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Spin-excitation anisotropy in the nematic state of detwinned FeSe

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The origin of the electronic nematicity in FeSe is one of the most important unresolved puzzles in the study of iron-based superconductors. In both spin- and orbital-nematic models, the intrinsic magnetic excitations at $Q_1 = (1,0)$ and $Q_2 = (0,1)$ of twin-free FeSe are expected to provide decisive criteria for clarifying this issue. Although a spin-fluctuation anisotropy below 10 meV between Q_1 and Q_2 has been observed by inelastic neutron scattering at low temperature, it remains unclear whether such an anisotropy also persists at higher energies and associates with the nematic transition T_s . Here we use resonant inelastic x-ray scattering to probe the high-energy magnetic excitations of detwinned FeSe. A prominent anisotropy between the magnetic excitations along the H and K directions is found to persist to $E \approx 200$ meV, which decreases gradually with increasing temperature and finally vanishes at a temperature around T_s . The measured high-energy spin excitations are dispersive and underdamped, which can be understood from a local-moment perspective. Taking together the large energy scale far beyond the d_{xz}/d_{yz} orbital splitting, we suggest that the nematicity in FeSe is likely spin-driven.

Intertwined order and fluctuations in high-transition-temperature (T_c) superconductors are pivotal for understanding the microscopic origin of superconducting electron pairing¹. Of particular interest is the electronic nematic state present in both cuprate and iron-based superconductors (FeSCs). Initially discovered through in-plane electronic anisotropy with C_2 symmetry in the paramagnetic orthorhombic state of detwinned Ba(Fe_{1-x}Co_x)₂As₂ (refs.²⁻⁴), the electronic nematic state (and its fluctuations) has been identified as a ubiquitous feature of FeSCs and is believed to be essential for the structural and magnetic transitions in FeSCs (refs.⁵⁻⁹), and may enhance the electron pairing for high- T_c superconductivity^{10–12}.

Iron selenide (FeSe) is a unique material among FeSCs because of its simple structure (Fig. 1a)¹³ and unusual electronic properties, such as strong anisotropy of the superconducting order parameter, extended electronic nematic phase and highly tunable T_c (ref. ⁹). In particular, unlike iron pnictide, which has a collinear antiferromagnetic (AF) ground state below the tetragonal-to-orthorhombic structural (nematic) phase transition T_s , although FeSe exhibits a similar nematic transition ($T_s \approx 90$ K)¹⁴, it has no static AF order, providing a broad temperature range below T_s as an ideal platform for investigating electronic nematicity and its interplay with superconductivity.

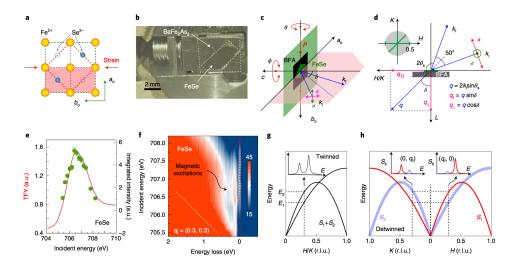


Fig. 1 Crystal structure, detwinning strategy, scattering geometry and incident energy-dependent RIXS. a, Structure of the FeSe layer. The FeSe crystal consists of stacked FeSe layers. The yellow-filled circles represent Fe²⁺ ions. The filled and open blue circles denote Se²⁻ ions above and below the Fe-Fe plane, respectively. The horizontal arrows mark the direction of the uniaxial strain. The red dashed diamond and green dashed square denote the tetragonal and orthorhombic unit cells, respectively. b, A mechanical detwinning device installed on the RIXS spectrometer, with a thin FeSe crystal glued on a pre-cleaved BaFe₂As₂ crystal that is pressured in the device. c.d. Scattering geometry for RIXS measurements. \mathbf{q}_{\parallel} and \mathbf{q}_{\perp} are projections of \mathbf{q} onto [H, K] plane and L direction. The sample rotation θ around the vertical axis controls the in-plane momentum transfer \mathbf{q}_{\parallel} and the rotation ϕ around the c axis can tune the scattering plane (pink area) (c). The grey-filled circle in d marks the momentum area accessible in this study ($|\mathbf{q}_{\parallel}| \lesssim 0.5$). The green lines in the grey area in **d** show the high-symmetry directions for RIXS measurements. P denotes uniaxial pressure; $k_{i/f}$, initial/final wave vector of photon; and BFA, BaFe₂As₂. e, The red curve is the total fluorescence yield (TFY) XAS spectrum of FeSe collected near the Fe-L₃ edge (left axis). The green-filled circles represent the integrated intensity (right axis) of the magnetic excitations shown in f. The error bars in e are estimated from the statistics of the data and the 95% confidence interval of the fittings. f, Incidentenergy dependence of the excitations of FeSe at $\mathbf{q}_{\parallel} = (0.3, 0.3)$, measured near the Fe-L₃ edge with π polarization at T = 20 K. The magnetic excitations are marked by a curved arrow. g,h, Schematics of magnetic excitation dispersions for twinned (g) and detwinned (h) FeSe assuming a difference between spin excitations S_2 and S_1 . Correspondingly, the insets in g and h show schematic RIXS energy spectra for twinned and detwinned FeSe.

