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Isolation and characterization of a high-spin mixed-valent iron dinitrogen complex†‡

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We report a rare example of a mixed-valence iron compound with an FeNNFe core, which gives insight into the structural, spectroscopic, and magnetic influences of single-electron reductions and oxidations. In the new compound, the odd electron is localized as judged from Mössbauer spectra at 80 K and infrared spectra at room temperature, and the backbonding into the N₂ unit is intermediate between diiron(II) and diiron(0) congeners. Magnetic susceptibility and relaxation studies on the series of FeNNFe compounds show significant magnetic anisotropy, but through-barrier pathways enable fairly rapid magnetic relaxation.

Both natural and industrial nitrogen fixation use catalysts with multiple iron atoms in their active sites. In nature, nitrogenase enzymes employ mixed-valence FeS clusters to carry out the reduction of N₂ with protons and reducing agents under ambient conditions.^{1–3} Industrial nitrogen fixation uses the Haber–Bosch process to combine N₂ with H₂, and the most common catalyst contains reduced iron that is formed *in situ* from iron oxide doped with potassium and aluminium.⁴ The iron atoms in the resulting promoted catalyst have an average oxidation state between 0 and +1.⁴ We aim to prepare well-characterized small molecules that capture the key properties, such as mixed iron oxidation states and N₂ binding.

To date, only a few examples of bridging Fe–N₂–Fe complexes in mixed-valent states have been characterized.⁵ The first example

was the complex [LFe–N₂–FeL][–] (L = PhB(CH₂PPh₂)₃[–]) reported by Peters *et al.*^{5a} Field and coworkers reported a second example, the mixed-valent iron(II)–iron(0) complex [(FeH(PP₃))(μ–N₂)(Fe(PP₃))] (PP₃ = P((CH₂)₂P(CH₃)₂)₃), which features an asymmetric ligand environment.^{5b} On the basis of ¹H and ³¹P NMR spectroscopy, it was assigned as Robin–Day Class I where the oxidation states are completely localized.⁶ Finally, we reported the series of mixed-valent triiron N₂ complexes M₂[L^{Me}Fe(N₂)]₃ (M = K, Rb, Cs; L^{Me} = the β-diketiminato ligand 2,4-bis(2,6-dimethylphenylimino)-3-methyl-pent-3-yl), which feature formal iron oxidation states Fe^IFe⁰₂.^{5c} For these trinuclear complexes, the presence of a single quadrupole doublet in the Mössbauer spectra suggested that the valence is delocalized (Robin–Day Class III) at 80 K. Herein, we report the synthesis of a bimetallic, mixed-valent iron–N₂ complex using a bulkier β-diketiminato L^{Me} = 2,4-bis(2,6-diisopropylphenylimino)-pent-3-yl and show that the valence is localized using Mössbauer and infrared spectroscopies.

Addition of 1 equiv. of potassium graphite (KC₈) to a purple solution of L^{Me}FeNNFeL^{Me} (**1**) and 18-crown-6 (18-c-6) in tetrahydrofuran (THF) resulted in a green mixture. Filtration of this solution through Celite[®] and vapor diffusion of pentane into the filtrate at –40 °C led to the formation of green crystals of the iron(II)–iron(0) dinitrogen complex salt [K(18-c-6)(THF)₂][L^{Me}Fe(μ–N₂)FeL^{Me}] (**2**) in 63% yield.⁷ X-ray crystallography of **2** revealed a structure with an N–N distance of 1.186(6) Å, which is significantly longer than that in free N₂ (1.09 Å), suggestive of substantial N₂ weakening (see Fig. 1 and 3). A comparison of the core bond distances and angles for **2** to those of the known analogues **1**, [K(18-c-6)(12-c-4)]₂[L^{Me}Fe(μ–N₂)FeL^{Me}] (**3**) and K₂[L^{Me}Fe(μ–N₂)FeL^{Me}] (**4**) (shown in Fig. 2) reveals that the metrical parameters for all four complexes are statistically indistinguishable (Table 1).⁷

In the IR spectrum of **2**, there is a weak band at 1749 cm^{–1}, which shifts to 1690 cm^{–1} when **2** is synthesized under ¹⁵N₂ (Fig. S6 and S7, ESI[†]). This 59 cm^{–1} shift matches well with the 61 cm^{–1} shift predicted using a harmonic oscillator model. The ν_{N–N} peak for **2** is significantly red-shifted compared to that of free N₂ (ν_{N–N} = 2359 cm^{–1}),⁸ consistent with the N–N bond

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† Dedicated to Richard A. Andersen, an amazing scholar and mentor, on the occasion of his 75th birthday.

