

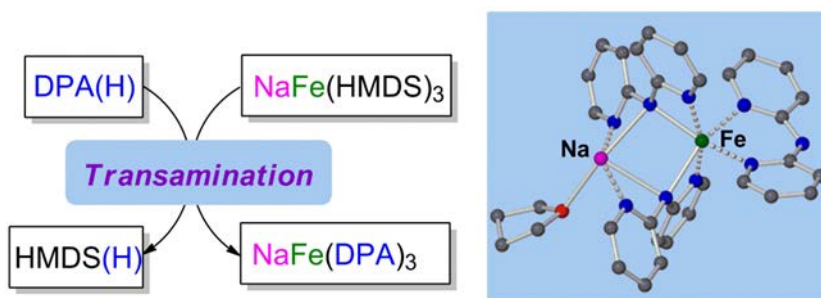
Synthetic, Structural and Magnetic Implications of Introducing 2,2'-Dipyridylamide to Sodium-Ferrate Complexes

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Establishing transamination as an efficient method to access novel heterobimetallic complexes, this study provides magnetic and structural insights of 2,2-dipyridylamide based sodium ferrate complexes.

Abstract

Using a transamination approach to access novel Fe(II) complexes, this study presents the synthesis, single crystal X-ray diffraction and magnetic characterisation of a series of new iron complexes containing the multifunctional 2,2-dipyridylamide (DPA) ligand using iron bis(amide) [$\{\text{Fe}(\text{HMDS})_2\}_2$] and sodium ferrate [$\{\text{NaFe}(\text{HMDS})_3\}_\infty$] (**1**) as precursors (HMDS = 1,1,1,3,3,3-hexamethyldisilazide). Reactions of DPA(H) with **1** show exceptionally good stoichiometric control, allowing access to heteroleptic [$(\text{THF})_2\cdot\text{NaFe}(\text{DPA})(\text{HMDS})_2$] (**3**) and homoleptic [$\{\text{THF}\cdot\text{NaFe}(\text{DPA})_3\}_\infty$] (**4**) by using 1 and 3 equivalents of DPA(H) respectively. Linking this methodology and co-complexation, which is a more widely used approach to prepare heterobimetallic complexes, **3** can also be prepared by combining NaHMDS with heteroleptic [$\{\text{Fe}(\text{DPA})(\text{HMDS})\}_2$] (**2**). In turn, **2** has been also synthesised and structurally defined by reacting [$\{\text{Fe}(\text{HMDS})_2\}_2$] with one equivalent of DPA(H). Structural studies demonstrate the coordination flexibility of the N-bridged bis(heterocycle) ligand DPA, with **2** and **3** exhibiting discrete monomeric motifs, whereas **4** displays a much more intricate supramolecular structure, with one of its DPA ligands coordinating in a *anti/anti* fashion (as opposed to **2** and **3** where DPA shows a *syn/syn* conformation), which facilitates propagation of the structure via its central amido N. Magnetic studies confirmed the high-spin electron configuration of the iron(II) centres in all three compounds and revealed the existence of weak ferromagnetic interactions in dinuclear compound **2** ($J=1.01\text{ cm}^{-1}$).

Introduction

Advances in the development of cooperative heterobimetallic compounds and their application in synthesis continue to attract widespread interest.¹⁻⁴ Exhibiting unique synergistic properties, mixed-metal complexes (also known as *ates*) can effectively execute key organic transformations such as deprotonative metallation,⁵ metal/halogen exchange⁶ and nucleophilic addition,⁷⁻⁹ demonstrating exceptional reactivity profiles, outperforming in many cases, traditional single-metal reagents.¹⁰⁻¹³ To date, most of this research activity has concentrated on complexes which combine an alkali-metal with an s/p block lower polarity metal such as Mg, Zn or Al.^{1-4,14,15} Some examples of the remarkable reactivity of these systems includes the regioselective alpha-metallation of THF and the *ortho-meta'* and *meta-meta'* di-magnesiatiations of a series of arenes with sodium magnesium template base $[\text{Na}_4\text{Mg}_2(\text{TMP})_6(^n\text{Bu})_2]$ (TMP = 2,2,6,6-tetramethylpiperidide).¹⁶

Efforts to expand upon the numerous successes observed with main-group systems and extend these synthetic approaches to earth-abundant transition metals have so far been limited.¹⁷⁻²⁰ Iron's inherent abundance and benign nature presents many economical and ecological benefits²¹⁻²⁴ but furthermore its open-shell character introduces a new dimension of interest not accessible to main-group systems, namely the potential to exhibit interesting magnetic behaviours.²⁵⁻³⁰ Heterobimetallic compounds containing iron, known as ferrate complexes,³¹⁻³⁷ have garnered interest as potential key intermediates in Fe-catalysed C-C bond forming processes³⁸⁻⁴¹ and have shown the ability to mediate other important synthetic processes. Indeed, Mongin has reported the metallation of aromatic and heteroaromatic substrates at ambient temperature using lithium ferrate complex $[\text{LiFe}(\text{TMP})_3]$.⁴² Closely related to this work, Knochel has demonstrated the synthesis of the Fe(II) complex $[(\text{TMP})_2\text{Fe}(\text{MgCl}_2)_2(\text{LiCl})_4]$, capable of metallating functionalised arenes to undergo subsequent nickel-catalysed cross-couplings with alkylhalides.⁴³ Though elegant applications of ferrate complexes in organic chemistry, their constitutions remain solely putative as the reports provide no structural or spectroscopic characterisation. However, Mulvey has reported the structurally well-defined ferrate complex $[\text{Na}_4\text{Fe}_2(\text{TMP})_6(\text{C}_6\text{H}_4)]$, which exhibits an unprecedented metallating power, promoting the direct diferration of benzene.⁴⁴

Other relevant structural studies include Layfield's work using $\text{Fe}(\text{HMDS})_2$ (HMDS = 1,1,1,3,3,3-hexamethyldisilazide) to access of homo- and heterometallic Fe(II) cage complexes⁴⁵ as well as homoleptic tris(amido) lithium ferrate complexes.⁴⁶ In addition, we

have also reported the ferration of N-heterocyclic carbene IPr (IPr = 1,3-bis(2,6-diisopropylphenyl)imidazol-2-ylidene) at its C4 position by sequentially reacting it with $\text{NaCH}_2\text{SiMe}_3$ and $\text{Fe}(\text{HMDS})_2$.⁴⁷ Moreover, our studies on co-complexation reactions have shown that $\text{Na}(\text{HMDS})$ and $\text{Fe}(\text{HMDS})_2$ undergo self-assembly in non-donor solvents such as hexane to form novel polymeric solvent-free $[\{\text{NaFe}(\text{HMDS})_3\}_\infty]$ (**1**) which exhibits an unusual polymeric chain structure.⁴⁷

Building on these initial studies, here we explore the reactivity of this heterobimetallic complex to undergo transamination reactions with 2,2'-dipyridylamine (DPA(H)) to access novel homo- and heteroleptic sodium ferrate complexes.

