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Tuning magnetoelectricity in a mixed-anisotropy antiferromagnet

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Ellen Fogh ^{1,2}✉, Bastian Klemke ³, Manfred Reehuis³, Philippe Bourges ⁴, Christof Niedermayer ⁵, Sonja Holm-Dahlin ^{5,6}, Oksana Zaharko ⁵, Jürg Schefer⁵, Andreas B. Kristensen², Michael K. Sørensen², Sebastian Paeckel ^{3,7}, Kasper S. Pedersen ⁸, Rasmus E. Hansen⁹, Alexandre Pages¹, Kimmie K. Moerner², Giulia Meucci², Jian-Rui Soh ¹, Alessandro Bombardi ¹⁰, David Vaknin ¹¹, Henrik. M. Rønnow ¹, Olav F. Syljuåsen ¹², Niels B. Christensen ²✉ & Rasmus Toft-Petersen ^{2,13}✉

Control of magnetization and electric polarization is attractive in relation to tailoring materials for data storage and devices such as sensors or antennae. In magnetoelectric materials, these degrees of freedom are closely coupled, allowing polarization to be controlled by a magnetic field, and magnetization by an electric field, but the magnitude of the effect remains a challenge in the case of single-phase magnetoelectrics for applications. We demonstrate that the magnetoelectric properties of the mixed-anisotropy antiferromagnet $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$ are profoundly affected by partial substitution of Ni^{2+} ions with Fe^{2+} on the transition metal site. This introduces random site-dependent single-ion anisotropy energies and causes a lowering of the magnetic symmetry of the system. In turn, magnetoelectric couplings that are symmetry-forbidden in the parent compounds, LiNiPO_4 and LiFePO_4 , are unlocked and the dominant coupling is enhanced by almost two orders of magnitude. Our results demonstrate the potential of mixed-anisotropy magnets for tuning magnetoelectric properties.

Utilizing the magnetoelectric (ME) effect to electrically control magnetic states has far-reaching prospects in next-generation electronics^{1,2}. Currently realised applications are based on heterostructure composites³, combining layers with distinct bulk properties. Proposals for application of the ME effect in heterostructures are abundant, including electric-field control of skyrmions⁴, magnetoelectric

spin-orbit logic devices^{5,6}, medical implants^{7–9} and low-power-consumption ME random access memory^{10–12}. As many of these proposals involve distinct ME layers, a fundamental understanding of the properties of single-phase magnetoelectrics is pivotal to their realization. While our understanding of the underlying mechanisms has been greatly improved since the discovery of the ME effect, the relatively

¹Laboratory for Quantum Magnetism, Institute of Physics, École Polytechnique Fédérale de Lausanne (EPFL), CH-1015 Lausanne, Switzerland. ²Department of Physics, Technical University of Denmark, DK-2800 Kongens Lyngby, Denmark. ³Helmholtz-Zentrum Berlin für Materialien und Energie, D-14109 Berlin, Germany. ⁴Université Paris-Saclay, CNRS, CEA, Laboratoire Léon Brillouin, 91191 Gif-sur-Yvette, France. ⁵Laboratory for Neutron Scattering and Imaging, Paul Scherrer Institute, Villigen CH-5232, Switzerland. ⁶Nano-Science Center, Niels Bohr Institute, University of Copenhagen, DK-2100 Copenhagen Ø, Denmark. ⁷Department of Physics, Arnold Sommerfeld Center for Theoretical Physics (ASC), Munich Center for Quantum Science and Technology (MCQST), Ludwig-Maximilians-Universität München, 80333 München, Germany. ⁸Department of Chemistry, Technical University of Denmark, DK-2800 Kongens Lyngby, Denmark. ⁹Department of Photonics Engineering, DK-2800 Kongens Lyngby, Denmark. ¹⁰Diamond Light Source Ltd., Harwell Science and Innovation Campus, Didcot, Oxfordshire OX11 0DE, UK. ¹¹Ames National Laboratory and Department of Physics and Astronomy, Iowa State University, Ames Iowa 50011, USA. ¹²Department of Physics, University of Oslo, P. O. Box 1048, Blindern N-0316 Oslo, Norway. ¹³European Spallation Source ERIC, P.O. Box 176, SE-221 00 Lund, Sweden. ✉ e-mail: ellen.fogh@epfl.ch; nbch@fysik.dtu.dk; rasp@fysik.dtu.dk

weak ME couplings are lingering barriers for applicability of single-phase magnetoelectrics.

The ME properties of a given single-phase material are a consequence of the magnetic point group symmetry inherent to its magnetically ordered state^{13,14}. More specifically, the absolute and relative orientation of the ordered moments dictate the non-zero elements of the ME tensor describing the coupling between electric and magnetic degrees of freedom^{14,15}. Mixing magnetic ions with incompatible, or mismatched, single-ion anisotropies gives rise to what can be thought of as a composite on the atomic level. This random site-dependent anisotropy in combination with the inter-species

exchange interaction creates frustration in the system and may result in what is known as an oblique antiferromagnetic phase. Here, the ordered moments are oriented away from any of the easy axes observed in the stoichiometric compounds^{16–18}.

A well-known family of isostructural magnetoelectric has chemical formula LiMPO_4 ($M = \text{Mn, Fe, Co, Ni}$) and space group $Pnma$ (No. 62)¹⁹ with the crystallographic unit cell illustrated in Fig. 1a. The compounds, LiNiPO_4 ($S=1$)^{20–22} and LiFePO_4 ($S=2$)^{23,24} order antiferromagnetically at 20.8 K and 50 K, respectively. Below their Néel temperatures, they display similar commensurate spin structures except for the orientation of the magnetic moments, which are predominantly along the