‡ Electronic supplementary information (ESI) available. CCDC 1866500 (**2**). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8cc07294a

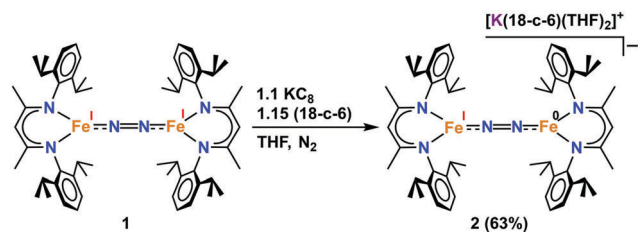


Fig. 1 Synthesis of $[L^{Me}Fe(\mu-N_2)FeL^{Me}]^-[K(18-c-6)(THF)_2]^+$ (**2**) from the reduction of **1** using KCl with 18-crown-6.

elongation in **2** as characterized by X-ray diffraction, demonstrating substantial reduction of the bound dinitrogen unit. The fact that the N–N stretching mode is observed in the infrared spectrum indicates that the molecule is not centrosymmetric, despite the inversion center in the X-ray crystal structure. In the previously characterized **1**, **3**, and **4** (as well as alkali-substituted congeners of **4**, analogues with larger β -diketiminato ligands, and Co and Ni analogues), the N–N stretching vibrations were infrared silent and thus were measured using resonance Raman spectroscopy.^{7,9} To our knowledge, the only IR-observable N–N vibrations in dinuclear β -diketiminato-supported MNNM systems have been in compounds with asymmetric cores: a spectroscopically characterized FeNNCo species^{9b} and a NiNNNi species with an asymmetrically coordinated potassium cation.¹⁰

Interestingly, complexes **1**, **2**, and **3** are congeners that span three oxidation states of the diiron core— $Fe^I Fe^I$, $Fe^I Fe^0$ and $Fe^0 Fe^0$ —with no other differences in the primary coordination environment. The sequential one-electron reductions of the core from **1** to **2** and **2** to **3** lead to bathochromic shifts of the N_2 stretch by 61 cm^{-1} and 66 cm^{-1} respectively. The red-shifting

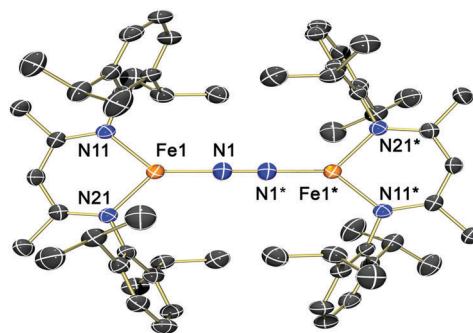


Fig. 3 X-ray crystal structure of $[L^{Me}FeNNFeL^{Me}]^-$ (**2**) with thermal ellipsoids drawn at the 50% probability level. Orange, blue, and black ellipsoids represent Fe, N, and C atoms, respectively. Hydrogen atoms, a THF molecule of crystallization, and the $[K(18-c-6)(THF)_2]^+$ counteranion are omitted for clarity. Selected bond distances (Å) and angles ($^\circ$): $Fe(1)-N(1) = 1.744(3)$; $Fe(1)-N(11) = 1.936(2)$; $Fe(1)-N(21) = 1.940(3)$; $N(1)-N(1)^* = 1.186(6)$; $N(1)-Fe(1)-N(11) = 131.7(1)$; $N(1)-Fe(1)-N(21) = 128.4(2)$; $N(11)-Fe(1)-N(21) = 98.6(1)$; $N(1)^*-N(1)-Fe(1) = 175.2(4)$.

of the N–N stretch results from the population of d orbitals that backbond into the π^* orbitals of the N_2 unit, decreasing the bond order of the N–N bond. The series **1–3** is distinctive, because the influence of the metal oxidation states can be discerned without coordinated alkali metal cations. This is important because alkali metal cations^{4b} and other Lewis acids¹¹ can further weaken the N–N bond.

