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Anisotropic spin wave excitations in a multiferroic BiFeO₃

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Polarized inelastic neutron scattering experiments have been performed to elucidate the anisotropic behavior of the low energy spin wave excitations in a multiferroic BiFeO₃, which shows a cycloidal spin structure below 640 K. With the neutron polarization analysis for single magnetic domained crystals, magnetic excitation modes in and out of the cycloidal plane below 6 meV were separated successfully. **The magnetic excitation spectra were analyzed using the linear spin wave theory.** The low-energy magnon density of states consist of several magnon modes, including the two anisotropic modes, Φ and Ψ modes, distributed in and out of the cycloidal plane, respectively, which were previously observed using optical spectroscopies. **Furthermore, there are other magnon modes that are not active in optical measurements. A model spin Hamiltonian, which reproduces the spin-wave frequencies observed using optical spectroscopies, explains the overall spectra reasonably well.**

I. INTRODUCTION

Bismuth ferrite, BiFeO₃, is one of the multiferroic materials that show ferroelectric polarization and an antiferromagnetic order above room temperature. The crystal structure is hexagonal $R3c$ with the lattice parameters of $a_{\text{hex}} \approx 5.58$ Å and $c_{\text{hex}} \approx 13.86$ Å below ~ 1100 K [1], as shown in Fig. 1(a). The structure can also be represented approximately by the pseudo-cubic unit cell with the lattice constant $a = a_{\text{pc}} \approx 3.96$ Å. This material primarily shows a large spontaneous ferroelectric polarization below 1100 K with an additional polarization below 640 K, where a cycloidal magnetic structure with the magnetic wave vector, $\mathbf{q}_m = [\delta, \delta, 3]_{\text{hex}}$ ($\delta \approx \frac{1}{222}$), develops [2–5]. Due to this multiferroic behavior occurring above room temperature, extensive studies have been performed to clarify the electromagnetic coupling in this material [6].

One of the interesting features in BiFeO₃ is the electromagnon, which is a hybrid mode of magnon and phonon activated by the magnetoelectric coupling. The electromagnon was studied extensively in one of the most celebrated multiferroic systems, $RMnO_3$ (R : rare-earth elements), using optical techniques [7]. The nature of the electromagnon depends on the energy range. Low and high energy electromagnons are activated by the inverse Dzyaloshinskii-Moriya (DM) coupling [8, 9] and exchange-striction mechanisms[10], respectively.

The electromagnon in BiFeO₃ was studied using Raman spectroscopy [11–14] and time-resolved THz spectroscopy measurements [15, 16]. Two kinds of magnon

modes (Φ and Ψ modes at lower and higher energies, respectively) were identified. The spectrum has a peaked structure at the crossing points of these magnon modes (Φ_n and Ψ_n with n integer) at the magnetic zone center due to the zone folding. The Φ and Ψ modes correspond to the spin wave excitations in and out of the cycloidal

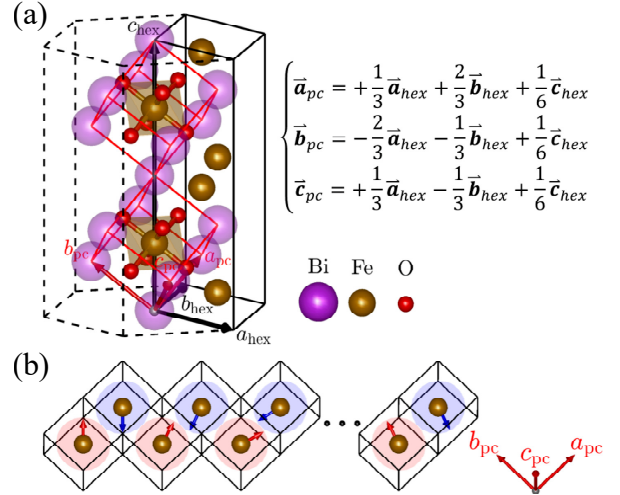


FIG. 1. Crystal and magnetic structures of BiFeO₃. (a) The nuclear structure of BiFeO₃, where the black arrows represent the hexagonal and pseudo-cubic unit cells, respectively and the violet, gold, and red spheres represent the Bi, Fe and O atoms, respectively. The coordinate transformation equations between the hexagonal and pseudo-cubic cells are listed. (b) The cycloidal magnetic structure of BiFeO₃ in the pseudo-cubic cell, where the magnetic wave vector is $\mathbf{q}_m = [\delta, \delta, 3]_{\text{hex}}$ with $\delta \approx \frac{1}{222}$ and the cycloidal plane is tilted by $\tau = 0.46^\circ$ from the $[H, \bar{H}, L]_{\text{hex}}$ plane. Both δ and τ are exaggerated by 20 times for a better demonstration.

