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Supramolecular chains of high nuclearity (Mn^{II}) barrel-like single molecule magnets
Supramolecular chains of high nuclearity \{\text{Mn}^{\text{III}}_{25}\} barrel-like single molecule magnets†

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The first application of 1-methyl-1H-pyrrole-2-carbaldehyde oxime as a ligand for the coordination of paramagnetic transition metal ions has afforded a new \(\text{Mn}^{\text{III}}_{25}\) barrel-like cluster linked via \text{Na}⁺ cations into a 1D polymeric topology that exhibits single-molecule magnetic behaviour.

The field of single-molecule magnets (SMMs) has grown considerably over the past two decades to include families of transition metal complexes incorporating paramagnetic ions with moderate-to-high oxidation states, as well as complexes assembled from trivalent lanthanide and actinide metal ions. Research efforts in this field have been largely driven by the two requirements for molecules to exhibit SMM behaviour namely, a bistable electronic ground state arising from an often large, ground state spin value (\(S\)), coupled with magnetic anisotropy, attributed to a negative zero-field splitting (ZFS) parameter (D). Such a combination affords an energy barrier (|\(D\)|\(|S^2\)) for reversal of the magnetization. Thus, at sufficiently low temperatures, these molecules function as nanoscale magnetic particles and straddle the classical/quantum interface by displaying not just classical magnetization hysteresis, but also quantum tunneling of magnetization (QTM) and quantum phase interference.5

In this context, SMMs have been proposed as promising candidates for potential applications in information storage and spintronics at the molecular level and for use as quantum bits in quantum computation.6 For SMMs to be incorporated into devices the molecules must be coupled to each other, and strategies to weakly couple SMMs while preserving their bistability are being actively pursued.7 Furthermore, in order for the magnetic properties of SMMs to be commercially viable, improvements in both their blocking temperatures and coercive fields are necessary. Strategies that enable the rational modification of the bridging units between SMMs represent a promising approach for maximizing these parameters.8,9 The self-assembly of SMMs into multi-dimensional architectures therefore remains a challenge for coordination chemists working at the interface of supramolecular chemistry and molecular magnetism.10

With respect to the preparation of novel families of SMMs, we and others have had success in developing high nuclearity 3d metal clusters based on 2-pyridyloxime ligands, \([\text{py})(\text{C}(R))\text{NOH}\) (\(R = \text{various}\); Scheme 1, left).8,11 More recently we have shown that the related 2-pyrrolyloximes (Scheme 1, middle) also act as bridging ligands for the coordination of paramagnetic metal centers. The simplest ligand initially chosen was pyrrole-2-carboxaldehyde oxime \(\text{praoH}_{2}(\text{R} = \text{Me, R}^0 = \text{H}; \text{Scheme } 1, \text{middle})\) which forms a family of Fe₃, Fe₆, Fe₁₀ and Fe₁₂ clusters, all with very small or zero \(S\) values.12 However, attempts to generate similar \text{Mn}^{\text{II}}⁠ complexes, which may offer large magnetic anisotropy, were thwarted by the presence of the pyrrole N–H functionality, a well-known reducing group. In order to inhibit such redox activity, we have turned our attention to the closely related ligand, 1-methyl-1H-pyrrole-2-carbaldehyde oxime \(\text{mpraoH}(\text{R} = \text{Me, R}^0 = \text{H}; \text{Scheme 1, middle})\). Here we show that the alkylated mprao⁻ ligand is robust with respect to redox activity and supports the formation of high nuclearity clusters. The title compound behaves as an SMM, and the diamagnetic

![Scheme 1](https://example.com/scheme1.png)

Scheme 1 The two families of 2-pyridyloxime (left) and 2-pyrrolyloxime (middle) ligands discussed in the text, and the crystallographically established coordination mode of mprao⁻ in complex 1 (right).
Na\(^+\) coordinated ions help organize the individual quantum spins on the SMMs into supramolecular chains.