The zero-field Mössbauer spectrum of **2** (Fig. 4) is best fit with two overlapping quadrupole doublets of identical intensity. A staggered fit yielded parameters $\delta_1 = 0.49\text{ mm s}^{-1}$ and $|\Delta E_Q|_1 = 1.71\text{ mm s}^{-1}$ and $\delta_2 = 0.63\text{ mm s}^{-1}$ and $|\Delta E_Q|_2 = 1.75\text{ mm s}^{-1}$ for the two components (Fig. 4; alternate fits shown in Fig. S2 and S3, ESI†). The isomer shift for component 1 (δ_1) is

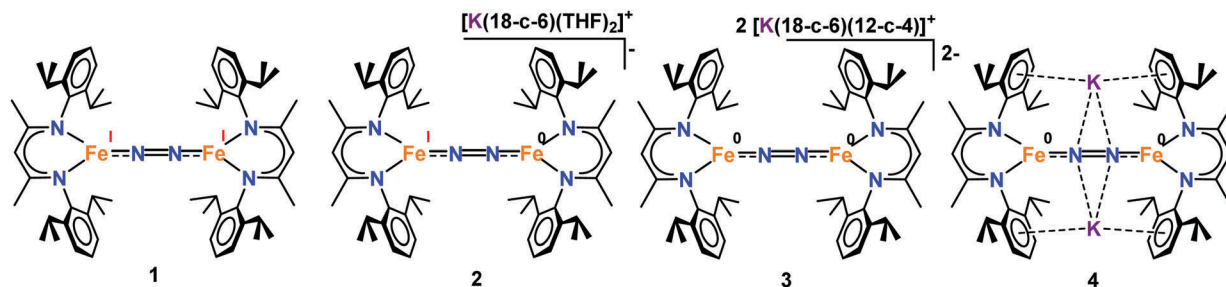


Fig. 2 ChemDraw representations of the structures of compounds **1–4**.

Table 1 Comparison of structural and spectral parameters for complexes **1–4**. Mössbauer spectral parameters were obtained on solid samples at 80 K unless otherwise noted

Complex	Formal Ox. State	N–N dist. (Å)	Fe– N_2 dist. (Å)	ν_{N-N} [$\nu_{15N-15N}$] (cm^{-1})	δ (mm s^{-1})	$ \Delta E_Q $ (mm s^{-1})
1	$Fe^I Fe^I$	1.186(7)	1.745(3)	1810 [1745] ^a	0.62 ^b	1.41 ^c
2	$Fe^0 Fe^I$	1.186(6)	1.744(3)	1749 [1690]	0.63	1.75
3	$Fe^0 Fe^0$	1.190(8)	1.758(4)	1683 [1632] ^a	0.49	1.71
4	$Fe^0 Fe^0$	1.215(6)	1.750(4), 1.755(5)	1625 [1566] ^a	0.45	1.74
					0.47	2.48

^a Values obtained from resonance Raman spectroscopy in ref. 7a and b. ^b From Mössbauer spectra of solid at 4.2 K from ref. 12. ^c From Mössbauer spectra of solid at 170 K from ref. 12.

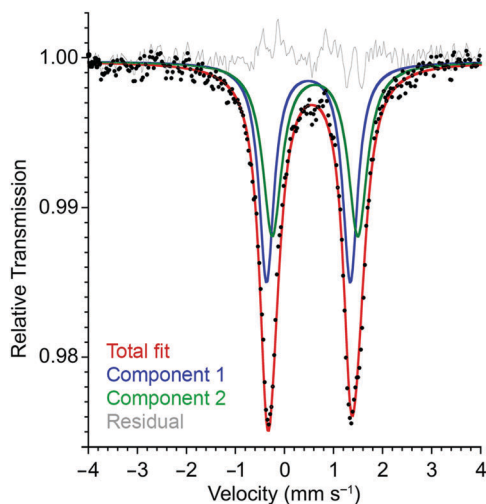


Fig. 4 Zero-field Mössbauer spectrum of solid **2** at 80 K. Fitting to one doublet gave poorer agreement (Fig. S3, ESI†).

similar to that in the diiron(i) analogue **1**, and δ_2 is similar to that in the diiron(0) analogues **3** and **4**, supporting the localization of distinct iron(i) and iron(0) sites in **2** (Table 1). An alternative nested fit (Fig. S2, ESI†) has very similar isomer shift values but different quadrupole splitting, which would not make sense for a delocalized system. Given the similarity of the Mössbauer parameters to very similar diiron(0) and diiron(i) analogues, it is most reasonable to use the staggered fit.

