# Magnetic Properties of High-Nuclearity $\mathrm{Fe}_{x}$-oxo ( $x=7,22,24$ ) Clusters Analyzed by a Multipronged Experimental, Computational, and Magnetostructural Correlation Approach 

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#### Abstract

The synthesis, structure, and magnetic properties of three related iron(III)-oxo clusters are reported, $\left[\mathrm{Fe}_{7} \mathrm{O}_{3}\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{9}(\mathrm{mda})_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)\right]$ (1), $\left[\mathrm{Fe}_{22} \mathrm{O}_{14}(\mathrm{OH})_{3}\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{6}\right]\left(\mathrm{ClO}_{4}\right)_{2}$ (2), and $\left[\mathrm{Fe}_{24} \mathrm{O}_{15}(\mathrm{OH})_{4}(\mathrm{OEt})\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{7}\right]\left(\mathrm{ClO}_{4}\right)_{2}$ (3), where $\mathrm{mdH}_{2}$ is N -methyldiethanolamine. 1 was prepared from the reaction of $\left[\mathrm{Fe}_{3} \mathrm{O}\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{6}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]\left(\mathrm{NO}_{3}\right)$ with mdaH ${ }_{2}$ in a 1:2 ratio in MeCN , whereas 2 and 3 were prepared from the reaction of $\mathrm{FeCl}_{3} / \mathrm{NaO}_{2} \mathrm{CMe} / \mathrm{mdaH}_{2}$ in a 2:~13:2 ratio and $\mathrm{FeCl}_{3} /$ $\mathrm{NaO}_{2} \mathrm{CMe} / \mathrm{mdaH}_{2} /$ pyridine in a $2: \sim 13: 2: 25$ ratio, respectively, both in EtOH. The core of 1 consists of a central octahedral $\mathrm{Fe}^{\text {III }}$ ion held within a nonplanar $\mathrm{Fe}_{6}$ loop by three $\mu_{3}-\mathrm{O}^{2-}$ and three $\mu_{2}-$  $\mathrm{RO}^{-}$arms from the three $\mathrm{mda}^{2-}$ chelates. The cores of the cations of 2 and $\mathbf{3}$ consist of an A:B:A three-layer topology, in which a central $\mathrm{Fe}_{6}(\mathbf{2})$ or $\mathrm{Fe}_{8}$ (3) layer $\mathbf{B}$ is sandwiched between two $\mathrm{Fe}_{8}$ layers $\mathbf{A}$. The $\mathbf{A}$ layers structurally resemble $\mathbf{1}$ with the additional Fe added at the center to retain virtual $\mathrm{C}_{3}$ symmetry. The central $\mathrm{Fe}_{6}$ layer $\mathbf{B}$ of $\mathbf{2}$ consists of a $\left\{\mathrm{Fe}_{4}\left(\mu_{4}-\mathrm{O}\right)_{2}\left(\mu_{3}-\mathrm{OH}\right)_{2}\right\}^{6+}$ cubane with an Fe on either side attached to cubane $\mathrm{O}^{2-}$ ions, whereas that of 3 has the same cubane but with an $\left\{\mathrm{Fe}_{3}\left(\mu_{3}-\mathrm{O}\right)(\mu-\mathrm{OH})\right\}$ unit attached on one side and a single Fe on the other. Variable-temperature dc and ac magnetic susceptibility studies revealed dominant antiferromagnetic coupling in all complexes leading to ground-state spins of $S=5 / 2$ for 1 and $S=0$ for 2 and 3 . All $\mathrm{Fe}_{2}$ pairwise exchange parameters $\left(J_{i j}\right)$ for $\mathbf{1 - 3}$ were estimated by two independent methods: density functional theory (DFT) calculations using broken symmetry methods and a magnetostructural correlation previously developed for high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ complexes. The two approaches gave satisfyingly similar $J_{i j}$ values, and the latter allowed rationalization of the experimental ground states by identification of the spin frustration effects operative and the resultant relative spin vector alignments at each $\mathrm{Fe}^{\text {III }}$ ion.


## INTRODUCTION

The beauty and utility of $\mathrm{Fe}^{\mathrm{III}}$ /oxo chemistry in diverse fields such as bioinorganic chemistry, molecular magnetism, and materials science have led to the discovery of a remarkable slew of clusters, spanning nuclearities from $\mathrm{Fe}_{2}{ }^{1-4}$ up to giant hexameric $\left\{\mathrm{Fe}_{28}\right\}_{6}$ nanocages. ${ }^{5}$ Dinuclear $\mathrm{Fe}^{\text {III }}$ complexes have served as both model systems to understand magnetic exchange couplings via magnetostructural correlations (MSCs) ${ }^{6}$ as well as synthetic analogues of di-iron biomolecules such as ribonucleotide reductase, ${ }^{7,8}$ hemerythrin, ${ }^{9}$ and the soluble methane monooxygenase, ${ }^{8,10}$ among others. ${ }^{11}$ Highernuclearity $\mathrm{Fe}^{\mathrm{III}}$ /oxo clusters are likewise highly coveted as potential models of intermediates in the growth of nanoscale $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O} / \mathrm{OH}$ units during the loading of the ferritin protein, ${ }^{12}$ as well as new magnetic molecules and single-molecule magnets (SMMs). ${ }^{13-18}$

Large, discrete $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}^{2-} / \mathrm{OH}^{-}$clusters remain highly prized since nuclearities $>12$ are rare despite considerable efforts over many years. In fact, the family of large $\mathrm{Fe}{ }^{\mathrm{III}} / \mathrm{O}^{2-} / \mathrm{OH}^{-}$clusters
with $n \geq 14$ only numbers 20 members, of which only 4 have $n$ $>19 .{ }^{19-21}$ Thus, there is still a great need for new synthetic routes to discrete, high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}^{2-} / \mathrm{OH}^{-}$clusters. One fruitful strategy in our group and others has been the use of polyalcohol chelating/bridging ligands in reactions with various $\mathrm{Fe}^{\mathrm{III}}$ salts or small $\mathrm{Fe} /$ oxo complexes, both in the presence and absence of carboxylates. ${ }^{19,20,22-25}$ For example, with $N$-methyldiethanolamine $\left(\mathrm{mdaH}_{2}\right)$, this gave $\left[\mathrm{Fe}^{\mathrm{III}}{ }_{22} \mathrm{O}_{14}(\mathrm{OH})_{3}\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{6}\right]^{2+19}$ and $\left[\mathrm{Fe}_{7} \mathrm{O}_{3}\left(\mathrm{O}_{2} \mathrm{C}^{t} \mathrm{Bu}\right)_{9}(\mathrm{mda})_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]^{25}$ from the reaction with $\mathrm{Fe}^{\mathrm{III}}$ salts in the presence of $\mathrm{MeCO}_{2}^{-}$or with $\left[\mathrm{Fe}_{3} \mathrm{O}-\right.$

[^0]

Table 1. Crystal Data and Structure Refinement Parameters for 1, 2, and 3

| parameter | 1 | $2 \cdot x$ (solv) | 3•x(solv) |
| :---: | :---: | :---: | :---: |
| formula ${ }^{\text {a }}$ | $\mathrm{C}_{78} \mathrm{H}_{80} \mathrm{Fe}_{7} \mathrm{~N}_{3} \mathrm{O}_{28}$ | $\mathrm{C}_{96} \mathrm{H}_{202} \mathrm{Cl}_{2} \mathrm{Fe}_{22} \mathrm{~N}_{6} \mathrm{O}_{91}$ | $\mathrm{C}_{87} \mathrm{H}_{170} \mathrm{Cl}_{2} \mathrm{Fe}_{24} \mathrm{~N}_{7} \mathrm{O}_{86.5}$ |
| FW, $\mathrm{g} \mathrm{mol}^{-1}$ | 1898.40 | 4192.24 | 4109.59 |
| space group | Pa $\overline{3}$ | C2/c | P $\overline{1}$ |
| a, $\AA$ | 24.9810(5) | 29.719(3) | 13.8505(7) |
| b, $\AA$ | 24.9810(5) | 35.321(4) | 21.9456(12) |
| c, $\AA$ | 24.9810(5) | 30.651(3) | 25.7420(14) |
| $\alpha$, deg | 90 | 90 | 81.118(2) |
| $\beta$, deg | 90 | 98.367(2) | 76.501(2) |
| $\gamma$, deg | 90 | 90 | 80.534(2) |
| $V, \AA^{3}$ | 15589.4(5) | 31832(6) | 7449.6(7) |
| $\lambda, \AA^{b}$ | 0.71073 | 0.71073 | 0.71073 |
| Z | 8 | 8 | 2 |
| T, K | 173(2) | 173(2) | 100(2) |
| $\rho_{\text {calc, }} \mathrm{g} / \mathrm{cm}^{3}$ | 1.618 | 1.751 | 1.832 |
| $R_{1}{ }^{c, d}$ | 0.0463 | 0.0680 | 0.0424 |
| $\mathrm{w} R_{2}{ }^{\text {e }}$ | 0.1166 | 0.1581 | 0.0954 |
| $a_{x}$ (solv) is $4 \mathrm{H}_{2} \mathrm{O} \cdot 4 \mathrm{EtOH} \cdot 4 \mathrm{Et}_{2} \mathrm{O}$ for 2, and ${ }^{3} /{ }_{2} \mathrm{Et}_{2} \mathrm{O} \cdot \mathrm{EtOH}$ for 3 . ${ }^{b}$ Including solvent molecules. ${ }^{c} I>2 \sigma(I) .{ }^{d} R_{1}=\sum\left(\left\\|F_{\mathrm{o}}\|-\| F_{\mathrm{c}}\right\\|\right) / \sum\left\|F_{\mathrm{o}}\right\| .{ }^{e} \mathrm{w} R_{2}=$ $\left[\sum\left[\mathrm{w}\left(F_{\mathrm{o}}{ }^{2}-F_{\mathrm{c}}{ }^{2}\right)^{2}\right] / \sum\left[\mathrm{w}\left(F_{\mathrm{o}}{ }^{2}\right)^{2}\right]\right]^{1 / 2}, w=1 /\left[\sigma^{2}\left(F_{\mathrm{o}}{ }^{2}\right) \pm(m \times p)^{2}+n \times p\right], p=\left[\max \left(F_{\mathrm{o}}{ }^{2}, 0\right)+2 F_{\mathrm{c}}{ }^{2}\right] / 3 ; m$ and $n$ are constants. |  |  |  |

$\left.\left(\mathrm{O}_{2} \mathrm{C}^{t} \mathrm{Bu}\right)_{6}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]^{1+}$, respectively. For over 15 years, the former was the largest $\mathrm{Fe}^{\text {III }} / \mathrm{O} / \mathrm{OH}$ cluster in the literature until the report in 2019 of a $\mathrm{Fe}_{34}$ cluster. ${ }^{21}$
Since oxo-bridged high-spin $\mathrm{Fe}^{\mathrm{III}}(S=5 / 2)$ pairs almost always exhibit antiferromagnetic (AF) exchange couplings, high-nuclearity clusters are often replete with spin frustration effects if triangular $\mathrm{AF} \mathrm{Fe}_{3}$ subunits are present, as they usually are. Spin frustration here follows the definition preferred by molecular chemists as the occurrence of competing exchange interactions, $J_{i j}$, of comparable magnitude that hinder (i.e., frustrate) the preferred spin alignments. ${ }^{26}$ As a result, spin frustration often leads to uncompensated ground-state spin, and if the latter is large enough, the $\mathrm{Fe}_{n}$ cluster may even function as an SMM, although $\mathrm{Fe}^{\text {III }}$ clusters typically do not possess substantial magnetic anisotropy. ${ }^{27}$
In the past, exchange interactions, $J_{i j}$, in such high-nuclearity $\mathrm{Fe}^{\text {III }} / \mathrm{O}$ clusters have been difficult to determine with confidence from experimental or computational methods owing to (i) the multitude of inequivalent $J_{i j}$ frequently observed as the metal nuclearity increases, (ii) the lack of a reliable MSC for high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ clusters that could provide reliable estimates of $J_{i j}$ from the metric parameters, and (iii) the increasing computational demands of theoretical methods as nuclearity increases. It was for reason (ii) that our group reported the Mitchell-Christou semiempirical polynuclear $\mathrm{Fe}^{\text {III }} / \mathrm{O}$ MSC procedure in $2016,{ }^{28}$ which enables the calculation of reliable, realistic $J_{i j}$ values in high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ clusters. The provided protocol also critically accounts for multiple types of oxo-bridging pathways between $\mathrm{Fe}^{\mathrm{III}}{ }_{2}$ pairs, a common occurrence in the $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ architecture of increasingly larger complexes. ${ }^{28}$

