Determination of the magnetic structure of manganese carbodiimide with diffraction experiments using polarized neutrons

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The magnetic structure of MnNCN, the nitrogen equivalent of MnO, and the first carbodiimide of a non-$d^{10}$ transition metal ever made, has been determined on the basis of spin-polarized neutron-diffraction data close to absolute zero temperature. The $S=5/2$ moments of the high-spin Mn$^{2+}$ ions lie within the $ab$ hexagonal sheets and exhibit an antiferromagnetic ordering between the sheets along the $c$ axis. Below the Néel point (28 K), a temperature-dependent ordering pattern exists due to magnetic frustration inside the sheets, characterized by a varying angle between the moments. A comparison with other magnetic structures reveals that the compact NCN$^{2-}$ group is capable of mediating significant spin density needed to ensure relatively strong magnetic communication in view of a large Mn$^{2+}$–Mn$^{2+}$ separation.

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I. INTRODUCTION

Developing synthetic routes for novel magnetic compounds is an everlasting challenge in inorganic solid-state chemistry. In order to succeed, coupling of magnetically active centers such as $d$- and $f$-configured ions with traditional bridging ligands, i.e., oxide, chalcogenide, or cyanide anions, has been pursued for many decades. As an alternative, the use of the corresponding base of $H_2NCN$, the NCN$^{2-}$ anion which is either named carbodiimide ($D_{ab}$ symmetry, as in $N=C=N$ with two double bonds) or cyanamide (potentially bent, as in $N=C=N^{2-}$ with one triple and one single bond), has recently come to a broader attention. Because of its two minus charge and the fact that NCN$^{2-}$ is electronically dominated by the terminal nitrogen atoms, NCN$^{2-}$ may be regarded a “divalent nitride” anion, thereby allowing us to repeat the entire oxide (O$^{2-}$) chemistry using a nitrogen species.

A good number of inorganic carbodiimides/cyanamides has been synthesized containing alkali$^{1-5}$ and alkaline-earth metals,$^{6-9}$ main-group elements,$^{10-14}$ $d^{10}$ transition metals,$^{15-21}$ and also rare-earth metals.$^{22-25}$ In 2005, the existence of quasibinary compounds containing non-$d^{10}$ magnetically active metals was first predicted using density-functional theory, indicating that all $MNCN$ ($M$=Mn, Fe, Co, Ni, and Cu) phases should be unstable with respect to both formation enthalpy $\Delta H_f$ and Gibbs formation energy $\Delta G_f$. In the meantime, all five synthetic targets have been synthesized, for example, through a metathesis route for MnNCN,$^{27}$ by the thermal decomposition of hydrogen-containing precursors for MNCN ($M$=Fe, Co, and Ni) (Refs. 28 and 29) and by the room-temperature oxidation of a Cu(I) precursor for CuNCN. $^{30}$

Despite its rhombohedral symmetry, the crystal structure of MnNCN (Fig. 1, most conveniently described using a hexagonal setting) resembles the rock-salt motif because of octahedral coordination for both Mn$^{2+}$ and NCN$^{2-}$. Magnetic-susceptibility measurements on this first carbodiimide of a magnetic transition metal ever made yielded antiferromagnetic interactions, a Néel point of about 28 K and a Curie-Weiss behavior above that temperature; the magnetic moment was determined as $\mu=5.84 \mu_B$ from the slope of the Curie–Weiss straight line, in accord with a high-spin Mn$^{2+}$ ion (five unpaired electrons, $S=5/2$) and an expected moment of $\mu=g\sqrt{S(S+1)}\mu_B=5.92 \mu_B$. In order to eventually

![FIG. 1. (Color online) The hexagonal setting of the rhombohedral crystal structure of manganese carbodiimide, MnNCN, with the Mn atoms given in red, N atoms in green, and C atoms in gray. The strictly linear NCN$^{2-}$ carbodiimide groups coincide with the $c$ axis and coordinate the Mn atoms through their terminal N atoms to yield nearly regular octahedra with $6\times Mn-N=2.26$ Å and $\angle (N,Mn,N)=84.1^\circ$ and $95.9^\circ$.](image_url)
disclose the magnetic structure of the nitrogen equivalent of manganese oxide, neutron powder-diffraction measurements have been carried out, both with and without spin-polarized neutrons.

