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Hydrogen bond mediated intermolecular magnetic coupling in mononuclear high spin iron(III) Schiff base complexes: synthesis, structure and magnetic study with theoretical insight†

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The crystal structure and magnetic properties of two mononuclear iron(III) Schiff base complexes, [FeL¹(NCS)₂] (**1**), HL¹ = 2-[1-[[2-[(2-aminoethyl)amino]ethyl]imino]ethyl]phenol and [FeL²(N₃)Cl] (**2**), HL² = 2-(-1-(2-(2-aminoethylamino)ethylimino)ethyl)-4-methylphenol are reported. Each complex contains a Fe(III) ion surrounded by a N₃O Schiff base ligand and two NCS⁻ ligands (in **1**) or one N₃⁻ and one Cl⁻ ligands (in **2**). The magnetic properties can be well reproduced with zero field splittings in the high spin *S* = 5/2 Fe(III) ions and weak intermolecular Fe–Fe interactions mediated by hydrogen bonds. This intermolecular antiferromagnetic interaction has been validated by using DFT calculations in complex **2**. Moreover, the interaction energies of the H-bonded dimers in both complexes have been estimated using DFT calculations and characterized using a combination of QTAIM and NCI plot computational tools. Complexes **1** and **2** constitute two rare examples of Fe(III) complexes with magnetic interactions through H-bonds.

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Introduction

The rational design and synthesis of iron(III) Schiff base complexes have attracted extensive interest due to their structural diversities and potential applications in molecular magnetism. These complexes represent an important class of spin crossover (SCO) systems because of the possibility of the co-existence of two different magnetic states under the same conditions.^{1,2} The phenomenon of spin cross-over is induced by external perturbations, such as pH, temperature, pressure, light irradiation, *etc.*^{3–7} The iron(III) sites in some heme proteins are also reported to exhibit SCO behaviors, which play key roles in their biological functions.^{8–10} Several iron(III) complexes of various geometries have been prepared using different chelating ligand,^{11–13} of which Schiff bases are special choices

due to the easy routes of their synthesis, the wide range of thermal stability and their very good chelating ability.^{14–19} Iron(III) complexes with N₃O₂-donor Schiff base (SB) ligands present a family of coordination compounds in which investigation of the impact of the structural influences on the resulting magnetic behavior, spin states and SCO properties is very appealing.^{20,21}

In the present work, two tetradentate N₃O half-salen type mono-condensed Schiff base ligands, {HL¹ = 2-[1-[[2-[(2-aminoethyl)amino]ethyl]imino]ethyl]phenol and HL² = 2-(-1-(2-(2-aminoethylamino)ethylimino)ethyl)-4-methylphenol}, have been used for the synthesis of two mononuclear iron(III) complexes, [Fe(L¹)(NCS)₂] (**1**) and [Fe(L²)(N₃)Cl] (**2**), respectively. Both complexes have been characterized by elemental and spectral analysis and the structures of the complexes have been confirmed by single crystal X-ray diffraction studies. The solid-state supramolecular interactions of both complexes have also been explored. Magnetic studies show that both complexes are essentially high spin *S* = 5/2 paramagnets with a zero field splitting and a very weak intermolecular interaction. This intermolecular interaction has been also proved by using DFT calculations and the broken symmetry approach. Moreover, the dimerization energies of the H-bonded dimers used to compute the magnetic coupling have been calculated and the interactions characterized using the quantum theory of “atoms-in-molecules” and the noncovalent interaction plot (NCI plot) index.

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Experimental

Materials

All chemicals were of reagent grade and used as purchased from Sigma-Aldrich without further purification.

Caution!!! Although no problems were encountered in this work, organic ligands in the presence of azide are potentially explosive. Only a small amount of the material should be prepared and it should be handled with care.

Preparation

Preparation of the ligands. The two N_3O tetradentate Schiff base ligands were prepared by reacting two different types of ketones (0.140 g of 2-hydroxyacetophenone or 0.150 g of 2-hydroxy-5-methylacetophenone) with 0.1 mL of diethyltri-amine in a 1 : 1 molar ratio in 15 mL of acetonitrile and refluxed during 1 hour to prepare the corresponding Schiff base ligands: 2-[1-[[2-[(2-aminoethyl)amino]ethyl]imino]ethyl]phenol (HL^1) and 2-[1-[[2-[(2-aminoethyl)amino]ethyl]imino]ethyl]-4-methylphenol (HL^2). The ligands, HL^1 and HL^2 , were not isolated and the acetonitrile solutions of these ligands were used for the syntheses of complexes **1** and **2**, respectively.

Preparation of the complexes

$[Fe(L^1)(NCS)_2]$ (**1**). An acetonitrile solution (10 mL) of iron(III) perchlorate hexahydrate (~1 mmol, 0.360 g) was added to the previously prepared acetonitrile solution of the Schiff base ligand (HL^1) under refluxing condition. An aqueous solution (15 mL) of sodium thiocyanate (2 mmol, 0.180 g) was then added to the resulting solution and refluxed further for *ca.* 30 min. The solution was then filtered and kept in open atmosphere at room temperature for 2 days. A dark crystalline product was collected by filtration. X-ray quality single crystals were collected from this crystalline product.

Yield: 0.282 g, ~72% (based on iron). Anal. calc. for $C_{14}H_{18}FeN_5OS_2$ (F.W. 392.32 g mol⁻¹): C, 42.86; H, 4.62; N, 17.85%. Found: C, 42.7; H, 4.5; N, 17.9%. FT-IR (KBr, cm⁻¹): 3225 (ν_{N-H}); 2884–2955 (ν_{C-H}); 2075 (ν_{NCS}); 2042 (ν_{NCS}); 1580 ($\nu_{C=N}$). λ_{max} (nm) [ϵ_{max} (M⁻¹ cm⁻¹)] (acetonitrile): 515 (2 × 10³), 329 (5 × 10³), 268 (1.3 × 10⁴), 230 (2.4 × 10⁴).

$[Fe(L^2)(N_3)Cl]$ (**2**). An acetonitrile solution (10 mL) of iron(III) chloride hexahydrate (1 mmol, 0.170 g) was added to the acetonitrile solution of the Schiff base ligand (HL^2) under refluxing condition. An aqueous solution (5 mL) of sodium azide (1 mmol, 0.070 g) was then added to the resulting solution and refluxed further for *ca.* 30 min. The solution was then filtered and kept in open atmosphere at room temperature for 2 days. A dark crystalline product was collected by filtration. X-ray quality single crystals were collected from this crystalline product.

Yield: 0.261 g, ~71% (based on iron). Anal. calc. for $C_{13}H_{20}ClFeN_6O$ (F.W. 367.65 g mol⁻¹): C, 42.47; H, 5.48; N, 22.86%. Found: C, 42.3; H, 5.3; N, 22.9%. FT-IR (KBr, cm⁻¹): 3225 (ν_{N-H}); 2893–2936 (ν_{C-H}); 2041 (ν_{N_3}); 1578 ($\nu_{C=N}$). λ_{max} (nm) [ϵ_{max} (lit mol⁻¹ cm⁻¹)] (acetonitrile): 520 (2.5 × 10³), 330 (5.4 × 10³), 269 (1.3 × 10⁴), 231 (2.5 × 10⁴).

