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Mesoscopic bar magnet based on ϵ -Fe₂O₃ hard ferrite

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Ferrite magnets have a long history. They are used in motors, magnetic fluids, drug delivery systems, etc. Herein we report a mesoscopic ferrite bar magnet based on rod-shaped ϵ -Fe₂O₃ with a large coercive field (>25 kOe). The ϵ -Fe₂O₃-based bar magnet is a single crystal with a single magnetic domain along the longitudinal direction. A wide frequency range spectroscopic study shows that the crystallographic a -axis of ϵ -Fe₂O₃, which corresponds to the longitudinal direction of the bar magnet, plays an important role in linear and non-linear magneto-optical transitions, phonon modes, and the magnon (Kittel mode). Due to its multiferroic property, a magnetic-responsive non-linear optical sheet is manufactured as an application using an ϵ -Fe₂O₃-based bar magnet, resin, and polyethylene terephthalate. Furthermore, from the viewpoint of the large coercive field property, we demonstrate that a mesoscopic ϵ -Fe₂O₃ bar magnet can be used as a magnetic force microscopy probe.

The first ferrite magnet, which was composed of magnetite, was discovered in Greece in the 7th century BC. Today ferrite bar magnets are common and economical materials; they are used in toys, stationaries, and crafts as well as technical applications. Ferrite magnets such as Fe₂O₃, Fe₃O₄, and BaFe₁₂O₁₉ have contributed to various industrial products^{1–9}. Fe₂O₃ has four known phases: α -, β -, γ -, and ϵ -Fe₂O₃. γ - and α -Fe₂O₃ are naturally existing minerals, whereas β -Fe₂O₃ and ϵ -Fe₂O₃ are uncommon and synthesized in the laboratory. γ -Fe₂O₃ has a spinel structure and is a ferrimagnet^{10–12}. α -Fe₂O₃ has a corundum structure and shows weak ferromagnetism^{13–21}. β -Fe₂O₃ has a bixbyite structure and exhibits antiferromagnetism^{22–24}. ϵ -Fe₂O₃ has an orthorhombic crystal structure^{23,25,26}. A pure phase of ϵ -Fe₂O₃ synthesized in 2004 was found to exhibit a large coercive field²⁷. Since then, various studies on its magnetic properties have been reported^{28–34}. The present study aims to investigate how large an ϵ -Fe₂O₃ single crystal can be grown, its magnetic property, and its magnetic domain structure. We prepare a bar magnet based on a mesoscopic ϵ -Fe₂O₃ rod and investigate its magnetic domain structure using atomic force microscopy (AFM) and magnetic force microscopy (MFM). Additionally, we measure various optical spectra over a wide frequency range from 75 GHz (wavelength λ = 4 mm) to 750 THz (λ = 400 nm) using Faraday spectroscopy, far-infrared (Far-IR) spectroscopy, terahertz (THz) time-domain spectroscopy, and non-linear magneto-optical Faraday spectroscopy. Combining the spectroscopic data with theoretical calculations for the magneto-optical transition moments, phonon modes, and the magnon, we clarify the linear and non-linear magneto-optical transitions, lattice vibrations, and the Kittel-mode magnon. Moreover, we prepare an ϵ -Fe₂O₃/resin on polyethylene terephthalate (PET) and propose a magnetically controlled non-linear optical sheet. Furthermore, since this bar magnet is resistant to external magnetic fields, we apply a mesoscopic ϵ -Fe₂O₃ rod as an MFM probe of a cantilever.

Results and Discussion

Morphology and crystal structure. ϵ -Fe₂O₃ rods were prepared by combining the reverse-micelle and sol-gel techniques. (See Methods and Supplementary Information.) The scanning electron microscopy (SEM) images show that the ϵ -Fe₂O₃ rods are on the mesoscopic scale (i.e., several hundred nanometers in the longitudinal direction). The powder X-ray diffraction (XRD) pattern with Rietveld analysis indicates that the crystal structure is orthorhombic $Pna2_1$ with lattice constants of a = 5.08923(16) Å, b = 8.7858(3) Å, and c = 9.4766(2) Å (Table S1). ϵ -Fe₂O₃ has four different Fe sites; the coordination geometries of Fe_AO₆ and Fe_BO₆ are distorted octahedral, Fe_CO₆ is regular octahedral, and Fe_DO₄ is tetrahedral (Fig. 1a). ϵ -Fe₂O₃ has a noncentrosymmetric