Although less prevalent throughout organometallic chemistry than HMDS,⁴⁸ amido DPA has been utilised in a number of varied branches of chemistry⁴⁹ including materials science,⁵⁰ catalysis,⁵¹ supramolecular chemistry⁵² and even in cooperative bimetallic chemistry.⁵³ Being the simplest of the poly 2,2'-pyridyl amides, DPA can potentially coordinate through three N sites; one central amido N and two neutral pyridyl N atoms.⁵⁴ Rotation around the two $\text{N}_{\text{amido}}\text{-C}$ bonds allows for DPA to adopt three different conformations; *syn/syn*, *syn/anti* and *anti/anti* (Fig. 1).⁵⁵ Within heterobimetallic chemistry, Mulvey has successfully prepared mixed sodium-zinc reagents containing this amide which can promote the remote alkylation of benzophenone at the para position of one of its phenyl groups.⁵³

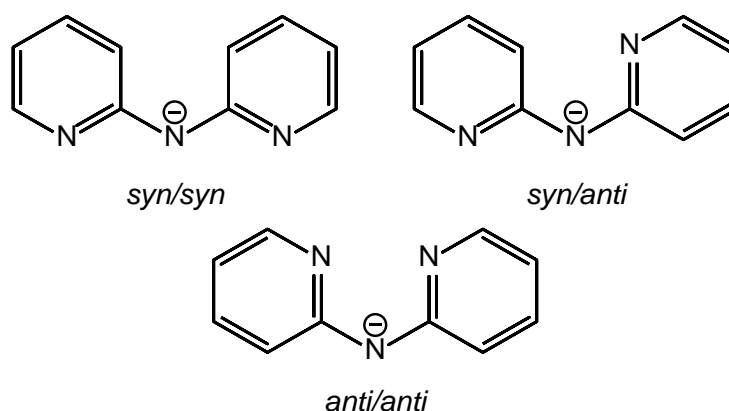


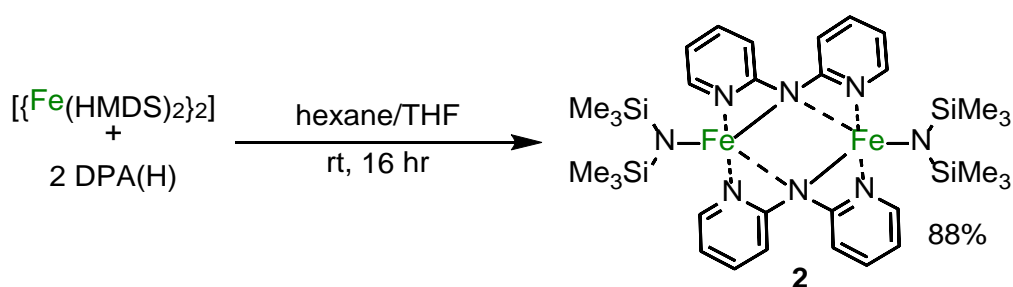
Fig. 1 The three conformational arrangements of 2,2'-dipyridylamide.

Extending the use of this amide to mixed sodium-iron chemistry, here we report our findings on the synthesis of a new family of sodium ferrates. Combining X-ray crystallographic studies with NMR spectroscopic and SQUID magnetisation investigations, new insights into the constitution and magnetic properties of these novel sodium ferrate complexes are presented.

Results and Discussion

Synthetic and structural studies

We started our investigations with single-metal amide $\text{Fe}(\text{HMDS})_2$, by reacting it with one molar equivalent of $\text{DPA}(\text{H})$ in hexane, which produced a brown solution with an off-white precipitate. Addition of THF afforded a homogeneous dark solution which upon cooling to -30°C deposited a crop of orange crystals of heteroleptic bis(amide) $[\{\text{Fe}(\text{HMDS})(\text{DPA})\}_2]$ (**2**) in a 88% yield (Scheme 1).



Scheme 1 Transamination of $\text{DPA}(\text{H})$ with $\text{Fe}(\text{HMDS})_2$.