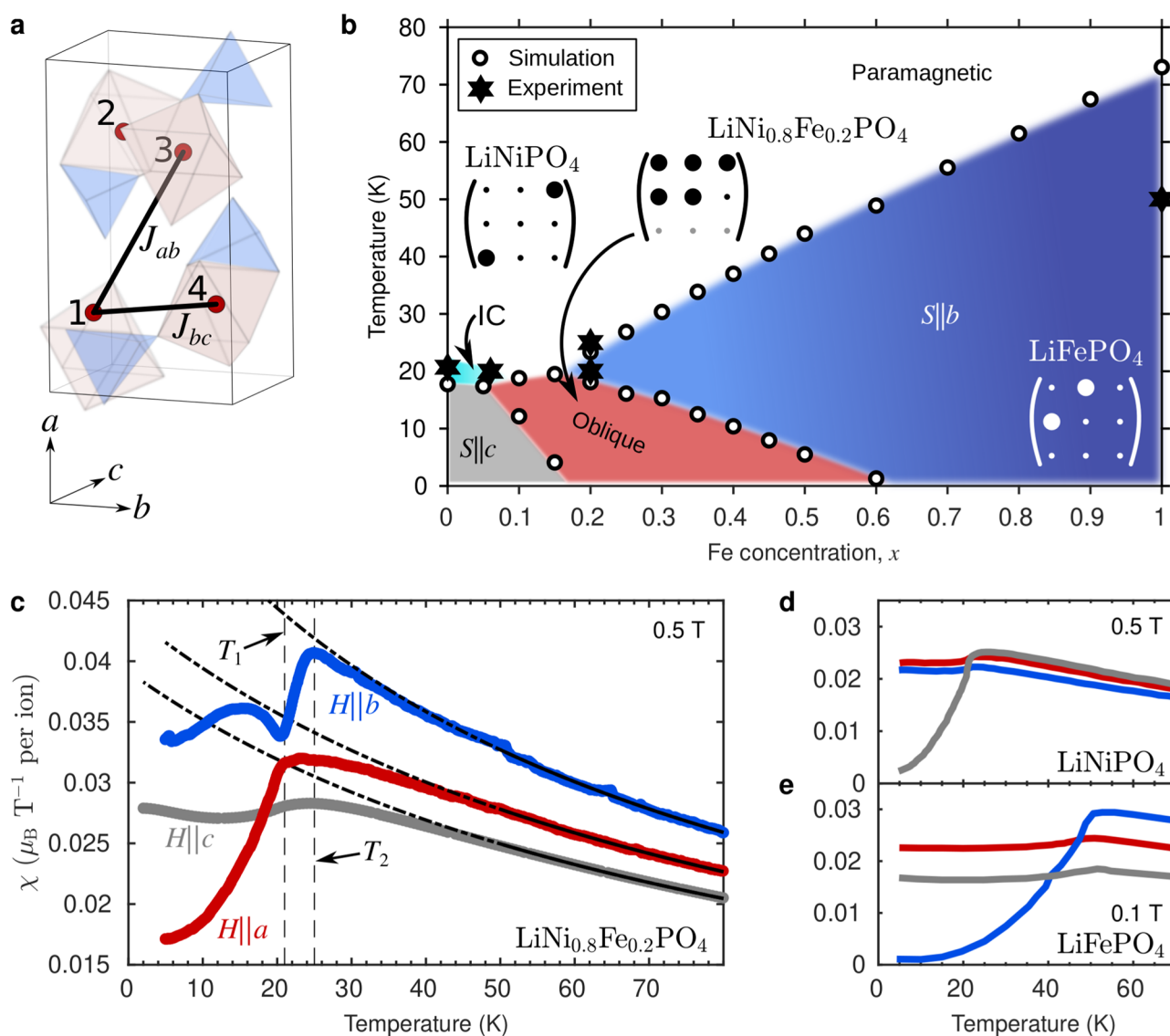


Fig. 1 | Unit cell, (x, T) phase diagram and magnetic susceptibility of $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$. **a** Crystallographic unit cell of LiMPO_4 with four magnetic ions (red numbered spheres) and the two most important exchange paths, J_{bc} and J_{ab} , shown. The MO_6 octahedra and PO_4 tetrahedra are illustrated with red and blue shading, respectively. **b** (x, T) phase diagram constructed from experimental data and simulation. The open circles correspond to phase transitions observed in the simulated specific heat. Filled stars represent phase transitions detected in magnetic susceptibility and neutron diffraction experiments for samples with $x = 0, 0.06, 0.20$ and 1. Both simulations and experiments reveal three phases: Commensurate phases with $S||c$ (grey shading) and $S||b$ (blue shading) are seen at small and large x , respectively, while an oblique phase is present in the range $0.1 < x < 0.6$ (red shading). For each phase, the observed form of the magnetoelectric tensor at low temperature is indicated. The gradient of the blue shading illustrates that the

ordered moment along b , $\langle S||b \rangle$, decreases when decreasing x while $\langle S||a \rangle \approx \langle S||c \rangle = 0$. For small x there exists an incommensurate (IC) phase in a narrow temperature interval above the commensurately ordered phase (cyan shading)^{21,22}. **c** Magnetic susceptibility of $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ measured with 0.5 T applied along the three crystallographic axes; a (red curve), b (blue curve) and c (grey curve). Fits to the Curie-Weiss law are shown with black lines. The solid parts of the lines indicate the fitted interval (50–300 K) and the dash-dotted parts are extrapolations to lower temperatures. The vertical dashed lines mark transitions at 21 and 25 K. **d, e** Corresponding susceptibilities for LiNiPO_4 and LiFePO_4 , measured with 0.5 and 0.1 T respectively. Reprinted with permission from refs. 32 and 54. Copyright (2023) by the American Physical Society. Error bars in all panels are smaller than symbol sizes.

crystallographic b and c axes for respectively LiFePO_4 and LiNiPO_4 . In LiNiPO_4 there exists in addition an incommensurate phase in a narrow temperature interval just above the Néel temperature^{21,22}. The static and dynamic properties of LiMPO_4 are well-described by the spin Hamiltonian

$$\hat{\mathcal{H}} = \sum_{(i,j)} J_{ij} \mathbf{S}_i \cdot \mathbf{S}_j + \sum_{i,\alpha} D_i^\alpha (S_i^\alpha)^2, \quad (1)$$

where the first sum accounts for the exchange interactions of magnitude J_{ij} between spins on sites i and j . The second sum over all sites i and three crystallographic directions, $\alpha = \{a, b, c\}$, reflects single-ion anisotropy energies, parameterized by the vector $\mathbf{D} = (D^a, D^b, D^c)$. This term is responsible for the distinct ordered moment direction selected upon ordering in stoichiometric LiNiPO_4 ²⁵ and LiFePO_4 ²⁶.

Here, we explore chemical tuning of mixed-anisotropy antiferromagnets as a novel route for tailoring the properties of single-phase magnetoelectrics. We have employed magnetic susceptibility and pyrocurrent measurements, neutron diffraction and Monte Carlo simulations to investigate the (x, T) phase diagram of $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$ (Fig. 1b). We observe three commensurate magnetic phases with propagation vector $\mathbf{k} = 0$. At low temperature and for $x < 0.2$, the spins order along c like in LiNiPO_4 . For $x > 0.6$, the spins order along b like in LiFePO_4 . For $x = 0.2$, two magnetic phases appear upon cooling²⁷. Neutron diffraction reveals ordered moments predominantly along the crystallographic b -axis below $T_2 = 25$ K, while below $T_1 = 21$ K, the moments partially reorient towards the a -axis in a low-temperature oblique phase. Our investigations of the field-induced polarization in these phases have uncovered a complex ME coupling scheme. The lowered magnetic symmetry of the oblique phase combined with the broken discrete translational symmetry, unlocks ME tensor elements that are otherwise forbidden in the parent compounds. Simulations show that the key factors responsible for the observed oblique phase are mismatched anisotropies combined with an inter-species exchange coupling creating competing exchange and single-ion anisotropy energy terms. This unusual mechanism is of general applicability and represents a promising approach to search for oblique ME phases in other families of compounds where the ME properties can be chemically tuned.

Results

Magnetic susceptibility

Figure 1c–e illustrate distinct differences in magnetic susceptibility between $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ and its parent compounds, LiNiPO_4 and LiFePO_4 . The susceptibility curves, χ_a , χ_b and χ_c , of both LiFePO_4 and LiNiPO_4 for fields along a , b and c display textbook behavior for antiferromagnets with easy axes along b and c , respectively. The component of χ parallel to the easy axis drops towards zero below the transition temperature while the two perpendicular components remain nearly constant. By contrast, the susceptibility of $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ shows clear evidence of two magnetic phase transitions. Below $T_2 = 25$ K, χ_b decreases while χ_a and χ_c remain constant. At a slightly lower temperature, $T_1 = 21$ K, χ_a begins to drop precipitously and the decrease of χ_b is interrupted, while χ_c remains approximately constant. These observations are indicative of a negligible c -axis component of the ordered moment at all temperatures, and of a rotation of the ordered moments from the b axis towards the a axis for temperatures lower than T_1 . These two transitions were previously reported and we compare our findings with those of the authors of ref. 27 later in the Results section. Note that overall the susceptibility of the mixed system is higher than for the parent compounds. This, together with the overall different temperature dependence of the susceptibility as compared to the parent compounds, is evidence that $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ is indeed a solid solution and we can exclude phase separation in the system.

Magnetic structures

To determine the magnetic structures in $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ we turn to neutron diffraction. At all temperatures below T_2 , the commensurate magnetic Bragg peaks were found to be resolution limited, implying long-range order (Supplementary Fig. 2d–f in the Supplementary Information). A representative selection of temperature-dependent integrated intensities as obtained at the diffractometer, E5, is shown in Fig. 2a. The intensity of each magnetic Bragg peak reflects different combinations of symmetry components of the magnetic order. In addition, it carries information about the spin orientation in the ordered states, because neutrons couple exclusively to components of the magnetic moment perpendicular to the scattering vector \mathbf{Q} (see Supplementary Table I). Our analysis indicates that the main magnetic structure component at all temperatures below T_2 is $(\uparrow\uparrow\downarrow\downarrow)$ with the numbering of spins defined in Fig. 1a. Rietveld refinement of the magnetic Bragg peak intensities at base temperature yields magnetic moments predominantly in the (a, b) -plane with major component along a . For $T_1 \leq T \leq T_2$, our data suggests moments aligned along b .

