

Cite this: *Chem. Sci.*, 2024, 15, 3003

All publication charges for this article have been paid for by the Royal Society of Chemistry

Metal–carbon bonding in early lanthanide substituted cyclopentadienyl complexes probed by pulsed EPR spectroscopy†

Lydia E. Nodaraki,^{‡,ab} Jingjing Liu,^{‡,a} Ana-Maria Ariciu,^{ab} Fabrizio Ortu,^{id a} Meagan S. Oakley,^{id a} Letitia Birnoschi,^a Gemma K. Gransbury,^{id a} Philip J. Cobb,^a Jack Emerson-King,^{id a} Nicholas F. Chilton,^{id *a} David P. Mills,^{id *a} Eric J. L. McInnes^{*ab} and Floriana Tuna^{id *ab}

We examine lanthanide (Ln)–ligand bonding in a family of early Ln³⁺ complexes [Ln(Cp^{tt})₃] (1-Ln, Ln = La, Ce, Nd, Sm; Cp^{tt} = C₅H₃^tBu₂-1,3) by pulsed electron paramagnetic resonance (EPR) methods, and provide the first characterization of 1-La and 1-Nd by single crystal XRD, multinuclear NMR, IR and UV/Vis/NIR spectroscopy. We measure electron spin T_1 and T_m relaxation times of 12 and 0.2 μ s (1-Nd), 89 and 1 μ s (1-Ce) and 150 and 1.7 μ s (1-Sm), respectively, at 5 K: the T_1 relaxation of 1-Nd is more than 10² times faster than its valence isoelectronic uranium analogue. ¹³C and ¹H hyperfine sublevel correlation (HYSCORE) spectroscopy reveals that the extent of covalency is negligible in these Ln compounds, with much smaller hyperfine interactions than observed for equivalent actinide (Th and U) complexes. This is corroborated by *ab initio* calculations, confirming the predominant electrostatic nature of the metal–ligand bonding in these complexes.

Received 17th November 2023

Accepted 12th January 2024

DOI: 10.1039/d3sc06175b

rsc.li/chemical-science

Introduction

Lanthanides (Ln) have numerous applications due to their unique physicochemical properties.¹ The text-book description of the electronic structure of Ln ions in the dominant 3+ oxidation state is that due to poor shielding the valence 4f orbitals are “core-like,” and as a result chemical bonding involving Ln ions is predominantly ionic. In contrast to Ln, the early actinides (An) exhibit variable oxidation states and participation in metal–ligand multiple bonding, which indicates greater involvement of the 5f (and 6d) valence orbitals in bonding regimes.² This is important because the differences in chemical bonding has direct impacts on the chemical behavior of Ln and An ions,³ and such differences are exploited in, for example, separations chemistry.^{4,5} Moreover, subtle variations in f-element–ligand bonding is important in determining electronic properties and hence applications in, for example, luminescence,⁶ paramagnetic NMR,⁷ and molecular magnetism.⁸ Use of relatively soft donor ligands, such as substituted cyclopentadienyls (Cp^R), can introduce interesting

bonding regimes, and their Ln/An complexes have been studied for many years.^{9–11} Such materials have found interest not just in chemical synthesis, but also due to their electronic structure leading to, for example, the stabilization of unprecedented metal oxidation states¹² and properties of single-molecule magnets.¹³ The archetypal examples are the [Ln/An(Cp)₃] type complexes, and indeed these have long been used as a computational test-bed to investigate trends in the nature of f-element bonding.^{14–18}

Since the majority of Ln³⁺ and An³⁺ ions are paramagnetic, electron paramagnetic resonance (EPR) spectroscopy should carry important information on metal–ligand bonding, particularly if resolution of ligand nuclear hyperfine interactions can be resolved. However, this is rarely the case in continuous wave (CW) EPR because of the intrinsically broad linewidths. This can in principle be addressed by pulsed EPR hyperfine methods.¹⁹ However, there are surprisingly few pulsed EPR studies on molecular Ln complexes,²⁰ with the notable exception of those of the ⁸S_{7/2} Gd³⁺ ion, and even fewer for An complexes.^{21–23} Most Ln³⁺ pulsed EPR studies have been performed on doped minerals or glasses, and have tended to focus on relaxation behaviour.²⁴ In 2011 Denning and co-workers reported pulsed EPR measurements on [Yb(Cp)₃], enabling quantification of the significant spin density at the ¹³C atoms of the Cp rings,²⁵ which was discussed in terms of mixing of the ²F_{7/2} ground term with low-lying charge-transfer states. More recently, some of us reported pulsed EPR studies of two An complexes, [An(Cp^{tt})₃] (1-An; An = Th, U; Cp^{tt} = C₅H₃^tBu₂-1,3),

^aDepartment of Chemistry, The University of Manchester, Oxford Road, Manchester, M13 9PL, UK. E-mail: floriana.tuna@manchester.ac.uk; nicholas.chilton@manchester.ac.uk; david.mills@manchester.ac.uk; eric.mcinnnes@manchester.ac.uk

^bPhoton Science Institute, The University of Manchester, Oxford Road, Manchester, M13 9PL, UK

† Electronic supplementary information (ESI) available. CCDC 2271676 and 2271677. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3sc06175b>

‡ These authors contributed equally.



again showing significant spin density on the Cp^{tt} ligands for both the U³⁺ and Th³⁺ analogues.²¹

Here we report studies, including pulsed EPR, on a family of early Ln³⁺ complexes [Ln(Cp^{tt})₃] (**1-Ln**, Ln = La, Ce, Nd, Sm). This allows: (i) investigation of trends in metal–arene interactions across the series, from the “parent” diamagnetic, f⁰ complex through the paramagnetic f^{1,3,5} analogues. The latter are chosen because they are all Kramers ions and hence are expected to be EPR-active; (ii) comparison of ligand spin density with the late Ln³⁺ complex [Yb(Cp)₃] studied by Denning *et al.*;²⁵ (iii) comparison of **1-Ln** with the early An³⁺ homologues **1-An**.²¹ The latter includes direct comparison of the valence isoelectronic Nd³⁺ (4f³) and U³⁺ (5f³) pair, which is the only comparison available between 4f and 5f M³⁺ ions that does not require designated radiochemical laboratories (although Ce³⁺ and Th³⁺ have the same number of valence electrons, the Th³⁺ ion often favours a 6d¹ rather than 5f¹ configuration).²⁶