FeSe consists of stacked charge-neutral FeSe layers (Fig. 1a). On cooling, it undergoes a nematic transition at $T_{\rm s}\approx 90$ K, below which twin domains form along two mutually perpendicular directions and exhibit macroscopic four-fold symmetry, impeding the study of the intrinsic electronic properties of the orthorhombic (nematic) state. Through detwinning of FeSe using uniaxial strain, resistivity and electronic structure measurements reveal strong electronic anisotropy in the nematic state ^{15,16}. At lower temperature, FeSe enters a superconducting ground state with $T_{\rm c}\approx 9$ K, in which a superconducting energy gap anisotropy has been observed via angle-resolved photoemission spectroscopy (ARPES) ^{17–19} and scanning tunnelling spectroscopy²⁰.

In the presence of orbital splitting (or orbital ordering) between d_{xz} and d_{yz} orbitals 16,21,22 , and the absence of magnetic order, the electronic nematic phase has been suggested to be driven by orbital fluctuations $^{23-26}$. On the other hand, experimental evidence, in particular the discovery of intense magnetic excitations and their correlation with the nematic transition $^{27-29}$, has emphasized the importance of the spin degree of freedom in driving electronic nematic order. In addition, various localized models based on quantum paramagnetism, spin frustration and magnetic quadrupolar order have been proposed to account for the nematic transition, magnetic excitations and the absence of AF

order $^{30-33}$.

It is proposed that resolving the intrinsic magnetic excitations in twin-free FeSe is key to clarifying the microscopic origin of the unusual electronic anisotropy in both the superconducting and nematic states^{5,6,34}. By employing BaFe₂As₂ as a substrate for applying uniaxial strain, some of us have recently measured low-energy spin fluctuations ($E \lesssim 10 \text{ meV}$) of detwinned FeSe using inelastic neutron scattering³⁵. These results have revealed anisotropic spin fluctuations in the normal state and a spin resonance appearing only at $\mathbf{Q}_1 = (1,0)$ below T_c , consistent with the picture of orbital-selective Cooper pairing^{20,36-38}. However, neutron scattering experiments were unable to determine what happens to the magnetic excitation anisotropy across the nematic transition due to enhanced background scattering from the large aluminium detwinning device on warming to T_s . In addition, the energy scale of the magnetic anisotropy is unknown because the background magnetic scattering from the BaFe₂As₂ substrate overwhelms the magnetic signal from FeSe for energies above 10 meV (ref. ³⁵).

An ideal method to probe the intrinsic magnetic excitations of FeSe is resonant inelastic X-ray scattering (RIXS) at the Fe-L₃ edge in combination with the aforementioned detwinning method (Fig. 1b)³⁹⁻⁴⁴. RIXS at transition-metal L edges has been widely used to study the (para)magnons of cuprate and FeSCs, as well as various elementary excitations including phonons, crystal-field excitations and plasmons³⁹⁻⁴⁹. Because Fe-L₃ X-rays (707 eV) penetrate less than 100 nm into FeSe, but the typical thickness of a cleaved FeSe single crystal is ~20 μm, RIXS studies of FeSe are free from signal contamination due to BaFe₂As₂ and provide a unique opportunity for measuring high-energy magnetic excitations on detwinned FeSe with high efficiency.

