



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

## Synthetic, structural, spectroscopic and theoretical study of a Mn(III)-Cu(II) dimer containing a Jahn-Teller compressed Mn ion

### Citation for published version:

Berg, N, Hooper, TN, Liu, J, Beedle, CC, Singh, SK, Rajaraman, G, Piligkos, S, Hill, S, Brechin, EK & Jones, LF 2013, 'Synthetic, structural, spectroscopic and theoretical study of a Mn(III)-Cu(II) dimer containing a Jahn-Teller compressed Mn ion', *Dalton Transactions*, vol. 42, no. 1, pp. 207-216.  
<https://doi.org/10.1039/c2dt31995k>

### Digital Object Identifier (DOI):

[10.1039/c2dt31995k](https://doi.org/10.1039/c2dt31995k)

### Link:

[Link to publication record in Edinburgh Research Explorer](#)

### Document Version:

Peer reviewed version

### Published In:

Dalton Transactions

### Publisher Rights Statement:

Copyright © 2012 by the Royal Society of Chemistry. All rights reserved.

### General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

### Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact [openaccess@ed.ac.uk](mailto:openaccess@ed.ac.uk) providing details, and we will remove access to the work immediately and investigate your claim.



Post-print of peer-reviewed article published by the Royal Society of Chemistry.

Published article available at: <http://dx.doi.org/10.1039/C2DT31995K>

Cite as:

Berg, N., Hooper, T. N., Liu, J., Beedle, C. C., Singh, S. K., Rajaraman, G., Piligkos, S., Hill, S., Brechin, E. K., & Jones, L. F. (2013). Synthetic, structural, spectroscopic and theoretical study of a Mn(III)-Cu(II) dimer containing a Jahn-Teller compressed Mn ion. *Dalton Transactions*, 42(1), 207-216.

Manuscript received: 30/08/2012; Accepted: 08/10/2012; Article published: 09/10/2012

## Synthetic, structural, spectroscopic and theoretical study of a Mn(III)–Cu(II) dimer containing a Jahn–Teller compressed Mn ion\*\*

Nelly Berg,<sup>1</sup> Thomas N. Hooper,<sup>2</sup> Junjie Liu,<sup>3</sup> Christopher C. Beedle,<sup>3</sup> Saurabh Kumar Singh,<sup>5</sup> Gopalan Rajaraman,<sup>5,\*</sup> Stergios Piligkos,<sup>6</sup> Stephen Hill,<sup>3,4,\*</sup> Euan K. Brechin,<sup>2,\*</sup> and Leigh F. Jones<sup>1,\*</sup>

<sup>[1]</sup>School of Chemistry, National University of Ireland, Galway, University Road, Galway, Ireland.

<sup>[2]</sup>EaStCHEM, School of Chemistry, Joseph Black Building, University of Edinburgh, West Mains Road, Edinburgh, EH9 3JJ, UK.

<sup>[3]</sup>National High Magnetic Field Laboratory, Florida State University, Tallahassee, USA.

<sup>[4]</sup>Department of Chemistry, Indian Institute of Technology, Powai, Mumbai, India.

<sup>[5]</sup>Department of Chemistry, University of Copenhagen, Universitetsparken 5, Denmark.

<sup>[6]</sup>Department of Physics, Florida State University, Tallahassee, USA.

[\*]Corresponding author; G.R. e-mail: [rajaraman@chem.iitb.ac.in](mailto:rajaraman@chem.iitb.ac.in); S.H. e-mail: [shill@magnet.fsu.edu](mailto:shill@magnet.fsu.edu); E.K.B. e-mail: [ebrechin@staffmail.ed.ac.uk](mailto:ebrechin@staffmail.ed.ac.uk); L.F.J. e-mail: [leigh.jones@nuigalway.ie](mailto:leigh.jones@nuigalway.ie)

[\*\*]We wish to thank the NUI Galway Millennium Fund (LFJ) and the Irish Research Council (IRCSET Embark Initiative (NB)). EKB wishes to thank the Leverhulme Trust and the EPSRC (ref. EP/H006737/1). Work at the National High Magnetic Field Laboratory is supported by NSF Cooperative Agreement No. DMR-0654118 and by the State of Florida. SH acknowledges the support of the NSF (CHE0924374) and SP the Danish Natural Science Research Council (FNU) for a Sapere Aude Fellowship (10-081659). GR would like to acknowledge financial support from the Government of India through Department of Science and Technology (SR/S1/IC-41/2010) and Indian Institute of Technology Bombay for computing time and a JRF position(SKS).

### Supporting information:

† Electronic supplementary information (ESI) available. CCDC 898371. For ESI and crystallographic data in CIF or other electronic format see <http://dx.doi.org/10.1039/C2DT31995K>

## Abstract

The heterobimetallic complex  $[\text{Cu(II)Mn(III)(L)}_2(\text{py})_4](\text{ClO}_4)\cdot\text{EtOH}$  (**1**) built using the pro-ligand 2,2'-biphenol ( $\text{LH}_2$ ), contains a rare example of a Jahn-Teller compressed Mn(III) centre. Dc magnetic susceptibility measurements on **1** reveal a strong antiferromagnetic exchange between the Cu(II) and Mn(III) ions mediated through the phenolate O-atoms ( $J = -33.4 \text{ cm}^{-1}$ ), with magnetisation measurements at low temperatures and high fields suggesting significant anisotropy. Simulations of high-field and high frequency powder EPR data suggest a single ion anisotropy  $D_{\text{Mn(III)}} = +4.45 \text{ cm}^{-1}$ . DFT calculations also yield an antiferromagnetic exchange for **1**, though the magnitude is overestimated ( $J_{\text{DFT}} = -71 \text{ cm}^{-1}$ ). Calculations reveal that the antiferromagnetic interaction essentially stems from the  $\text{Mn}(d_{x^2-y^2})\text{-Cu}(d_{x^2-y^2})$  interaction. The computed single-ion anisotropy and cluster anisotropy also correlates well with experiment. A larger cluster anisotropy for the  $S = 3/2$  state compared to the single ion anisotropy of Mn(III) is rationalised on the basis of orbital mixing and various contributions that arise due to the spin-orbit interaction.

## Introduction

Coordination chemistry regularly provides examples of fascinating homo- and heterometallic molecules with potential applications in disparate fields. In bioinorganic chemistry certain heterometallic manganese complexes have been proposed as biomimetic models for energy and electron transfer processes - one such stimulus being the heterobimetallic  $[\text{Mn}_4\text{CaO}_4]$  cubane-like catalytic unit within photosystem II (PSII).<sup>1</sup> Mn has also played a pivotal role in the field of molecular magnetism: the anisotropic nature of the Mn(III) ion means that it is regularly selected as the metal of choice in the synthesis of Single-Molecule Magnets (SMMs)<sup>2</sup> and Single-Chain Magnets (SCMs),<sup>3</sup> whilst the isotropic Mn(II) ion can be employed in the construction of molecular magnetic refrigerants.<sup>4</sup> Molecular Nanomagnetism has also seen a renaissance in the synthesis of heterometallic  $3d/4f$  cluster compounds, driven, in the main, by the tuneable anisotropy of the lanthanide ions,<sup>5,6</sup> offering the chemist the opportunity to vary the physical properties of a molecule without significantly altering structure. Similarly the systematic synthesis of the extensive family of heterometallic  $[\text{Cr}_7\text{M}]$  wheels and their many siblings has made elegant in-roads into the field of quantum information processing *via* molecular qubits.<sup>7</sup>

The dissemination of these wonderfully elaborate and complex polynuclear heterometallic architectures was of course preceded by fundamental studies of the magnetic exchange between metal ions in very simple complexes, such as homo- and heterometallic dimers. An early example of this was the heterobimetallic complex  $[\text{CuVO}(\text{fsa})_2\text{en}]$  (where  $(\text{fsa})_2\text{en}^{4-}$  is the ligand derived from N, N'-(2-hydroxy-3-carboxybenzylidene)-1,2-diaminoethane) in which the nature of the intramolecular

ferromagnetic exchange was explained by the orthogonality of the magnetic orbitals centred on the Cu(II) and V(IV) ions.<sup>8</sup> More recently the heterodinuclear complex [Cu(II)Mn(III)Cl(5-Br-sap)<sub>2</sub>(MeOH)] (where 5-Br-sap = 5-Bromo-2-salicylideneamino-1-propanol) was shown to exhibit ferromagnetic exchange, with the magnetic anisotropy stemming from the presence of the single Jahn-Teller elongated Mn(III) ion conferring Single-Molecule Magnet (SMM) behaviour.<sup>9</sup>

