

LA-UR- 08-5530

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Intended for: Physical Review Letters



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Magnetic exchange interaction between rare-earth and Mn ions in multiferroic hexagonal manganites

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(Dated: August 4, 2008)

Abstract

We report a study of magnetic dynamics in multiferroic hexagonal manganite HoMnO_3 by far-infrared spectroscopy. Low-temperature magnetic excitation spectrum of HoMnO_3 consists of magnetic-dipole transitions of Ho ions within the crystal-field split $J = 8$ manifold and of the triangular antiferromagnetic resonance of Mn ions. We determine the effective spin Hamiltonian for the Ho ion ground state. The magnetic-field splitting of the Mn antiferromagnetic resonance allows us to measure the magnetic exchange coupling between the rare-earth and Mn ions.

Superexchange is one of the main concepts of magnetism in solids. This exchange interaction underpins an enormous variety of magnetic and related phenomena, including the differently ordered magnetic states and strong electron correlation effects. Recently, superexchange was found to mediate magnetoelectric (ME) coupling in multiferroics - materials with coexisting magnetic and ferroelectric orders¹⁻³. ME coupling may find important technological applications, for example, in an electric-write magnetic-read memory element⁴. Multiferroic hexagonal manganites display robust room-temperature ferroelectricity and remarkable ME behavior⁵ that allow technological exploitation⁶. In these multiferroics, the superexchange interaction between the rare-earth and manganese ions is responsible for the strong ME coupling that allows the control of ferromagnetism by an electric field². The details of the rare-earth/manganese superexchange coupling have heretofore remained unknown. Here, we report the measurement of the rare-earth/manganese superexchange in multiferroic hexagonal HoMnO_3 via the detection of an antiferromagnetic resonance by far-infrared (far-IR) spectroscopy. These results provide new insights into the interplay between magnetism and ferroelectricity in hexagonal manganites with magnetic rare earth ions.

HoMnO_3 (HMO) crystallizes in a hexagonal lattice, space group $P6_3cm$. The crystal structure consists of layers of corner-sharing trigonal MnO_5 bipyramids separated by layers of Ho^{3+} ions². Ferroelectricity sets in at $T_{FE} = 875$ K with the polarization along the hexagonal c axis and results from electrostatic and size effects that lead to the buckling of MnO_5 bipyramids and the displacement of the Ho^{3+} ions out of the a - b plane^{2,7}. The magnetic structure of HMO is formed by Mn^{3+} ions in a two-dimensional triangular network in the a - b plane coupled by antiferromagnetic (AF) exchange and also by Ho^{3+} ($4f^{10}$, ground state $J = 8$) ions located at $2a$ and $4b$ lattice sites with point symmetries C_{3v} and C_3 , respectively^{8,9}. The AF exchange between Mn^{3+} magnetic moments leads to their 120° ordering at $T_N = 72$ K^{10,11}, and easy plane anisotropy confines the Mn spins to the a - b plane. Two in-plane spin reorientation transitions occur at temperatures $T_{SR1} \approx 38$ K and $T_{SR2} \approx 5$ K associated with changes of magnetic symmetry to $P6'_3cm'$ and to $P6_3cm$, respectively^{8,10,12}. The spin reorientation at $T_{SR2} \approx 5$ K is accompanied by antiferromagnetic ordering of the Ho^{3+} ions located at the $4b$ lattice sites⁸. Several manifestations of ME coupling in HMO have been documented, one of the most prominent being the presence of the reentrant phase with a strong magnetodielectric response^{2,13-16}.

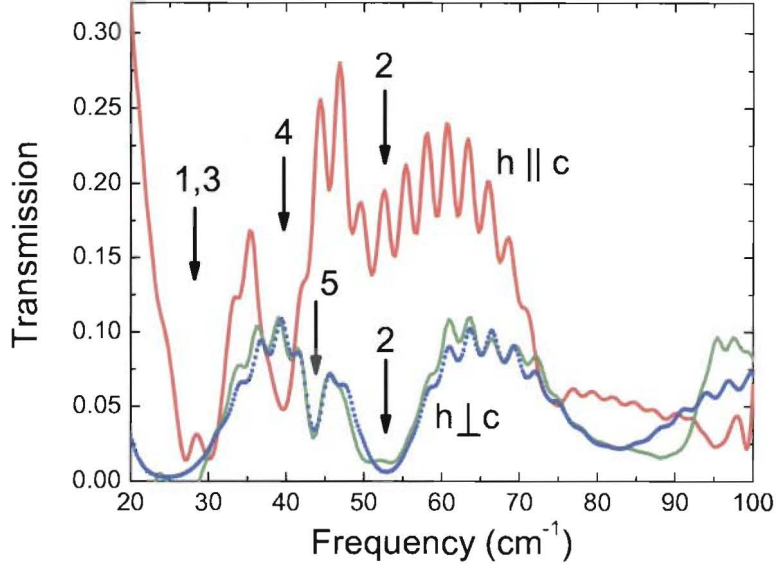


FIG. 1: Power transmission spectra of HoMnO₃ with polarized far-IR light incident along the crystal's [110] direction at T=10 K. Arrows indicate magnetic absorption lines. The dotted line illustrates the simulation of the transmission spectrum using Eqs. (1) and (2).