As described above, the previously reported mixed-valent Fe–N₂–Fe complexes were assigned as either Robin-Day Class I mixed-valent with full localization^{5b} or Robin-Day Class III with full delocalization down to 80 K.^{5b,c} For **2**, the presence of an N₂ stretching vibration in the IR spectrum and two doublets in the Mössbauer spectrum together indicate localization of the additional electron on one of the two iron sites. However, the room temperature ¹H NMR spectrum of **2** collected in THF-*d*₈ exhibits 8 resonances, with integrations that are most consistent with *D*_{2d} or *D*_{2h} symmetry (Fig. S4 and S5, ESI†). Therefore, the electron can sample both of the iron sites on the NMR timescale in solution. This is most consistent with Robin-Day Class II mixed valency, though there may be structural differences in solution (see below).⁶

Using the Evans method, the room-temperature solution magnetic moment of **2** in THF was $\mu_{\text{eff}} = 6.6 \mu_{\text{B}}$. Solid-state magnetic susceptibility data were also collected using a SQUID magnetometer, and above 50 K, the molar magnetic susceptibility times temperature ($\chi_{\text{M}}T$) for **2** collected under applied fields of 0.1–7 T each approach $7 \text{ cm}^3 \text{ K mol}^{-1}$ above 50 K (Fig. S11, ESI†), consistent with an overall $S = 5/2$ ground state. A sextet ground state can be rationalized either with ferromagnetic coupling between high-spin iron(i) ($S = 3/2$) and high-spin iron(0) ($S = 1$), or with a Fe^{II}–N₂^{2–}–Fe^I three-spin model where high-spin iron(ii) and iron(i) sites are spin-aligned ($S_{\text{Fe}} = 7/2$) and anti-ferromagnetically coupled to a triplet N₂^{2–} bridge ($S_{\text{N}_2} = 1$) to give a total $S = 5/2$ ground state.^{7,12} The latter model is more compatible with previous computations on compound **1**, which predicted an Fe^{II}–N₂^{2–}–Fe^{II} core with two high-spin iron(ii) sites

that are antiferromagnetically coupled to a triplet N₂^{2–} bridge to give a $S = 3$ ground state.^{7,12} These three-coordinate iron(ii) sites are known^{12,13} to have large negative zero-field splitting parameters (D) that suggest the potential for slow magnetic relaxation.¹⁴ Accordingly, we sought to further evaluate the magnetization behaviour of the N₂-bridged compounds.

The solid-state $\chi_{\text{M}}T$ values of **1**, **2**, and **4** at room temperature correspond to $S = 3$, $5/2$, and 2 systems, having isotropic g values of 2.38, 2.60, and 2.28. These spin states agree with solution magnetic moments,^{7,9} and the elevated g values suggest significant spin-orbit coupling. Low temperature magnetization data for **1** and **2** were collected to examine the magnetic anisotropy and the data were fit to a phenomenological zero-field splitting Hamiltonian. The data for **1** could be fit well with multiple sets of parameters and all fits suggested significant axial anisotropy ($D = -45 \text{ cm}^{-1}$ from the best fit) as well as significant transverse anisotropy ($|E/D| = 0.24$ from the best fit). The fit for **2** suggested that there is significantly less axial anisotropy ($D = -4.5 \text{ cm}^{-1}$), again with a substantial contribution from transverse anisotropy ($|E/D| = 0.29$). In contrast, X-band EPR spectra collected from a frozen 50 : 1 mixture of THF/2-methylTHF at 4.2 K exhibit several signals from $S = 5/2$ states with positive D , $D \gg h\nu$ (microwave quantum), with the majority form having $E/D = 0.097$ (Fig. S24, ESI†). This difference in the sign of D from the solid-state susceptibility measurements indicates a change in the electronic structure in frozen solution, possibly caused by interactions with the solvent.

