



Cite this: *RSC Adv.*, 2019, 9, 37740

A new member of a class of rod-like Mn₁₂ single molecule magnets using 2-(pyridine-2-yl)propan-2-ol†

En-Che Yang,^a Shi-Yi Huang,^a Wolfgang Wernsdorfer,^b Ling-Xuan Hong,^a Marko Damjanovic,^b Lukas Niekamp^b and Gene-Hsiang Lee^c

This paper reports on the synthesis, structure and magnetic properties of a new type of rod like Mn₁₂ metal cluster, [Mn₁₂O₇(OH)₂(OMe)₂(dmhmp)₄(O₂CPh)₁₁(H₂O)] (6) where the ligand (dmhmpH) is 2-(pyridine-2-yl)propan-2-ol. Compound (6) was obtained by reacting MnCl₂·4H₂O with dmhmpH in the presence of benzoic salt and Et₃N. The resulting crystalline material is assigned to the triclinic space group *P* $\bar{1}$. Although compound (6) displays ferromagnetic and antiferromagnetic competition behavior, this does not prevent the molecule from functioning as a single-molecule magnet (SMM). The SMM behavior is evidenced by observing frequency dependent out-of-phase ac signals as well as magnetization hysteresis loops at low temperatures in a micro-SQUID study. A brief comparison between all rod-like Mn₁₂ materials is also given in the manuscript.

Received 13th August 2019
Accepted 12th November 2019

DOI: 10.1039/c9ra06280g

rsc.li/rsc-advances

Introduction

Because of their natural abundance and the fact that they are involved in numerous biologically related processes, 1st row transition metal ions have attracted the interest of the scientific community.¹ For example, manganese plays a key role in photo system II (PSII).² Oxo-bridged manganese clusters are of great interest because of their reactivity, redox properties, ferroelectricity as well as their magnetochemistry.³ In applications in magnetochemistry, the most important branch is single-molecule magnets (SMMs).⁴ Single molecule magnets (SMMs) are metal clusters that simultaneously have large molecular spins and significant negative magneto-anisotropy. Upon combining these two factors, an energy barrier for spin flip is created. The manganese ion, especially Mn(III), due to its high spin number *S* = 2, and the negative anisotropy parameter resulting from the elongation of Jahn–Teller distortion, make them ideal candidates for constructing SMMs.⁵ Since the discovery of the first SMMs [Mn₁₂O₁₂(O₂CR)₁₆(H₂O)_{*x*}] (*x* = 3 or 4) in 1993,⁶ a dynasty of SMMs has arisen, not only concerning their magnetism but also metal-oxo chemistry.

To control the structure for SMMs, one of the most important factors is the geometry of the ligand. The most widely used ligand for the construction of SMMs, except for the carboxylic acid, is hydroxymethylpyridine (hmpH).⁷ By using hmpH and hydroxyethylpyridine (hepH), Christou and co-workers successfully prepared a new type of “rod-like” Mn₁₂: [Mn₁₂O₈Cl₄(O₂CPh)₈(hmp)₆] (1), [Mn₁₂O₈Cl₄(O₂CPh)₈(hep)₆] (2) and [Mn₁₂O₈Br₄(O₂CPh)₈(hep)₆] (3) in 2002.⁸ In 2009, they changed the ligands from hmpH to dmhmpH dimethylhydroxymethylpyridine. Afterward, [Mn₇O₃(OH)₃(O₂CtBu)₇(dmhmp)₄] (4) and [Mn₁₂O₇(OH)(MeO)₂(O₂CPh)₁₂(dmhmp)₄(H₂O)] (5) were prepared by two very different reaction procedures.⁹ At first glance, (4) and (5) appear to have very different structures, which can be attributed to very different reaction conditions used. In the process of modifying the reaction conditions for preparing Mn₇ (4), but switching the carboxylate ligand from pivalic to benzoic acid and the use of a more MeOH rich solution system, quite surprisingly, we obtained a new Mn₁₂ complex. This compound has the formula [Mn₁₂O₇(OH)₂(OMe)₂(dmhmp)₄(O₂CPh)₁₁(H₂O)] (6) which is different from compound (5). In this paper, we report on a detailed study of this accidentally prepared compound and also compare it with the other rod-like Mn₁₂ complexes.

Results and discussion

Synthesis and structures

Compound 6 was produced by treating dmhmpH with MnCl₂·4H₂O, NaO₂CPh, in the presence of Et₃N in a solution of MeCN/MeOH (1/29 mL). It is important to use a ratio of MeCN/

^aDepartment of Chemistry, Fu Jen Catholic University, Hsinchuang, New Taipei City, 24205, Taiwan, Republic of China. E-mail: 071549@mail.fju.edu.tw; Tel: +886-2-2905-3571

^bInstitute of Nanotechnology (INT), Karlsruhe Institute of Technology (KIT), Hermann-von-Helmholtz-Platz 1, D-76344 Eggenstein-Leopoldshafen, Germany

^cInstrumentation Centre, College of Science, National Taiwan University, Taipei, 10672, Taiwan, Republic of China

† Electronic supplementary information (ESI) available. CCDC 1945849. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9ra06280g



MeOH that is rich in MeOH, otherwise desired compound **6** is not produced in high yields.