Reaction of mpraoH with Mn(ClO\(_4\))\(_2\)-6H\(_2\)O, NEt\(_3\) and NaN\(_3\) in a 1:1:1:1 molar ratio in MeCN-MeOH (2:1 v/v) in air afforded a dark brown solution indicative of the presence of Mn\(^{III}\) and/or Mn\(^{IV}\) ions. Layering this solution with Et\(_2\)O afforded, after four days, X-ray quality, dark brown plate-like crystals of the 1D coordination polymer [Mn\(_{25}\)NaO\(_3\)(OMe)\(_2\)(hmp)\(_6\)(pdm)\(_6\)(pdmH)\(_6\)]\(^{8+}\) (pdmH\(_2\) = 2,6-pyridinedimethanol; hmpH = 2-(hydroxymethyl)pyridine), 13 the oxidation states of Mn\(^{III}\) and/or Mn\(^{IV}\) ions, as well as the deprotonated oximato arms of twelve \(\eta^1: \eta^2: \mu_2\) mprao \(^-\) groups (Fig. 1, top). The Mn\(_{25}\) core can be dissected into five parallel layers of three types with an ABCBA arrangement (Fig. 1, bottom). Layer A is a Mn\(^{III}\) triangular unit with a capping \(\mu_2\)-O\(^2-\) ion; layer B is a Mn\(^{III}\) hexagonal triangle that can be described as three corner-fused Mn\(^{III}\) triangular moieties each capped by a \(\mu_3\)-O\(^2-\) ion; and layer C is a disk-like Mn\(^{III}\) unit, reminiscent of the known Anderson-type structure. Each layer is held together and linked to its neighbours by a combination of oxido, methoxido, and/or oximato bridges. Whilst the core geometry is similar to that previously observed for the mixed-valence Mn\(^{III}\)\(_6\)Mn\(^{III}\)\(_8\)Mn\(^{IV}\)\(_8\) barrels [Mn\(_{25}\)O\(_{18}\)(OH)\(_2\)(N\(_3\))\(_{12}\)(pdm)\(_6\)(pdmH)\(_6\)]\(^{19+}\) and [Mn\(_{25}\)O\(_{18}\)(OH)-(OMe)(hmp)\(_6\)(pdm)\(_6\)(pdmH)\(_6\)]\(^{19+}\) (pdmH\(_2\) = 2,6-pyridinedimethanol; hmpH = 2-(hydroxymethyl)pyridine), 13 the oxidation states of the Mn ions in 1 are all assigned to Mn\(^{III}\). The oxidation states were determined qualitatively by inspection of metric parameters and detection of Mn\(^{III}\) Jahn-Teller (JT) elongation axes for all crystallographically independent Mn ions, as well as quantitatively by bond valence sum (BVS)\(^{14}\) calculations (Table S1, ESI†). In the case of the central Mn\(^{III}\) atom (Mn\(_0\)), which lies on a crystallographic inversion centre, both the intermediate for a Mn\(^{III}\)/Mn\(^{III}\) ion BVS value and the absence of a clear JT axis are assigned to the static disorder among the three symmetry-equivalent O(15,19,22)-Mn(6)-O(15′,19′,22′) axes.\(^{15}\)

Peripheral ligation about the Mn\(_{25}\) core is provided by twelve terminally bound N\(_3\) ions, four of which assemble end-to-end linking the Mn\(_{25}\) units to a Na\(^+\) ion at each end, affording a 1D linear chain (Fig. 2). The coordination sphere of the octahedral Na\(^+\) cations is completed by two axially bonded H\(_2\)O molecules. The shortest Mn···Mn separation between the Mn\(_{25}\) magnetic units is appreciably large (11.23(5) Å), while there are no other significant intermolecular interactions between the 1D chains. The overall barrel shape and nanometer size of the Mn\(_{25}\) unit in 1 are emphasized in the space-filling plots shown in Fig. S1 and S2 (ESI†); the cluster has a length of ~18.5 Å and a diameter of ~17.2 Å, excluding the H-atoms.

