu₄N][BF₄] solution in THF at $\nu = 100$ mV s^{–1}.

indicated that these ligands are not redox active in this potential range,⁵² so it is likely that the process corresponds to a Fe(II) to Fe(I) reduction. Cyclic voltammograms recorded at 100 mV s^{–1} show a redox couple for complexes 2 and 4, with properties characteristic of a diffusion controlled process,⁵⁶ but little to no return wave for 1, 3, and 5 (Fig. 2, Fig. S15–S19, Table S3†). At higher scan rates (>200 mV s^{–1}), more pronounced return waves for 1, 3, and 5 are observed, suggesting that the reduced complexes are prone to decomposition. A linear relationship between the cathodic peak current ($I_{p,c}$) and the square-root of the scan rate ($\nu^{1/2}$) is observed for 1–5 (Fig. S20†), which suggests the reduction is diffusion controlled for all complexes.⁵⁶ It is unclear why certain R groups better stabilise the reduced species since there is no obvious correlation with the steric bulk or electron withdrawing capability of the substituent. However, the cathodic peak potentials ($E_{p,c}$) of 1–5 do correlate with σ_{para} ⁵⁴ (Fig. S14†) such that more electron donating groups shift $E_{p,c}$ to more negative voltages, indicative of a linear free energy relationship across the series.

Electronic structure calculations

The electronic structures of 1–5 were studied with complete active space self-consistent field spin-orbit (CASSCF-SO) calculations using OpenMolcas⁵⁷ on the geometries determined by X-ray crystallography (see ESI, section S6.1† for details). The first four electronic states are grouped as two pseudo-doublets, with an energy separation that ranges between 159 and 173 cm^{–1} (Table S5†). These are well-isolated from the fifth electronic state, which is consistently predicted at *ca.* 250 cm^{–1}. Assuming a ground multiplet with an effective $S = 2$ spin, the anisotropy of the complexes can be modelled as a second-order effect *via* a ZFS Hamiltonian according to eqn (1):⁵⁸

$$H = D \left(S_z^2 - \frac{S(S+1)}{3} \right) + E(S_x^2 - S_y^2) \quad (1)$$

with the axial and rhombic terms defined as $D = (3/2)D_z$ and $E = (D_x - D_y)/2$, respectively. The resulting *g*-tensors and ZFS parameters (Table S6†) are in agreement with previously reported Fe(II) compounds.²⁶ A comparison between the experimental, calculated (CASSCF-SO), and fitted (*fit1*, eqn (2), see below) $\chi_m T$ plots for 1–5 is shown in Fig. S21 and S22.† We observe no correlation between the ZFS parameters (either



calculated or fitted) and the electron donating strength of the R groups or the C–Fe–C bond angles of the crystal structures. Given the predicted energy separation between the first two pseudo-doublets (159–173 cm⁻¹) one would expect a large effective energy barrier (U_{eff}) associated to an Orbach relaxation in 1–5 comparable to the linear Fe(II) complexes Fe[C(SiMe₃)₃]₂ ($U_{\text{eff}} = 146$ cm⁻¹) and Fe[N(SiMe₃)(Dipp)]₂ ($U_{\text{eff}} = 181$ cm⁻¹). However, this could not be observed experimentally in 1–5 due to efficient QTM (in zero-field) and Raman and direct (in field) relaxation processes (see below). For completeness, CASPT2 calculations were also performed for 1 (see ESI, section S6.1† for details), which showed the largest deviation between CASSCF predicted and experimental magnetic data. However, the CASPT2 results were very similar to the CASSCF (Tables S7 and S8†), suggesting that deviation from experiment does not arise from a lack of dynamical correlation as in work by Neese *et al.*,^{24,25,59} and thus further CASPT2 calculations were not deemed worthwhile.

Static magnetic characterisation

Magnetisation curves at 2, 3, 5 and 10 K up to 40 kOe (Fig. S24†), together with variable temperature magnetic susceptibility data under a 1000 Oe applied field (Fig. 3), were collected on ground, polycrystalline samples of 1–5. Complexes 1–5 show $\chi_{\text{m}}T$ values at 300 K (Table 2) far in excess of the predicted spin-only values for Fe(II) (3.00 cm³ K mol⁻¹ for $S = 2$,

$g = 2.00$). While 1 and 5 give the lowest and highest values for $\chi_{\text{m}}T$ at 300 K respectively (4.07 and 5.03 cm³ K mol⁻¹), there is not a clear correlation between $\chi_{\text{m}}T$ and σ_{para} ($R^2 = 0.370$, Fig. S21†). The $\chi_{\text{m}}T$ data for the 1–5 bear strong similarities to a previously reported series of linear and near-linear 2-coordinate Fe(II) complexes,²⁴ particularly the presence of a maximum in $\chi_{\text{m}}T$ at *ca.* 75 K (Fig. 3). This maximum indicates that the orbital angular momentum is unquenched due to the axiality of the systems (Table 1). This agrees with previous studies in which only markedly non-linear two-coordinate Fe(II) complexes failed to display this maximum, *i.e.* the bent Fe[N(H)Dmp]₂ (Dmp = 2,6-Mes₂C₆H₃; N–Fe–N = 140.9(2)°).²⁴ For 1–5, starting at 300 K, the increase of $\chi_{\text{m}}T$ on cooling suggests depopulation of states with a lower magnetic moment due to anisotropy,^{24,60} followed by a decrease in $\chi_{\text{m}}T$ at the lowest temperatures.

