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ABSTRACT

The spin orientation in synthetic hematite-ilmenite samples and in a sample of natural hematite was studied from room temperature to above the antiferromagnetic-paramagnetic phase transition (the Néel temperature; \( T_N \approx 600–950 \) K) by neutron powder diffraction and at room temperature by Mössbauer spectroscopy. The usually assumed magnetic structure of hematite within this temperature range is antiferromagnetic with the spins confined to the basal plane of the hexagonal structure; however, an out-of-plane spin component is allowed by the symmetry of the system and has been observed in recent studies of synthetic hematite samples. We find the spins in the antiferromagnetic sublattices to be rotated out of the basal plane by an angle between 11(2)° and 22.7(5)° in both synthetic hematite-ilmenite samples and in the natural hematite sample. The spin angle remains tilted out of the basal plane in the entire temperature range below the Néel temperature and does not depend systematically on Ti-content. The results indicate that the out-of-plane spin component is an intrinsic feature of hematite itself, with an origin not yet fully understood, but consistent with group theory. This represents a major shift in understanding of one of the two main mineral systems responsible for rock magnetism.

Keywords: Hematite, magnetic properties, spin orientation, neutron scattering, Mössbauer spectroscopy

INTRODUCTION

The ilmenite-hematite \([x\text{FeTiO}_3-(1-x)\text{Fe}_2\text{O}_3]\) solid-solution series has been studied extensively because of its complex and interesting magnetic and electronic properties. Intermediate compositions are magnetic semiconductors (Ishikawa and Akimoto 1957; Ishikawa 1958) and could conceivably be utilized in spintronics devices (Butler et al. 2003; Fujii et al. 2004). The \( \text{FeTiO}_3-\text{Fe}_2\text{O}_3 \) solid-solution series further serves as a model system for natural ilmenite-hematite minerals, which are studied because of their remarkable magnetic properties and because of their importance as a source of anomalies in the magnetic field of the Earth (McEnroe et al. 2001; Kletetschka et al. 2002), and possibly of other planets like Mars (McEnroe et al. 2004). Interesting magnetic properties of natural ilmeno-hematite samples include giant exchange bias (McEnroe et al. 2007; Fabian et al. 2008) and large and stable remanent magnetization that cannot be explained by the individual properties of hematite and ilmenite alone (McEnroe et al. 2002; Robinson et al. 2002, 2004). There is strong evidence that these properties are associated with fine scale exsolution structures observed in natural hematite-ilmenite samples (Robinson et al. 2002; Fabian et al. 2008; Brok et al. 2014). Solid solution \( \text{FeTiO}_3-\text{Fe}_2\text{O}_3 \) does not exhibit these well-developed exsolution structures and can therefore be used as a baseline for determining if the observed properties are tied to the exsolution structure.

Between the Néel temperature of \( T_N \approx 955 \) K (Morris 1994) and the Morin temperature of \( T_M \approx 264 \) K (Besser et al. 1967; Morin 1950) hematite is a canted antiferromagnet with \( \text{Fe}^{3+} \) spins aligned ferromagnetically within the basal plane of the hexagonal structure while spins in adjacent planes are aligned antiferromagnetically apart from a small canting of about 0.065° that gives rise to a small net magnetization. The spin orientation is usually assumed to be in or very close to the basal plane (Morris 1994; Shull et al. 1951; Flanders 1972) even though the symmetry of the structure allows for an out-of-plane spin component (Dzyaloshinsky 1958). At \( T_N \) hematite undergoes the so-called Morin transition in which the spins rotate 90° to form a perfectly antiferromagnetic structure (no canting) with spins in the two sublattices parallel and antiparallel to the hexagonal [001] axis. The Morin transition is known to be suppressed by impurities as...
well as by finite crystallite size and the canted antiferromagnetic structure with spins close to the basal plane is maintained even at the lowest temperatures in nanoparticles smaller than 20 nm and in samples doped with even small amounts (<1%) of Ti (Bødker et al. 2000; Besser et al. 1967; Morrish 1994).