II. EXPERIMENT

Samples of green-colored transparent powderous manganese carbodiimide were prepared via a metathesis reaction between MnCl₂ and ZnNCN to yield MnNCN and ZnCl₂ in quartz ampoules at 650 °C. Single-crystal data confirmed MnNCN to exhibit rhombohedral symmetry (space group R3m) with the hexagonal lattice parameters a = 3.3583(4) Å and c = 14.247(2) Å at room temperature; the atoms are located on the Wyckoff sites 3a (Mn), 3b (C), and 6b (N) with z = 0.5855(3). The first neutron powder-diffraction data of MnNCN were collected at Forschungszentrum Jülich by means of the SV7 diffractometer with λ = 1.0952(4) Å in the 2θ range 4° ≤ 2θ ≤ 90° at temperatures of T = 4.5 and 10 K, between T = 15 and 35 K in steps of 2 K, and finally at room temperature. In addition, neutron measurements with spin-polarization analysis were carried out at FRM II, Garching, with the help of the most advanced diffuse neutron-scattering (DNS) instrument and λ = 4.74 Å in the 2θ range 5° ≤ 2θ ≤ 105° at temperatures of T = 4.5 and 20 K. Although DNS typically serves as a “cold” time-of-flight spectrometer, its use in diffraction mode (with a constant wavelength) but with polarization analysis served the problem of magnetic structure determination quite well. All numerical Rietveld refinements were carried out using the FULLPROF program suite.  

III. RESULTS

The nonpolarized neutron data collected at the SV7 instrument already corroborated the existence of an antiferromagnetic ordering below 28 K (Fig. 2, top) with a complicated temperature-dependent behavior that came quite unexpectedly. In Fig. 3, this phenomenon is more carefully analyzed for the case of the first two SV7 magnetic reflections. Figure 3 also allows a visual comparison with the previously published magnetic-susceptibility data. In particular, cooling below 30 K causes a strong rise in intensity for the first reflection (top) while the second one (middle) is hardly affected. Below 20 K, however, the second peak increases considerably whereas the first one slightly weakens. Indeed, the magnetic-susceptibility mirrors that temperature dependence, highlighted by straight lines (Fig. 3, bottom) which are simple guides to the eye. Obviously, a set of different magnetic orderings exists. Nonetheless, because of insufficient statistics at the SV7 instrument, a definite structure model for the spin structure could not be developed in the very beginning. Thus, additional polarized neutron data had to be collected using the DNS instrument, and these yielded six well-shaped magnetic reflections (see Fig. 2, bottom) below the Néel point, two of which overlap substantially.

These six reflections could be indexed by doubling the unit cell in all directions. The first reflection was indexed as (003), hence the orientation of the spin moments can be excluded to point along the c axis; consequently, the rhombohedral symmetry is entirely destroyed. Instead, the magnetic indexing refers to a triclinic symmetry and all calculations concerning the magnetic structure were performed in space group P̅1. To do so, only the magnetic DNS reflections were refined using the lattice parameters, the Gaussian peak profile, and the angle between the spin moments (S = 5/2) as refinable parameters. The best accordance between the observed and calculated intensities is depicted in Fig. 4 and Table I contains all relevant Rietveld parameters. Table II offers the Bragg positions and indices of the magnetic reflections.

In addition to the aforementioned structural characterization, we independently determined the value of the Mn²⁺ magnetic moment by a simultaneous refinement of the magnetic and the crystallographic reflections on the basis of the SV7 data because such a procedure requires the intensities of the nuclear reflections for scaling purposes. Table III provides all relevant information and the graphical result is depicted in Fig. 5. The magnetic moment based on neutron data arrives at 4.72(15)μ₆, to be compared with the expectation value of the free Mn²⁺ ion which is μₑ = gSμₑ = 5μₑ (g = 2, S = 5/2) for the neutron case; both values are identical within two sigma.
The magnetic structure(s) of MnNCN (Fig. 6) can be described by an antiferromagnetic ordering of the magnetic moments within the $ab$ sheets of the manganese ions stacked along the $c$ axis; despite the triclinic symmetry of the magnetic structure, we still refer to the hexagonal metric of the rhombohedral unit cell for reasons of simplicity. Inside these $ab$ sheets, the spins are orientated perpendicular to the $c$ axis and enclose an angle whose size depends on the temperature, thereby leading to the observed differences in intensities. Because all distances between the manganese ions in the sheets are equal, magnetic spin frustration seemingly occurs. With respect to a (hypothetical) ferromagnetic ordering within one sheet (i.e., $\varphi=0$), the spins are tilted against each other with an angle of $46^\circ$ at 20 K and $100.8^\circ$ at 4.5 K. At $T=20$ K, the (003) reflection gains its maximum intensity, thereby also indicating maximum attainable precision for the magnetic structure; at the lowest measured temperature of $T=4.5$ K, the angle $\varphi$ widens. For a better understanding, Fig. 6 depicts the entire magnetic unit cells at $T=4.5$ and 20 K. We also note that the spin structure at $T=4.5$ K (Fig. 6, top left) alludes to the possibility of a structural distortion but this is clearly beyond the resolution limit of our data.

