Spontaneous decays of magneto-elastic excitations in non-collinear antiferromagnet \((\text{Y,Lu})\text{MnO}_3\)

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Magnons and phonons are fundamental quasiparticles in a solid and can be coupled together to form a hybrid quasi-particle. However, detailed experimental studies on the underlying Hamiltonian of this particle are rare for actual materials. Moreover, the anharmonicity of such magnetoelastic excitations remains largely unexplored, although it is essential for a proper understanding of their diverse thermodynamic behaviour and intrinsic zero-temperature decay. Here we show that in non-collinear antiferromagnets, a strong magnon-phonon coupling can significantly enhance the anharmonicity, resulting in the creation of magnetoelastic excitations and their spontaneous decay. By measuring the spin waves over the full Brillouin zone and carrying out anharmonic spin wave calculations using a Hamiltonian with an explicit magnon-phonon coupling, we have identified a hybrid magnetoelastic mode in \((\text{Y,Lu})\text{MnO}_3\) and quantified its decay rate and the exchange-striction coupling term required to produce it.
Spin and lattice constitute two of the four fundamental degrees of freedom in the solid: the other two are charge and orbital. In linearized models that account for many current understandings of the solid, excitations of spin and lattice, magnon of spin waves and phonon of lattice vibrations, are two principal examples of such quasiparticles. Although there have been some experimental observations of cross-coupling between magnon and phonon, it is rather rare to actually observe and, more importantly, quantify the magnon–phonon coupling for real materials. Nevertheless, it is essential for a proper understanding of their diverse thermodynamic behaviour and intrinsic zero-temperature decay. Furthermore, it is generally believed that the magnon–phonon coupling is important for materials such as multiferroic compounds, geometrically frustrated systems, spin-Peierls systems and Invar materials, to name only a few.

Quasiparticles like magnon and phonon, the cornerstone of modern condensed matter physics, are fundamentally the byproducts of linearized theories that ignore all the higher-order terms than quadratic terms and neglect any conceivable interaction among the quasiparticles themselves. As such, they are considered to be stable, except for very few exceptions. For example, for classical spin systems without strong quantum fluctuations, magnon breakdown is thought to be unlikely for most of purposes. Therefore, observing and understanding how the otherwise stable quasiparticles break down in these unusual cases are the central theme of condensed matter physics.

One route leading to the breakdown of magnon and phonon is the cubic anharmonicities. Despite the general belief that this nonlinear magnon–magnon interaction is rather weak in real materials, recent insights gained mainly through theoretical studies suggest that things should be drastically different for certain cases, namely for non-collinear antiferromagnetic structures. Unlike collinear magnetic structures that forbid the cubic anharmonicities, it was shown that such interaction is in principle allowed for non-collinear magnetic structures, such as the canonical 120° spin pattern in a two-dimensional (2D) triangular lattice. There have since been several experimental reports supporting these theoretical postulates. Nevertheless, the full details of the nonlinear interaction still need to be worked out, especially from experiments. We should also point out that this non-collinear magnetic order, in principle, allows a hitherto forbidden magnon–phonon coupling that has been less recognized among the community: the first-order variation of the nonlinear interaction among the quasiparticles themselves. As such, they are considered to be stable, except for very few exceptions.

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Here we report the direct observation of the magnon–phonon coupling and the spontaneous decay of magneto-elastic excitations in the triangular antiferromagnets (Y,Lu)MnO₃. The full magnetic excitation spectra measured by inelastic neutron-scattering experiments show clear deviations from the linear spin wave theory without the magnon–phonon coupling: an additional mode at high energies and the downward shift of the bottom mode at the Brillouin zone boundary. This is the most direct and stark evidence of the linear coupling of magnons and phonons, which, in turn, leads to enhancement of the anharmonic effects. We demonstrate that these experimental anomalies can only be fully resolved by incorporating the magnon–phonon coupling and carrying out the nonlinear spin wave analysis. We further reveal that the magneto-elastic excitation leads to significant broadening of the magnon spectra at the zone boundaries, originating from the decay of the magneto-elastic excitations into the two-magnon continuum.