Complexes **1-Ln** are readily prepared by reacting LnCl₃ with three equivalents of KCp^{tt}; **1-Ce**²⁷ and **1-Sm**²⁸ have been prepared previously by alternative synthetic routes, whilst **1-La** and **1-Nd** are structurally characterised here for the first time. We report the CW and pulsed EPR data for **1-Ce**, **1-Nd** and **1-Sm**, along with the NMR data of diamagnetic **1-La**. We quantify the hyperfine interaction in paramagnetic **1-Ln** with ligand nuclei, originating from interaction with the Ln³⁺ centres, and show that we can model these data using a simple point-dipole model. This is in agreement with fully-*ab initio* complete active space self-consistent field spin-orbit (CASSCF-SO) calculations that directly report hyperfine coupling parameters,²⁹ as well as spin densities. We find, using either the simple or *ab initio* methods, that there is negligible spin density on the Cp^{tt} ligands in **1-Ln**, in contrast to unequivocal spin delocalisation in U³⁺ and Th³⁺ analogues [An(Cp^{tt})₃],²¹ clearly highlighting the differences between early 4f and early 5f bonding in *tris*-Cp complexes.^{14–18} Since the ability to perform such studies are limited by electron spin relaxation times, we also report relaxation data for **1-Ln** by pulsed EPR methods. We find that 4f **1-Ln** relax orders of magnitude faster than their 5f An analogues,²¹ despite preconceptions that relaxation rates increase for heavier elements due to increased spin-orbit coupling.

Results and discussion

Synthesis

Complexes **1-Ln** (Ln = La, Ce, Nd, Sm) were prepared from the parent LnCl₃ and three equivalents of KCp^{tt} (Scheme 1) by modification of the reported syntheses of **1-Sm** using SmI₃ and



Scheme 1 Synthesis of **1-Ln** (Ln = La, Ce, Nd, Sm).

KCp^{tt},²⁸ and **1-Ce** from Ce(OTf)₃ and LiCp^{tt}.²⁷ The crystalline yields for **1-La**, **1-Ce**, **1-Nd** and **1-Sm** were 41%, 54%, 34% and 52%, respectively. We attempted to extend the **1-Ln** series to smaller Ln, but in our hands we found that applying the same synthetic methodology to GdCl₃ gave no isolable products, likely due to the higher Lewis acidity of the smaller Ln³⁺ ions.¹ NMR spectroscopy was used to analyse C₆D₆ solutions of **1-Ln** (ESI, Fig. S1–S5†). ¹H NMR spectra were fully assigned for all **1-Ln**; in each case three signals were observed in a ratio of 54 : 6 : 3 that correspond to the ^tBu groups and the two unique environments of the Cp^{tt} ring protons, respectively. The paramagnetism of **1-Ce**, **1-Nd** and **1-Sm** precluded assignment of their ¹³C{¹H} NMR spectra, however, for diamagnetic **1-La** this could be interpreted, with the two ^tBu group resonances seen at 32.77 and 33.75 ppm and the three Cp^{tt} ring carbon environments found at 110.57 (CH-Cp), 110.69 (CH-Cp) and 143.45 (C-Cp) ppm. Although NMR spectroscopy showed few protic impurities, elemental analysis results for **1-Ln** consistently gave low carbon values; this was previously seen for [An(Cp^{tt})₃],²¹ and is ascribed to carbide formation leading to partial combustion, a common issue with these experiments.³⁰ In the case of **1-Sm** elemental analysis results were not in accord with expected values. However, microcrystalline samples of **1-Ln** gave essentially superimposable ATR-IR spectra (ESI, Fig. S6–S10†), and when these samples were suspended in a minimum amount of fomblin they gave poorly resolved powder XRD patterns (Fig. S13–S16†) that are similar to each other and show peaks in common with predicted patterns from single crystal XRD data (see below). Together, the characterisation data obtained show the bulk purities of samples of **1-Ln**.

Crystallography

The solid-state structures of **1-Ln** were determined by single crystal XRD (**1-Nd** is depicted in Fig. 1, see ESI, Fig. S11 and S12† for selected distances and angles of **1-La** and **1-Nd**). The structural data of **1-La** and **1-Nd** are reported here for the first time, whilst **1-Ce**²⁷ and **1-Sm**²⁸ have been reported previously; although **1-Ln** are isostructural they are not all isomorphous. As expected, the structures of **1-Ln** are trigonal planar with respect to the η⁵-Cp^{tt} centroids, with the three C2 atoms (Fig. 1) in the plane defined by the Ln³⁺ ion and the three Cp^{tt} centroids. The three Cp^{tt} ligands adopt the same orientation to form a “picket-fence” motif with three ^tBu groups above and three below the trigonal plane. Complexes **1-Ln** do not show high symmetry in



Fig. 1 (a) Crystal structure of **1-Nd**; Nd = cyan, C = grey. Displacement ellipsoids are at the 30% probability level. Hydrogen atoms are omitted for clarity. (b) Schematic of C₃ symmetry of the structures of **1-Ln**.



the solid state, consistent with the solid-state structures of $[M(\text{Cp}^{\text{tt}})_3]$ ($M = \text{Th},^{21} \text{U},^{21} \text{Yb}^{31}$). The mean $\text{Ln}\cdots\text{Cp}_{\text{centroid}}$ distances decrease regularly across the lanthanide series [mean $\text{La}\cdots\text{Cp}_{\text{centroid}} = 2.635 \text{ \AA}$; $\text{Ce}\cdots\text{Cp}_{\text{centroid}} = 2.596 \text{ \AA}$; $^{27} \text{Nd}\cdots\text{Cp}_{\text{centroid}} = 2.56 \text{ \AA}$; $\text{Sm}\cdots\text{Cp}_{\text{centroid}} = 2.531 \text{ \AA}$; $^{28} \text{Yb}\cdots\text{Cp}_{\text{centroid}} = 2.47 \text{ \AA}$]. Additionally, the mean $\text{M}\cdots\text{Cp}_{\text{centroid}}$ distances in **1-Nd** are shorter than those seen in $[\text{Th}(\text{Cp}^{\text{tt}})_3]$ [2.570 \AA],²¹ which has an analogous electronic configuration ($\text{Nd}^{3+}: [\text{Xe}]4f^3; \text{U}^{3+}: [\text{Rn}]5f^3$). Conversely, in **1-Ce**, the mean $\text{M}\cdots\text{Cp}_{\text{centroid}}$ distances are longer than those seen in $[\text{Th}(\text{Cp}^{\text{tt}})_3]$ [2.566 \AA],²¹ which has a different valence electronic configuration ($\text{Ce}^{3+}: [\text{Xe}]4f^1; \text{Th}^{3+}: [\text{Rn}]6d^1$) due to the near-degeneracy of the 5f and 6d orbitals for Th^2 and the stabilization of the $6d_{z^2}$ orbital in trigonal ligand environments.^{26,32}