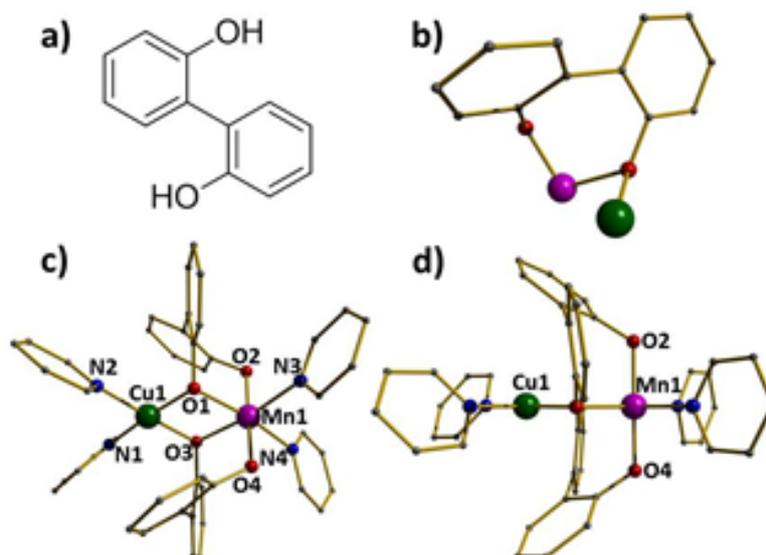
Herein we present the synthesis, structure, magnetism and theoretical study of the rather unusual heterobimetallic complex [Cu(II)Mn(III)(L)<sub>2</sub>(py)<sub>4</sub>](ClO<sub>4</sub>)·EtOH (**1**·EtOH) constructed using the pro-ligand 2,2'-biphenol (LH<sub>2</sub>) which contains a rare example of a Jahn-Teller compressed Mn(III) ion.<sup>10</sup>

## Results and Discussion

Complex **1** is readily prepared *via* the reaction of Mn(II)(ClO<sub>4</sub>)<sub>2</sub>·6H<sub>2</sub>O, anhydrous Cu(I)Cl, 2,2'-biphenol (LH<sub>2</sub>) and NEt<sub>4</sub>(OH) in a EtOH / pyridine solvent mixture. The resultant green / black solution, upon filtration and slow evaporation, gives rise to X-ray diffraction quality crystals of **1** in ~20% yield (see Table S1 for crystallographic data). Crystals were in a triclinic cell and structure solution was performed in the space group *P*-1.

The structure of the cation in **1** (Figure 1) comprises a single distorted square planar Cu(II) ion (Cu1) linked to a single six coordinate Jahn-Teller compressed Mn(III) ion (Mn1) by two bridging O<sub>phen</sub> donor atoms (Mn1-O1-Cu1, 99.72°; Mn1-O3-Cu1, 101.28°) belonging to two distinct doubly deprotonated 2,2'-biphenolate ligands. Their remaining O<sub>phen</sub> atoms (O2 and O4) are terminally bonded, providing the short axial bonds to the sole Mn(III) ion that define the Jahn-Teller compressed axis (Mn1-O2, 1.880 Å; Mn1-O4, 1.872 Å). The longer equatorial bonds range between 2.050 Å (Mn1-O3) and 2.203 Å (Mn1-N4). The “free” axial sites of the square planar Cu(II) ion are blocked by the presence of the phenyl rings of the rather twisted L<sup>2-</sup> ligands (Cu···Ph(centroid) = 3.560-3.579 Å); indeed the dihedral angles between the Ph rings on these same ligands are 49.86° [C5-C8] and 51.60° [C17-C20] respectively. Four terminal pyridine ligands complete the coordination spheres at the six coordinate Mn(III) centre (N3 and N4) and at the four coordinate Cu(II) site (N1 and N2) affording {MnO<sub>4</sub>N<sub>2</sub>} and {CuO<sub>2</sub>N<sub>2</sub>} coordination spheres, respectively.

**(turn to next page →)**



**Figure 1.** (a) The pro-ligand 2,2'-biphenol ( $LH_2$ ) and (b) its coordination mode upon double deprotonation. Molecular structure of the heterobimetallic cation in **1** as viewed off-set (c) and perpendicular (d) to the Jahn-Teller compressed axis of the Mn(III) ion. Colour code: green (Cu), purple (Mn), red (O), blue (N) and grey (C). Hydrogen atoms and counter anions have been omitted for clarity.

The resultant  $\{Cu(II)Mn(III)(L)_2(py)_4\}^+$  cation is charge balanced by the presence of a single  $ClO_4^-$  ion. The latter H-bonds to a EtOH molecule of crystallisation ( $O8 \cdots O9$ , 2.827 Å), with the closest contacts to the cluster complex being  $O5 \cdots H28 = 2.569$  Å and  $O5 \cdots H29 = 2.599$  Å. The ethanol molecule of crystallisation also hydrogen bonds to an aromatic proton (H26) of a nearby pyridine ligand ( $O9 \cdots H26$ , 2.759 Å). In the crystal the individual  $\{Cu(II)Mn(III)(L)_2(py)_4\}^+$  cluster units pack in superimposable rows along the  $a$  unit cell direction and are separated at a  $M \cdots M$  distance of  $\sim 12$  Å. These 1D rows pack in the common brickwork formation in the  $bc$  plane (Fig. S1).

Although a number of Cu(II)-Mn(II) dinuclear complexes are known in the literature,<sup>11</sup> a CSD search reveals that **1** is one of very few examples to show the Cu(II)-Mn(III) oxidation state distribution,<sup>9,12</sup> and is indeed the first to possess a Mn(III) ion exhibiting a Jahn-Teller compression. Complex **1** is structurally very similar to the aforementioned  $[Cu(II)Mn(III)Cl(5-Br-sap)_2(MeOH)]$  (**2**). Both contain  $[Cu(II)(OR)_2Mn(III)]$  magnetic cores (albeit with different R-groups), resulting in very similar Cu(II)-O(R)-Mn(III) bond angles ( $99.72$  and  $101.28^\circ$  in **1**) and ( $102.1^\circ$  in **2**). Indeed the only difference in terms of magnetic structure is the presence of a Jahn-Teller compressed Mn(III) ion in **1** and a Jahn-Teller elongated Mn(III) ion in **2**. The synthesis of **1** therefore represents an ideal opportunity to investigate the magneto-structural relationship in this family of complexes (*vide infra*).

## ***SQUID magnetometry***

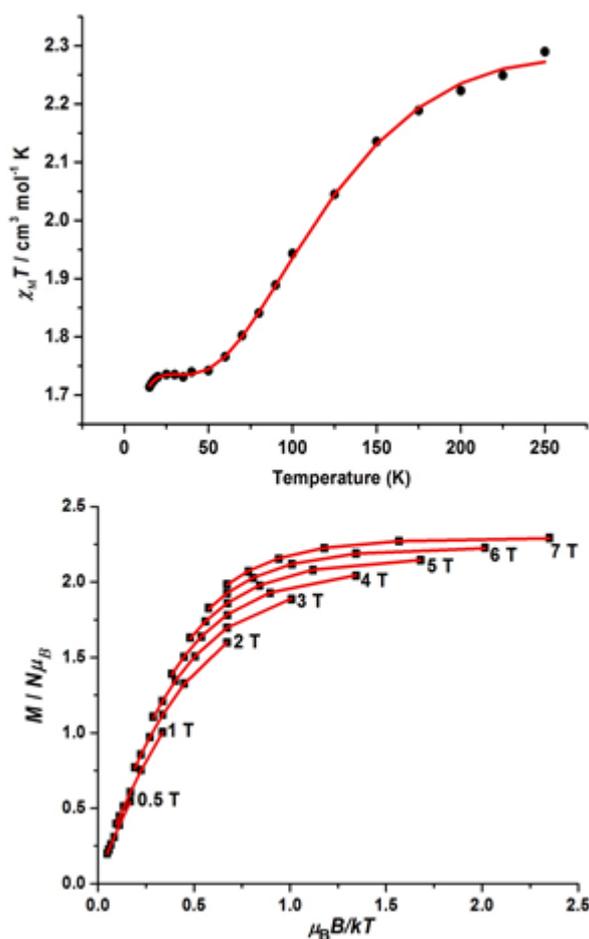
Dc magnetic susceptibility measurements were performed on a powdered microcrystalline sample of **1** in an applied magnetic field of 0.1 T and in the temperature range 250 to 5 K. The results are plotted as the  $\chi_M T$  product vs.  $T$  in the top panel of Figure 2. The high temperature  $\chi_M T$  value of 2.270 cm<sup>3</sup> K mol<sup>-1</sup> is lower than that expected from the spin-only value (3.375 cm<sup>3</sup> K mol<sup>-1</sup>) for non-interacting high-spin Mn(III) ( $3d^4$ ) and Cu(II) ( $3d^9$ ) ions, assuming  $g = 2$ . This reflects the existence of a strong intramolecular antiferromagnetic interaction in **1**. The  $\chi_M T$  product decreases in a sigmoidal-like fashion with decreasing temperature until it reaches a plateau of around 1.75 cm<sup>3</sup> K mol<sup>-1</sup> in the 45 to 20 K temperature interval. The  $\chi_M T$  product at this low temperature plateau is consistent with the existence of a thermally isolated  $S = 3/2$  spin-state. For the modelling of the  $\chi_M T$  product we used spin-Hamiltonian (1):

