Spin-lattice coupling mediated multiferroicity in (ND₄)₂FeCl₅·D₂O

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We report a neutron diffraction study of the multiferroic mechanism in (ND₄)₂FeCl₅·D₂O, a molecular compound that exhibits magnetically induced ferroelectricity. This material exhibits two successive magnetic transitions on cooling: a long-range order transition to an incommensurate (IC) collinear sinusoidal spin state at $T_N$=7.3 K, followed by a second transition to an IC cycloidal spin state at $T_{FE}$=6.8 K, the later of which is accompanied by spontaneous ferroelectric polarization. The cycloid structure is strongly distorted by spin-lattice coupling as evidenced by the observations of both odd and even higher-order harmonics associated with the cycloid wave vector, and a weak commensurate phase that coexists with the IC phase. The appearance of the 2nd-order harmonic coincides with the onset of the electric polarization, thereby providing unambiguous evidence that the induced electric polarization is mediated by the spin-lattice interaction. Our results for this system, in which the orbital angular momentum is expected to be quenched, are remarkably similar to those of the prototypical TbMnO₃, in which the magnetoelectric effect is attributed to spin-orbit coupling.

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“Improper multiferroics” (also referred as type-II magnetic multiferroics) are a unique group of materials that exhibit direct coupling between magnetism and electric polarization. In these magnetically induced multiferroics, the magnetoelectric (ME) effect is strong and the onset of ferroelectricity arises directly from magnetic order that breaks spatial inversion symmetry. Due to the strong ME effect, there has been enormous interest in these materials motivated by their potential applications in novel multifunctional devices. However, natural single-phase magnetically driven multiferroics are rare, only a few transition-metal oxides such as TbMnO₃, MnWO₄, Ni₃V₂O₈, CuO₂, LiCuVO₄, and CaCoMnO₃ are currently known to exhibit such effect. It is thus of great interest to discover and investigate new materials that will shed light on the underlying multiferroic mechanism. Recent efforts to search for new multiferroics have been extended to molecular compounds and metal-organic framework materials (MOFs). In particular, several ionic salts containing NH₄ have been reported to exhibit ferroelectricity and the incorporation of NH₄ has been used as a strategy to search for new multiferroics. In this paper, we report a neutron diffraction study of the multiferroic mechanism in (NH₄)₂FeCl₅·H₂O. Using both polarized and unpolarized neutrons, we show that the induced ferroelectricity in (NH₄)₂FeCl₅·H₂O is mediated via spin-lattice coupling mechanism strikingly similar to TbMnO₃.

(NH₄)₂FeCl₅·H₂O belongs to the erythrosiderite-type compounds $A_2[FeX_2\cdotH_2O]$, where $A$ is an alkali metal or ammonium, and $X$ is a halide ion. It crystallizes in an orthorhombic structure at room temperature (space group $Pnma$) with a crystal structure consisting of distorted [FeCl₅·H₂O]²⁻ octahedra linked by a network of hydrogen bonds as illustrated in Fig. 1(a). The orbital angular momentum of the magnetic Fe³⁺ (3d⁵, high spin state) ion is expected to be completely quenched. The

![FIG. 1. (Color online) (a) Crystal structure of (NH₄)₂FeCl₅·H₂O. (b) IC collinear sinusoidal spin structure at 7 K in the paraelectric phase; (c) IC cycloidal-spiral spin structure at 4 K in the ferroelectric phase, as viewed along the b-axis (three unit cells along c-axis are plotted). Only magnetic Fe ions are shown in (b) and (c) for the purpose of clarity.](image-url)
magnetic interactions in these materials are mediated via multiple superexchange pathways, such as Fe-Cl···Cl-Fe, Fe-O···Cl-Fe, and Fe-O-H···Cl-Fe involving hydrogen bonds, suggesting the presence of magnetic frustration in the system\(^{22}\). (NH\(_4\))\(_2\)FeCl\(_5\)-H\(_2\)O\(^{24}\) is the only compound that exhibits spontaneous electric polarization in the A\(_2\)[Fe\(_x\)X\(_2\)-H\(_2\)O] series. In sharp contrast to other isostructural counterparts, such as (K, Rb)\(_2\)FeCl\(_5\)-H\(_2\)O which undergo a single magnetic transition adopting a collinear aniferromagnetic (AFM) spin structure with moments along the \(a\)-axis\(^{22}\) and have no spontaneous electric polarization, (NH\(_4\))\(_2\)FeCl\(_5\)-H\(_2\)O exhibits two magnetic transitions at \(T_N\sim7.3\) K and \(T_{FE}\sim6.8\) K, respectively. A disorder-order transition also occurs at \(T_s\sim79\) K associated with the motion of the NH\(_4\) group\(^{25}\). Unlike other NH\(_4\) contained materials investigated so far\(^{12,23}\) where ferroelectricity is induced via the disorder-order transition at high temperature decoupled with the low temperature magnetic order, which is characteristic of the so-called type-I “proper multiferroics”\(^{1,2}\). (NH\(_4\))\(_2\)FeCl\(_5\)-H\(_2\)O remains paraelectric below 79 K and spontaneous electric polarization only appears below \(T_{FE}\). Consequently, it is crucial to know the magnetic structure and microscopic interactions to unveil the underlying mechanism responsible for the multiferroic properties in this compound. However, the determination of the full magnetic structure has been hampered due to the large amount of hydrogen atoms (40 H atoms per unit cell) in (NH\(_4\))\(_2\)FeCl\(_5\)-H\(_2\)O.