Given the evidence of magnetic anisotropy in both **1** and **2** from the dc susceptibility measurements, we collected ac magnetic susceptibility data to probe for slow magnetic relaxation. For **1**, **2**, and **4**, non-zero signal was observed in the out-of-phase magnetic susceptibility (χ'') at the highest frequencies. However, in each case a small bias field was required to see fully resolved peaks within the measured frequency range of 1–1500 Hz (see Fig. 5 for the data for **1** and **2**), consistent with systems having both axial and transverse anisotropy. Temperature-dependent magnetic relaxation times (τ) were extracted for **1** and **2** from Cole-Cole fits to the in- and out-of-phase magnetic susceptibility data (the peaks for **4** were too broad for further analysis). Plots of the natural log of τ versus temperature for **1** and **2** are shown in Fig. S14 and S18 (ESI†), respectively, and the pronounced curvature in both sets of data indicate that thermally-activated Orbach relaxation is not dominant over the entire measured temperature and frequency range. However, a linear fit to the high temperature data gives lower limits for the spin-reversal barriers, U_{eff} , and upper limits for the attempt times, τ_0 . For **1** these limits are $U_{\text{eff}} \geq 56 \text{ cm}^{-1}$ and $\tau_0 \leq 6.8 \times 10^{-6} \text{ s}$, and for **2** they are $U_{\text{eff}} \geq 27 \text{ cm}^{-1}$ and $\tau_0 \leq 3.6 \times 10^{-7} \text{ s}$. Thus, although both of these multinuclear systems exhibit significant magnetic anisotropy, their relaxation behavior is very similar to typical mononuclear single-molecule magnets, in that both sets of complexes are strongly affected by through-barrier pathways.

In conclusion, we have described the preparation and characterization of the mixed-valent diiron compound $[\text{K}(\text{18-c-6})(\text{THF})_2][\text{L}^{\text{Me}}\text{Fe}(\mu\text{-N}_2)\text{FeL}^{\text{Me}}]$, which contains a rare example of an N₂-bridged complex that can be analyzed in several

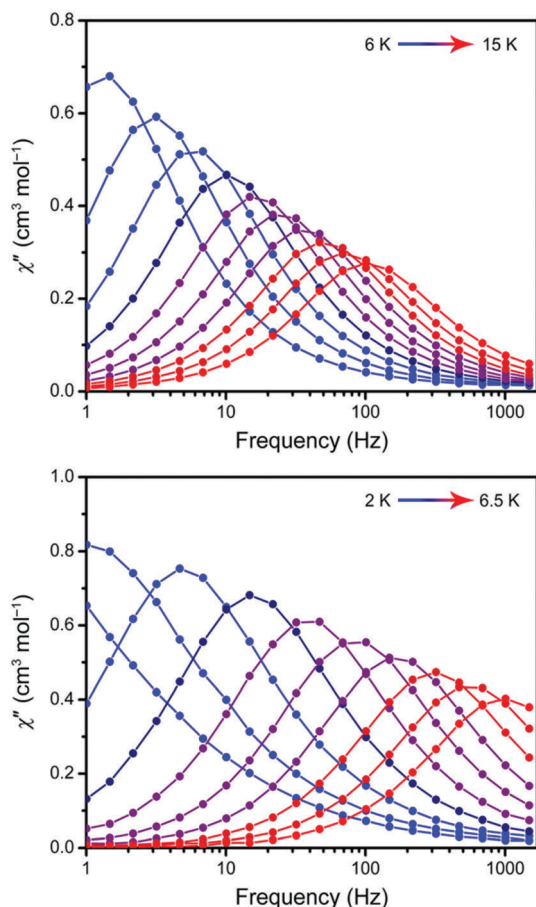


Fig. 5 (top) Out-of-phase (χ'') magnetic susceptibility data for solid $L^{\text{Me}}\text{FeNNFe}L^{\text{Me}}$ (**1**) collected under a 500 Oe dc field over temperatures ranging from 6 to 15 K. (bottom) Out-of-phase magnetic susceptibility data for solid $[\text{K}(18\text{-c-}6)][L^{\text{Me}}\text{FeNNFe}L^{\text{Me}}]$ (**2**) collected under a 1000 Oe dc field over temperatures ranging from 2 to 6.5 K. Solid lines are guides for the eye. In-phase (χ') magnetic susceptibility and additional magnetic data are shown in the ESI.†

oxidation levels that vary by one electron.¹⁵ Probing the electronic structure and magnetism in greater detail will be the basis for interesting future studies.

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Conflicts of interest

There are no conflicts to declare.