Solution phase optical properties

The electronic spectra of 0.5 mM toluene solutions of **1-Ln** (Fig. 2 and ESI, Fig. S17–S21†) are as expected;¹ **1-La** with a closed shell La^{3+} ion is essentially colourless, with charge-transfer (CT) bands restricted to the UV region, whilst **1-Ce**, **1-Nd** and **1-Sm** are pale purple, pale green, and pale orange, respectively. Due to their Laporte-forbidden nature, even spin-allowed f-f transitions are relatively weak ($\epsilon < 400 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1}$),¹ so CT bands tailing in from the UV region dominate the spectra. The f-f transitions in **1-Nd** were clearly observed, with the most intense absorption at $\tilde{\nu}_{\text{max}} = 16750 \text{ cm}^{-1}$ ($\epsilon = 390 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1}$) likely arising from the $^4\text{I}_{9/2} \rightarrow ^4\text{G}_{5/2}/^4\text{G}_{7/2}$ transition.³³ In contrast, **1-Ce** shows a strong broad absorption at $\tilde{\nu}_{\text{max}} = 17400 \text{ cm}^{-1}$ ($\epsilon = 230 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1}$) due to $[\text{Xe}]4f^1 \rightarrow [\text{Xe}]4f^05d^1$ transitions, which are formally allowed by electric dipole selection rules. The electronic spectrum of **1-Sm** shows a weak set of f-f transitions in the near-IR region around 7000 cm^{-1} ($\epsilon < 50 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1}$), likely arising from the $^6\text{H}_{5/2} \rightarrow ^6\text{F}_J$ transitions.³³

Magnetism

The measured χT values of 1.45 and $0.75 \text{ cm}^3 \text{ mol}^{-1} \text{ K}$ at 300 K for **1-Nd** and **1-Ce**, respectively (Fig. 3), are close to the expected values of 1.63 and $0.80 \text{ cm}^3 \text{ mol}^{-1} \text{ K}$ for Nd^{3+} ($^4\text{I}_{9/2}$) and Ce^{3+} ($^2\text{F}_{5/2}$).¹ The χT product decreases gradually upon cooling until

below 50 or 60 K , where it drops to reach 0.44 and $0.30 \text{ cm}^3 \text{ mol}^{-1} \text{ K}$ at 1.8 K , respectively. For **1-Sm**, χT is $0.20 \text{ cm}^3 \text{ mol}^{-1} \text{ K}$ at 300 K , which is larger than expected for an isolated free-ion $^6\text{H}_{5/2}$ term of $0.09 \text{ cm}^3 \text{ mol}^{-1} \text{ K}$, owing to a combination of thermal population of the low-lying $^6\text{H}_{7/2}$ term and temperature independent paramagnetism. The decrease of χT in all cases is due to the depopulation of the crystal field states. Isothermal magnetisation measurements of all compounds deviate from values expected for pure Ising-like m_j states (ESI, Fig. S22–S24†),³⁴ indicating significant mixing of m_j states. To corroborate these results, we performed CASSCF-SO calculations using a minimal n -electrons in 7 orbital active space (where $n = 1$ for Ce^{3+} , $n = 3$ for Nd^{3+} and $n = 5$ for Sm^{3+} ; see ESI† for details) and the solid-state XRD geometries (Tables S10–S13†). These give excellent reproductions of the magnetic data (Fig. 3 and S22–S24†) and confirm substantially mixed m_j states (Tables S10–S13†); the ground Kramers doublet for **1-Ce** is dominated by $|m_j| = 1/2$, with the first excited state at 105 – 135 cm^{-1} , while for **1-Nd** the ground Kramers doublet is dominated by $m_j = \pm 5/2$ mixed with $\mp 7/2$, with a first excited state at 26 cm^{-1} . For **1-Sm** the ground Kramers doublet is dominated by $m_j = \pm 3/2$, with a first excited state at 400 cm^{-1} .

EPR spectroscopy

CW EPR spectra of **1-Nd**, **1-Ce** and **1-Sm** are observed at temperatures below $\sim 30 \text{ K}$ (Fig. 4 and ESI, Fig. S25–S27†). Much sharper spectra are observed from powders (such that it is difficult to remove polycrystalline effects) than from frozen solutions (toluene/hexane $9:1 \text{ v/v}$), indicating some relaxation and strain of the structures in the latter medium. The spectra are dominated in each case by the rhombic or axial effective g -values arising from transitions within a thermally isolated ground Kramers doublet of the ground Russell-Saunders state, which can be treated as an effective spin-1/2. In this approximation, the frozen solution EPR spectrum for **1-Nd** (Fig. 4a) can be modeled with the spin Hamiltonian eqn (1).



Fig. 2 UV/vis/NIR spectra of **1-La**, **1-Ce**, **1-Nd**, **1-Sm**, from 6000 – 30000 cm^{-1} (inset 6000 – 20000 cm^{-1}) recorded as 0.5 mM solutions in toluene at room temperature.



Fig. 3 Temperature dependence of χT for **1-Nd** (green squares), **1-Ce** (blue crosses), and **1-Sm** (red circles) recorded at 0.1 , 0.1 and 0.5 T . Calculated temperature dependence of χT using CASSCF-SO is given with black lines using the XRD structure for each molecule; two traces for **1-Ce** arise from each of the non-equivalent molecules in the crystal structure.





Fig. 4 X-Band CW EPR spectra, frozen toluene : hexane (9 : 1) solutions of (a) **1-Nd** at 8 K, and (b) **1-Ce** at 10 K. The resonator cavity is marked with (*), and the simulation parameters are provided in the text and Table S2.

$$\hat{\mathcal{H}} = \mu_B B \cdot \vec{g} \cdot \hat{S} + \hat{I} \cdot \vec{A} \cdot \hat{S} \quad (1)$$

This gives $g_{x,y,z} = 3.33, 1.22, 0.56$, and hyperfine coupling constants of $A_x = 1280$ MHz (A_y and A_z unresolved; we have assumed \vec{g} and \vec{A} are collinear) to the $I = 7/2$ nuclear spin of Nd (natural abundance: 12.2% ^{143}Nd and 8.3% ^{145}Nd). Similarly, frozen solution spectra of **1-Ce** and **1-Sm** can be simulated with $g_{x,y,z} = 3.15, 1.88, 0.64$ for **1-Ce** (Fig. 4b; note there are no isotopes of Ce with $I \neq 0$), and $g_x = g_y = 0.92$ for **1-Sm** (Fig. S27†). The g_z component of **1-Sm** is not well resolved in CW spectra but is more clearly defined in pulsed spectra giving $g_z = 0.45$ (Fig. S28;† see below). CW spectra of the related $[\text{Nd}(\text{Cp}'')_3]$ ($\text{Cp}'' = \text{C}_5\text{H}_3(\text{SiMe}_3)_2-1,3$) have been reported,³⁵ and although the precise g -values differ the pattern is similar.