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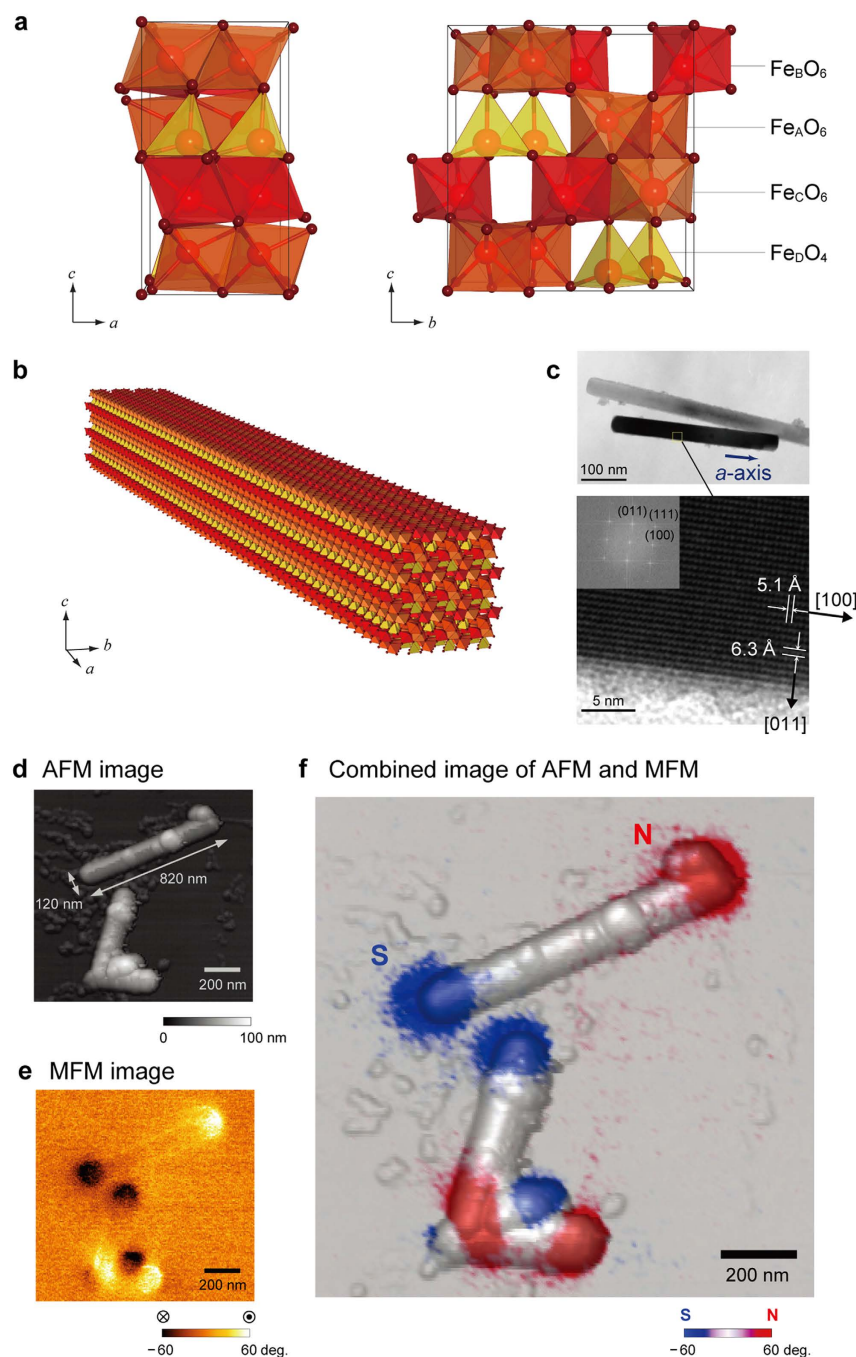


Figure 1. Crystal structure and AFM and MFM images of the mesoscopic ϵ -Fe₂O₃ rod. (a) Crystal structure of the ϵ -Fe₂O₃ unit cell (orthorhombic, $Pna2_1$) viewed along the b -axis (left) and a -axis (right). Red, dark red, orange octahedra, and the yellow tetrahedra indicate the four nonequivalent Fe sites for Fe_AO₆, Fe_BO₆, Fe_CO₆, and Fe_DO₄, respectively. (b) Schematic illustration of a bar magnet based on a mesoscopic ϵ -Fe₂O₃ rod. (c) HRTEM images of the mesoscopic ϵ -Fe₂O₃ rod showing that the longitudinal direction is along the crystallographic a -axis of ϵ -Fe₂O₃. Lower figure is an enlarged image, and the inset shows the Fourier transformed image. (d) AFM image of the ϵ -Fe₂O₃ rods. Color scale indicates the height. One of the mesoscopic ϵ -Fe₂O₃ rods observed in the upper part of the image has longitudinal- and short-axes of 820 nm and 120 nm, respectively. (e) MFM image of the mesoscopic ϵ -Fe₂O₃ rods. Color scale indicates the vectors of the magnetic flux going out of (white) and into (black) the page, which correspond to the N-pole and S-pole, respectively. (f) Combined AFM and MFM image for the ϵ -Fe₂O₃ rods. Red and blue indicate the N-pole and the S-pole, respectively.

crystal structure with an electric polarization along the crystallographic c -axis. The high-resolution transmission electron microscopy (HRTEM) measurements show that the longitudinal direction of the ϵ -Fe₂O₃ rod is the crystallographic a -axis (Fig. 1b,c).

Single magnetic domain structure. ϵ -Fe₂O₃ rods with sizes near 1 μ m were spread on a carbon plate for the AFM measurements as shown in the SEM images of Figure S1. Figure 1d shows the AFM topograph of an ϵ -Fe₂O₃ rod on a carbon plate. The longitudinal- and short-axes of this rod are 820 nm and 120 nm, respectively, and the height is 120 nm.

The same area was also measured by MFM. Figure 1e shows the magnetic response image. The combined AFM and MFM image shows that the north (N) and south (S) poles are at the edges of the mesoscopic ϵ -Fe₂O₃ rods (Fig. 1f), indicating that the ϵ -Fe₂O₃ rod has a single magnetic domain structure. The direction of the magnetic pole (i.e., the magnetic easy-axis) is along the longitudinal direction, which corresponds to the crystallographic *a*-axis. Such a mesoscopic scale single crystal rod with a single magnetic domain structure has not been reported for a ferrite magnet.

Magneto-optical and optical properties. The magneto-optical effect of a mesoscopic ϵ -Fe₂O₃ rod was measured using Faraday spectroscopy. The mesoscopic ϵ -Fe₂O₃ rod exhibits Faraday ellipticities around 570 nm and 480 nm in the visible region (Fig. 2a, upper). To understand these Faraday ellipticities, the magneto-optical transitions of ϵ -Fe₂O₃ were calculated by first-principles calculations using a Vienna ab initio simulation package. (See Methods.) The density of states of the electronic structure shows ϵ -Fe₂O₃ has a wide band gap, and the calculation well reproduces the Faraday ellipticities (Fig. 2a, lower). Both the top of the valence band and the bottom of the conduction band are polarized by up-spins (Fig. 2b).

The magneto-optical transition at 570 nm is an up-spin \rightarrow up-spin transition (denoted by 1* in Fig. 2a), which is a transition from the valence band composed of the 2p orbitals of O1 and O3 to the conduction band composed of the 3d orbitals on the Fe_C site. The direction of the magneto-optical transition moment is mainly along the crystallographic *a*-axis (Fig. 2c and Movie 1). As for the ellipticity at 480 nm, the optical transition (denoted by 2*) is also assigned to the up-spin \rightarrow up-spin transition from the 2p orbitals of O1 and O3 to the 3d orbitals of Fe_B and Fe_C. On the other hand, the down-spin \rightarrow down-spin transition, which is observed around 460 nm (denoted by 3*), is assigned to the optical transition from the 2p orbitals of O2, O4, O5, and O6 to the 3d orbitals of Fe_A and Fe_D. The real (ϵ') and imaginary (ϵ'') parts of the dielectric functions for each crystallographic axis were also calculated (Figure S2). The observed Faraday spectrum and the anisotropic dielectric functions show that the Faraday effect in the visible region appears when light is irradiated from the end-surface of the bar magnet in the crystallographic *a*-axis direction.