$$\hat{H} = -2J\hat{S}_{Cu} \cdot \hat{S}_{Mn} + D_{Mn}[\hat{S}_{z,Mn}^2 - S_{Mn}(S_{Mn} + 1)/3] + \mu_B \sum_{i=Cu,Mn} \{g_i \bar{B}_i \cdot \hat{S}_i\} \quad (1)$$

where  $J$  is the Cu(II)-Mn(III) isotropic exchange interaction parameter,  $\hat{S}$  is a spin-operator,  $D_{Mn}$  is the single-ion uniaxial anisotropy of Mn(III),  $S = 2$  is the single-ion spin of Mn(III),  $\mu_B$  is the Bohr magneton,  $g_{Mn} = 2.0$  and  $g_{Cu} = 2.1$  are the employed isotropic  $g$ -factors of the Mn(III) and Cu(II) ions, respectively, and  $\bar{B}$  is the applied magnetic field vector. The  $\chi_M T$  product of **1** was numerically fitted, by use of the simplex algorithm,<sup>13</sup> to spin-Hamiltonian (1) by numerical diagonalisation of the spin-Hamiltonian matrix of dimension 10 by 10. While fitting the  $\chi_M T$  product, it became apparent that inclusion of the Mn(III) single-ion anisotropy term does not significantly improve the quality of fit, as measured by the  $\chi^2$  statistics. This is as expected since anisotropy effects are predominant at very low temperatures and thus, do not weight significantly in the fitting of the  $\chi_M T$  product. The best-fit parameter of (1) to the  $\chi_M T$  product of **1** was  $J = -33.4$  cm<sup>-1</sup>, affording an  $S = 3/2$  spin ground state with the  $S = 5/2$  excited state some 167 cm<sup>-1</sup> higher in energy. In addition, inter-molecular interactions were taken into account in the frame of mean-field theory, by use of the Curie–Weiss temperature,  $\theta$ . A Curie-Weiss constant of  $\theta = -0.9$  K, was necessary to reproduce the small drop of the  $\chi_M T$  product below  $\sim 20$  K. The determined strong antiferromagnetic interaction is consistent with the above qualitative description of the thermal dependence of the  $\chi_M T$  product of **1**.

In order to determine, by magnetisation measurements, the single-ion axial anisotropy parameter for the Mn(III) centre in **1**, variable-temperature-variable-field magnetisation measurements were performed in the  $T = 2.0$ – $7.0$  K and  $B = 0.5$ – $7.0$  T temperature and dc magnetic field ranges, respectively. These experimental data are presented as reduced magnetisation ( $M/N\mu_B$  vs.  $\mu_B B/kT$ , with  $N$  being Avogadro's number and  $k$  the Boltzmann constant) in the lower panel of Figure 2. They were again numerically fitted, by use of the simplex algorithm,<sup>13</sup> to spin-Hamiltonian (1), by numerical diagonalisation of the 10 by 10 spin-Hamiltonian matrix. The best fit parameter, keeping  $J$

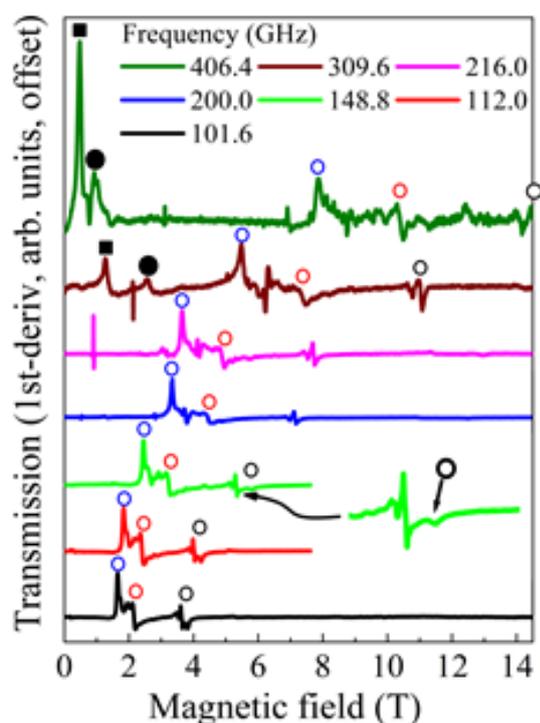
fixed to  $-33.4 \text{ cm}^{-1}$  (as determined by fitting the  $\chi_M T$  product), was  $D_{\text{Mn}} = +4.95 \text{ cm}^{-1}$ . Forcing the  $D_{\text{Mn}}$  parameter to vary only in the negative-values semi-axis, leads to  $\chi^2$  values that are two orders of magnitude higher than the ones obtained by letting  $D_{\text{Mn}}$  vary freely to give  $D_{\text{Mn}} = +4.95 \text{ cm}^{-1}$ . The positive value of  $D_{\text{Mn}}$  is in good agreement with the axially compressed nature of the coordination sphere of the Mn(III) ion.



**Figure 2.** Upper panel: Plot of the  $\chi_M T$  product of **1** vs.  $T$  in the 250 - 5 K temperature range in an applied field of 0.1 T. The solid red line is the best-fit of the data with  $J = -33.4 \text{ cm}^{-1}$  as described in the text. Lower panel: Plot of reduced magnetisation of **1** in the 2.0 to 7.0 K temperature range and at the indicated field strengths. The solid red lines are the best-fit of the data with  $J$  fixed to  $-33.4 \text{ cm}^{-1}$  and  $D_{\text{Mn}} = +4.95 \text{ cm}^{-1}$ , as described in the text.

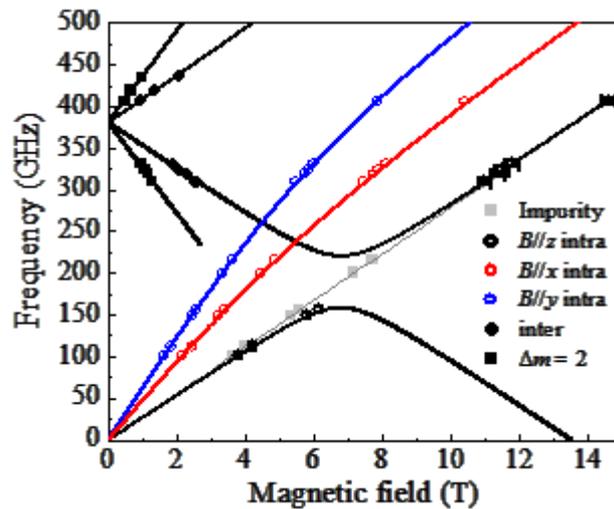
Powder EPR data were collected for complex **1** in order to determine the magnetic anisotropy of the dimer. Given the spin states of the constituent ions ( $S = 2$  and  $1/2$  for Mn(III) and Cu(II) respectively), one expects a half integer ground state spin of either  $S = 3/2$  or  $5/2$  for the dimer. The low-energy

spectrum should thus consist of either two or three [*i.e.*  $(2S + 1)/2$ ] Kramers doublets, split by any associated magnetic anisotropy. Figure 3 displays representative frequency-dependent powder EPR spectra for complex **1** collected between 100.6 and 406 GHz at a relatively low temperature of 5 K. All of the labelled resonances persist to the lowest temperatures investigated (not shown), indicating that they involve transitions either within or from the lowest Kramers doublet. At frequencies below 200 GHz, the spectra are dominated by three strong resonance branches that we assign to the x, y and z-transitions within the ground-state Kramers doublet; these are marked by open red, blue and black circles, respectively. As can be seen in the frequency versus field map in Figure 4, the peak positions corresponding to these transitions each lie on simulated curves (*vide infra*) that intersect the origin, as expected for intra-Kramers transitions. At frequencies above 200 GHz, additional resonance branches appear at low fields (solid black squares and circles). As can be seen in Figure 4, these resonances all lie on lines/curves that meet at a unique finite-frequency offset on the zero-field axis. We thus assign them as inter-Kramers transitions, and the intercept on the ordinate corresponds exactly to the zero-field energy splitting between the two lowest Kramers doublets. We note that this splitting of  $381 \pm 5$  GHz is rather substantial for a complex comprised of just Mn(III) and Cu(II) ions. The solid circles in Figure 4 correspond to the usual  $\Delta m_s = \pm 1$  perpendicular-mode EPR transitions, while the solid squares correspond to  $\Delta m_s = \pm 2$  resonances (see below). The simulations assume that the applied field,  $B$ , is parallel to z. However, all three inter-Kramers components (*i.e.*  $B//x$ , y and z) generate  $\Delta m_s = \pm 1$  and  $\pm 2$  resonances at similar locations, provided the applied field is relatively weak ( $< 3$  T), which may account for the appreciable intensity of the lowest field 406.4 GHz peak in Figure 3.



