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Two $[\text{Mn}_3(\mu_3\text{-O})]^{7+}$ based single chain magnets with different solvent ligation[†]

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ABSTRACT

$[\text{Mn}^{\text{III}}_3(\mu_3\text{-O})]^{7+}$ based EE-azido-bridged 1D chains $[\{\text{Mn}_3\text{O}(\text{5-Br-salox})_3(\text{N}_3)(\text{H}_2\text{O})_4\}.3\text{H}_2\text{O}\}]_n$ (**1**) and $[\{\text{Mn}_3\text{O}(\text{5-Br-salox})_3(\text{N}_3)(\text{H}_2\text{O})_3(\text{DMF})\}.2\text{H}_2\text{O.DMF}\}]_n$ (**2**) (5-Br-saloxH₂ = 2-hydroxy-5-bromo-benzaldehyde oxime) have been synthesized and characterized by single crystal X-ray diffraction analysis. The complex **1** was synthesized by the reaction between ligand and metal precursor in the presence of azide ion in MeOH, while complex **2** can be obtained in two ways: in the first method, the same reaction as in **1**, with only difference in solvent composition – a mixture of solvents DMF/MeOH (2:3/ v/v) was used. In another method, **2** was isolated simply by the recrystallization of **1** from DMF. Magnetic studies on these complexes reveal that both **1** ($\Delta E = 45.9 \text{ K}$; $\tau_0 = 6.6 \times 10^{-10}$) and **2** ($\Delta E = 54$ and 49.7 K ; $\tau_0 = 1.4 \times 10^{-12} \text{ s}$ and $7.6 \times 10^{-12} \text{ s}$) exhibit SCM behaviour with a slight difference in magnetic properties due to the only difference in coordinated/non-coordinated solvents.

1.Introduction

The development of molecule-based magnetic materials like Single Molecule Magnets (SMMs) and Single Chain Magnets (SCMs) is now an important research topic in the contemporary chemistry and physics[1] due to their characteristic features that include quantum phenomena[2] and finite size effects[3] and their potential applications in information storage at the molecular level [4]. The design of SCMs requires large uniaxial anisotropy, strong intrachain magnetic interactions between the high-spin magnetic units, and good isolation of the chains in order to prevent two-dimensional (2D) and three- dimensional (3D) ordering. This has opened up a new strategy to create SCM with exciting new perspectives to store information in low-dimensional materials. However, it took almost forty years to get the first experimental evidence of such behaviour in a real one-dimensional SCM compound $[\text{Co}(\text{hfac})_2(\text{NITPhOMe})]$ [5]. The 1D chains with oxide-centered trinuclear manganese units $[\text{Mn}^{\text{III}}_3(\mu_3\text{-O})]^{7+}$ as the building block and N_3^- , RCOO^- etc as linkers have been well known in recent years[6]. It was interesting to observe that magnetic

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