The lattice vibration mode (phonon mode) of the mesoscopic ϵ -Fe₂O₃ rod was measured using Far-IR spectroscopy. Figure 2d shows that the low-frequency range lattice vibrations are observed at 2.62 THz (87.3 cm⁻¹), 3.31 THz (110.4 cm⁻¹), 3.65 THz (121.8 cm⁻¹), 3.87 THz (129.1 cm⁻¹), 4.45 THz (148.4 cm⁻¹), 4.73 THz (157.9 cm⁻¹), and 5.21 THz (173.8 cm⁻¹) in the frequency range of 2.2 THz ($\lambda = 136 \mu$ m, 73 cm⁻¹)–5.4 THz ($\lambda = 56 \mu$ m, 180 cm⁻¹). The Far-IR spectrum in the range of 5.4–12 THz is shown in Figure S3a.

To assign the observed Far-IR spectrum, a phonon mode calculation was carried out using the Phonon code. (See Methods.) The calculated phonon modes (Fig. 2d, lower and 2e) show similar spectral patterns to the observed Far-IR spectrum. The observed lowest frequency lattice vibration of 2.62 THz is assigned to the optical phonon mode (calc. 2.51 THz) due to the Fe atom vibration along the crystallographic *a*-axis with A₁ symmetry (Fig. 2f and Movie 2). The calculated phonon mode spectrum between 5.4 and 12 THz and the phonon density of states are shown in Figure S3b. Table S2 lists all of the phonon modes accompanied by their symmetries and IR/Raman activities.

The Kittel-mode magnon was measured by THz time-domain spectroscopy in the millimeter wave region of 75 GHz (0.075 THz, $\lambda = 4.0$ mm) – 300 GHz (0.3 THz, $\lambda = 1.0$ mm). (See Methods.) The temporal waveform of the transmitted THz pulse is shown in Fig. 2g (inset). The Fourier-transformed spectrum shows that the mesoscopic ϵ -Fe₂O₃ rod possesses a millimeter wave absorption peak at 181 GHz (0.181 THz, $\lambda = 1.66$ mm) with fringe patterns of 20 GHz cycles (Fig. 2g, upper). This absorption peak is due to the resonance of the Kittel-mode magnon, originating from the precession of bulk magnetization around the magnetic easy-axis (crystallographic *a*-axis). This type of zero-magnetic-field ferromagnetic resonance is called a natural resonance. The accompanied fringe pattern is due to the interference between the front and back sample surfaces; the fringe pattern depends on the sample thickness (Fig. 2g, lower).

Considering this interference effect, the real (μ') and imaginary parts (μ'') of the magnetic permeability (Fig. 2h) were evaluated on the basis of the Landau-Lifshitz analysis^{35,36} (Fig. 2i, Movie 3, and Methods). In the present ferrite bar magnet, the Kittel-mode magnon frequency is remarkably high and approaches the phonon mode frequency. The Kittel-mode magnon frequency of 0.181 THz is as high as 7% of the lowest frequency optical phonon mode of 2.62 THz. This is due to the strong magnetic anisotropy of ϵ -Fe₂O₃. Furthermore, the Kittel-mode resonance frequency in the present bar magnet at 181 GHz has almost the same value as that of the spherical ϵ -Fe₂O₃ nanoparticle at 182 GHz in our previous report²⁸, indicating that the resonance frequency is independent of the material shape. Thus, the high-frequency Kittel-mode magnon originates from the intrinsic magnetic anisotropy of ϵ -Fe₂O₃, and the contribution of the shape magnetic anisotropy is insignificant.

Polar-crystal ferrite bar magnet. The electric conductivity of the mesoscopic ϵ -Fe₂O₃ rod was investigated by impedance measurements with a precision impedance analyzer in the frequency range from 40 Hz to 110 MHz using the four-terminal pair method. Figure 3a shows the real (Z') and imaginary (Z'') parts of the complex plane impedance. The Cole-Cole circular arc fitting plot, which considers a Gaussian distribution of the conductivity, gives a conductivity value (σ) of 9×10^{-8} S cm⁻¹. Therefore, the mesoscopic ϵ -Fe₂O₃ rod has an insulating character.

The crystal structure of this material is a polar crystal with the *Pna*2₁ space group. In fact, hybrid functional calculations suggest an electric polarization along the *c*-axis as shown in the charge polarization map of Fig. 3b