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plane, respectively [17]. The Heisenberg model with a single ion anisotropy and two kinds of DM interactions successfully explain the electromagnon [$\Psi_1^{(1)}$] and magnetic enhanced [Ψ_0 , $\Psi_1^{(2)}$, $\Phi_1^{(1)}$, and $\Phi_2^{(1)}$] modes [18], observed with optical spectroscopies. Due to the electromagnetic coupling, the energies of the magnon modes were found to shift with changing electric field even at room temperature, indicating the control of spin wave excitations using electric field [19].

In principle, the magnon modes reported in the optical spectroscopy measurements can be observed using the inelastic neutron scattering measurements, which has an advantage of measuring magnon and phonon dispersions over the full range of momentum space, whereas only modes around the zone center can be measured with optical spectroscopies. Therefore, it is important to perform detailed inelastic neutron scattering measurements in BiFeO₃ to study the magnon modes relevant to the electromagnon, although the enhancement of the electromagnon by a photon's electric field cannot be measured. In fact, low-energy magnon modes corresponding to the electromagnon were identified in DyMnO₃ and TbMnO₃ using inelastic neutron scattering [20, 21]. However, there has been no neutron scattering study in BiFeO₃ in this respect. One of the reasons is that the instrumental energy resolution is not sufficient enough to separate the magnon modes. Even without excellent energy resolution, observing anisotropy of the magnetic excitation modes, **which could be feasible using the polarized neutron scattering technique**, would give useful information in identifying magnon modes associated with the electromagnon. However, a crucial issue is the difficulty of identifying the anisotropy of the magnon modes even using the polarized neutron technique because there exist three magnetic domains that are integrated over due to the instrumental Q resolution broader than the split of the magnetic peaks. As a result, the anisotropic effect is averaged out completely.

Due to the large spin value and exchange interactions, the magnetic excitation band extends up to ~ 70 meV [22, 23]. However, the low energy magnon modes below ~ 6 meV are extremely important to study the magnon modes relevant to the electromagnon in BiFeO₃. The low energy magnetic excitations were measured and analyzed to determine the single ion anisotropy and the DM interactions in Ref. [24]. Since the neutron scattering study was performed using a multi-magnetic-domained sample with unpolarized neutrons, the magnetic anisotropy of the magnon modes was not evaluated.

In this study, two changes to the experimental conditions were made in order to identify the low energy magnon modes accurately. In particular, we could successfully align the three magnetic domains under high magnetic fields and observe the anisotropy of the magnon modes, i.e. in and out of the cycloidal plane, using the polarized neutron technique. **Another important point is that the inelastic neutron scattering can observe magnon modes that are not active in optical measurements. We**

found that a model spin Hamiltonian, which reproduces the spin-wave frequencies observed using optical spectroscopies [25], explains the overall spectra, including the anisotropy of the magnetic excitations, reasonably well.

II. EXPERIMENTAL DETAILS

A single crystal of BiFeO₃ was grown using the traveling solvent floating zone (TSFZ) method, as described in Ref. [26]. As described in Ref. 27 and 28, as-grown crystals have three magnetic domains and the domains can be aligned to be almost single by applying a magnetic field of 6 T. The domain distribution is preserved as long as the sample is stored at ambient temperature and field. We comounted two single magnetic domained crystals with the dimensions of $3\phi \times 10$ mm³ in the $(HHL)_{\text{hex}}$ scattering plane, which corresponds to the cycloidal plane approximately with a small amount of tilt out of the plane, as shown in Fig. 1(b).

Polarized inelastic neutron scattering measurements were performed on the hybrid neutron spectrometer HYSPEC [29] installed at the Spallation Neutron Source at Oak Ridge National Laboratory. The incident neutron energy, E_i , was fixed at 7.5 meV to cover the major low energy part of the magnetic excitations of BiFeO₃. The energy resolution is ~ 0.4 meV at the elastic position. The incident neutrons were polarized using Heusler (111) crystals and the polarization of the scattered neutrons was analysed using wide-angle-arrayed supermirrors. A pair of magnetic coils was used to align the neutron polarization perpendicular to the scattering plane at the sample position. The Mezei spin flipper was used to optionally flip the incident neutron spin. With this setup, the non-spin flip (NSF) and spin flip (SF) components were measured with the two-dimensionally arrayed ³He tube detectors. The flipping ratio of this setup was ~ 10 , **estimated using nuclear Bragg peaks, which is consistent with the typical polarization of the instrument with the same configuration**. In this configuration, the NSF and SF components correspond to the magnetic contributions out of and in the scattering plane, respectively [30]. **The sample table was rotated by ± 23 degrees around $(0, 0, 3)_{\text{hex}}$ to measure the excitations in a wide Q range**. Although the phonon signal can be observed at the NSF channel, purely magnetic excitations are expected at the momentum and energy ranges in this study. Due to the vertical focusing of the incident beam, the instrumental Q resolution out of the scattering plane is broad, which is much broader than the tilt of the cycloidal plane in the sample. The observed intensities were corrected with the correction factors for the flipping ratio and the transmission of the supermirrors.