In a natural sample of ilmeno-hematite with nanoscale exsolution lamellae of ilmenite in the parent hematite structure Harrison et al. (2010) found the spins to be rotated 29.5° out of the basal plane and in a nanocomposite of hematite particles interacting with particles of NiO (Frandsen et al. 2011) measured an out-of-plane angle as large as 70°. These experiments show that exchange interaction across interfaces can change the spin orientation in hematite. In studies of Al-substituted hematite it has been proposed that intermediate structures (spins not parallel or perpendicular to [001]) exist close to the Morin transition (Vandenberghe et al. 2001, 2002). Even in pure hematite at ambient conditions out-of-plane spin-angles of about 18° have been measured by Parise et al. (2006) and Klotz et al. (2013). The spin orientation in hematite between $T_{NM}$ and $T_N$ is thus not always confined to the basal plane. The spin orientation could depend on factors such as impurities and strain and could thus be sample dependent. In particular, the spin orientation in Ti-substituted hematite could deviate from that of pure hematite, and may potentially vary with Ti-content.

The degree of Fe-Ti cation order in hematite-ilmenite solid solutions depends greatly on the composition and thermal history of the sample. Quenched samples containing less than ~40% ilmenite exhibit no long-range Fe-Ti cation order, and adopt the space group $R3c$. Quenched samples containing more than ~40% ilmenite exhibit an increasing degree of short- and then long-range Fe-Ti cation order with increasing ilmenite content, leading to a loss of symmetry from $R3c$ to $R3$ (Harrison and Redfern 2001). Quenched samples with intermediate compositions are infamous for their ability to acquire self-reversed thermoremanent magnetization, linked to chemical and ordering heterogeneities at the nanoscale (Fabian et al. 2011; Robinson et al. 2011, 2012, 2014).

Here we investigate the spin-orientation in synthetic samples of solid solution $\left[\text{Fe}_x\text{TiO}_3 - (1-x)\text{Fe}_2\text{O}_3\right]$ with compositions $x \leq 0.40$ and in a natural hematite sample ($x \approx 0$). We use Rietveld refinement of neutron powder diffraction data, taken at temperatures from room temperature to $T_{NM}$, to determine the average out-of-plane spin angle $\alpha$. We use Mössbauer spectroscopy to investigate the distribution of $\alpha$ in each sample. We find that the spins in all hematite-ilmenite samples have significant out-of-plane components in the range from $\alpha = 11(2)^\circ$ to $\alpha = 22.7(5)^\circ$. All given uncertainties are the estimated standard deviations ($\sigma$). Even the natural hematite sample has a significant out-of-plane spin-angle of $\alpha = 18.1(6)^\circ$. Our results show that $\alpha$ does not vary systematically with Ti-content.

**GROUP THEORY AND CRYSTALLOGRAPHY**

Magnetic ordering in hematite at temperatures $T_{NM} < T < T_N$ can be described by a two-dimensional primary magnetic order parameter ($Q_1$, $Q_2$) corresponding to the active irreducible representation ($m\Gamma$) of the parent space group ($R3c$). Different combinations of $Q_1$ and $Q_2$ lead to three possible magnetic space groups: $C2/c$ for $Q_1 \neq 0, Q_2 = 0$; $C2'/c'$ for $Q_1 = 0, Q_2 \neq 0$; and $P\Gamma$ for $Q_1 \neq 0, Q_2 \neq 0$. The magnetic structure observed by experiment is the $C2/c$ structure with $Q_1 \neq 0$ and $Q_2 = 0$. This structure is a canted antiferromagnet with parallel alignment of the spins within each (001) layer and nearly antiparallel alignment of spins in neighboring (001) layers (indices refer to the parent hexagonal unit cell). The primary order parameter $Q_1$ places spins parallel to the (001) basal plane and nearly perpendicular to the diad (i.e., nearly perpendicular to an $a$ crystallographic axis of the parent hexagonal unit cell). A degree of canting within the basal plane is permitted by the $C2/c$ symmetry, such that spins are rotated by $\approx -0.065^\circ$ to create a weak ferromagnetic (WF) moment along the diad (Morrish 1994). A secondary irreducible representation ($m\Gamma_1$) is permitted within $C2/c$. By itself, $m\Gamma_1$ would give a strictly antiferromagnetic alignment of spins normal to the (001) basal plane (as observed for $T < T_{NM}$). In combination with $m\Gamma_1$, the antiferromagnetic spins are rotated about the diad so that they lie at an angle, $\alpha$, to the basal plane, as has been observed in recent experiments (Frandsen et al. 2011; Harrison et al. 2010; Klotz et al. 2013; Parise et al. 2006). Note that even when $\alpha \neq 0$, the canted WF moment still lies parallel to the diad and therefore in the basal plane. If we assign a secondary order parameter $Q_2$, to $m\Gamma_1$, then $Q_1$ and $Q_2$ are proportional to the in-plane and out-of-plane spin components, respectively. Given the symmetry relationship between $Q_1$ and $Q_2$, it is permitted for these order parameters to couple bilinearly in the expansion of Gibb’s free energy. In this case we would expect that $Q_2$ would vary linearly with $Q_1$. This proposition can be tested directly by Rietveld refinement of high-temperature powder neutron diffraction data.