The HMO single crystals were grown using an optical floating zone furnace and characterized by static magnetization, electric polarization and dielectric constant measurements¹¹. The material's magnetic phase diagram¹¹ at temperatures above 5 K displays the reentrant phase discovered by Lorenz *et al*¹⁴. Polarized far-IR transmission of a 0.4 mm thick, (110) oriented crystal was measured as a function of temperature and applied magnetic field using a Bruker 66 Fourier-transform spectrometer coupled to an 8 Tesla split-coil superconducting magnet¹⁷. Static magnetic field B was applied both parallel and perpendicular to the c axis.

Figure 1 shows far-IR power transmission spectra of the HMO crystal at low temperatures and zero magnetic field for two polarizations of the light incident along the [110] direction - the magnetic h -field of the lightwave parallel ($h||c$) and perpendicular ($h\perp c$) to the c axis. The spectra in Fig. 1 are normalized to the transmission through an empty aperture of the same size as the sample. The fringes with the period of $\approx 3\text{cm}^{-1}$ seen in the spectra are due to multiple reflections of light within the sample. The transmission minima indicated by arrows

correspond to magnetic resonance frequencies. The broad absorption in both polarizations at frequencies above 70 cm^{-1} is of non-magnetic origin, as it exhibits no magnetic field dependence. The dotted line in Fig. 1 shows the simulated transmission, assuming that the magnetic and dielectric response of HMO consists of a collection of lorentzian oscillators:

$$\epsilon(\omega) = \epsilon' + i\epsilon'' + \frac{f_0}{\omega_0^2 - \omega^2 - i\gamma_0\omega}, \quad (1)$$

$$\mu(\omega) = 1 + \sum_j \frac{f_j}{\omega_j^2 - \omega^2 - i\gamma_j\omega}, \quad (2)$$

where ω_j , γ_j , and f_j are frequency, relaxation rate, and oscillator strength of magnetic resonances. ϵ' and ϵ'' are constants, and ω_0 , γ_0 , and f_0 are introduced to account for the non-magnetic absorption above 70 cm^{-1} . The simulation shown by the dotted line in Fig. 1 uses three magnetic resonances that are centered at 25.1, 43.4, and 52.1 cm^{-1} .

Color-coded absorption maps in Fig. 2 illustrate the magnetic field dependence of the resonance frequencies at $T=10 \text{ K}$. To create the maps, we recorded the transmission spectra in 1 T steps between 0 and 8 T. The darker color in Fig. 2 indicates stronger absorption and the white color indicates where the absorption is the weakest; the symbols specify the positions of magnetic resonances extracted from the numerical simulation of the far-IR transmission. In the remainder of this Letter, we show that the magnetic field dependence of the resonance frequencies carries an unambiguous signature of the superexchange interaction between magnetic Ho^{3+} and Mn^{3+} ions that allows us to quantify the interaction.

We attribute the absorption lines labeled 1-4 in Figs. 1 and 2 to crystal field excitations of Ho^{3+} ions. The crystal fields of C_{3v} and C_3 symmetry have qualitatively similar effects on the Ho ion. According to the work of Elliott and Stevens¹⁸⁻²¹, the crystal field potential V_{CF} can be written in terms of spin operators J , J_z , and J_{\pm} for the manifold of states of constant J . The diagonal part of V_{CF} depends only on J and J_z . The off-diagonal part of V_{CF} has non-zero matrix elements only between the states m and m' that differ by $\Delta m = 6$, $m' = m \pm 6$, in the field of C_3 symmetry and by $\Delta m = 3, 6$, $m' = m \pm 3, 6$, in the field of C_{3v} symmetry. In zero applied magnetic field, the degenerate $J = 8$ manifold splits into a set of singlet and doublet states that have the form $|\pm 6, \pm 3, 0\rangle$ and $|\pm 8, \pm 5, \pm 2, \mp 1, \mp 4, \mp 7\rangle$, respectively. The full quantitative description of Ho crystal field levels would rely on the knowledge of crystal field parameters. The measured transition energies and their magnetic field dependence in principle allow the determination of those parameters. Here, we restrict