← **Figure 3.** Representative frequency dependent powder EPR spectra collected at 5 K: the red, blue and black open circles denote the intra-Kramers x, y and z-components of the spectra, while the solid circles ( $\Delta m_s = \pm 1$ ) and squares ( $\Delta m_s = \pm 2$ ) denote transitions between Kramers doublets; the inset displays an expanded portion of the 200 GHz spectrum in the vicinity of the z-component (see main text for explanation). The extremely sharp features seen in some of the spectra (*e.g.* at  $\sim 1$  T in the 216 GHz spectrum) correspond to known signals from oxygen adsorbed in the sample holder.

While analysis of susceptibility data enables reliable estimates of the exchange coupling within a simple spin system, especially a dimer, EPR measurements provide much more robust constraints on both the sign and magnitude of the magneto-anisotropy. The EPR data presented in Figure 4 clearly reveal the presence of a pair of low-energy Kramers doublets. However, they do not rule out the possibility of a third higher-lying doublet. Thus, a determination of the ground state spin from the EPR data is not entirely straightforward. This would require careful analysis of much weaker transitions observed at elevated temperatures, and we do not pursue this here because the susceptibility measurements clearly indicate that the coupling within the dimer is antiferromagnetic. Therefore, in the following analysis, we assume that the ground state spin value is  $S = 3/2$ . In doing so, we will see that the obtained anisotropy is quite consistent with expectations, thus providing indirect confirmation of the  $S = 3/2$  ground state spin value.



**Figure 4.** Plot of the frequency dependent EPR peak positions as a function of the applied magnetic field strength. The solid curves (same colour coding as the data) correspond to the simulations described in the main text. Note that the grey data points and the grey line correspond to an impurity signal with  $g = 2.003(2)$ .

We first turn our attention to the sign of the second-order axial zero-field splitting parameter  $D_S$  associated with the  $S = 3/2$  ground state. In the presence of easy-plane type anisotropy ( $D_S > 0$ ), the  $B//x$ - and  $y$ -components of the spectrum associated with the lowest Kramers doublet (red and blue circles, respectively, Figure 4) possess effective  $g$ -values that are substantially larger than 2.00, *i.e.* slopes that substantially exceed 28 GHz/T as  $B \rightarrow 0$ ; meanwhile, the  $B//z$ -component has an effective  $g$ -value (slope) very close to 2.00 (or 28 GHz/T, denoted by the grey line in Figure 4). The opposite

holds for the easy-axis case ( $D < 0$ ).<sup>14,15</sup> Thus, the powder EPR data clearly support the presence of an easy-plane type anisotropy ( $D > 0$ ), which one would expect in a JT compressed Mn(III) system, since this represents the main source of anisotropy within the dimer. The magnitude of  $D_S$  is approximately constrained by the intercept of the inter-Kramers resonance branches. For a purely axial system, this intercept corresponds exactly to  $2D_S$ , hence  $D_S \approx 381/2 \text{ GHz} = 6.35 \text{ cm}^{-1}$ . However, as can clearly be seen from Figures 3 and 4, there is a considerable splitting between the x and y components of the spectrum, signifying a rhombic distortion and, hence, a rhombic  $E_S$  parameter. In such situations,  $D_S$  can only be determined precisely *via* diagonalisation of the following effective spin Hamiltonian:

$$\hat{H}_S = D_S \hat{S}_z^2 + E_S (\hat{S}_x^2 - \hat{S}_y^2) + \mu_B \vec{B} \cdot \mathbf{g}_S \cdot \hat{S} \quad (2)$$

where  $\hat{S}$  represents the total spin operator and  $\hat{S}_i$  ( $i = x, y, z$ ) its components, the subscript  $S = 3/2$  defines the total spin-state on which the effective spin-Hamiltonian (2) is applied and  $\mathbf{g}_S$  is the Landé tensor of the total spin-state  $S$ . The best simulation of the data in Figure 4 was obtained with the following parameters:  $D_S = +6.27 \text{ cm}^{-1}$ ,  $E_S = \pm 0.57 \text{ cm}^{-1}$ ,  $g_{S,x} = 1.97$ ,  $g_{S,y} = 1.97$  and  $g_{S,z} = 1.98$ . The main constraint on the  $E_S$  parameter is the splitting between the x- and y-components of the intra-Kramers transitions (red and blue curves in Figure 4), *i.e.* its value is well constrained. While one might expect a corresponding difference between  $g_{S,x}$  and  $g_{S,y}$ , this leads to an over parameterization of the model. Therefore, we have assumed these parameters to have the same value.

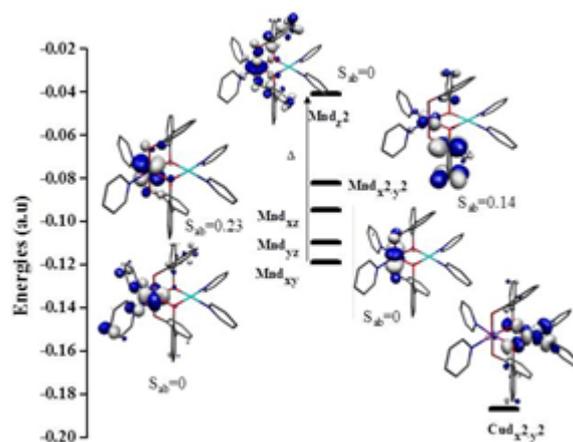
The transverse anisotropy ( $E$ ) leads naturally to an avoided-crossing between the  $m_S = -3/2$  and  $m_S = -1/2$  components of the two Kramers doublets, which meet at  $\sim 7 \text{ T}$  when  $B//z$ . It is this interaction that is the reason for the  $\sim 60 \text{ GHz}$  ( $2 \text{ cm}^{-1}$ ) gap between the inter- and intra-Kramers z-component resonance branches at  $\sim 7 \text{ T}$  in Figure 4 (solid black lines). Surprisingly, spectra recorded at frequencies close to 200 GHz (*e.g.* the blue curve in Fig. 3) still show weak resonances in this gapped region of the  $B//z$  simulations. Closer inspection of the spectra reveals that the z component appears to consist of a narrow derivative (*i.e.* a sharp peak followed by a sharp minimum) superimposed on a broad dip; this contrasts the x and y components that each consist of a single feature, *i.e.* a peak for y and a shoulder for x, as expected for such a powder spectrum. The narrow and broad z-components are separately plotted in Figure 4 as grey squares and open black circles. The sharp components lie on a straight (grey) line passing through the origin, having a slope corresponding to  $g = 2.003(2)$ . At most frequencies, the broad dip lies very close to this line. However, one clearly sees that the broad component moves appreciably to the high-field side of the  $g = 2.003$  position (the sharp peak) as the frequency approaches the gapped region; the relevant portion of the 148.8 GHz spectrum has been enlarged in Figure 3 (green curve) to emphasize this point. Indeed, if one assumes that it is this broad dip that corresponds to the z-component of the dimer spectrum, one finds excellent agreement with the simulations in Figure 4. Meanwhile, we believe that the sharp component corresponds to an impurity phase within the powder, possibly containing isotropic Mn(II); this is often found to be the

case in Mn-containing polynuclear clusters, and explains why a resonance is still observed in the gapped region of the  $B//z$  spectrum. In fact, it is notable that the 200 GHz spectrum exhibits only a sharp component, the broad dip appears to be absent, which is why an open circle is not included in Figure 4 for this frequency.