Physical measurements

Elemental analysis (C, H and N) was performed using a Perkin-Elmer 240C elemental analyser. IR spectrum in KBr (4500–500 cm⁻¹) was recorded with a Perkin-Elmer Spectrum Two spectrophotometer. Electronic spectra in DMF were recorded on a Shimadzu UV-1700 spectrophotometer. Variable temperature magnetic susceptibility measurements were performed on polycrystalline samples of both complexes (with masses of 37.447 and 36.265 mg, for **1** and **2**, respectively) with a Quantum Design MPMS-XL-5 SQUID susceptometer in the temperature range 2–400 K with an applied magnetic field of 0.1 T. The data were corrected for the sample holder and for the diamagnetic contribution of the salts using Pascal's constants ($\chi_{dia} = -197.3 \times 10^{-6}$ and -204.7×10^{-6} cm³ mol⁻¹ for **1** and **2**, respectively).²²

X-ray crystallography

Suitable crystals of both complexes were used for data collection using a 'Bruker D8 QUEST area detector' diffractometer equipped with graphite-monochromated Mo K_{α} radiation ($\lambda = 0.71073$ Å). Molecular structures were solved by direct methods and refined by full-matrix least squares on F^2 using the SHELXL-14 package.^{23a} Non-hydrogen atoms were refined with anisotropic thermal parameters. The hydrogen atoms attached to nitrogen atoms were located by difference Fourier maps and were kept at fixed positions. All other hydrogen atoms were placed in their geometrically idealized positions and constrained to ride on their parent atoms. Multi-scan empirical absorption corrections were applied to the data using the program SADABS.^{23b} In complex **2** there are two disordered chlorine atoms taking two close positions which were refined with occupation factors x and $1 - x$, with x refining to 0.52(3).

Table 1 Crystal data and refinement details of complexes **1** and **2**

Complex	1	2
Formula	$C_{14}H_{18}FeN_5OS_2$	$C_{13}H_{20}ClFeN_6O$
Formula weight	392.32	367.65
Temperature (K)	273	273
Crystal system	Monoclinic	Monoclinic
Space group	$P2_1/n$	$P2_1/n$
a (Å)	11.6962(10)	11.143(3)
b (Å)	12.7639(10)	13.276(3)
c (Å)	12.7562(10)	11.981(3)
α (°)	90	90
β (°)	112.592(2)	108.225(6)
γ (°)	90	90
Z	4	4
d_{cal} (g cm ⁻³)	1.482	1.451
μ (mm ⁻¹)	1.105	1.064
$F(000)$	812	764
Total reflection	24 912	16 196
Unique reflections	3891	2923
Observe data [$I > 2\sigma(I)$]	3167	2234
$R(int)$	0.036	0.074
R_1, wR_2 (all data)	0.0498, 0.0371	0.1011, 0.0787
R_1, wR_2 [$I > 2\sigma(I)$]	0.1144, 0.1053	0.2336, 0.2050



Details of crystal data and refinement are given in Table 1. Important bond lengths and bond angles are gathered in Tables 2 and S1 (ESI†) respectively.

Theoretical methods

All calculations included in this study have been performed using the ORCA program, version 4.1.2.^{24a} For the calculation of the magnetic coupling constants we have used the TPPSh functional^{24b} combined with the def2-TZVP basis set.^{24c} This level of theory has been recommended for the calculation of J values accurately.^{24d} For the interaction energies, we have used the B3LYP/def2-TZVP level of theory. The B3LYP/def2-TZVP wave function obtained from the ORCA program was used to generate a spin density cube that was represented using the Gaussview software.^{24e} The QAIM/NCI plot representation has been performed at the same level using the ORCA wave function and the AIMAll program.^{24f}

Results and discussion

Synthesis

Schiff base ligands, HL¹ and HL², were prepared by 1 : 1 condensation of diethyltriamine with 2-hydroxy-acetophenone and 2-hydroxy-5-methylacetophenone in acetonitrile, respectively.^{25b} Acetonitrile solution of both ligands, HL¹ and HL², were reacted with iron(III) perchlorate and iron(III) chloride hydrate under reflux. Around 2 mmol of sodium thiocyanate was used as coligand in complex **1** and 1 mmol of sodium azide was used for complex **2**. Both complexes are stable at room temperature. The synthetic routes for both complexes have been shown in Scheme 1.

Crystal structure of [Fe(L¹)(NCS)₂] (**1**)

Complex **1** crystallizes in the monoclinic space group $P2_1/n$. The asymmetric unit consists of a discrete mononuclear unit [Fe(L¹)(NCS)₂]. A perspective view of this complex with selected atom-numbering scheme is shown in Fig. 1. The iron(III) center, Fe1, is octahedrally coordinated by a deprotonated Schiff base ligand and two terminal thiocyanate group. The equatorial plane consists of two amine nitrogen atoms, N1 and N2, one phenoxido oxygen atom, O1 of a deprotonated Schiff base ligand (L¹)⁻ and one nitrogen atom, N4 of a terminal thiocyanate group. The axial positions are occupied by one imine nitrogen atom, N3 of the deprotonated Schiff base and one nitrogen atom, N5 of another terminal thiocyanate group. The deviations of the coordinating atoms N1, N2, N4 and O1 from

the least-square mean plane through them are $-0.175(3)$ Å, $0.198(2)$ Å, $-0.152(2)$ Å and $-0.184(16)$ Å, respectively, and that of Fe1 from the same plane is $-0.055(4)$ Å. The trans angles, N1–Fe1–N4, N2–Fe1–O1 and N3–Fe1–N5, around Fe1 are $165.1(1)^\circ$, $163.15(9)^\circ$ and $175.36(9)^\circ$, respectively (Table S1, ESI†). Both five-membered chelate rings [Fe1–N1–C1–C2–N2] and [Fe1–N2–C3–C4–N3] have half-chair conformation, with puckering parameters,²⁶ $q = 0.489(3)$ Å, $\varphi = 66.1(3)^\circ$ and $q = 0.465(3)$ Å, $\varphi = 53.4(3)^\circ$, respectively.