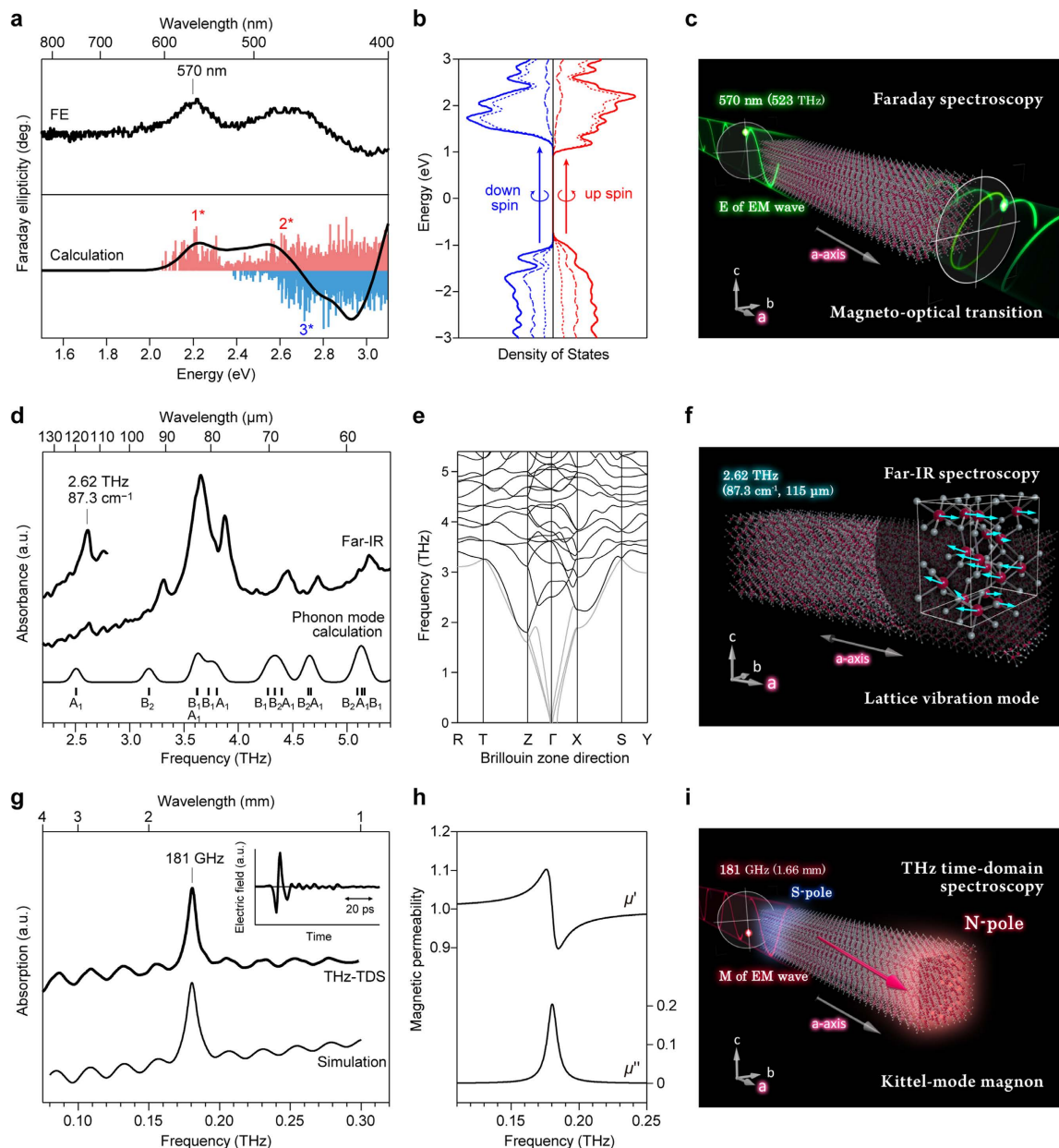


Figure 2. Faraday spectrum, Far-IR spectrum, and Kittel-mode magnon spectrum. (a) Experimentally obtained Faraday ellipticity (FE) spectrum (upper), and calculated magneto-optical transition probability by first-principles calculation (lower). Pink sticks, blue sticks, and black line in the lower panel indicate the transition probabilities of the up-spin \rightarrow up-spin transitions, down-spin \rightarrow down-spin transitions, and their sum, respectively. (b) Density of states of the electronic structure for ϵ -Fe₂O₃ obtained by first-principles calculation. Red and blue indicate the up- and down-spins, respectively, while the solid, dotted, and dashed lines indicate the density of states for the total, Fe, and O, respectively. (c) Linearly polarized light is converted into rotated elliptical light by the optical transition of up-spin \rightarrow up-spin transition at 570 nm along the longitudinal axis (parallel to the crystallographic *a*-axis), resulting in the Faraday effect. *E* is the electric field of the electromagnetic (EM) wave. (d) Far-IR spectrum (thick lines) and the calculated spectrum (thin line) by phonon mode calculation of ϵ -Fe₂O₃. Tick marks indicate the calculated positions and the vibrational symmetries of the phonon modes. (e) Phonon dispersion of ϵ -Fe₂O₃. Black and gray lines indicate the optical and the acoustic phonon modes, respectively. (f) Atomic movement of the lowest frequency optical phonon mode of 2.62 THz (calc. 2.51 THz) with *A*₁ symmetry. The Fe atoms are vibrating in the longitudinal direction of the rod (parallel to the crystallographic *a*-axis). (g) Kittel-mode magnon spectrum of the mesoscopic ϵ -Fe₂O₃ rod measured by THz time-domain spectroscopy (THz-TDS) (thick line) and the fitted spectrum by the Landau-Lifshitz analysis considering the interference effect between the front and back surfaces of the sample (thin line). The inset is the temporal waveform of the transmitted THz pulse. (h) Real (μ') and imaginary part (μ'') of the magnetic permeability obtained by Landau-Lifshitz analysis. (i) Kittel-mode magnon at 181 GHz (0.181 THz) caused by the precession of the bulk magnetization (magenta arrow) around the magnetic easy-axis along the longitudinal direction (parallel to the crystallographic *a*-axis). *M* is the magnetic field of the EM wave.