Assuming a ground multiplet with an effective $S = 2$ spin, the anisotropy of the complexes can be described as a second-order effect. As such, the magnetic data can be modelled through a combination of a crystal field and a Zeeman Hamiltonian (eqn (2)),⁵⁸ as implemented in PHI.⁶¹

$$\hat{H} = \hat{H}_{\text{CF}} + \hat{H}_{\text{Zee}} = B_2^0 \theta_2 \hat{O}_2^0 + B_2^2 \theta_2 \hat{O}_2^2 + \mu_{\text{B}} \vec{S} \cdot \vec{g} \cdot \vec{B} \quad (2)$$

where B_k^q are the crystal-field parameters, θ_2 the operator equivalent factor, \hat{O}_2^0 the operator equivalent, and \vec{g} the g -tensor. Within this formalism, $D = 3B_2^0 \theta_2$ and $E = B_2^2 \theta_2$ describe the axial and rhombic terms of a zero-field splitting Hamiltonian, respectively.

Initially, a global fitting of both the susceptibility and magnetisation was attempted for 1–5.⁶² To avoid over-parameterisation, the CASSCF-SO values were used as a starting point, with $g_x = g_y \neq g_z$ and B_2^2 fixed to the CASSCF-SO value (see above and Table S6†). However, the obtained fit was biased towards the magnetisation curves, which drastically worsened the model's agreement with the susceptibility traces. This is tentatively assigned to the way residuals are calculated in a global fitting. Further data analysis revealed that both the high temperature susceptibility and magnetisation curves tend towards an anisotropic model, whereas low temperature susceptibility suggested otherwise. To overcome this apparent inconsistency, $\chi_{\text{m}}T$ was first fit using D and $g_x = g_y \neq g_z$ as parameters, biasing toward the high temperature susceptibility data with PHI's "Residual" keyword. Next, both the $\chi_{\text{m}}T$ and M vs. H data were fit, fixing the previously optimised parameters, and allowing E to vary as well as introducing a mean-field parameter (zJ) to model intermolecular interactions. This was needed to reproduce the drop in $\chi_{\text{m}}T$ at low temperatures. The values obtained from this fit are labelled as *fit1* (Table 3, Fig. 3). Finally, a global fitting without any constraints was performed, labelled *fit2* (Table S10†). Both approaches predict small antiferromagnetic interactions for all complexes, compatible with the reported intermolecular interactions. A comparison between experimental and fitted values with both models is presented in Fig. S24 and S25,† highlighting that a sizeable change in parameters between *fit1* and *fit2* leads to a very

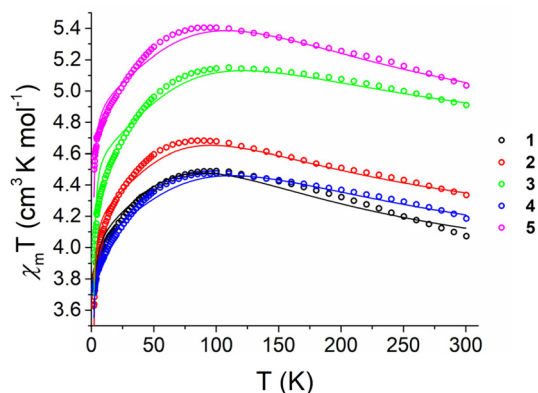


Fig. 3 Temperature dependence of $\chi_{\text{m}}T$ values for powdered polycrystalline samples of 1–5 measured under a 1000 Oe field between 1.8 and 300 K (circles) with calculated fittings (*fit1*) according to eqn (2) (lines).

Table 2 $\chi_{\text{m}}T$ at 300 K and low temperature M_{sat} values for complexes 1–5 as determined by DC magnetometry. M_{sat} determined at 4 T and 1.8 K

Sample	$\chi_{\text{m}}T$ (cm ³ K mol ⁻¹)	M_{sat} (μ_{B})
1	4.07	2.7
2	4.34	2.8
3	4.91	2.9
4	4.19	2.6
5-(C ₆ H ₁₄) _{0.5}	5.03	3.3



Table 3 Zero-field splitting Hamiltonian parameters for 1–5 determined using the procedure *fit1* as described in the main text. Fitting conducted using PHI⁶¹

Compound	$g_x = g_y$	g_z	D (cm ⁻¹)	E (cm ⁻¹)	zJ (cm ⁻¹)
1	1.80	2.87	-59.14	2.94	-1.6×10^{-2}
2	1.90	2.91	-59.91	-6.13	-7.3×10^{-3}
3	2.08	3.03	-68.54	7.19	-2.1×10^{-3}
4	1.82	2.86	-73.07	2.69	-1.7×10^{-2}
5	1.98	3.15	-72.39	4.69	-9.9×10^{-3}

modest improvement in M vs. H but no significant improvement in $\chi_m T$. Based on this, and because the values obtained from *fit1* are much closer to the CASSCF-SO values than *fit2* (Table S6†), we conclude that the much more anisotropic values of *fit1* are the most representative. Still, it is worth mentioning that the so-obtained E values are at least one order of magnitude larger than the CASSCF-SO ones, significantly reducing the E/D ratio, and one obtains the same fitting regardless of its sign. Fitting with $g_x \neq g_y$ or including magnetic impurities does not improve the fittings.