The ORTEP plot for compound **6** is shown in Fig. 1. The metal–ligand bond lengths are listed in Table 1. On calculating the charge by their structural data, and combining these with the BVS result,¹⁰ unlike compound **1–3** where the cation has a charge of +34, compound **6**, with a charge +33, is more similar to compound **5**. Although, unlike compound **1–3** which possess inversion symmetry, all of these complexes have similar structural characteristics with a central $[\text{Mn}_4\text{O}_6]$ unit of uncompleted dicubane connected with two $[\text{Mn}_4\text{O}]$ tetrahedral units on both two ends of the rod. Compounds **6** and **5** have no intramolecular symmetry. All of compounds **1–3** contain two Mn(II) (Mn(6) and Mn(6') in ref. 8) ions located on the two tips on the end of the rod. In compounds **5** and **6**, there are three Mn(II) ions, one of which being located on the tip of one end, the other two being located on a lateral of the tetrahedral on the other end but does not occupy the tip position. In a comparison of compounds **5** and **6**, their ligands are obviously different. Compound **5** has twelve PhCOO^- units, while compound **6** has eleven PhCOO^- units. A detailed study of the structure of **6** revealed that the whole metal-oxo core is enclosed in a package formed from four dmhmp^- ligands and eleven PhCOO^- units, where each

dmhmp^- binds a Mn(III) ion. The PhCOO^- units have three different binding modes: one monodentate η^1 type coordinated with one manganese ion, another type μ_2 that bridges two metal ions in a $\eta^1:\eta^1$ fashion. The final μ_3 bridges three metal ions in a $\eta^1:\eta^2$ manner. (For detailed information on the binding modes, see Fig. S1.†)

Magnetic susceptibility study of compound **6**

Variable temperature dc magnetic susceptibility measurements were carried on a polycrystalline sample of compound **6** restrained by parafilm under a 1000 G magnetic field in the temperature range of 2.0–300 K. The $\chi_{\text{M}}T$ vs. T data are shown in Fig. 2. It can be seen that the $\chi_{\text{M}}T$ value is $27.3 \text{ cm}^3 \text{ K mol}^{-1}$ at 300 K, which is significantly smaller than the spin-only value: $40.13 \text{ cm}^3 \text{ K mol}^{-1}$ for nine uncoupled Mn(III) and three Mn(II) ions. This phenomenon indicates that antiferromagnetic coupling plays an important role at room temperature. The $\chi_{\text{M}}T$ values then steadily decrease to $20.0 \text{ cm}^3 \text{ K mol}^{-1}$ as the temperature is cooled down to 50 K and then is nearly constant down to 4 K followed by a drop to $16.8 \text{ cm}^3 \text{ K mol}^{-1}$ at 2 K. The abnormal feature of the $\chi_{\text{M}}T$ values indicate a ferromagnetic and antiferromagnetic competition behavior in compound **6**. Nevertheless, the $\chi_{\text{M}}T$ value down to 2.0 K roughly gives a $S = 11/2$ ground state with a g value of 1.95. The g value deviating from 2.0 can be attributed to intermolecular antiferromagnetic coupling and/or zero-field splitting.

The spin number as well as the zero-field splitting parameters were then obtained by variable fields, variable temperatures reduced magnetization measurements. Based on the known information regarding compound **6**, it is a ferromagnetic and antiferromagnetic competition case in which low lying high spin excited states may approach to the ground state in energy. To avoid the interference of high spin excited states and to obtain the parameter of the low spin ground state, a lower magnetic field should be applied.¹¹ Therefore, in this experiment, magnetic fields in the range from 1000 G to 9000 G were applied in the temperature range of 2–4 K. The experimental results are illustrated in Fig. 3, where the solid red lines are the fittings based on eqn (1).

$$\hat{H} = g\beta\hat{S} + D \left[\hat{S}_z^2 + \frac{1}{3}S(S+1) \right] \quad (1)$$

It can be seen that the eqn (1) gives a reasonably good fitting and reasonable spin parameters are obtained which gives $S = 11/2$, $g = 2.0$, $D = -0.34 \text{ K}$. Based on this set of parameters, we postulate that compound **6** can function as a single-molecule magnet (SMM).

a.c. magnetic susceptibility

To further confirm that compound **6** is, in fact, a SMM, a.c. magnetic susceptibility measurements were carried. In the first attempt, we applied typical measurements on the sample which applies to an a.c. magnetic field 3.5 G in the temperature range 1.9–9.0 K with the frequencies below 1000 Hz. Although frequency-dependent out-of-phase signals were seen, none of