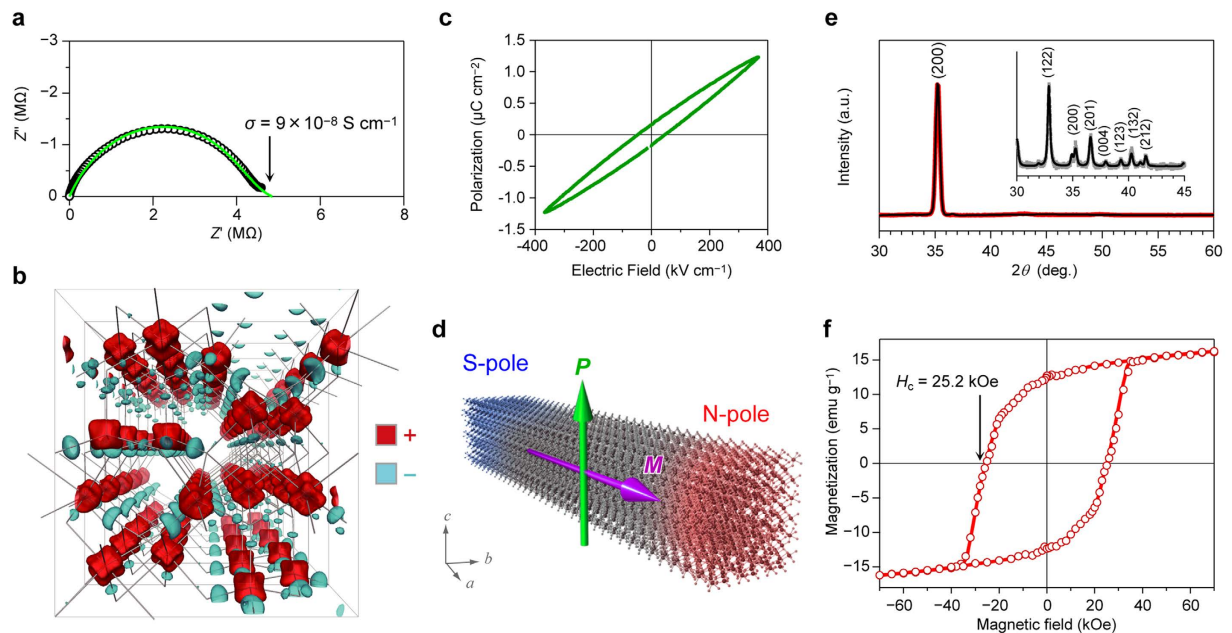


Figure 3. Polar-crystal ferrite bar magnet and thin film of oriented ϵ -Fe₂O₃ rods. (a) Z'' versus Z' plots of the mesoscopic ϵ -Fe₂O₃ rod measured at room temperature. Green line shows the Cole-Cole arc fitting curve. (b) Charge polarization map of ϵ -Fe₂O₃. This figure is viewed along the a -axis. Green and light blue surfaces denote the positively and negatively charged densities with the isosurface levels of $\pm 0.035 a_0^{-3}$ (a_0 : Bohr radius), respectively. (c) Ferroelectric hysteresis loop measured at 77 K. (d) Schematic image of the mesoscopic ϵ -Fe₂O₃ rod with a single magnetic domain. The magnetic polarization (M) of ϵ -Fe₂O₃-based bar magnet is along the crystallographic a -axis (magenta arrow), which corresponds to the longitudinal direction of the rod, while the electric polarization (P) is parallel to the c -axis (green arrow). (e) XRD pattern of the film based on crystallographically oriented ϵ -Fe₂O₃ rods and XRD pattern of the nonoriented ϵ -Fe₂O₃ rods (inset). Red line, gray line, and black lines indicate the observed pattern of oriented ϵ -Fe₂O₃ rods, observed pattern of nonoriented ϵ -Fe₂O₃ rods, and calculated patterns, respectively. (f) Magnetization versus external field plot of the ϵ -Fe₂O₃-rod oriented film measured at room temperature along a single in-plane direction.

(see Methods). In this map, the positively and negatively charged areas are attributed to the Fe cations and O anions, respectively.

Additionally, the ferroelectric hysteresis loop was measured in this mesoscopic ϵ -Fe₂O₃ rod (Fig. 3c). The result shows that the mesoscopic ϵ -Fe₂O₃ rod exhibits ferroelectricity, although the remanent electric polarization and the electric coercive field are slightly lower than the reported values for ϵ -Fe₂O₃ nanocrystals deposited on a SrTiO₃ substrate by pulsed laser deposition³¹. Such a mesoscopic ferrite bar magnet based on a single-magnetic-domain polar-crystal (Fig. 3d) has yet to be reported.

Thin film based on an oriented ferrite bar magnet. To investigate the magnetic hysteresis loop of the mesoscopic ϵ -Fe₂O₃ rods, we prepared a crystallographically oriented ϵ -Fe₂O₃ film using the following procedure; a dispersion solution containing ϵ -Fe₂O₃ rods, polyurethane, and polyvinylchloride was initially dropped onto a PET film. Then the coated film was dried at room temperature under an external magnetic field of 3 Tesla. The XRD pattern of the obtained film shows one sharp peak of (2 0 0). The orientation degree of the crystallographic a -axis in the in-plane direction of the film was evaluated by Rietveld refinement (Fig. 3e). The crystallographic a -axis (longitudinal direction) of the ϵ -Fe₂O₃ rod is remarkably well oriented along a single direction in the film plane. The Lotgering factor is 0.96. The magnetic hysteresis measurements using a superconducting quantum interference device (SQUID) show an H_c value of 25.2 kOe along the a -axis at 300 K and a saturation magnetization value of 16.2 emu g⁻¹ at 7 T. The H_c value of 25.2 kOe is the largest among the H_c values of ϵ -Fe₂O₃ reported to date (Fig. 3f).

Magnetically controlled non-linear magneto-optical film. The correlation between the magnetic polarization along the a -axis and the electric polarization along the c -axis can induce a non-linear Faraday effect. Therefore, we conducted second-order non-linear optical measurements using a sheet of ϵ -Fe₂O₃ rod/resin on PET. Figure 4a shows the optical coordinates of the measurement system, in which the crystallographic a -axis of the ϵ -Fe₂O₃ rod is oriented along the horizontal direction. The sample was irradiated with horizontally polarized incident light ($\lambda = 775$ nm femtosecond laser). The rotation angular dependence of the output second harmonic light intensity $I_{SH}(\theta)$ in the transmission mode shows that the $I_{SH}(\theta)$ value follows $\sin^2\theta$ curve where it is zero at $\theta = 0^\circ$, gradually increases to a maximum at 90° , decreases to zero at 180° , increases again to a maximum at 270° , and finally returns to zero at 360° (Fig. 4b). The clear contrast between light and shade shows a double loop plot in the polar figure; that is, when the ϵ -Fe₂O₃ mesoscopic rod is irradiated with polarized light parallel to the