For **1-Ce** and **1-Nd**, CASSCF-SO calculations (Tables S10–S13; † note there are two non-equivalent molecules in the XRD structure of **1-Ce**) give effective g -values for the lowest Kramers doublets as $g_{x,y,z} = 2.1, 3.0, 0.7$ and $2.2, 2.4, 1.1$, respectively, with the numerically smallest value (g_z) corresponding to the orientation of the pseudo- C_3 axis (Fig. 1); the anisotropy patterns agree with the experiments. For **1-Sm**, CASSCF-SO calculations give $g_{x,y,z} = 0.4, 0.4, 0.6$ for the lowest Kramers doublet. This has the opposite sense of anisotropy ($g_{x,y} < g_z$) to the experimental data ($g_{x,y} > g_z$), but given the numerically similar and very low (all < 1) effective g -values, such a switch in anisotropy will be very sensitive to subtle state mixing effects. Nevertheless, given the axially symmetric form of the EPR spectrum, g_z must be associated with the pseudo- C_3 axis.

Pulsed EPR spectroscopy

Echo-detected field-swept spectra (EDFS) of **1-Nd**, **1-Ce** and **1-Sm** (in toluene-hexane) are observed up to 10 K (see ESI† for all pulsed EPR spectra). These were recorded by integration of the Hahn echo generated with the standard pulse sequence $\pi/2-\tau-\pi$ -echo, where $\pi/2$ and π are microwave pulses and τ is the inter-pulse delay.¹⁹ The EDFS spectra at 5 K for **1-Nd**, **1-Ce** and **1-Sm** (Fig. 5, S28 and S29†) are consistent with the CW spectra, Q-band EDFS are consistent with the rhombic g -values of **1-Nd** and **1-Ce** (Fig. S30†).

Measurements of the spin-lattice (T_1) and phase-memory (T_m) relaxation times were performed at X-band on 10 mM frozen solution samples of **1-Nd**, **1-Ce** and **1-Sm** at 5 K. T_1 data were recorded with an inversion recovery pulse sequence, and the data fitted with a bi-exponential function (see ESI eqn (3);† Fig. S34 and S35†), where the fastest process is assigned to the spectral diffusion and the slow process is attributed to the spin-lattice relaxation.³⁶ The T_1 value for **1-Nd** is field-dependent and reaches a maximum value of 12 μs at the highest field g -value (12 360 G, Table S3†). Measured at the maximum of the EDFS (6040 G), T_1 is 5 μs . Much longer maximum T_1 times of 89 and 150 μs were obtained for **1-Ce** (at the high-field g -value, 11 228 G) and **1-Sm** (at the low-field g -value, 6936 G), respectively (Tables S4 and S5†). For comparison, T_1 measured at the EDFS maxima are 78 and 118 μs for **1-Ce** and **1-Sm**, respectively. We note that the measured T_1 data are anisotropic (Fig. S34 and S35, Tables S3–S5†), and the anisotropy (evaluated as the ratio of $T_{1,\parallel}/T_{1,\perp}$)³⁷ is most significant for **1-Nd** which also has the greatest g -anisotropy.

T_m was determined by fitting Hahn echo decays to a stretched exponential function (see ESI eqn (1); Fig. S31–S33†).³⁸ The maximum T_m relaxation times observed are 0.7, 1.0 and 1.7 μs for **1-Nd**, **1-Ce** and **1-Sm**, respectively (Tables S3–S5†), in each case measured at the highest field g -value. T_m measured at the EDFS maxima are 0.5, 0.6 and 1.6 μs , respectively.

An interesting comparison can be made between **1-Nd** and its valence isoelectronic An^{3+} analog **1-U**. Complex **1-U** has T_1 and T_m times of 860 and 0.8 μs , measured at the EDFS



Fig. 5 X-band (9.7 GHz) EDFS spectra of (a) **1-Nd** and (b) **1-Ce**, measured at 5 K from toluene : hexane (9 : 1) solution. Arrows indicate the observer positions at which T_1 (blue), T_m (blue) and Hyscore (red) were measured.



maximum under similar conditions,²¹ compared to 5 and 0.5 μs for **1-Nd**. Hence, T_1 for the $\text{Ln}^{3+} 4f^3$ complex is more than two orders of magnitude shorter than for its $\text{An}^{3+} 5f^3$ analogue. In general, electron spin-lattice relaxation times are expected to decrease going down the Periodic table, as modulation of the electronic structure by vibrational modes (which is a dominant factor for T_1 relaxation in immobilised samples²⁴) impacts the orbital angular momentum. This is connected to the spin *via* spin-orbit coupling (SOC), and SOC increases with atomic number;³⁹ the values for Nd^{3+} and U^{3+} are 900 and *ca.* 1700 cm^{-1} , respectively.⁴⁰ It seems reasonable to assume that the slower T_1 relaxation of **1-U** is due to the partial quenching of the orbital angular momentum by the larger crystal field interaction for U^{3+} *cf.* Nd^{3+} . This is promising for future pulsed EPR studies of An-containing materials. The $4f^1$ complex **1-Ce** also has a much shorter T_1 than its An^{3+} analogue **1-Th** (T_1 *ca.* 21 ms under similar conditions):²¹ this is due to the orbital singlet $6d^1$ configuration of **1-Th**,²⁶ compared to $4f^1$ for **1-Ce**.