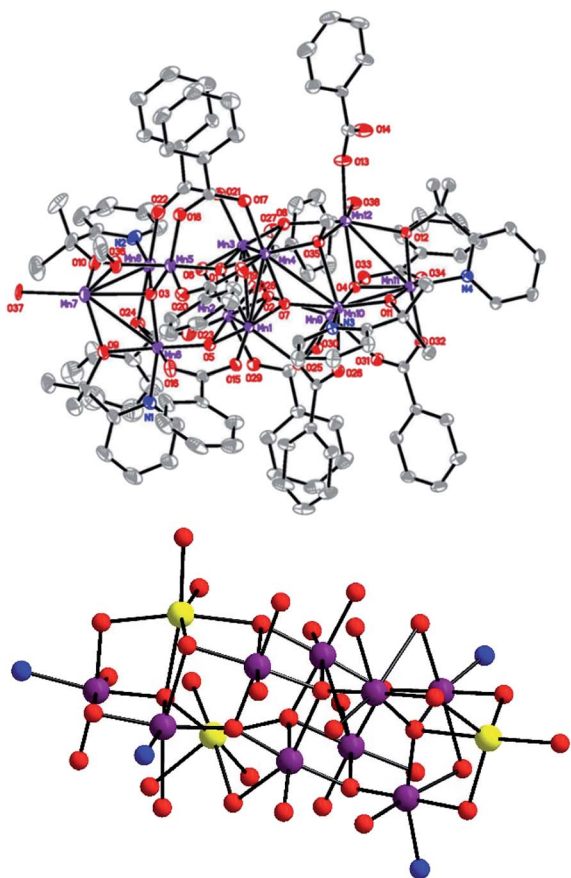


Fig. 1 (Top) ORTEP plot of complex **6** at the 30% probability level (Mn, purple; O, red; N, blue; C, gray). (Bottom) Framework of compound **6** (Mn^{II} yellow, Mn^{III} purple, O red, N blue).

Table 1 Metal–Ligand bond lengths of compound 6

Mn(1)–O(5)	1.877(2)	Mn(5)–O(1)	1.850(2)	Mn(9)–O(30)	2.138(2)
Mn(1)–O(7)	1.899(2)	Mn(5)–O(36)	1.894(2)	Mn(9)–O(4)	2.1532(19)
Mn(1)–O(15)	1.968(2)	Mn(5)–O(3)	1.930(2)	Mn(9)–O(33)	2.157(2)
Mn(1)–O(2)	1.968(2)	Mn(5)–O(18)	2.015(2)	Mn(9)–O(31)	2.195(2)
Mn(1)–O(25)	2.212(2)	Mn(5)–O(20)	2.101(3)	Mn(9)–O(2)	2.279(2)
Mn(1)–O(1)	2.271(2)	Mn(6)–O(5)	1.849(2)	Mn(9)–O(25)	2.321(2)
Mn(2)–O(6)	1.929(2)	Mn(6)–O(9)	1.896(2)	Mn(10)–O(7)	1.873(2)
Mn(2)–O(29)	1.953(2)	Mn(6)–O(3)	1.900(2)	Mn(10)–O(11)	1.877(2)
Mn(2)–O(23)	2.004(3)	Mn(6)–N(1)	2.020(3)	Mn(10)–O(4)	1.906(2)
Mn(2)–O(2)	2.007(2)	Mn(6)–O(16)	2.117(3)	Mn(10)–N(3)	2.039(3)
Mn(2)–O(28)	2.069(3)	Mn(6)–O(24)	2.479(3)	Mn(10)–O(26)	2.214(2)
Mn(2)–O(5)	2.083(2)	Mn(7)–O(36)	2.088(3)	Mn(10)–O(35)	2.377(2)
Mn(3)–O(6)	1.882(2)	Mn(7)–O(9)	2.142(3)	Mn(11)–O(12)	1.889(2)
Mn(3)–O(8)	1.923(2)	Mn(7)–O(10)	2.150(3)	Mn(11)–O(4)	1.894(2)
Mn(3)–O(2)	1.932(2)	Mn(7)–O(37'')	2.190(8)	Mn(11)–O(32)	1.942(2)
Mn(3)–O(21)	1.972(2)	Mn(7)–O(37')	2.253(9)	Mn(11)–N(4)	2.045(3)
Mn(3)–O(27)	2.186(3)	Mn(7)–O(3)	2.301(2)	Mn(11)–O(34)	2.120(2)
Mn(3)–O(1)	2.329(2)	Mn(7)–O(37)	2.341(4)	Mn(11)–O(11)	2.277(2)
Mn(4)–O(7)	1.880(2)	Mn(8)–O(6)	1.874(2)	Mn(12)–O(13)	2.132(2)
Mn(4)–O(1)	1.884(2)	Mn(8)–O(10)	1.885(3)	Mn(12)–O(12)	2.134(2)
Mn(4)–O(35)	1.940(2)	Mn(8)–O(3)	1.913(3)	Mn(12)–O(8)	2.159(2)
Mn(4)–O(17)	1.970(2)	Mn(8)–N(2)	2.018(3)	Mn(12)–O(38)	2.226(3)
Mn(4)–O(19)	2.158(2)	Mn(8)–O(22)	2.187(3)	Mn(12)–O(4)	2.269(2)
Mn(4)–O(8)	2.303(2)	Mn(8)–O(24)	2.301(3)	Mn(12)–O(35)	2.289(2)

the signal showed peak feature above 1.9 K. We therefore examined a wider scope of oscillating frequencies from 10 to 10 000 Hz in the temperature range 1.9–6.1 K. Fig. 4 presents the results of temperature dependent and frequency dependent ($\log \omega$) scans, which show two peaks above 1.9 K. The energy barrier for spin flipping was estimated by the Arrhenius formula:¹²

$$\ln(\tau) = \ln(\tau_0) + U_{\text{eff}}/kT \quad (2)$$

which is shown in the inset of Fig. 4 to give a value of $U_{\text{eff}} = 9.2$ K. This result is somewhat lower than the theoretical value $D(S^2 - 0.25) = 10.2$ K. This implies that somewhat quantum tunnelling of magnetization (QTM)¹³ occurs at this moment.