**Results**

**Failure of standard spin waves calculation.** Hexagonal rare-earth manganite RMnO₃ represents a good model system for geometrical frustration on a 2D triangular lattice: the nearest-neighbour antiferromagnetic interaction between $S = 2$ ($\text{Mn}^{3+}$) spins dominate, whereas the interlayer interaction is relatively weak. Although there have been some experimental observations of a spontaneous magnon decay and a remarkably large spatial anisotropy for Mn$^{3+}$ ions in the exchange interactions for LuMnO₃. This was attributed to a structural distortion in which groups of three Mn atoms become more closely bound, such that the intratramer $J_1$ exchange constants may differ from the intertrimer $J_2$ (see Fig. 1). Similar interpretations were shared by other groups too. However, we should note that the large $J_1/J_2$ ratio of 6.4 obtained from fitting the data is inconsistent with the value of 1.15, obtained from first-principles calculations that used the experimental atomic positions as reported by neutron diffraction. This realization prompted us to handle both magnon and phonon, as well as their cross-coupling on an equal footing and to go beyond the linear spin wave analysis.

Figure 1 shows the spin waves measured at the MAPS beamline of the ISIS facility together with the theoretical dispersion relation calculated from the spin Hamiltonian by a linear spin wave theory using the following parameters (see Supplementary Note 2): for YMnO₃: $J_1 = 1.4, J_2 = 1.8, D_1 = -0.28, D_2 = 0.02$ meV; for $Y_0.5Lu_0.5$MnO₃: $J_1 = 12.5, J_2 = 0.97, D_1 = 0.18, D_2 = -0.018$ meV; for LuMnO₃: $J_1 = 9, J_2 = 1.4, D_1 = 0.28, D_2 = 0.02$ meV. Despite the apparent success of the linear spin wave calculations, there lies a critical failure: first, the downward curvature along the AB direction and, second, the additional peaks at ~19 meV indicated by a red box in Fig. 1c–e. However, most importantly, here we have to use an unphysically large $J_1/J_2$ ratio, to explain the additional high-energy peaks. Apart from the large $J_1/J_2$ ratio, this analysis of the linear spin waves has another drawback: which is that the calculated dynamical structure factor using the linear spin wave theory as shown in Fig. 1f–h always produces stronger intensity at the top mode of the spin waves than at the middle one, in marked contrast with the experimental data.

This discrepancy requires us to adopt a radically different approach to explain the measured full spin waves and go beyond the standard linear spin wave theory. One clue for how to address this problem can be taken from the physical properties: for example, our earlier neutron diffraction data revealed a giant spin–lattice coupling for (Y,Lu)MnO₃ (ref. 5). This observation was subsequently corroborated by independent measurements on rare-earth hexagonal RMnO₃ (refs 18–25). More importantly, ultrasound measurements on YMnO₃ found marked softening in $C_{11}$ and $C_{44}$, supporting our view that there is a strong in-plane deformation below $T_N$. This observation naturally indicates the importance of magnon and in-plane phonon coupling. This conclusion is also backed up by theoretical calculations.

**Calculation of magneto-elastic excitations.** Following this idea of a large spin–lattice coupling in RMnO₃, we took a first-principles approach to the magnon–phonon coupling.
First, we construct the following full model Hamiltonian, which couples the in-plane manganese vibrations directly to the spin system:

\[ H = H_{\text{spin}} + \hbar \sum_{i=1}^{90} \epsilon_{ij} b_i^+ b_j + \tilde{z} \sum_{g} \left( e_{ij} \cdot u_i + e_{ij} \cdot u_j \right) S_i \cdot S_j \]

(1)

where \( e_{ij} \) denotes the unit vector connecting the \( i \)-th manganese atom and the neighbouring oxygen atoms between the \( i \)-th and \( j \)-th manganese atoms as shown in Supplementary Fig. 2, \( \tilde{z} \) is the exchange striction, \( \tilde{z} = \frac{a_d}{d} \) which is naturally made into a dimensionless exchange–striction constant \( a = \tilde{z} \cdot 2d / J^2 \), and \( d \) is the Mn–O bond length at the equilibrium. Therefore, our Hamiltonian takes into account the modulations of the Mn–O bond length as a function of Mn displacements.