Complex **1** presents some interesting non-covalent interactions that give rise to a supramolecular network. Two types of non-covalent interactions have been observed: (i) hydrogen bonding and (ii) C–H $\cdots\pi$ interactions. There are two unconventional hydrogen bonding interactions in **1**. On one side, the hydrogen atom H1N, attached to N1, is involved in intermolecular hydrogen bonding with the sulphur atom S2. On the other side, the hydrogen atom H2N, attached to N2, is involved in intermolecular hydrogen bonding with the sulphur atom S1. These inter-molecular hydrogen bonding interactions originate a one-dimensional array as shown in Fig. 2. The second type of interaction ((C–H $\cdots\pi$) is formed between the hydrogen atom H9, attached to C3, and the symmetry related $(1/2 + x, 3/2 - y, 1/2 + z)$ chelate ring, Cg4 [C7–C8–C9–C10–C11–C12] and between the atom H14, attached to C6, and the symmetry related $(1 - x, 1 - y, -z)$ chelate ring, Cg4 [C7–C8–C9–C10–C11–C12]. These intermolecular C–H $\cdots\pi$ interactions give rise to layers as shown in Fig. 3. Details of the geometric features of the C–H $\cdots\pi$ interactions are given in Table 3. Details of H-bonding interactions are given in Table 4.

Crystal structure of [Fe(L²)(N₃)Cl] (**2**)

Complex **2** crystallizes in the monoclinic space group $P2_1/n$. The asymmetric unit consists of a discrete mononuclear unit [Fe(L²)(N₃)Cl]. A perspective view of this complex with selected atom-numbering scheme is shown in Fig. 4.

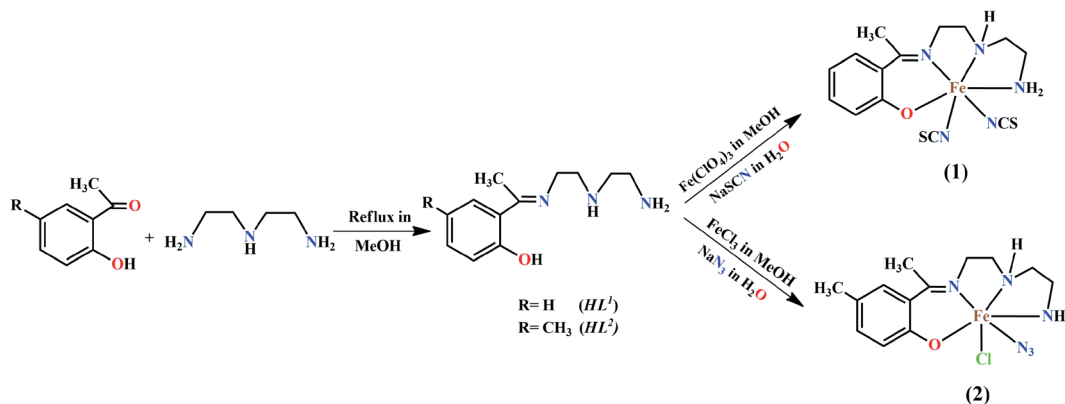
The iron(III) center, Fe1, is octahedrally coordinated by a deprotonated Schiff base ligand, a Cl atom and a terminal azide ligand. The equatorial plane contains two amine nitrogen atoms, N1 and N2, one phenoxido oxygen atom, O1, of a deprotonated Schiff base ligand (L²)⁻ and one chlorine atom, Cl1. The axial positions are occupied by one imine nitrogen atom, N3, of the deprotonated Schiff base ligand (L²)⁻ and by one nitrogen atom, N4, of a terminal azide group. The deviations of the coordinated atoms N1, N2, O1 and Cl1 from the least-square mean plane through them are $0.216(8)$ Å, $-0.225(5)$ Å, $-0.217(5)$ Å and $0.170(3)$ Å, respectively. Fe1 deviation from the same plane is $-0.055(9)$ Å. The trans angles N1–Fe1–Cl1, N2–Fe1–O1 and N3–Fe1–N4, around Fe1 are $167.3(2)^\circ$, $162.3(2)^\circ$ and $172.5(2)^\circ$, respectively (Table S1†). Both saturated five-membered chelate rings [Fe1–N1–C1–C2–N2] and [Fe1–N2–C3–C4–N3] have envelop conformation, with puckering parameters,²⁶ $q = 0.485(8)$ Å, $\varphi = 245.1(8)^\circ$ and $q = 0.471(6)$ Å, $\varphi = 227.1(7)^\circ$, respectively.

Complex **2** shows two types of hydrogen bonds: on one side, hydrogen atom H2, attached to N1, is involved in intermolecular hydrogen bonding with Cl1 and on the other side,

Table 2 Selected bond lengths (Å) in complexes **1** and **2**

Complex	1	2
Fe1–O1	1.8602(19)	1.868(4)
Fe1–N1	2.164(3)	2.177(7)
Fe1–N2	2.162(2)	2.174(5)
Fe1–N3	2.143(2)	2.162(5)
Fe1–N4	2.067(2)	2.070(7)
Fe1–N5	2.036(3)	—
Fe1–Cl1	—	2.367(4)





Scheme 1 Synthetic route for ligand and complexes 1 and 2.

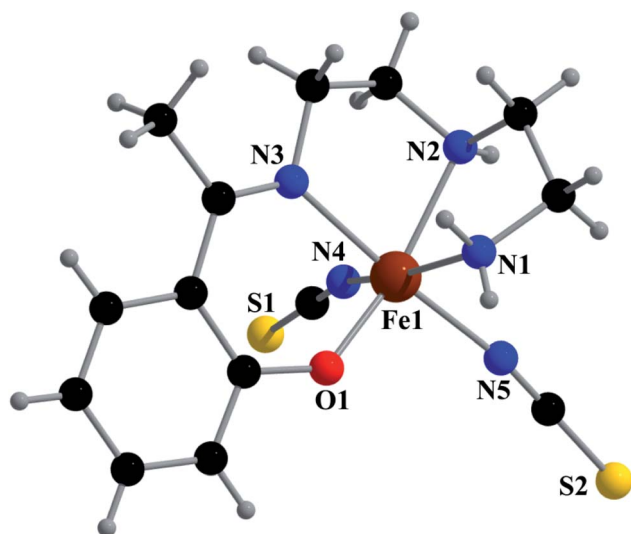


Fig. 1 Perspective view of complex 1 with selected numbering scheme.

hydrogen atom H2N, attached to N2, is involved in intermolecular hydrogen bonding with N6. These inter-molecular hydrogen bonding interactions give rise to a two-dimensional structure as shown in Fig. 5. Details of the geometric features of these hydrogen bonding interactions are given in Table 4.