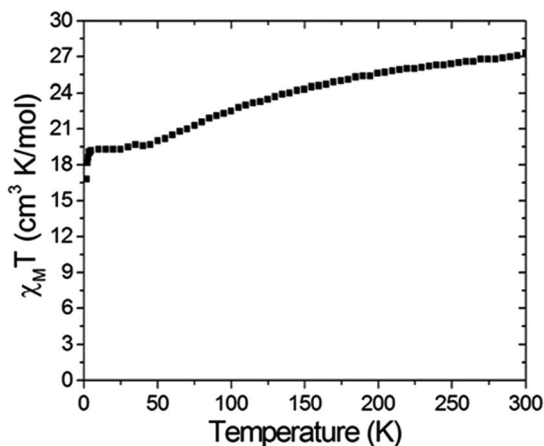


Fig. 2 A $\chi_M T$ versus T plot of complex 6 measured under a 1 kG magnetic field in the temperature range 2–300 K.

Armed with this data, it can be concluded that compound 6 is a SMM.

Further detail a.c. magnetic susceptibility examinations regarding magnetization relaxation process was analysed by the Debye model of Argand's plot (χ'_M vs. χ''_M plot) which is also known as Cole–Cole plot.¹⁴ The Debye model is described as eqn (3) and (4):

$$\chi'(\omega) = \chi_S + \frac{(\chi_T - \chi_S) \left[1 + (\omega\tau)^{1-\alpha} \sin\left(\frac{\alpha\pi}{2}\right) \right]}{1 + 2(\omega\tau)^{1-\alpha} \sin\left(\frac{\alpha\pi}{2}\right) + (\omega\tau)^{2(1-\alpha)}} \quad (3)$$

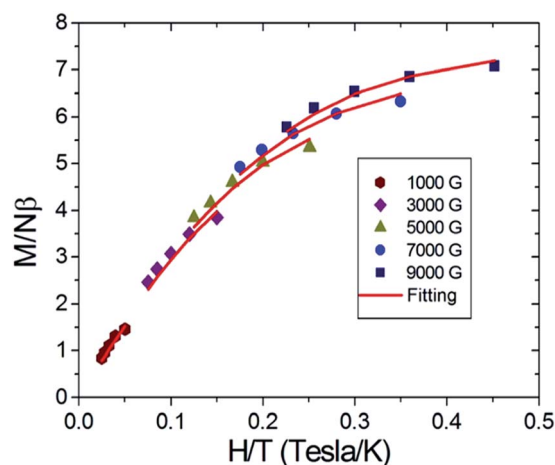


Fig. 3 Reduced magnetization of complex 6 under magnetic field 0.1–0.9 T in the temperature range 2–4 K. Red solid lines denote the best fitting to the experimental data which give: $S = 11/2$, $g = 2.0$, and $D = -0.34$ K.

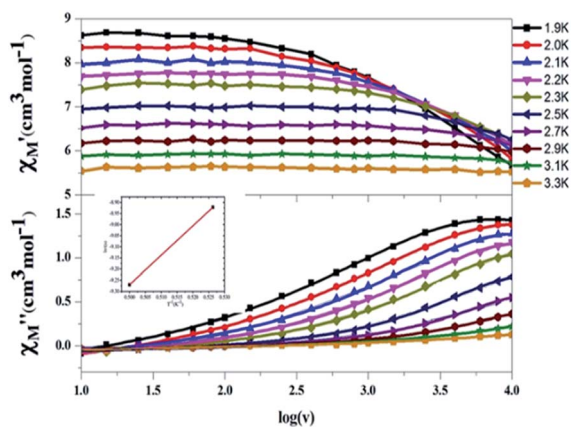


Fig. 4 AC magnetic susceptibility data for complex **6** collected in a 3.5 G oscillating field with frequencies of 10–10 000 Hz in the temperature range 1.9–3.3 K. The in-phase signals χ_M' are presented in the top, and the out-of-phase signals χ_M'' are in the bottom. Inset: Arrhenius plot. The red solid line represents the least-squares fit.

$$\chi''(\omega) = \frac{(\chi_T - \chi_S)(\omega\tau)^{1-\alpha} \cos\left(\frac{\alpha\pi}{2}\right)}{1 + 2(\omega\tau)^{1-\alpha} \sin\left(\frac{\alpha\pi}{2}\right) + (\omega\tau)^{2(1-\alpha)}} \quad (4)$$

where χ_M' and χ_M'' are the in-phase and out-of-phase a.c. magnetic susceptibilities. Both are the functions of scanning frequencies ω . The parameters χ_S represents the adiabatic susceptibility, χ_T the isothermal susceptibility and τ the magnetization relaxation time. The value for parameter α varies in the range 0–1 and provide an estimation of the distribution of relaxations. An Argand's plot for compound **6** at 1.9 K is shown in Fig. 5. The parameters in eqn (3) and (4) obtained from the best fitting are given as $\chi_T = 11.64 \text{ cm}^3 \text{ mol}^{-1}$, $\chi_S = 4.09 \text{ cm}^3 \text{ mol}^{-1}$, $\alpha = 0.35$, and $\tau = 2.03 \times 10^{-3} \text{ s}$. From this result, we can conclude a distribution of single relaxation for compound **6**.