Due to the strong Cu(II)-Mn(III) exchange interaction ( $J \sim -33 \text{ cm}^{-1}$ ), we can make a reliable estimation of the anisotropy parameters associated with the individual ions using the projection method.<sup>16</sup> According to Kramers' theorem, any zero-field splitting interaction associated with the Cu(II) ion ( $S_{\text{Cu}} = 1/2$ ) is strictly forbidden; hence, the zero-field splitting parameters of the molecule are determined only by the anisotropy parameters of the Mn(III) ion. The projection method gives  $\ddot{D}_{S=3/2} = 1.4\ddot{D}_{\text{Mn}}$ , where  $\ddot{D}_{S=3/2}$  and  $\ddot{D}_{\text{Mn}}$  are the anisotropy tensors of the ground spin-state and the Mn(III) ion, respectively. Therefore, we estimate that  $D_{\text{Mn}} = +4.45 \text{ cm}^{-1}$  and  $E_{\text{Mn}} = \pm 0.41 \text{ cm}^{-1}$ . We note that these zero-field interaction parameters obtained *via* EPR data are in excellent agreement with the parameters obtained *via* the reduced magnetisation measurements, and are consistent with values found for related single-ion Mn(III) compounds (see Table S3 and references therein). Finally, we comment on the  $g$ -values obtained from the EPR measurements. Again, using projection methods, one can estimate that  $\vec{g}_{S=3/2} = 1.4\vec{g}_{\text{Mn}} - 0.4\vec{g}_{\text{Cu}}$ . Consequently, the obtained  $g$ -values for the  $S = 3/2$  ground spin-state should be dominated by the Mn(III) ion, which is consistent with the obtained values that are very close to 2.00.

### ***DFT Analysis***

Density functional studies have been performed on complex **1** to compute the  $J$  value and to explore the origin of the nature of the interaction observed experimentally. The B3LYP/TZV combination (see computational details below) yields a  $J$  value of  $-71 \text{ cm}^{-1}$ . Although the sign of  $J$  has been correctly reproduced, the magnitude is overestimated compared to experiment ( $J_{\text{exp}} = -33.4 \text{ cm}^{-1}$ ). Although the employed methodology generally offers good numerical estimates of  $J$  values, there are instances where overestimation of the  $J$  has been noted.<sup>17,18</sup> Calculations performed incorporating the counter-anion did not lead to any significant improvement ( $-71 \text{ cm}^{-1}$  vs.  $-83 \text{ cm}^{-1}$ ; Table S5). Since our aim is to probe the origin of antiferromagnetic coupling in the {MnCu} pair, we have analysed the wave function and the magnetic orbitals of the Mn(III) and Cu(II) ions (Figure 5).



**Figure 5.** DFT calculated magnetic orbitals of Mn(III) and Cu(II) with computed  $S_{ab}$  (overlap integral) values. The isodensity surface represented corresponds to a value of  $0.05 \text{ e}^-/\text{bohr}^3$ .

The Cu(II) ion is in a square planar environment, with the unpaired electron residing in the  $d_{x^2-y^2}$  orbital ( $b_{1g}$ ). For Mn(III) the following configuration has been detected from the computed wave function:  $b_{2g}(d_{xy})^1 e_g(d_{xz})^1 e_g(d_{yz})^1 b_{1g}(d_{x^2-y^2})^1$  where the  $a_{1g}(d_z^2)$  orbital is found to be empty and lies 1.12 eV higher in energy. The spin density values on the Mn(III) (1.663) and Cu(II) (0.59) ions, clearly indicates strong spin delocalisation to other atoms. The spin densities on the bridging O-atoms (O1 and O2) are found to be positive (0.11 and 0.12) while the N-atoms coordinated to Cu(II) and Mn(III) are also found to have positive spin densities. On the other hand, the spin densities on the O-atoms lying in the axial direction are found to be negative (Figure 6). This is in line with our expectation where the empty  $d_z^2$  orbital propagates spin polarization rather than spin delocalization. Such a mixture of delocalization and polarization has been reported previously.<sup>13c,21</sup>

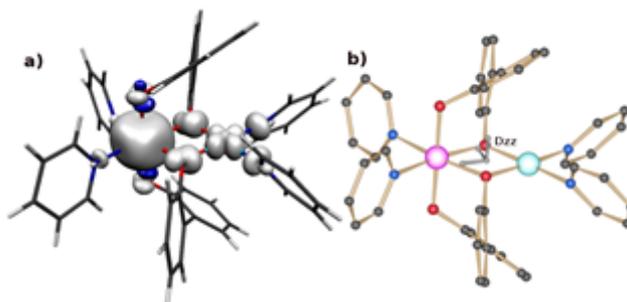
energy than the  $b_{1g}$  orbital. This splitting is related to the strength of the distortion and significant splitting demonstrates that the Mn(III) ion is undergoing relatively strong JT compression.<sup>19,20</sup>

Qualitatively  $\Delta(E_{d_{xy}} - E_{d_z^2})$  – calculated to be  $\sim 2.12 \text{ eV}$  for **1** (Figure 5) - is related to the anisotropy by the following equation for a Jahn-Teller compressed Mn(III) ion:

$$D = D_{zz} - \frac{(D_{xx} + D_{yy})}{2} \approx + \frac{3\zeta_{\text{eff}}^2}{16\Delta} \quad (3)$$

where  $\zeta_{\text{eff}}$  is the effective covalently reduced one-electron spin-orbit coupling (SOC) constant of the metal ion under investigation, and  $\Delta$  is as defined in Figure 5. Since in both the metal ions the  $b_{1g}(d_{x^2-y^2})$  orbital is singly occupied, a significant interaction between these two magnetic orbitals is expected, leading to antiferromagnetic coupling. To prove this hypothesis, overlap integrals between pairs of magnetic orbitals have been computed. Our overlap integral analysis reveals a significant overlap

between  $\text{Mn}(d_{x^2-y^2})$  and  $\text{Cu}(d_{x^2-y^2})$  (Table S6). Additionally  $\text{Mn}(d_{xz})$  has non-negligible overlap with the  $\text{Cu}(d_{x^2-y^2})$  orbital. The  $\text{Mn}(d_{x^2-y^2})$ - $\text{Cu}(d_{x^2-y^2})$  interaction is very significant (Figure 5) and this leads to a moderate antiferromagnetic interaction in complex **1**. The other dinuclear  $\{\text{MnCu}\}$  (**2**) complex reported in the literature has ferromagnetic coupling.<sup>9a</sup> In that complex the Mn(III) is Jahn-Teller elongated, the  $d_{x^2-y^2}$  orbital is empty and the significant  $\text{Mn}(d_{x^2-y^2})$ - $\text{Cu}(d_{x^2-y^2})$  interaction present in **1**, is absent, leading to moderate ferromagnetic coupling. This pair of complexes illustrates rather nicely how the magnetic coupling can be tuned from ferromagnetic to antiferromagnetic by altering the nature of the distortion (elongation vs. compression) on the Mn(III) site.



**Figure 6.** a) DFT computed spin density plot of **1** for the  $S = 2$  state. The isodensity surface represented corresponds to a value of  $0.005 e^-/\text{bohr}^3$ . The white and blue regions indicate positive and negative spin populations, respectively. b) DFT calculated orientation of  $D$ -tensor in **1**. H-atoms have been omitted for clarity.