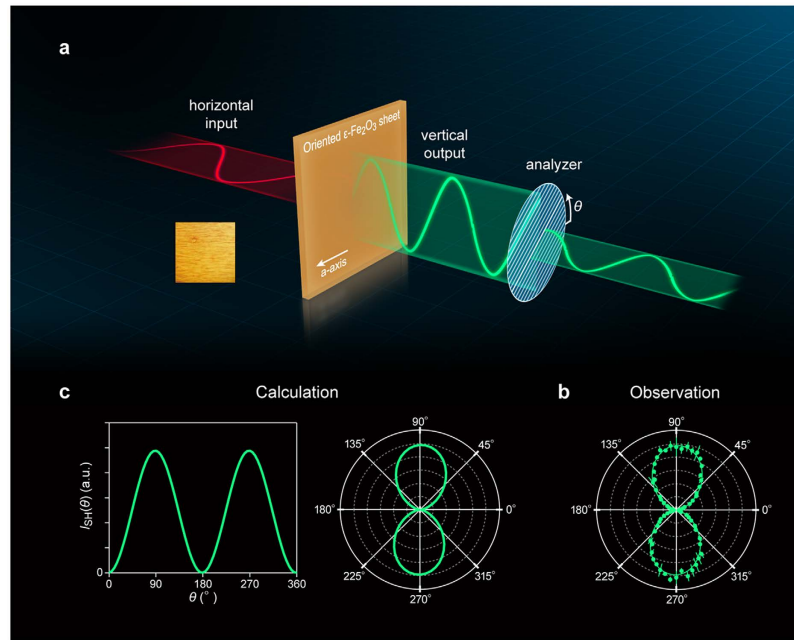


Figure 4. Angular dependence of the non-linear Faraday effect on the oriented ϵ -Fe₂O₃ /resin PET sheet. (a) Optical coordinates of the experimental setup of magneto-optical non-linear Faraday effect, where the crystallographic a -axis of the ϵ -Fe₂O₃ rod is oriented along the horizontal direction in the sheet. Input light shown in red is horizontally polarized 775 nm fundamental light, and output light shown in green is vertically polarized 388 nm second harmonic light. Left inset shows a photograph of the crystallographically oriented ϵ -Fe₂O₃/resin PET sheet with a size of 1 mm \times 1 mm. (b) Observed $I_{SH}(\theta)$ versus θ polar figure. (c) Calculated $I_{SH}(\theta)$ versus θ curve (left) and calculated $I_{SH}(\theta)$ versus θ polar figure (right) based on the theory of non-linear Faraday effect in $Pna2_1$ magnetic space group (see Methods).

longitudinal direction along the crystallographic a -axis, a vertically polarized second harmonic light is emitted (Supplementary Movie 4).

The observed effect can be understood as follows. The second-order non-linear magneto-optical tensor in the $Pna2_1$ magnetic space group shows that $I_{SH}(\theta)$ in the present optical coordinates is expressed as

$$I_{SH}(\theta) \propto 1/2[|\chi_{caa}^{cry}|^2 + |\chi_{baa}^{mag}(M)|^2]E_X^4 \sin^2 \theta, \quad (1)$$

where χ_{caa}^{cry} and $\chi_{baa}^{mag}(M)$ represent the crystallographic and the magnetic terms of the second harmonic susceptibility, respectively. (See Methods for details). The calculated angular dependence agrees well with the observed plots as shown in Fig. 4c.

Tilting the sample sheet by a tilted angle (ϕ) of $+30^\circ$ towards the direction of the incident light shifts the maximum and minimum values by $+3.3^\circ$. On the other hand, a -30° tilt shifts the maximum and minimum values of $I_{SH}(\theta, \phi)$ by -2.5° . The expected $I_{SH}(\theta, \phi)$ and the observed data are shown in Supplementary Information S8 and Supplementary Figures S4 and S5, respectively. If the second harmonic signal only has a crystallographic term, then the absolute value of the \pm shift should be the same. Therefore, the difference in the shift angles originates from the magnetic term of $\chi_{baa}^{mag}(M)$, demonstrating that the oriented ϵ -Fe₂O₃ mesoscopic rod/resin on PET is a magnetically controlled non-linear optical rotation sheet.

Application to an MFM probe on a cantilever. The ϵ -Fe₂O₃ mesoscopic bar magnet has a gigantic H_c value, indicating that it resists an external magnetic flux. Therefore, it should be useful as an MFM probe³⁷. We prepared an MFM probe using an ϵ -Fe₂O₃ rod mounted onto the tip of the non-magnetic cantilever. Figure 5a shows a schematic illustration and SEM image of the ϵ -Fe₂O₃-MFM probe.

Using this ϵ -Fe₂O₃-MFM probe, we measured the magnetic bit cells on a commercially available hard disc, Microdrive of IBM, which is composed of a cobalt magnetic layer. The AFM image of the surface of this hard disc shows that the surface is very flat with a roughness less than 0.4 nm (Fig. 5b). The same area was measured using the ϵ -Fe₂O₃-MFM probe, and we obtained an MFM image with a striped pattern due to magnetic flux leakage from the magnetic bit cells (Fig. 5c). In this MFM image, the black and orange areas indicate the directions of the magnetic flux on the surface of the hard disc medium going out of and into the page, which correspond to the N-pole and S-pole, respectively. The intervals between the stripes vary because the magnetic pole direction of each magnetic bit cell depends on the recorded information. Strong color is observed at the S-S or N-N borders, while the contrast is weak at the N-S or S-N borders. Figure 5d schematically shows the magnetic bit pattern, which is inferred from the observed MFM image. Gray and white areas denote the magnetic bit cells with the magnetic pole directions as shown in the lower left enlarged illustration of Fig. 5d, which are expressed here