It should be noted that while the simulated (*fit1*) M vs. H curves match the experimental data well for 2, 3, and 5; the agreement is less good for 1 and 4 (Fig. S24†). The reason for this is unclear, but could arise from small errors in the mass (a ca. 7% error in the mass of 1 would give good agreement between the simulated and experimental curves) or from structural variations, as indicated by the large α values from the generalised Debye model.⁶³ Nevertheless, the experimental profile of the M vs. H curves is reproduced for all simulations, and the shape of the $\chi_m T$ plots is well captured.

Plots of M vs. H/T for 1–5, recorded at 10, 20, 30, and 40 kOe over a temperature range of 1.8–5.0 K, show similar behaviour to other reported 2-coordinate 3d complexes^{24,28,34} with data that are non-superimposable on a single Brillouin curve (Fig. S26–S30†). This provides further evidence of strong magnetic anisotropy in 1–5.^{24,64} M_{sat} values for 1–5 of 2.6–3.3 μ_B (Table 2) are far lower than the spin-only 4.0 μ_B expected for a $S = 2$ centre, however, these values are comparable to those obtained for other linear Fe(II) complexes.²⁴

Dynamic magnetic characterisation

AC magnetic susceptibility measurements were performed on 1–5 to investigate their low temperature magnetic relaxation. For all AC susceptibility measurements, magnetic relaxation times (τ) were determined by a global fitting of the in phase (χ') and out-of-phase susceptibilities (χ'') vs. frequency using a generalised Debye model⁶⁵ as implemented in the CC-FIT2 software package.⁶³

For 1–5, no out-of-phase susceptibility signal (χ'') was detected at zero-field over the frequency range 1–1400 Hz. This is likely due to rapid quantum tunnelling of magnetisation (QTM), which is commonly observed in mononuclear Fe(II) species.^{20–24,26,27} However, 1–5 exhibit slow magnetic relaxation under an applied DC field. To determine the dependence of τ on field strength, AC susceptibility measurements were

performed on 1–5 at 2 K (1–1400 Hz switching frequency) over a range of applied DC fields ($H = 100$ –5000 Oe). For 1–5, τ initially increases with increasing applied field as QTM is reduced, reaches a maximum at a certain field strength (H_{max}), and then decreases (Fig. S36 and S37†) following the onset of a direct process. This behaviour is similar to that observed for previously reported Fe(II) field induced SMMs.^{20–24,26,27} Estimates of H_{max} for 1–5 are provided in Table 4, and fall in the range 1000–2000 Oe. At 2 K and an applied field of 1500 Oe, τ is observed to increase in the order $5 < 4 < 2 < 3 \ll 1$. This trend approximately parallels the electron donating/withdrawing power of the R group on the *m*-terphenyl ligand, with the electron-donating *t*Bu (1) showing by far the longest relaxation time (Fig. S36†), and the electron-withdrawing CF₃ (5) the shortest (Fig. S37†). It should be noted that τ values were not determined for 1 at $H = 100$ –500 Oe or 3 at $H = 100$ –200 Oe as the χ' vs. χ'' data could not be well fit to the generalised Debye model.

Subsequently, AC susceptibility measurements were performed for 1–5 at a fixed DC field (H_{max} , Table 4) over a range of temperatures (1.8–16.0 K) and frequencies (1–1400 Hz) to determine the temperature dependence of τ . Fig. 4a shows the out-of-phase susceptibility (χ'') of 1 as a function of frequency, while plots of χ'' as a function of frequency (Fig. S38–S42†), temperature (Fig. S43–S47†) and Cole–Cole plots (Fig. S48–S52†) for 1–5 are provided in the ESI. 1–5 all show a maximum in χ'' which shifts to higher frequencies with increasing temperature. These data were fit to the general Debye model (Tables S11–S15†) and \log_{10} – \log_{10} plots of τ^{-1} vs. T , henceforth referred to as relaxation profiles, were constructed. The relaxation profiles for 1–5 are presented in Fig. 4b. Note that for 2, 4, and 5 relaxation rates are not given below 2.4 K; this is because the Cole–Cole plots for these complexes (Fig. S49, S51 and S52†) could not be well fit to the general Debye model due to the presence of a low frequency process. There were insufficient low frequency data points to satisfactorily model these according to a 2-process equation.