The two transitions observed in our susceptibility measurements have clear signatures in the diffraction data: The $(0, 0, -1)$ and $(3, 0, -1)$ reflections grow linearly with decreasing temperature below $T_2 \approx 25$ K. By contrast, the $(0, 1, 0)$ peak appears only below $T_1 \approx 21$ K where in addition, there is a kink in the temperature profile of the $(3, 0, -1)$ intensity. The temperature dependencies of all recorded peaks are well described by a combination of a linear function and a power law, reflecting the existence of two order parameters, below T_2 and T_1 , respectively (solid lines in Fig. 2a and in Supplementary Fig. 1). Simultaneous fits to all data sets yield transition temperatures $T_2 = 25.7(2)$ K and $T_1 = 20.8(1)$ K respectively, in good agreement with refs. 27 and 28. We note that the critical exponents for the two order parameters are clearly different. Below T_2 , the neutron intensity increases linearly with decreasing temperature which means a critical exponent of $\frac{1}{2}$ as assumed fixed in the fit. This corresponds to the critical exponent resulting from long-range interactions or from a secondary order parameter. At T_1 , $(0, 1, 0)$ displays a power law behavior with $\beta = 0.32(3)$ which is comparable to the critical exponent of a 3D Heisenberg, XY or Ising system.

To unambiguously determine the spin orientations, we performed a polarized neutron diffraction experiment using the triple axis spectrometer 4F1 and with scattering vector $\mathbf{Q} = (0, K, L)$ in the horizontal scattering plane. Uniaxial polarization analysis allows the two spin components perpendicular to \mathbf{Q} to be individually addressed. This is done by measuring spin-flip (SF) and non spin-flip (NSF) intensities for the neutron beam polarization along the scattering vector ($P||x$), perpendicular to \mathbf{Q} in the horizontal scattering plane ($P||y$), and along the direction perpendicular to the scattering plane ($P||z$). The temperature-dependencies of the resulting six cross sections were collected for the $(0, 1, 0)$, $(0, 0, 1)$ and $(0, 1, 2)$ reflections. The SF cross sections carry information on spin components perpendicular to both \mathbf{Q} and the neutron beam polarization \mathbf{P} . The NSF cross sections reveal spin components perpendicular to \mathbf{Q} but parallel to \mathbf{P} in addition to any finite nuclear Bragg peak intensity.

Noting that the $(0, 1, 0)$ magnetic peak exclusively reflects $(\uparrow\uparrow\downarrow\downarrow)$ symmetry components (Supplementary Table I), Fig. 2b, c show that the magnetic structure below T_1 involves sizeable spin components along a , but only negligible c -axis components. Spin components parallel to b do not contribute to magnetic scattering at $\mathbf{Q} = (0, 1, 0)$, but can be probed at $\mathbf{Q} = (0, 0, 1)$ or $(0, 1, 2)$. Figure 2d, e confirm the involvement of an a -axis spin components below T_1 , and show that the scattering is dominated by spins oriented along b in the range $T_1 \leq T \leq T_2$. Note that here we plot only data for $(0, 1, 0)$ and $(0, 0, 1)$ as their interpretation is straightforward. The data for $(0, 1, 2)$ is shown in Supplementary Fig. 2a, b. A comparison of the observed intensities to the structure factors for the magnetic symmetry components contributing to the $(0, 1, 0)$, $(0, 0, 1)$ and $(0, 1, 2)$ peaks makes it clear

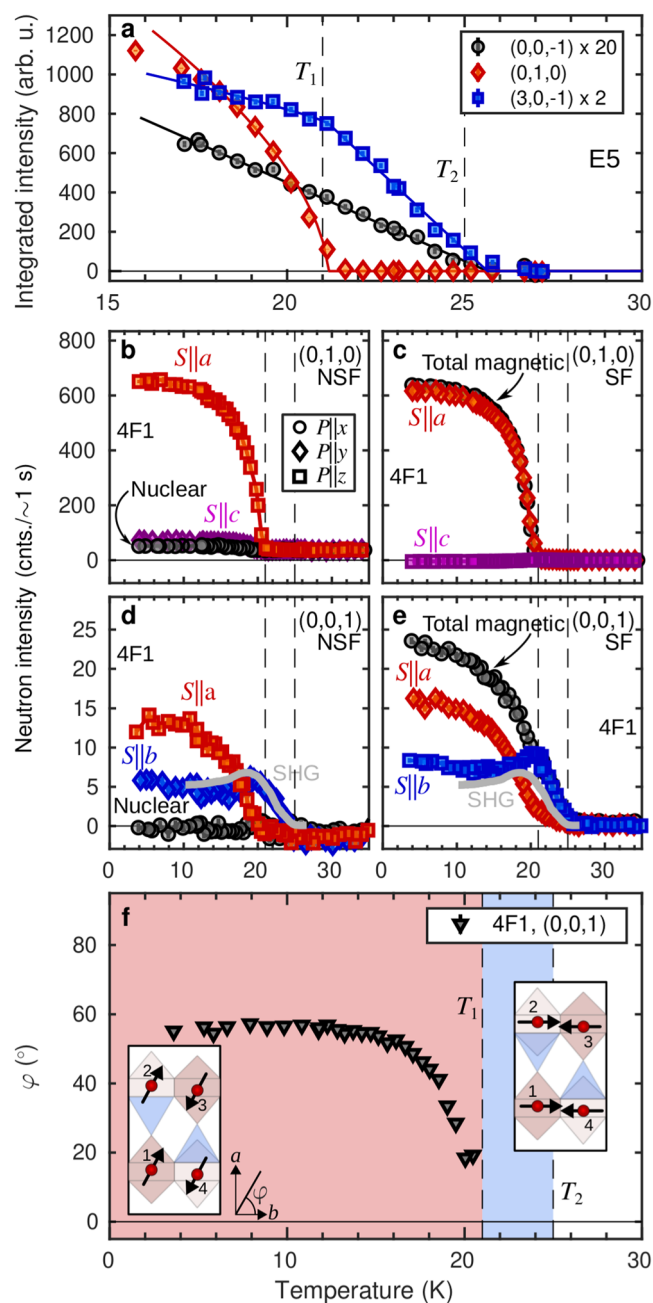


Fig. 2 | Neutron diffraction. **a** Temperature profiles of neutron diffraction intensities for three magnetic Bragg peaks for $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$. Circle, diamond and square symbols denote $(0, 0, -1)$, $(0, 1, 0)$ and $(3, 0, -1)$ respectively. Intensities are scaled as indicated in the legend. The solid lines show the fits described in the main text. Peak intensities obtained with polarized neutrons in the NSF (**b** and **d**) and SF (**c** and **e**) channels after correcting for non-perfect beam polarization⁵⁵. Circle, diamond and square symbols are for neutron polarizations along x , y and z respectively. The symbols are coloured black for nuclear or total magnetic scattering, red for $S||a$, blue for $S||b$ and purple for $S||c$. The crystal was oriented with the a axis vertical such that NSF intensities for $P||z$ reflect a axis spin components for both $(0,1,0)$ and $(0,0,1)$. Grey curves in **d**, **e** show optical second harmonic generation measurements reprinted with permission from ref. 27. Copyright (2023) by the American Physical Society. **f** Moment rotation angle, φ , as a function of temperature with insets showing projections in the $(0, 0, 1)$ plane of the magnetic structures for $T \leq T_1$ and $T_1 \leq T \leq T_2$. The red and blue shadings illustrate the extends of the two respective phases. The two transitions at T_1 and T_2 are marked by vertical dashed lines in all panels. Error bars of the neutron counts, N , follow Poisson statistics as \sqrt{N} and the errors in **f** are propagated from the neutron counts. Error bars are smaller than symbol sizes in all panels.

that the dominant symmetry component for $T_1 \leq T \leq T_2$ is also $(\uparrow\uparrow\downarrow\downarrow)$. The scattering from b -axis spin components, reflected by the NSF, $P||y$ and SF, $P||z$ cross sections in Fig. 2d, e increases monotonically for temperatures in the range $T_1 \leq T \leq T_2$ and levels off to a finite value at our experimental base temperature. The rotation angle, φ , in the (a, b) -plane may be calculated from the ratio of $P||y$ and $P||z$ data in Fig. 2d, e leading to the conclusion that the angle between the moments and the b axis approaches $\varphi = 60^\circ$ at low temperatures (Fig. 2f).