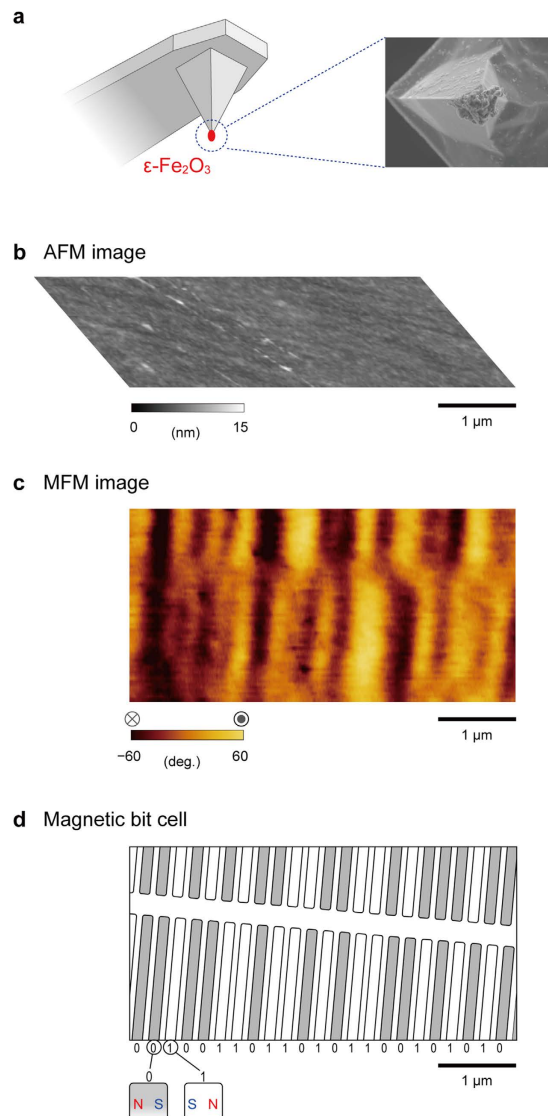


Figure 5. Demonstration of the mesoscopic ϵ -Fe₂O₃ rod as an MFM probe. (a) Schematic illustration of the ϵ -Fe₂O₃ rod mounted on a cantilever (left) and SEM image of the cantilever tip (right). (b) AFM image of a hard disc medium measured using the ϵ -Fe₂O₃-MFM probe. (c) MFM image of a hard disc medium measured using the ϵ -Fe₂O₃-MFM probe. Color scale indicates the vector of the magnetic flux going out of (orange) and into (black) the page, which correspond to the N-pole and S-pole, respectively. (d) Schematic image of the magnetic bit pattern for the observed area of the hard disc medium. Gray and white areas denote the magnetic bit cells with the magnetic pole directions as shown in the lower left enlarged illustration, which are expressed as 0 and 1, respectively.

as 0 and 1, respectively. In the present MFM image, the bit cells are magnetically recorded as the bit pattern of 00100110110101100101010 (Fig. 5d). Hence, the ϵ -Fe₂O₃-MFM probe successfully acquires an MFM image.

The ϵ -Fe₂O₃-MFM probe has the following advantages. (i) The magnetic poles of the ϵ -Fe₂O₃-MFM probe are resistant to a strong magnetic flux from the measured magnetic objects or an applied magnetic field. (ii) Since ϵ -Fe₂O₃ is an insulator, the ϵ -Fe₂O₃-MFM probe does not disorder the magnetization states of the measured magnetic object due to electric current leakage from the cantilever.

Conclusion

In this paper, we report a mesoscopic single crystal bar magnet composed of ϵ -Fe₂O₃ that withstands strong magnetic fields and electric current flow. Additionally, it does not rust. Typical bar magnets are produced by hot pressing a magnetic powder under an external magnetic field. However, the present type of ferrite bar magnet (i.e., an ideal single crystal ferrite bar magnet with a single magnetic domain structure) has not been previously reported because most ferrite magnets exhibit a weak magnetic anisotropy, preventing a single magnetic domain structure.

Concerning the magnetic anisotropy energy density of ϵ -Fe₂O₃, the value is considered to be 7×10^6 erg cm⁻³³², which is much higher than the values for other ferrite magnets. Spectroscopic measurements over a wide frequency range (0.075–750 THz) show that (i) the magneto-optical transition moment of the Faraday effect is along the longitudinal direction of the bar magnet, (ii) the atomic movement of the lowest frequency phonon mode vibrates along the direction of the bar magnet, (iii) bulk magnetization precesses around the longitudinal direction of the bar magnet, and (iv) a non-linear magneto-optical Faraday effect, which depends on the direction of the bar magnet, is observed (Supplementary Movie 5).

We also demonstrate a “magnetization-responsive organic-inorganic composite non-linear optical sheet” as a new application for the ϵ -Fe₂O₃ rod/resin on PET. The mesoscopic ϵ -Fe₂O₃ rod is resistant to the magnetic flux of any magnetic material, and this bar magnet exhibits insulating properties. As an example of an application of ϵ -Fe₂O₃, we demonstrated a cantilever probe for magnetic force microscopy. The “ ϵ -Fe₂O₃-MFM probe” has advantages for measuring strong permanent magnets and avoiding turbulence in the image by an electric current due to accidental contact between the probe and the measured magnetic object.

Methods

Material. Reverse-micelle solution containing iron(III) nitrate and barium(II) nitrate aqueous solution was mixed with reverse-micelle solution containing ammonium hydroxide. Then tetraethyl orthosilicate was added and stirred to form a silica matrix. The obtained precursor was sintered in air at 980 or 1025 °C. The silica matrix was etched according to the modified technique of our previous method²⁷. Details are described in the Supplementary Information.

Physical measurements. SEM and TEM images were acquired using a JEOL JSM-7001F and JEOL JEM 2000EX, respectively. XRD patterns were measured by a Rigaku Ultima IV and Rigaku SmartLab. AFM and MFM measurements were conducted by a Hitachi High-Tech Science AFM5000II/AFM5300E MFM. For the Faraday effect and Far-IR measurements, a JASCO E-250 magneto-optical meter and a JASCO 6100 spectrometer were used, respectively. The magnetic measurements were carried out using a Quantum Design MPMS SQUID magnetometer. The impedance measurements were conducted with an Agilent 4294A precision impedance analyzer. Ferroelectric property was measured using a TF Analyzer 1000, aixACCT electric polarization hysteresis meter.

Magneto-optical transition moment calculation. The magneto-optical transition of ϵ -Fe₂O₃ was calculated by Vienna ab initio Simulation Package³⁸. The magnetic spins were treated by a spin-polarized model. The Brillouin zone was integrated with $7 \times 5 \times 5$ k -mesh and first order Gaussian smearing with a width of 0.1 eV. The magneto-optical transition probabilities of the up-spin and down-spin were obtained from the calculated optical matrix elements, while the sum was calculated using the Gaussian waveform for up-spin \rightarrow up-spin transitions and down-spin \rightarrow down-spin transitions.

Phonon mode calculation. The phonon modes were calculated by the Phonon code³⁹ in a Material Design MedeA package. The atomic positions of ϵ -Fe₂O₃ were optimized with an energy cutoff of 400 eV and $3 \times 3 \times 3$ k -mesh until satisfying a 10^{-5} eV pm⁻¹ force tolerance. The optimized structures were used to calculate the phonon modes of ϵ -Fe₂O₃, which were determined by the direct method implemented in Phonon code with 2-pm displacements using the optimized structures.