In this Article we use RIXS to measure the intrinsic spin excitations of FeSe and BaFe₂As₂ along the high-symmetry directions H, K and [H, H], denoted by $S_h(q_{\parallel})$, $S_k(q_{\parallel})$ and $S_{hh}(q_{\parallel})$, respectively (Figs. 2 and 3). To facilitate discussions, we define the spin excitations associated with $\mathbf{Q}_1 = (1, 0)$ as $S_1(\mathbf{q}, E)$ and those associated with $\mathbf{Q}_2 = (0, 1)$ as $S_2(\mathbf{q}, E)$. The ratio between $S_h(q_{\parallel})$ and $S_k(q_{\parallel})$, $\psi(q_{\parallel}) = S_h(q_{\parallel})/S_k(q_{\parallel})$, directly probes the spin-excitation unbalance between S_1 and S_2 , which is commonly referred to as nematic spin correlations in the nematic ordering and fluctuating region^{34,50,51}. We denote the momentum transfer in reciprocal lattice units (r.l.u.; see Experimental set-ups in the Methods).

Our results reveal that the spin-excitation anisotropy in detwinned FeSe manifests over a large energy range up to 200 meV. It persists up to a temperature slightly above T_s , before fading away at a temperature well above T_s . Its comparison with the intrinsic spin-wave anisotropy of BaFe₂As₂ establishes strong nematic spin correlations in both energy scale and amplitude in FeSe. This strong spin-excitation anisotropy builds a direct connection with the nematic phase, suggesting that the nematic order is primarily spin-driven because the energy range of the nematic-phase-induced spin-excitation anisotropy is much larger than that of orbital splitting^{22–24}. Furthermore, the RIXS results identify dispersive high-energy spin excitations that are underdamped, which is highly peculiar for a paramagnet, but can be understood from a local-moment-based model with antiferroquadrupolar (AFQ) order³¹. As such, our results provide much-needed new insights into the mechanism for the nematicity of FeSe.

Figure 1b shows an FeSe crystal that has been prepared for RIXS measurements, and is glued onto

a square-shaped BaFe₂As₂ sample with the same orientation. Uniaxial pressure is applied on the BaFe₂As₂ along the tetragonal [110] direction (orthorhombic *b* axis). On cooling, the BaFe₂As₂ will be detwinned below $T_s \approx 138$ K and generates an orthorhombic distortion $\delta = (a - b)/(a + b) = 0.36\%$ that can be transferred to and thereby detwin FeSe below its structural transition at $T_s \approx 90$ K. Figure 1c,d illustrates the scattering geometry, the substantial area of the first Brillouin zone accessible with Fe-L₃ RIXS, and calculations of the in-plane momenta q_{\parallel} .

We first carried out incident-energy-dependent RIXS (energy detuning) measurements for both BaFe₂As₂ and FeSe around their resonating energies, which were determined by X-ray absorption spectroscopy (XAS; Fig. 1e and Supplementary Fig. 2). Figure 1f presents a RIXS map for an unstrained FeSe sample measured at $\mathbf{q}_{\parallel} = (0.3, 0.3)$ with π polarization and T = 20 K. Although fluorescence and particle-hole excitations dominate the scattering signal above $E \approx 0.5$ eV, a clear intrinsic elementary excitation (Raman mode) is observed at $E \approx 160$ meV that is well separated from the fluorescence peak setting above $E \approx 200$ meV. The integrated intensity of this Raman mode (Fig. 1e, green circles) follows the XAS (Fig. 1e, red curve), indicating that the cross-section is enhanced near the Fe-L₃ edge. In addition, phonon contributions to this mode can also be excluded because of their much smaller energy scale ($E \approx 40$ meV)⁴³. Taking together the consistency in energy dispersion between the excitations in Figs. 2 and 3 and those reported in previous studies^{39,43,44}, we can safely attribute these dispersive excitations to single spin-flip magnetic excitations.

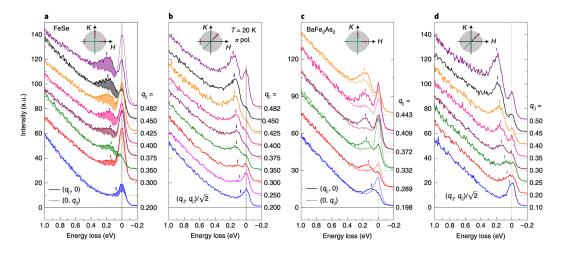


Fig. 2 Summary of RIXS results on detwinned FeSe and BaFe₂As₂. a,b, Momentum-dependent RIXS spectra of FeSe along the H (solid lines) (a) and K (dashed lines) directions (a) and twinned FeSe along the [H, H] direction (b). The shaded areas mark the difference between $S_h(q_{\parallel})$ and $S_k(q_{\parallel})$. c,d, Momentum-dependent RIXS spectra of detwinned BaFe₂As₂ along the H/K (c) and [H, H] (d) directions. The coloured dots in the insets mark the momenta where spectra were collected.