We have grown deuterated (ND\(_4\))\(_2\)FeCl\(_5\)-D\(_2\)O single crystals suitable for neutron scattering by solution method using starting materials (DCl, FeCl\(_3\), and ND\(_4\)Cl) similar to that reported in Ref. \(^{24}\). The saturated solution was sealed and kept at 38°C using a sample environment chamber to allow slow evaporation. Large deuterated crystals were obtained and later characterized by magnetic susceptibility and specific heat measurements. No significant deuteration-induced effects were observed since specific heat data indicates no transition temperature changes compared to the hydrogenated sample\(^{22,24}\). Elastic neutron scattering experiments were carried out using the HB1, HB1A triple-axis spectrometer (TAS), and HB3A Four-Circle Diffractometer located at the High Flux Isotope Reactor (HFIR), and the Elastic Diffuse Scattering Spectrometer (CORELLI) at the Spallation Neutron Source (SNS) at Oak Ridge National Laboratory (ORNL). The magnetic structures at 4 K and 7 K were determined by refining the data collected at HB3A. HB1A and CORELLI were used to investigate the temperature and \(q\) dependence of the observed higher-order harmonics associated with the cycloid order. The nature of the higher-order harmonics were further clarified by polarized neutron experiments using the HB1 polarized TAS. The polarization of the incident beam and the polarization analysis of the scattered beam were produced by Heusler crystals, and a Mezei spin flipper was used for reversing the neutron polarization vector.

Neutron data show that (ND\(_4\))\(_2\)FeCl\(_5\)-D\(_2\)O undergoes an incommensurate (IC) AFM long range order (LRO) transition at \(T_N\sim7.3\) K. The primary magnetic satellite peak can be indexed with a propagation vector \(k_1=(0\ 0\ k_{\parallel})\), \(k_{\parallel}\sim0.23\) at 1.5 K, consistent with the recent study by Rodríguez-Velamazín et al.\(^{25}\). To determine the magnetic structures associated with both the paraelectric phase between \(T_{FE}<T<T_N\) and the ferroelectric phase below \(T_{FE}=6.8\) K, sets of nuclear and magnetic reflections were collected at 7 K and 4 K at HB3A (wave length 1.546 Å). FullProf refinement of 4 K data confirms an IC cycloidal spiral spin structure with moments mainly confined in the \(ac\)-plane (moment size \(\sim4.08\ \mu_B\)) consistent with Ref. \(^{25}\). The magnetic structure at 7 K is found to be an IC collinear sinusoidal spin state with moments along the \(a\)-axis, with a moment size of \(\sim2.17\ \mu_B\). The schematic magnetic structures at 7 K and 4 K are illustrated in Fig. 1 (b) and (c), respectively. This indicates a magnetic structure change from collinear sinusoidal to cycloidal spiral at \(T_{FE}=6.8\) K that suggests an
The cycloid structure is strongly distorted via spin-lattice coupling. Furthermore, weak commensurate peaks with a propagation vector $k_2=(0\ 0\ k_{z2})$ ($k_{z2}=0.25$) are also observed at 1.5 K. As depicted in Fig. 2(b), both (2 0 0.23) IC and (2 0 0.25) commensurate peaks are observed in the L-scan along (2 0 L). This suggests the coexistence of commensurate and incommensurate phases at 1.5 K. The inset illustrates the temperature dependence of (2 0 0.23) and (2 0 0.25). Both reflections show increased intensity with decreasing temperature suggesting they are magnetic in origin. The observed spin texture (both even and odd-order harmonics associated with $k_1$ and coexistence of the $k_2$ commensurate phase) indicates the cycloidal spiral spin structure is strongly distorted since no higher-order harmonics should be observed for an ideal cycloid spiral. Furthermore, as discussed in Ref. 26, odd-order harmonics and even order harmonics are expected to be magnetic and nuclear in origin, respectively. Therefore, the observation of the 2nd-order harmonic provides direct evidence of a lattice modulation associated with the cycloid order with wave vector $k_1$.