Given that the data has been collected in field, the relaxation profiles for 1–5 were fit by considering a combination of Raman and direct terms, according to eqn (3).⁶³

$$\tau^{-1} = 10^R T^n + 10^A T \quad (3)$$

where $10^R = C$ and n are the Raman parameters, and $10^A = B$ is the direct term that implicitly accounts for the effect of the external field. We note that the α values associated with the

Table 4 Summary of AC magnetic data for 1–5. Parentheses indicate the error on the associated parameter

Compound	H_{max} (Oe)	$\log_{10}[C/(s^{-1} K^{-n})]$	n	$\log_{10}[A/(s^{-1} \text{Oe}^{-1} K^{-1})]$
1	1000	-1.12(2)	4.98(2)	0.839(6)
2	1750	-0.58(7)	5.19(9)	2.062(9)
3	1000	-0.73(2)	4.74(2)	1.712(4)
4	2000	-0.1(1)	4.6(1)	2.22(1)
5	2000	-1(1)	5(1)	2.82(2)



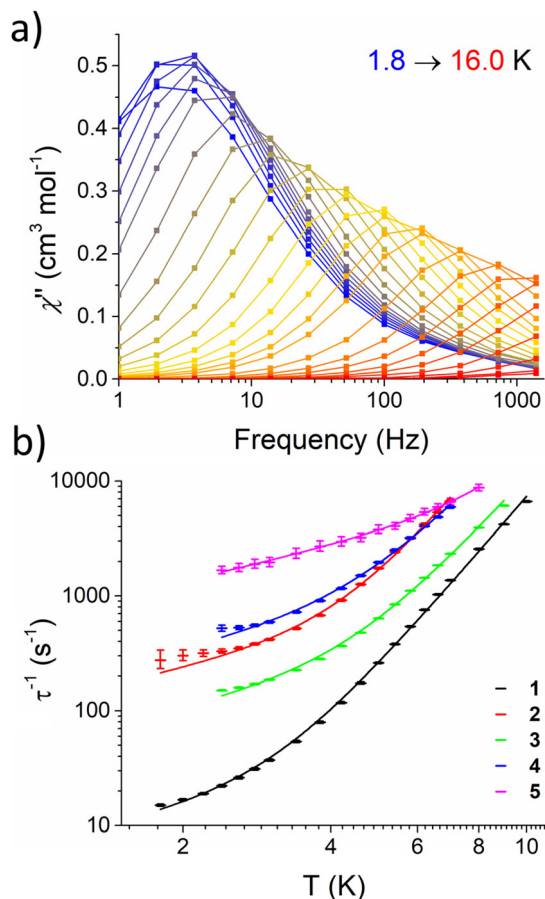


Fig. 4 (a) Frequency dependence of χ'' for **1** over the temperature range 1.8–16.0 K, recorded on a MPMS magnetometer at 1000 Oe applied field. Lines are intended as guides for eyes. (b) Relaxation profile for **1–5** under an applied field (H_{max} , Table 4) with fittings (solid lines) according to eqn (3). Error bars represent error in the fitting of τ from Cole–Cole plots.

generalised Debye model are large, likely due to considerable structural variations, leading to large estimated standard deviations (ESDs)⁶³ on τ and hindering a meaningful comparison between **1–5** (see Table S16[†]). As such, we have performed the fitting of the relaxation profile without considering the ESDs. We note, however, that consideration of the α parameters is uncommon for 3d SMMs; many recent publications do not provide them.^{18,19,27} Those we found that do provide them show similar α parameters to those reported here, but do not make it clear if they were considered in fitting the relaxation profiles.^{66,67} It is thus difficult to compare the large uncertainties obtained from consideration of α parameters here to other literature examples.

Nevertheless, the optimised parameters for a given compound are virtually identical with and without considering ESDs (see Fig. S53–S62, Table S7[†]). The relaxation parameters (C , n , A) for **1–5** are presented in Table 4. The parameters associated to the Raman process are in agreement with those found for other Fe(II) complexes²⁶ and, importantly, the characteristic n term is consistently close to 5. The data was

also fitted considering an Orbach process, but this model was discarded as it led to effective barriers one order of magnitude smaller than the CASSCF predicted first excited state (see above). While this model affords a satisfactory fit of the data, recent studies have shown that spin–phonon coupling mediated by low energy phonons can be important for rationalising under-barrier relaxation processes in SMMs.^{68–70} We thus considered two different models which account for spin–lattice relaxation published by Wu (eqn (S1)[†]),⁶⁹ and Lunghi (eqn (S2)[†]).⁶⁸ In both cases, a reasonable fit of the data for complex **1** (Wu and Lunghi models, Fig. S63 and S65[†]) and **2** (Wu model only, Fig. S64[†]) could only be obtained through the use of physically unreasonable parameters, with spin phonon couplings several orders of magnitude larger than the expected values of 0.1–1.0 cm^{-1} (Tables S17 and S18[†]). We thus concluded that eqn (3) is the most appropriate model for the relaxation rates of **1–5**.