The small but finite nuclear intensity for $P||x$ in Fig. 2b and Supplementary Fig. 2a may be due to a change of the lattice symmetry which could be caused by magnetostriction. Magnetostriction is common in magnetoelectrics and for LiFePO_4 this effect has been observed when applying magnetic fields²⁹. Future synchrotron X-ray studies will uncover the evolution of the crystal lattice and symmetry as a function of temperature.

The solid grey lines in Fig. 2d, e represent the intensity of the second harmonic generation (SHG) susceptibility tensor element, χ_{xxx} , from ref. 27. Here the first subscript signifies the component of the non-linear polarization induced by an electric field with components denoted by the last two subscripts. The similarity of the SHG signal with the NSF, $P||y$ and SF, $P||z$ cross sections is clear evidence that these two observations are intimately related. The SHG data was interpreted by the authors of ref. 27 as a signature of spin rotation from the easy b axis of stoichiometric LiFePO_4 towards the easy c axis of stoichiometric LiNiPO_4 , upon cooling below T_1 . Our polarized neutron diffraction results only allow for a small spin component along c and show instead a sizeable component along a . This picture is consistent with the susceptibility data in Fig. 1c. The physical mechanism for this surprising reorientation away from the easy axes of the two parent compounds is explored in our Monte Carlo simulations to be presented further on, but first we look into its profound consequences for the ME coupling.

Magnetoelectric effect

The linear ME effect is described by the relation $\mathbf{P}^E = \alpha \mathbf{H}$ between the components of the induced electrical polarization, \mathbf{P}^E , and those of the applied magnetic field, \mathbf{H} . A related equation, $\mu_0 \mathbf{M} = \alpha^T \mathbf{E}$, connects the components of the induced magnetization, \mathbf{M} , to those of the applied electric field, \mathbf{E} . For systems invariant to integer lattice vector translations, the allowed elements of the ME tensor α are imposed by the point group symmetry of the magnetically ordered state^{14,15}. Specifically, for the stoichiometric parent compounds LiNiPO_4 and LiFePO_4 , the reported magnetic structures imply that the elements which may be non-zero are α_{ac} , α_{ca} and α_{ab} , α_{ba} , respectively.

The ME response of $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ was probed with measurements of the pyrocurrent produced by a temperature change (see Methods and Supplementary Information for details). Our results for $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ are shown in Fig. 3 and are compared to the ME response of the parent compounds, LiNiPO_4 ¹⁹ and LiFePO_4 ³⁰. Note that in the following analysis we assume space group $Pnma$ although it was recently shown that LiFePO_4 may display a lower symmetry³⁰. The pyrocurrent for $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ for two orthogonal orientations of the electric poling field, \mathbf{E} , and three directions of the magnetic field shows clear signatures of two ME phase transitions slightly below T_2 and T_1 , see Fig. 3a–c. The evidence is in the form of spikes in the pyrocurrent, which following a geometrical correction can be integrated to obtain the temperature dependent polarization components, P_i^E . A signal is thus observed for all probed couplings except α_{bc} .

The electric polarization corresponding to the tensor elements α_{ab} and α_{ba} together with that corresponding to α_{ac} are shown in Fig. 3d. As mentioned above, these components are known to be non-zero for stoichiometric LiFePO_4 and LiNiPO_4 ^{23,31}. When comparing the ME response of $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ to that of LiFePO_4 and LiNiPO_4 measured under identical conditions (blue and grey dashed lines in Fig. 3d), it is apparent that the polarizations induced along a and b are

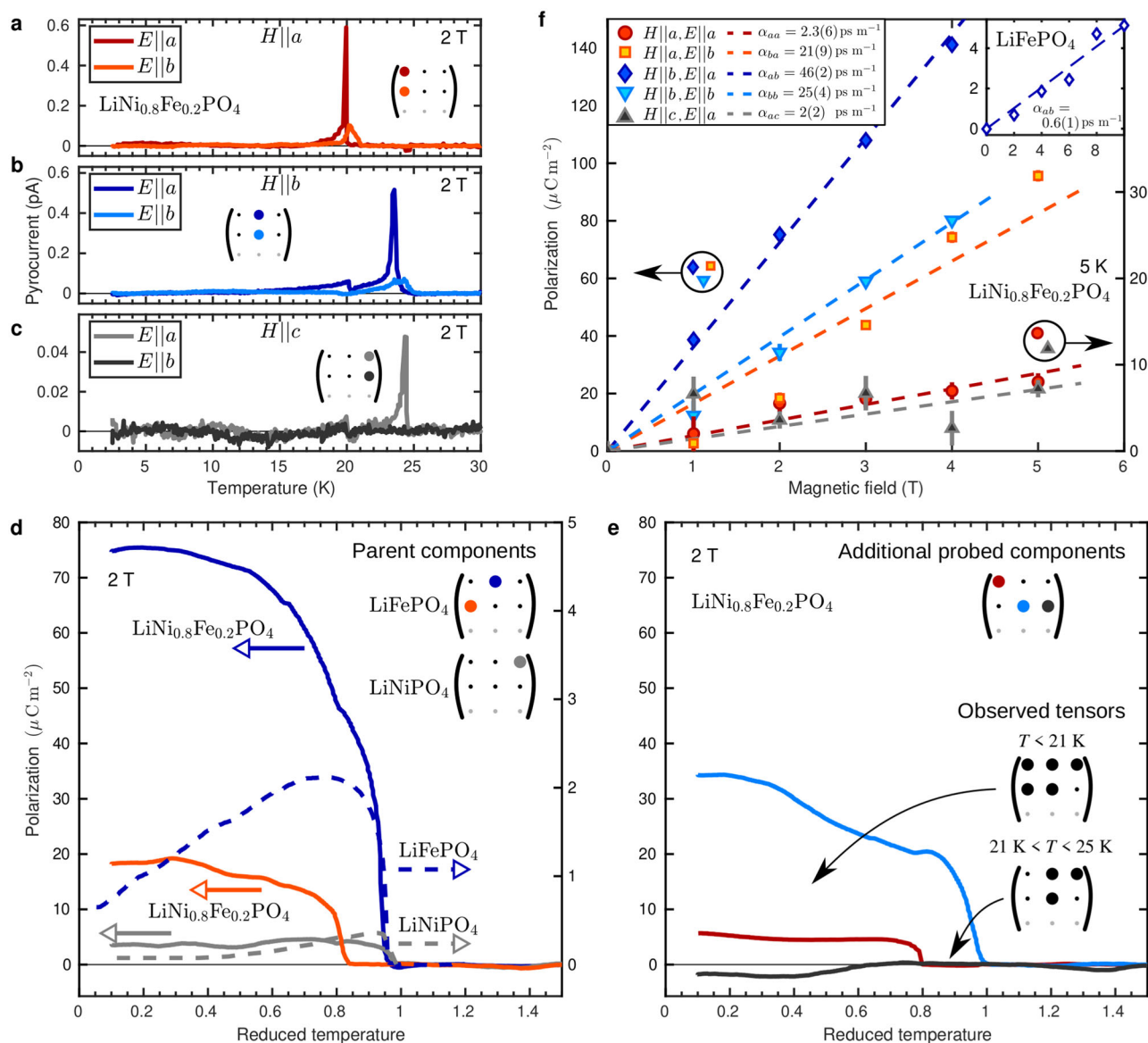


Fig. 3 | Pyrocurrent and magnetoelectric effect. Panels **a–c** show the pyrocurrent for $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ as a function of temperature for magnetic fields applied along *a* (red and orange curves), *b* (dark and light blue curves) and *c* (grey and black curves), respectively. The insets indicate which elements of the ME tensor, α , were probed. The error on the measured current is of the order of 5 fA. The colour codes given in **a–c** are followed in the remaining panels of this figure. **d** Electric polarization as a function of the reduced temperature with transition temperatures 21, 25 and 50 K at zero field for LiNiPO_4 , $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ and LiFePO_4 , respectively. Note the two y-axes: the left one for the data for the mixed system (solid lines) and the right one for the parent compounds (dashed lines). The curve shown for LiNiPO_4 is from ref. 19 and that for LiFePO_4 is reprinted with permission from ref. 30. Copyright (2023) by the American Physical Society. **e** Temperature dependency of the