The MSC approach coupled with the continually improving power of computational methods and experimental magnetic susceptibility measurements now provide a realistic opportunity to better understand the nature of the various exchange couplings within high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ clusters from a threepronged approach involving MSC, density functional theory (DFT), and experimental magnetochemistry methods. While application of three-pronged approaches is common in smallernuclearity systems up to $\mathrm{Fe}_{12}$ or so, ${ }^{26 c, 29,30}$ our objective more recently has been to tackle clusters of a higher nuclearity at
which a detailed study of the exchange couplings and the resulting origin of the magnetic properties is almost nonexistent. In the present work, we report a three-pronged approach to the study of three $\mathrm{Fe}_{x}$ clusters, the new $\left[\mathrm{Fe}_{7} \mathrm{O}_{3}\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{9}(\mathrm{mda})_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)\right]$ (1), previously reported $\left[\mathrm{Fe}_{22} \mathrm{O}_{14}(\mathrm{OH})_{3}\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{6}\right]-\left(\mathrm{ClO}_{4}\right)_{2} \quad(2)$ mentioned above, and the newly discovered $\left[\mathrm{Fe}_{24} \mathrm{O}_{15}(\mathrm{OH})_{4}(\mathrm{OEt})\right.$ $\left.\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{7}\right]\left(\mathrm{ClO}_{4}\right)_{2}(3) ; \mathrm{Fe}_{7}$ complex 1 is a structural subunit of the larger clusters. We herein report (a) the syntheses of 1 and 3, (b) the results from a detailed study of the exchange-coupling $\left(J_{i j}\right)$ parameters of $1-3$ by broken symmetry DFT calculations and use of the polynuclear MSC method, (c) the $J_{i j}$ parameters for 1 from fits of experimental $\chi_{\mathrm{M}} T$ versus $T$ data using the program $\mathrm{PHI}^{31}$ and (d) the analysis and rationalization of both the spin frustration effects operative and the resulting experimentally determined groundstate spins of $\mathbf{1 - 3}$.

## - EXPERIMENTAL SECTION

Syntheses. All manipulations were performed under aerobic conditions. Reagent grade solvents were used without further purification. Organic reagents were used as received, and $\left[\mathrm{Fe}_{3} \mathrm{O}-\right.$ $\left.\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{6}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]\left(\mathrm{NO}_{3}\right)$ was prepared as described elsewhere; ${ }^{32}$ mdaH ${ }_{2}=\mathrm{N}$-methyldiethanolamine.
$\left[\mathrm{Fe}_{7} \mathrm{O}_{3}\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{9}(\mathrm{mda})_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)\right](1)$. To a stirred orange solution of $\left[\mathrm{Fe}_{3} \mathrm{O}\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{6}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]\left(\mathrm{NO}_{3}\right)(0.50 \mathrm{~g}, 0.49 \mathrm{mmol})$ in $\mathrm{MeCN}(25$ $\mathrm{mL})$ was added $\mathrm{mdaH}_{2}(0.21 \mathrm{~g}, 1.8 \mathrm{mmol})$, causing the solution color to turn brown. The solution was stirred for a further 4 h and filtered, and the sealed filtrate was maintained undisturbed at ambient temperature. Brown plate crystals of 1 slowly grew over a few days and were either maintained in the mother liquor for single-crystal Xray crystallography or collected by filtration, washed with MeCN , and dried in vacuo for other studies. The yield was 0.095 g ( $24 \%$ based on Fe ). The vacuum-dried material was analyzed as $\mathbf{1} \cdot \mathrm{H}_{2} \mathrm{O}$. Anal. Calcd (found) for $\mathrm{C}_{78} \mathrm{H}_{82} \mathrm{Fe}_{7} \mathrm{~N}_{3} \mathrm{O}_{29}$ : C, 48.89 (48.64); $\mathrm{H}, 4.31$ (4.36); N , 2.19 (2.59). Selected IR data ( KBr disk, $\mathrm{cm}^{-1}$ ): 3418 (b, m), 2971 (m, sh), 2989 (m, sh), 2860 (m, sh), 1599 ( s, sh), 1557 ( $\mathrm{s}, \mathrm{sh}$ ), 1519 (m, sh), 1498 ( m, sh), 1447 (m sh), 1401 ( s$), 1262(\mathrm{w}, \mathrm{sh}), 1175$ (m, sh), 1104 (m), 1061 (m, sh), $1025(\mathrm{~m}, \mathrm{sh}), 999(\mathrm{~m}, \mathrm{sh}), 900(\mathrm{~m}, \mathrm{sh})$, 865 (m, sh), 833 ( w, sh), 760 ( w ), 719 ( s$), 686$ ( $\mathrm{s}, \mathrm{sh}$ ), 673 ( $\mathrm{s}, \mathrm{sh}$ ), 618 (m, sh), $580(\mathrm{~m}), 530(\mathrm{~m}), 466(\mathrm{~m})$.
$\left[\mathrm{Fe}_{22} \mathrm{O}_{14}(\mathrm{OH})_{3}\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{6}\right]\left(\mathrm{ClO}_{4}\right)_{2}$ (2). The procedure below is an improvement to the one reported previously. ${ }^{19}$ To a
stirred solution of $\mathrm{FeCl}_{3} \cdot 6 \mathrm{H}_{2} \mathrm{O}(0.30 \mathrm{~g}, 1.1 \mathrm{mmol})$ and $\mathrm{NaO}_{2} \mathrm{CMe}$ $(0.60 \mathrm{~g}, 7.3 \mathrm{mmol})$ in $\mathrm{EtOH}(10 \mathrm{~mL})$ was added $\mathrm{mdaH}_{2}(0.14 \mathrm{~g}, 1.1$ mmol ), giving rise to a rapid color change from bright orange to dark red-orange. To this solution, $\mathrm{NaClO}_{4}(0.07 \mathrm{~g}, 0.57 \mathrm{mmol})$ was added, and then the solution was stirred for 15 min . A colorless precipitate was removed by filtration and discarded. Dark red-orange crystals grew over 4 weeks after layering the filtrate with $\mathrm{Et}_{2} \mathrm{O}$, and these were collected by filtration, washed with $\mathrm{Et}_{2} \mathrm{O}$, and dried in vacuo. The vacuum-dried material was analyzed as $2 \cdot 4 \mathrm{H}_{2} \mathrm{O}$. Anal. Calcd (found) for $\mathrm{C}_{72} \mathrm{H}_{140} \mathrm{Cl}_{2} \mathrm{Fe}_{22} \mathrm{~N}_{6} \mathrm{O}_{83}$ : C, 23.26 (23.55); H, 3.80 (3.75); N, 2.26 (2.00). The yield was $0.037 \mathrm{~g}(20 \%$ based on Fe$)$. Crystals suitable for X-ray crystallography were grown in small vials and maintained in mother liquor until mounted for data collection. Selected IR data ( KBr disk, $\mathrm{cm}^{-1}$ ): 3420 (b, m), 2972 (m, sh), 2926 (m, sh), 2867 (m, sh), 2361 ( w , sh), 2337 ( w , sh), 1576 (v.st, sh), 1540 (v.st, sh), 1436 (v.st, sh), 1351 (w, sh), 1277 (w, sh), 1248 (w, sh), 1096 (st, sh), 1025 (m, sh), 999 (m, sh), 903 (m, sh), 876 (m, sh), 761 (w, sh), 708 ( $\mathrm{w}, \mathrm{sh}$ ), 654 (m, sh), 623 (m, sh), 583 ( st, sh), 538 ( st, sh), 428 (m).
$\left[\mathrm{Fe}_{24} \mathrm{O}_{15}(\mathrm{OH})_{4}(\mathrm{OEt})\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{7}\right]\left(\mathrm{ClO}_{4}\right)_{2}$ (3). To a stirred solution of $\mathrm{FeCl}_{3} \cdot 6 \mathrm{H}_{2} \mathrm{O}(0.30 \mathrm{~g}, 1.1 \mathrm{mmol})$ and $\mathrm{NaO}_{2} \mathrm{CMe}(0.60 \mathrm{~g}$, $7.3 \mathrm{mmol})$ in $\mathrm{EtOH}(10 \mathrm{~mL})$ was added $\mathrm{mdaH}_{2}(0.14 \mathrm{~g}, 1.1 \mathrm{mmol})$ giving rise to a rapid color change from bright orange to dark redorange. Pyridine ( $1.0 \mathrm{~mL}, 12.4 \mathrm{mmol}$ ) was added followed by $\mathrm{NaClO}_{4}$ $(0.07 \mathrm{~g}, 0.57 \mathrm{mmol})$, and then the solution was stirred for 15 min . The colorless precipitate was removed by filtration and discarded. Dark red-orange crystals grew over 6 weeks after layering the filtrate with $\mathrm{Et}_{2} \mathrm{O}$, and these were collected by filtration, washed with EtOH , and dried in vacuo. The vacuum-dried material was analyzed as 3.5EtOH. Anal. Calcd (found) for $\mathrm{C}_{89} \mathrm{H}_{179} \mathrm{Cl}_{2} \mathrm{Fe}_{24} \mathrm{~N}_{7} \mathrm{O}_{89}$ : C, 25.56 (26.17); H, 4.31 (3.84); N, 2.34 (2.66). The yield was $0.038 \mathrm{~g}(20 \%$ based on Fe ). Crystals suitable for X-ray crystallography were grown in small vials and maintained in mother liquor until mounted for data collection. Selected IR data ( KBr disk, $\mathrm{cm}^{-1}$ ): 3394 (b, m), 2972 (m, sh), 2926 (m, sh), 2867 (m, sh), 2677 ( $\mathrm{w}, \mathrm{sh}$ ), 1552 ( v.st, sh), 1428 (v.st, sh), 1370 (w, sh), 1288 (w, sh), 1241 (w, sh), 1094 (st, sh), 1024 (m, sh), 999 (m, sh), 902 (m, sh), 876 (m, sh), 760 (w, sh), 712 ( $\mathrm{w}, \mathrm{sh}$ ), 655 (m, sh), 634 (m, sh), 583 (st, sh), 534 (st, sh), 431 (m).

X-ray Crystallography. The structures were refined using fullmatrix least-squares cycles. The non-H atoms were refined with anisotropic displacement parameters, and all H atoms were placed in calculated, idealized positions and refined riding on their parent atoms. The refinements were carried out by minimizing the $\mathrm{w} R_{2}$ function using $F^{2}$ rather than $F$ values. $R_{1}$ was calculated to provide a reference to the conventional $R$ value, but its function was not minimized. Crystal data and refinement parameters are collected in Table 1.