Magnetic properties

The thermal variation of the product of the molar magnetic susceptibility times the temperature ($\chi_m T$) per Fe(III) ions in both complexes is, as expected, very similar. They show room temperature values of *ca.* $4.4 \text{ cm}^3 \text{ K mol}^{-1}$ ($\mu_{\text{eff}} \approx 5.93 \mu_B$), very close to the spin only value for a $S = 5/2$ Fe(III) ion. When the temperature decreases, $\chi_m T$ remains constant in the range down to *ca.* 30 K in complex 1 and *ca.* 80 K in complex 2 (Fig. 6). Below these temperatures $\chi_m T$ shows a progressive decrease to reach values of *ca.* 3.1 and $1.2 \text{ cm}^3 \text{ K mol}^{-1}$ in 1 and 2, respectively. This behaviour indicates that both complexes behave essentially as paramagnets with a weak antiferromagnetic intermolecular interaction and/or a zero Field Splitting

(ZFS) of the $S = 5/2$ spin ground state. Accordingly, we have fit the magnetic properties with a $S = 5/2$ monomer including a ZFS and an inter-molecular exchange interaction (zJ) using the PHI software.²⁷ This model reproduces very satisfactorily the magnetic properties of both complexes in the whole temperature range with $g = 2.043(6)$, $|D| = 1.23(3) \text{ cm}^{-1}$ and $zJ = -0.03(1) \text{ cm}^{-1}$ in 1 and $g = 2.126(2)$, $|D| = 1.6(1) \text{ cm}^{-1}$ and $zJ = -0.23(1) \text{ cm}^{-1}$ in 2 (solid lines in Fig. 6). Note that the sign of D cannot be determined with powder magnetic measurements.

As expected, both complexes show very weak antiferromagnetic inter-molecular exchange interactions. The larger absolute value found in complex 2 agrees with the most important decrease observed in $\chi_m T$ in this complex and can be easily explained by the stronger H-bonds observed in complex 2, with much shorter D...A distances (3.062 and 3.391 \AA in 2 compared to 3.551 and 3.787 \AA in 1) and also shorter H...A distances (2.21 and 2.48 \AA in 2 compared to 2.85 and 2.87 \AA in 1, Table 4). On the other side, the D values are within the normal range observed for other high spin $S = 5/2$ Fe(III) monomers.²⁸

Note that, although there are a few examples of ferromagnetic interactions and even long range ferro- and antiferromagnetic ordering, in most cases the magnetic interactions through hydrogen bonds are weak and antiferromagnetic (Table 5). Theoretical calculations performed by several groups also confirm this experimental observation.²⁹ Interestingly, most of the reported complexes^{30–53} showing magnetic interactions through H-bonds are Cu(II) complexes with Cu–O–H...O–Cu pathways (Table 5). As can be seen in Table 5, there are very few examples containing other metals as Ni(II), Mn(II), Mn(III) and Fe(III). Therefore, complexes 1 and 2 are expected to show very weak antiferromagnetic interactions through the H-bonds and C–H– π interactions, in agreement with the experimental data.

In order to verify this explanation (stronger H-bonded dimer in 2) we have performed a DFT study using the RIJCOSX-B3LYP/def2-TZVP level of theory. Fig. 7 shows the combined QTAIM/NCI Plot analysis of the H-bonded dimers commented above (see Fig. 2 and 5 for details) and the dimerization energies. In the dimer of 1, the QTAIM shows an intricate combination of bond critical points and bond paths connecting the monomers. That is, in addition to the N–H...S H-bonds (marked with



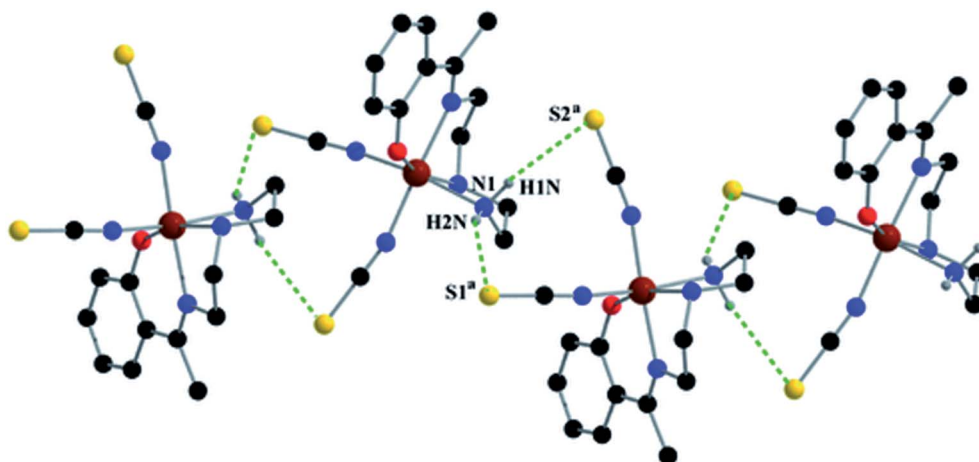


Fig. 2 1D supramolecular array in **1** formed by N–H...S hydrogen bonding interactions. Symmetry transformations: $a = 1/2 - x, 1/2 + y, 3/2 - z$.

Table 3 Hydrogen bond distances (Å) and angles (°) of complexes **1** and **2**^a

Complex	D–H...A	D–H	H...A	D...A	∠D–H...A
1	N(1)–H(1N)–S(2) ^a	0.96(3)	2.85(3)	3.787(3)	165(3)
	N(1)–H(2N)–S(1) ^a	0.76(4)	2.87(4)	3.551(4)	151(4)
2	N(2)–H(2)–Cl(1) ^b	0.9800	2.48(5)	3.391(5)	154(1)
	N(1)–H(2N)–N(6) ^c	0.91(8)	2.21(8)	3.062(9)	155(7)

^a D = donor; H = hydrogen; A = acceptor. Symmetry transformations: $a = 1/2 - x, 1/2 + y, 3/2 - z$; $b = 1 - x, 1 - y, -z$ and $c = 3/2 - x, -1/2 + y, 1/2 - z$.

Table 4 Geometric features (distances in Å and angles in °) of the C–H...π interactions in **1**^a

C–H...Cg(ring)	H...Cg (Å)	C–H...Cg (°)	C...Cg (Å)
C3–H9...Cg(4) ^d	3.00	138	3.773(3)
C6–H14...Cg(4) ^b	2.98	143	3.787(4)

^a Symmetry transformations: $d = 1/2 + x, 3/2 - y, 1/2 + z$; $b = 1 - x, 1 - y, -z$; Cg(4) = centre of gravity of the ring [C7–C8–C9–C10–C11–C12].

asterisk symbols in Fig. 7a), the combined QTAIM/NCI plot analysis also reveals the existence of several C–H...S and C–H...N H-bonds (marked with arrows). All these H-bonds are

characterized by a bond critical point (CP) and bond path connecting the H-atom to the electron rich atom (S, N). These interactions are further characterized by green NCI plot iso-surfaces located between the interacting atoms. The green colour is indicative of weak interaction,⁵⁴ which is further