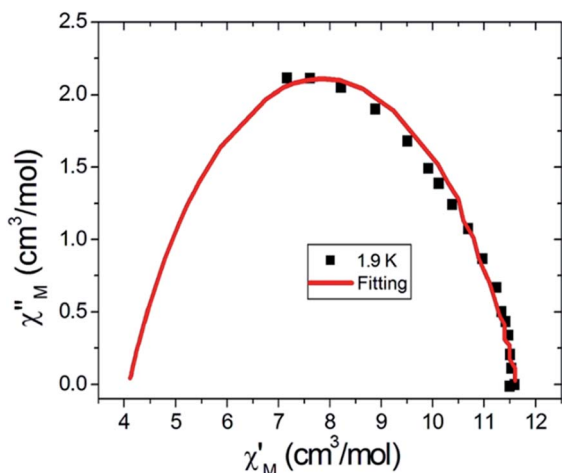


Fig. 5 Cole–Cole plot at 1.9 K for complex **6**. The red solid line is the least-square fit of the data to eqn (3) and (4).

This was also observed in many other manganese based single molecule magnets (SMMs).

Magnetization hysteresis studies

Magnetization hysteresis measurements were then carried on a single crystal of **6** in a micro-SQUID array with the magnetic field aligned with the easy axis of the crystal. Fig. 6(a) presents the hysteresis loops observed in the temperature range 0.025–0.9 K. Obviously, the hysteresis loops can be seen at temperatures below 0.9 K providing an indication of magnet type behavior. Upon cooling, the magnitude for the coercivity of each loop increases until the lowest reachable temperature 25 mK, which reflects the fact that slower magnetization relaxation occurs at lower temperatures. Since the magnetization relaxation process appears to be dominated by quantum tunnelling of the magnetization (QTM) at 25 mK, to explore the nature of the QTM, we examined the hysteresis loops when the temperature is maintained at 25 mK but varying the scan rate in the range of 0.004 T s^{-1} to 0.128 T s^{-1} . Several points are worth mentioning in Fig. 6(b). First, all the loops show a giant transition at zero magnetic field that can be attributed to the QTM process between the ground spin states. The fact that this occurs at zero field indicates that there is no significant intermolecular

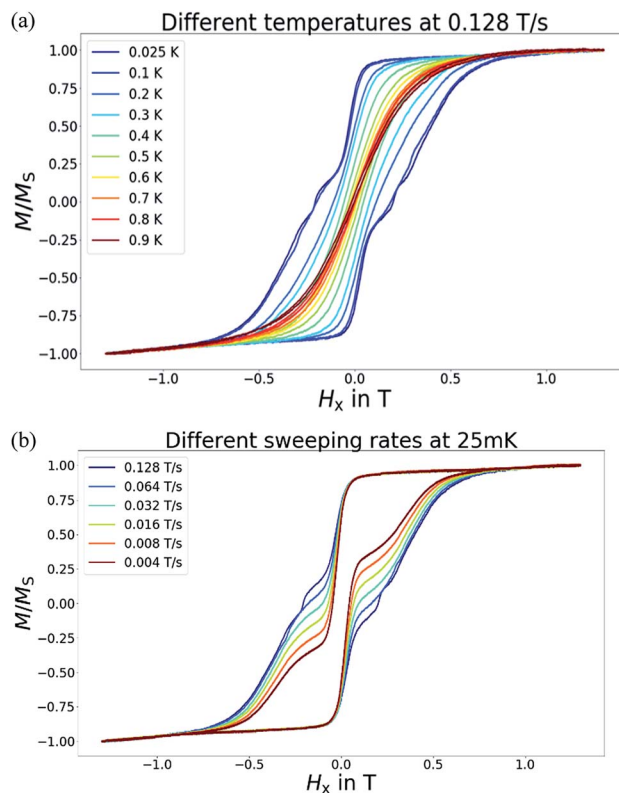


Fig. 6 Magnetization hysteresis measurements for compound **6**. (a) Experiments were performed at a scanning rate of 0.128 T s^{-1} in the temperature range of 0.025–0.9 K. (b) Hysteresis loops were collected at a constant temperature (25 mK) with scanning rates in the range 0.004 – 0.128 T s^{-1} .

interaction to shift the degeneracy of the ground states of spin multiplicity. In another words, no significant intermolecular exchange bias¹⁵ occurs in this crystal system. Secondly, the step occurring at zero field increase in size when the scanning rate is slowed. This seems to be the result of a longer time interval the molecule system stays in the QTM window the larger is the QTM amplitude and therefore bigger transition steps are seen. Compared with most typical manganese SMMs [$\text{Mn}_{12}\text{O}_{12}(\text{O}_2\text{-CR})_{16}(\text{H}_2\text{O})_{3\sim 4}$],¹⁶ the step feature for compound **6** is significantly larger, we therefore conclude that compound **6** belongs to the category of SMMs with a fast QTM. Of course, compound **6** possesses the molecular spin $S = 11/2$, a half integer spin that might hinder the QTM process. Nevertheless, a combination of a hyperfine interaction of ⁵⁵Mn ($I = 5/2$, 100%) as well as the lack of symmetry axis of C_3 and/or above which introduces a transverse field, can still trigger a fast QTM in compound **6**.¹⁷