For 1 and $2 \cdot 4 \mathrm{H}_{2} \mathrm{O} \cdot 4 \mathrm{EtOH} \cdot 4 \mathrm{Et}_{2} \mathrm{O}$, data were collected at 173 K on a Siemens SMART PLATFORM equipped with a CCD area detector and a graphite monochromator utilizing $\mathrm{Mo} \mathrm{K} \alpha$ radiation ( $\lambda=$ $0.71073 \AA$ ). For 1, the cell parameters were refined using 8192 reflections. A full sphere of data (1850 frames) was collected using the $\omega$-scan method ( $0.3^{\circ}$ frame width). The first 50 frames were remeasured at the end of data collection to monitor instrument and crystal stability (maximum correction on $I$ was $<1 \%$ ). Absorption corrections by integration were applied based on measured indexed crystal faces. The structure was solved by direct methods in SHELXTL18. ${ }^{33}$ The asymmetric unit consists of a $1 / 3 \mathrm{Fe}_{7}$ cluster with two disorders: (i) The C13 benzoate Ph ring displays a resolved twofold rotational disorder with $58: 42 \%$ occupancies and atom C14 common to both rings; (ii) The second disorder involves the C21 benzoate ligand being a bidentate chelate on Fe 3 in one instance and a monodentate ligand in the other with a $\mathrm{H}_{2} \mathrm{O}$ (O8) ligand at the second coordination site. The occupancies were $2: 1$ corresponding to the overall formula $\left[\mathrm{Fe}_{7} \mathrm{O}_{3}\left(\mathrm{O}_{2} \mathrm{CPh}\right)_{9}(\mathrm{mda})_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)\right]$. To check that this disorder is not introduced by choosing the higher symmetry space group $P a \overline{3}$, the structure was refined in the non-centrosymmetric space group as well as the orthorhombic space group Pbca. In both cases, the disorders were observed, thus proving that $P a \overline{3}$ is the correct space group. A total of 366 parameters were refined in the
final cycle of refinement using 5590 reflections with $I>2 \sigma(I)$ to yield $R_{1}$ and $\mathrm{w} R_{2}$ of 4.63 and $11.66 \%$, respectively.

For $2 \cdot 4 \mathrm{H}_{2} \mathrm{O} \cdot 4 \mathrm{EtOH} \cdot 4 \mathrm{Et}_{2} \mathrm{O}$, a preliminary search of reciprocal space revealed a set of reflections with a monoclinic lattice. Initial choice of space group $C 2 / c$ was subsequently confirmed by the successful solution of the structure. The asymmetric unit consists of the $\mathrm{Fe}_{22}$ cluster, two $\mathrm{ClO}_{4}{ }^{-}$anions, and four each of $\mathrm{H}_{2} \mathrm{O}, \mathrm{EtOH}$, and $\mathrm{Et}_{2} \mathrm{O}$ solvent molecules. Charge balance considerations require a +2 charge on the cluster to counter the two $\mathrm{ClO}_{4}^{-}$anions in the asymmetric unit, and after bond valence sum calculations on both metals and ligands and close examination of the bond lengths, it was concluded that there is a proton situated on O28. This proton could not be located in a difference Fourier map however, and it was not included in the final refinement. A total of 1618 parameters were refined in the final cycle of refinement using 7571 reflections with $I>2 \sigma(I)$ to yield $R_{1}$ and $\mathrm{w} R_{2}$ of 6.80 and $15.81 \%$, respectively.

For $3 \cdot{ }^{3} /{ }_{2} \mathrm{Et}_{2} \mathrm{O} \cdot \mathrm{EtOH}$, data were collected at 100 K on a Bruker Dual micro source D8 Venture diffractometer and a PHOTON III detector running the APEX3 software package of programs and using Mo $\mathrm{K} \alpha$ radiation $(\lambda=0.71073 \AA)$. The data frames were integrated, and multiscan scaling was applied in APEX3. Intrinsic phasing structure solution provided all non-H atoms. ${ }^{22}$ The asymmetric unit consists of the $\mathrm{Fe}_{24}$ cluster, two $\mathrm{ClO}_{4}^{-}$counterions, two $\mathrm{Et}_{2} \mathrm{O}$ solvent molecules, one of which is disordered and refined in three parts, and an EtOH molecule disordered over two positions. In the final cycle of refinement, 53,101 reflections [of which 42,129 are observed with $I>$ $2 \sigma(I)$ ] were used to refine 1952 parameters, and the resulting $R_{1}, \mathrm{w} R_{2}$, and $S$ (goodness of fit) were $4.24 \%, 9.54 \%$, and 1.040 , respectively.

DFT Calculations. DFT calculations were performed for $\mathbf{1}$ and the cations of 2 and 3 using single-crystal X-ray crystallographic coordinates. We considered a total of 12,62 , and 68 distinct $J_{i j}$ first-neighbor exchange couplings for 1 and the cations of 2 and 3, respectively. These couplings were determined from DFT calculations by mapping broken-symmetry solutions to Ising-type spin configurations $\{S\}$. These configurations were chosen as one high-spin, all single-spin inversions, and all first-neighbor two-spin inversions, giving a total of 20,85 , and 93 broken-symmetry solutions for 1 and the cations of 2 and 3, respectively. The energies of these configurations are expressed in terms of a sum over spin interactions (eq 1)

$$
\begin{equation*}
E(\{S\})=E_{0}-2 \sum_{\langle i j\rangle} J_{i j} S_{i} \cdot S_{j} \tag{1}
\end{equation*}
$$

where $\langle i j\rangle$ stands for all neighboring $i j$ pairs, $S_{k}= \pm{ }^{5} / 2$ for $\mathrm{Fe}^{\text {III }}$, and $E_{0}$ is a constant introduced to match the spin model with the DFT energies.

To determine the exchange couplings, $J_{i j}$, the energies of all configurations $\{S\}$ resulting from broken spin-symmetry DFT calculations were used to perform a linear fit of the Ising-type energy expression in eq 1 . This approach has been successfully used in the literature to determine exchange couplings in multicenter transition metal complexes. ${ }^{34-36}$ The $R^{2}$ in our case for the linear regression differs from 1 by less than $10^{-6}$, which indicates that the magnetization is well localized at the magnetic centers, and thus the broken-symmetry DFT solutions are reliable representations of the Ising-type model configurations. The atomic spin populations from the DFT calculations are consistent with the expected broken spin symmetry configuration for all cases.

All DFT calculations were performed using the hybrid Perdew-Burke-Ernzerhof (PBEh) density functional approximation, which admixes $25 \%$ of exact (Hartree-Fock-type) exchange and $75 \%$ of PBE exchange. This choice of functional is known to perform well for magnetic exchange couplings, ${ }^{37}$ yielding for the particular case of oxobridged $\mathrm{Fe} \cdots \mathrm{Fe}$ couplings an RMS error of approximately $10 \%$, as shown for a set of 11 dinuclear $\mathrm{Fe}^{\mathrm{III}}$ complexes. ${ }^{38}$ Pople's all-electron $6-311+\mathrm{G}^{* *}$ basis was used for Fe atoms and the $6-31 \mathrm{G}^{* *}$ for lighter elements ${ }^{39}$ with relativistic effects neglected in all calculations. All broken-symmetry DFT energies were obtained using an in-house version of the Gaussian 16 program ${ }^{40}$ that allows for a simple spin


Figure 1. (a) Structure of 1 with all but the ipso-C atoms of the Ph rings removed for clarity; (b) labelled core and a side view along the $\mathrm{Fe} 2-\mathrm{Fe} 1 \mathrm{~B}$ vector emphasizing the nonplanarity of the $\mathrm{Fe}_{7}$ unit; and (c) a stereopair of the complete molecule. H atoms are omitted for clarity. Color code: $\mathrm{Fe}^{\mathrm{III}}$ lime green, O red, N powder blue, C gray.
inversion of the magnetic centers to produce a suitable initial guess for self-consistent calculations. No symmetry was assumed at any point in the model or the DFT calculations. A self-consistency convergence threshold of $10^{-6} \mathrm{Ha}=0.2 \mathrm{~cm}^{-1}$ in the energy and $10^{-8}$ in the RMS changes in the density matrix was used in all calculations.

Other Studies. Infrared spectra were collected in the solid state ( KBr pellets) in the $400-4000 \mathrm{~cm}^{-1}$ range using a Nicolet Nexus 670 FTIR spectrometer. Direct current (dc) and alternating current (ac) magnetic susceptibility measurements were performed on vacuumdried polycrystalline solids, embedded in eicosane to prevent torquing, with a Quantum Design MPMS-XL SQUID magnetometer capable of operating with applied dc fields up to 7 T in the $1.8-400 \mathrm{~K}$ range. Magnetization vs. field and temperature data were fit using the program MAGNET, ${ }^{41}$ and $D$ versus $g$ fit error surfaces were generated using the program GRID. ${ }^{42}$ Pascal's constants were used to estimate the diamagnetic correction, ${ }^{43}$ and contributions from the eicosane and gel capsule were measured as a blank. These values were subtracted
from the experimental susceptibility to give the molar paramagnetic susceptibility. Dc studies were carried out in the $5.0-300.0 \mathrm{~K}$ range in a $0.1 \mathrm{~T}(1000 \mathrm{Oe})$ magnetic field and ac studies in the $1.8-15.0 \mathrm{~K}$ range in a 3.5 G ac field at an oscillation frequency of 1000 Hz . Elemental analyses (C, H, N) were performed on vacuum-dried samples at Atlantic Microlab, Inc.

## RESULTS AND DISCUSSION

Syntheses. Our group's original investigation of the use of $\mathrm{mdaH}_{2}$ in $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ carboxylate chemistry had led to the discovery of $\mathbf{2}^{19}$ and $\left[\mathrm{Fe}_{7} \mathrm{O}_{3}\left(\mathrm{O}_{2} \mathrm{C}^{t} \mathrm{Bu}\right)_{9}(\mathrm{mda})_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right](4)^{25}$ from somewhat similar procedures involving the reaction of $\mathrm{mdaH}_{2}$ with $\left[\mathrm{Fe}_{3} \mathrm{O}\left(\mathrm{O}_{2} \mathrm{CR}\right)_{3}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]^{+}$salts either preformed or generated in situ from $\mathrm{FeCl}_{3} / \mathrm{NaO}_{2} \mathrm{CR}$ mixtures. In the present work, we have explored this reaction scheme in more detail,
concentrating initially on the carboxylate and solvent employed because (i) 2 and 4 are obtained with $R=M e$ and ${ }^{t} B u$, respectively, of significantly different bulkiness, (ii) 2 and 4 are obtained from EtOH and MeCN reaction solvents, respectively, and (iii) the $\mathrm{Fe}_{7}$ unit of 4 is similar to the $\mathrm{Fe}_{8}$ units at each end of 2 , differing in the addition of another $\mathrm{Fe}^{\text {III }}$ (vide infra) -we thus wondered whether the large bulk of ${ }^{t} \mathrm{BuCO}_{2}{ }^{-}$ groups was trapping $\mathrm{Fe}_{7}$ products by shutting down further aggregation to $\mathrm{Fe}_{22}$. We thus employed the $\mathrm{R}=\mathrm{Ph}$ complex in MeCN in the present work and this again gave an $\mathrm{Fe}_{7}$ product, complex 1, with an identical core structure with 4 . When the R $=\mathrm{PhCO}_{2}^{-}$and ${ }^{t} \mathrm{BuCO}_{2}^{-}$reactions were instead performed in EtOH , the products were again the $\mathrm{Fe}_{7}$ products 1 and 4, respectively, as confirmed by single-crystal unit cell measurements. In contrast, the $\mathrm{R}=\mathrm{Et}$ reaction in EtOH gave a product that was concluded to be the propionate $\mathrm{Fe}_{22}$ cluster from IR spectral comparison with 2 . Thus, it seems that the carboxylate bulk is indeed the determining factor in giving an $\mathrm{Fe}_{7}$ or $\mathrm{Fe}_{22}$ product. This conclusion is rationalized by the space-filling view of the $\mathrm{Fe}_{22}$ cation (Figure S 1 ), which shows closely spaced acetate groups in the center of the molecule that could not accommodate larger $\mathrm{PhCO}_{2}{ }^{-}$and ${ }^{t} \mathrm{BuCO}_{2}{ }^{-}$groups, thus rationalizing formation instead of 1 and 4.
The synthesis of 2 involves an excess of $\mathrm{mdaH}_{2}$, and we suspected that it also functions as a base. We thus repeated the reaction with less $\mathrm{mdaH}_{2}$ and/or with added pyridine to see what effect these would have. Small changes in the $\mathrm{mdaH}_{2}$ amount did not change the product, but the presence of an excess of pyridine in the procedure to 2 gave instead $\mathrm{Fe}_{24}$ complex 3. The assembly of the cation is summarized in eq 2 . Other bases such as $\mathrm{NEt}_{3}$ also gave 3 but in lower yields.