To further understand the magnetic structure of MnNCN, a comparison with those seen in MnO and also the isostructural $\alpha$-MnS is appropriate. Note that the magnetic structure of MnO serves as the archetype, simply because (a) MnO is the oxide equivalent of MnNCN and (b) the MnO structure was the very first magnetic structure ever investigated and solved by neutron scattering. In 1951, Shull performed neutron-diffraction studies on MnO and suggested a structural distortion but this is clearly beyond the resolution limit of our data.

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| TABLE I. Structure and profile parameters for the refined neutron-diffraction data of MnNCN collected at the DNS instrument at temperatures of $T=4.5$ and 20 K. $\varphi$ refers to the angle between two neighboring spin moments; for illustration, see also Fig. 6. Due to the limited number of reflections, $U$, $V$, and $W$ could not be refined simultaneously such that standard deviations cannot be specified for those values. |
|-----------------|-----------------|-----------------|
|                | $T=4.5$ K       | $T=20$ K        |
| Lattice parameters: (Å) | $a=6.68(1)$ | $a=6.67(5)$ |
|                 | $c=28.68(7)$   | $c=28.66(7)$   |
| $\varphi$ (deg) | 100.8(4)        | 46(2)          |
| $U$, $V$, $W$   | 2.24, -2.15, 2.72 | 2.75, -0.34, 2.48 |
| $R_p$, $R_{df}$ | 0.106, 0.035   | 0.084, 0.019   |
structure model with a cell whose edge is twice the length of
the crystallographic cell and which exhibits ferromagnetic
sheets parallel to the (111) plane. Neighboring planes are
coupled in an antiferromagnetic fashion and the magnetic
moments were suggested to run parallel to the [001] axis.34
Seven years later, the model was revised by Roth, with a
corrected ferromagnetic spin orientation lying within the
(111) plane and this model is covered in many solid-state
physics textbooks. The in-plane orientation of the magnetic
moments with respect to the crystallographic axes, however,
was indeterminable from powder data35 due to overlapping
reflections because of the cubic symmetry. In 1988, a low-
temperature high-resolution neutron study indicated that the
true symmetry of MnO at 8 K must be lower than rhombohedral.36
Total neutron-scattering data eventually yielded, in 2006, that the ferromagnetically aligned spins
within the (111) sheets show a slight preference for spin
alignment along the (112) directions.37
Contrasting this magnetic structure with our aforementioned model for MnNCN, one finds similarities as well as
dissimilarities. For both MnO and MnNCN, the magnetic
structure can be described by ferromagnetic (MnO) and almost
ferromagnetic (MnNCN) layers of Mn$^{2+}$ ions which are
stacked antiferromagnetically along the crystallographically threefold axis, namely, [111] for MnO and [001] for
MnNCN. Nonetheless, the MnNCN structure contains one additional degree of freedom for the orientations of
the spins. The ferromagnetic orientation within one sheet of Mn$^{2+}$ ions corresponds to a constant $\varphi=0^\circ$ scenario (MnO case) between the magnetic moments. For MnNCN, how-

| TABLE III. Structure and profile parameters for the refined neutron-diffraction data of MnNCN collected at the SV7 instrument at $T=4.5$ K. |
|-----------------|-----------------|-----------------|
| Crystallographic | Magnetic |
| Number of reflections: | 55 | 147 |
| Lattice parameters: (Å) | $a=3.329(3)$ | $a=6.66(1)$ |
| | $c=14.247(15)$ | $c=28.49(3)$ |
| Magnetic moment: | 4.72(15)$\mu_B$ |
| $R_p$, $R_{Bragg}$ | 0.070, 0.108 | 0.070, 0.246 |
DETERMINATION OF THE MAGNETIC STRUCTURE OF...

V. CONCLUSION

At lowest temperatures, MnNCN, the nitrogen equivalent of MnO, exhibits significantly different and temperature-dependent magnetic structures due to magnetic frustration in the $ab$ sheets of the rhombohedral structure. Thus, the phenomenon nicely fits into the picture of those magnetic lattices which are geometrically or topologically prone to frustration because of being based on layered triangular motifs. Similar to the case of MnO, the spin layers are antiferromagnetically stacked on top of each other along the crystallographically threefold axis. The numerical data reveal that the carbodiimide group is capable to ensure strong magnetic coupling despite relatively wide Mn$^{2+}$–Mn$^{2+}$ distances between the sheets.

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31. The minor reduction below the expectation $\mu$ value might be attributable by a small diamagnetic impurity due to leftover Zn$^{2+}$ instead of Mn$^{2+}$ cations; unfortunately, the metathesis reaction cannot be made 100% quantitative.