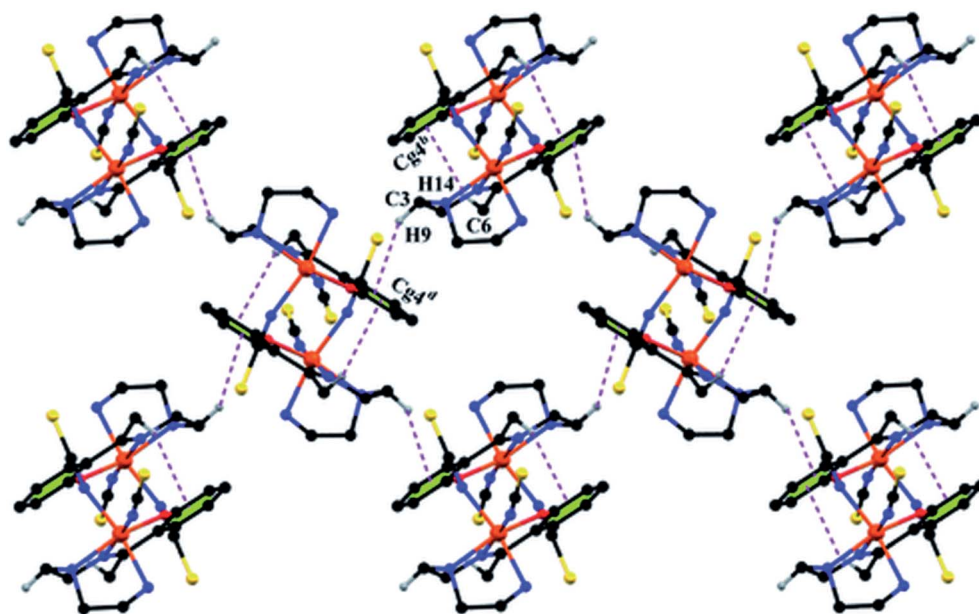


Fig. 3 Supramolecular 2D structure in **1** formed by intermolecular C–H...π interactions. Symmetry transformations: $b = 1 - x, 1 - y, -z$; $d = 1/2 + x, 3/2 - y, 1/2 + z$.



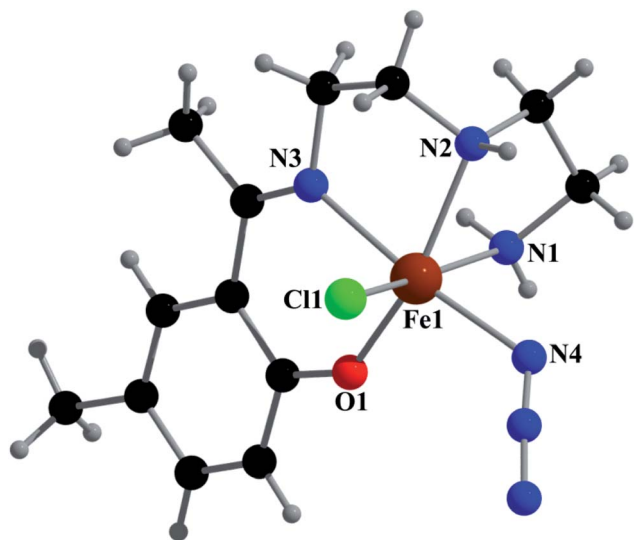


Fig. 4 Perspective view of complex 2 with selected atom numbering scheme.

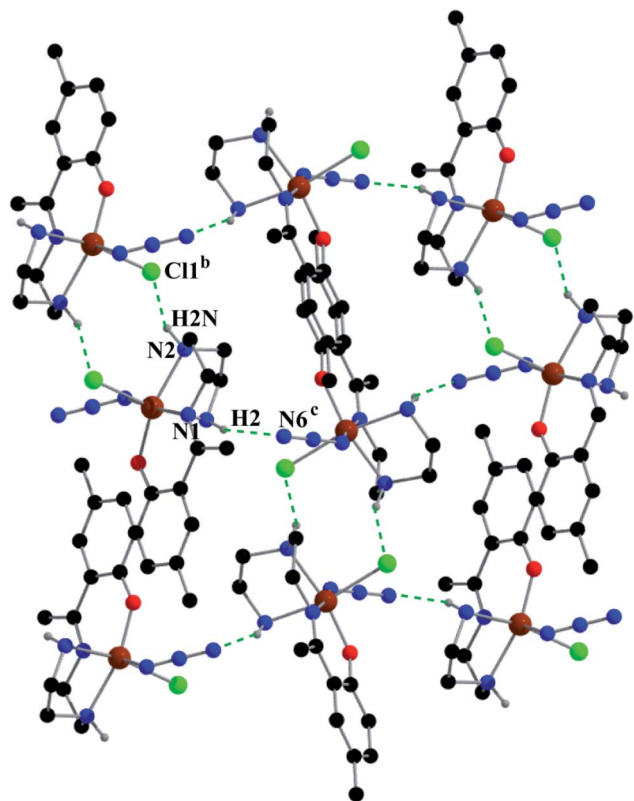


Fig. 5 Supramolecular 2D structure in 2 formed by intermolecular hydrogen bonding interactions. Symmetry transformations: $b = 1 - x, 1 - y, -z$; $c = 3/2 - x, -1/2 + y, 1/2 - z$.

validated by the modest dimerization energy ($\Delta E_1 = -8.3 \text{ kcal mol}^{-1}$) obtained for 1, taking into consideration the number of H-bonds that are established between the monomers. The QTAIM distribution of bond CPs and bond paths in complex 2 (see Fig. 7b) shows two symmetrically related bond

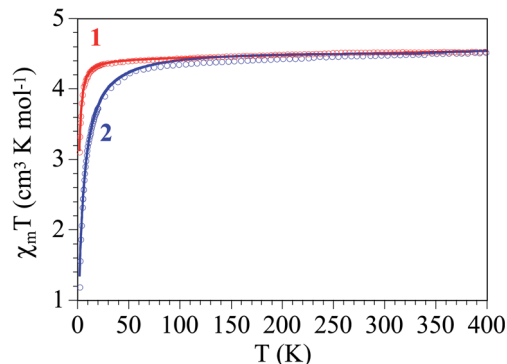


Fig. 6 Thermal variation of $\chi_m T$ for complexes 1 and 2. Solid lines are the best fit to the model (see text).

CPs and bond paths that connect the N-H groups to the chlorido ligands. Moreover, two additional and symmetrically related bond CPs and bond path connect two C-H bonds two the N-atoms of the azido ligands. The N-H \cdots Cl bond are also characterized by bluish isosurfaces, indicated moderately strong H-bonds. Consequently, the H-bonded dimer of complex 2 presents a stronger interaction ($\Delta E_2 = -25.6 \text{ kcal mol}^{-1}$) compared to 1, in agreement with the stronger antiferromagnetic coupling constant observed experimentally.