$$
\begin{align*}
24 \mathrm{Fe}^{3+}+ & 21 \mathrm{MeCO}_{2}^{-}+7 \mathrm{mdaH}_{2}+49 \mathrm{py}+19 \mathrm{H}_{2} \mathrm{O} \\
+\mathrm{EtOH} & \rightarrow 49 \mathrm{pyH}^{+} \\
\quad+ & {\left[\mathrm{Fe}_{24} \mathrm{O}_{15}(\mathrm{OH})_{4}(\mathrm{OEt})\left(\mathrm{O}_{2} \mathrm{CMe}\right)_{21}(\mathrm{mda})_{7}\right]^{2+} } \tag{2}
\end{align*}
$$

The reaction time and temperature were also explored. However, when the procedures affording 2 and 3 were run at room temperature for 48 h , at $60^{\circ} \mathrm{C}$ for 4 h , and up to $120^{\circ} \mathrm{C}$ in a microwave reactor for 1 h , all gave the same products rather than higher-nuclearity species. Similarly, use of phenyldiethanolamine $\left(\mathrm{pdaH}_{2}\right)$, which was reported to react with $\left[\mathrm{Fe}_{3} \mathrm{O}\left(\mathrm{O}_{2} \mathrm{C}^{t} \mathrm{Bu}\right)_{6}\left(\mathrm{H}_{2} \mathrm{O}\right)_{3}\right]\left(\mathrm{NO}_{3}\right)$ in MeCN to yield an analogous $\mathrm{Fe}_{7}$ cluster to $4,{ }^{44}$ did not lead to a characterizable product. We note, however, that Baca et al. recently reported a pair of $\mathrm{Fe}_{22}$ clusters using ${ }^{i} \mathrm{PrCO}_{2} \mathrm{H}$ and either $\mathrm{mdaH}_{2}$ or the ${ }^{n} \mathrm{Bu}$ analogue bdaH${ }_{2}$; these $\mathrm{Fe}_{22}$ products are similar to 2 but nevertheless significantly different in that they do not have a central $\left\{\mathrm{Fe}_{4} \mathrm{O}_{4}\right\}$ cubane. ${ }^{20}$

Description of Structures. The complete structure, labeled core, and stereopair of $\mathbf{1}$ are shown in Figure 1. Selected interatomic distances and angles are listed in Table S1. 1 crystallizes in the trigonal space group $\mathrm{Pa} \overline{3}$ with the cluster on a $C_{3}$ symmetry axis. The structure consists of a central $\mathrm{Fe}^{\text {III }}(\mathrm{Fe} 2)$ held within a nonplanar $\mathrm{Fe}_{6}$ loop ( $\mathrm{Fe} 1, \mathrm{Fe} 3$, and their symmetry partners) by three $\mu_{3}-\mathrm{O}^{2-}$ (O7) ions and three $\mu_{2}$-alkoxide (O2) arms from the three mda ${ }^{2-}$ ligands. All metals are $\mathrm{Fe}^{\text {III }}$ with near-octahedral geometry, and the deprotonated nature of the bridging $\mathrm{O}^{2-}$ and $\mathrm{RO}^{-}$groups was confirmed by O bond valence sums (Table S2). ${ }^{45}$ The Fe 1 Fe 2 pairs are bridged by both $\mathrm{O}^{2-}$ and $\mathrm{RO}^{-}$ions, whereas
the Fe 3 Fe 2 pairs are only bridged by $\mathrm{O}^{2-}$, and this will have important consequences for the magnetic properties (vide infra). Peripheral ligation is completed by (i) a chelating mda ${ }^{2-}$ on each Fe 1 , with one alkoxide arm bridging to the central Fe 2 (as mentioned above) and the other bridging to an adjacent Fe3, (ii) a pair of syn,syn $\eta^{1}: \eta^{1}: \mu-\mathrm{PhCO}_{2}{ }^{-}$ligands bridging each of three symmetry-related Fe 1 Fe 3 pairs, and (iii) a chelating $\mathrm{PhCO}_{2}{ }^{-}$ligand on two Fe 3 ions, whereas the third Fe 3 has a monodentate $\mathrm{PhCO}_{2}^{-}$hydrogen-bonded to a terminal $\mathrm{H}_{2} \mathrm{O}$ (O8) ligand ( $\mathrm{O} 8 \cdots \mathrm{O}^{\prime} 0^{\prime}=2.539(5) \AA$ ). The resulting disorder about the $C_{3}$ axis is described in the Experimental Section.

Complex 2 has been reported elsewhere ${ }^{19}$ and is included here for comparison with 3 (vide infra) because their cations are structurally very similar. The complete structure of the cation of 3 as a stereopair and its labeled core are shown in Figure 2, and the selected metric parameters are listed in Table S3. 3 crystallizes in the triclinic space group $P \overline{1}$ with the cation in a general position. Its core consists of three $\mathrm{Fe}_{8}$ layers with


Figure 2. (a) Labelled core of the $\mathrm{Fe}_{24}$ cation of 3 and (b) stereopair of the complete cation. H atoms are omitted for clarity. Color code: octahedral $\mathrm{Fe}^{\text {III }}$ lime green, square-pyramidal $\mathrm{Fe}^{\mathrm{III}}$ lavender, O red, N powder blue, C gray.
an A:B:A pattern: the central $\mathrm{Fe}_{8}$ unit $\mathbf{B}$ is strikingly asymmetric and comprises an $\left\{\mathrm{Fe}_{4}\left(\mu_{3}-\mathrm{OH}\right)_{2}\left(\mu_{4}-\mathrm{O}\right)_{2}\right\}^{6+}$ cubane $\left(\mathrm{OH}^{-}\right.$ions are O 32 and O 34$)$ with a triangular $\left\{\mathrm{Fe}_{3}\left(\mu_{3}-\mathrm{O}\right)\right\}$ unit ( $\mathrm{Fe} 9, \mathrm{Fe} 23, \mathrm{Fe} 24$ ) attached on one side via two $\mu_{2}-\mathrm{OH}^{-}$ (O37, O39) ions and cubane O29, and a single Fe ( Fe 14 ) on the other side attached via $\mu_{2}-\mathrm{OEt}^{-} \mathrm{O} 42$ and cubane O35 (Figures 2 and S 2 ); in the $\mathrm{Fe}_{22}$ cation of 2, the central layer is a more symmetric $\mathrm{Fe}_{6}$ unit with a single Fe on either side of the cubane (vide infra). $\mathrm{Fe}^{\mathrm{III}}$ oxidation states and core O protonation levels were confirmed by BVS calculations ${ }^{45}$ (Tables S4 and S5, respectively). Twenty-two $\mathrm{Fe}^{\mathrm{III}}$ are octahedral, while the other two ( Fe 9 and Fe 14 ) are squarepyramidal; the latter are at the analogous positions as those in 2. ${ }^{19}$ The two $\mathrm{Fe}_{8}$ units $\mathbf{A}$ at each end of the cation of $\mathbf{3}$ are nearly isostructural with a three-blade propeller topology that can be described as essentially the $\mathrm{Fe}_{7}$ structure of $\mathbf{1}$, with an additional Fe atom on the virtual $\mathrm{C}_{3}$ axis to give the axle ( Fe 2 Fe 7 and Fe 11 Fe 21 ) of each propeller (Figure S2). One $\mathrm{Fe}_{8}$ unit $\mathbf{A}$ is connected to the central one by three $\mu_{3}-\mathrm{O}^{2-}$ ions ( $\mathrm{O} 16 / \mathrm{O} 20 / \mathrm{O} 24$ ), an alkoxide arm ( O 21 ) of the unique $\mu_{3}$ -$\mathrm{mda}^{2-}$ chelate that has no counterpart at the other end of the molecule, and two $\mu-\mathrm{MeCO}_{2}{ }^{-}$groups. The other $\mathrm{Fe}_{8}$ unit $\mathbf{A}$ is thus connected to the central one by three $\mu_{3}-\mathrm{O}^{2-}$ ions (O38/ $\mathrm{O} 41 / \mathrm{O} 44)$ and three $\mu$ - $\mathrm{MeCO}_{2}^{-}$groups. The peripheral ligation about the cation is completed by the remainder of the $\mu$ - $\mathrm{MeCO}_{2}{ }^{-}$groups and three $\mu_{3}$ - $\mathrm{mda}^{2-}$ chelates on each $\mathrm{Fe}_{8}$ unit A . Interestingly, all $\mathrm{MeCO}_{2}{ }^{-}$groups except one bridge in their common syn,syn $\eta^{1}: \eta^{1}: \mu$ mode, the exception being the one bridging Fe 14 Fe 23 in the rare syn, anti $\eta^{1}: \eta^{1}: \mu$ mode.