Comparing the relaxation data for the **1-Ln** series, it could be inferred that **1-Nd** has the shortest T_1 because it has the largest ground state orbital angular momentum ($L = 6$) and indeed the largest total angular momentum ($J = 9/2$). However, comparison with T_1 data for these Ln^{3+} ions at similar concentrations and temperatures in water/ethanol glasses gives a different trend, with Nd^{3+} having the longest T_1 .⁴¹ This emphasises the importance of the crystal field on the relaxation behaviour of Ln^{3+} ions. In our case, of the three complexes studied, **1-Nd** has, by some margin, the lowest energy excited state and indeed the smallest energy spread of m_j states despite having the largest multiplicity (Tables S10–S13†). More densely packed electronic states renders the states more sensitive to perturbations (following textbook perturbation theory arguments), and so we can speculate that this exposes **1-Nd** to more influences from spin-vibration coupling, and thus causes a shorter T_1 .

The T_m values for **1-Ln** complexes studied are all similar, around the 1 μs mark. These are ample to allow further investigation of the complexes by multi-pulse microwave sequences. In order to quantify the weak hyperfine interactions between the electron spin(s) and surrounding ^1H and ^{13}C nuclei, we employed two-dimensional hyperfine sub-level correlation (HYSCORE) spectroscopy, which uses a four-pulse spin echo envelope modulation sequence; $\pi/2 - \tau - \pi/2 - t_1 - \pi - t_2 - \pi/2 - \text{echo}$, with t_1 and t_2 independently varied.⁴² In the HYSCORE experiment the first two $\pi/2$ pulses generate nuclear coherences, which are then transferred between electron spin states by the π pulse. In the 2D frequency domain spectrum, cross-peaks appear for weakly-coupled nuclei ($2|\nu_n| > |A|$) in the (+, +) quadrant straddling the nuclear Larmor frequencies (ν_n). Ridges in the spectra are due to the anisotropic hyperfine couplings.⁴³

X-band HYSCORE spectra measured at static magnetic fields corresponding to orientations in the molecular *xy* plane for **1-Nd** and **1-Ce** ($B_0 = 353.0$ and 348.2 mT, respectively) reveal signals from ^{13}C and ^1H nuclei (Fig. 6). HYSCORE signals for **1-Sm** were too weak to detect when measured under similar conditions; this does not appear to be relaxation limited given the similar T_m (see above), and may be a function of the very low effective *g*-values.²⁴ The ^{13}C features for **1-Nd** and **1-Ce** are

similar, consisting of a ridge extending beyond the ^{13}C Larmor frequency by *ca.* ± 1 MHz. The ^1H features are also similar, with a ridge extending $\nu_n \pm 2.5$ MHz.

In a first attempt to model the HYSCORE data of **1-Ce** and **1-Nd**, we used a simple point dipolar approach assuming the electron spin density to be located at the Ln^{3+} ion, calculated (ESI, eqn (4)†) using the experimental *g*-values and the atomic coordinates from XRD (thus neglecting structural relaxation in solution). The calculated point dipolar hyperfine values for the Cp-ring hydrogen and carbon atoms (Tables S8 and S9†) were then used with EasySpin^{44,45} to calculate HYSCORE spectra. The calculated spectra show excellent agreement with the experimental data (Fig. 6).

Comparing the HYSCORE data of $4f$ **1-Ce** and **1-Nd** to their $5f$ analogues **1-An**, ^{13}C HYSCORE spectra measured for **1-Th** under equivalent conditions have ridges that spread $\nu_n \pm 2.4$ MHz,²¹ substantially larger than for **1-Ce**. We did not have sufficient signal to measure the full ^{13}C hyperfine pattern for **1-U**, but we did measure well-resolved ^1H ridges in the $g_{x,y}$ region:²¹ these spanned $\nu_n \pm 2.5$ MHz, which is much larger than for **1-Nd**. The spectra for **1-Th** and **1-U** could not be modelled on a simple point dipole basis, requiring significant additional contributions to the hyperfine interactions.

We further confirmed these results using CASSCF-SO calculations (see ESI†). To compare with the HYSCORE experiment, we approximate solution phase structures by optimising the molecular geometry of each complex in the gas-phase using density-functional theory (DFT) methods (see ESI† for details), and then use the optimised geometries to perform another set of minimal CASSCF-SO calculations; note that these calculations explicitly include CF, SO and interelectronic repulsion



Fig. 6 X-Band HYSCORE spectra for **1-Nd** (a and b; static field $B_0 = 353.0$ mT) and **1-Ce** (c and d; $B_0 = 348.2$ mT) at 5 K. (a and c) ^{13}C region (blue) for **1-Nd** and **1-Ce**, respectively, and calculated spectra (red) based on a dipolar model including C1–C5 (Tables S8 and S9†); (b and d) ^1H region (blue) for **1-Nd** and **1-Ce**, respectively, and calculated spectra (red) based on a dipolar model including H2, H4 and H5 (Tables S8 and S9†). The red dashed anti-diagonal lines indicate the nuclear Larmor frequency for ^{13}C (a and c) or ^1H (b and d).



effects of the f-shell ground configuration. We then use the Hyperion package to calculate the relativistic hyperfine coupling tensors directly for the nuclei of interest (Tables S14 and S15[†]);²⁹ note that the Hyperion method implicitly includes all through space dipolar and Fermi contact terms, as well as relativistic paramagnetic spin-orbit terms. Inspection of the computed hyperfine values show the C(Cp) and H(Cp) atoms couple more strongly than any other Cp^{tt} ligand atoms. Hence, we simulate ¹³C and ¹H HYSCORE spectra using five C and three H nuclei from one Cp^{tt} ligand, again using EasySpin; these calculations provide excellent reproductions of the experimental data (Fig. S36[†]). The calculated Mulliken spin populations (Table S16[†]), show that only ~0.1% of the total spin density is transferred to the C1–C5 atoms of the three Cp^{tt} ligands. However, we note that the density does increase slightly with increasing Ln³⁺ atomic number (**1-Ce** < **1-Nd** < **1-Sm**).

Conclusions

We have synthesised a family of Ln³⁺ complexes [Ln(Cp^{tt})₃] (**1-Ln**; Ln = La, Ce, Nd, Sm) using a salt metathesis route. The solid-state structures of **1-Ln** reveal that the distances between the Ln³⁺ centres and the Cp^{tt} centroids decrease regularly from La³⁺ to Sm³⁺ due to the ionic radii of Ln³⁺ ions decreasing across the series. Complexes **1-La** and **1-Nd** exhibit *pseudo*-trigonal planar geometries, which is consistent with the previously reported structures of **1-Ce**,²⁷ **1-Sm**²⁸ and **1-Yb**.³¹ Continuous wave and pulsed EPR studies were performed on **1-Nd**, **1-Ce**, and **1-Sm**. The CW spectra show rhombic systems with anisotropic *g*-values for both frozen solution and powder samples for **1-Nd** and **1-Ce**. HYSCORE spectroscopy shows resonances for ¹³C and ¹H regions for both **1-Nd** and **1-Ce**, and point-dipole simulations provide excellent agreement with experimental data. CASSCF-SO calculations effectively reproduce magnetic data, and fully-*ab initio* simulations of the HYSCORE spectra are also in excellent agreement with experiment, confirming minimal spin density at ligand nuclei. The larger ligand hyperfine interactions observed for **1-Th** (ref. 21) *cf.* **1-Ce** are consistent with the 6d¹ vs. 4f¹ configurations of Th³⁺ and Ce³⁺, respectively, and the significant radial extent of the 6d orbital(s). Comparing the results for **1-Nd** with **1-U** shows larger ligand hyperfines observed for the latter, highlighting the difference between 4f and 5f valence orbitals.