Before going into detailed discussion, we would like to make a general remark on the related issue. In cases when the spin rotational symmetry is broken completely in the ground state, that is, when the spin structure is non-collinear, the Heisenberg term provides a coupling of the transverse and longitudinal modes on neighbouring sites. That is, in terms of the local site-dependent preferred spin direction of the ordered state, the coupling terms take the form of the type \( S_i^z S_j^z \) and so on. In the magnon language, they are quantized into the ‘odd’ terms, producing linear \( (a + a') \) and cubic \( (a' a a') \) and so on) contributions. In equilibrium, the linear magnon term must vanish, leaving the anharmonic cubic magnon coupling to be the sole outcome, which is important for magnon decays. However, in the presence of coupling to phonons, the linear \((a + a')\) terms is ‘activated’, as the local atomic displacements \( u_i \) violate the equilibrium conditions locally, hence the ‘direct’ coupling of magnons and phonons.

To calculate the full dispersions of all 90 phonon modes for the unit cell with six formula units, we used a first-principles density functional theory (DFT). We show the full phonon dispersion curves for the three compounds as dashed lines in Fig. 2. We note that the calculated phonon density of states (DOS) is in good agreement with the phonon spectra we measured using powder YMnO\(_3\) and LuMnO\(_3\) at the AMATERAS beamline of J-PARC (see Supplementary Fig. 1). We then calculated the dynamical spin structure factor within the linear approximation by using the full Hamiltonian above with the explicit magnon–phonon coupling: we used the dimensionless exchange–striction coefficients of \( a \approx 16\)–20.

The exchange–striction constant can also be estimated by using the pressure dependence of the crystal structure and the antiferromagnetic transition reported for YMnO\(_3\) (refs 29,30). Using the experimental data reported in refs 9,10, we came to an estimate of the dimensionless exchange striction \( a \approx 14 \) with the following formula: \( a = \frac{\partial(\rho)}{\partial P} \). Here, \( \rho \) is average Mn–O bond length, which is approximately one-third of lattice constant \( a \). The experimental parameters used in our estimate are \( \frac{\partial(\rho)}{\partial P} = 3 \text{ K GPa}^{-1} \) (ref. 29) and \( \frac{\partial(\rho)}{\partial P} = 0.0057 \text{ Å GPa}^{-1} \).
In fact, an estimate of the exchange striction from the splitting of the high-energy hybrid modes in Fig. 2 yields the values in the same range, \( \alpha \approx 10–20 \) (see Supplementary Note 3). We also note that our polarized neutron-scattering data are in good agreement with our calculations (see Supplementary Fig. 4 and Supplementary Note 4).

**Spontaneous decay of the hybrid mode**. In addition to the magnon–phonon hybridization, non-collinear spin structures allow three magnon interactions as discussed above, which can lead to spontaneous magnon decay into two magnon states when the kinematic conditions are satisfied. The magneto-elastic excitations have, by definition, both magnon and phonon characters. Therefore, the above mechanism can also lead to the decay of magneto-elastic excitations inside the two quasi-particles continuum of magnon. Indeed, we observe significant broadening of the top mode in LuMnO\(_3\) near the \( B \) and \( D \) points as shown in Fig. 3 and Supplementary Fig. 3: less strong broadening has been seen for other two compounds.