Comparison of the $\mathrm{Fe}_{22}$ and $\mathrm{Fe}_{24}$ Cations. The cations of 2 and 3 are compared in Figure 3 and in the space-filling


Figure 3. Comparison of the $\mathrm{Fe}_{22}$ and $\mathrm{Fe}_{24}$ cations of 2 (left) and 3 (right), respectively, using essentially the same viewpoint for 3 as Figure 2. Color code: octahedral $\mathrm{Fe}^{\mathrm{III}}$ lime green, new octahedral $\mathrm{Fe}^{\mathrm{III}}$ in 3 dark green, square-pyramidal $\mathrm{Fe}^{\mathrm{III}}$ lavender, O red, $\mathrm{OH}^{-}$sky blue, N and C in $\mathrm{mda}^{2-}$ ligands powder blue, C in $\mathrm{MeCO}_{2}{ }^{-}$gray, and C in $\mathrm{EtO}^{-}$purple.
mode in Figure S3. They have dimensions of $\sim 1.6 \times \sim 0.7 \times$ $\sim 0.8 \mathrm{~nm}$, excluding C and N atoms. Both possess an $\mathrm{A}: \mathrm{B}: \mathrm{A}$ three-layer topology and are overall fairly similar, with the main differences being in the differing nuclearity of the $\mathbf{B}$ layer, which are $\mathrm{Fe}_{6}$ and $\mathrm{Fe}_{8}$, respectively, and the presence of a seventh $\mathrm{mda}^{2-}$ chelate in 3 . The two additional $\mathrm{Fe}^{\text {III }}$ in 3 ( $\mathrm{Fe} 23 / \mathrm{Fe} 24$ ) are shown in darker green in Figure 3 and are
located in the $\left\{\mathrm{Fe}_{3}\left(\mu_{3}-\mathrm{O}\right)\right\}$ unit attached on one side of the central cubane, as mentioned earlier, with the seventh mda ${ }^{2-}$ chelating on Fe 24 . As a result, the cation of 3 is distinctly more asymmetric than that of 2 , which has a near- $C_{2}$ virtual symmetry, and a root-mean-square deviation (RMSD) analysis gave a weighted RMSD of $0.602 \AA$ (Table S6 and Figure S4) arising from differences in the relative attachment of the $\mathbf{A}$ layers to the central B layer in 3 as well as the structural changes in the central B layer caused by incorporation of Fe 23 and Fe 24 . Referring to the viewpoint of Figure 3, the top $\mathrm{Fe}_{8}$ layer $\mathbf{A}$ in $\mathbf{2}$ is near-isostructural with the analogous top layer in 3, except for a few minor changes to the ligation sphere (Figure 4). However, the bottom $\mathrm{Fe}_{8}$ layer in $\mathbf{2}$ is the mirror image of the one in 3 (Figure S5), that is, the orientations of the 'propeller blades' are opposite. In addition, the top and bottom $\mathrm{Fe}_{8}$ layers A are closer to being parallel in 2 than in 3: for example, the two mean planes through the outermost $\mathrm{Fe}_{3}$ triangles in each $\mathrm{Fe}_{8}$ give dihedral angles of 7.21 and $19.44^{\circ}$ for 2 and 3, respectively (Figure S6). This can be assigned to the structural distortions resulting from addition of the seventh $\mathrm{mda}^{2-}$ and the extra $\mathrm{Fe} 23 / \mathrm{Fe} 24$ ions and their attendant $\mathrm{O}^{2-} /$ $\mathrm{HO}^{-}$ions onto one side of the cation of 3.

Notwithstanding the differences in the overall structure, the $\mathrm{Fe}_{8}$ layers A of the cations of $\mathbf{2}$ and $\mathbf{3}$ are very similar to each other and to the $\mathrm{Fe}_{7}$ unit in 1 , except that the former two have an extra Fe on the virtual $\mathrm{C}_{3}$ axis, as mentioned earlier, and some changes to the ligation resulting from attachment to the central layer B. The comparison in Figure 4 emphasizes this overall similarity, including in the positions and binding modes of the three $\mathrm{mda}^{2-}$ groups. Finally, we note the differences between the cations of 2 and 3 and the $\mathrm{Fe}_{22}$ clusters with ${ }^{i} \mathrm{PrCO}_{2}{ }^{-}$by Baca et al.; the latter do not have a cubane in the central layer but instead an $\left\{\mathrm{Fe}_{4}\left(\mu_{3}-\mathrm{O}^{2-}\right)_{2}\right\}$ rhombus. ${ }^{20}$

Magnetic Susceptibility Studies. Variable-temperature dc magnetic susceptibility $\left(\chi_{\mathrm{M}}\right)$ data were collected on crushed vacuum-dried microcrystalline samples in the $5.0-300 \mathrm{~K}$ range in a $0.1 \mathrm{~T}(1 \mathrm{kG})$ applied dc field and plotted as $\chi_{\mathrm{M}} T$ versus $T$. For $1 \cdot \mathrm{H}_{2} \mathrm{O}, \chi_{\mathrm{M}} T$ decreases steadily from $6.8 \mathrm{~cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}$ at 300 K to a nearly constant $4.4 \mathrm{~cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}$ at 5.0 K (Figure 5a). The 300 K value is much lower than the value for seven noninteracting high-spin $\mathrm{Fe}^{\text {III }}$ ions with $g=2\left(30.6 \mathrm{~cm}^{3} \mathrm{~K}\right.$ $\mathrm{mol}^{-1}$ ), indicating dominant AF interactions. For 35 EtOH , $\chi_{\mathrm{M}} T$ decreases steadily from $29.6 \mathrm{~cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}$ at 300 K to 3.6 $\mathrm{cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}$ at 5.0 K (Figure 6). The 300 K value is much lower than the $105.0 \mathrm{~cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}(g=2)$ value for noninteracting $\mathrm{Fe}^{\mathrm{III}}$ ions, again indicating dominant AF interactions. This is analogous to the $\chi_{\mathrm{M}} T$ versus $T$ data for 2 (included for comparison in Figure 6), which also exhibits dominant AF interactions. ${ }^{19}$

The near-plateau value for $\mathbf{1} \cdot \mathrm{H}_{2} \mathrm{O}$ at 5.0 K suggests an $S=$ $5 / 2$ ground state, and this was supported by ac in-phase ( $\chi_{\mathrm{M}}{ }^{\prime}$ ) magnetic susceptibility data collected in the 1.8 to 15 K range in a 3.5 G ac field with a 1000 Hz oscillation frequency and plotted as $\chi_{\mathrm{M}}{ }^{\prime} T$ versus $T$ in Figure 5 b. The latter exhibited an essentially constant $\chi_{\mathrm{M}}{ }^{\prime} T \approx 4.5 \mathrm{~cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}$ from 15 K down to 1.8 K , confirming only the population of a well isolated $S=$ $5 / 2$ ground state in this temperature range. As further confirmation, magnetization $(M)$ versus field $(H)$ and $T$ data were collected in the $0.1-7.0 \mathrm{~T}$ and $1.8-10.0 \mathrm{~K}$ ranges and plotted as $M / N \mu_{\mathrm{B}}$ versus $H / T$ in Figure 7. The data were fit using the program MAGNET, ${ }^{41}$ which assumes that only the ground state is occupied, includes axial zero-field splitting


Figure 4. Comparison of 1 with the $\mathrm{Fe}_{8}$ layers A of the cations of 2 and 3: (a) $\mathrm{Fe}_{7}$ unit of $\mathbf{1}$ with all but the ipso-C atoms of the Ph rings removed for clarity; (b) $\mathrm{Fe}_{8}$ layer $\mathbf{A}$ of 2 in Figure 3 viewed along the $\mathrm{Fe} 2-\mathrm{Fe} 7$ vector; (c) top $\mathrm{Fe}_{8}$ layer $\mathbf{A}$ of 3 in Figure 3 viewed along the $\mathrm{Fe} 2-\mathrm{Fe} 7$ vector and including the fourth $\mathrm{mda}^{2-}$ chelate. H atoms are omitted for clarity. Color code: octahedral $\mathrm{Fe}^{\mathrm{III}}$ lime green, O red, C and N atoms of $\mathrm{mda}^{2-}$ powder blue, and C gray.


Figure 5. (a) $\chi_{M} T$ vs $T$ for $1 \bullet \mathrm{H}_{2} \mathrm{O}$ in a 0.1 T dc field. The simulation using the $J_{\text {MSC }}$ or $J_{\text {DFT }}$ values is shown as a solid blue line, and the $J_{\text {PHI }}$ fit as a solid red line; (b) ac in-phase $\chi_{\mathrm{M}}{ }^{\prime} T$ vs $T$ at a 1000 Hz oscillation frequency.


Figure 6. a) $\chi_{\mathrm{M}} T$ vs $T$ in a 0.1 T dc field, and (b) $\chi_{\mathrm{M}}{ }^{\prime} T$ vs $T$ with a 1000 Hz oscillation frequency for $\mathbf{2} \cdot 4 \mathrm{H}_{2} \mathrm{O}$ (red $\bigcirc$ ) and $3 \cdot 5 \mathrm{EtOH}$ (blue $\bigcirc$ ).


Figure 7. Reduced magnetization $\left(M / N \mu_{\mathrm{B}}\right)$ vs $H / T$ for $1 \bullet \mathrm{H}_{2} \mathrm{O}$ in the $1.8-10.0 \mathrm{~K}$ range at the indicated applied dc fields. The solid lines are the fit of the data; see the text for the fit parameters.
(zfs), $D \hat{S}_{z}{ }^{2}$, and the Zeeman interaction, and incorporates a full powder average; the corresponding spin Hamiltonian is given by eq 3 .

$$
\begin{equation*}
\hat{H}=D \hat{S}_{z}^{2}+g \mu_{\mathrm{B}} \mu_{0} \hat{S} \cdot H \tag{3}
\end{equation*}
$$

An excellent fit (solid lines in Figure 7) was obtained with $S$ $=5 / 2, g=1.99(1)$, and $D=-0.37(3) \mathrm{cm}^{-1}$. As expected, an excellent fit was also obtained with a positive $D$, giving $S=5 / 2$, $g=1.99(1)$, and $D=+0.45(4) \mathrm{cm}^{-1}$. Both fits are visible in the $g$ versus $D$ error surface generated using the program GRID, ${ }^{42}$ and they are of essentially equal quality (Figure S7) with a slightly greater softness in the positive $D$ value. As magnetization fits are not very sensitive to the sign of $D$, a more sensitive technique such as EPR would be required to reliably determine the sign and magnitude of $D$.
The dc $\chi_{\mathrm{M}} T$ versus $T$ profile for 35 EtOH at low $T$ suggests an $S=0$ ground state, and this was confirmed by the ac $\chi_{\mathrm{M}}{ }^{\prime} T$ versus $T$ data (Figure 6b), which decreases from $9.1 \mathrm{~cm}^{3} \mathrm{~K}$ $\mathrm{mol}^{-1}$ at 15.0 K to $1.2 \mathrm{~cm}^{3} \mathrm{~K} \mathrm{~mol}^{-1}$ at 1.8 K clearly heading for zero at 0 K . This is the same as previously reported for 2 . $4 \mathrm{H}_{2} \mathrm{O}$, whose data are included in Figure 6b for comparison. ${ }^{19}$ The decreasing $\chi_{\mathrm{M}}{ }^{\prime} T$ with decreasing $T$ is consistent with lowlying excited states with $S$ greater than the ground state, as expected for a high-nuclearity complex with dominant AF couplings. No out-of-phase $\chi_{\mathrm{M}}{ }^{\prime \prime}$ signals were observed for $\mathbf{1 - 3}$ (Figure S8).