The results show that the interactions between Ln³⁺ ions and these cyclopentadienyl ligands in these early Ln organometallic complexes **1-Ln** can be considered as being largely ionic in the sense of minimal transfer of spin density from the 4f to ligand orbitals. The results are consistent with early CW EPR studies of U³⁺ and Nd³⁺ doped fluorite⁴⁶ (and later studies on other inorganic lattices),⁴⁷ where superhyperfine coupling to ¹⁹F was observed for the former but not the latter. These are mineral lattices with hard ligands, and we have now shown related effects in molecular species with soft arene ligand sets. The results on **1-Ln** also contrast with HYSCORE studies of the closely related [Yb(Cp)₃] complex,²⁵ a late Ln complex (4f¹³), which showed far more significant transfer of spin density from 4f to arene and can hence be described as being “more covalent”

in this sense, despite the smaller ionic radius of the Yb³⁺ ion. This is consistent with Denning's conclusion²⁵ that this “increased covalency” is due to the low reduction potential of Yb³⁺ combined with an electron-rich ligand leading to greater metal-to-ligand charge transfer character in the ground state. This highlights the importance of ligand substituents and Ln ion charge density in controlling the magnitude of 4f metal-ligand interactions, and shows that pulsed EPR spectroscopy is a sensitive probe to study these effects.

Experimental

Materials and methods

General comments. All manipulations were carried out using standard Schlenk line and glove box techniques under dry argon. Solvents were passed through columns containing alumina or were dried by refluxing over K, and were stored over K mirrors or 4 Å molecular sieves (THF) and degassed before use. For NMR spectroscopy C₆D₆ was dried by refluxing over K, degassed by three freeze-pump-thaw cycles, and vacuum-transferred before use. Anhydrous LnCl₃ were purchased from Alfa Aesar and were used as received. KCp^{tt} was synthesised by literature methods,⁴⁸ whilst **1-Ln** were prepared by modification of literature procedures.^{27,28} General synthetic procedures for **1-Ln** are given below; full details can be found in the ESI.[†] ¹H (400 and 500 MHz) and ¹³C{¹H} (100 and 125 MHz) NMR spectra were obtained on Avance III 400 or 500 MHz spectrometers at 298 K. ¹H NMR spectra were measured from 0 to +10 ppm for diamagnetic **1-La** and from –200 to +200 ppm for paramagnetic **1-Ce**, **1-Nd** and **1-Sm**. ATR-IR spectra were recorded as microcrystalline powders using a Bruker Tensor 27 spectrometer. Elemental analyses were performed by Mrs Anne Davies and Mr Martin Jennings at The University of Manchester School of Chemistry Microanalysis Service, Manchester, UK. UV/Vis/NIR spectroscopy was performed on samples in Youngs tap-appended 10 mm path length quartz cuvettes on an Agilent Technologies Cary Series UV/Vis/NIR spectrophotometer at 175–3300 nm. CW EPR measurements were carried out on a Bruker EMX300 spectrometer; pulsed EPR X-band studies were performed on a Bruker ElexSys E580 spectrometer. The primary Hahn-echo sequence ($\pi/2-\tau-\pi-\tau$ -echo) was used for the two-pulse electron spin echo measurements, with initial $\pi/2$ and π pulse of 16 and 32 ns, respectively. For the relaxation time measurements, *T*_m studies were made by incrementing the τ time in the Hahn-echo sequence (longer pulses were used to suppress the ¹H modulation), *T*₁ was measured by the inversion recovery sequence ($\pi-t-\pi/2-t_1-\pi-t_2-\pi/2$ -echo) $\pi/2$ and π pulse of 16 and 32 ns, respectively, with a fixed $\tau = 300$ ns. HYSCORE measurements were performed using the four-pulse sequence ($\pi/2-\tau-\pi/2-t_1-\pi-t_2-\pi/2$ -echo), $\pi/2$ and π pulse of 16 and 32 ns, respectively, initial times *t*_{1,2} = 0.1 μs and τ values of 136 and 200 ns.

General synthesis of 1-Ln. THF (30 mL) was added to a pre-cooled (–78 °C) ampoule containing LnCl₃ (2 mmol) and KCp^{tt} (6 mmol). The reaction mixture was allowed to reflux for 16 hours. The solvent was removed *in vacuo* and toluene (30 mL)



was added. The reaction mixture was allowed to reflux for 40 hours. The resultant suspension was allowed to settle for 3 hours and filtered. The solution was concentrated to *ca.* 2 mL and stored at 8 °C to afford crystals of **1-Ln**.

1-La: colourless crystals (0.550 g, 41%). Anal calcd (%) for C₃₉H₆₃La: C, 69.81; H, 9.47. Found (%): C, 67.03; H, 9.50. ¹H NMR (C₆D₆, 400 MHz, 298 K): δ = 1.35 (s, 54H, C(CH₃)₃), 6.21 (s, 3H, Cp-H), 6.28 (s, 6H, Cp-H) ppm. ¹³C{¹H} NMR (C₆D₆, 100 MHz, 298 K): δ = 32.77 (C(CH₃)₃), 33.75 (C(CH₃)₃), 110.57 (CH-Cp ring), 110.69 (CH-Cp ring), 143.45 (C-Cp ring) ppm. FTIR (ATR, microcrystalline): $\tilde{\nu}$ = 2960 (s), 2899 (w), 2862 (w), 1459 (m), 1388 (w), 1356 (m), 1252 (s), 1198 (w), 1163 (w), 1088 (br, s), 1018 (s), 927 (w), 803 (s), 736 (s), 660 (w), 605 (w) cm⁻¹.