Notes and references

- 1 K. Burgess and D. J. Lowe, *Chem. Rev.*, 1996, **96**, 2983–3012.
- 2 (a) B. M. Hoffman, D. Lukoyanov, Z.-Y. Yang, D. R. Dean and L. C. Seefeldt, *Chem. Rev.*, 2014, **114**, 4041; (b) B. M. Hoffman, D. Lukoyanov, D. R. Dean and L. C. Seefeldt, *Acc. Chem. Res.*, 2013, **46**, 587.
- 3 (a) T. Spatzal, J. Schlesier, E.-M. Burger, D. Sippel, L. Zhang, S. L. A. Andrade, D. C. Rees and O. Einsle, *Nat. Commun.*, 2016, **7**, 10902; (b) R. Björnsson, F. Neese and S. DeBeer, *Inorg. Chem.*, 2017, **56**, 1470–1477.
- 4 (a) R. Schlögl, *Handbook of Heterogeneous Catalysis*, 2nd edn, Wiley-VCH, Weinheim, 2008; (b) G. P. Connor and P. L. Holland, *Catal. Today*, 2017, **286**, 21–40.
- 5 (a) T. A. Betley and J. C. Peters, *J. Am. Chem. Soc.*, 2003, **125**, 10782–10783; (b) L. D. Field, R. W. Guest and P. Turner, *Inorg. Chem.*, 2010, **49**, 9086–9093; (c) K. Grubel, W. W. Brennessel, B. Q. Mercado and P. L. Holland, *J. Am. Chem. Soc.*, 2014, **136**, 16807–16816.
- 6 M. B. Robin and P. Day, in *Advances in Inorganic Chemistry and Radiochemistry*, ed. H. J. Emeléus and A. G. Sharpe, Academic Press, 1968, vol. 10, pp. 247–422.
- 7 (a) J. M. Smith, A. R. Sadique, T. R. Cundari, K. R. Rodgers, G. Lukat-Rodgers, R. J. Lachicotte, C. J. Flaschenriem, J. Vela and P. L. Holland, *J. Am. Chem. Soc.*, 2006, **128**, 756–769; (b) S. F. McWilliams, E. Bill, G. Lukat-Rodgers, K. R. Rodgers, B. Q. Mercado and P. L. Holland, *J. Am. Chem. Soc.*, 2018, **140**, 8586–8598.
- 8 T. Shimanouchi, Molecular Vibrational Frequencies, in *NIST Chemistry WebBook*, ed. P. J. Linstrom and W. G. Mallard, NIST Standard Reference Database Number 69, National Institute of Standards and Technology, Gaithersburg, MD, <http://webbook.nist.gov>, accessed Aug 01, 2018.
- 9 (a) J. M. Smith, R. J. Lachicotte, K. A. Pittard, T. R. Cundari, G. Lukat-Rodgers, K. R. Rodgers and P. L. Holland, *J. Am. Chem. Soc.*, 2001, **123**, 9222–9223; (b) K. Ding, A. W. Pierpont, W. W. Brennessel, G. Lukat-Rodgers, K. R. Rodgers, T. R. Cundari, E. Bill and P. L. Holland, *J. Am. Chem. Soc.*, 2009, **131**, 9471–9472; (c) S. Pfirrmann, S. Yao, B. Ziemer, R. Stösser, M. Driess and C. Limberg, *Organometallics*, 2009, **28**, 6855–6860; (d) B. Horn, S. Pfirrmann, C. Limberg, C. Herwig, B. Braun, S. Mebs and R. Metzinger, *Z. Anorg. Allg. Chem.*, 2011, **637**, 1169–1174; (e) S. F. McWilliams, K. R. Rodgers, G. Lukat-Rodgers, B. Q. Mercado, K. Grubel and P. L. Holland, *Inorg. Chem.*, 2016, **55**, 2960–2968.
- 10 S. Pfirrmann, C. Limberg, C. Herwig, R. Stößer and B. Ziemer, *Angew. Chem., Int. Ed.*, 2009, **48**, 3357–3361.
- 11 J. B. Geri, J. P. Shanahan and N. K. Szymczak, *J. Am. Chem. Soc.*, 2017, **139**, 5952–5956.
- 12 S. A. Stoian, J. Vela, J. M. Smith, A. R. Sadique, P. L. Holland, E. Münck and E. L. Bominaar, *J. Am. Chem. Soc.*, 2006, **128**, 10181–10192.
- 13 H. Andres, E. Bominaar, J. M. Smith, N. A. Eckert, P. L. Holland and E. Münck, *J. Am. Chem. Soc.*, 2002, **124**, 3012–3025.
- 14 G. Christou, D. Gatteschi, D. N. Hendrickson and R. Sessoli, *MRS Bull.*, 2000, **25**, 66–71.
- 15 (a) D. E. Richardson, J. P. Sen, J. D. Buhr and H. Taube, *Inorg. Chem.*, 1982, **21**, 3136–3140; (b) W. A. Hoffert, A. K. Rappé and M. P. Shores, *Inorg. Chem.*, 2010, **49**, 9497–9507.