We have also computed the theoretical zJ value using the broken symmetry (BS) approach and the equation proposed by Ruiz and coworkers ($J = -(E_{\text{HS}} - E_{\text{BS}})/(S_{\text{max}}(S_{\text{max}} + 1))$), where the energies of the high spin (HS) and low spin (BS) states are used,⁵⁵ as implemented in the ORCA package. In case of 1, the experimental J value is too small to be reliably reproduced by the DFT calculations (-0.03 cm^{-1} is within the accuracy of the DFT method). Therefore, we have focused the DFT study on the calculation of complex 2. Interestingly, the calculation predicts a zJ value of -0.18 cm^{-1} , which is in reasonable agreement with the experimental value (-0.23 cm^{-1}) and confirms the very weak antiferromagnetic coupling of the Fe metal centers in the H-bonded dimer of 2. However, the $|D|$ value is somewhat underestimated by the DFT calculation (0.53 cm^{-1}). The spin density plot of the high spin configuration of the dimer of 2 is represented in Fig. 8. The observed spherical distribution of the spin density around the Fe atoms is typical of a d^5 configuration with a single occupation of each d orbital. It can be also observed that some spin is delocalized onto the atoms of the ligands directly bonded to the metal. It is also worth mentioning that some of the delocalized spin density is located to the heteroatoms involved in the strong N-H \cdots Cl hydrogen bonds.

Conclusion

We have shown a convenient way to prepare two high spin Fe(III) complexes with two different Schiff base ligands and different pseudohalides as counterions. The magnetic properties of both complexes can be very well reproduced with a weak intermolecular antiferromagnetic interaction (mediated through the inter-molecular H-bonds) and a zero field splitting of the



Table 5 Structural and magnetic parameters of compounds showing magnetic coupling through hydrogen bonds^b

Complex	CSD code/CCDC no.	J (cm ⁻¹)	M	H-Bond type	D-H...A (Å)	Ref.
[Cu ₂ (L ^a) ₂]	—	≈ -100	Cu(II)	O-H...O	2.29	30
[Cu ₂ (L ^a) ₂]	HEAICU10	-94	Cu(II)	O-H...O (x2)	2.31	31
				O-H...O (x2)	2.33	
[Cu ₃ (L ^b) ₂ (C ₆ H ₅ COO) ₂ Cl]Cl	DEPSAT	-0.4(2)	Cu(II)	N-H...Cl	3.163	32
[Cu ₂ (μ ₂ -H ₂ O)(L ^c) ₂ (H ₂ O) ₂](ClO ₄) ₂ ·2H ₂ O	KEDNIR	-13.23(1)	Cu(II)	O-H...O (x2)	2.627	33
[Cu ₂ (L ^d) ₂ (H ₂ O) ₂ (ClO ₄)](ClO ₄)·H ₂ O	FUTCON	-19.8(2)	Cu(II)	O-H...O	2.685	34
				O-H...O	2.601	
[Cu(L ^e) ₂ (H ₂ O)]	BEYRAY	+0.23(1)	Cu(II)	O-H...O (x2)	2.695	35
[Zn ^{II} (H ₂ O) ₆][Cu ^{II} (L ^f) ₂ (H ₂ O) ₂]	HULMOQ	-0.220(1)	Cu(II)	O-H...O (x2)	2.771	36
[Cu ₂ (L ^g) ₂ (H ₂ O) ₂]·2H ₂ O	MATLOJ	-1.6(1)	Cu(II)	O-H...O (x2)	2.721	37
[Cu(L ^h)(H ₂ O)(NO ₃)]	NUQKOZ01	-26.6	Cu(II)	O-H...O (x2)	2.685	38
[Cu(L ⁱ) ₂ (H ₂ O) ₂] _n	FAHNAE	-3.26	Cu(II)	O-H...O (x2)	2.71	39
[Cu(L ^j)(H ₂ O)]·4H ₂ O	SAGLAC	-2.19(2)	Cu(II)	O-H...O (x2)	2.757	40
[NiCl ₂ (L ^k) ₂]	FUJQOQ	-1.72(1)	Ni(II)	N-H...Cl (x2)	3.256	41
		-0.22(1)		N-H...Cl	3.275	
[Cu(HL ^l)(L ^l)(H ₂ O)] ₂ NO ₃	AETCUB	-70	Cu(II)	O-H...O (x2)	2.516	42
[Cu(HL ^m)(L ^m) ₂ (NO ₃) ₂]	AETCUA	-56	Cu(II)	O-H...O (x2)	2.452	42
				O-H...O (x2)	2.434	
[{Cu(H ₂ L ⁿ)}{Cu(L ⁿ)}]BF ₄	ODALAG	-7	Cu(II)	O-H...O (x2)	2.44	43
				O-H...O (x2)	2.66	
[{Cu(H ₂ L ⁿ)} ₂](BF ₄) ₂	ODALEK	-21	Cu(II)	O-H...O (x2)	2.60	43
[Cu(HL ^o)(L ^o)]PF ₆	MASQIJ	-9.91(2)	Cu(II)	O-H...O	2.412	44
[Cu(HL ^p)(L ^p)]BF ₄ ·2H ₂ O	YUKCOX	-23.1	Cu(II)	O-H...O	2.425	45
[Cu(L ^q) ₂ (L ^q)(H ₂ O) ₂]	BUQLIJ	-6.25	Cu(II)	O-H...O (x2)	2.692	46
<i>cis</i> -[Cu(L ^s) ₂ (H ₂ O) ₂]	NEDPAO	-3.0	Cu(II)	O-H...O (x2)	2.767	47
				O-H...O (x2)	2.713	
<i>trans</i> -[Cu(L ^s) ₂ (H ₂ O) ₂]·H ₂ O	PAXTUE	-4.0	Cu(II)	O-H...O (x2)	2.651	47
[Ni ₃ (L ^t)(CO ₃)(H ₂ O)(py) ₇]	GIDNAK	-39.6	Ni(II)	O-H...O (x2)	2.71	48
					2.73	
[{Mn(bpy)(H ₂ O)}(L ^u) ₂ (μ-O){Mn(bpy)(ClO ₄)}]ClO ₄	AGOJOY	-9.2	Mn(II)	O-H...O	2.074	49
					2.797	
[{Mn(bpy)(H ₂ O)}(L ^u) ₂ (μ-O){Mn(bpy)(NO ₃)}]NO ₃	AGOJUE	-27.3	Mn(II)	O-H...O	2.746	49
					2.843	
[Fe(L ^v)Cl(H ₂ O)]·MeOH	AZOXAO	$T_N = 3.2$ K	Fe(III)	O-H...O	2.582	50
				O-H...O	2.610	
				O-H...O	2.827	
				N-H...O	2.901	
[(Ni(L ^w) ₂) ₃ (Fe(CN) ₆) ₂]·7H ₂ O	ROQCAB	$T_C = 23$ K	Fe ^{III} ₃ Ni ^{II} ₂	O-H...N (x4)	2.76–3.09	51
				O-H...O (x4)	2.78–3.08	
[{Mn(OH)(OAc) ₂ ·AcOH·H ₂ O}] _n	HUWHOW	$T_N = 6.1$ K	Mn(III)	O-H...O	2.550	52
				O-H...O	2.691	53
[FeL ¹ (NCS) ₂]	2036380	-0.03(1) ^a	Fe(III)	N-H...S (x2)	3.551	This work
					3.787	
[FeL ² (N ₃)Cl]	2036381	-0.23(1) ^a	Fe(III)	N-H...Cl	3.391	This work
				N-H...N	3.062	

^a A zero field splitting of the Fe(III) ion is included in the fit, resulting in reduced J value. ^b L^{a-w} = ligand names have been given in Table S2 (ESI).