Comparisons with other rod-like Mn_{12} complexes

According to the literature, four rod-like Mn_{12} complexes have been reported in which two are SMMs. Combining with the compound **6**, all five compounds were synthesized using a hydroxyl-alkyl-pyridine type of ligand. Changing the alkyl group slightly causes significant differences in the magnetic properties of these complexes. In compound **1**, pure hmp ligands with alkyl = methylene leads to a SMM with $S = 7$ (or 6), all of the ac frequencies above 10 Hz have their out-of-phase signals above 1.9 K. Compounds **2** and **3**, both contain hep ligands with alkyl = ethyl groups. No SMMs behavior was observed in this series compounds which have molecular spin $S = 0$. Compound **5** and the current work **6** contain dmhmp with alkyl = isopropyl groups. Although both exhibit SMM behavior, their spin parameters are somewhat different. Compound **5** has molecular spin $S = 13/2$ and all of its ac out-of-phase signals show only a tail above 1.9 K. However, promising magnetization hysteresis loops have been reported at temperatures below 0.7 K. Thus, compound **5** is a SMM. The current compound **6** also contains alkyl = isopropyl. However, its peripheral ligands are arranged slightly different from those for compound **5** with two coordinated hydroxide ion and a dangling benzoic acid ligand. In this compound, frequency-dependent out-of-phase signals are seen, and two peaks above 1.9 K are seen with frequencies in the interval 1000–10 000 Hz. Solid magnetization hysteresis loops are seen below 0.9 K. The obtained molecular spin for compound **6** was $S = 11/2$. Hysteresis measurements indicates that this molecular system is a fast QTM of SMM. Details regarding the comparison of these five compounds are given in Table 2 with their plots in the ESI (Fig. S2†).

In conclusion, the Mn_{12} rod-like complex reported in this study exhibits competing ferromagnetic and antiferromagnetic exchange interactions. As expected because Mn(II) ions are present in its structure these interactions are very weak leading to the existence of several low-lying excited states, as was also evidenced from variable field-variable temperature magnetization studies. Minor structural changes in the structure of such complexes can lead to small changes of the exchange interactions between the Mn ions, thus leading to a different spin

ground state value. This explains why the S values of compounds **5** and **6** are different.

Experimental

All chemicals used in this study were of commercial grade and were used without further purification. The ligand dmhmpH was synthesized according to a literature report.¹⁸

$[\text{Mn}_{12}(\text{dmhmp})_4(\text{PhCO}_2)_{11}(\text{CH}_3\text{O})_2(\text{H}_2\text{O})(\text{OH})_2\text{O}_7]$ (**6**)

In a 100 mL beaker, the ligand dmhmpH 0.13 g (1 mmol) and triethylamine Et_3N 0.14 mL (1 mmol) were dissolved in a mixed solvent MeCN/MeOH (1/29 mL). $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$ 0.2 g (1 mmol) and NaO_2Ph 0.29 g (2 mmol) were added and the mixture was then stirred overnight. After filtration, diethylether was allowed to diffuse into the filtrate. Dark black crystals suitable for X-ray crystallographic analysis was obtained after two weeks. Selected IR data (cm^{-1}): 3435(br), 2973(m), 1602(s), 1557(s), 1482(s), 1392(s), 1122(s), 1052(m), 978(s), 884(s), 840(s), 781(s). Anal. Calcd (found) for $\text{C}_{111}\text{H}_{104}\text{N}_4\text{ClMn}_{12}\text{O}_{37}$: C, 47.92 (47.75); H, 3.77 (3.44); N, 2.01 (1.89).

X-ray crystallography

X-ray analysis data for **6** was collected on a Bruker D8 Venture diffractometer with Mo $K\alpha$ radiation ($\lambda = 0.71073 \text{ \AA}$). The temperature was cooled to 200(2) K. The corresponding data was collected by Bruker APEX3 program, and simplified by Bruker SAINT program. Data were corrected by the empirical absorption SADABS program.¹⁹ The SHELXTL program on a PC computer was used for the structure analysis. The structure was solved by direct method of the Shelxs program²⁰ and refinement was done by the Shelxl program²¹ based on the full-matrix least squares on F^2 values. Non-hydrogen atoms were refined anisotropically. Hydrogen atoms were fixed at the calculated positions and refined using a riding mode. Table 3 summarizes the structure refinement parameters for complex **6**.

Physical property measurements

NMR spectra were collected on a Bruker AV-300 spectrometer. Infrared spectra were obtained using KBr pellets on a PerkinElmer 1600 spectrometer in the 450–4000 cm^{-1} range. DC magnetic measurements were collected using a Quantum Design MPMS7 system. Samples were restrained with parafilm to prevent torqueing. The magnetic background caused by the gel cap and the parafilm were subtracted by blank measurements. The diamagnetism correction was estimated from Pascal's constant.²² Elemental analyses (C, H, N) and dc magnetic susceptibility measurements were carried by the National Taiwan University Instrument Centre, College of Science. AC magnetic susceptibility data were collected using the Quantum Design MPMS XL7 system of the National Chiao Tung University. Oriented single crystal magnetization hysteresis loops were measured by employing a micro-SQUID array that was described elsewhere.²³ A single crystal was put onto the array and external field is oriented parallel to the crystal easy axis.

Table 2 Comparisons of all rod-like Mn₁₂ complexes^a

Complex	S	D (cm ⁻¹)	U _{eff} (K)
(1) [Mn ₂ ^{II} Mn ₁₀ ^{III} O ₈ Cl ₄ (O ₂ CPh) ₈ (hmp) ₆]	6(7)	-0.40	29.1
(2) [Mn ₂ ^{II} Mn ₁₀ ^{III} O ₈ Cl ₄ (O ₂ CPh) ₈ (hep) ₆]	0	—	—
(3) [Mn ₂ ^{II} Mn ₁₀ ^{III} O ₈ Br ₄ (O ₂ CPh) ₈ (hep) ₆]	0	—	—
(5) [Mn ₃ ^{II} Mn ₉ ^{III} O ₇ (OH)(OMe) ₂ (O ₂ CPh) ₁₂ (dmhmp) ₄ (H ₂ O)]	13/2	-0.18	11.5
(6) [Mn ₃ ^{II} Mn ₉ ^{III} O ₇ (OH) ₂ (OMe) ₂ (O ₂ CPh) ₁₁ (dmhmp) ₄ (H ₂ O)]	11/2	-0.24	9.2

^a hmpH 2-(hydroxymethyl)pyridine – hepH 2-(hydroxyethyl)pyridine – dmhmpH 2-(pyridine-2-yl)propan-2-ol.