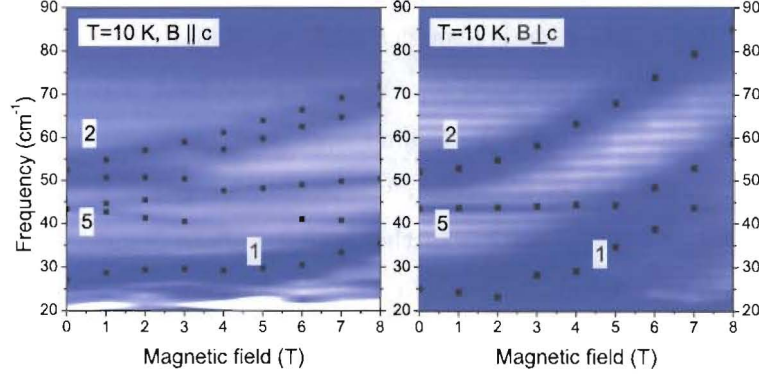


FIG. 2: Absorption maps for the ($h \perp c$) polarization where darker color indicates higher absorption. Left panel - static field $B \parallel c$, right panel - $B \perp c$. Symbols represent magnetic resonance positions extracted from fitting the measured transmission using Eqs. (1) and (2). Resonances are labeled by the same numbers used in Fig. 1.

our discussion to a qualitative description of the observed crystal field transitions, as the frequency range of $20 - 80 \text{ cm}^{-1}$ in our far-IR measurements does not allow us to characterize all magnetic-dipole active crystal field transitions. For example, the 12.5 cm^{-1} transition observed by inelastic neutron scattering¹⁰ is outside of our accessible frequency range.

The magnetic-dipole selection rule for the ($h \perp c$) polarization follows from the action of perturbation $-\gamma \hbar h_{\perp} J_{\pm} \cos \omega t$ on the crystal field eigenstates and allows only transitions with $\Delta m = \pm 1$. This rule forbids transitions between singlets, but allows transitions between singlets and doublets and between different doublets. Transitions 1 and 2 are strong in the ($h \perp c$) polarization (Fig. 2) but are very faint in the ($h \parallel c$) polarization, which suggests that they are allowed in the former, but forbidden in the latter, polarization. Since we observe transition 2 in the highest applied magnetic fields, it must be the transition from the ground state. The final state of the transition could be either a doublet or a singlet. With a magnetic field B applied along the c axis (left panel of Fig. 2) the transition splits linearly, a characteristic of the Zeeman splitting of a doublet. The lower branch of the Zeeman-split transition disappears when the applied field increases beyond 3 T, which is indicative of a thermal depopulation of the upper branch of the Zeeman-split ground state doublet. The

slope of the upper branch of transition 2 allows an estimation of the g -factor in the effective spin Hamiltonian $\mathcal{H}_{eff} = g\mu_B B \tilde{S}$, $\tilde{S} = 1/2$, that describes the ground state. Assuming that the final state of the transition is a singlet with energy independent of the applied field (at sufficiently low fields), we get $g_z \approx 8.8$. The magnetic field B perpendicular to the c axis produces no Zeeman splitting of the ground state doublet. Therefore, neglecting the mixing of the ground state with higher energy levels at sufficiently low field B , we set $g_x = g_y = 0$. The upper branch of transition 2 is further split by about 5 cm^{-1} in magnetic fields along the c axis, which we attribute to hyperfine interactions.

We now turn to the magnetic response of Mn ions in HMO and assign the sharp absorption line at 43.4 cm^{-1} in the $(h \perp c)$ polarization to the antiferromagnetic resonance (AFMR) of the triangular lattice of Mn magnetic moments (transition 5 in Figs. 1, 2). This assignment is supported by the observation of the triangular AFMR at the same frequency ($\approx 43 \text{ cm}^{-1}$) and light polarization in hexagonal YMnO_3 ²², a related compound with the same Mn magnetic structure and similar exchange and anisotropy parameters²³. The AFMR corresponds to the gapped $k = 0$ magnon detected by inelastic neutron scattering at 47 cm^{-1} in HMO¹⁰. The similarity between the $k = 0$ magnon frequency and the position of the AFMR in our data further supports the assignment of the resonance. The magnetic field dependence of the AFMR agrees qualitatively with the expected linear splitting induced by the field applied along the c axis ($B \parallel c$) (left panel of Fig. 2). A slight increase in the AFMR frequency and no splitting of the resonance are observed in fields up to 5 T applied perpendicular to the c axis ($B \perp c$). In fields of 5 T and higher, the AFMR line is hard to identify, as it becomes a part of the broad absorption band due to the crystal field levels of Ho ions (right panel of Fig. 2). The same qualitative behaviour, *i.e.*, linear splitting with $B \parallel c$ and no splitting with $B \perp c$, in fields up to 5.4 T was recorded for the AFMR in YMnO_3 ²².