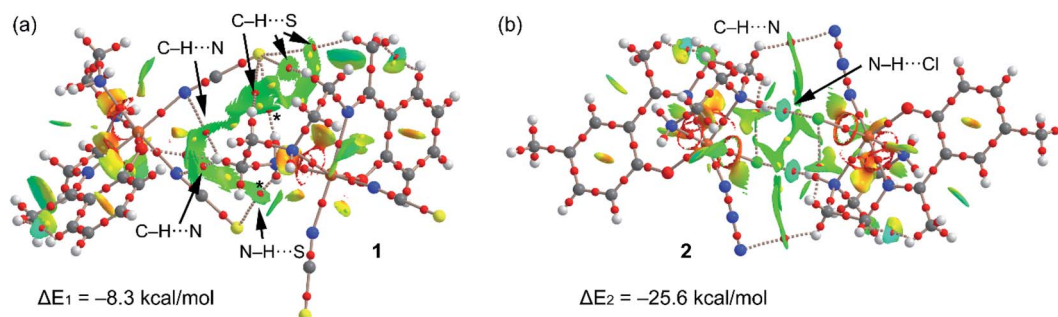


Fig. 7 Combined QTAIM (bond CPs in red and ring CP in yellow) and NCIPlot analyses of the H-bonded dimers of **1** (a) and **2** (b) at the RIJCOSX-B3LYP/def2-TZVP level of theory. The dimerization energies are also indicated.



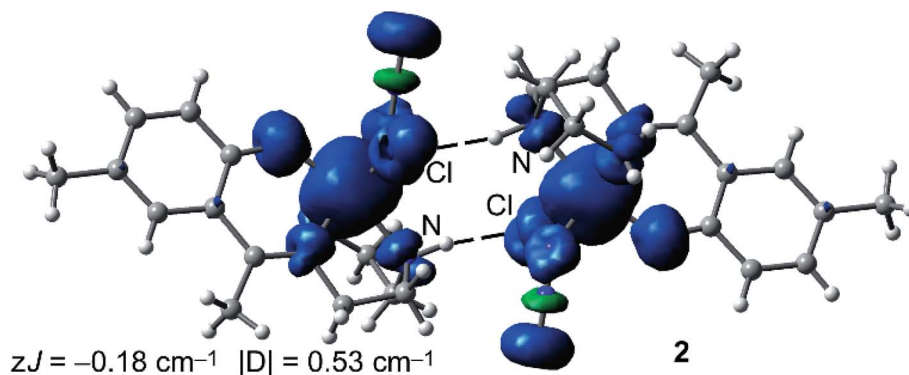


Fig. 8 Spin density plot (isosurface = 0.004 a.u.) of the high spin configuration of the H-bonded dimer of **2** at the RIJK-TPSSH/def2-TZVP level of theory.

high spin $S = 5/2$ ground state. The DFT study shows that H-bonded dimer of complex **2** is significantly stronger than **1**, in agreement with its larger antiferromagnetic coupling zJ value. The antiferromagnetic coupling in **2** has been confirmed by DFT calculations and rationalized by the spin density plot.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 *Top. Curr. Chem.*, ed. P. Gütllich and H. A. Goodwin, 2004, pp. 233–235.
- 2 S. Jana, A. Bhattacharyya, B. N. Ghosh, K. Rissanen, S. Herrero, R. Jiménez-Aparicio and S. Chattopadhyay, *Inorg. Chim. Acta*, 2013, **453**, 715–723.
- 3 H. A. Goodwin, *Coord. Chem. Rev.*, 1976, **18**, 293–325.
- 4 E. König, *Prog. Inorg. Chem.*, 1987, **35**, 527–622.
- 5 P. Gütllich, Y. Garcia and T. Woike, *Coord. Chem. Rev.*, 2001, **219–221**, 839–879.
- 6 P. Gütllich, A. Hauser and H. Spiering, *Angew. Chem., Int. Ed.*, 1994, **33**, 2024–2054.
- 7 P. Gütllich, Y. Garcia and H. A. Goodwin, *Chem. Soc. Rev.*, 2000, **29**, 419–427.
- 8 A. H. Ewald, R. L. Martin, I. G. Ross and A. H. White, *Proc. R. Soc. London, Ser. A*, 1984, **280**, 235–257.
- 9 G. Harris, *Theor. Chim. Acta*, 1966, **5**, 379–397.
- 10 M. Zerner, M. Gouterman and H. Kobayashi, *Theor. Chim. Acta*, 1966, **6**, 363–400.
- 11 T. Basak, K. Ghosh, C. J. Gómez-García and S. Chattopadhyay, *Polyhedron*, 2018, **146**, 42–54.
- 12 S. Jana, S. Chatterjee and S. Chattopadhyay, *Inorg. Chim. Acta*, 2012, **48**, 189–198.
- 13 K. Santo, M. Hirotsu and I. Kinoshita, *Dalton Trans.*, 2015, **44**, 4155–4166.
- 14 T. Basak, D. Das, P. P. Ray, S. Banerjee and S. Chattopadhyay, *CrystEngComm*, 2020, **22**, 5170–5181.
- 15 T. Basak, A. Frontera and S. Chattopadhyay, *Inorg. Chim. Acta*, 2021, **516**, 120081.
- 16 K. C. Gupta and A. K. Sutar, *Coord. Chem. Rev.*, 2008, **252**, 1420–1450.
- 17 T. Basak, A. Frontera and S. Chattopadhyay, *Dalton Trans.*, 2020, **22**, 5731.
- 18 T. Basak, K. Ghosh and S. Chattopadhyay, *Polyhedron*, 2018, **146**, 81–92.
- 19 M. Orío, O. Jarjayes, H. Kanso, C. Philouze, F. Neese and F. Thomas, *Angew. Chem., Int. Ed.*, 2010, **49**, 4989–4992.
- 20 C. Faulmann, K. Jacob, S. Dorbes, S. Lampert and I. Malfant, *Inorg. Chem.*, 2007, **46**, 8548–8559.
- 21 C. Faulmann, J. Chahine, L. Valade, G. Chastanet, J.-F. Létard and D. de Caro, *Eur. J. Inorg. Chem.*, 2013, 1058–1067.
- 22 G. A. Bain and J. F. Berry, *J. Chem. Educ.*, 2008, **85**, 532–536.
- 23 (a) G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122; (b) G. M. Sheldrick, *SADABS, V2014/5, Software for Empirical Absorption Correction*, University of Göttingen, Institute für Anorganische Chemie der Universität Göttingen Germany, 1999–2003.
- 24 (a) F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2017, **8**, e1327; (b) V. N. Staroverova and G. E. Scuseria, *J. Chem. Phys.*, 2003, **119**, 12129–12137; (c) A. Schäfer, C. Huber and R. Ahlrichs, *J. Chem. Phys.*, 1994, **100**, 5829–5835; (d) M. Orío, D. A. Pantazis, T. Petrenko and F. Neese, *Inorg. Chem.*, 2009, **48**(15), 7251–7260; (e) R. Dennington, T. Keith and J. Millam, *GaussView, Version 6.1.1*, Semichem Inc., Shawnee Mission, KS, 2019; (f) T. A. Keith, *AIMAll (Version 19.10.12)*, TK Gristmill Software, Overland Park KS, USA, 2019, <http://aim.tkgristmill.com>.