electric polarization originating from tensor elements not present in the parent compounds. For $T < T_1$ and $T_1 < T < T_2$, all observed non-zero ME tensor elements in $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ are indicated. The measurements shown in **a–e** were carried out with an applied magnetic field strength of 2 T where the ME effect is still linear. The errors on the polarization are of the order of $1 \mu\text{C m}^{-2}$. **f** Field dependency of the average of the induced electric polarization for $T < 5$ K for non-zero couplings. The error bars are estimated from the variations observed in the temperature profiles of the polarization (see Supplementary Fig. 4). Note that strong and weak ME components are plotted on two different y-axes as illustrated with encircled symbols and arrows. The dashed lines are linear fits, $P_i^E = \alpha_{ij}H_j$, to the data with the obtained ME coefficients, α_{ij} , listed in the legend. The inset shows the corresponding data for LiFePO_4 with $H \parallel b$, $E \parallel a$.

significantly larger in $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ at all temperatures below the transition temperature. Most strikingly, in the limit $T \rightarrow 0$, the polarization due to the dominant tensor component, α_{ab} is increased by almost two orders of magnitude compared to LiFePO_4 . A second remarkable observation is that the onset temperatures of α_{ab} and α_{ba} are different. α_{ba} vanishes in the range $T_1 \leq T \leq T_2$ whereas α_{ab} is finite already below T_2 and displays a kink at T_1 .

Finally, in Fig. 3e we probe tensor components that are by symmetry not allowed for LiNiPO_4 and very small for LiFePO_4 ³⁰. Similarly, we observe α_{bb} below T_2 while α_{aa} is finite only below T_1 . For the last tensor

element measured, α_{bc} , there is no spike to be seen in the pyrocurrent and we conclude that this element is either very weak or zero.

By measuring the pyrocurrent at different magnetic field strengths, we obtain the electric polarization as a function of field as shown in Fig. 3f. The values of the polarization shown here are the mean values for temperatures below 5 K. For the corresponding polarization curves at the different field strengths, see Supplementary Fig. 4. The measured polarization is linear with field for most couplings, except notably for α_{ba} . Interestingly, α_{ba} is exactly the component with a different onset temperature compared to α_{ab} , underlining that the behavior of the ME

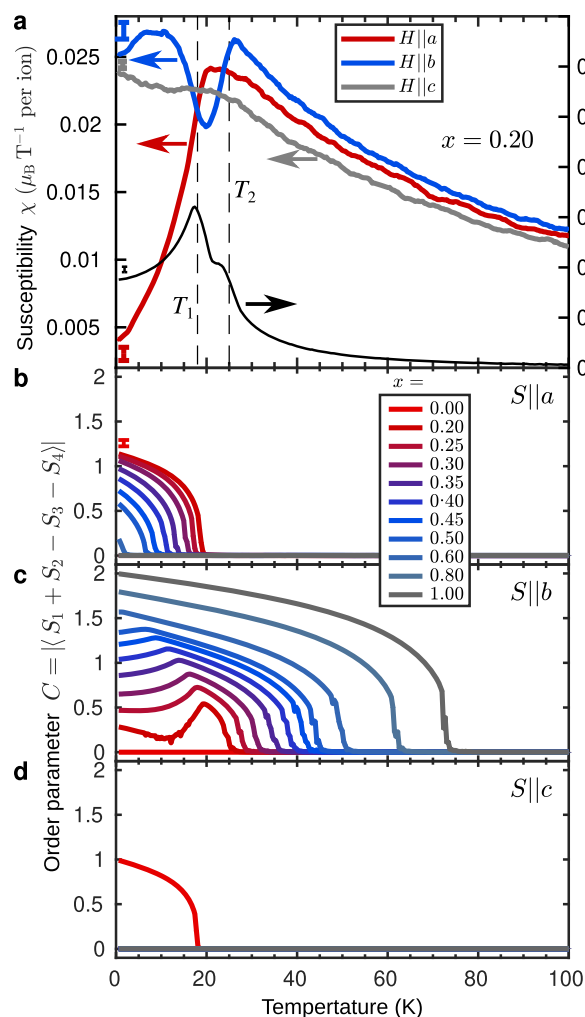


Fig. 4 | Monte Carlo simulations. **a** Magnetic susceptibility (left axis) for fields along a (red curve), b (blue curve) and c (grey curve) and specific heat (right axis, black curve) for $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$ with $x = 0.20$. **b–d** The absolute value of the C-type order parameter plotted as a function of temperature for spin components along the three crystallographic axes and for different values of x . The colours of the curves illustrate the Fe concentration from red ($x = 0$) over blue to grey ($x = 1$). Note that in **d** the curves for $x \geq 0.20$ coincide. The maximum error bar sizes for each of the variables are indicated at the lowest temperatures in **a** and **b**.

effect in $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$ is complex. Nevertheless, it is clear from our measurements that the effect is much stronger in the mixed system compared to the parent compounds and that at low field strengths, the system is in the linear regime.

Monte Carlo simulations. We now show that classical Monte Carlo simulations reproduce the salient features of the susceptibility and diffraction results for $x = 0.20$. In the calculations, we chose $J_{bc} = 1$ meV and $J_{ab} = 0.3$ meV for all corresponding pairs of sites (see Fig. 1a), irrespective of their occupancy by Ni or Fe. For the single-ion anisotropies we used $(D^a, D^b, D^c) = (0.3, 1.8, 0)$ meV for Ni with $S = 1$ and $(D^a, D^b, D^c) = (0.6, 0, 1.6)$ meV for Fe with $S = 2$. This choice of parameters looks like a dramatic simplification when compared to the full set of experimentally determined parameters^{25,26,32} for LiNiPO_4 and LiFePO_4 as given in Supplementary Table IV. Nevertheless, the set of simplified parameters yields results in good agreement with our experimental neutron diffraction and susceptibility measurements for $x = 0.20$. In turn, this agreement justifies the use of the Hamiltonian from Equation (1) with the chosen parameters for an exploration of the full phase diagram of $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$.

In Fig. 4a, we plot the calculated magnetic susceptibility and specific heat for $x = 0.20$ as a function of temperature. Two phase transitions are observed near 25 and 20 K. The transition temperatures as well as the temperature-dependencies of the three components of χ are in excellent agreement with the experimental results shown in Fig. 1c. The accuracy of the simulations is further illustrated by comparing the calculated and measured susceptibilities for $x = 0.06$ with the corresponding experimental data (Supplementary Fig. 5).

Next, we study the simulated C-type order parameter $C = |\langle S_1 + S_2 - S_3 - S_4 \rangle|$ for spin components along a , b and c , respectively. Figure 4b–d show the temperature dependencies of C for the full range of Fe concentrations, x . Focusing on $x = 0.20$, the resemblance with the polarized neutron diffraction data in Fig. 2 is striking. Note that the neutron diffraction intensity is proportional to the moment squared. The first phase transition at 25 K corresponds to spins ordering along b . The growth of the corresponding order parameters is interrupted at 20 K where a rotation towards a starts and the oblique low-temperature phase is entered. The c -axis component remains zero at all temperatures. From the a and b components of the simulated order parameter we arrive at a rotation angle of 76° at low temperature, which compares reasonably well with the value of $\approx 60^\circ$ obtained from the experimental data (Fig. 2f). We use the transition temperatures derived from the simulated specific heat and order parameters data to construct the (x, T) phase diagram shown in Fig. 1b. The simulations underestimate the transition temperatures for small x compared to the measured values (star symbols for $x = 0, 0.06$ and 0.20), but the ratios of simulated to observed transition temperatures are relatively constant with x in this range.