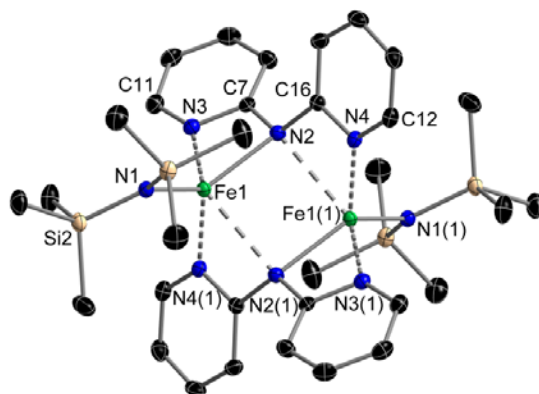


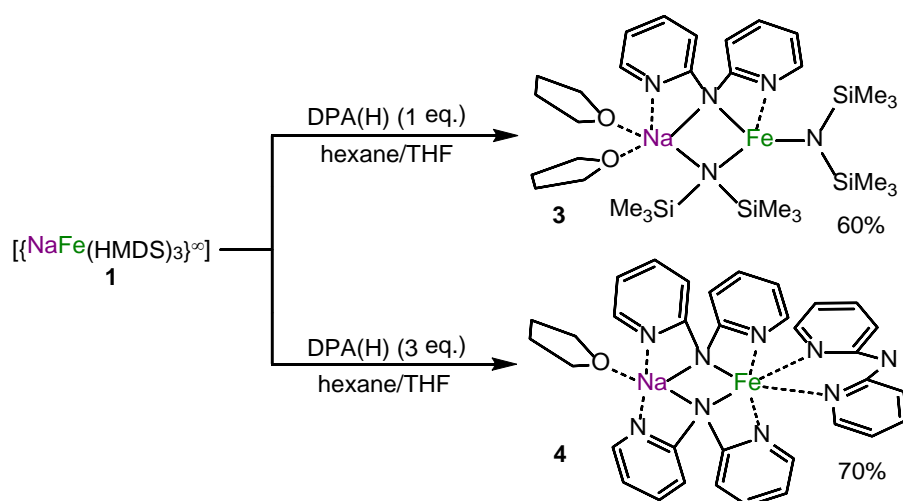
Fig. 2 X-ray crystal structure of complex **2**. Hydrogen atoms omitted for clarity. Thermal ellipsoids displayed at 50% probability level. Selected bond distances (\AA) and angles ($^\circ$): Fe1-N1 1.9583(14), Fe1-N2 2.1642(14), Fe1-N3 2.2112(15), Fe1-N(4) 2.1044(14), $\text{Fe1}\cdots\text{N2(1)}$ 2.546(1), $\text{Fe1}\cdots\text{Fe1(1)}$ 3.3609(1); N1-Fe1-N2 134.76(6), N1-Fe1-N3 119.34(6), N1-Fe1-N4(1) 111.04(6), N2-Fe1-N3 61.70(5), N2-Fe1-N4(1) 113.54(5), N3-Fe1-N4(1) 97.63(5), $\text{N1-Fe1}\cdots\text{Fe1(1)}$ 134.078(1).

As determined by X-ray crystallography, **2** displays a dimeric structure with a novel eight-membered $\{\text{FeNCNFeNCN}\}$ core (Fig. 2). Two opposing Fe centres are each coordinated by HMDS in terminal positions whilst the DPA ligands assume bridging positions between the metal centres. The DPA ligands maximise coordination to the Fe centres through their neutral

ring nitrogens with the pyridyl rings adopting a *syn/syn* conformation (a linear array of the N atoms) with an interplanar Py-N_{amido}-Py (Py = pyridyl) angle of 50.458°. Fe1 is formally five-coordinate when considering the long contact to the opposing central amido nitrogen (Fe1---N2(1) 2.546(1) Å); though much shorter bond distances are observed between Fe1 and amido nitrogens N1 and N2 (1.9583(14) and 2.1642(14) Å, respectively) and to pyridyl nitrogens N3 and N4(1) (2.2112(15) and 2.1044(14) Å, respectively). Discounting N2(1), a distorted tetrahedral geometry is present around Fe1 (average N-Fe-N angle = 106.36°, range 61.70(5)° to 134.76(6)°, excluding N2(1)). The Fe1---Fe1(1) separation in **2** is 3.3609(1) Å, considerably elongated from that of the equivalent Fe---Fe separation in [{Fe(HMDS)₂}₂] at a distance of 2.663(2) Å.⁵⁶ Exhibiting a good solubility in C₆D₆, the paramagnetic character of **2** was evidenced in the five broad paramagnetically shifted resonances observed in its ¹H NMR spectrum, ranging from 47.15 to −14.31 ppm and accounting for all protons of DPA along with a distinct broad singlet at 17.01 ppm integrating for 18H which can be assigned to the SiMe₃ groups. This chemical shift is significantly different to that found for the Fe(HMDS)₂ precursor (at 60.27 ppm).⁴⁷ The solution phase magnetic moment of **2** was found to be 11.28 μ_B (determined by Evans method)^{57,58} which is consistent with previous examples found in the literature where two high-spin Fe(II) centres are connected by a bridging ligand as for example **xxxx** (Ivana, any examples here?).

Whilst several Fe complexes containing the amine DPA(H) have been structurally defined,^{59–61} there are only two examples where Fe is bonded to amide DPA.^{62,63} The structure of **2** is reminiscent to that recently reported for [{Fe(Mes)(DPA)}₂] resulting from the metallation of DPA(H) by bis(aryl) complex [{Fe(Mes)₂}₂], where the DPA ligands also coordinate in a *syn-syn* fashion, bridging the Fe centres while the mesityl groups are bound terminally.⁶² Interestingly, despite the fact that 2 eq. of DPA(H) are employed in the reaction, only one of mesityl groups can be replaced by the amide DPA, which contrasts with divergent reactivity observed with other first row transition metals (Cr, Co, Ni) which under the same conditions are able to form bis(amide) complexes [M₂(DPA)₄].^{62,64–68}

Encouraged by the successful transamination reaction using Fe(HMDS)₂, we next assessed the incorporation of DPA into sodium ferrate scaffolds, using a similar approach, by treating homoleptic **1** with variable amounts of the amine DPA(H) (Scheme 2).



Scheme 2 Transamination of 1 and 3 equivalents of DPA(H) with sodium ferrate **1**.

Addition of one molar equivalent of DPA(H) to a solution of **1** in hexane immediately afforded a brown suspension which could be solubilised by introducing THF (Scheme 2, top). Orange crystals obtained at -30°C were found to be mixed-amide sodium ferrate $[(\text{THF})_2 \cdot \text{NaFe}(\text{DPA})(\text{HMDS})_2]$ (**3**), recovered in a 60% yield. Interestingly, complex **3** was found to be accessible by the combination of equimolar equivalents of mixed-amido iron complex **2** and NaHMDS in C_6D_6 (see Supporting Information).