To calculate the decay rate directly and compare it with the experimental data, we simplify the problem by assuming a dispersionless optical phonon mode at \( \omega \approx 20 \) meV, where the strongest coupling has been observed in our data. Next, our model Hamiltonian reads as follows:

\[
H = J \sum_{ij} \left( s_i^x s_j^x + s_i^y s_j^y + J^S Z_i Z_j \right) + \hbar \omega_0 \sum_i b_i^\dagger b_i + \frac{\hbar}{2} \sum_{ijkl} (a_{ijkl} \cdot \mathbf{u}_i + a_{ijkl}^* \cdot \mathbf{u}_j) s_i^+ s_j^-.
\]

(2)

First, we calculate the dynamical spin structure factor by using a standard method. As shown in Fig. 3a, despite the simplification, the calculation results reproduce well the experimental intensity along the \( \Gamma-B-D \) direction. For the calculations, we used the following set of the parameters: for YMnO\(_3\): \( J = 2.7 \) meV, \( \gamma = 0.93, \ h\omega_0 = 17.5 \) meV, \( \alpha = 7.2 \); for LuMnO\(_3\): \( J = 3.2 \) meV, \( \gamma = 0.93, \ h\omega_0 = 19.5 \) meV, \( \alpha = 8 \). Next, the decay rate of the high-energy mode was calculated using an anharmonic spin wave theory within the \( 1/S \) approximation. The calculated results summarized in Fig. 3 also show the significant linewidth broadening for the top mode near the \( B \) and \( D \) points only for LuMnO\(_3\), consistent with the experimental results.

The reason for this is that in LuMnO\(_3\), the combination of the higher energies of the magnon and magnetoelastic modes means that more decay channels, including two quasiparticle emission, are kinematically allowed. The different decay channels have different boundaries in the reciprocal space, which also corresponds to logarithmic singularities in the decay rate\(^{33}\), and the largest broadening is observed at momentum transfers where the single-quasiparticle dispersion crosses these boundaries, such as at the \( B \) point. In the case of YMnO\(_3\), no such crossing occurs; thus, there are fewer decay channels available explaining why the observed linewidths remain narrow. Here we should stress that the single-magnon branches do not cross the line of singularities, whereas the magnetoelastic modes do; thus, a pure magnon decay is forbidden in this case. Similarly, the intrinsic decay rate of phonons is usually small due to a weak cubic anharmonicity. Thus, the strong hybridization of magnons and phonons provides a new mechanism to enhancing the magnon decays.

**Discussion**

Our studies using an exchange-striction model indicates that the deviations from the linear spin wave theory must be a common feature for other triangular antiferromagnets with a non-collinear magnetic order. Looking beyond the 2D triangular antiferromagnets, we believe that the idea of magnon–phonon coupling can also be important in the studies of a wide variety of...
the 3d transition-metal magnetic compounds. For example, similar analysis might shed a light on the investigation of spin phonon coupling mechanism and anharmonic effects in many other important non-collinear magnets that exhibit a rather large spin–lattice coupling, such as spinel34 and invar materials35, which should be dominated by the exchange–striction as discussed in ref. 36. This is in contrast to the previous studies on magnon–phonon coupling focusing on materials with strong spin–orbit coupling such as rare-earth elements37.

To summarize, we mapped out the spin waves and phonon excitations of (Y,Lu)MnO3 over the Brillouin zone. By carrying out the spin wave calculations using the full Hamiltonian with both magnons and phonons on an equal footing and their coupling, we have not only demonstrated in our inelastic neutron-scattering data a clear sign of magnon–phonon coupling, but also have quantified the coupling strength directly. Our work provides the rare experimental test and quantification of magnon–phonon coupling focusing on materials with strong spin–orbit coupling such as rare-earth elements37.