Solid-state, dc magnetic susceptibility (\(\chi_M\)) data for an air-dried sample of 1 were collected in the temperature range 5–300 K and in an applied field of 0.1 T. The data reveal that \(\chi_M\) steadily decreases from 73.48 cm\(^3\) K mol\(^{-1}\) at 300 K to 20.62 cm\(^3\) K mol\(^{-1}\) at 5.0 K (Fig. S3, ESI†). The 300 K value is slightly less than the spin-only (\(g = 2\)) value of 75 cm\(^3\) K mol\(^{-1}\) for 25 Mn\(^{III}\) non-interacting ions, indicating the presence of dominant antiferromagnetic exchange interactions within the cage. The 5 K value suggests that 1 possesses a fairly large ground state spin value of possibly \(S = 6\); the spin-only (\(g = 2\)) value for \(S = 6\) is 21 cm\(^3\) K mol\(^{-1}\). Given the size of the Mn\(_{25}\) molecule, and the resulting number of inequivalent exchange constants, it is not possible to determine the individual pairwise Mn\(_2\) exchange interaction parameters. Thus, we concentrated instead on characterizing the ground state spin, \(S\), and the zfs parameter, \(D\), by performing magnetization (\(M\)) vs. dc field measurements in a magnetic field and temperature range 1–70 kG and 1.8–10.0 K, respectively. Unfortunately, we were unable to obtain an acceptable fit for the data collected over the whole field range. This is a common problem in many large Mn clusters due to the population of low-lying excited states even at ultra-low temperatures, especially if some have an \(S\) value greater than that of the ground state.

A powerful complement to dc studies for determining the ground state of a system, and also to study magnetization dynamics, is ac magnetic susceptibility measurements which preclude any complications arising due to the presence of a dc field. These were performed in a 3.5 G ac field oscillating at different frequencies. The in-phase susceptibility (\(\chi'\))
shown as $\chi M/T$ vs. $T$ in Fig. S4 (ESI†) (top), and reveals several pertinent features: (i) $\chi M'/T$ decreases linearly with decreasing temperature in the 4–15 K range, indicating depopulation of a high density of excited states with spin $S$ greater than that of the ground state, which is in agreement with the conclusion of the dc studies; (ii) extrapolation of the $\chi M'/T$ data from above ~4.0 K to 0 K gives a value of ~21 cm$^3$ K mol$^{-1}$, which is indicative of an $S = 6$ ground state with $g = 2$; (iii) below ~4 K, there is a frequency-dependent decrease in $\chi M'/T$ and a concomitant appearance of frequency-dependent $\chi M'$ signals (Fig. S4, bottom, ESI†); such signals are an indication of the superparamagnetic-like slow relaxation of an SMM.

To confirm this last observation, magnetization vs. applied dc field data were collected on a single-crystal of 1.8MeOH down at 0.03 K on a micro-SQUID. Hysteresis loops are seen below ~0.8 K whose coercivities increase with increasing sweep rate (Fig. 3, top) and with decreasing temperature (Fig. 3, bottom), confirming 1 to be an SMM. These loops do not show the steps characteristic of quantum tunnelling, as expected for large SMMs which are more susceptible to various step-broadening effects.\textsuperscript{10,13,16}

It should be noted that complex 1 is simply too complicated to rationalize the $S = 6$ ground state on the basis of its structure alone. There are extensive spin frustration effects operating within the many fused Mn$_3$ triangular units, which render any spin assignment inaccurate and superficial.

In conclusion, we have established that the $N$-alkylation of a 2-pyrrolyloxime stabilizes the molecule with respect to oxidation. The resultant mpao$^-$ ligand is therefore able to bridge Mn$^{II}$ centres affording a large polynuclear barrel-shaped [Mn$_{10}$Cl$_{35}$] cluster that is structurally related to the examples of mixed-valence Mn$_{15}$ barrel-like SMMs reported in the literature.\textsuperscript{13} To date, this system represents the highest nuclearity Mn cluster which is organized into a 1D polymer through chelation with diamagnetic metal centers.\textsuperscript{17} Given the apparent stability of these barrel-like clusters in multiple oxidation states, the redox activity of 1 is being actively pursued. Furthermore, strategies aimed at the selective replacement of the diamagnetic Na$^+$ cations with paramagnetic 4f-metal ions, targeting the rational assembly of 1 into new classes of SCMs, are currently under investigation.

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Notes and references

\textsuperscript{5} W. Wernsdorfer and R. Sessoli, \textit{Science}, 1999, 284, 133.