Ground-State Spin Rationalization Using MSC and DFT Methods. Rationalization of the experimental $S=5 / 2$ and $S=0$ ground states of $\mathbf{1}$ and the cations of $\mathbf{2}$ and $\mathbf{3}$ requires determination of the constituent $\mathrm{Fe}_{2}$ pairwise exchange coupling parameters, $J_{i j}$. We thus sought these via two independent methods, a semiempirical MSC developed for
high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ clusters ${ }^{28}$ and broken-symmetry DFT calculations. $J_{i j}$ values from the MSC ( $J_{\mathrm{MSC}}$ ) were calculated following the protocol delineated by Mitchell et al. ${ }^{28}$ and the use of eq 4, where $r$ is the average $\mathrm{Fe}-\mathrm{O}$ bond length and $\varphi$ is the $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}$ angle

$$
\begin{equation*}
J_{i j}=\left(1.23 \times 10^{9}\right)\left(-0.12+1.57 \cos \varphi+\cos ^{2} \varphi\right) \mathrm{e}^{-8.99 r} \tag{4}
\end{equation*}
$$

within each bridged $\mathrm{Fe}_{2}$ pair. For $\mathrm{Fe}_{2}$ pairs with multiple monoatomic bridges, the longer average $\mathrm{Fe}^{\mathrm{III}}-\mathrm{O}$ bond was used in accordance with the protocol. ${ }^{28}$

For 1, $J_{\text {MSC }}$ and $J_{\text {DFT }}$ are collected in Table 2. The crystallographic $C_{3}$ symmetry gives four independent $J_{i j}$ parameters, $J_{1}$ and $J_{2}$ in the outer $\mathrm{Fe}_{6}$ loop and $J_{3}$ and $J_{4}$ to the central Fe 2 (Figure 8). The $J_{1}-J_{4}$ determined from the DFT and MSC calculations are satisfyingly in agreement in both sign and magnitude, with the weakest interaction being $J_{3}$, as expected for $\mathrm{Fe}_{2}$ pairs with two monoatomic bridges owing to their smaller $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}$ angles ( 96.6 and $99.9^{\circ}$ ); the other $\mathrm{Fe}_{2}$ pairs all have only one monoatomic bridge and consequently larger $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}$ angles (119.5-135.1 ${ }^{\circ}$ ). Since all interactions are AF and the $\mathrm{Fe}_{7}$ topology comprises six edge-fused $\mathrm{Fe}_{3}$ triangles, the complex is expected to experience spin frustration, here defined in the way most appropriate for molecular systems as competing exchange interactions that prevent (frustrate) the preferred spin vector alignments. $J_{1}$ and $J_{4}$ are both strong relative to $J_{2}$ and $J_{3}$, which are intermediate and weak, respectively, so the former two dominate leading to antiparallel alignments both between Fe 2 and the $\mathrm{Fe} 3 / \mathrm{Fe} 3 \mathrm{~A} /$ Fe 3 B set and between the latter and the $\mathrm{Fe} 1 / \mathrm{Fe} 1 \mathrm{~A} / \mathrm{Fe} 1 \mathrm{~B}$ set. This gives an alternating "spin-up, spin-down" alignment pattern around the outer $\mathrm{Fe}_{6}$ loop corresponding to $m_{s}= \pm{ }^{5} / 2$ $z$-components of spin and "spin-up" on the central Fe 2 (Figure 8 B ). This situation also satisfies the $J_{2}$ interactions, but the resulting parallel alignments between Fe 2 and the $\mathrm{Fe} 1 / \mathrm{Fe} 1 \mathrm{~A} /$ Fe1B set frustrate $J_{3}$ (red dashed lines in Figure 8b), which is AF but much too weak to compete with $J_{1}$ and $J_{4}$. This is the reason that a "spin-up"/"spin-down" alignment corresponding to $m_{s}= \pm{ }^{5} / 2 z$-components of spin results, rather than any intermediate spin vector alignments. Thus, the total spin of $\mathbf{1}$ is $S=20 / 2-15 / 2=5 / 2$, rationalizing the experimentally observed ground state. Note that the same spin vector alignments and the overall $S=5 / 2$ are obtained when either the $J_{\mathrm{MSC}}$ or $J_{\mathrm{DFT}}$ values for $J_{1}-J_{4}$ are used in this analysis.

The $J_{\mathrm{MSC}}$ values also give an excellent simulation using the program $\mathrm{PHI}^{31}$ of the experimental $\chi_{\mathrm{M}} T$ versus $T$ data (blue line in Figure 5a) indicating an $S=5 / 2$ ground state and an $S=$ $7 / 2$ first excited state at $157 \mathrm{~cm}^{-1}$ above it. Interestingly, when the $J_{\mathrm{MSC}}$ or $J_{\mathrm{DFT}}$ values were used as inputs to fit the data in the

Table 2. Exchange Interactions $J_{1}-J_{4}$ for 1 from MSC Calculations, Broken Symmetry DFT Methods, and PHI Fitting of the Experimental dc Data

| $J^{a}$ | $\mathrm{Fe}-\mathrm{O}^{b} \mathrm{~b}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{\text {b }}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $J_{\text {DFT }}{ }^{\text {c, }{ }^{\text {d }}}$ | $\mathrm{JPHI}^{c}{ }^{\text {c,e }}$ | $J_{0}{ }^{c}{ }_{2}$ | $J_{-10}{ }^{c}{ }^{\text {f }}$ | $J_{-20}{ }^{c}{ }^{\text {f }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $J_{1}$ | 1.857 | 119.5 | -45.0 | -57.0 | -28.7(9) | -21.3(3) | -55.6(4) | -23.4(1) |
| $J_{2}$ | 2.009 | 128.6 | -12.6 | -16.3 | -30.0(8) | -44.3(8) | -47.4(9) | -56.7(9) |
| $J_{3}$ | 1.987 | 96.6 | -6.2 | -4.9 | -13.5(8) | -24.5(8) | +67.5(9) | +24.9(7) |
| $J_{4}$ | 1.915 | 135.1 | -30.1 | -33.4 | -42.6(6) | -7.3(8) | -25.7(3) | -40.5(8) |
| TIP ${ }^{g}$ |  |  |  |  | 700 | 700 | 700 | 700 |

[^1]

Figure 8. (a) Core of 1 showing the labeling scheme for the constituent exchange interactions. (b) Diagrammatic representation of the core showing the $J_{\text {MSC }}$ exchange interactions $\left(\mathrm{cm}^{-1}\right)$ from Table 2, and the $m_{s}= \pm^{5} / 2$ spin alignments predicted by both the $J_{\text {MSC }}$ and the $J_{\mathrm{DFT}}$ values. Satisfied interactions are blue dashed lines, and frustrated pathways are red dashed lines.
$10.0-300 \mathrm{~K}$ range (data below 10 K were excluded to avoid effects such as zero-field splitting), a slightly improved fit was obtained (red line in Figure 5a) but with significantly altered $J_{\text {PHI }}$ values (Table 2). The fitted $J_{\text {PHI }}$ values support the $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ in that $J_{3}$ is again the weakest coupling constant, but given the significant changes in absolute values and an apparent "switch" in the values of $J_{1}$ and $J_{4}$, we suspect that the $J_{\mathrm{PHI}}$ fit is an artifact of overparameterization. Thus, since $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ are similar and the simulation is already excellent, we conclude that they are a much better indication of the true $J$ values. As a control, fitting of the data with three other sets of input values were also tested: $J_{1}=J_{2}=J_{3}=J_{4}=0$, -10 , or $-20 \mathrm{~cm}^{-1}$. Excellent fits were obtained for each, but the $J_{0}, J_{-10}$, and $J_{-20}$ values were unreasonable, especially in the strongly AF and/or F values for $J_{3}$ (Table 2). We thus conclude that the MSC and DFT approaches provide a synergistic duo of distinct methods for determining the exchange couplings in high-nuclearity $\mathrm{Fe}^{\mathrm{II}} / \mathrm{O}$ clusters where fits of data become unreliable due to overparameterization. We thus now applied them to the $\mathrm{Fe}_{22}$ and $\mathrm{Fe}_{24}$ cations of 2 and 3.

To our knowledge, a multipronged analysis using the MSC and theoretical methods of high-nuclearity $\mathrm{Fe} / \mathrm{O}$ systems has never been attempted before, although the MSC was used in conjunction with experimental fits for complexes of nuclearity up to $\mathrm{Fe}_{12}{ }^{29 \mathrm{a}}$ As there are 62 and 68 symmetry-inequivalent $J_{i j}$ couplings in 2 and 3, respectively, attempts to fit the experimental $\chi_{\mathrm{M}}{ }^{T}$ versus $T$ data were never considered. Instead, we implemented the MSC and DFT methods to estimate the $J_{i j}$ values and then tried to rationalize the experimentally determined $S=0$ ground states of both cations by identifying, if possible, spin frustration effects and the resulting spin vector alignments at each $\mathrm{Fe}^{\mathrm{III}}$, that is, an analogous analysis to that for 1 . The resulting $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ values for 2 and 3 are collected in Tables 3-5 covering $J_{i j}$ values (i) within the central $\mathrm{Fe}_{8}$ layers $\mathbf{B}$, (ii) at the interface between layers $\mathbf{A}$ and $\mathbf{B}$, and (iii) within the top $\mathrm{Fe}_{8}$ layers $\mathbf{A}$

Table 3. $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ Exchange Interactions in the Top $\mathrm{Fe}_{8}$ Layer $A$ in the Cations of 2 and 3

| $J^{a}$ | 2 |  |  |  | 3 |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{Fe}-\mathrm{O}^{b}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{b}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $J_{\text {DFT }}{ }^{\text {c }}$ | $\mathrm{Fe}-\mathrm{O}^{\text {b }}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{\text {b }}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $J_{\text {DFT }}{ }^{\text {c }}$ |
| $J_{1,2}$ | 1.996 | 97.5 | -6.1 | -3.0 | 1.992 | 97.2 | -6.2 | -3.5 |
| $J_{1,5}$ | 1.930 | 116.7 | -22.4 | -30.6 | 1.961 | 117.2 | -17.1 | -23.1 |
| $J_{1,6}$ | 1.977 | 129.9 | -16.8 | -20.3 | 2.001 | 127.6 | -12.4 | -16.8 |
| $J_{1,7}$ | 2.052 | 135.9 | -8.8 | -17.7 | 1.973 | 127.6 | -17.1 | -21.0 |
| $J_{2,3}$ | 1.995 | 97.3 | -6.0 | -2.3 | 2.011 | 95.9 | -4.7 | -0.5 |
| $J_{2,4}$ | 1.996 | 97.9 | -6.3 | -4.5 | 2.013 | 95.6 | -4.5 | +1.1 |
| $J_{2,5}$ | 2.038 | 135.2 | -9.9 | -17.5 | 2.024 | 134.4 | -11.2 | -17.7 |
| $J_{2,6}$ | 2.052 | 134.0 | -8.8 | -15.5 | 2.07 | 139.0 | -7.5 | -14.8 |
| $J_{2,7}$ | 2.160 | 82.7 | +0.4 | +6.0 | 2.11 | 83.6 | +0.4 | +5.9 |
| $J_{2,8}$ | 2.044 | 134.7 | -9.4 | -16.3 | 2.038 | 136.4 | -10.0 | -17.6 |
| $J_{3,5}$ | 1.991 | 129.6 | -14.9 | -19.3 | 2.005 | 127.0 | -12.9 | -18.1 |
| $J_{3,7}$ | 2.030 | 136.4 | -10.7 | -19.5 | 2.026 | 134.5 | -11.0 | -19.2 |
| $J_{3,8}$ | 1.946 | 116.3 | -19.3 | -25.8 | 1.943 | 116.7 | -19.9 | -26.6 |
| $J_{4,6}$ | 1.964 | 117.3 | -16.6 | -23.2 | 1.978 | 114.1 | -14.1 | -20.0 |
| $J_{4,7}$ | 1.997 | 129.4 | -14.1 | -21.4 | 2.016 | 136.6 | -12.2 | -22.1 |
| $J_{4,8}$ | 1.979 | 128.8 | -16.4 | -20.4 | 1.997 | 127.9 | -13.9 | -18.6 |
| $J_{5,7}$ | 2.094 | 90.7 | -1.1 | -1.6 | 2.036 | 94.3 | -3.2 | -3.3 |
| $J_{6,7}$ | 2.053 | 92.3 | -2.2 | -1.4 | 2.073 | 91.9 | -1.7 | -1.7 |
| $J_{7,8}$ | 2.078 | 90.4 | -1.3 | -1.3 | 2.053 | 91.9 | -2.0 | -1.9 |