Here we follow the crystallographic conventions and refinement procedures described in detail by Harrison et al. (2010). For technical reasons, refinements were performed in a non-standard $A2/a$ setting of the monoclinic $C2/c$ phase described above. For ease of comparison with the high-temperature hexagonal phase the unit-cell parameters are presented here using a monoclinic pseudocell with volume equal to that of the high-temperature hexagonal cell: $a_m = (1/\sqrt{3})[210]_{\text{hex}}, b_m = [010]_{\text{hex}}, c_m = [001]_{\text{hex}}, \beta_m = 90^\circ$, where $[abc]_{\text{hex}}$ refers to the axes of the hexagonal cell. Similarly the refined magnetic moments $M_r$ and $M_r$ in the monoclinic cell are converted to components parallel ($M_r$) and perpendicular ($M_r$) to the basal plane. Figure 1 shows a representation of the chemical and magnetic structure of hematite with a spin rotation of $\alpha = 20^\circ$ away from the basal plane.

**METHODS**

**Samples**

Synthetic powder samples of $\left[\text{Fe}_x\text{TiO}_3 - (1-x)\text{Fe}_2\text{O}_3\right]$ with nominal compositions of $x = 0, 0.13, 0.20, 0.35$, and 0.40 were prepared by heating in sealed silica tubes followed by rapid cooling of mixtures of FeO and TiO$_2$ as described in Burton et al. (2008). A natural sample of hematite was obtained from the Sedgwick Museum of Earth Sciences, Cambridge, U.K. The hematite sample was crushed and ground into a fine powder. We refer to these samples as ilm13, ilm20, ilm35, ilm40, and hem.

**Neutron powder diffraction**

We performed neutron powder diffraction experiments at the OSIRIS instrument at the ISIS spallation neutron source, Oxfordshire, U.K., and at the D20 instrument at the reactor source at Institut Laue Langevin, Grenoble, France. In both experiments the samples were mounted inside an evacuated furnace providing a temperature range from room temperature to 1373 K (1100 °C). The Néel temperatures of the Ti-containing samples are expected to be lower (Besser et al. 1967) than the 955 K of pure hematite, and the available temperature range should
The magnetite impurity was added to the refinements for ilm20 and hem where the iron belonging to the structure of magnetite. Because of this magnetite impurity, a small amount of the sample was ground to a fine powder in an agate mortar and 30 mg of it was mixed with boron nitride and placed in a plastic holder. The Mössbauer spectra were obtained at Department of Physics, Technical University of Denmark, Kgs. Lyngby, Denmark using a constant acceleration Mössbauer spectrometer with a source of 57Co in rhodium. The magnetically split spectra were fitted to sextets of Lorentzians with appropriate peak shape functions for time-of-flight (OSIRIS) and constant wavelength (D20) data. In all refinements the cations were assumed to be fully disordered. The validity of the assumption of cation disorder can be verified by the absence of reflections from the \( R \) structure of FeTiO\(_3\) in the diffraction data. Long-range cation order would give rise to a nonzero intensity at the position of the magnetic hematite peaks above \( T_m \) that was not observed.