The magnetic transitions occur within a narrow temperature range at $T_N\approx7.3$ K and $T_{FE}\approx6.8$ K. To clarify the nature and exact transition temperature of the primary satellite and higher-order harmonic peaks, we measure both (0 0 0.77) and (0 0 2.46) (2nd-order harmonic) as a function of temperature by performing $L$ scans. The obtained order parameters and peak center (indexed using $k_{z1}$) versus T are plotted in Fig. 3 in comparison with the specific heat data (Fig. 3(a)). The integrated intensity was obtained by fitting the scan at each temperature to a Gaussian function with a constant background. The order parameter of (0 0 0.77) (Fig. 3(b)) shows a transition at 7.3 K corresponding to $T_N$ and depicts no clear anomaly at $T_{FE}$. The order parameter of (0 0 2.46) (Fig. 3(c)) reveals that the appearance of the 2nd-order harmonic coincides with the onset of electric polarization. Fitting the order parameter to a power-law $I(T)=I_0[(T_{SE}-T)/T_N]^{\beta}$ yield $T_{FE}\approx7.34\pm0.02$ K and $\beta=0.194\pm0.015$ for (0 0 0.77), and $T_{FE}\approx6.68\pm0.03$ K and $\beta=0.245\pm0.002$ for (0 0 2.46), respectively. The fitting results are plotted in Fig. 3(b) and (c) as red solid curves. The fitting was performed over the temperature ranges 2 K $< T < T_N$ and 2 K $< T < T_{FE}$, respectively. These critical exponent $\beta$ values are much smaller than the theoretical value of 0.36 predicted for a three-dimensional Heisenberg antiferromagnet (de Jongh and Miedema 1974). This suggests a lower magnetic dimensionality of (ND$_4$)$_2$FeCl$_5$·D$_2$O as proposed in Ref. 21.

Temperature dependence of the IC propagation vector $k_{z1}$, determined by fitting the L-scans of (0 0 0.77) and (0 0 2.46), is plotted in Fig. 3(d). Good agreement is obtained between 2 K $< T < 6$ K, the discrepancy above 6 K can be attributed to the significant broadening of both reflections near $T_{FE}$ and $T_N$. The wave vector (0 0 $k_{z1}$) continues to vary with decreasing temperature through-
out the ferroelectric phase and it shows no evidence of an incommensurate-commensurate (IC-C) lock-in transition down to 1.5 K.

To clarify the origin of the higher-order harmonics and \( k_2 \)-type commensurate peaks, a polarized neutron diffraction experiment was carried out using the HB1 polarized triple-axis spectrometer. Fig. 4(a) plots the polarized data analyzed using the method discussed in Ref. 27–29. The magnetic origin of the commensurate peak and the odd-order harmonics associated with the cycloidal order is verified by the stronger scattering observed in the spin-flip (SF, -+) channel in Fig. 4(a). The remaining scattering in the non-spin-flip (NSF, +++) channel can be attributed to finite instrumental flipping ratio which is estimated to be \( \sim 1/10 \) by comparing the integrated intensity of SF and NSF data of the (0 0 2.46) 2nd-order harmonic peak. Comparison of 1.5 K and 10 K data of (0 0 2.46) from (c) SF channel and (d) NSF channel, respectively.

FIG. 4. (Color online) Polarized neutron data measured in the horizontal field configuration, \( P_0 \parallel Q \). (a) L-scan along [0 1 L] measured at 1.5 K comparing the spin-flip (SF, -+) and non-spin-flip (NSF, +++) scatterings with the intensity plotted on a logarithmic scale. (b) Comparison of the 1.5 K SF and NSF data suggests the hybrid nature of the (0 0 2.46) 2nd-order harmonic peak. Comparison of 1.5 K and 10 K data of (0 0 2.46) from (c) SF channel and (d) NSF channel, respectively.