[^2]Table 4. $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ Exchange Interactions in the Interface Between the Top and Bottom $\mathrm{Fe}_{8}$ Layers A and the Central Layer $B$ in the Cations of 2 and 3

|  | 2 |  |  |  | 3 |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| $J^{a}$ | $\mathrm{Fe}-\mathrm{O}^{b}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{\text {b }}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $\mathrm{JDFT}^{\text {c }}$ | $\mathrm{Fe}-\mathrm{O}^{\text {b }}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{\text {b }}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $J_{\text {DFT }}{ }^{c}$ |
| $J_{5,10}$ | 1.918 | 130.9 | -28.7 | -32.4 | 1.886 | 130.1 | -38.3 | -40.8 |
| $J_{6,9}$ | 1.880 | 132.9 | -41.0 | -43.8 | 1.891 | 135.6 | -37.2 | -41.6 |
| $J_{7,9}$ | 1.890 | 125.4 | -35.6 | -38.0 | 1.889 | 120.8 | -34.3 | -35.5 |
| $J_{7,10}$ | 1.891 | 119.7 | -33.2 | -36.4 | 1.890 | 128.0 | -36.3 | -36.3 |
| $J_{7,11}$ | 1.901 | 124.1 | -32.1 | -36.2 | 1.911 | 121.5 | -28.4 | -31.5 |
| $J_{8,11}$ | 1.914 | 127.8 | -29.2 | -34.1 | 1.916 | 132.0 | -29.4 | -32.5 |
| $J_{12,16}$ | 1.900 | 131.2 | -33.8 | -33.8 | 1.900 | 130.8 | -33.8 | -33.6 |
| $J_{12,17}$ | 1.892 | 123.3 | -34.3 | -37.6 | 1.901 | 125.5 | -32.5 | -32.7 |
| $J_{13,17}$ | 1.879 | 127.7 | -40.2 | -41.4 | 1.894 | 121.5 | -33.2 | -35.0 |
| $J_{13,18}$ | 1.901 | 130.9 | -33.5 | -37.7 | 1.900 | 131.6 | -34.0 | -33.3 |
| $J_{14,17}$ | 1.880 | 124.8 | -39.0 | -42.4 | 1.897 | 122.7 | -32.8 | -37.6 |
| $J_{14,19}$ | 1.894 | 132.7 | -36.1 | -38.6 | 1.898 | 127.4 | -33.7 | -40.6 |
| $J_{6,24}$ |  |  |  |  | 2.031 | 128.2 | -10.2 | -14.7 |

${ }^{a}$ See Figures 10 and 11 for 2 and 3, respectively. ${ }^{b}$ Average in $\AA$, and deg. ${ }^{c} \mathrm{~cm}^{-1} ; \bar{H}=-2 J_{i j} \bar{S} \cdot \bar{S}$ convention.
Table 5. $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ Exchange Interactions for Central Layer B in the Cations of 2 and 3

| $J^{a}$ | 2 |  |  |  | 3 |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{Fe}-\mathrm{O}^{b}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{b}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $J_{\text {DFT }}{ }^{\text {c }}$ | $\mathrm{Fe}-\mathrm{O}^{b}$ | $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}^{\text {b }}$ | $J_{\text {MSC }}{ }^{\text {c }}$ | $J_{\text {DFT }}{ }^{\text {c }}$ |
| $J_{9,10}$ | 2.048 | 94.5 | -3.0 | -0.3 | 2.018 | 114.8 | -9.8 | -14.0 |
| $J_{9,11}$ | 1.950 | 141.9 | -22.2 | -28.9 | 2.021 | 120.9 | -10.5 | -14.7 |
| $J_{9,12}$ | 2.004 | 108.9 | -9.7 | -8.2 | 2.037 | 122.6 | -9.3 | -11.9 |
| $J_{10,11}$ | 2.102 | 96.4 | -2.2 | -3.7 | 2.104 | 94.8 | -1.8 | -2.5 |
| $J_{10,12}$ | 2.105 | 102.5 | -3.1 | -6.1 | 2.087 | 100.8 | -3.3 | -3.5 |
| $J_{10,13}$ | 2.095 | 103.4 | -3.5 | -5.4 | 2.077 | 104.0 | -4.2 | -7.0 |
| $J_{11,12}$ | 2.086 | 94.7 | -2.1 | -1.0 | 2.083 | 95.2 | -2.3 | +0.3 |
| $J_{11,13}$ | 2.125 | 101.5 | -2.4 | -4.4 | 2.116 | 100.9 | -2.6 | -6.3 |
| $J_{11,14}$ | 2.029 | 116.4 | -9.1 | -11.3 | 1.998 | 113.5 | -11.5 | -13.1 |
| $J_{12,13}$ | 2.082 | 95.8 | -2.5 | -2.3 | 2.095 | 96.4 | -2.3 | -1.6 |
| $J_{12,14}$ | 1.995 | 128.3 | -14.2 | -22.8 | 1.981 | 138.5 | -16.7 | -24.6 |
| $J_{13,14}$ | 2.015 | 113.8 | -9.9 | -16.8 | 2.012 | 97.8 | -5.4 | -3.4 |
| $J_{9,23}$ |  |  |  |  | 1.897 | 135.3 | -35.4 | -39.1 |
| $J_{9,24}$ |  |  |  |  | 1.996 | 95.5 | -5.2 | -11.6 |
| $J_{11,23}$ |  |  |  |  | 1.976 | 125.7 | -16.5 | -16.3 |
| $J_{12,23}$ |  |  |  |  | 1.964 | 133.2 | -19.2 | -20.49 |
| $J_{23,24}$ |  |  |  |  | 1.866 | 121.8 | -42.7 | -45.4 |

${ }^{a}$ See Figures 10 and 11 for 2 and 3, respectively. ${ }^{b}$ Average in $\AA$, and deg. ${ }^{c} \mathrm{~cm}^{-1} ; \bar{H}=-2 J_{i j} \bar{S}_{i} \cdot \bar{S}_{j}$ convention.
(from the viewpoint of Figure 4), respectively. $J_{i j}$ for the bottom $\mathrm{Fe}_{8}$ layers $\mathbf{A}$ are tabulated in Table S7.

Even with such large $\mathrm{Fe}_{22} / \mathrm{Fe}_{24}$ nuclearity clusters, $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ are in very good agreement in terms of both sign and magnitude. For convenience, we consider $J_{i j}$ to be of three types: weak, intermediate, and strong, with approximate $\left|J_{i j}\right|$ of $<10 \mathrm{~cm}^{-1}, \sim 10$ to $\sim 25 \mathrm{~cm}^{-1}$, and $>25 \mathrm{~cm}^{-1}$, respectively. As stated, $\left|J_{i j}\right|<10 \mathrm{~cm}^{-1}$ are characteristic of $\mathrm{Fe}_{2}$ pairs with two monoatomic bridges, whereas stronger couplings are expected with only one monoatomic bridge. DFT calculations are on the complete cations and are affected by the total nuclearity, whereas MSC calculations are independent of the nuclearity, being localized at each $\mathrm{Fe}_{2}$ pair in turn. So, the very satisfying agreement in $J_{\text {MSC }}$ and $J_{\text {DFT }}$ for essentially every interaction emphasizes the power and utility of the DFT approach even for such complicated systems. The $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ only differ in sign for two couplings each in 2 and 3: $J_{\mathrm{MSC}} / J_{\mathrm{DFT}}$ for $J_{15,21}=$ $-4.9 \mathrm{~cm}^{-1} /+1.1 \mathrm{~cm}^{-1}$ and $J_{17,18}=-2.1 \mathrm{~cm}^{-1} /+0.8 \mathrm{~cm}^{-1}$ for 2 , and $J_{2,4}=-4.5 \mathrm{~cm}^{-1} /+1.1 \mathrm{~cm}^{-1}$ and $J_{11,12}=-2.3 \mathrm{~cm}^{-1} /+0.3$
$\mathrm{cm}^{-1}$ for 3 . This sign discrepancy is not surprising given that these couplings are so weak.

Since the top $\mathrm{Fe}_{8}$ layers A are similar to $\mathbf{1}$, we shall discuss them first. The $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ values are listed in Table 3, and diagrammatic representations are provided in Figure 9 showing the $J_{\mathrm{MSC}}$ locations; note that use of the $J_{\mathrm{DFT}}$ values in Figure 9 would lead to the same conclusions reached below. The spin alignments in the $\mathrm{Fe}_{8}$ propeller unit in 2 and 3 can be determined by identifying the spin frustrated pathways, as was done for $\mathbf{1}$. Notwithstanding that the extra central Fe 7 forming the axle with Fe 2 causes some structural perturbation relative to 1 , a similar spin frustration pattern results between the $\mathrm{Fe}_{8}$ layer A of $\mathbf{2}$ and $\mathbf{3}$ and that of $\mathbf{1}$, that is, the weakest $\mathrm{AF} J_{i j}$ for bis-monoatomically bridged $\mathrm{Fe}_{2}$ pairs $\left(J_{1,2}, J_{2,3}, J_{2,4}, J_{5,7}, J_{6,7}\right.$, and $J_{7,8}$ ) is frustrated by the two stronger $J_{i j}$ in each $\mathrm{Fe}_{3}$ triangular unit, leading to parallel alignments in these $\mathrm{Fe}_{2}$ pairs as the outer spin vectors align in an antiparallel up-down fashion around the outer hexagon. This also leads to the spin vectors $\mathrm{Fe} 2 / \mathrm{Fe} 7$ being aligned antiparallel, which means that $J_{2,7}$ is also frustrated if it really is F , as indeed $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ indicate in


Figure 9. Diagrammatic representation of the cores of the top $\mathrm{Fe}_{8}$ layer $\mathbf{A}$ of (a) 2 and (b) 3 showing the $J_{\mathrm{MSC}}$ exchange interactions $\left(\mathrm{cm}^{-1}\right)$ from Table 3, and the $m_{\mathrm{s}}= \pm^{5} / 2$ spin alignments predicted by both the $J_{\mathrm{MSC}}$ and the $J_{\mathrm{DFT}}$ values. Satisfied interactions are blue dashed lines, and frustrated pathways are red dashed lines.
both 2 and 3 (Table 3). This is reasonable given that the $\mathrm{Fe} 2 /$ Fe 7 axle pair is tris-monoatomically bridged by three $\mu_{4}-\mathrm{O}^{2-}$ ions with very acute $\mathrm{Fe}-\mathrm{O}-\mathrm{Fe}$ angles in the $82.7-86.5^{\circ}$ range. Overall, the $\mathrm{Fe}_{8}$ unit is thus predicted to have an $S=10-10=$ 0 ground state. A similar analysis of the bottom $\mathrm{Fe}_{8}$ layers A of 2 and 3 leads to an analogous alignment pattern and a predicted $S=0$ ground state (Figure S9).
A similar analysis can now be carried out on the interfaces between the layers $\mathbf{A}$ and the central $\mathrm{Fe}_{6}$ and $\mathrm{Fe}_{8}$ layers $\mathbf{B}$ of the cations of 2 and 3 , respectively, and then on the central layers themselves. We shall describe the former first because the $J_{i j}$ in the central layer are almost all weak so that their spin alignments are strongly dominated by the $J_{i j}$ at the interfaces. $J_{\mathrm{MSC}}$ and $J_{\text {DFT }}$ at the latter in the cation of 2 (Table 4 and Figure 10) show that all $J_{i j}$ between $\mathrm{Fe} 5,6,7,8$ of top layer A and $\mathrm{Fe} 9,10,11$ of the central layer B are strongly AF , and since the spin vectors of the former are all parallel due to the spin frustration effects described above, this aligns the spin vectors of $\mathrm{Fe} 9,10,11$ all parallel to each other and antiparallel to those of $\mathrm{Fe} 5,6,7,8$, frustrating the weaker AF interactions $J_{9,10}, J_{9,11}$, and $J_{10,11}$ (red dashed lines in Figure 10): $J_{9,10}$ and $J_{10,11}$ are very weak (Table 5), as expected for bis-monoatomically bridged $\mathrm{Fe}_{2}$ pairs, and easily frustrated, whereas $J_{9,11}$ is of intermediate strength $\left(J_{\mathrm{MSC}} / J_{\mathrm{DFT}}=-22.2 /-28.9 \mathrm{~cm}^{-1}\right)$ since it has a single monoatomic bridge, but it is competing with the
four strong $J_{6,9}, J_{7,9}, J_{7,11}$, and $J_{8,11}$ in the ranges $J_{\mathrm{MSC}}=-29.2$ to $-41.0 \mathrm{~cm}^{-1}$ and $J_{\mathrm{DFT}}=-34.1$ to $-43.8 \mathrm{~cm}^{-1}$.
The same situation is seen at the interface of the bottom $\mathrm{Fe}_{8}$ layer A of $\mathbf{2}$ with the central layer $\mathbf{B}$. $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ again show that all $J_{i j}$ between Fe16,17,18,19 of the bottom layer A and Fe12,13,14 of the central layer B are strongly AF (Table 4 and Figure 10), and since the spin vectors of the former are all parallel, this again aligns the spin vectors of $\mathrm{Fe} 12,13,14$ all parallel to each other and antiparallel to those of Fe16,17,18,19, frustrating the weaker AF interactions $J_{12,13}$, $J_{12,14}$, and $J_{13,14}$ (red dashed lines in Figure 10); $J_{12,14}$ is of intermediate strength $\left(J_{\mathrm{MSC}} / J_{\mathrm{DFT}}=-14.2 /-22.8 \mathrm{~cm}^{-1}\right)$ (Table 5), but it is competing with the four strong $J_{12,16}, J_{12,17}, J_{14,17}$, and $J_{14,19}$ in the ranges $J_{\mathrm{MSC}}=-33.8$ to $-39.0 \mathrm{~cm}^{-1}$ and $J_{\mathrm{DFT}}=$ -33.8 to $-42.4 \mathrm{~cm}^{-1}$. The remaining step is to assess the situation in the central $\mathrm{Fe}_{6}$ layer B of 2, and it is clear from Table 5 that almost all $J_{i j}$ are AF and weak except the two mentioned already, that is, $J_{9,11}$ and $J_{12,14}$. However, even though all $J_{i j}$ between the $\mathrm{Fe} 9,10,11$ and $\mathrm{Fe} 12,13,14$ planes are weak, they are not competing and thus none are frustrated by the resulting antiparallel alignments of the spin vectors of these two planes (Figure 10).