Neutron powder diffraction

The room-temperature diffraction patterns from the samples hem and ilm13 are shown in Figure 2. The positions of the (101) and (003) magnetic peaks and the most intense of the peaks from the magnetite impurity are indicated. The much better resolution in the data from OSIRIS as compared to D20 is clearly seen by comparing the width of the peaks in the two diffraction patterns. The refined model represents the data reasonably well, and the disagreements between model and measurement originate primarily from misfits of the peak profiles, which is a problem that can never be completely eliminated. There was some variation in background between samples and in some cases the background was somewhat nonuniform and not that well modeled. However, it is important to note that for all samples the background was well modeled in the \( d \)-spacing range close to the two main magnetic peaks (101) and (003), which is the area of the diffraction pattern that is most significant for obtaining information about the magnetic structure of the sample, in particular the out-of-plane spin-angle \( \alpha \).
A three-dimensional representation (d-spacing, temperature, intensity) of the complete set of measurements of the ilm13 sample from room temperature (295 K) to above 1240 K is displayed in Figure 3. The magnetic phase transition is identified by the disappearance of the two main magnetic peaks [(101) at \(d = 4.2\) Å and (003) at \(d = 4.6\) Å] at a temperature of \(~800\) K. The weak line at \(d = 4.9\) Å can be identified as a magnetic reflection from magnetite and it completely disappears at the Curie temperature of magnetite (\(\approx 850\) K). Lines pertaining to the magnetite crystal structure can also be identified (e.g., at \(d = 3.0\) Å) and these achieve a significant gain in intensity at temperatures around 1100 K, indicating that in addition to a small initial impurity of magnetite part of the sample is being reduced to magnetite during heating. When magnetite is produced by reduction of the sample the composition of the remaining hematite-ilmenite becomes more Ti-rich resulting in an enlargement of the unit cell seen as a shift of the diffraction lines to lower scattering angles (larger d-spacings).

The refined unit-cell volumes and the determined compositions are given in Table 1 together with the room-temperature values of the magnetic moments \(M_p, M_s, M_{tot}\), and \(\alpha\). The compositions deviate in some cases from the nominal values. In all cases \(M_p\) is significantly larger than \(M_s\), as expected for hematite between \(T_N\) and \(T_N\). However, there is a significant out-of-plane moment for all samples. The size of \(\alpha\) varies from 11(2)\(^\circ\) for ilm13 to 22.7(5)\(^\circ\) for ilm35 and it does not depend systematically on Ti-content. For the hem sample \(\alpha = 18.1(6)\(^\circ\)\).

In Figure 4, we show \(M_p, M_s\), and the total moment \(M_{tot}\) as function of temperature for the samples hem, ilm13, ilm20, and ilm35. As the temperature increases toward \(T_N\), both components of the moment decrease at the same rate until close to \(T_N\). For the ilm13 and ilm20 samples \(M_s\) becomes zero before \(M_p\), indicating a reorientation of the spins close to \(T_N\). However, the relative uncertainties on the refined moments (especially \(M_s\)) become very large close to \(T_N\) because the magnetic peaks are weak. The Néel temperatures determined from fitting the \(M_{tot}\) data to a power law

\[
M = M_0 \left(\frac{T_N - T}{T_N}\right)^\beta
\]

are given in Table 2.\(^1\) The fits represent the data well until very close to \(T_N\) where the order parameter is expected to deviate from the power law because of critical behavior of the system. Because of the critical behavior a finite intensity remains in the magnetic peaks at temperatures a few K above the fitted \(T_N\). The critical exponent \(\beta\) determined from the fit to \(M_{tot}\) was in the range 0.3–0.4 for all samples, which is in the expected range for a phase transition in a three-dimensional system.