Figure 3 | Linewidth broadening of magneto-elastic excitation. (a) The neutron-scattering data along the CBD direction (middle) and the calculated dynamical structure factor within the linear spin wave theory (bottom). Observed linewidth broadening of the top mode (square) together with the calculated result from the 1/5 approximation (line) and the experimental resolution (dashed line) (top) for YMnO3 (left) and LuMnO3 (right). (b) Observed linewidth of the top mode for LuMnO3 (contour map) and (c) calculated intrinsic broadening of the top mode using the model Hamiltonian, equation (2). The experimental linewidth of the top mode was estimated by using multi-Gaussian functions and the typical results are shown in Supplementary Fig. 3.

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Methods

Sample preparation. We synthesized powder samples using a solid-state reaction method following the recipe as described in the literature38. We then grew single crystals of Y1_xLum,MnO3 (with typical size of 5 × 5 × 40 mm3) by using a commercial optical floating zone furnace (Crystal Systems, Japan). Our subsequent powder and single crystal XRD confirmed that all our samples are prepared in high quality. We also measured the bulk properties (susceptibility and heat capacity) of all the samples to further confirm the quality by using a commercial set-up (MPMS5XL and PPMS9, Quantum Design USA).

Inelastic neutron scattering. Inelastic neutron-scattering experiments have been performed on single crystal samples using the MAPS time-of-flight spectrometer at ISIS, UK, and a triple axes spectrometer with polarization analysis at Chalk River, Canada. In the time-of-flight experiments, incident energies were chosen at 40 meV for LuMnO3, 35 meV for Y0.5Lu0.5MnO3, and 30 meV for YMnO3 to adjust to the slight variations in the energy scales for each samples. The chopper frequency has been set to 250 Hz, which gave us a full width at half maximum energy resolution of 0.43 ~ 1.36 meV depending on the energy transfer. The measurements have been performed at 4 K for YMnO3 and Y0.5Lu0.5MnO3, and 13 K for LuMnO3. We used the Horace programme for our data analysis39. In the triple axes spectrometer experiments, spin-polarized neutrons have been produced by using vertical focusing Heusler monochromator and analyser with the final energy fixed at 13.7 meV. To measure phonon DOS, inelastic neutron-scattering experiments have also been performed on the powder samples with the incident energy of 42 meV using the AMATERAS beamline at J-PARC, Japan.

Theoretical calculations. We carried out first-principles calculations of phonon using a DFT + U method with U = 4 eV. We used the PHONOPY code based on the force constant method40. In addition, the force constants were constructed by means of a supercell approach based on the density functional perturbation theory41, implemented in the VASP code42. Detailed discussion is given in the Supplementary Note 1.

For the spin waves calculations, we used a rotating framework with the direction of easy axis anisotropy being rotated from parallel to perpendicular to the crystallographic axes. To make our calculations simpler and transparent, we ignored the interlayer exchange coupling as it is known to be more than 100 times smaller than the in-plane coupling63,44. Using this approximation, we can have the following minimal Hamiltonian:

\[ H = J_1 \sum_{\text{intra}} S_i \cdot S_j + J_2 \sum_{\text{inter}} S_i \cdot S_j + D_1 \sum_i (S_i^z)^2 + D_2 \sum_i (S_i^y)^2 \]

where \( J_1 \) and \( J_2 \) represent intra- and inter-trimer exchange constants and \( D_1 \) and \( D_2 \) are two magnetic anisotropies. We then calculated the spin wave dispersion using the standard linear spin wave theory45. We give detailed description of our spin wave calculations for the full Hamiltonian with the magnon and phonon coupling and the magnon–magnon nonlinear interaction in the Supplementary Notes 2 and 3.

Data availability. All relevant data that support the findings of this study are available from the corresponding author on request.
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Authors contributions

J.G.P. conceived and supervised the project. H.S., Y.Y., H.E. and S.W.C. grew the single crystals. The work of ALC was supported by the U.S. DOE, Office of Science, Basic Energy Sciences under Award Number DE-FG02-04ER46174. The work at Rutgers University was supported by the DOE under Grant Number DOE: DE-FG02-07ER46382.

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