Comparing now the $J_{\mathrm{MSC}} / J_{\mathrm{DFT}}$ values for 2 versus 3 in Tables 3-5 leads to the conclusion that the spin vector alignments in the larger $\mathrm{Fe}_{24}$ cation of $\mathbf{3}$ are analogous to those in 2 (Figure 11). The main difference is, of course, the extra Fe 23 and Fe 24 ions in the now $\mathrm{Fe}_{8}$ central layer $\mathbf{B}$, and these have entered one each into what were the two $\mathrm{Fe}_{3}$ planes to make them $\mathrm{Fe}_{4}$. Otherwise, the same pattern of spin vector alignments is seen, that is, strongly AF interactions in the interface region between layers $\mathbf{A}$ and $\mathbf{B}$, frustrating the weaker interactions within the $\mathrm{Fe}_{4}$ planes and thus leading to parallel and antiparallel spin alignments within and between the $\mathrm{Fe}_{4}$ planes, respectively. $J_{12,14}$ and $J_{12,23}$ are again of intermediate strength but are nevertheless frustrated by the multiple stronger interactions with which they are competing, $J_{12,16}$, $J_{12,17}, J_{9,23}$, and $J_{23,24}$.

The overall picture that emerges is that in all areas of the cations of 2 and 3, there is complete frustration of one interaction in each $\mathrm{Fe}_{3}$ triangular unit either because the other two interactions are much stronger, or because strong interactions in neighboring $\mathrm{Fe}_{3}$ triangular units are enforcing the frustration, or both. In effect, the $\mathrm{Fe}_{3}$ units are magnetically near-isosceles, and this is primarily due, of course, to the common occurrence in higher-nuclearity $\mathrm{Fe}^{\mathrm{II}} / \mathrm{O}$ clusters of bis-monoatomically bridged $\mathrm{Fe}_{2}$ pairs giving characteristically very weak $J_{i j}$ values. This also leads in the cations of 2 and 3 to a "spin-up"/"spin-down" alignment of the spin vectors corresponding to $m_{s}= \pm{ }^{5} / 2 z$-components of spin, as summarized in Figure 12. The spin alignments described above lead to an $S=0$ ground state for the three A:B:A layers, and so the overall spin is predicted to be $S=0$ for both cations, in agreement with the experimental susceptibility data. As a consistency check for the interpretation of the $S=0$ ground state, we also determined the lowest-energy configuration of the cations of 2 and 3 by explicitly evaluating the $2^{22}$ and $2^{24}$ energies given by eq 1 using both the $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ sets of couplings. The resulting lowest-energy configuration was found in all cases to be the same as shown in Figure 12.

## - CONCLUSIONS

The use of $\mathrm{mdaH}_{2}$ in $\mathrm{Fe}^{\text {III }} / \mathrm{O}$ carboxylate cluster chemistry has led to three products with $\mathrm{Fe}_{7}, \mathrm{Fe}_{22}$, and $\mathrm{Fe}_{24}$ nuclearities, the


Figure 10. Diagrammatic representation of the core of the central $\mathrm{Fe}_{6}$ layer $\mathbf{B}$ of $\mathbf{2}$ plus selected atoms from the two $\mathrm{Fe}_{8}$ layers $\mathbf{A}$, showing the $J_{\mathrm{MSC}}$ exchange interactions $\left(\mathrm{cm}^{-1}\right)$ from Tables 4 and 5 in (left) the interface regions between $\mathbf{A}$ and $\mathbf{B}$ and (right) the central layer $\mathbf{B}$. For clarity, $J_{11,14}$ is shown on the left. Also shown are the ground-state $m_{s}= \pm^{5} / 2$ spin alignments predicted by both $J_{\mathrm{MSC}}$ and the $J_{\mathrm{DFT}}$ values. Satisfied interactions are blue dashed lines, and frustrated pathways are red dashed lines.


Figure 11. Diagrammatic representation of the core of the central $\mathrm{Fe}_{8}$ layer $\mathbf{B}$ of $\mathbf{3}$ plus selected atoms from the two $\mathrm{Fe}_{8}$ layers $\mathbf{A}$, showing the $J_{\mathrm{MSC}}$ exchange interactions $\left(\mathrm{cm}^{-1}\right)$ from Tables 4 and 5 in (left) the interface regions between $\mathbf{A}$ and $\mathbf{B}$ and (right) the central layer $\mathbf{B}$. For clarity, $J_{11,14}$ and $J_{11,23}$ are shown on the left. Also shown are the ground-state $m_{s}= \pm \frac{5}{2}$ spin alignments predicted by both $J_{\mathrm{MSC}}$ and the $J_{\mathrm{DFT}}$ values. Satisfied interactions are blue dashed lines, and frustrated pathways are red dashed lines.
different products resulting from a systematic screening of different carboxylic acids and the presence or absence of additional base. There is also a clear structural relationship between them; the $\mathrm{Fe}_{22}$ and $\mathrm{Fe}_{24}$ are overall very similar, and the $\mathrm{Fe}_{7}$ is related to the $\mathrm{Fe}_{8}$ units at each end of the larger clusters. As is usual in high-nuclearity $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ clusters, there are many bis-monoatomically bridged $\mathrm{Fe}_{2}$ pairs with resulting
weak $J_{i j}$ exchange coupling, and these are completely frustrated by competing strong interactions, leading to Ising-like $m_{s}=$ $\pm 5 / 2$ spin alignments and thus a ready rationalization of the overall ground-state $S$ values for the three clusters. The synergistic use of the polynuclear $\mathrm{Fe}^{\mathrm{III}} / \mathrm{O}$ MSC and DFT methods represents a powerful complement to experimental magnetic studies, providing an overall three-pronged analytical


Figure 12. Summary of the ground-state $m_{s}= \pm^{5} / 2$ spin alignments for the complete cations of (left) 2 and (right) 3 predicted by both the $J_{\text {MSC }}$ and the $J_{\mathrm{DFT}}$ values, and also the energy calculations by DFT. Color code: octahedral $\mathrm{Fe}^{\mathrm{III}}$ lime green, square-pyramidal $\mathrm{Fe}^{\mathrm{III}}$ lavender, O red, and $\mathrm{OH}^{-}$ sky blue.
approach of great potential for application to a variety of cluster nuclearities and topologies. This is well evidenced in the present work, which has led to a highly satisfying elucidation of the ground-state spin configuration in such high-nuclearity systems. Further studies at even higher $\mathrm{Fe}_{x}$ nuclearities are in progress.

## ASSOCIATED CONTENT

## (s) Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.inorgchem.2c01371.

Selected bond distances, bond valence sums, RMSD analyses, $J_{\mathrm{MSC}}$ and $J_{\mathrm{DFT}}$ exchange interactions, various structural figures, $g$ versus $D$ error surface for $\mathbf{1}$, and $\chi_{\mathrm{M}}{ }^{\prime \prime}$ versus $T$ plots for $\mathbf{1}, \mathbf{2}$, and $\mathbf{3}$ (PDF)

## Accession Codes

CCDC 274154, 2163470 , and 2163471 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/ cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223336033.

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## Notes

The authors declare no competing financial interest. An anonymous referee analyzed the $\mathrm{Fe}_{7}$ cluster $\mathbf{1}$ by employing the irreducible tensor approach. The total number of magnetic states $N=6^{7}=279,936$ in the partition function can be processed using the $M=24,017$ zero-field states. Such a matrix can be divided into blocks of the same molecular spin: for $S=$ $1 / 2$ to $S={ }^{35} / 2$, the size of each block is $1050,1974,2666,3060$, 3150, 2975, 2604, 2121, 1610, 1140, 750, 455, 252, 126, 56, 21,6 , and 1 . Using the four exchange coupling constants $J_{1}-J_{4}$ obtained from the MSC approach (Figure 8), the calculated ground state is $S=5 / 2$ separated from an $S=7 / 2$ first excited state by $157 \mathrm{~cm}^{-1}$ and from the second excited state of a degenerate $S=3 / 2$ pair by $190 \mathrm{~cm}^{-1}$. The complete energy spectrum is shown in Figure S11. We thank the referee for
taking the time to do this and the interest in our work that it reflects, which we take as a very nice compliment.

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[^1]:    ${ }^{a}$ See labeling in Figure 8. ${ }^{b}$ Average in $\AA$, and deg. ${ }^{c} \mathrm{~cm}^{-1} ; \bar{H}=-2 J_{i j} \bar{S}_{i} \cdot \bar{S}_{j}$ convention. ${ }^{d}$ The DFT calculations gave all $J_{i j}$ values independently, so only one is shown for symmetrically equivalent sets. For $J_{1}$ and $J_{3}$, all three were identical. For $J_{2}$ and $J_{4}$, one value differed by $0.1 \mathrm{~cm}^{-1}$. ${ }^{e}$ The input values were the MSC values. ${ }^{f} J_{\#}$ indicates all $J_{1}-J_{4}$ input values (\#) for the fits with PHI. ${ }^{g}$ Temperature-independent paramagnetism, $\times 10^{-6} \mathrm{~cm}^{3} \mathrm{~mol}^{-1}$.

[^2]:    ${ }^{a}$ See Figure $9 .{ }^{b}$ Average in $\AA$, and deg. ${ }^{c} \mathrm{~cm}^{-1} ; \bar{H}=-2 J_{i j} \bar{\delta} \cdot \bar{S}$ convention.