In Figure 5 (left) the total moment is displayed as function of the reduced temperature \((T/T_N)\) for all samples. In Figure 5 (right) the out-of-plane tilt angle \(\alpha\) is displayed as a function of the reduced temperature. There is no systematic dependence of \(\alpha\) on the Ti-concentration. The spin reorientation toward the basal plane with increasing temperature is systematic for all Ti-containing samples with the exception of two points very close to \(T_N\) \((0.995 \ T_N < T < T_N)\) for the ilm35 sample. The figure also illustrates that the uncertainty on \(\alpha\) becomes large near \(T_N\). For sample hem there is a significant change in \(\alpha\) between the first and the second point, which are measured at room temperature and at 673 K \((400\) °C), respectively.

To illustrate the relationship between the primary order parameter \(Q_1\) and the secondary order parameter \(Q_2, M_s\) is plotted against \(M_p\) in Figure 6. For the hem sample the linear fit represents the data well. For the ilm13 and ilm20 samples, \(M_s\) turns to zero before \(M_p\) ending the linear relationship between \(Q_1\) and \(Q_2\) shortly before \(T_N\). For ilm35 the trend is the same as for the other Ti-containing samples if the three anomalous points closest to \(T_N\) are disregarded from the fit.

In the experiment at D20 on sample ilm13 we obtained sufficient data above and below \(T_N\) to follow the transformation of the

\(^1\)Deposit item AM-17-65792, CIF and Tables. Deposit items are free to all readers and found on the MSA web site, via the specific issue’s Table of Contents (go to http://www.minsocam.org/MSA/AmMin/TOC/2017/June2017_data/June2017_data.html).
monoclinic unit cell to the high-symmetry hexagonal cell occurring simultaneously with the magnetic phase transition. The change in the unit-cell parameters as a function of temperature can be seen in Figure 7. The unit-cell parameters for the low-temperature monoclinic phase, $a_m$, $b_m$, and $c_m$, are given in the reference frame of the monoclinic pseudo-cell and can thus be compared directly to the unit-cell parameters of the high-temperature hexagonal cell, $a_h$ and $c_h$. As the temperature increases $a_m$, $b_m$ converge toward the value of $a_h$ at $T_N$. The slope of the curve does not change significantly at the phase transition, indicating

**Figure 4.** Refined in-plane ($M_{\parallel}$), out-of-plane ($M_{\perp}$), and total ($M_{\text{tot}}$) moment as a function of temperature for samples hem, ilm13, ilm20, and ilm35. The solid lines are power law fits. The error bars represent one standard deviation ($\sigma$).

**Figure 5.** (a) Total magnetic moment as function of the reduced temperature. The solid lines are the same fits as in Figure 4. (b) Out-of-plane tilt-angle $\alpha$ as function of the reduced temperature. The ilm13 data has been binned so that each data point represents a 17 K temperature interval. The error bars represent one standard deviation ($\sigma$).
that the strain associated with the phase transition is small. The same is true for the transformation of $c_m$ to $c_h$. At high temperatures (above 1000 K) the slope of the $a_h$ and $c_h$ curves change. This change happens in the same temperature range where the intensity of the magnetite lines start to increase (see Fig. 2) and is probably associated with the reduction of some of the ilm13 to magnetite and the resulting change in hematite-ilmenite composition. The transformation of the crystallographic $b$-angle from monoclinic $b_m \neq 90^\circ$ to hexagonal $b_h = 90^\circ$ can be followed in Figure 8. At room-temperature $b_m$ is $90.039(7)^\circ$. As the temperature increases $b_m$ decreases toward $90^\circ$.

Mössbauer spectroscopy

The Mössbauer spectra of the hem, ilm20, ilm35, and ilm40 samples are shown in Figure 9. Also shown are model fits to the data.