- 25 (a) P. Masarova, P. Zoufaly, J. Moncol, I. Nemeč, J. Pavlik, M. Gembický, Z. Travnicek, R. Boca and I. Salitros, *New J. Chem.*, 2015, **39**, 508–519; (b) T. Basak, D. Das, P. P. Ray, S. Banerjee and S. Chattopadhyay, *CrystEngComm*, 2020, **22**, 5170–5181; (c) T. Basak, A. Frontera and S. Chattopadhyay, *CrystEngComm*, 2020, **22**, 5731–5742.
- 26 D. Cremer and J. A. Pople, *J. Am. Chem. Soc.*, 1975, **97**, 1354–1358.
- 27 N. F. Chilton, R. P. Anderson, L. D. Turner, A. Soncini and K. S. Murray, *J. Comput. Chem.*, 2013, **34**, 1164–1175.
- 28 R. Boca, *Coord. Chem. Rev.*, 2004, **248**, 757–815.
- 29 (a) N. A. G. Bandeira and B. Le Guennic, *J. Phys. Chem. A*, 2012, **116**, 3465–3473; (b) C. Desplanches, E. Ruiz, A. Rodríguez-Forteza and S. Alvarez, Exchange Coupling of Transition-Metal Ions through Hydrogen Bonding: A Theoretical Investigation, *J. Am. Chem. Soc.*, 2002, **124**, 5197–5205; (c) M. Perić, M. Zlatar, S. Grubišić and M. Gruden-Pavlović, *Polyhedron*, 2012, **42**, 89–94; (d) J. Tercero, C. Diaz, J. Ribas, E. Ruiz, J. Mahía and M. Maestro, *Inorg. Chem.*, 2002, **41**, 6780–6789.
- 30 J. A. Bertrand and F. T. Helm, *J. Am. Chem. Soc.*, 1973, **95**, 8184–8185.
- 31 J. A. Bertrand, T. D. Black, P. Eller, P. Gary and F. T. Helm, *Inorg. Chem.*, 1976, **15**, 2965–2970.
- 32 S. Sarkar, A. Mondal, D. Chopra, J. Ribas and K. K. A. Rajak, *Eur. J. Inorg. Chem.*, 2006, 3510–3516.
- 33 P. Talukder, S. Sen, S. Mitra, L. Dahlenberg, C. Desplanches and J. P. Sutter, *Eur. J. Inorg. Chem.*, 2006, 329–333.
- 34 J. Tang, J. S. Costa, A. Golobic, B. Kozlevcar, A. Robertazzi, A. V. Vargiu, P. Gamez and J. Reedijk, *Inorg. Chem.*, 2009, **48**, 5473–5479.
- 35 C. L. Klein, R. J. Majeste, L. M. Trefonas and C. J. Oconnor, *Inorg. Chem.*, 1982, **21**, 1891–1897.
- 36 Y. Rodríguez-Martin, J. Sanchiz, C. Ruiz-Perez, F. Lloret and M. Julve, *CrystEngComm*, 2002, 631–637.
- 37 H. Núñez, L. Soto, J. J. Server-Carrió, J. García-Lozano, A. Sancho, R. Acerete and E. Escrivá, *Inorg. Chem.*, 2005, **44**, 4644–4655.
- 38 P. Baran, R. Boca, M. Breza, H. Elias, H. Fuess, V. Jorik, R. Klement and I. Svoboda, *Polyhedron*, 2002, **21**, 1561–1571.
- 39 S. Salunke-Gawali, S. Y. Rane, V. G. Puranik, C. Guyard-Duhayon and F. Varret, *Polyhedron*, 2004, **23**, 2541–2547.
- 40 W. Estes and W. E. Hatfield, *Inorg. Chem.*, 1978, **17**, 3226–3231.
- 41 R. D. Willett, C. J. Gómez-García, B. Twamley, S. Gomez-Coca and E. Ruiz, *Inorg. Chem.*, 2012, **51**, 5487–5493.
- 42 J. A. Bertrand, E. Fujita and D. G. VanDerveer, *Inorg. Chem.*, 1980, **19**, 2022–2028.
- 43 W. Plass, A. Pohlmann and J. Rautengarten, *Angew. Chem., Int. Ed.*, 2001, **40**, 4207–4210.
- 44 T. Hamaguchi, T. Nagata, S. Hayami, S. Kawata and I. Ando, *Dalton Trans.*, 2017, **46**, 6196–6201.
- 45 A. Okazawa and T. Ishida, *Chem. Phys. Lett.*, 2009, **480**, 198–202.
- 46 Z. Vasková, J. Moncol, M. Korabik, D. Valigura, J. Švorec, T. Lis, M. Valko and M. Melník, *Polyhedron*, 2010, **29**, 154–163.
- 47 B. Kozlevčar, N. Kitanovski, Z. Jagličić, N. A. G. Bandeira, V. Robert, B. Le Guennic and P. Gamez, *Inorg. Chem.*, 2012, **51**, 3094–3102.
- 48 G. A. Craig, O. Roubeau, J. Ribas-Ariño, S. J. Teat and G. Aromí, *Polyhedron*, 2013, **52**, 1369–1374.
- 49 J. M. Pages, L. Escriche-Tur, M. Font-Bardia, G. Aullon and M. Corbella, *CrystEngComm*, 2018, **20**, 6629–6639.
- 50 R. J. Staples, W. M. Reiff and E. V. Rybak-Akimova, *Inorg. Chem.*, 2004, **43**, 3930–3941.
- 51 K. Van Langenberg, S. R. Batten, K. J. Berry, D. C. R. Hockless, B. Moubaraki and K. S. Murray, *Inorg. Chem.*, 1997, **36**, 5006–5015.
- 52 D. J. Price, S. R. Batten, B. Moubaraki and K. S. Murray, *Polyhedron*, 2003, **22**, 2161–2167.
- 53 A. J. Tasiopoulos, N. C. Harden, K. A. Abboud and G. Christou, *Polyhedron*, 2003, **22**, 133–143.
- 54 E. R. Johnson, S. Keinan, P. Mori-Sanchez, J. Contreras-Garcia, A. J. Cohen and W. Yang, *J. Am. Chem. Soc.*, 2010, **132**, 6498–6506.
- 55 E. Ruiz, J. Cano, S. Alvarez and P. Alemany, *J. Comput. Chem.*, 1999, **20**, 1391–1400.