Here, the relation between the hexagonal and pseudo-cubic structure is described. The hexagonal unit cell of BiFeO₃ is shown as the black cell and arrows in Fig. 1 (a), where the violet, gold and red spheres represent the Bi, Fe and O atoms, respectively. The pseudo-cubic unit

cell is shown as the red cells and arrows. Each pseudo-cubic unit cell contains one chemical formula unit, while the hexagonal cell contains six. The lattice vectors of the hexagonal and pseudo-cubic cells can be converted using the following equations,

$$\begin{aligned} \mathbf{a}_{\text{pc}} &= +\frac{1}{3}\mathbf{a}_{\text{hex}} + \frac{2}{3}\mathbf{b}_{\text{hex}} + \frac{1}{6}\mathbf{c}_{\text{hex}}, \\ \mathbf{b}_{\text{pc}} &= -\frac{2}{3}\mathbf{a}_{\text{hex}} - \frac{1}{3}\mathbf{b}_{\text{hex}} + \frac{1}{6}\mathbf{c}_{\text{hex}}, \\ \mathbf{c}_{\text{pc}} &= +\frac{1}{3}\mathbf{a}_{\text{hex}} - \frac{1}{3}\mathbf{b}_{\text{hex}} + \frac{1}{6}\mathbf{c}_{\text{hex}}, \end{aligned} \quad (1)$$

where \mathbf{a}_{pc} , \mathbf{b}_{pc} , \mathbf{c}_{pc} and \mathbf{a}_{hex} , \mathbf{b}_{hex} , \mathbf{c}_{hex} represent the lattice vectors for the pseudo-cubic and hexagonal cells, respectively.

Similarly, we can also obtain the transformation equations for the reciprocal lattice vectors,

$$\begin{aligned} \mathbf{a}_{\text{pc}}^* &= +\mathbf{b}_{\text{hex}}^* + 2\mathbf{c}_{\text{hex}}^*, \\ \mathbf{b}_{\text{pc}}^* &= -\mathbf{a}_{\text{hex}}^* + 2\mathbf{c}_{\text{hex}}^*, \\ \mathbf{c}_{\text{pc}}^* &= +\mathbf{a}_{\text{hex}}^* - \mathbf{b}_{\text{hex}}^* + 2\mathbf{c}_{\text{hex}}^*, \end{aligned} \quad (2)$$

where \mathbf{a}_{pc}^* , \mathbf{b}_{pc}^* , \mathbf{c}_{pc}^* and $\mathbf{a}_{\text{hex}}^*$, $\mathbf{b}_{\text{hex}}^*$, $\mathbf{c}_{\text{hex}}^*$ represent the reciprocal lattice vectors for the pseudo-cubic and hexagonal cells, respectively.

Therefore, the $[1, 1, 0]_{\text{hex}} - [0, 0, 1]_{\text{hex}}$ scattering plane corresponds to the $[1, -1, 0]_{\text{pc}} - [1, 1, 1]_{\text{pc}}$ plane and the polarization direction of $[1, -1, 0]_{\text{hex}}$ corresponds to $[-1, -1, 2]_{\text{pc}}$. The hexagonal notation will mostly be used in this paper and the equivalent values for the pseudo-cubic notation will be shown at some places. We will also use \mathbf{x}' , \mathbf{y}' , and \mathbf{z}' , which are defined as unit vectors along $[1, 1, 0]_{\text{hex}}$, $[1, -1, 0]_{\text{hex}}$, and $[0, 0, 1]_{\text{hex}}$, respectively.

III. RESULTS AND DISCUSSION

A. Polarized Inelastic Neutron Scattering Experiments

Figure 2 shows the polarized inelastic neutron scattering spectra of BiFeO₃ measured at 300 K along three orthogonal reciprocal directions in the vicinity of $(0, 0, 3)_{\text{hex}}$, which is close to the magnetic wave vector $\mathbf{q}_m = [\delta, \delta, 3]_{\text{hex}}$. Although the incommensurate magnetic peaks are present at $(\delta, \delta, 3)_{\text{hex}}$, the instrumental Q resolution is not sufficient enough to resolve the detailed structure in the momentum space. The magnetic excitation spectra clearly show that both the NSF and SF components exhibit a gap at $\hbar\omega \approx 1.5$ meV and the NSF component mainly contributes to the magnetic excitations above the gap, while the SF signals are mainly distributed below the gap.

In order to show the anisotropic behavior of the magnetic excitations more clearly, the NSF and SF components of the magnetic excitations are displayed as a function of excitation energy around $(0, 0, 3)_{\text{hex}}$ in Fig.