**Hematite sample.** The spectrum of the hem sample consists of a well-defined sextet as expected for pure hematite, as well as a small component (10% of the spectral area) that can be ascribed to the magnetite impurity, also observed in the neutron data. The room-temperature Mössbauer parameters of the hematite sextet are: Hyperfine field $B_{hf} = 51.50(2)$ T, isomer shift $\Delta IS = 0.370(4)$ mm/s and quadrupole shift $\varepsilon = -0.094(4)$ mm/s independent of whether the magnetite impurity is included in the fit. Here the uncertainties on the hyperfine parameters are estimated from the uncertainties on the calibration with respect to $\alpha$-Fe. The Lorentzian linewidths are 0.311, 0.288, and 0.249 mm/s for lines 1 and 6, 2 and 5, and 3 and 4, respectively. Because the linewidths are close to the instrumental line width of 0.260 mm/s fitting with a more complicated model with a distribution of hyperfine parameters is not justified.

The quadrupole shift ($\varepsilon$) in the Mössbauer spectrum of hematite is sensitive to $\alpha$, and the fact that the spectrum is well fitted with one symmetric sextet rather than two sextets with different $\varepsilon$ indicates that the directions of the Fe$^{3+}$ spins in the sample are grouped around one value of $\alpha$, rather than being distributed into distinct domains with different spin-orientations. In particular, we can rule out a scenario with significant proportions of the spins being in the basal plane and along the c-axis, respectively. In principle the average value of $\alpha$ can be found from the quadrupole shift of the spectrum from (Morrish 1994)

$$\varepsilon = \varepsilon_0 [3\cos^2(90^\circ - \alpha) - 1]/2,$$

where $\varepsilon_0$ depends on the quadrupole moment of the iron nucleus in the nuclear spin $= 3/2$ state, and on the electric field gradient...
along the c-axis. From the literature (Artman et al. 1968; Tobler et al. 1981) $e_0$ can be estimated to $e_0 = 0.215(5)$ mm/s and $\alpha$ can be determined from the measured quadrupole shift $-0.094(4)$ mm/s to be $\alpha = 12(2)^\circ$.

Mössbauer spectra of the hem sample were also collected at 80 and 20 K (not shown). These spectra taken below $T_M$ have quadrupole shifts of $0.190(4)$ and $0.189(4)$ mm/s, respectively, corresponding to an imperfect Morin transition with the spins making an angle of $16(2)^\circ$ with [001] even at 20 K.

**Ti-containing samples.** The room-temperature Mössbauer spectra of the samples ilm20, ilm35, and ilm40 have much broader lines than the hem spectrum. Each spectrum is fitted with four sextets, three with isomer shifts close to 0.4 mm/s, corresponding to Fe$^{3+}$, and one with isomer shift closer to 0.6 mm/s, corresponding to Fe$^{2+}$. The hyperfine fields of the Fe$^{3+}$ sextets are in the range 47–50 T and thus a bit lower than expected for pure hematite at room temperature (probably due to Ti$^{4+}$ substitution in the hematite structure). The hyperfine field of the Fe$^{2+}$ sextet is 37–44 T in all samples. The quadrupole shifts of the Fe$^{3+}$ sextets are in the range between $-0.110$ and $-0.076$ mm/s. The ilm35 spectrum contains a central doublet with isomer shift $0.377(4)$ mm/s and quadrupole splitting $0.663(4)$ mm/s, suggesting that it could originate from a paramagnetic Fe$^{3+}$ phase. The quadrupole shifts in the Ti-containing samples cannot be used to calculate $\alpha$ from Equation 2 because $e_0$ depends on the electric field gradient along the hexagonal c-axis, which is expected to change when Ti is substituted for Fe in the hematite structure. While it is not possible to calculate $\alpha$ the fact that the quadrupole shifts of all the Fe$^{3+}$ sextets are close to $-0.1$ mm/s suggests that, like in the hem sample, the spin orientations are grouped around some average angle, rather than two distinct populations of in-plane and out-of-plane spins. The room-temperature hyperfine parameters obtained from the sextet fits can be seen in Table 3.