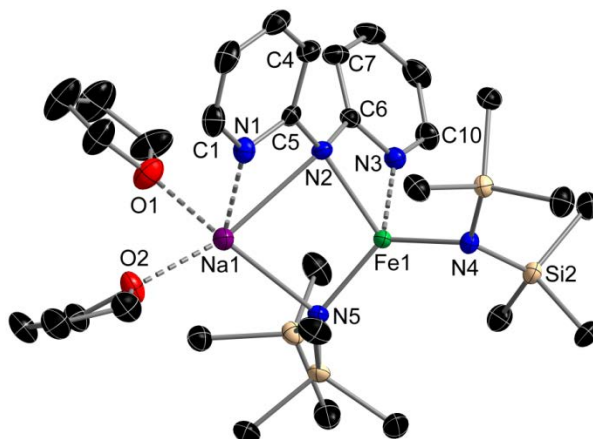


Fig. 3 Molecular structure of complex **3**. Hydrogen atoms and disorder present in the two THF molecules and one SiMe₃ unit omitted for clarity. Thermal ellipsoids displayed at 50% probability level. Selected bond distances (Å) and angles (°): Fe1-N2 2.1723(13), Fe1-N3 2.1844(14), Fe1-N4 1.9769(14), Fe1-N5 2.0171(14), Fe1---Na1 3.2253(7), Na1-N1 2.3822(17), Na1-N2 2.6710(15), Na1-N5 2.5464(15), Na1-O1 2.3479(15), Na1-O2 2.3425(14); N2-Fe1-N3 61.71(5), N2-Fe1-N4 118.24(6), N2-Fe1-N5 105.93(5), N3-Fe1-N4 103.30(6), N3-Fe1-N5 121.04(6), N4-Fe1-N5 128.42(6), Na1-N2-Fe1 82.82(5), Na1-N5-Fe1 89.16(5), Na1---Fe1-N4 145.19(4), N1-Na1-N2 53.62(5), N1-Na1-N5 119.96(5), N1-Na1-O1 91.84(6), N1-Na1-O2 98.80(6), N2-Na1-N5 79.73(5), N2-Na1-O1 94.22(5), N2-Na1-O2 152.28(5), N5-Na1-O1 131.07(6), N5-Na1-O2 118.20(5), O1-Na1-O2 88.84(5).

X-ray crystallographic studies of **3** confirmed successful transamination with HMDS and the incorporation of one DPA ligand into the ferrate structure which resides in a *syn/syn* conformation, acting as a bridge between Na1 and Fe1 to generate a monomeric contacted ion-pair structure (Fig. 3). DPA's central amido nitrogen, N2, bridges between Fe1 and Na1 at distances of 2.1723(13) and 2.6710(15) Å, respectively, whilst neutral pyridyl nitrogens N1 and N3 provide additional coordination to Na1 and Fe1, respectively. The interplanar Py-N_{amido}-Py angle displayed by the DPA ligand in **3** is 44.281°, marginally more acute than in **2**. As previously described for other mixed-metal systems, including sodium ferrate **1**,⁴⁷ *anchoring* and *ancillary* bonding modes are present in the molecular architecture of compound **3**.⁶⁹ The Fe(II) centre forms shorter and more covalent Fe-N_{amide} bonds [ranging from 1.9769(14) to 2.1723(13) Å], providing the foundation for the {Fe(DPA)(HDMS)₂}⁻ anion to which the {Na(THF)₂}⁺ cation is affixed by a combination of weaker Na-N ancillary bonds involving N2, N5 and N1. Notably a close inspection of the different metal-N(DPA) distances shows that while for iron both Fe-N_{amide} and Fe-N_{pyridyl} are comparable [2.1723(13) and 2.1844(5) Å, respectively], in the case of Na, the interaction with the N of the pyridyl ring (N1 in Fig. 3) is stronger than that with N_{amide} (N2 in Fig. 3) [2.3822(17) and 2.6710(15) Å, respectively] which is consistent with significant delocalisation of the negative charge of the

amido ligand within the pyridyl rings. As mentioned above, although alkali-metal amido ferrates have already shown interesting applications in synthesis, the number of structurally defined complexes is scarce. Related to **2**, Layfield has reported mixed lithium-iron (II) complex $[\{\text{LiFe}(\text{BTA})(\text{HMDS})_2\}_2]$ (BTA(H) = benzotriazole), which displays a dimeric structure with a central $\{\text{Li}(\text{BTA})\}_2$ core capped on each end by a $\text{Fe}(\text{HMDS})_2$ unit and has been prepared by salt-metathesis of $\text{Li}(\text{BTA})$ with FeBr_2 .⁴⁵

The ^1H NMR spectrum of **3** in C_6D_6 reveals one very broad resonance at 6.39 ppm and two marginally sharper resonances at 4.97 and 2.12 ppm. The extremely broad nature of the resonance at 6.39 ppm which also overlaps with the residual solvent signal and another resonance at 4.97 ppm, precludes a meaningful integration and assignment of the spectrum. The solution phase magnetic moment of **3** was found to be $4.93 \mu_{\text{B}}$ (determined by Evans method) which is close to the expected value ($4.90 \mu_{\text{B}}$) for a high-spin ($S = 2$) $\text{Fe}(\text{II})$ centre.

Introduction of 3 molar equivalents of DPA(H) to a hexane solution of **1** and stirring overnight generated a mustard coloured suspension in a brown solution (Scheme 2, bottom). Recrystallisation from toluene/THF at -30°C furnished yellow plate-like crystals whose structure was established by X-ray crystallography to be polymeric $[\{\text{THF} \cdot \text{NaFe}(\text{DPA})_3\}_\infty]$ (**4**) (Fig. 4), isolated in an 70% crystalline yield.