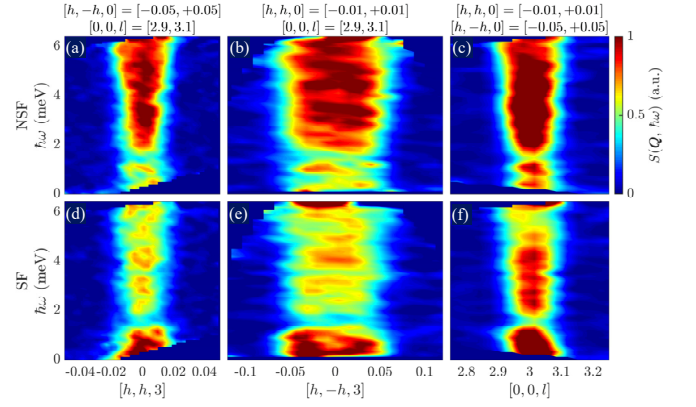


FIG. 2. Contour plots of the polarized inelastic neutron scattering spectra of BiFeO₃ measured at 300 K. Panels (a)-(c) and (d)-(f) show the neutron intensity, $S(\mathbf{Q}, \hbar\omega)$, as a function of the momentum transfer \mathbf{Q} and the energy transfer $\hbar\omega$ for the non-spin flip (NSF) and spin flip (SF) components, respectively, along three orthogonal directions, $[h, h, 3]_{\text{hex}}$ [(a) and (d)], $[h, -h, 3]_{\text{hex}}$ [(b) and (e)], and $[0, 0, l]_{\text{hex}}$ [(c) and (f)]. In panels (a) and (d), $S(\mathbf{Q}, \hbar\omega)$ is integrated over $[h, -h, 0]_{\text{hex}}$ for $-0.05 \leq h \leq +0.05$ and $[0, 0, l]_{\text{hex}}$ for $2.9 \leq l \leq 3.1$; in panels (b) and (e), the integration is applied over $[h, h, 0]_{\text{hex}}$ for $-0.01 \leq h \leq +0.01$ and $[0, 0, l]_{\text{hex}}$ for $2.9 \leq l \leq 3.1$; while in panels (c) and (f), the integration is applied over $[h, h, 0]_{\text{hex}}$ for $-0.01 \leq h \leq +0.01$ and $[h, -h, 0]_{\text{hex}}$ for $-0.05 \leq h \leq +0.05$. **The counting time for each of SF and NSF channels is approximately 40 hours.** The wider excitations in panels (b) and (e) originate from the broader instrumental Q resolution due to the vertical focusing. The intrinsic width is considered to be similar to that along the two other directions. The observed intensities were corrected with the transmission and the flipping ratio factors.

3. The experimental intensity $S_{\text{exp}}(\hbar\omega)$ is integrated over $[h, h, 0]_{\text{hex}}$ for $-0.01 \leq h \leq +0.01$, $[h, -h, 0]_{\text{hex}}$ for $-0.05 \leq h \leq +0.05$, and $[0, 0, l]_{\text{hex}}$ for $2.9 \leq l \leq 3.1$. The NSF and SF components mainly contribute to the magnetic excitations above and below ~ 1.5 meV, respectively. This indicates that the out-of-plane and in-plane spin wave modes are largely populated above and below the gap, respectively. The shape of the overall low-energy magnetic excitations are similar to that measured with unpolarized neutrons in Ref. 24, considering that the result in Ref. 24 corresponds to the sum of the NSF and SF components.

B. Linear Spin Wave Calculations

The spin Hamiltonian and ground state for the spin state of BiFeO₃ are described in [18]. The Hamiltonian \mathcal{H} has five terms: the nearest-neighbor (NN), next-nearest-neighbor (NNN) and anisotropic exchange interactions, J_1 , J_2 , and K , respectively, and two Dzyaloshinskii-Moriya (DM) interactions with $\mathbf{D} \parallel [1, -1, 0]_{\text{hex}}$ and