The oblique antiferromagnetic state is relatively robust. The simulations show that the only requirements are an inter-species exchange interaction as well as competing single-ion anisotropies with opposite easy and hard axes for the parent compounds but a common intermediate axis off any of these easy and hard axes. It is this frustration between exchange and single-ion anisotropy energies that generates the oblique state. In the analysis of the neutron diffraction data we assumed a collective behavior of all spins regardless of species. The simulations show that indeed the ensemble average of the moments give a collectively ordered picture. However, we also find local fluctuations between Ni and Fe sites (Supplementary Fig. 6). The ordered moment for the oblique phase is therefore lower when calculating the average over the entire system than when considering individual sites, not only due to thermal fluctuations but also due to site specific differences in the moment orientation. This effect and the general consequences of violation of discrete translational symmetry for the ME effect are an interesting topic of future theoretical investigations.

Discussion

The effects on the magnetic ground state of a quenched random distribution of ions in mixed anisotropy antiferromagnets have been extensively studied by renormalization group theory and in various mean field models^{16–18,33–36}. The resulting phase diagrams generically contain one or more oblique phases at intermediate compositions, in which the ordered moments are oriented away from the easy axes of the parent compounds. Depending on the details of the exchange and anisotropy terms in the spin Hamiltonian, the oblique ground state may involve ordered moments in a plane spanned by the easy axes of the two parent compounds, or perpendicular to both in the particular case where the easy and hard axis of one species coincide with the hard and easy axes of the other species¹⁷. These predictions of a chemically tunable magnetic ground state are broadly consistent with our experimental observations and Monte Carlo simulations for $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$, and have in the past been found to agree well with experimental studies of other mixed-anisotropy antiferromagnets^{37–42}.

In the lithium orthophosphates, the magnetic C-type structure is the dominant structure component for all stoichiometric family

members. The single-ion anisotropy plays a crucial role for the ME coupling, as the allowed tensor elements derive from the magnetic point group, ipso facto the direction of the ordered moments. Since $S||b$ in LiFePO_4 , α_{ab} and α_{ba} are allowed, while as $S||c$ in LiNiPO_4 , α_{ac} and α_{ca} are allowed. When $S||a$, the diagonal elements are allowed (α_{aa} , α_{bb} , $\alpha_{cc} \neq 0$). It follows from our results, however, that such simple rules do not apply in the mixed systems, where discrete translational symmetry is broken and the local spin anisotropy is site dependent. Between 21 and 25 K, the predominant spin orientation in $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ is $S||b$. Nevertheless, the observed diagonal tensor element α_{bb} is almost as strong as the expected α_{ab} -component, while $\alpha_{ba} = 0$. Below 21 K, the ME-tensor is more complex and most tensor elements are observed. This is due to the off-axis direction of the ordered moments. However, the fact that some expected tensor elements are extinct between 21 and 25 K, while some unexpected elements are not, is a strong hint that while discrete translational symmetry is broken, the existence of the ME coupling is still governed by magnetic point group symmetry, yet not in the same manner as in the stoichiometric systems. This is an interesting point in itself and should be subject to further theoretical study.

The most intriguing observation is the almost hundredfold increase in the strength of the ME coupling for $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ observed in the pyrocurrent measurements. In LiNiPO_4 and LiFePO_4 , the effect is believed to arise from exchange striction^{22,26,43,44}. Specifically, a microscopic model connects the electric polarization caused by a displacement, x_i , of PO_4 tetrahedra along i , to the component of the applied magnetic field along j as follows:

$$P_i^E \propto x_i = \frac{\lambda_i}{\epsilon_i} \langle S \rangle^2 \chi_j H_j. \quad (2)$$

Here $\langle S \rangle$ is the order parameter, χ_j the magnetic susceptibility for fields along j , ϵ_i the elastic energy constant associated with tetrahedron displacement ($\mathcal{E}_i = \epsilon_i x_i^2$) and λ_i reflects the strength of the exchange striction ($J_{H \neq 0} \rightarrow J_{H=0} + \lambda_i x_i$). In addition to the general increase in magnetic susceptibility in $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ as compared to the parents, both a reduction of the elastic displacement energy and an increase in exchange striction could cause stronger ME couplings. A lowering of crystal symmetry may indeed result in lower energy cost for displacing the PO_4 tetrahedra as well as more allowed options for displacement directions, i.e. a softer lattice. Moreover, the local number of nearest neighbors of a given species, and variations thereof, could bring exchange striction terms into play which would otherwise cancel out (in the parent compounds only interactions between ion pairs (1,2) and (3,4) contribute to the ME effect²²).

Tuning of the magnetic symmetry was also recently achieved in the olivine series of compounds, $\text{Li}_{1-x}\text{Fe}_x\text{Mn}_{1-x}\text{PO}_4$ ⁴⁵, as well as in Mn and Co doped LiNiPO_4 ⁴⁶. These studies do not report on the corresponding consequences for the ME effect but they further illustrate that the lithium orthophosphate family harbour many possibilities for tailoring the magnetic and consequently also ME material properties. In general, the existence of ME and multiferroic oblique antiferromagnets is unlikely to be limited to this family. Notably, our Monte Carlo simulations show that competing single-ion anisotropies with a common intermediate axis combined with a significant inter-species exchange coupling are the decisive ingredients to produce an oblique magnetoelectric state, whereas details of the exchange coupling scheme play almost no role. Generally, transition metal ions exhibit complex single-ion anisotropies in octahedral environments, and will likely share an intermediate anisotropy axis in many families of compounds. An oblique ME phase may therefore also exist in other classes of materials. In future studies, Monte Carlo simulations similar to those performed in this work, possibly combined with DFT calculations to determine exchange constants, could precede time-demanding material synthesis in order to predict the viability of the candidate family to produce oblique ME phases. An interesting family of

compounds for future studies of this type is for example $\text{Mn}_{1-x}\text{M}_x\text{WO}_4$ ($M = \text{Fe, Co, Cu, Zn}$)^{47–50}.

In summary, we studied $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ experimentally using magnetometry, neutron diffraction and pyrocurrent measurements and theoretically through Monte Carlo simulations. We have identified an oblique low-temperature phase over an extended range of compositions. In this phase the spins rotate away from the distinct easy axes of the parent compounds, LiNiPO_4 and LiFePO_4 , towards the direction of the common intermediate axis. The magnetoelectric properties correlate with the observed magnetic phase transitions, but the form of the magnetoelectric tensor departs from theoretical expectations based on discrete translational invariance and magnetic point group symmetries. Most dramatically, we observe a strong enhancement of almost two orders of magnitude for the dominant magnetoelectric tensor element compared to the parent compounds. These data in combination with our Monte Carlo simulation results suggest that the observations have broader implications and that chemical tuning of oblique magnetoelectric phases represents a promising path for tailoring magnetoelectric material properties.

Methods

Magnetic susceptibility

Magnetization measurements were performed using a Cryogenic Ltd. cryogen free vibrating sample magnetometer. A magnetic field of 0.5 T was applied along a and b for temperatures in the range 2–300 K. For measurements with field along c we used a Physical Property Measurement System (PPMS) from Quantum Design. The sample for all measurements of $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ was a 25 mg box-shaped single crystal.

Neutron diffraction

Unpolarized neutron diffraction data on $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ were collected at the E5 diffractometer at the Helmholtz-Zentrum Berlin using an 4-circle cradle, a neutron wavelength of $\lambda = 2.38 \text{ \AA}$ selected by a pyrolytic graphite (PG) monochromator and a 2D position sensitive neutron detector of size $9 \times 9 \text{ cm}^2$. A PG filter before the sample position was used for suppression of second order neutrons ($\lambda/2$) from the monochromator.