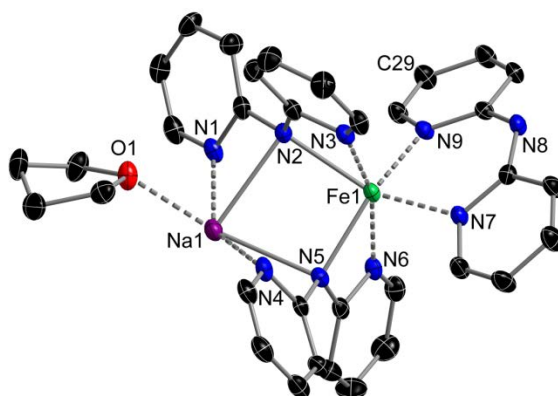


Fig. 4 Asymmetric unit of complex **4**. Hydrogen atoms and co-crystallised disordered toluene omitted for clarity. Thermal ellipsoids displayed at 30% probability level. Selected bond distances (Å) and angles (°): Fe1-N2 2.286(4), Fe1-N3 2.177(4), Fe1-N5 2.159(3), Fe1-N6 2.262(4), Fe1-N7 2.127(3), Fe1-N9 2.101(3), Fe1---Na1 3.4879(17), Na1-N1 2.512(4), Na1-N2 2.571(4), Na1-N4 2.602(4), Na1-N5 2.512(4), Na1-O1 2.362(4), Na1-N8(1) 2.480(4); N2-Fe1-N3 60.34(13), N2-Fe1-N5 92.97(13), N2-Fe1-N6 112.41(13), N2-Fe1-N7 152.05(14), N2-Fe1-N9 93.73(14), N3-Fe1-N5 103.05(14), N3-Fe1-N6 162.71(12), N3-Fe1-N7 92.74(14), N3-Fe1-N9 102.07(14), N5-Fe1-N6 60.46(13), N5-Fe1-N7 101.03(14), N5-Fe1-N9 154.03(14), N6-Fe1-N7 95.53(14), N6-Fe1-N9 93.87(13), N7-Fe1-N9 84.21(14), Na1-N2-Fe1 91.61(13), Na1-N5-Fe1 96.31(14), N1-Na1-N2 53.13(12), N1-Na1-N4 147.38(13), N1-Na1-N5 110.86(13), N1-Na1-O1 87.01(14), N1-Na1-N8(1) 103.22(13), N2-Na1-N4 94.31(12), N2-Na1-N5 78.73(12), N2-Na1-O1 94.62(13), N2-Na1-N8(1) 153.95(15), N4-Na1-N5 52.85(12), N4-Na1-O1 98.97(14), N4-Na1-N8(1) 108.15(14), N5-Na1-O1 149.71(14), N5-Na1-N8(1) 104.14(13), O1-Na1-N8(1) 94.72(13).

Complex **1** has undergone a three-fold transamination to release three equivalents of HMDS(H), incorporating three DPA ligands to furnish a new homoleptic sodium ferrate. Two DPA units bridge between Na1 and Fe1, residing in *syn/syn* conformations as seen in **3**. Contrastingly, the terminal DPA ligand adopts an *anti/anti* conformation, thus Fe1 is coordinated by the two pyridyl nitrogens N7 and N9 (2.127(3) and 2.101(3) Å, respectively), whilst bridgehead amido N8 points away to a sodium atom of a second monomer unit [at a distance of 2.480(4) Å] to give a novel 1D polymeric chain (Fig. 5).

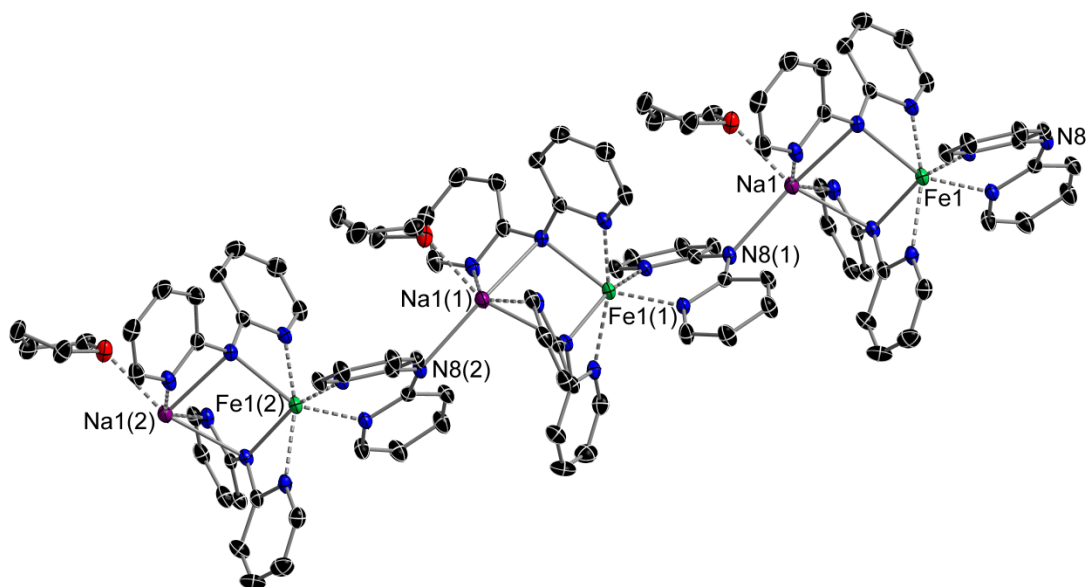


Fig. 5 Section of polymeric chain in **4** showing propagation and selected atom labelling, Na1-N8(1) 2.480(4) Å. Hydrogen atoms and co-crystallised disordered toluene omitted for clarity. Thermal ellipsoids displayed at 30% probability level.

The Fe(II) centre is hexacoordinated residing coordinating to 6 N atoms in a highly distorted octahedral environment [NFeN angles ranging from 60.34(13)° to 162.71(12)°]. Along with the pseudo-terminal DPA ligand with an *anti/anti* conformation (vide supra), Fe1 completes its coordination by bonding to amido N2 and N5 at distances of 2.286(4) and 2.159(3) Å, respectively and two further pyridyl N3 and N6 at distances of 2.177(4) and 2.262(4) Å, respectively. Looking at the sodium coordination, it binds the two bridging DPA ligands thorough their amido N and one of their pyridyl rings (N2, N5 and N1, N4 respectively) giving rise to interactions of similar strength despite the different types of N atoms involved in the bonding [Na-N distances ranging from 2.512(4) to 2.602(4) Å]. Coordinative saturation is achieved by binding to a molecule of THF and by interacting with the amido N of a DPA unit from a neighbouring unit (N8) which allows the propagation of the polymeric structure (vide supra, Fig. 5). Noticeably this supramolecular Fe-N interaction [2.480(4) Å] is shorter than those observed within the asymmetric unit of **4** (average Fe-N 2.549 Å).

In addition, the bridging DPA ligands in **4** effectively mirror one another by significant twisting of the pyridyl rings from the {Na1-N2-Fe1-N5} core plane, maximising N_{pyridyl} coordination to both metal centres (Fig. 6). Incorporating two bridging DPA groups in *syn/syn* conformations, translates in an appreciably larger Na---Fe separation [3.4879(17) Å] in **4** when compared to sodium ferrates **1** [3.0131(13) Å] and **3** [3.2253(7) Å]. Whilst the interplanar Py-N_{amido}-Py angle of 50.465° for N2 DPA ligand is of similar size to the equivalent angles

observed in complexes **2** and **3**, the N5 bridging DPA ligand displays a far more acute interplanar angle of 39.444° , whilst an acuter still angle of 24.384° is observed for the N8 terminal DPA ligand in **4**.