Fig. 4b shows that magnetic relaxation rates increase in order $1 < 3 < 2 < 4 < 5$ over the direct-dominated temperature range (1.8–5.0 K), and that **1** and **5** remain the slowest and fastest relaxing across the whole temperature range, respectively. This is likely because the direct mechanism dominates at $T < 5$ K; with **5**, **4**, and **2** being measured under the largest external fields. Nonetheless, for molecules measured under the same external field (Table 4), more electron withdrawing groups lead to faster relaxation rates: *i.e.* **5** relaxes *ca.* 2 \times faster than **4** and **3** relaxes *ca.* 10 \times faster than **1**. At higher temperatures ($T > 6.0$ K) the relaxation profiles begin to converge and the difference in their relaxation rates becomes less pronounced. This indicates that, despite a change in the phonon density of states associated with the different R groups, the Raman mechanism remains largely unaffected in the studied temperature range. This is somewhat surprising, as **1–5** crystallise in different structure types with different space groups, and one might expect to observe different low-energy phonon modes, changing the efficacy of the Raman mechanism.

Hysteresis measurements were also performed on **1–5** at 1.8 K and a sweep rate of 67 Oe s^{-1} . Under these conditions, slight open hysteresis was observed for complexes **1** and **3** (Fig. S31 and S33[†]), but not for **2**, **4**, or **5** (Fig. S32, S34, and S35[†]). This is in line with the AC susceptibility data, which shows that **1** and **3** have the slowest rates of magnetic relaxation.

Conclusions

The reaction of the *m*-terphenyl lithium salts [4-R-2,6-Xyl₂C₆H₂Li]₂ (R = *t*Bu, SiMe₃, H, Cl, CF₃) and FeCl₂·(thf)_{1.5} affords Fe(II) complexes **1–5** as yellow-green crystalline solids. Single crystal X-ray diffraction demonstrates that **1–5** all feature two-coordinate Fe(II) ions in near-linear geometries. DC magnetometry measurements on **1–5** show maxima in the $\chi_m T$ plots at *ca.* 75 K, $\chi_m T$ values which exceed the spin-only value, and discontinuity in plots of M vs. H/T at low temperatures, indicating substantial orbital angular momentum contri-



butions leading to large magnetic anisotropy. These observations are corroborated by the highly axial values obtained from fitting the magnetic data to the ZFS Hamiltonian. *Ab initio* calculations on the crystal structures afford computed ZFS Hamiltonian parameters which are also indicative of a highly axial ligand field. Complexes 1–5 all display field-induced SMM behaviour, with the rate of relaxation dependent upon the identity of the R group. The relaxation profiles were fit to a combination of direct and Raman relaxation processes and show that the most electron donating group (1) relaxes considerably slower than the most electron withdrawing counterpart (5) across the whole investigated temperature range.

While 1–5 exhibit modest SMM properties, this work demonstrates that the *m*-terphenyls are a tuneable and versatile class of ligand capable of inducing highly axial ligand fields upon a metal centre. These ligands have not been explored in this context, and this study opens exciting opportunities to apply them to other metal ions and thus establish magneto-structural correlations for the development of SMMs with improved relaxation behaviour.

Author contributions

AJV developed the synthesis and characterisation for 1, 2, 4, and 5, which was continued by LJ and WT. AMG and TJB performed synthesis and initial magnetic measurements on complex 3. MJC helped to perform MPMS measurements and assisted with data analysis and interpretation. ESD performed CV measurements on 1–5. SPA and WL helped with the solution and refinement of single-crystal X-ray diffraction data for 1–5. JM provided guidance and support to AV and LJ throughout the project. LJ performed the MPMS measurements on 1–5 presented in this manuscript, analysed the magnetic data, prepared the initial draft of the manuscript, and coordinated with other authors. DR performed CASSCF calculations and fittings of the magnetic data and wrote these sections for the manuscript. He also helped interpret the magnetic data and with the development of the manuscript. DLK was the PI for this research, conceived of the project, helped with writing, and provided input throughout.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

We acknowledge the EPSRC [grant number EP/R004064/1]; the Leverhulme Trust [grant numbers RPG-2014-317; RF-2021-102]; and the University of Nottingham [Hobday Bequest] for financial support of this research. We would also like to thank the National Mass Spectrometry Facility at Swansea University for mass spectrometry measurements and the Microanalysis

Service at London Metropolitan University for elemental microanalysis measurements. We thank Dr María Giménez López (University of Santiago de Compostela) for preliminary discussions, and Prof. Mark Murrie and Dr Emma Regincós Martí (University of Glasgow) for helpful discussions about AC susceptibility data.