To quantify the magnetic-field behaviour of the AFMR, we use the following free energy:

$$F = \lambda \sum_{i \neq j} \mathbf{M}_i \cdot \mathbf{M}_j + K \sum_i (M_i^z)^2 - B \sum_i M_i^z, \quad (3)$$

where M_i , $i = 1, 2, 3$, are the three sublattice magnetizations within the $a - b$ plane, λ describes the antiferromagnetic exchange between the sublattices, and K represents the easy-plane anisotropy. The $B = 0$ excitation spectrum of the free energy (3) consists of a zero-frequency mode corresponding to global rotations of the sublattices about the c axis and a doubly-degenerate gapped mode whose degeneracy is lifted by the application of $B \parallel c$.

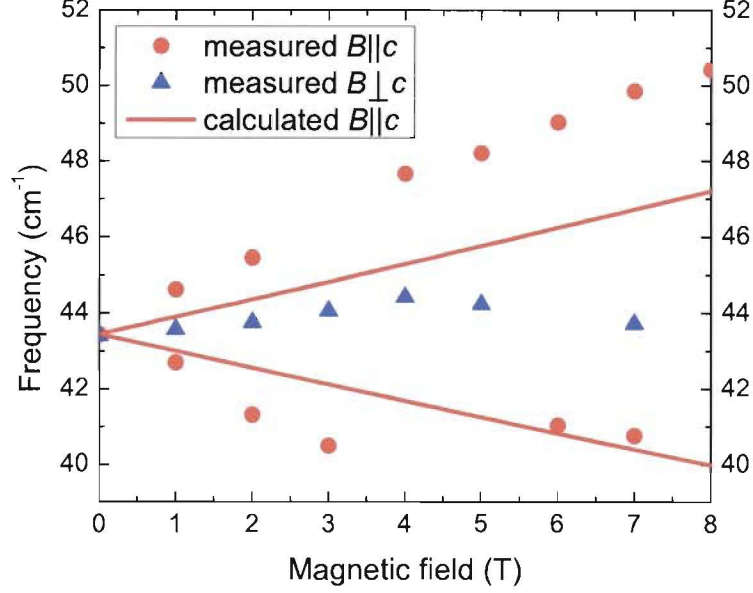


FIG. 3: Field dependence of triangular AFMR frequencies at T=10 K. The measured splitting of the resonance by magnetic field is about twice the splitting calculated using the free energy of Eq. (3).

The analytical expression for the frequency of the gapped mode in $B||c$ was given by Palme *et al*²⁴:

$$\omega_{\pm}^2 = \frac{ab}{2} - \frac{b(a-b)}{2(a+b)^2}B^2 \pm \frac{bB}{2(a+b)^2} \sqrt{B^2b(b-2a) + 2ab(a+b)^2}, \quad (4)$$

where $a = 2H_d$, $b = 3H_{ex}$, the exchange field $H_{ex} = \lambda M_0$, the anisotropy field $H_d = KM_0$, and M_0 is the sublattice magnetization. Equation (3) is a molecular-field version of the Hamiltonian used by Vajk *et al* to describe the spinwave dispersion in HMO¹⁰. In this picture, $H_{ex} = 3SJ = 127$ T and $H_d = SD = 5.75$ T, where J and D are the exchange and anisotropy parameters measured by inelastic neutron scattering and S is the total spin of Mn ions. The value of H_d was lowered slightly to reproduce the zero-field AFMR frequency observed in our far-IR study. The field dependence of the AFMR calculated using Eq. (4) and the g -factor $g = 2$ is shown in Fig. 3. The calculated frequencies largely disagree with the measured ones as the measured splitting is about twice the calculated splitting. For

comparison, the splitting of the AFMR in YMnO₃ is in good agreement with our calculation: the measured splitting of 4.8 cm⁻¹ at ≈ 5.4 T ($g = 1.9 \pm 0.1$)²² agrees with the calculated value of 4.9 cm⁻¹. Here we use exchange and anisotropy fields measured by inelastic neutron scattering²³ in YMnO₃²⁵.