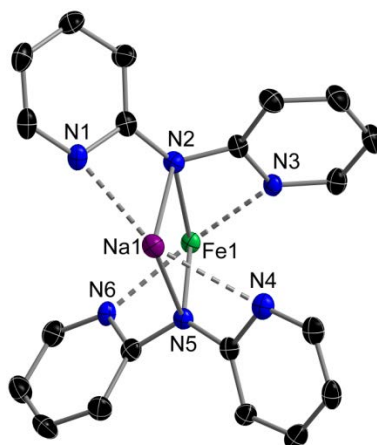


Fig. 6 Alternate view of the Na/Fe core and two bridging DPA ligands of **4**. Hydrogen atoms omitted for clarity. Thermal ellipsoids displayed at 30% probability level.

Similarly to **3**, the ^1H NMR spectrum of **4** in C_6D_6 displayed a number of poorly resolved, broad and overlapping signals (between 0 and 30 ppm) which precluded a meaningful assignment of the resonances. The solution-phase magnetic moment of **4** ($5.30 \mu_{\text{B}}$) was determined using the Evans method and is consistent with a high-spin $S = 2$ configuration.

As far as can be ascertained, complexes **3** and **4** constitute the first examples of ferrate chemistry which incorporate DPA in their constitution which have been prepared and structurally defined. Within mixed-metal chemistry it should be noted Mulvey's work on sodium zincates which include the synthesis of $[(\text{TMEDA})_2\text{Na}_2(\mu\text{-DPA})_2\text{Zn}(\text{tBu})_2]$ and $[\text{Na}(\text{THF})_6]^+[\text{Zn}(\text{tBu})_2(\text{DPA})\text{Zn}(\text{tBu})_2]^-$.⁵³ In the former structure, one DPA bridges *syn/syn* between two Na atoms and another bridges in *anti/anti* conformation with N_{amido} centred between the Na atoms and the pyridyl N atoms coordinating to Zn. Whilst in the latter solvent separated ion pair, a single unit of DPA is sandwiched between two Zn centres in an *anti/anti* conformation. From a more general perspective, Coronado has described the design of several mixed-metal chains containing oxalate ligands, including trimetallic complex $[\text{K}(18\text{-crown-6})]^+[\text{Co}(\text{DPA})\text{Fe}(\text{ox})_3]^-$ (ox = oxalate),⁷⁰ which exhibits interesting magnetic properties although the DPA ligand in this system coordinates exclusively to Co.

Magnetic Studies of DPA Complexes

Can we perhaps comment on the different magnetic properties between the new ferrates and those reported by us for {NaFe(HMDS)3} in *Inorg. Chem.* 2015, 54, 9201

The electronic structure of Fe(II) in complexes **2**, **3** and **4** was studied through bulk magnetisation measurements and (for **2**) EPR spectroscopy. Thus, molar paramagnetic susceptibility (χ_M) data were collected on microcrystalline samples from 2 to 300 K, under a constant magnetic field of 0.5 T. Additionally, field dependent (0 to 5 T) magnetization measurements at 2 K were performed.

The $\chi_M T$ vs T plot for **2** (Fig. 7) at 300 K features a value of $\chi_M T$ of $6.79 \text{ cm}^3 \text{ K mol}^{-1}$, which is higher than expected for two non-interacting high-spin (HS; $S = 2$) Fe(II) centres ($6.00 \text{ cm}^3 \text{ K mol}^{-1}$ if $g = 2.0$). The anomalous tail in this temperature region is attributed to marginal decomposition of the sample upon warming. A g factor of 2.13, revealing the coupling of an unquenched angular momentum to the electronic spin is estimated using the Curie Law near room temperature. This point is reached from a maximum of $9.37 \text{ cm}^3 \text{ K mol}^{-1}$ at 9 K, which is followed by a sharp decrease that becomes smoother after 50 K. The maximum is attained through a sharp increase from a $\chi_M T$ value of $6.88 \text{ cm}^3 \text{ K mol}^{-1}$ at 2 K. The latter pattern is ascribed to the effects of the zero-field splitting (ZFS). These effects are corroborated by the $M/N\mu_B$ vs H curve, which does not reach saturation at the highest magnetic field (it reaches $6.02 \mu_B$ at 5 T, while the expected value is $8 \mu_B$ for $g = 2.0$ and $S = 4$ or two $S = 2$).

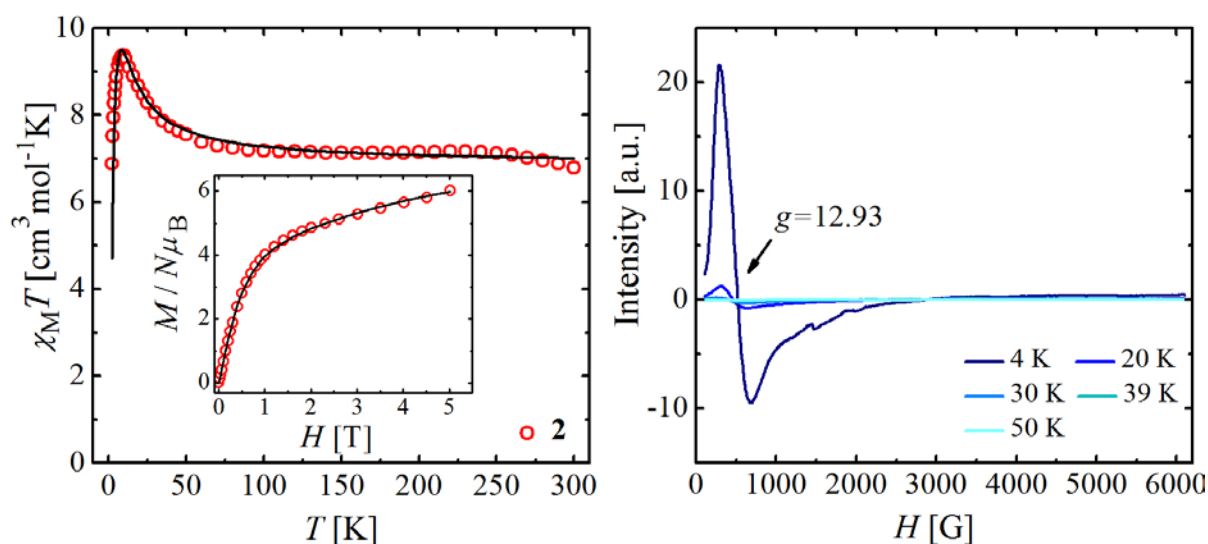


Fig. 7 Left: $\chi_M T$ vs T and $M/N\mu_B$ vs H (inset) curves of compound **2** with the best fit (solid line, see text for details). Right: Variable temperature X-band EPR spectra of a powdered sample of complex **2**.