1-Ce: blue crystals (0.726 g, 54%). Anal calcd (%) for C₃₉H₆₃Ce: C, 69.68; H, 9.45. Found (%): C, 67.49; H, 9.43. ¹H NMR (C₆D₆, 500 MHz, 298 K): δ = -5.01 (s, 54H, C(CH₃)₃), 17.14 (s, 6H, Cp-H), 26.30 (s, 3H, Cp-H) ppm. The paramagnetism of **1-Ce** precluded assignment of its ¹³C{¹H} NMR spectrum. FTIR (ATR, microcrystalline): $\tilde{\nu}$ = 2951 (br, s), 2899 (w), 2863 (w), 1459 (s), 1388 (m), 1356 (s), 1298 (w), 1251 (s), 1198 (m), 1164 (s), 1051 (m), 1021 (m), 927 (s), 806 (s), 738 (s), 674 (s), 659 (s), 604 (w), 556 (br, w), 480 (w), 422 (m) cm⁻¹.

1-Nd: green crystals (0.460 g, 34%). Anal calcd (%) for C₃₉H₆₃Nd: C, 69.26; H, 9.40. Found (%): C, 65.81; H, 9.30. ¹H NMR (C₆D₆, 400 MHz, 298 K): δ = -9.06 (s, 54H, C(CH₃)₃), 12.68 (s, 6H, Cp-H), 34.47 (s, 3H, Cp-H). The paramagnetism of **1-Nd** precluded assignment of its ¹³C{¹H} NMR spectrum. FTIR (ATR, microcrystalline): $\tilde{\nu}$ = 2950 (s), 2899 (w), 2863 (w), 1459 (s), 1388 (w), 1356 (s), 1251 (s), 1164 (m), 1060 (br, w), 1021 (w), 927 (s), 806 (s), 737 (s), 659 (s), 605 (w), 423 (w) cm⁻¹.

1-Sm: orange crystals (0.716 g, 52%). ¹H NMR (C₆D₆, 400 MHz, 298 K): δ = -1.58 (s, 54H, C(CH₃)₃), 18.66 (s, 6H, Cp-H), 21.19 (s, 3H, Cp-H). The paramagnetism of **1-Sm** precluded assignment of its ¹³C{¹H} NMR spectrum. FTIR (ATR, microcrystalline): $\tilde{\nu}$ = 2952 (s), 2901 (m), 2864 (w), 1460 (s), 1390 (s), 1366 (s), 1298 (s), 1250 (s), 1199 (s), 1164 (s), 1085 (w), 1060 (w), 1022 (m), 927 (s), 807 (s), 741 (s), 700 (s), 661 (s), 606 (m), 560 (m), 519 (w), 483 (w), 426 (s) cm⁻¹.

Data availability

Additional experimental details for physical and computational data associated with this manuscript (PDF). Additional research data supporting this publication are available from Figshare at <https://figshare.com/10.6084/m9.figshare.23302253>.

Author contributions

F. T., E. J. L. M. and D. P. M. provided the original concept. L. N. and A.-M. A. collected and modelled EPR data. J. L., F. O., P. J. C. and J. E.-K. synthesized and characterized the complexes and collected, solved and refined the single crystal and powder XRD data. F. O. further refined the single crystal XRD data and finalized cifs. L. N. and G. K. G. collected and analyzed magnetic data. M. S. O. and L. B. performed calculations. N. F. C. supervised the calculations, D. P. M. supervised the synthesis and characterization of compounds, and F. T. and E. J. L. M.

supervised EPR spectroscopy. L. N., J. L., N. F. C., D. P. M., E. J. L. M. and F. T. wrote the manuscript, with contributions from all other authors.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the Engineering and Physical Sciences Research Council (studentship to L. N. and EP/P002560/1), the China Scholarship Council (studentship to J. L.), The European Commission Marie Curie Initial Training Network 'Magic' (studentship to A. M. A), the Ministry of Defence (studentship to L. B.), the Royal Society (fellowship URF191320 to N. F. C.), Natural Sciences and Engineering Research Council of Canada (fellowship 557458 to M. S. O.), the European Research Council (StG-851504 to N. F. C., CoG-816268 to D. P. M.) The University of Manchester Single Crystal X-ray Diffraction Service (EP/P001386/1), The University of Manchester Computational Shared Facility for computational resources, and the EPSRC UK EPR National Research Facility (EP/W014521/1, EP/V035231/1) for supporting this work.

References

- 1 *The Rare Earth Elements: Fundamentals and Applications*, ed. D. A. Atwood, John Wiley and Sons Ltd., Chichester, 2012.
- 2 *The Chemistry of the Actinide and Transactinide Elements*, ed. L. R. Morss, N. M. Edelstein and J. Fuger, Springer, Dordrecht, 3rd edn, 2006.
- 3 M. L. Neidig, D. L. Clark and R. L. Martin, *Coord. Chem. Rev.*, 2013, **257**, 394–406.
- 4 H. H. Dam, D. N. Reinhoudt and W. Verboom, *Chem. Soc. Rev.*, 2007, **36**, 367–377.
- 5 K. L. Nash, *Solvent Extr. Ion Exch.*, 1993, **11**, 729–768.
- 6 L. Ungur, B. Szabo, Z. A. AlOthman, A. A. S. Al-Khatani and L. F. Chibotaru, *Inorg. Chem.*, 2022, **61**, 5972–5976.
- 7 D. Parker, E. A. Suturina, I. Kuprov and N. F. Chilton, *Acc. Chem. Res.*, 2020, **53**, 1520–1534.
- 8 M. Brigante, G. F. Garcia, J. Jung, R. Sessoli, B. Le Guennic and F. Totti, *Chem. Sci.*, 2019, **10**, 7233–7245.
- 9 H. Schumann, J. A. Meese-Marktscheffel and L. Esser, *Chem. Rev.*, 1995, **95**, 865–986.
- 10 O. Walter, *Chem.-Eur. J.*, 2019, **25**, 2927–2934.
- 11 M. Ephritikhine, *Organometallics*, 2013, **32**, 2464–2488.
- 12 W. J. Evans, *Organometallics*, 2016, **35**, 3088–3100.
- 13 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.
- 14 B. E. Bursten, L. F. Rhodes and R. J. Strittmatter, *J. Am. Chem. Soc.*, 1989, **111**, 2756–2758.
- 15 R. J. Strittmatter and B. E. Bursten, *J. Am. Chem. Soc.*, 1991, **113**, 552–559.