Table 3 Crystallographic data for complex 6^a

Complex 6	
Formula	C ₁₁₅ H ₁₁₂ Mn ₁₂ N ₄ O ₃₉
Fw, g mol ⁻¹	2833.36
T/K	200(2)
Space group	P $\bar{1}$
a/Å	15.8858(4)
b/Å	17.4589(4)
c/Å	27.1441(7)
α (°)	96.7381(6)
β (°)	103.1034(5)
γ (°)	107.7821(6)
Volume/Å ³	6839.3(3)
Z, Z'	2
F(000)	2884
Density (calcd) mg m ⁻³	1.376
Absorption coefficient mm ⁻¹	1.140
Absorption correction	Semi-empirical from equivalents
Reflns, measured	61 548
Reflns, independent	31 323 [R(int) = 0.0303]
Data/restraints/parameters	31 323/87/1632
Goodness-of-fit on F ²	1.028
R indices [I > 2 σ (I)]	R ₁ = 0.0474, wR ₂ = 0.1312
R indices (all data)	R ₁ = 0.0675, wR ₂ = 0.1458

^a R₁ = $\frac{\sum ||F_0| - |F_c||}{\sum |F_0|}$; wR₂ = $[\frac{\sum w(F_0^2 - F_c^2)^2}{\sum w(F_0^2)^2}]^{1/2}$;

GOF = $\sqrt{\frac{\sum w\Delta^2}{m - n}}$.

Conclusions

Many factors in a synthetic procedure can influence the geometries of the resulting products. For example, when hmpH ligands were used, a more symmetric geometry of compound 1 was obtained. SMM type behavior was observed in this compound. While hepH was employed, two Mn₁₂ clusters (2 and 3) with S = 0 were obtained and none were found to be SMMs. Upon switching the backbone of the hmpH ligand to isopropyl (dmhmpH), the situation becomes more sophisticated. When MnCl₂ and tBuCO₂⁻ are used as the reactants in a MeCN rich solution, a Mn₇ cluster (4) was isolated which is not a SMM. When Mn(O₂CPh)₂ was used in a CH₂Cl₂ rich solution a more distorted rod like Mn₁₂ (5) was isolated which was shown to be a SMM. Attempts to explore other geometries of manganese clusters by utilizing MnCl₂ and PhCO₂⁻ as the reactants in a MeOH rich solution, a new type Mn₁₂ (6) was

obtained. According to the work herein, we conclude that this compound is a SMM. Obviously, a number of factors, including peripheral ligands and solvent systems play important roles in determining the geometry of the manganese cluster and so do their magnetic properties. Although without the ability to carefully control the parameters that affect the nature of the product, this study clearly shows that approaches are available to significantly tune the magnetic properties of the metal clusters.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors wish to thank Professor Hui-Lien Tsai in National Cheng Kung University of Taiwan for assistance with measuring the ac magnetic susceptibilities. This work was supported by the Ministry of Science and Technology of Taiwan (MOST 106-2113-M-030-008).

Notes and references

- M. Guo, T. Corona, K. Ray and W. Nam, *ACS Cent. Sci.*, 2019, 5(1), 13–28.
- G. C. Dismukes, R. Brimblecombe, G. A. N. Felton, R. S. Pryadun, J. E. Sheats, L. Spiccia and G. F. Swiegers, *Acc. Chem. Res.*, 2009, 42(12), 1935–1943.
- (a) G. E. Kostakis, A. M. Ako and A. K. Powell, *Chem. Soc. Rev.*, 2010, 39, 2238–2271; (b) L. K. Thompson, *Coord. Chem. Rev.*, 2002, 233–234, 193–206; (c) C.-M. Liu, R.-G. Xiong, D.-Q. Zhang and D.-B. Zhu, *J. Am. Chem. Soc.*, 2010, 132, 4044.
- (a) G. Christou, D. Gatteschi, D. N. Hendrickson and R. Sessoli, *MRS Bull.*, 2000, 25(11), 66–71; (b) D. Gatteschi and R. Sessoli, *Angew. Chem., Int. Ed.*, 2003, 42, 268; (c) G. Christou, *Polyhedron*, 2005, 24(16–17), 2065–2075; (d) G. Aromi and E. K. Brechin, *Struct. Bonding*, 2006, 122, 1; (e) A. K. Powell, *Nat. Chem.*, 2010, 2, 351.
- (a) H. J. Gerristen and E. S. Sabisky, *Phys. Rev.*, 1963, 132, 1507–1512; (b) C. Boskovic, W. Wernsdorfer, K. Folting, J. C. Hoffman, D. N. Hendrickson and G. Christou, *Inorg. Chem.*, 2002, 41, 5107; (c) C. J. Milios, A. Vinslava, W. Wernsdorfer, S. Moggach, S. Parsons, S. P. Perlepes,