A simultaneous fit of both curves was carried out using the program PHI⁷¹ by matrix diagonalisation of the (perturbative) anisotropic spin Hamiltonian defined in Equation 1:

$$\hat{H} = D \sum_i \left(\hat{S}_{iz}^2 - \frac{1}{3} \hat{S}_i^2 \right) + E \sum_i \left(\hat{S}_{ix}^2 - \hat{S}_{iy}^2 \right) + g\mu_B B \sum_i \hat{S}_i - 2J(\hat{S}_1 \hat{S}_2) \quad (1)$$

In Eq. 1, J is the exchange constant, \hat{S}_{ij} ($i = 1, 2; j = x, y, z$) is the total spin operator of the individual Fe(II) ions, B is the magnetic induction and μ_B is the Bohr magneton, while D and E stand for axial and rhombic ZFS parameters, respectively. To avoid the overparameterization of the Hamiltonian, the isotropic g factor was fixed at 2.14. The best fit produced $J = 1.01 \text{ cm}^{-1}$, $D = 7.31 \text{ cm}^{-1}$ and $|E| = 1.36 \text{ cm}^{-1}$, together with a small intermolecular interaction $zJ = 0.02 \text{ cm}^{-1}$. Considering that negative values of D have been reported for other trigonal-pyramidal Fe(II) complexes,^{72,73} a second set of parameters with negative D was explored, yielding $J = 0.93 \text{ cm}^{-1}$, $D = -5.59 \text{ cm}^{-1}$ and $|E| = 1.45 \text{ cm}^{-1}$. The latter presents slightly higher deviations from the experiment at low temperature. In both cases, large values of the rhombic ZFS parameter E reflect significant distortions of the basal FeN_3 plane from the ideal three-fold symmetry. However, a positive sign of D can be anticipated by the strong distortion of the coordination geometry of **2** with respect to the highly regular trigonal-pyramidal symmetry reported by Long *et al.*^{72,73} Instead, the ligand field in **2** brings it closer to a very distorted tetrahedral geometry, which in d^6 ions is expected to cause positive D values.⁷⁴

The magnetic exchange between both Fe centres of the complex is consistent with the short distance (3.3611(6) Å) mediating between them. The most likely mechanism of this exchange is the spin polarization of ligand centred electronic clouds spread over each of the N–C–N moieties of DPA bridging the metals, and orthogonal to some of the magnetic orbitals of the latter. The unusual coordinating mode seen in **2** was also reported for the compound $[\{\text{Fe}(\text{Mes})(\text{DPA})\}_2]$.⁶² In the absence of magnetic studies, DFT calculations confirmed the HS state of the Fe(II) ions and local spin densities of 3.62 at these centres (thus showing the $S = 4$ ground state). Ferromagnetic coupling within Fe(II) dinuclear complexes incorporating similar bridging motifs was also reported for $[\text{Fe}_2\text{L}_4](\text{ClO}_4)_4$ (L = 1,13,14-triaza-

dibenz[*a,j*]anthracene)⁷⁵ and [Fe₂Br₃(trop₂AM)] (H-trop₂AM= *N,N'*-bis-trop-phenylamidine).⁷⁶

Variable temperature EPR (Fig. 7) mirrors the magnetic behaviour of **2** described above, thus confirming the Fe oxidation state of +2. The latter is obvious from the fact that the sample is EPR silent above 40 K. Down to 20 K, one resonance becomes apparent in the low-field region (*g* value of 12.93) increasing in intensity upon cooling. The observed spectral feature is related to the *S* = 4 state of the Fe(II) dimer where a forbidden transition occurs between the *M_s* levels +4 and −4 ($\Delta M_s = 8$) which are split in zero magnetic field by $\sim E^2/D$.⁷⁷

For compounds **3** and **4**, the $\chi_M T$ value at 300 K is (in the **3/4** format) 3.682/3.417 cm³Kmol^{−1} (Fig. 8), higher than expected for an isolated HS (*S*=2) Fe(II) centre (calculated as 3.00 cm³ K mol^{−1} for *g* = 2.0) because of the effect of unquenched angular momentum. This yields a Curie Law estimated *g* factor of 2.22/2.13. The $\chi_M T$ vs *T* plot shows a nearly constant value with only a slightly positive slope as a result of temperature independent paramagnetism (TIP) estimated as 330/250 x10^{−6} cm³mol^{−1}. Below 40 K, the curve drops abruptly down to 1.52/1.83 cm³Kmol^{−1} at 2 K. The cause of this decline is the ZFS of the metal ions, which is also evident from the isothermal (2K) *M/Nμ_B* vs *H* curves (Fig. 8), since the values from the latter at the highest magnetic fields are far from saturation (with a measured value of 3.01/2.57 μ_B compared to the expected of 4 μ_B for *g*=2 and *S*=2). Simultaneous fitting of both sets of data using the spin Hamiltonian defined in Eq. 2, yielded the parameters *D*=6.70/−10.48 cm^{−1} and |*E*|= 0.67/0.79 cm^{−1} as well as weak intermolecular interaction constants *zJ*=0.02/0.01 cm^{−1}. In these fits, an isotropic *g* factor and a TIP value were fixed at 2.17/2.10 and 330/250 x10^{−6} cm³mol^{−1}, respectively.

$$\hat{H} = D \left(\hat{S}_z^2 - \frac{1}{3} \hat{S}^2 \right) + E \left(\hat{S}_x^2 - \hat{S}_y^2 \right) + \mu_B \hat{S} g B \quad (2)$$

For both compounds, attempts to simulate the data employing opposite signs for *D* were unsuccessful. The positive *D* value of **3** is consistent with that obtained for **2** given the large similarity of their coordination geometries.