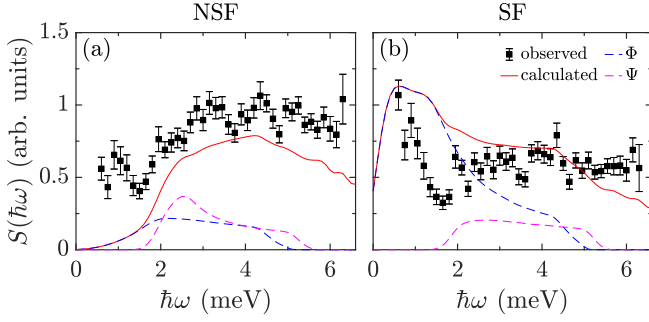


FIG. 3. Energy dependence of the magnetic excitations of BiFeO₃ measured with polarized neutrons. The NSF and SF components correspond to the magnetic excitations out of and in the cycloidal plane, respectively. Panels (a) and (b) compare the experimental and calculated spin wave intensities for the NSF and SF component, respectively. The filled squares show the \mathbf{Q} -integrated neutron intensity, $S_{\text{exp}}(\hbar\omega) = \int d^3\mathbf{Q} S_{\text{exp}}(\mathbf{Q}, \hbar\omega)$, obtained from the experimental results shown in Fig. 2, where the integration is applied over $[h, h, 0]_{\text{hex}}$ for $-0.01 \leq h \leq +0.01$, $[h, -h, 0]_{\text{hex}}$ for $-0.05 \leq h \leq +0.05$, and $[0, 0, l]_{\text{hex}}$ for $2.9 \leq l \leq 3.1$. The red solid line represents the calculated $\hbar\omega$ dependence of spin wave intensity, $S_{\text{calc}}(\hbar\omega)$. The $S_{\text{calc}}(\hbar\omega)$ is integrated over the \mathbf{Q} points in the vicinity of $(0, 0, 3)_{\text{hex}}$ considering the experimental \mathbf{Q} resolution, which integrates most of the spin-wave excitations in the energy range. The blue and magenta dashed lines show the spin wave mode contribution of $S_{\text{calc}}(\hbar\omega)$ from the Φ and Ψ modes, respectively [18]. **The excitation signal below 0.5 meV containing elastic component is not shown.**

$$D' \parallel [0, 0, 1]_{\text{hex}},$$

$$\begin{aligned} \mathcal{H} = & -J_1 \sum_{\text{NN}} \mathbf{S}_i \cdot \mathbf{S}_j - J_2 \sum_{\text{NNN}} \mathbf{S}_i \cdot \mathbf{S}_j - K \sum_i S_{iz'}^2 \\ & - D \cdot \sum_{\langle i,j \rangle} (\mathbf{S}_i \times \mathbf{S}_j) \\ & - D' \cdot \sum_{\langle i,j \rangle} (-1)^{6R_{iz'}/c} (\mathbf{S}_i \times \mathbf{S}_j), \end{aligned} \quad (3)$$

where \mathbf{R}_i and \mathbf{S}_i are the spatial position and the spin orientation of the i -th spin, respectively. The sums for the two DM terms run over the NNs. For the second DM term, the sign changes from layer n to layer $n+1$ along the c axis. **The first DM term gives rise to the cycloid modulation. The second DM term causes a small spin density wave component along $[1, -1, 0]_{\text{hex}}$, which tilts the cycloidal plane. Since the Fe³⁺ spin is large ($S=5/2$), a $1/S$ linear expansion about the classical limit is justified for BiFeO₃. This expansion produces a spectrum of observed in-plane and out-of-plane cycloidal modes that agree very well with spectroscopic measurements [25].**

The magnetic ground state of the Fe³⁺ spins shows a cycloidal structure with $\mathbf{q}_m = [\delta, \delta, 3]_{\text{hex}} = [0.5 + \delta, 0.5 - \delta, 0.5]_{\text{pc}}$ ($\delta \approx \frac{1}{222}$), in which the cycloidal plane lies approximately in the $\mathbf{x}' - \mathbf{z}'$ plane with a tilting angle $\tau (=0.46^\circ)$. The spin components of the Fe³⁺ at \mathbf{R} , $S(\mathbf{R})$, can be expressed as

$$\begin{aligned} S_{x'}(\mathbf{R}) &= (-1)^{6R_{z'}/c} \cos \tau \sqrt{S^2 - S_{z'}(\mathbf{R})^2} \\ &\quad \times \text{sgn}[\sin(2\pi\delta R_{x'}/a)], \\ S_{y'}(\mathbf{R}) &= \sin \tau \sqrt{S^2 - S_{z'}(\mathbf{R})^2} \text{sgn}[\sin(2\pi\delta R_{x'}/a)], \\ S_{z'}(\mathbf{R}) &= (-1)^{6R_{z'}/c} S \sum_{m=0}^{\infty} C_{2m+1} \cos[(2m+1)2\pi\delta R_{x'}/a], \end{aligned} \quad (4)$$

where C_{2m+1} are the odd-order cosine coefficients, which satisfy $\sum_{m=0}^{\infty} C_{2m+1} = 1$.