- G. Christou and E. K. Brechin, *J. Am. Chem. Soc.*, 2007, **129**, 2754; (d) Y. Sunatsuki, Y. Kishima, T. Kobayashi, T. Yamaguchi, T. Suzuki, M. Kojima, J. Krzystek and M. R. Sundberg, *Chem. Commun.*, 2011, **47**(32), 9149–9151; (e) J. Krzystek, G. J. Yeagle, J.-H. Park, R. D. Britt, M. W. Meisel, L.-C. Brunel and J. Telsler, *Inorg. Chem.*, 2003, **42**(15), 4610–4618.
- 6 (a) R. Sessoli, D. Gatteschi, A. Caneschi and M. A. Novak, *Nature*, 1993, **365**, 141–143; (b) R. Sessoli, H.-L. Tsai, A. R. Schake, S. Y. Wang, J. B. Vincent, K. Folting, D. Gatteschi, G. Christou and D. N. Hendrickson, *J. Am. Chem. Soc.*, 1993, **115**, 1804–1816.
- 7 (a) J. Yoo, A. Yamaguchi, M. Nakano, J. Krzystek, W. E. Streib, L.-C. Brunel, H. Ishimoto, G. Christou and D. N. Hendrickson, *Inorg. Chem.*, 2001, **40**(18), 4604–4616; (b) M. A. Bolcar, S. M. J. Aubin, K. Folting, D. N. Hendrickson and G. Christou, *Chem. Commun.*, 1997, 1485–1486; (c) E. C. Sañudo, W. Wernsdorfer, K. A. Abboud and G. Christou, *Inorg. Chem.*, 2004, **43**(14), 4137–4144; (d) P. L. Feng, C. C. Beedle, C. Koo, W. Wernsdorfer, M. Nakano, S. Hill and D. N. Hendrickson, *Inorg. Chem.*, 2008, **47**(8), 3188–3204; (e) E.-C. Yang, W. Wernsdorfer, L. N. Zakharov, Y. Karaki, A. Yamaguchi, R. M. Isidro, G. D. Lu, S. A. Wilson, A. L. Rheingold, H. Ishimoto and D. N. Hendrickson, *Inorg. Chem.*, 2006, **45**(2), 529–546; (f) W. R. Yu, G. H. Lee and E.-C. Yang, *Dalton Trans.*, 2013, **42**, 3941.
- 8 (a) C. Boskovic, E. K. Brechin, W. E. Streib, K. Folting, J. C. Bollinger, D. N. Hendrickson and G. Christou, *J. Am. Chem. Soc.*, 2002, **124**, 3725–3736; (b) C. Boskovic, E. K. Brechin, W. E. Streib, K. Folting, D. N. Hendrickson and G. Christou, *Chem. Commun.*, 2001, 467.
- 9 T. Taguchi, W. Wernsdorfer, K. A. Abboud and G. Christou, *Inorg. Chem.*, 2010, **49**, 199–208.
- 10 I. D. Brown, *Chem. Rev.*, 2009, **109**, 6858–6919.
- 11 (a) M. Soler, W. Wernsdorfer, K. Folting, M. Pink and G. Christou, *J. Am. Chem. Soc.*, 2004, **126**, 2156; (b) P. King, W. Wernsdorfer, K. A. Abboud and G. Christou, *Inorg. Chem.*, 2005, **44**, 8659.
- 12 A. J. van Duynveldt, in *Magnetic Molecular Materials*, D. Gatteschi, O. Kahn, J. Miller, and F. Palacio, Kluwer Academic Publishers, London, 1991.
- 13 M. A. Novak, and R. Sessoli, in *Quantum Tunnelling of Magnetization, QTM '94*, ed. L. Gunther, and B. Barbar, Kluwer, Dordrecht, The Netherlands, 1995, pp. 171–188.
- 14 (a) S. M. Aubin, Z. Sun, L. Pardi, J. Krzystek, K. Folting, L. J. Brunel, A. L. Rheingold, G. Christou and D. N. Hendrickson, *Inorg. Chem.*, 1999, **38**, 5329–5340; (b) K. S. Cole and R. H. Cole, *J. Chem. Phys.*, 1941, **9**, 341.
- 15 W. Wernsdorfer, N. Allaga-Alcalde, D. N. Hendrickson and G. Christou, *Nature*, 2002, **416**(6879), 406–409.
- 16 L. Thomas, F. Lioni, R. Ballou, D. Gatteschi, R. Sessoli and B. Barbara, *Nature*, 1996, **383**, 145–147.
- 17 (a) W. Wernsdorfer, S. Bhaduri, C. Boskovic, G. Christou and D. N. Hendrickson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2002, **65**, 180403; (b) W. Wernsdorfer, N. E. Chakov and G. Christou, *Phys. Rev. Lett.*, 2005, **95**, 037203.
- 18 M. J. Ridd, D. J. Gakowski, G. E. Sneddon and F. R. Keene, *Dalton Trans.*, 1992, 1949.
- 19 Bruker APEX3, SAINT and SADABS, Bruker AXS Inc., Madison, Wisconsin, USA, 2016.
- 20 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015a, **71**, 3–8.
- 21 G. M. Sheldrick, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015b, **71**, 3–8.
- 22 (a) O. Kahn, *Molecular Magnetism*, VCH Publishers, Inc, 1993; (b) R. L. Carlin, *Magnetochemistry*, Springer-Verlag., 1986.
- 23 W. Wernsdorfer, *Adv. Chem. Phys.*, 2001, **118**, 99–190.