References

- D. Gatteschi, R. Sessoli and J. Villain, *Molecular Nanomagnets*, Oxford University Press, 2006.
- M. J. Giansiracusa, G. K. Gransbury, N. F. Chilton and D. P. Mills, in *Encyclopedia of Inorganic and Bioinorganic Chemistry*, John Wiley & Sons, Ltd., 2021, p. eibc2784.
- M. Shiddiq, D. Komijani, Y. Duan, A. Gaita-Ariño, E. Coronado and S. Hill, *Nature*, 2016, **531**, 348–351.
- R. Vincent, S. Klyatskaya, M. Ruben, W. Wernsdorfer and F. Balestro, *Nature*, 2012, **488**, 357–360.
- R. Sessoli, H. Tsai, A. R. Schake, S. Wang, J. B. Vincent, K. Folting, D. Gatteschi, G. Christou and D. N. Hendrickson, *J. Am. Chem. Soc.*, 1993, **115**, 1804–1816.
- N. Ishikawa, M. Sugita, T. Ishikawa, S. Y. Koshihara and Y. Kaizu, *J. Am. Chem. Soc.*, 2003, **125**, 8694–8695.
- N. Ishikawa, M. Sugita, T. Okubo, N. Tanaka, T. Iino and Y. Kaizu, *Inorg. Chem.*, 2003, **42**, 2440–2446.
- C. A. P. Goodwin, F. Ortu, D. Reta, N. F. Chilton and D. P. Mills, *Nature*, 2017, **548**, 439–442.
- K. R. McClain, C. A. Gould, K. Chakarawet, S. J. Teat, T. J. Groshens, J. R. Long and B. G. Harvey, *Chem. Sci.*, 2018, **9**, 8492–8503.
- F. S. Guo, B. M. Day, Y. C. Chen, M. L. Tong, A. Mansikkamäki and R. A. Layfield, *Science*, 2018, **362**, 1400–1403.
- V. S. Parmar, D. P. Mills and R. E. P. Winpenny, *Chem. – Eur. J.*, 2021, **27**, 7625–7645.
- C. A. Gould, K. R. McClain, D. Reta, J. G. C. Kragoskow, D. A. Marchiori, E. Lachman, E. S. Choi, J. G. Analytis, R. D. Britt, N. F. Chilton, B. G. Harvey and J. R. Long, *Science*, 2022, **375**, 198–202.
- G. A. Craig and M. Murrie, *Chem. Soc. Rev.*, 2015, **44**, 2135–2147.
- P. P. Samuel, K. C. Mondal, N. Amin Sk, H. W. Roesky, E. Carl, R. Neufeld, D. Stalke, S. Demeshko, F. Meyer, L. Ungur, L. F. Chibotaru, J. Christian, V. Ramachandran, J. Van Tol and N. S. Dalal, *J. Am. Chem. Soc.*, 2014, **136**, 11964–11971.
- J. M. Zadrozny, D. J. Xiao, M. Atanasov, G. J. Long, F. Grandjean, F. Neese and J. R. Long, *Nat. Chem.*, 2013, **5**, 577–581.
- W. M. Reiff, A. M. LaPointe and E. H. Witten, *J. Am. Chem. Soc.*, 2004, **126**, 10206–10207.
- W. M. Reiff, C. E. Schulz, M. H. Whangbo, J. I. Seo, Y. S. Lee, G. R. Potratz, C. W. Spicer and G. S. Girolami, *J. Am. Chem. Soc.*, 2009, **131**, 404–405.