In order to understand the nature of the magnetic excitations and to reproduce the observed polarized spectra, the spin wave energies and intensities of BiFeO₃ are calculated, using the equations-of-motion technique for non-collinear spins [31] on a pseudo-commensurate magnetic super cell containing $M = 2/\delta = 2 \times 222$ pseudo-cubic cells [18]. The exchange interactions used for our analysis are $J_1 = -5.32$ meV, $J_2 = -0.24$ meV, $K = 0.005$ meV, $D = 0.18$ meV, and $D' = 0.085$ meV, which reproduce the spin-wave frequencies in zero and finite magnetic fields [25]. Based on these parameters, the magnetic ground state then can be determined: the cycloidal plane tilting angle $\tau = 0.46^\circ$ and the cosine coefficients $C_1 = 1.049275$, $C_3 = -0.051811$, $C_5 = 0.002536$, where the higher order coefficients are fixed at zero. The linear spin wave calculations in the vicinity of \mathbf{q}_m are shown in Figs. 4 and 5, where the \mathbf{Q} space centers are located at $(0, 0, 3)_{\text{hex}}$ and $(\delta, \delta, 3)_{\text{hex}}$, respectively.

As expected from the cycloidal spin structure, the magnetic dispersions has a periodicity of 2δ along the $[h, h, 0]_{\text{hex}}$ direction, whereas the dispersions perpendicular to the $[h, h, 0]_{\text{hex}}$ direction are more straightforward. The magnetic excitations emerging from $(0, 0, 3)_{\text{hex}}$, shown in Fig. 4, indicate that both the NSF and SF components are weak in the gapless mode and the gapped modes with minimum at ~ 2 meV are intense both in the NSF and SF components. On the other hand, the magnetic excitations emerging from $(\delta, \delta, 3)_{\text{hex}}$, shown in Fig. 5, indicate that the gapless mode is more intense in the SF component than in the NSF component and the gapped mode with minimum at ~ 2.5 meV is more intense in the NSF component than in the SF component. As described in Sec. III A, those magnon modes at $(0, 0, 3)_{\text{hex}}$ and $(\delta, \delta, 3)_{\text{hex}}$ cannot be resolved and the magnetic excitations integrated over finite momentum and energy ranges are observed experimentally.

The calculated $S_{\text{calc}}(\hbar\omega)$, as represented with solid and dashed curves in Fig. 3, is integrated over the \mathbf{Q} points in the vicinity of $(0, 0, 3)_{\text{hex}}$ in the following way. As seen from Figs. 4 and 5, $S(\mathbf{Q}, \hbar\omega)$ shows an approximate cylindrical symmetry, where the longitudinal axis is along the \mathbf{x}' axis $\{[1, 1, 0]_{\text{hex}}\}$, and the reference plane aligns with $\mathbf{y}' - \mathbf{z}' = [1, -1, 0]_{\text{hex}} - [0, 0, 1]_{\text{hex}}$ plane. Therefore, the integrated intensity, $S_{\text{calc}}(\hbar\omega)$, can be approximated as,