The spin texture (in the form of higher-order harmonics) observed in (\( \text{ND}_4 \))\(_2\)\(\text{FeCl}_5\)\(_2\)\(\text{D}_2\)\(\text{O}\) is strikingly similar to that of Tb\(\text{MnO}_3\), a well studied type-II multiferroic system\(5–9\). Three transitions are observed in Tb\(\text{MnO}_3\) at 41 K, 27 K, and 7 K. The first one at 41 K is associated with the IC magnetic LRO transition of \( \text{Mn}^{3+} \), and the third one at 7 K is due to the ordering of \( \text{Tb}^{3+} \) moments, typical for rare-earth compounds. The second transition at 27 K is particularly interesting and is accompanied by a dielectric anomaly and electric polarization. It has been clarified by a neutron diffraction study\(5–2\) that this 27 K transition is associated with a magnetic structure change from an IC sinusoidally modulated collinear magnetic structure to an IC noncollinear spiral. Strong odd-order harmonics are observed indicating the spiral spin configuration is strongly distorted. In particular, the observation of a 2nd-order harmonic associated with chiral wave vector\(5–2\) ignited a debate of the FE mechanism between the “pure electronic” model by Katsura, Nagaosa, and Balatsky\(10,11\) that suggests the electric polarization can be induced without the involvement of lattice degrees of freedom, and the “ion displacement” model that suggests a lattice distortion via spin-orbit coupling is essential for the FE polarization\(5–2\). Resonant x-ray diffraction study\(9\) reveals the melting of the chiral order associated with the onset of the electric polarization at 27 K, in which the 2nd-order harmonic is unambiguously detected with an order parameter behaving similarly to what we observed in (\( \text{ND}_4 \))\(_2\)\(\text{FeCl}_5\)\(_2\)\(\text{D}_2\)\(\text{O}\). The intensity of the 2nd-order harmonic increases sharply at 27 K coinciding with the onset of the ferroelectricity. These experimental results combined with theories\(5–2\) have classified Tb\(\text{MnO}_3\) as a prototypical material in which a cycloidal-spin structure generates an electric polarization via the spin-orbit interaction.

Comparing our results to Tb\(\text{MnO}_3\), it is quite remarkable that almost identical behavior (spin texture) is observed in a molecular compound like (\( \text{ND}_4 \))\(_2\)\(\text{FeCl}_5\)\(_2\)\(\text{D}_2\)\(\text{O}\) in which the orbital angular momentum of \( \text{Fe}^{3+} \) is expected to be completely quenched. Both even and odd-order harmonics are observed. Similarly, the magnetic structure of (\( \text{ND}_4 \))\(_2\)\(\text{FeCl}_5\)\(_2\)\(\text{D}_2\)\(\text{O}\) changes from IC collinear sinusoidal to IC cycloidal-spiral at \( T_{FE} \) without an IC-C lock-in. The material is paraelectric between \( T_{FE} \) \( T < T_N \) and ferroelectricity only sets in below \( T_{FE} \), where the 2nd-order harmonic appears indicating lattice distortion resulting from spin-lattice coupling. Hence, our neutron diffraction study provides unambiguous evidence that spin-lattice coupling plays a key role in its FE mechanism. Moreover, the observed coexistence of incommensurate and commensurate phases adds additional complexity in this material compared with Tb\(\text{MnO}_3\). A preliminary inelastic neutron scattering study suggests they are both energetically favorable and the compe-
tion between these two phases is responsible for the rich magnetic field versus temperature phase diagram of \((\text{ND}_4)_2\text{FeCl}_5\cdot\text{D}_2\text{O}\).

In summary, our neutron diffraction study on \((\text{ND}_4)_2\text{FeCl}_5\cdot\text{D}_2\text{O}\) reveals that the appearance of ferroelectricity in this molecular magnet is mediated by spin-lattice coupling. We find remarkable similarities with TbMnO\(_3\). Extensive experimental and theoretical studies\(^{4,5}\) indicate that the subtle distorted spin texture observed in TbMnO\(_3\) plays a critical role in the FE mechanism for this prototypical system. Our study suggests \((\text{ND}_4)_2\text{FeCl}_5\cdot\text{D}_2\text{O}\) is a rare molecular analogue to TbMnO\(_3\). Although the incorporation of NH\(_4\) has been used as a strategy to search for new multiferroics, however, to our knowledge only type-I multiferroics have been discovered through this approach so far.\(^{17,19}\) The spin-lattice coupling mediated FE mechanism revealed in this system certainly opens an avenue for searching new multiferroics and calls for more thorough theoretical investigations.

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