We have also computed the ZFS parameter of complex **1** using the methodology established by Neese *et al.*<sup>20</sup> We have performed calculations on complex **1** and also on a fictitious  $\{\text{MnZn}\}$  model complex of **1** to estimate the  $S = 3/2$  ZFS of **1** and the  $S = 2$  single-ion anisotropy of the Mn(III) ion, in order to compare to the experimentally extracted parameters. The calculated single ion anisotropy of the Mn(III) ion is  $+1.94 \text{ cm}^{-1}$ , while the cluster anisotropy is computed to be  $+2.35 \text{ cm}^{-1}$ . The HF-EPR estimates of the ZFS are  $+4.45$  and  $+6.27 \text{ cm}^{-1}$ , respectively. Our DFT calculations reproduce the sign and the trend correctly, but clearly underestimate the magnitude quite considerably. Such underestimation by DFT methods has been well documented: accurate reproduction demands inclusion of excited state contributions and this is possible only by means of *ab initio* calculations.<sup>20</sup>

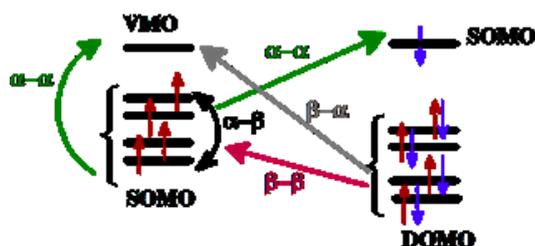
Analysis of the contributions to the net  $D$ -tensor suggest the main source arises from spin-orbit coupling ( $D_{\text{SOC}}$ ), while the spin-spin contribution ( $D_{\text{SS}}$ ) is found to be small, and significantly less than that reported for complexes in which the Mn(III) ion is Jahn-Teller elongated.<sup>19,20</sup> The significant contributions to the ZFS are listed in Table 1. For  $D_{\text{SOC}}$  the spin-flip excitations account for nearly

65% of the contribution. Multiple contributions to the  $D$ -tensor listed in Table 1 reveal some distinct differences between the single-ion and the cluster anisotropy. The  $\alpha \rightarrow \beta$  (spin-up to spin-down) excitations are pronounced for the Mn(III) single-ion ( $S = 2$  state) anisotropy, while other contributions are very small. However for the dinuclear  $\{\text{MnCu}\}$  compound ( $S = 3/2$  ground state), very large  $\beta \rightarrow \alpha$  and  $\alpha \rightarrow \alpha$  contributions were detected, in addition to the expected  $\alpha \rightarrow \beta$  spin flip.

spin flip excitations	single ion anisotropy ( $\text{cm}^{-1}$ )	cluster anisotropy ( $\text{cm}^{-1}$ )
<b>SOMO-SOMO</b> $\alpha \rightarrow \beta$	1.215	1.484
<b>DOMO-SOMO</b> $\beta \rightarrow \beta$	0.028	0.059
<b>SOMO -VMO</b> $\alpha \rightarrow \alpha$	0.339	-2.729
<b>DOMO-VMO</b> $\beta \rightarrow \alpha$	0.002	3.554

**Table 1.** DFT computed spin flip/spin-conserved excitations for the  $\{\text{MnCu}\}$  and  $\{\text{MnZn}\}$  complexes.

Analysing the wave function (see Figure 5 and related discussion on  $\Delta$ ), reveals that the main reason for the difference is that the Mn(II) ion has one empty  $\alpha$  type orbital (for the  $S = 3/2$  ground state) and this leads to large singly occupied molecular orbital (SOMO) to virtual molecular orbitals (VMO) contribution. There is also a significant contribution due to doubly occupied orbitals (DOMO) to VMO due to the presence of Cu(II). These two contributions are minimal for the mononuclear Mn(III) ions. This is illustrated in Figure 7. It is apparent from the break-up contributions given for the  $\{\text{MnCu}\}$  complex that the DOMO  $\rightarrow$  VMO excitation which has the largest positive value determines the net sign of the anisotropy. Since this parameter is related to the mixing of Mn and Cu  $d$ -orbitals, efforts to vary the ligand field are expected to significantly affect the magnitude of the anisotropy. The computed  $D$ -tensor orientation for the  $\{\text{MnCu}\}$  complex is shown in Figure 6. As expected the  $D_{zz}$  axis is aligned along JT axis with a deviation of approximately  $5^\circ$ .



**Figure 7.** Schematic illustrating the different contribution to anisotropy in complex 1.

Complex	$D$ total ( $\text{cm}^{-1}$ )	$D_{\text{soc}}$ ( $\text{cm}^{-1}$ )	$D_{\text{ss}}$ ( $\text{cm}^{-1}$ )	$E/D$	$g$ -tensor
DFT on <b>1</b> ( $S=3/2$ )	2.35	2.369	-0.009	0.06	1.98
DFT on {MnZn}	1.94	1.58	0.35	0.06	1.99
HF-EPR on <b>1</b>	6.27			0.09	----
HF-EPR $\text{Mn}^{\text{III}}$ ion	4.78			0.08	1.98

**Table 2.** Experimental versus DFT computed single-ion and cluster anisotropies with SOC and SS contributions.

## Conclusions

Reaction of anhydrous  $\text{CuCl}$ ,  $\text{Mn}(\text{ClO}_4)_2 \cdot 6\text{H}_2\text{O}$  and 2,2-biphenol ( $\text{LH}_2$ ) in an EtOH / pyridine solvent mixture produces the heterobimetallic complex  $[\text{Cu}(\text{II})\text{Mn}(\text{III})(\text{L})_2(\text{py})_4](\text{ClO}_4) \cdot \text{EtOH}$  (**1**·EtOH). Complex **1** is a rare example of a dimeric  $[\text{Cu}(\text{II})\text{Mn}(\text{III})]$  cluster compound and the first example to exhibit a Jahn-Teller compressed Mn(III) centre. Magnetic susceptibility and magnetisation studies conclude that the exchange in **1** is antiferromagnetic resulting in an isolated  $S = 3/2$  ground spin state, with  $J = -33.4 \text{ cm}^{-1}$ ,  $g = 2.0$  and  $D_{\text{Mn}(\text{III})} = +4.95 \text{ cm}^{-1}$ . High-field, high-frequency EPR studies on powdered samples of **1** corroborate these findings, giving rise to the spin Hamiltonian parameters  $D_{\text{Mn}(\text{III})} = +4.45 \text{ cm}^{-1}$  and  $E_{\text{Mn}(\text{III})} = +0.41 \text{ cm}^{-1}$ . DFT calculations offer insights into the mechanism of magnetic coupling where the origin of the antiferromagnetic interaction is related to the nature of the Jahn-Teller distortion. Calculations also yield a reasonable estimate of the anisotropy for complex **1** and explain how the mixing of the Mn(III) and Cu(II)  $d$ -based orbitals leads to dramatic changes in the sign as well as the magnitude of the anisotropy for **1**.

## Experimental Section

### *Physical measurements*

Infra-red spectra were recorded on a Perkin Elmer FT-IR *Spectrum One* spectrometer equipped with a Universal ATR Sampling accessory (NUI Galway). Elemental analysis was carried out by the School of Chemistry microanalysis service at NUI Galway. Variable-temperature, solid-state direct current (dc) magnetic susceptibility data down to 5 K were collected on powdered samples using a Quantum Design MPMS-XL SQUID magnetometer equipped with a 7 T dc magnet (University of Edinburgh). Diamagnetic corrections were applied to the observed paramagnetic susceptibilities using Pascals' constants. Complex **1** was set in eicosane to avoid torquing of the crystallites. Powder EPR data were collected in a transmission-type spectrometer employing a 17 T superconducting magnet. A phase-locked oscillator, in conjunction with a series of multipliers and amplifiers, was employed as a

microwave source capable of providing quasi-continuous frequency coverage up to 600+ GHz; a cold bolometer was used for detection.<sup>22</sup>

### ***Computational Details***

DFT calculations combined with the Broken Symmetry (BS) approach<sup>23</sup> were employed to compute the *J* values. The BS method has a proven record of yielding good numerical estimates of *J* constants for a variety of complexes.<sup>24,25</sup> A detailed technical discussion of the computational details can be found elsewhere.<sup>26,27</sup> Here, we have performed most of our calculations using the Gaussian 09 suite of programs with the fragment approach.<sup>28</sup> We have employed a hybrid B3LYP functional<sup>29</sup> with TZV basis sets on all atoms.<sup>30</sup> A very tight SCF convergence was employed throughout. DFT calculations for the estimation of the *D*-tensor used the ORCA suite of programmes,<sup>31</sup> employing the non-hybrid BP86 functional<sup>32</sup> using quasi-degenerate theory<sup>33</sup> with CP approach.<sup>34,35</sup> The Alhrichs TZVPPP basis set was used for the Mn(III) and Cu(II) ions, while for the remaining atoms we have used the TZVP basis set. The RI approximations were considered during calculation with auxiliary TZV/J cumbic fitting basis set.<sup>36</sup> Increased integration grids (Grid 5 in ORCA convention) along with tight SCF convergence were used.

### ***Materials and syntheses***

All reactions were performed under aerobic conditions and all reagents and solvents were used as purchased. **Caution:** Although we encountered no problems care should be taken when using the potentially explosive perchlorate salts. The NEt<sub>4</sub>(OH) solution employed was a 40% by weight aqueous solution which was used as purchased.