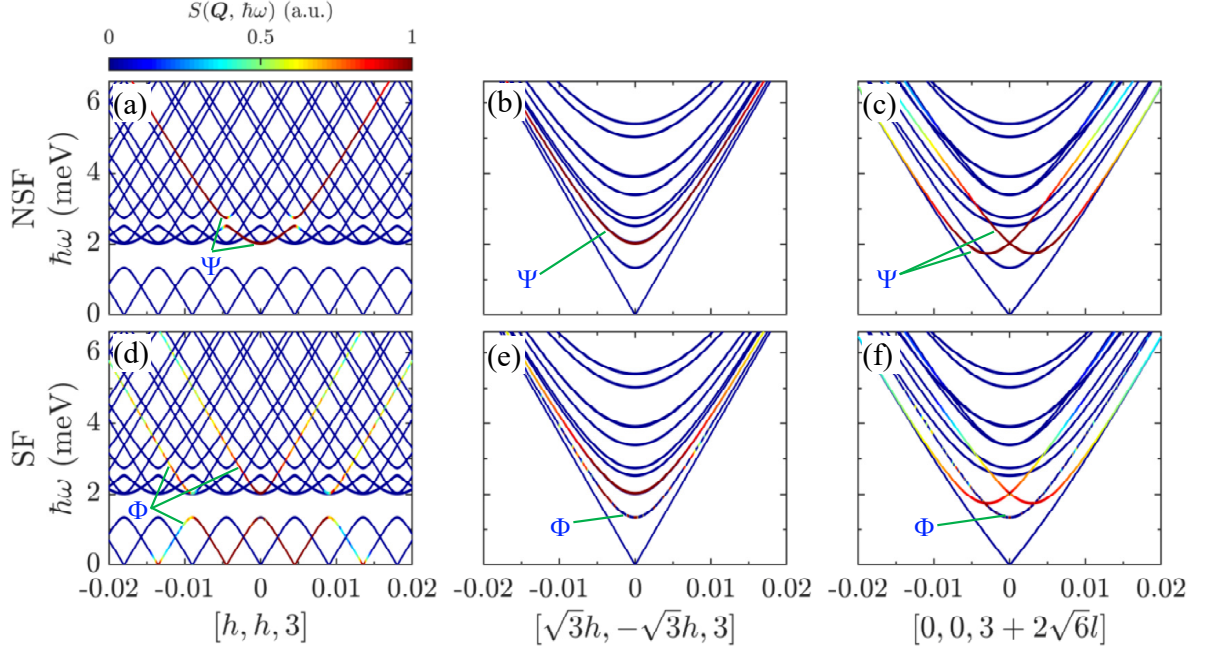


FIG. 4. Magnetic dispersion relations in the vicinity of $(0,0,3)_{\text{hex}}$ calculated using the linear spin wave theory for BiFeO_3 . The spin wave energies and intensities were calculated using the equations-of-motion technique for non-collinear spins [31] on a pseudo-commensurate magnetic super cell containing $M = 2 \times 222$ pseudo-cubic cells [18]. Panels (a)-(c) and (d)-(f) show the calculated spin wave intensity, $S(\mathbf{Q}, \hbar\omega)$, as a function of the momentum transfer \mathbf{Q} and the energy transfer $\hbar\omega$ for the NSF and SF components, respectively, along three orthogonal directions, $[h, h, 3]_{\text{hex}}$, $[\sqrt{3}h, -\sqrt{3}h, 3]_{\text{hex}}$, and $[0, 0, 3 + 2\sqrt{6}l]_{\text{hex}}$, which are consistent with the geometry used in Fig. 2. The wave vector components, h and l are rescaled so that they display the equivalent unit in the horizontal axis.

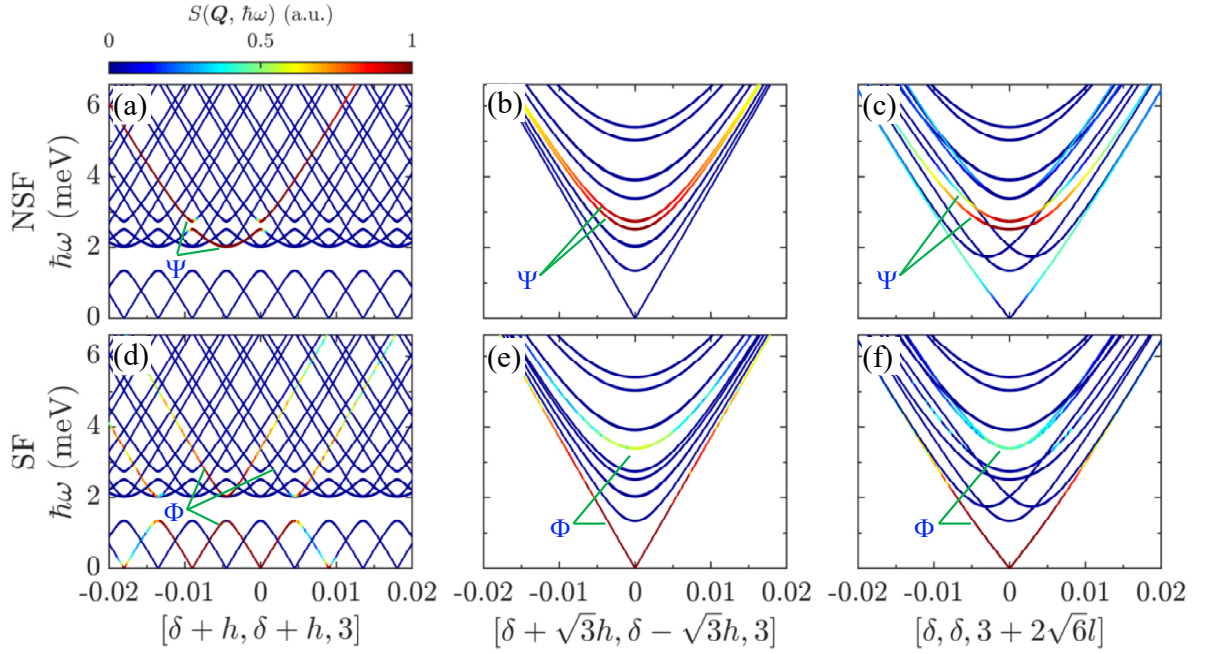


FIG. 5. Magnetic dispersion relations in the vicinity of $(\delta, \delta, 3)_{\text{hex}}$ calculated using the linear spin wave theory for BiFeO_3 . Panels (a)-(c) and (d)-(f) show the calculated spin wave intensity, $S(\mathbf{Q}, \hbar\omega)$, for the NSF and SF components, respectively, along three orthogonal directions, $[\delta + h, \delta + h, 3]_{\text{hex}}$, $[\delta + \sqrt{3}h, \delta - \sqrt{3}h, 3]_{\text{hex}}$, and $[\delta, \delta, 3 + 2\sqrt{6}l]_{\text{hex}}$.