The polarized neutron diffraction data were obtained on the 4F1 triple-axis spectrometer at the Laboratoire Léon Brillouin, using a wavelength of $\lambda = 2.44 \text{ \AA}$. The monochromatic incident beam was polarized using a bending mirror and a PG filter after the bender reduced second order contamination of the incident beam. A spin flipper was placed before the sample position. In combination with a Heusler analyzer this allowed for probing both spin flip and non spin-flip scattering. A set of Helmholtz-like coils around the sample position enabled polarization of the neutron beam along x , y or z in the coordinate system described in the Supplementary Information. The same high-quality 250 mg $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ single crystal was used in both polarized and unpolarized neutron diffraction experiments. At 4F1, it was oriented with $(0, K, L)$ in the horizontal scattering plane. Flipping ratios $F = 40$ and 25 were deduced from measurements on the structural (020) and (002) reflections, and used to correct for non-ideal beam polarization.

Preliminary studies of the magnetic structures of $\text{LiNi}_{1-x}\text{Fe}_x\text{PO}_4$ were carried out at the triple-axis spectrometer RITA-II and diffractometer TriCS at the Paul Scherrer Institute.

Pyrocurrent measurements

The quasi-static method⁵¹ was used to perform pyrocurrent measurements with a Quantum Design PPMS at the Helmholtz-Zentrum Berlin. The custom-build insert allows for sample rotations with respect to the vertical magnetic field and thus enables probing off-diagonal ME tensor elements^{30,52}. Plate-shaped $\text{LiNi}_{0.8}\text{Fe}_{0.2}\text{PO}_4$ single crystals with gold sputtered faces perpendicular to a and b , and sample thickness 0.5 and 0.9 mm, respectively, were used. Single crystals of LiNiPO_4 and LiFePO_4

were similarly prepared with faces perpendicular to a and thickness 0.6 and 0.5 mm, respectively. The precision of the cuts were within 0.5° which determines the uncertainty of the field direction with the respect to the axes perpendicular to the surface. For the other field directions, the crystals were aligned by eye to obtain a sample alignment within 2° . A potential of 100 V was applied as well as a magnetic field while cooling to obtain a single ferroelectric domain. The potential was switched off at the experimental base temperature. The measurement was then performed upon heating at a constant temperature rate. Magnetic fields were applied along the a , b and c directions.

Monte Carlo simulations

Classical Monte Carlo simulations were carried out using the spin Hamiltonian [Equation (1)] and employing the Metropolis algorithm⁵³. The system was of size $10 \times 10 \times 10$ crystallographic unit cells (i.e. 4000 magnetic sites) and with Fe and Ni ions randomly distributed. The simulation was run for 10^5 Monte Carlo steps for each temperature in the range 1–100 K with step size 0.35 K and decreasing temperature. For each temperature we use the final configuration from the previous temperature step as a starting point. This procedure mimics the process of slowly lowering the temperature in the physical experiment. All simulations were run at zero field.

For each value of the Fe concentration, x , we simulated 5 distinct configurations from which we calculated the magnetic susceptibility, specific heat and order parameter. The susceptibility is calculated as $\chi = \beta(\langle M^2 \rangle - \langle M \rangle^2)$, where M denotes the total magnetization of the system, $\beta = \frac{1}{k_B T}$ and k_B is the Boltzmann constant. The brackets, $\langle \rangle$, denote the ensemble average over system configurations. In Fig. 4a and in Supplementary Fig. 5b we show χ normalized per magnetic site. The specific heat is calculated from the energy dissipation theorem, $C_V = k_B \beta^2 (\langle E^2 \rangle - \langle E \rangle^2)$, where E is the total energy of the system. The order parameter is calculated as an average over all unit cells, each containing four magnetic sites, i.e. $C = |\langle S_1 + S_2 - S_3 - S_4 \rangle|$. The curves shown in Fig. 4b–d are then the average quantities over the 5 configurations.

Data availability

The data that support the findings of this study are available at <https://doi.org/10.5281/zenodo.7515107> and from E.F. upon request. Source data are provided with this paper.

Code availability

The Monte Carlo code used in this study is available from E.F. upon reasonable request.

References

- Spaldin, N. A. & Ramesh, R. Advances in magnetoelectric multiferroics. *Nat. Mater.* **18**, 203 (2019).
- Liang, X., Chen, H. & Sun, N. X. Magnetoelectric materials and devices. *APL Mater.* **9**, 041114 (2021).
- Pradhan, D. K., Kumari, S. & Rack, P. D. Magnetoelectric composites: applications, coupling mechanisms, and future directions. *Nanomaterials* **10**, 2072 (2020).
- Ba, Y. et al. Electric-field control of skyrmions in multiferroic heterostructure via magnetoelectric coupling. *Nat. Commun.* **12**, 322 (2021).
- Manipatruni, S. et al. Scalable energy-efficient magnetoelectric spin-orbit logic. *Nature* **565**, 35 (2019).
- Trier, F. et al. Oxide spin-orbitronics: spin-charge interconversion and topological textures. *Nat. Rev. Mater.* **7**, 258 (2022).
- Nan, T. et al. Acoustically actuated ultra-compact nems magnetoelectric antennas. *Nat. Commun.* **8**, 296 (2017).
- Gu, Y. et al. Mini review on flexible and wearable electronics for monitoring human health information. *Nanoscale Res. Lett.* **14**, 263 (2019).
- Dong, C. et al. A portable very low frequency (vlf) communication system based on acoustically actuated magnetoelectric antennas. *IEEE Antennas Wirel. Propag. Lett.* **19**, 398 (2020).
- Hu, J.-M., Li, Z., Chen, L.-Q. & Nan, C.-W. High-density magnetoresistive random access memory operating at ultralow voltage at room temperature. *Nat. Commun.* **2**, 553 (2011).
- Liu, M. et al. Voltage tuning of ferromagnetic resonance with bistable magnetization switching in energy-efficient magnetoelectric composites. *Adv. Mater.* **25**, 1435–1439 (2013).
- Matsukura, F., Tokura, Y. & Ohno, H. Control of magnetism by electric fields. *Nat. Nanotechnol.* **10**, 209 (2015).
- Eerenstein, W., Mathur, N. D. & Scott, J. F. Multiferroic and magnetoelectric materials. *Nature* **442**, 759 (2006).
- Rivera, J.-P. A short review of the magnetoelectric effect and related experimental techniques on single phase (multi-) ferroics. *Eur. Phys. J. B* **71**, 299–313 (2009).
- Schmid, H. Some symmetry aspects of ferroics and single phase multiferroics. *J. Phys.: Condens. Matter* **20**, 43 (2008).
- Fishman, S. & Aharony, A. Phase diagrams of multicritical points in randomly mixed magnets. i. mixed anisotropies. *Phys. Rev. B* **18**, 3507 (1978).
- Oguchi, T. & Ishikawa, T. Theory of a mixture of two anisotropic antiferromagnets with different easy axis. *J. Phys. Soc. Jpn.* **45**, 1213 (1978).
- Mano, H. Possible phase diagrams of systems with competing anisotropies. *Prog. Theo. Phys. Suppl.* **101**, 597 (1990).
- Mercier, M. Ph.d. thesis. (Université de Grenoble, 1969).
- Santoro, R. P., Segal, D. J. & Newnham, R. E. Magnetic properties of LiCoPO_4 and LiNiPO_4 . *J. Phys. Chem. Solids* **27**, 1192 (1966).
- Vaknin, D., Zarestky, J. L., Rivera, J.-P. & Schmid, H. Commensurate-incommensurate magnetic phase transition in magnetoelectric single crystal LiNiPO_4 . *Phys. Rev. Lett.* **92**, 207201 (2004).
- Jensen, T. B. S. et al. Field-induced magnetic phases and electric polarization in LiNiPO_4 . *Phys. Rev. B* **79**, 092413 (2009).
- Santoro, R. P. & Newnham, R. E. Antiferromagnetism in LiFePO_4 . *Acta Cryst.* **22**, 344 (1967).
- Li, J., Garlea, V. O., Zarestky, J. L. & Vaknin, D. Spin-waves in antiferromagnetic single-crystal LiFePO_4 . *Phys. Rev. B* **73**, 024410 (2006).
- Jensen, T. B. S. et al. Anomalous spin waves and the commensurate-incommensurate magnetic phase transition in LiNiPO_4 . *Phys. Rev. B* **79**, 092413 (2009).
- Toft-Petersen, R. et al. Anomalous magnetic structure and spin dynamics in magnetoelectric LiFePO_4 . *Phys. Rev. B* **92**, 024404 (2015).
- Zimmermann, A. S., Sondermann, E., Li, J., Vaknin, D. & Fiebig, M. Antiferromagnetic order in $\text{Li}(\text{Ni}_{1-x}\text{Fe}_x)\text{PO}_4$ ($x = 0.06, 0.20$). *Phys. Rev. B* **88**, 014420 (2013).
- Li, J. et al. Tweaking the spin-wave dispersion and suppressing the incommensurate phase in LiNiPO_4 by iron substitution. *Phys. Rev. B* **79**, 174435 (2009).
- Werner, J. et al. High magnetic field phase diagram and failure of the magnetic Grüneisen scaling in LiFePO_4 . *Phys. Rev. B* **99**, 214432 (2019).
- Fogh, E. et al. The magnetoelectric effect in LiFePO_4 – revisited. *Phys. B* **648**, 414380 (2022).
- Mercier, M., Bauer, P. & Fouilleux, B. Mesures magnétoélectriques sur LiFePO_4 . *C. R. Acad. Sc. Paris B* **267**, 1345 (1968).
- Toft-Petersen, R. et al. High-field magnetic phase transitions and spin excitations in magnetoelectric LiNiPO_4 . *Phys. Rev. B* **84**, 054408 (2011).
- Aharony, A. Tetracritical points in mixed magnetic crystals. *Phys. Rev. Lett.* **34**, 590 (1975).
- Lindgaard, P.-A. Theory of random anisotropic magnetic alloys. *Phys. Rev. B* **14**, 4074 (1976).