Charge polarization map calculation. The charge polarization map in the unit cell of ϵ -Fe₂O₃ was obtained by screened Coulomb hybrid functional calculation of HSE06⁴⁰. The positive and negative charge distributions were calculated by the difference between the charge distribution of ϵ -Fe₂O₃ and the charges of the Fe and O atoms.

Magnon (Kittel mode) measurement. The resonance of the mesoscopic ϵ -Fe₂O₃ rod due to Kittel-mode magnon in the millimeter wave region was measured by THz time-domain spectroscopy. A THz pulse wave was generated using a mode-locked Ti: sapphire pulse laser as a source and a dipole type low-temperature-grown GaAs photoconductive antenna as an emitter. The transmitted THz electric field was detected with a bowtie type low-temperature-grown GaAs photoconductive antenna, which was recorded with variable time delays as a temporal waveform in the time domain using a pump-probe system.

Landau-Lifshitz analysis with interference effect. Kittel-mode magnon spectrum was analysed by considering multiple reflection within the sample. Magnon absorption (A) is described by:

$$A = -20 \log \left| \left(\left\{ 1 - \left[(\sqrt{\mu_r/\epsilon_r} - 1)/(\sqrt{\mu_r/\epsilon_r} + 1) \right]^2 \right\} \exp(-2i\pi f d \sqrt{\mu_r/\epsilon_r}/c) \right) / \left\{ 1 - \left[(\sqrt{\mu_r/\epsilon_r} - 1)/(\sqrt{\mu_r/\epsilon_r} + 1) \exp(-2i\pi f d \sqrt{\mu_r/\epsilon_r}/c) \right]^2 \right\} \right|, \quad (2)$$

where d , μ_r and ϵ_r are thickness, relative magnetic permeability, and relative dielectric constant of the ϵ -Fe₂O₃ pellet sample, respectively, i is the imaginary unit, and c is the speed of light. μ_r , which is a function of frequency (f), is derived from the Landau-Lifshitz equation^{36,37} describing the motion of the magnetization against the intrinsic magnetic anisotropy field of the magnet and the magnetic field of the THz wave;

$$\mu_r' = \mu_r''_{\max} \sin \phi \cos \phi + 1 \quad (3)$$

$$\mu_r'' = \mu_r''_{\max} \sin^2 \phi \quad (4)$$

$$\phi = \arctan[\Delta f / 2(f_r - f)], \quad (5)$$

where f_r is the resonance frequency, $\mu_r''_{\max}$ is the maximum value of μ_r'' , and Δf is the full width at half maximum of μ_r'' peak.

Angular dependence of non-linear Faraday effect. Because the sample is magnetized along the magnetic easy axis of the a -axis, the magnetic space group of the oriented film of ϵ -Fe₂O₃ is $Pna2_1$. In this case, the non-zero tensor elements are $\chi_{baa}^{\text{mag}}(M)$, $\chi_{bbb}^{\text{mag}}(M)$, $\chi_{bcc}^{\text{mag}}(M)$, χ_{bbc}^{cry} , χ_{caa}^{cry} , χ_{cbb}^{cry} , χ_{ccc}^{cry} , $\chi_{cbc}^{\text{mag}}(M)$, χ_{aca}^{cry} , and $\chi_{aab}^{\text{mag}}(M)$, where a , b , and c denote the crystal axes of ϵ -Fe₂O₃. When θ is the angle from the horizontal direction, which is also the analyzer angle, the polarization of the output SH light ($P(\theta)$) is

$$P(\theta) = [\chi_{baa}^{\text{mag}}(M) \cos \varphi \sin \theta \exp\{-i(\alpha + \delta_{baa}^{\text{mag}})\} + \chi_{caa}^{\text{cry}} \sin \varphi \sin \theta \exp(-i\alpha)] E_X^2, \quad (6)$$

where X , Y , and Z are the coordinates of the measurement system, φ is the angle between the b -axis (or c -axis) of the ϵ -Fe₂O₃ mesoscopic rod and the Y -axis (or Z -axis) of the optical coordinate, α is the phase shift due to light propagation, and $\delta_{baa}^{\text{mag}}$ is the phase shift between the magnetic term and the crystallographic term. Thus, $I_{\text{SH}}(\theta)$ is described as

$$I_{\text{SH}}(\theta) \propto P(\theta)P(\theta)^* = [|\chi_{baa}^{\text{mag}}(M)|^2 \cos^2 \varphi + |\chi_{caa}^{\text{cry}}|^2 \sin^2 \varphi + 2\chi_{baa}^{\text{mag}}(M)\chi_{caa}^{\text{cry}} \sin \varphi \cos \varphi \cos \delta_{baa}^{\text{mag}}] E_X^4 \sin^2 \theta. \quad (7)$$

By integrating $I(\theta)$ with respect to φ from 0 to 2π , $I_{\text{SH}}(\theta)$ of the sample with a random orientation in the bc plane is expressed as

$$I_{\text{SH}}(\theta) \propto 1/2[|\chi_{baa}^{\text{mag}}(M)|^2 + |\chi_{caa}^{\text{cry}}|^2] E_X^4 \sin^2 \theta, \quad (8)$$

which is proportional to $\sin^2 \theta$.

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Author Contributions

S.O. designed and coordinated this study, contributed to all measurements and calculations, and wrote the paper. A.N. contributed to sample synthesis, TEM measurements, and THz analysis. T.Y. conducted the AFM and MFM measurements. M.Y. contributed to the THz measurements, analysis of the phonon mode calculations, and charge density calculations. K.I. conducted the magnetic and SHG measurements of the oriented sample and the Faraday spectra measurements. T.N. contributed to the first-principles calculations. S.A. prepared and measured the magnetically oriented sample. Y.U. measured the Far-IR spectrum. K.N. performed the conductivity measurements and ferroelectricity measurements. H.T. coordinated this study and conducted the XRD, SEM, AFM and MFM measurements. All authors commented on the manuscript.

Additional Information

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