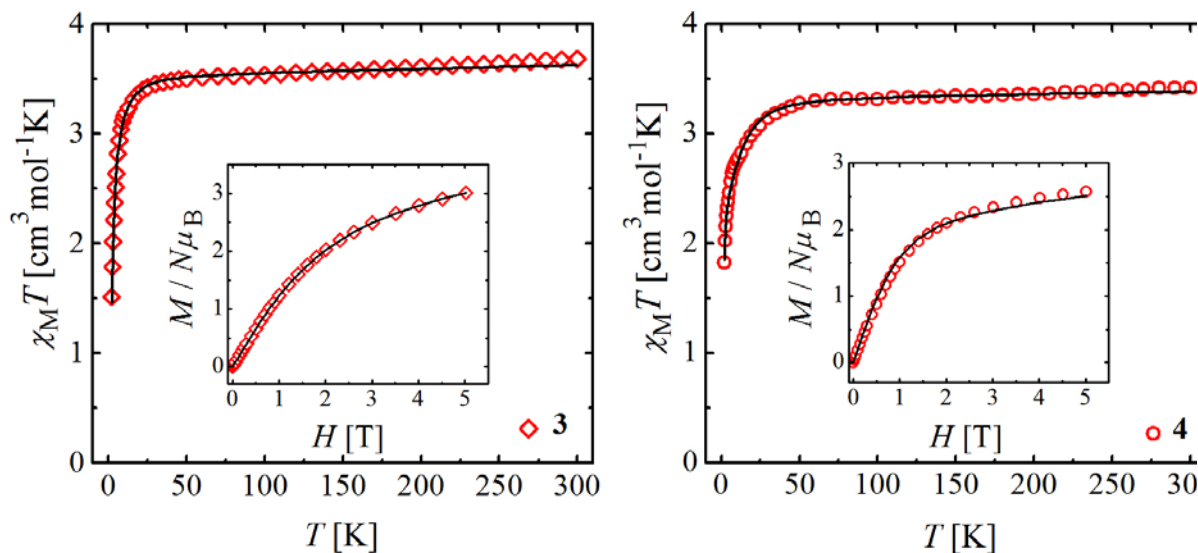


Fig. 8 $\chi_M T$ vs T and $M/N\mu_B$ vs H (inset) curves of compounds **3** and **4** with their best fit (solid line, see text for details).

Although the N_6 coordination environment around the metal centre in **4** could be appropriate for the appearance of thermally induced spin-crossover, the experimental results show that this ion centre stays trapped in the HS state, as indicated by the crystal structure (Fe–N bond lengths at 123 K $> 2.1 \text{ \AA}$). The likely explanation is that the large distortions from the ideal octahedral geometry imposed by the ligands cause the putative LS state to possess higher enthalpy than the HS state.⁷⁸ The evaluation of the local distortion from the ideal octahedron at the Fe(II) ion in **4** using the parameters Σ and Θ gave $142.4(5)^\circ$ and $503.7(10)^\circ$, respectively, which fall into the reported ranges typical for the HS state.^{79–82} These findings are also consistent with reported magnetic data for other compounds with identical coordination geometry.^{83, 84}

The possibility that compounds **2** to **4** exhibit slow relaxation of the magnetization was evaluated by means of dynamic magnetization measurements. Thus, experiments under an oscillating (AC) field of 4 Oe were performed under zero or 1000 Oe applied constant field (Figs. S5 to S7). These measurements yielded, for all three compounds, superimposed curves for the in-phase magnetic susceptibility (χ_M') and no signal for the out-of-phase component (χ_M''). The lack of single molecule magnet (SMM) behaviour for **4** (which could be anticipated, given its large and axial ZFS parameter; $D = -10.48 \text{ cm}^{-1}$) can be rationalised with the existence of significant rhombic anisotropy, which likely accelerates the quantum tunnelling of magnetization, as is known for the non-Kramers ions such as high-spin Fe(II).

Conclusions

On studying transamination reactions of sodium ferrate [$\{\text{NaFe}(\text{HMDS})_3\}_\infty$] (**1**) and its parent iron bis(amido) $\text{Fe}(\text{HMDS})_2$ with 2,2'-dipyridylamine DPA(H), three new iron complexes have been isolated and structurally defined containing the N-bridged bis(heterocycle) ligand DPA. In stoichiometrically controlled processes, reacting **1** with 1 or 3 equivalents of DPA(H) accesses the sodium ferrates $[(\text{THF})_2\cdot\text{NaFe}(\text{DPA})(\text{HMDS})_2]$ (**3**) and $[\{\text{THF}\cdot\text{NaFe}(\text{DPA})_3\}_\infty]$ (**4**) respectively. Interestingly **3** can also be prepared using an alternative co-complexation approach by combining heteroleptic $[\{\text{Fe}(\text{HMDS})(\text{DPA})\}_2]$ (**2**) with the sodium amide NaHMDS. Highlighting the coordination flexibility of this multifunctional amido ligand, structural studies revealed that while in **2** and **3** DPA adopts a *syn/syn* conformation, acting as a bridge between two metal centers, in homoleptic complex **4**, two DPA ligands present the same type of conformation, whereas a remaining DPA group binds in an *anti/anti* fashion, facilitating formation of a novel intricate polymeric chain structure. SQUID magnetisation measurements confirmed the structural findings by establishing the presence of ferromagnetically coupled (compound **2**, $J=1.01\text{ cm}^{-1}$) and isolated high-spin Fe(II) centres ($S=2$) in compounds **3** and **4**. Evaluation of the ZFS parameters yielded positive D values for the spin carriers in **2** and **3**, with similar five-coordination geometries, and negative for compound **4**, consistent with its the octahedral geometry. Despite its negative $D=-10.48\text{ cm}^{-1}$ parameter, this compound does not show slow relaxation of magnetization.

Collectively these findings advance the synthesis of alkali-metal ferrates and the understanding of the intriguing structural/reactivity/magnetic correlations in this class of mixed-metal reagents. While alkali-metal ferrates have already shown great promise in synthesis, the number of methods available to access these systems remains scarce. These findings reveal the potential that transamination approaches offer, using tris(amido) sodium ferrate **1** as a precursor, to access novel homo- and heteroleptic heterobimetallic complexes.

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