What causes the large discrepancy between the measured and calculated splitting of the AFMR line in HMO? Our calculation describes well the AFMR in YMnO₃, a material that is largely similar to HMO, but with one important difference: the Y³⁺ ions are not magnetic. Therefore, we conclude that the magnetic Ho ions in HMO are responsible for the discrepancy. The Mn ions in HMO interact with the surrounding Ho ions via an exchange coupling of the form $J'_{ij} \mathbf{J}_i^{Ho} \cdot \mathbf{S}_j^{Mn}$, where \mathbf{S}^{Mn} and \mathbf{J}^{Ho} are the angular momenta of Mn and Ho ions. Using the equivalence of matrix elements $g_J J_z = g_z \tilde{S}_z$, where $g_J = 5/4$ is the Lande factor, g_z and \tilde{S}_z are the effective g-factor and spin for the Ho ion ground state, we can write the exchange Hamiltonian as¹⁸

$$\mathcal{H}_{ex} = \mathcal{J}_x \tilde{S}_x S_x^{Mn} + \mathcal{J}_y \tilde{S}_y S_y^{Mn} + \mathcal{J}_z \tilde{S}_z S_z^{Mn}, \quad (5)$$

where $\mathcal{J}_z = (g_z/g_J)J'_{ij}$, etc. We found earlier that $g_z \approx 8.8$ and $g_x = g_y = 0$, which reduces the Hamiltonian to $\mathcal{H}_{ex} = \mathcal{J}_z \tilde{S}_z S_z^{Mn}$. In magnetic field B applied along the c axis, we can replace \tilde{S}_z by its thermal average $\langle \tilde{S}_z \rangle = \chi B / g_z \mu_B$ and write $\mathcal{H}_{ex} = (\mathcal{J}_z \chi) / (g_z \mu_B) B S_z^{Mn}$, where χ is the magnetic susceptibility of Ho ions. The last expression shows that the Ho-Mn (HM) exchange interaction is equivalent to an effective magnetic field in the direction of the c axis acting on Mn spins. To describe the effect of the HM exchange on the AFMR, we add the term

$$F_{HM} = \frac{\mathcal{J}_z \chi N}{g_z g \mu_B^2} B \sum_i M_i^z = \lambda_{HM} B \sum_i M_i^z, \quad (6)$$

where $N=6$ is the number of Ho ions neighboring a Mn ion, to the free energy of the Eq. (3). The total magnetic field acting on Mn ions is then the sum of the external static field and the effective field of Ho ions, $B^{tot} = (1 - \lambda_{HM})B$, and can be larger or smaller than the external field, thus enhancing or mitigating its effect on the AFMR. Comparing the measured AFMR splitting and the splitting calculated using the Eq. (3), we find that $\lambda_{HM} = -1$, which corresponds to ferromagnetic HM coupling.

The ferromagnetic HM exchange that we have measured causes the electric-field induced ferromagnetic ordering of Ho magnetic moments along the c axis reported by Lottermoser *et al*². The HM interaction may also be responsible for the multitude of low-temperature ($T < 8$

K) magnetic phases and the two Mn spin reorientation transitions in HMO¹⁰. The reentrant phase that is induced in HMO when the magnetic field is applied along the c axis displays a strong magnetodielectric effect¹⁴. Such ME behaviour was explained by the formation of magnetic domain walls during spin reorientation, which reduces the local magnetic symmetry and allows the coupling between magnetic and ferroelectric polarizations^{13,14}. Theoretical descriptions of the domain-wall magnetoelectricity must include domain wall contributions to the free energy from lattice distortions and magnetic superexchange interactions. Such descriptions are incomplete without the contribution of the magnetic exchange between the rare earth and Mn ions. Thus, we believe that the rare earth-Mn interaction plays a significant role in all aspects of the physics of multiferroic hexagonal manganites. Our measurement of the ferromagnetic HM exchange provides an important contribution to the modeling of the interplay between magnetism and ferroelectricity in hexagonal manganites with magnetic rare earth ions.

I. ACKNOWLEDGMENTS

We would like to thank Darryl Smith for illuminating theoretical discussions and Jonathan Gigax for the numerical simulation of the Ho crystal field eigenstates. The work at Los Alamos National Laboratory was supported by the LDRD program and the Center for Integrated Nanotechnologies.

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- ²⁵ Sato *et al.*²³ find that the chemical triangular lattice of Mn ions is slightly trimerized with exchange strength J_1 between the nearest neighbors belonging to the same trimer and exchange strength J_2 between the ions belonging to neighboring trimers. The zero wave-vector magnon spectrum of the trimerized chemical lattice is calculated using Eq. (4) and the substitution

$$J \rightarrow (J_1 + 2J_2)/3.$$