**[Mn(III)Cu(II)(L)<sub>2</sub>(py)<sub>4</sub>](ClO<sub>4</sub>).MeOH (1):** Mn(ClO<sub>4</sub>)<sub>2</sub>.6H<sub>2</sub>O (0.25 g, 0.98 mmol) and anhydrous Cu(I)Cl (0.39 g, 3.94 mmol) were dissolved in 35 cm<sup>3</sup> EtOH along with 1 cm<sup>3</sup> of pyridine. 2,2'-biphenol (LH<sub>2</sub>) (0.73 g, 3.94 mmol) was then added along with 2 drops of NEt<sub>4</sub>(OH). The resultant solution was stirred for 5-7 minutes before being filtered to give a green / black solution. Brown block-like crystals of **1** were obtained in ~20% yield upon slow evaporation of the mother liquor over a period of 24 hours. *Note: Attempts at repeating the reaction employing Cu(II) salts have all been unsuccessful.* Elemental analysis calculated (found) (%) for C<sub>45</sub>H<sub>40</sub>N<sub>4</sub>O<sub>9</sub>ClMnCu (**1**.EtOH): C: 57.82 (57.98), H: 4.31 (3.81), N: 5.99 (6.05). FT-IR (cm<sup>-1</sup>): 3508.2(w), 3043.5(w), 2970.4(w), 1605.5(m), 1585.3(w), 1484.2(m), 1465.7(m), 1446.4(m), 1432.4(m), 1288.4(w), 1272.5(w), 1250.0(m), 1213.6(m), 1153.7(w), 1080.7(vs), 1066.8(vs), 1043.2(s), 1015.6(m), 956.7(w), 940.4(w), 869.4(w), 852.8(m), 833.7(m), 756.0(vs), 731.3(m), 695.1(vs).

### *X-ray crystallography*

The structure of **1** was collected on an Xcalibur S single crystal diffractometer (Oxford Diffraction) using an enhanced Mo source. Each data reduction was carried out on the CrysAlisPro software package. For more detailed refinement information please consult the ESI. Full details can also be found in the CIF file: CCDC 898371.

## References

- [1] See for example: (a) V. K. Yachandra, V. J. DeRose, M. J. Latimer, I. Mukerji, K. Sauer and M. P. Klein., *Science*, 1993, **260**, 675. (b) V. J. DeRose, I. Mukerji, M. J. Latimer, V. K. Yachandra, K. Sauer and M. P. Klein, *J. Am. Chem. Soc.* 1994, **116**, 5239. (c) G. Aromí, M. W. Wemple, S. J. Aubin, K. Folting, D. N. Hendrickson and G. Christou., *J. Am. Chem. Soc.*, 1998, **120**, 5850. (d) T. Wydrzynski and S. Satoh., *Photosystem II: The Light-Driven Water: Palstpoquinone Oxidoreductase*; Springer: Dordrecht, The Netherlands, 2005. (e) J. Yano and V. K. Yachandra., *Inorg. Chem.*, 2008, **47**, 1711.
- [2] (a) G. Aromí and E. K. Brechin, *Struct. Bonding*, 2006, **122**, 1. (b) M. Murrie and D. J. Price, *Annu. Rep. Prog. Chem., Sect. A*, 2007, **103**, 20. (c) A. K. Boudalis, Y. Sanakis, C. P. Raptopoulou, V. Psycharis, *Magnetism and Superconductivity in Low-Dimensional Systems: Utilization in Future Applications*, D. Stamopoulos (Ed.) Nova Science Publishers Inc., New York, 2008, pp. 1-77; (d) G. Christou, *Polyhedron*, 2005, **24**, 2065. (e) *Molecular Cluster Magnets*, R. E. P. Winpenny (Ed.), World Scientific Books, Singapore 2011.
- [3] (a) C. Coulon, H. Miyasaka and R. Clérac, *Struct. Bonding*, 2006, **122**, 163. (b) L. Bogani, A. Vindigni, R. Sessoli and D. Gatteschi, *J. Mater. Chem.*, 2008, **18**, 4750. (c) H.-L. Sun, Z.-M. Wang and S. Gao, *Coord. Chem. Rev.*, 2010, **254**, 1081. (d) R. Lescouëzec, L. M. Toma, J. Vaissermann, M. Verdaguer, F. S. Delgado, C. Ruiz-Perez, F. L. Lloret and M. Julve, *Coord. Chem. Rev.*, 2005, **249**, 2691. (e) R. Clérac, H. Miyasaka, M. Yamashita and C. Coulon, *J. Am. Chem. Soc.*, 2002, **124**, 12837. (f) M. Ferbinteanu, H. Miyasaka, W. Wernsdorfer, K. Nakata, K. Sugiura, M. Yamashita, C. Coulon and R. Clérac, *J. Am. Chem. Soc.*, 2005, **127**, 3099.
- [4] (a) M. Evangelisti and E. K. Brechin, *Dalton Trans.*, 2010, **39**, 4672. (b) M. Evangelisti, F. Luis, L. J. de Jongh and M. Affronte, *J. Mater. Chem.*, 2006, 2534.
- [5] See for example: (a) R. Sessoli and A. K. Powell, *Coord. Chem. Rev.*, 2009, **253**, 2328. (b) T. C. Stamatatos, S. J. Teat, W. Wernsdorfer and G. Christou., *Angew. Chem. Int. Ed.*, 2009, **48**, 521. (c) M. N. Akhtar, V. Mereacre, G. Novitchi, J.-P. Tuchagues, C. E. Anson, A. K. Powell, *Chem. Eur. J.*, 2009, **15**, 7278. (d) J.-L. Liu, F.-S. Guo, Z.-S. Meng, Y.-Z. Zheng, J.-D. Leng, M.-L. Tong, L. Ungur, L. F. Chibotaru, K. J. Heroux and D. N. Hendrickson. *Chem. Sci.*, 2011, **2**, 1268. (e) J.-P. Costes, F. Dahan and A. Dupuis, *Inorg. Chem.*, 2000, **39**, 5994. (f) M. Andruh, J.-P. Costes, C. Diaz and S. Gao, *Inorg. Chem.*, 2009, **48**, 3342. (g) G. Karotsis, S. Kennedy, S. J. Teat, C. M. Beavers, D. A. Fowler, J. J. Morales, M. Evangelisti, S. J. Dalgarno and E. K. Brechin, *J. Am. Chem. Soc.*, 2010, **132**, 12983. (h) E. Colacio, J. Ruiz-Sanchez, F. J. White and E. K. Brechin, *Inorg. Chem.*, 2011, **50**, 7268.

- [6] (a) J. Rinck, G. Novitchi, W. Van den Heuvel, L. Ungur, Y. Lan, W. Wernsdorfer, C. E. Anson, L. F. Chibotaru and A. K. Powell. *Angew. Chem. Int. Ed.*, 2010, **49**, 7583. (b) A. Yamashita, A. Watanabe, S. Akine, T. Nabeshima, M. Nakano, T. Yamamura and T. Kajiwara. *Angew. Chem. Int. Ed.*, 2011, **50**, 1. (c) T. Peristeraki, M. Samios, M. Siczek, T. Lis and C. J. Milios. *Inorg. Chem.*, 2011, **50**, 5175. (d) T. N. Hooper, J. Schnack, S. Piligkos, M. Evangelisti and E. K. Brechin. *Angew. Chem. Int. Ed.*, 2012, DOI: 10.1002/aine.201200072. (e) C. E. Burrow, T. J. Burchell, P.-H. Lin, F. Habib, W. Wernsdorfer, R. Clérac and M. Murugesu, *Inorg. Chem.*, 2009, **48**, 8051.
- [7] (a) M. Affronte, S. Carretta, G. A. Timco and R. E. P. Winpenny. *Chem. Commun.*, 2007, 1789. (b) G. A. Timco, T. B. Faust, F. Tuna and R. E. P. Winpenny. *Chem. Soc. Rev.*, 2011, **40**, 3067. (c) T. B. Faust, V. Bellini, A. Candini, S. Carretta, G. Lorusso, D. R. Allan, L. Carthy, D. Collison, R. J. Docherty, J. Kenyon, J. Machin, E. J. L. McInnes, C. A. Muryn, H. Nowell, R. G. Pritchard, S. J. Teat, G. A. Timco, F. Tuna, G. F. S. Whitehead, W. Wernsdorfer, M Affronte and R. E. P. Winpenny. *Chem. Eur. J.*, 2011, **17**, 14020-14030. (d) C.F-Lee, D. A. Leigh, R. G. Pritchard, D. Schultz, S. J. Teat, G. A. Timco and R. E. P. Winpenny, *Nature*, 2009, **458**, 314. (e) G. A. Timco, S. Carretta, F. Troiani, F. Tuna, R. J. Pritchard, C. A. Muryn, E. J. L. McInnes, A. Ghirri, A. Candini, P. Santini, G. Amoretti, M. Affronte and R. E. P. Winpenny, *Nature Nanotechnology*, 2009, **4**, 173.
- [8] (a) O. Kahn, *Inorg. Chim. Acta.*, 1982, **62**, 3. (b) O. Kahn, J. Galy, Y. Journaux, J. Jaud and I. Morgenstern-Badarau, *J. Am. Chem. Soc.*, 1982, **104**, 2165. (c) O. Kahn. *Molecular Magnetism*, VCH, Weinheim. 1993.
- [9] (a) H. Oshio, M. Nihei, A. Yoshida, H. Nojiri, M. Nakano, A. Yamaguchi, Y. Karaki and H. Ishimoto. *Chem. Eur. J.*, 2005, **11**, 843. (b) M. Nihei, A. Yoshida, S. Koizumi and H. Oshio, *Polyhedron*, 2007, **26**, 1997.
- [10] For examples of the coordination chemistry of 2,2'-biphenol (LH<sub>2</sub>) see: (a) N. Berg, S. M. Taylor, A. Prescimone, E. K. Brechin and L. F. Jones, *CrystEngComm.*, 2012, **14**, 2732. (b) N. Berg, T. Rajeshkumar, S. M. Taylor, E. K. Brechin, G. Rajaraman and L. F. Jones, *Chem. Eur. J.*, 2012, **18**, 5906. (c) N. Berg and L. F. Jones, *CrystEngComm*, 2011, **13**, 5510. (d) A. R. Schake, E. A. Schmitt, A. J. Conti, W. E. Strieb, J. C. Huffman, D. N. Hendrickson and G. Christou, *Inorg. Chem.*, 1991, **30**, 3192. (e) D. P. Goldberg, A. Caneschi, C. D. Delfs, R. Sessoli and S. J. Lippard, *J. Am. Chem. Soc.*, 1995, **117**, 5789.
- [11] (a) C. J. O'Connor, D. P. Freyberg and E. Sinn, *Inorg. Chem.*, 1979, **18**, 1077. (b) H. Okawa, J. Nishio, M. Ohba, M. Tadokoro, N. Matsumoto, M. Koikawa, S. Kida and D. E. Fenton, *Inorg. Chem.*, 1993, **32**, 2949. (c) C. Fraser, R. Ostrander, A. L. Rheingold, C. White and B. Bosnich,