$$\begin{aligned}
S_{\text{calc}}(\hbar\omega) &= \int d^3\mathbf{Q} S_{\text{calc}}(\mathbf{Q}, \hbar\omega) \\
&= \int d\eta \int d\xi S_{\text{calc}}(\xi, \eta, \hbar\omega),
\end{aligned} \tag{5}$$

where η , representing the coordinate along the longitudinal axis, is along the \mathbf{x}' , and ξ represents the coordinate along an arbitrary polar axis ξ , which is within the reference plane $\mathbf{y}' - \mathbf{z}'$. $S_{\text{calc}}(\hbar\omega)$ is averaged over three ξ orientations, i.e., \mathbf{z}' , $\mathbf{y}' + \mathbf{z}'$, and $\mathbf{y}' - 2\mathbf{z}'$. Then, the integral is taken over $-0.02 \leq \eta \leq +0.02$ and $0 \leq \xi \leq +0.02$ to include most of the spin-wave excitations below 6 meV (Figs. 4 and 5), as the instrumental \mathbf{Q} resolution integrates experimentally. Finally, the instrumental energy resolution is convoluted with the \mathbf{Q} integrated excitation data.

As shown in Fig. 3, the observed NSF and SF components are more intense above and below ~ 1.5 meV, respectively. The calculated NSF and SF components reproduce the overall feature of the observed ones reasonably well, although the detailed structure cannot be fully reproduced. In particular, a dip observed around 1.8 meV, possibly corresponding to the gap around 1.8 meV shown in Figs. 5(a) and 5(d), is not reproduced. This is partly because the calculated intensities are not integrated perfectly well. The interaction parameters might need to be adjusted in order to obtain better agreement between the experimental and calculated results. However, the integration of the calculated intensities over the wide momentum and energy ranges requires intense effort. Therefore, the refinement of the parameters is beyond the scope of this study. Another puzzling feature is that the NSF intensities observed below ~ 1.5 meV are much larger than those calculated, as shown in Fig. 3(a). Although there may be some contributions, originating from a small fraction of misoriented magnetic domains, the intensities are supposed to be minor. In principle, there can be phonon contribution in the NSF channel. However, the phonon contribution is considered to be absent since the $(\delta, \delta, 3)_{\text{hex}}$ Bragg peaks are purely magnetic in origin and acoustic phonons at low energies are not expected. A further study is required to clarify this issue.

Here, let us look into the magnetic excitations in more detail. The observed spin excitations of BiFeO₃ break down into two classes. The class Φ are in-plane cycloidal modes that roughly correspond to rotations of the spin in the cycloidal plane defined by the propagation vector \mathbf{q}_m and the polarization direction \mathbf{z}' . This class includes a Goldstone mode that requires no energy. The other class Ψ corresponds to out-of-plane rotations that correspond to oscillations primarily in the direction $\mathbf{q}_m \times \mathbf{z}'$.

As described in Sec. I, the two magnon modes, Φ and Ψ , were reported from the optical spectroscopy measurements [11, 15–17]. In Figs. 4 and 5, Φ and Ψ modes are indicated among the calculated magnon modes. The Φ mode emerges from the zero energy and has a gap around 1.8 meV. The Ψ mode is a gapped mode with a gap energy of ~ 2 meV. In Figs. 3(a) and 3(b), the calculated Φ and Ψ mode intensities are exhibited separately. The Φ mode is predicted mostly in the SF channel, corresponding to the easy plane anisotropy within the cycloidal plane. On the other hand, the Ψ mode, which emerges above ~ 2 meV, has larger contribution to the NSF channel, corresponding to the easy axis anisotropy out of the cycloidal plane. As shown in Figs. 4(c) and 4(f), this mode has nonnegligible contribution to the SF channel. Above ~ 2.5 meV, there is a large amount of contributions to both the SF and NSF channels from other excitation modes, which are predicted to be isotropic.

$\Psi_1^{(1)}$ at ~ 2.7 meV was identified to be the electromagnon in Ref. [18]. In our inelastic neutron measurements, a major part of the Ψ mode corresponds to the shoulder around 2.5 meV in the NSF channel. Therefore, this out-of-plane mode is considered to be enhanced by the photon's electric field. Besides the anisotropic Φ and Ψ modes, which have electromagnon and magnetic resonance characters, other magnon modes are found to be isotropic.

IV. CONCLUSION

Using the polarized inelastic neutron scattering technique with single magnetic domained crystals, the magnetic excitation modes distributed in and out of the cycloidal plane have been observed separately in BiFeO₃. The theoretical calculations using the linear spin wave theory identified the anisotropic and isotropic spin wave excitation modes, which can explain the overall experimental results reasonably well. Our results provide important information on how the magnetic excitations, in particular Φ and Ψ modes, observed by optical and neutron scattering spectroscopies are related.

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