- 18 P. Wang, M. R. Saber, P. E. Vannatta, G. P. A. Yap, C. V. Popescu, C. C. Scarborough, M. T. Kieber-Emmons, K. R. Dunbar and C. G. Riordan, *Inorg. Chem.*, 2021, **60**, 6480–6491.
- 19 Y. Li, J. Xi, J. Ferrando-Soria, Y. Q. Zhang, W. Wang, Y. Song, Y. Guo, E. Pardo and X. Liu, *Dalton Trans.*, 2022, **51**, 8266–8272.
- 20 D. E. Freedman, W. H. Harman, T. D. Harris, G. J. Long, C. J. Chang and J. R. Long, *J. Am. Chem. Soc.*, 2010, **132**, 1224–1225.
- 21 D. Weismann, Y. Sun, Y. Lan, G. Wolmershäuser, A. K. Powell and H. Sitzmann, *Chem. – Eur. J.*, 2011, **17**, 4700–4704.
- 22 X. Feng, C. Mathonière, I. R. Jeon, M. Rouzières, A. Ozarowski, M. L. Aubrey, M. I. Gonzalez, R. Clérac and J. R. Long, *J. Am. Chem. Soc.*, 2013, **135**, 15880–15884.
- 23 C. Mathonière, H. J. Lin, D. Siretanu, R. Clérac and J. M. Smith, *J. Am. Chem. Soc.*, 2013, **135**, 19083–19086.
- 24 J. M. Zadrozny, M. Atanasov, A. M. Bryan, C. Y. Lin, B. D. Rekker, P. P. Power, F. Neese and J. R. Long, *Chem. Sci.*, 2013, **4**, 125–138.
- 25 M. Atanasov, J. M. Zadrozny, J. R. Long and F. Neese, *Chem. Sci.*, 2013, **4**, 139–156.
- 26 T. Bodenstern and A. Eichhöfer, *Dalton Trans.*, 2019, **48**, 15699–15712.
- 27 Q. Q. Su, Q. Yuan, X. F. Wu, S. H. Chen, J. Xiang, X. X. Jin, L. X. Wang, B. W. Wang, S. Gao and T. C. Lau, *Chem. Commun.*, 2021, **57**, 781–784.
- 28 X. N. Yao, J. Z. Du, Y. Q. Zhang, X. B. Leng, M. W. Yang, S. Da Jiang, Z. X. Wang, Z. W. Ouyang, L. Deng, B. W. Wang and S. Gao, *J. Am. Chem. Soc.*, 2017, **139**, 373–380.
- 29 T. Jurca, A. Farghal, P. H. Lin, I. Korobkov, M. Murugesu and D. S. Richeson, *J. Am. Chem. Soc.*, 2011, **133**, 15814–15817.
- 30 V. V. Novikov, A. A. Pavlov, Y. V. Nelyubina, M. E. Boulon, O. A. Varzatskii, Y. Z. Voloshin and R. E. P. Winpenny, *J. Am. Chem. Soc.*, 2015, **137**, 9792–9795.
- 31 S. Gómez-Coca, A. Urtizberea, E. Cremades, P. J. Alonso, A. Camón, E. Ruiz and F. Luis, *Nat. Commun.*, 2014, **5**, 1–8.
- 32 F. Habib, O. R. Luca, V. Vieru, M. Shiddiq, I. Korobkov, S. I. Gorelsky, M. K. Takase, L. F. Chibotaru, S. Hill, R. H. Crabtree and M. Murugesu, *Angew. Chem., Int. Ed.*, 2013, **52**, 11290–11293.
- 33 S. Gomez-Coca, E. Cremades, N. Aliaga-Alcalde and E. Ruiz, *J. Am. Chem. Soc.*, 2013, **135**, 7010–7018.
- 34 P. C. Bunting, M. Atanasov, E. Damgaard-Møller, M. Perfetti, I. Crassee, M. Orlita, J. Overgaard, J. Van Slageren, F. Neese and J. R. Long, *Science*, 2018, **362**, 1378.
- 35 R. Ruamps, R. Maurice, L. Batchelor, M. Boggio-Pasqua, R. Guillot, A. L. Barra, J. Liu, E. E. Bendeif, S. Pillet, S. Hill, T. Mallah and N. Guihéry, *J. Am. Chem. Soc.*, 2013, **135**, 3017–3026.
- 36 J. Vallejo, A. Pascual-Álvarez, J. Cano, I. Castro, M. Julve, F. Lloret, J. Krzystek, G. De Munno, D. Armentano, W. Wernsdorfer, R. Ruiz-García and E. Pardo, *Angew. Chem., Int. Ed.*, 2013, **52**, 14075–14079.
- 37 B. Twamley, S. T. Haubrich and P. P. Power, in *Advances in Organometallic Chemistry*, ed. R. West and A. F. Hill, Academic Press, 1999, vol. 44, pp. 1–65.
- 38 J. A. C. Clyburne and N. McMullen, *Coord. Chem. Rev.*, 2000, **210**, 73–99.
- 39 D. L. Kays, in *Organometallic Chemistry*, ed. I. J. S. Fairlamb and J. S. Lynam, Royal Society of Chemistry, Cambridge, 2010, vol. 36, pp. 56–76.
- 40 D. L. Kays and A. R. Cowley, *Chem. Commun.*, 2007, 1053–1055.
- 41 C. Ni and P. P. Power, *Chem. Commun.*, 2009, 5543–5545.
- 42 C. Ni, T. A. Stich, G. J. Long and P. P. Power, *Chem. Commun.*, 2010, **46**, 4466–4468.
- 43 C. Ni, H. Lei and P. P. Power, *Organometallics*, 2010, **29**, 1988–1991.
- 44 H. R. Sharpe, A. M. Geer, T. J. Blundell, F. R. Hastings, M. W. Fay, G. A. Rance, W. Lewis, A. J. Blake and D. L. Kays, *Catal. Sci. Technol.*, 2018, **8**, 229–235.
- 45 H. R. Sharpe, A. M. Geer, W. Lewis, A. J. Blake and D. L. Kays, *Angew. Chem., Int. Ed.*, 2017, **56**, 4845–4848.
- 46 A. J. South, A. M. Geer, L. J. Taylor, H. R. Sharpe, W. Lewis, A. J. Blake and D. L. Kays, *Organometallics*, 2019, **38**, 4115–4120.
- 47 H. R. Sharpe, A. M. Geer, H. E. L. Williams, T. J. Blundell, W. Lewis, A. J. Blake and D. L. Kays, *Chem. Commun.*, 2017, **53**, 937–940.
- 48 H. R. Sharpe, A. M. Geer, L. J. Taylor, B. M. Gridley, T. J. Blundell, A. J. Blake, E. S. Davies, W. Lewis, J. McMaster, D. Robinson and D. L. Kays, *Nat. Commun.*, 2018, **9**, 3757.
- 49 R. Wolf, C. Ni, T. Nguyen, M. Brynda, G. J. Long, A. D. Sutton, R. C. Fischer, J. C. Fettinger, M. Hellman, L. Pu and P. P. Power, *Inorg. Chem.*, 2007, **46**, 11277–11290.
- 50 Y. Peng, R. C. Fischer, W. A. Merrill, J. Fischer, L. Pu, B. D. Ellis, J. C. Fettinger, R. H. Herber and P. P. Power, *Chem. Sci.*, 2010, **1**, 461–468.
- 51 A. J. Valentine, A. M. Geer, L. J. Taylor, A. M. Teale, K. E. Wood, H. E. L. Williams, W. Lewis, S. P. Argent, J. McMaster and D. L. Kays, *Dalton Trans.*, 2021, **50**, 722–728.
- 52 A. J. Valentine, L. J. Taylor, A. M. Geer, C. D. Huke, K. E. Wood, W. Tovey, W. Lewis, S. P. Argent, A. M. Teale, J. McMaster and D. L. Kays, *Organometallics*, 2022, **41**, 1426–1433.
- 53 Y. S. Meng, L. Xu, J. Xiong, Q. Yuan, T. Liu, B. W. Wang and S. Gao, *Angew. Chem., Int. Ed.*, 2018, **57**, 4673–4676.
- 54 C. Hansch, A. Leo and R. W. Taft, *Chem. Rev.*, 1991, **91**, 165–195.
- 55 G. Parigi, E. Ravera and C. Luchinat, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2019, **114–115**, 211–236.
- 56 N. Elgrishi, K. J. Rountree, B. D. McCarthy, E. S. Rountree, T. T. Eisenhart and J. L. Dempsey, *J. Chem. Educ.*, 2018, **95**, 197–206.
- 57 I. F. Galván, M. Vacher, A. Alavi, C. Angeli, F. Aquilante, J. Autschbach, J. J. Bao, S. I. Bokarev, N. A. Bogdanov, R. K. Carlson, L. F. Chibotaru, J. Creutzberg, N. Dattani,