- Inorg. Chem.*, 1994, **33**, 324. (d) A. Hori, Y. Mitsuka, M. Ohba and H. Okawa, *Inorg. Chim. Acta.*, 2002, **337**, 113. (d) L. Bo, Z. Hong, P. Zhi-Quan, S. You, W. Cheng-Gang, Z. Han-Ping, H. Jing-Dong and C. Ru-An, *J. Coord. Chem.*, 2006, **569**, 1271.
- [12] (a) F. Birkelbach, M. Winter, U. Florke, H.-J. Haupt, C. Butzlaff, M. Lengen, E. Bill, A. X. Trautwein, K. Wiegardt and P. Chaudhuri, *Inorg. Chem.*, 1994, **33**, 3990. (b) Z.-W. Li, P.-P. Yang, L.-C. Li and D.-Z. Liao, *Inorg. Chim. Acta.*, 2009, **362**, 3381.
- [13] W. H. Press, S. A. Teukolsky, W. T. Vetterling, B. P. Flannery, "Numerical Recipes in C: The Art of Scientific Computing". Second Edition, Cambridge, Cambridge University Press, 1992.
- [14] J. M. Zadrozny, J. Liu, N. A. Piro, C. J. Chang, S. Hill and J. R. Long, *J. R. Chem. Commun.* **2012**, 48, 3927.
- [15] X. Feng, J. Liu, T. D. Harris, S. Hill and J. R. Long, *J. Am. Chem. Soc.* **2012**, 134, 7521.
- [16] A. Bencini and D. Gatteschi, *Electron Paramagnetic Resonance of Exchange Coupled Clusters* (Springer, Berlin, 1990).
- [17] E. Ruiz, S. Alvarez, A. Rodriguez-Forteza, P. Alemany, Y. Pouillon and C. Massobrio, in: J.S. Miller, M. Drillon (Eds.), *Magnetism: Molecules to Materials*, vol. II, Wiley-VCH, Weinheim, 2001.
- [18] (a) A. Abu-Nawwas, J. Cano, P. Christian, T. Mallah, G. Rajaraman, S. J. Teat, R. E. P. Winpenny and Y. Yasuhiko, *Chem. Commun.*, 2004, 314. (b) G. Rajaraman, J. Cano, E. K. Brechin and E.J.L. McInnes, *Chem. Commun.*, 2004, 1476. (c) G. Rajaraman, M. Murugesu, M. Soler, W. Wernsdorfer, M. Helliwell, S.J. Teat, G. Christou and E. K. Brechin, *J. Am. Chem. Soc.*, 2004, **126**, 15445.
- [19] J. Cirera, E. Ruiz, S. Alvarez, F. Neese and J. Kortus, *Chem. Eur. J.*, 2009, 15, 4078.
- [20] (a) D. Maganas, S. Sottini, P. Kyritsis, E. J. J. Groenen and F. Neese, *Inorg. Chem.*, 2011, **50**, 8741. (b) C. Duboc, D. Ganyushin, K. Sivalingam, M.-N. Collomb and F. Neese, *J. Phys. Chem. A.*, 2010, **114**, 10750. (c) F. Neese, *J. Am. Chem. Soc.*, 2006, **128**, 10213.
- [21] L. F. Jones, G. Rajaraman, J. Brockman, M. Murugesu, J. Raftery, S. J. Teat, W. Wernsdorfer, G. Christou, E. K. Brechin and D. Collison, *Chem. Eur. J.*, 2004, **10**, 5180.
- [22] R. Herchel, R. Boča, J. Krzystek, A. Ozarowski, M. Dura and J. van Slageren, *J. Am. Chem. Soc.* **2007**, 129, 10306.
- [23] L. Noodleman, *J. Chem. Phys.*, 1981, **74**, 5737.
- [24] (a) E. Ruiz, S. Alvarez, J. Cano and P. Alemany, *J. Comput. Chem.*, 1999, **20**, 1391. (b) E. Ruiz, A. R. Forteza, J. Cano, S. Alvarez and P. Alemany, *J. Comput. Chem.*, 2003, **24**, 982. (c) E. Ruiz, J. Cano, S. Alvarez, A. Caneschi and D. Gatteschi, *J. Am. Chem. Soc.*, 2003, **125**, 6791. (d) G. Rajaraman, J. Cano, E. K. Brechin and E. J. L. McInnes, *Chem. Commun.*, 2004, 1476.
- [25] S. Piligkos, G. Rajaraman, M. Soler, N. Kirchner, J. van Slageren, R. Bircher, S. Parsons, H. Güdel, J. Kortus, W. Wernsdorfer, G. Christou and E. K. Brechin, *J. Am. Chem. Soc.*, 2005, **127**, 5572.

- [26] P. Christian, G. Rajaraman, A. Harrison, M. Helliwell, J. J. W. McDouall, J. Raftery and R. E. P. Winpenny, *Dalton Trans.*, 2004, 2550.
- [27] E. Ruiz, S. Alvarez, A. Rodriguez-Forteza, P. Alemany, Y. Pouillon and C. Massobrio, in “*Magnetism: Molecules to Materials*”, Vol. II ed J. S. Miller, M. Drillon, Wiley-VCH, Weinheim, 2001, pp-227.
- [28] *Gaussian 09*, Revision A.1, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, Ö. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski, and D. J. Fox, Gaussian, Inc., Wallingford CT, 2009.
- [29] A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648.
- [30] A. Schafer, C. Huber and R. Ahlrichs, *J. Chem. Phys.*, 1994, **100**, 5829.
- [31] ORCA 2.9.0 F. Neese, 2012.
- [32] (a) J. P. Perdew, *Phys. Rev. B* 1986, **33**, 8822. (b) A. D. Becke, *Phys. Rev. A* 1988, **38**, 3098.
- [33] D. Ganyushin and F. Neese, *J. Chem. Phys.*, 2006, **125**, 024103.
- [34] F. Neese, *J. Chem. Phys.*, 2007, **127**, 164112.
- [35] K. Ray, A. Begum, T. Weyhermuller, S. Piligkos, J. van Slageren, F. Neese, K. Wieghardt, *J. Am. Chem. Soc.*, 2005, **127**, 4403.
- [36] F. Weigend, *Phys. Chem. Chem. Phys.*, 2006, **8**, 1057.