**DISCUSSION**

The Rietveld refinement of the neutron powder diffraction data tells us that there is a significant out-of-plane component of the hematite spins in all hematite-ilmenite samples and even in the natural hematite sample. While there is a variation in the measured $\alpha$ between samples it does not seem to depend systematically on Ti-content. There is no pronounced temperature dependence of $\alpha$, however, for the Ti-containing samples the spins tend to reorient toward the basal plane close to $T_N$, with the exception of a few points very close to $T_N$ for ilm35. It should be noted that very close to the phase transition the refinement of the magnetic structure is susceptible to systematic errors and the parameters extracted from the refinements should not be given too much weight.

From the quadrupole shift in the room-temperature Mössbauer spectrum of the hem sample the spin-angle can be determined to $\alpha = 12(2)^\circ$, which is not consistent with the neutron result [$\alpha = 18.1(6)^\circ$], although it does confirm that there is a significant out-of-plane component of the spin in the natural hematite sample. The value of $\alpha$ determined from the quadrupole shift in the Mössbauer spectrum may not be taken as an accurate number because the hyperfine parameters (most importantly $e_0$) may depend delicately on crystallographic strain and other sample conditions.
dependent factors such as impurities in the structure. Because of this uncertainty in the determination of $\alpha$ from Mössbauer spectroscopy we regard the value obtained with neutron diffraction as more accurate. From the Mössbauer spectrum taken at 20 K we find the spins to be rotated by 16(2)$^\circ$ from the c-axis. As soon as the spins are observed to be rotated away from the c-axis, a canted moment becomes symmetry-allowed and could contribute to the defect moment sometimes observed for hematite below $T_{M}$. The spectrum also reveals that the spin orientation is not groupd in distinct populations, e.g., one with $\alpha = 0^\circ$ and one with $\alpha = 90^\circ$ corresponding to the spin orientation usually assumed above and below the Morin transition, but are grouped around some average value. The Mössbauer spectra of ilm20, ilm35, and ilm40 suggest that this is the case also in Ti-substituted hematite.

We followed the structural phase transition at $T_{C}$ for a sample (ilm13) with $x = 0.176(12)$ and observed a gradual transition from the low symmetry monoclinic cell to the high symmetry hexagonal cell. Within the resolution of D20 there is no strain associated with the phase transition.

The 30$^\circ$ out-of-plane spin-angle found by Harrison et al. (2010) was believed to be linked to the exsolution microstructure in the natural sample, and it is surprising that we find appreciable out-of-plane spin-angles in our samples that should not exhibit pronounced exsolution microstructure. While the neutron diffraction data exclude a long-range ordering of the Ti-cations it is not possible to determine whether some short-range cation ordering exists. Frandsen et al. (2010) suggested that small clusters of ordered hematite exist inside an ilmenite matrix for samples with $x > 0.5$. They conclude that the size of the hematite clusters is only on the order of 1–2 nm. Spins around the interfaces of such clusters could be rotated due to exchange interaction between Fe$^{2+}$ and Fe$^{3+}$ species. This would, however, not explain why we observe an out-of-plane spin orientation in the natural hematite sample. The out-of-plane spin angle of 30$^\circ$ observed by Harrison et al. (2010) is significantly larger than the spin angles we find in our samples. This suggests that while some out-of-plane spin orientation may be intrinsic to the hematite magnetic structure exsolution could enhance the effect.

It is well known that the magnetic structure of hematite does not prohibit an out-of-plane spin-component, however, it is generally assumed to be much smaller than the 18.1(6)$^\circ$ found in this work (Dzyaloshinsky 1958). The spin orientation in hematite above and below the Morin transition is determined by a fine balance between competing anistropies that may depend on factors such as impurities, strain, and applied pressure (Morrish et al. 1963; Artman et al. 1965; Besser et al. 1967; Klotz et al. 2013). In numerous neutron studies it has been observed that there is a finite intensity in the magnetic (003) reflection at temperatures far below the Morin transition (Morrish et al. 1963; Krén et al. 1974; Sváb and Krén 1979; Morrish 1994), proving that the spins are not perfectly aligned along the [001] axis below $T_{M}$. Our results show that the spin alignment above the Morin transition can also be imperfect, with the spins rotated away from the (001) plane by a significant angle.