- M. G. Delcey, S. S. Dong, A. Dreuw, L. Freitag, L. M. Frutos, L. Gagliardi, F. Gendron, A. Giussani, L. González, G. Grell, M. Guo, C. E. Hoyer, M. Johansson, S. Keller, S. Knecht, G. Kovačević, E. Källman, G. Li Manni, M. Lundberg, Y. Ma, S. Mai, J. P. Malhado, P. Å. Malmqvist, P. Marquetand, S. A. Mewes, J. Norell, M. Olivucci, M. Oppel, Q. M. Phung, K. Pierloot, F. Plasser, M. Reiher, A. M. Sand, I. Schapiro, P. Sharma, C. J. Stein, L. K. Sørensen, D. G. Truhlar, M. Ugandi, L. Ungur, A. Valentini, S. Vancoillie, V. Veryazov, O. Weser, T. A. Wesolowski, P. O. Widmark, S. Wouters, A. Zech, J. P. Zobel and R. Lindh, *J. Chem. Theory Comput.*, 2019, **15**, 5925–5964.
- 58 A. Abragam and B. Bleaney, *Electron Paramagnetic Resonance of Transition Ions*, Oxford University Press, 1970.
- 59 M. Atanasov, D. Ganyushin, D. A. Pantazis, K. Sivalingam and F. Neese, *Inorg. Chem.*, 2011, **50**, 7460–7477.
- 60 M. Kotani, *J. Phys. Soc. Jpn.*, 1949, **4**, 293–297.
- 61 N. F. Chilton, R. P. Anderson, L. D. Turner, A. Soncini and K. S. Murray, *J. Comput. Chem.*, 2013, **34**, 1164–1175.
- 62 S. Mugiraneza and A. M. Hallas, *Commun. Phys.*, 2022, **5**, 95.
- 63 D. Reta and N. F. Chilton, *Phys. Chem. Chem. Phys.*, 2019, **21**, 23567–23575.
- 64 C. Benelli and D. Gatteschi, *Introduction to Molecular Magnetism: From Transition Metals to Lanthanides*, Wiley-VCH, 2015.
- 65 K. S. Cole and R. H. Cole, *J. Chem. Phys.*, 1941, **9**, 341–351.
- 66 A. Cornia, A. L. Barra, V. Bulicanu, R. Clérac, M. Cortijo, E. A. Hillard, R. Galavotti, A. Lunghi, A. Nicolini, M. Rouzières, L. Sorace and F. Totti, *Inorg. Chem.*, 2020, **59**, 1763–1777.
- 67 J. Acharya, A. Sarkar, P. Kumar, V. Kumar, J. Flores Gonzalez, O. Cador, F. Pointillart, G. Rajaraman and V. Chandrasekhar, *Dalton Trans.*, 2020, **49**, 4785–4796.
- 68 A. Lunghi, F. Totti, R. Sessoli and S. Sanvito, *Nat. Commun.*, 2017, **8**, 14620.
- 69 L. Gu and R. Wu, *Phys. Rev. Lett.*, 2020, **125**, 117203.
- 70 M. Briganti, F. Santanni, L. Tesi, F. Totti, R. Sessoli and A. Lunghi, *J. Am. Chem. Soc.*, 2021, **143**, 13633–13645.