35. Matsubara, F. & Inawashiro, S. Mixture of two anisotropic antiferromagnets with different easy axis. *J. Phys. Soc. Jpn.* **42**, 1529 (1977).
36. Mano, H. Pair spin theory of a random mixture with competing ising and xy anisotropies. *J. Phys. Soc. Jpn.* **55**, 908 (1986).
37. Wong, P., Horn, P. M., Birgeneau, H. J., Safinya, C. H. & Shirane, G. Competing order parameters in quenched random alloys: $\text{Fe}_{1-x}\text{Co}_x\text{Cl}_2$. *Phys. Rev. Lett.* **45**, 1974 (1980).
38. Katsumata, K., Kobayashi, M., Sato, T. & Miyako, Y. Experimental phase diagram of a random mixture of two anisotropic antiferromagnets. *Phys. Rev. B* **19**, 2700 (1979).
39. Tawaraya, T., Katsumata, K. & Yoshizawa, H. Neutron diffraction experiment on a randomly mixed antiferromagnet with competing spin anisotropies. *J. Phys. Soc. Jpn.* **49**, 1299 (1980).
40. Perez, F. A. et al. Phase diagram of a three-dimensional antiferromagnet with random magnetic anisotropy. *Phys. Rev. Lett.* **114**, 097201 (2015).
41. Bevaart, L., Frikkee, E., Lebesque, J. & de Jongh, L. J. Magnetic and neutron scattering experiments on the antiferromagnetic layer-type compounds $\text{K}_2\text{Mn}_{1-x}\text{M}_x\text{F}_4$ ($M = \text{Fe}, \text{Co}$). *Phys. Rev. B* **18**, 3376 (1978).
42. Wegner, F. On the magnetic phase diagram of $(\text{Mn}, \text{Fe})\text{WO}_4$. *Solid State Comm.* **12**, 785 (1973).
43. Toft-Petersen, R. et al. Field-induced reentrant magnetoelectric phase in LiNiPO_4 . *Phys. Rev. B* **95**, 064421 (2017).
44. Fogh, E. et al. Magnetic structures and quadratic magnetoelectric effect in LiNiPO_4 beyond 30T. *Phys. Rev. B* **101**, 024403 (2020).
45. Diethrich, T. J., Gnewuch, S., Dold, K. G., Taddei, K. M. & Rodriguez, E. E. Tuning magnetic symmetry and properties in the olivine series $\text{Li}_{1-x}\text{Fe}_x\text{Mn}_{1-x}\text{PO}_4$ through selective delithiation. *Chem. Mater.* **34**, 5039 (2022).
46. Semkin, M. A. et al. Magnetic phase transitions in the $\text{LiNi}_{0.9}\text{M}_{0.1}\text{PO}_4$ ($M = \text{Mn}, \text{Co}$) single crystals. *Phys. Scr.* **97**, 025707 (2022).
47. Ye, F. et al. Magnetic switching and phase competition in the multiferroic antiferromagnet $\text{Mn}_{1-x}\text{Fe}_x\text{WO}_4$. *Phys. Rev. B* **78**, 193101 (2008).
48. Ye, F. et al. Magnetic order and spin-flop transitions in the cobalt-doped multiferroic $\text{Mn}_{1-x}\text{Co}_x\text{WO}_4$. *Phys. Rev. B* **86**, 094429 (2012).
49. Patureau, P. et al. Incorporation of Jahn-Teller Cu^{2+} Ions into Magnetoelectric Multiferroic MnWO_4 : Structural, Magnetic, and Dielectric Permittivity Properties of $\text{Mn}_{1-x}\text{Cu}_x\text{WO}_4$ ($x \leq 0.25$). *Inorg. Chem.* **54**, 10623 (2015).
50. Chaudhury, R. P., Lorenz, B., Wang, Y. Q., Sun, Y. Y. & Chu, C. W. Reentrant ferroelectricity and the multiferroic phase diagram of $\text{Mn}_{1-x}\text{Fe}_x\text{WO}_4$. *N. J. Phys.* **11**, 033036 (2009).
51. Chynoweth, A. G. Dynamic method for measuring the pyroelectric effect with special reference to barium titanate. *J. Appl. Phys.* **27**, 78 (1956).
52. Paেকে, S. *Setup of a measurement system to determine the electric polarization*, Bachelor's thesis. (Technische Universität Berlin, 2010).
53. Metropolis, N., Rosenbluth, A. W., Rosenbluth, M. N. & Teller, A. H. Equation of state calculations by fast computing machines. *J. Chem. Phys.* **21**, 1087 (1953).
54. Liang, G. et al. Anisotropy in magnetic properties and electronic structure of single-crystal LiFePO_4 . *Phys. Rev. B* **77**, 064414 (2008).
55. Stewart, J. R. et al. Disordered materials studied using neutron polarization analysis on the multi-detector spectrometer, D7. *J. Appl. Cryst.* **42**, 69 (2009).

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Author contributions

The experimental project and theoretical framework was conceived by E.F., R.T.-P. and N.B.C. The crystals were grown by D.V. Magnetic susceptibility measurements were performed by E.F., K.S.P. and K.K.M. Diffraction experiments were performed by E.F., R.E.H., A.B.K., M.K.S., G.M., A.B., J.-R.S. M.R., P.B., S.H.-D., O.Z., J.S., Ch.N. R.T.-P. and N.B.C. Pyrocurrent measurements were carried out by E.F., B.K., S.P. and R.T.-P. Sample preparation for the pyrocurrent measurements was performed by E.F. and A.P. E.F. implemented and performed Monte Carlo simulations with support from O.F.S. Data analysis and figure preparation were performed by E.F. Interpretation and manuscript writing was done by E.F., R.T.-P., N.B.C. and H.M.R. with contributions from all authors.

Competing interests

The authors declare no competing interests.

Additional information

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Correspondence and requests for materials should be addressed to Ellen Fogh, Niels B. Christensen or Rasmus Toft-Petersen.

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