- 16 N. Kaltsoyannis and B. E. Bursten, *J. Organomet. Chem.*, 1997, **528**, 19–33.
- 17 I. Kirker and N. Kaltsoyannis, *Dalton Trans.*, 2011, **40**, 124–131.
- 18 N. Kaltsoyannis, *Inorg. Chem.*, 2013, **52**, 3407–3413.
- 19 A. Schweiger and G. Jeschke, *Principles of Pulse Electron Paramagnetic Resonance*, Oxford University Press, Oxford, 2001.
- 20 For recent examples, see: (a) M. Shiddiq, D. Komijnai, Y. Duan, A. Gaita-Ariño, E. Coronado and S. Hill, *Nature*, 2016, **531**, 348–351; (b) K. Kundu, J. R. K. White, S. A. Moehring, J. M. Yu, J. W. Ziller, F. Furche, W. J. Evans and S. Hill, *Nat. Chem.*, 2022, **14**, 392–397; (c) A.-M. Ariciu, D. H. Woen, D. N. Huh, L. E. Nodaraki, A. K. Kostopoulos, C. A. P. Goodwin, N. F. Chilton, E. J. L. McInnes, R. E. P. Winpenny, W. J. Evans and F. Tuna, *Nat. Commun.*, 2019, **10**, 3330; (d) K. S. Pedersen, A.-M. Ariciu, S. McAdams, H. Weihe, J. Bendix, F. Tuna and S. Piligkos, *J. Am. Chem. Soc.*, 2016, **138**, 5801–5804.
- 21 A. Formanuik, A.-M. Ariciu, F. Ortu, R. Beekmeyer, A. Kerridge, F. Tuna, E. J. L. McInnes and D. P. Mills, *Nat. Chem.*, 2017, **9**, 578–583.
- 22 G. Rao, A. B. Altman, A. C. Brown, L. Z. Tao, T. A. Stich, J. Arnold and R. D. Britt, *Inorg. Chem.*, 2019, **58**, 7978–7988.
- 23 L. Barluzzi, N. Jori, T. He, T. Rajeshkumar, R. Scopelliti, L. Maron, P. Oyala, T. Agapie and M. Mazzanti, *Chem. Commun.*, 2022, **58**, 4655–4658.
- 24 For a recent review, see: J. E. McPeak, S. S. Eaton and G. R. Eaton, *Methods Enzymol.*, 2021, **651**, 63–101.
- 25 R. G. Denning, J. Harmer, J. C. Green and M. Irwin, *J. Am. Chem. Soc.*, 2011, **133**, 20644–20660.
- 26 W. K. Kot, G. V. Shalimov, N. M. Edelstein, M. A. Edelman and M. F. Lappert, *J. Am. Chem. Soc.*, 1988, **110**, 986–987.
- 27 C. D. Sofield and R. A. Andersen, *J. Organomet. Chem.*, 1995, **501**, 271–276.
- 28 N. A. Pushkarevsky, I. Y. Ilyin, P. A. Petrov, D. G. Samsonenko, M. R. Ryzhikov, P. W. Roesky and S. N. Konchenko, *Organometallics*, 2017, **36**, 1287–1295.
- 29 L. Birnoschi and N. F. Chilton, *J. Chem. Theory Comput.*, 2022, **18**, 4719–4732.
- 30 R. E. H. Kuveke, L. Barwise, Y. van Ingen, K. Vashisth, N. Roberts, S. S. Chitnis, J. L. Dutton, C. D. Martin and R. L. Melen, *ACS Cent. Sci.*, 2022, **8**, 855–863.
- 31 A. V. Khvostov, B. M. Bulychev, V. K. Belsky and A. I. Sizov, *J. Organomet. Chem.*, 1998, **568**, 113–119.
- 32 F. Ortu, A. Formanuik, J. R. Innes and D. P. Mills, *Dalton Trans.*, 2016, **45**, 7537–7549.
- 33 V. A. Volkovich, A. B. Ivanov, S. M. Yakimov, D. V. Tsarevskii, O. A. Golovanova, V. V. Sukhikh and T. R. Griffiths, *AIP Conf. Proc.*, 2016, **1767**, 020023.
- 34 J.-L. Liu, Y.-C. Chen and M. L. Tong, *Chem. Soc. Rev.*, 2018, **47**, 2431–2453.
- 35 W. W. Lukens, M. Speldrich, P. Yang, T. J. Duignan, J. Autschbach and P. Kögerler, *Dalton Trans.*, 2016, **45**, 11508–11521.
- 36 S. S. Eaton and G. R. Eaton, in *Biological Magnetic Resonance Volume 19: Distance Measurements in Biological Systems by EPR*, ed. L. J. Berliner, G. R. Eaton and S. S. Eaton, Kluwer Academic/Plenum Publishers, New York, 2002, pp. 29–154.
- 37 N. P. Kazmierczak and R. G. Hadt, *J. Am. Chem. Soc.*, 2022, **144**, 20804–20814.
- 38 K. Bader, M. Winkler and J. van Slageren, *Chem. Commun.*, 2016, **52**, 3623–3626.
- 39 R. Husted, J.-L. Du, G. R. Eaton and S. S. Eaton, *Magn. Reson. Chem.*, 1995, **33**, S66–S69.
- 40 A. Abragam and B. Bleaney, *Electron Paramagnetic Resonance of Transition Ions*, Oxford University Press, Oxford, 1970.
- 41 S. S. Eaton and G. R. Eaton, *J. Magn. Reson.*, 2023, **348**, 107392.
- 42 S. Van Doorsleer, *eMagRes*, 2017, **6**, 51–69.
- 43 J. Harmer, G. Mitrikas and A. Schweiger, *Advanced Pulse EPR Methods for the Characterization of Metalloproteins*, In *High Resolution EPR*, ed. G. Hanson and L. Berliner, Springer Science, New York, 2009, vol. 28, pp. 13–61.
- 44 S. Stoll and A. Schweiger, *J. Magn. Reson.*, 2006, **178**, 42–55.
- 45 S. Stoll and R. D. Britt, *Phys. Chem. Chem. Phys.*, 2009, **11**, 6614–6625.
- 46 B. Bleaney, P. M. Llewellyn and D. A. Jones, *Proc. R. Soc. B*, 1956, **69**, 858–860.
- 47 L. K. Aminov, I. N. Kurkin and B. Z. Malkin, *Phys. Solid State*, 2013, **55**, 1343–1363.
- 48 F. Jaroschik, F. Nief, X. F. Le Goff and L. Ricard, *Organometallics*, 2007, **26**, 3552–3558.