All of the samples investigated in this work have impurities of magnetite in amounts that give rise to visible diffraction peaks but that except for the hem sample are not detectible by Mössbauer spectroscopy due to overlapping lines. While the magnetite impurities are likely to be important for the macroscopic magnetic properties of the sample (the bulk magnetization) inclusions of magnetite in the structure may not be important for the spin structure in the hematite-ilmenite phase. Other impurities or defects may, however, be important for the spin orientation and could explain the variation in spin angle between samples.

It may seem surprising that we find a significant out-of-plane spin component in hematite but it is consistent with the experiments of Parise et al. (2006) and Klotz et al. (2013). While our sample is a natural sample, which contains impurities that could affect its magnetic structure, the samples investigated by both Parise et al. (2006) and Klotz et al. (2013) were high-purity synthetic samples and impurities are thus unlikely to be the sole cause of the observations of out-of-plane spin orientation in hematite. Klotz et al. (2013) ascribe the spin orientation as an artifact of the Rietveld method’s insensibility to changes in $\alpha$. It is true that the quality of the Rietveld-fit, as quantified by the $\chi^2$-value or any other measure of the sum of residuals, does not depend strongly on the spin orientation but it is clear that the model with $\alpha = 20.0^\circ$ (close to the refined value $\alpha = 18.1^\circ$) represents the data much better than $\alpha = 0^\circ$, as can be seen in Figure 10, where the (101) and (003) magnetic peaks at room temperature are plotted together with models with different values of $\alpha$. However, the insensitivity of the fit to variations in $\alpha$, when $\alpha$ is close to 0$^\circ$ has the consequence that the experiment becomes very susceptible to systematic errors. We have not identified any systematic errors in this experiment that could give rise to the observed out-of-plane spin orientation.

The insensitivity to variations in the spin angle is not unique to the Rietveld method, but is a consequence of the small change in 1–sin$^2(\alpha)$ (which is essentially the measured quantity) when $\alpha$ is small. To completely eliminate this problem one would have to perform a polarized neutron diffraction
experiment on a single crystal of hematite.

We have studied the spin orientation in synthetic samples of hematite-ilmenite as well as a natural sample of hematite with neutron powder diffraction and Mössbauer spectroscopy. We find that the spin has a significant component out of the basal plane in all samples. The nonzero tilt-angle of the antiferromagnetic sublattices is consistent with other studies but we cannot at present give a detailed explanation of the origin of this deviation from the usually assumed magnetic structure with the spins confined to the basal plane. The tilt angles in our samples are in the range $11.0(9)-22.7(5)^\circ$, and seem not to depend systematically on Ti-content. Furthermore, the out-of-plane angle does not require an exsolution structure as seen in natural samples.

**Implications**

The implications of the present results can best be appreciated by reference to the theoretical paper on hematite magnetization by Dzyaloshinsky (1958), which showed both the individual layer spin orientations and the spin-canted ferromagnetic moments parallel to the (001) crystallographic plane. This understanding persisted for the next half century and was enshrined in the major book Canted Antiferromagnetism by Morrrish (1994). The first warnings that this might be incorrect came from Monte Carlo modeling of low-temperature giant magnetic exchange bias in titanohematite with nanoscale ilmenite exsolution lamellae (Harrison et al. 2007), and evidence that such exchange bias could be produced when similar natural samples were cooled in zero field to low temperature before the hysteresis experiment (Fabian et al. 2008). These studies implied that an unexpected out-of-plane component of spin should exist in the hematite, even at room temperature and above. This was later observed directly in a similar exsolved sample (Harrison et al. 2010), but was then attributed to a localized effect of Fe$^{2+}$ cations at interfaces. The present results indicate instead, for the first time, that the out-of-plane spin component is an intrinsic feature of hematite itself, with an origin not yet fully understood, but consistent with group theory. Probably earlier workers mistook the crystallographic position of the spin-canted ferromagnetic moment as also the position of the individual layer spin components. The present work thus establishes a major shift in understanding of one of the two mineral systems responsible for rock magnetism.

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