To quantitatively determine the spin-excitation anisotropy in FeSe, we measured the spin excitations of both detwinned FeSe and BaFe₂As₂ samples (Figs. 2 and 3), taking the latter as a reference; this can be compared to previous neutron scattering studies of detwinned BaFe₂As₂ (ref.⁵⁰). Before discussing the results, we illustrate in Fig. 1g,h the principle for resolving the inherent spin-wave anisotropy and nematic spin correlations in the first Brillouin zone (Supplementary Fig. 3). In a twinned sample with anisotropic excitations, RIXS measurements generate $S_h = S_k$ (black curve in the inset of

Fig. 1g). Both S_h and S_k consist of two spin-excitation branches from S_1 and S_2 in twin domains, where we assume S_1 and S_2 have the same energy dispersion for simplicity. For a detwinned sample with local-moment AF order, the spin waves emanate only from the AF vector $\mathbf{Q}_1 = (1, 0)$ ($S_2 = 0$), and S_h/S_k determines the inherent spin-wave anisotropy of S_1 . In a detwinned system with nematic spin correlations ($S_1 \neq S_2 \neq 0$), S_1 and S_2 will be present along both H and K directions (Fig. 1h) with different spectral weights, for which the ratio between S_h and S_k reflects the nematic spin correlations (Supplementary Fig. 3).

Momentum-dependent RIXS spectra of FeSe and BaFe₂As₂ collected at T=20 K < T_s are summarized in Fig. 2. In both samples, highly dispersive magnetic excitations along three high-symmetry directions H, K and [H, H] are resolved. Figure 2a (2c) displays the intrinsic spin excitations of FeSe (BaFe₂As₂) along the H and K directions, and Fig. 2b (2d) along the [H, H] direction. Although twinned FeSe and BaFe₂As₂ are expected to show four-fold symmetric magnetic excitations 39,52 [$S_h(q_{\parallel}) = S_k(q_{\parallel})$], we find that the detwinned samples exhibit highly anisotropic excitations with $S_h(q_{\parallel}) > S_k(q_{\parallel})$. The substantial difference between $S_h(q_{\parallel})$ and $S_k(q_{\parallel})$ for FeSe persists at all q_{\parallel} measured (Fig. 2a, shaded areas), thus demonstrating the existence of a high-energy spin-excitation anisotropy (nematic spin correlations) in the nematic state of detwinned FeSe. For BaFe₂As₂, the spin-excitation anisotropy is a manifestation of the inherent difference in the spin-wave branches along the H and K directions 50 . The spin-wave anisotropy of BaFe₂As₂ increases with q_{\parallel} up to 0.409 but decreases at higher $q_{\parallel} = 0.443$ (Fig. 2c), revealing a non-monotonic momentum dependence. In comparison, the spectral weight difference in FeSe (Fig. 2a, shaded area) retains a large amplitude at an even higher $q_{\parallel} = 0.482$.

To achieve a quantitative characterization of the spin-excitation anisotropy in FeSe and BaFe₂As₂, we use a general damped harmonic oscillator model^{39–41,43}

$$S(q,E) = A \frac{E_0}{1 - e^{-\beta E}} \frac{2\gamma E}{\left(E^2 - E_0^2\right)^2 + (\gamma E)^2}$$
(1)

to fit the magnetic excitations, in which E_0 is the undamped energy, γ the damping factor, $\beta = \frac{1}{k_B T}$ (k_B is the Boltzmann constant) and A is a fitting coefficient. The elastic peak can be fitted with a Gaussian function, and the fluorescence contributions below $E \approx 1$ eV can be described with a quadratic polynomial (Supplementary Figs. 5 and 6).

The fitting results for FeSe and BaFe₂As₂, the undamped energy dispersion ($E_0(q)$), the energies for the intensity maxima ($E_m(q)$) and the damping factor ($\gamma/2$) are summarized in Fig. 3. Figure 3a-c shows a simulation of the spin waves in BaFe₂As₂ overlaid by the energy dispersions and the damping factors. The simulation is based on an anisotropic Heisenberg $J_{1a}-J_{1b}-J_2$ model as described in ref. ⁵² (J_{1a} (J_{1b}) is the nearest-neighbour exchange interaction along a (b) axis. J_2 is the next-nearest-neighbour exchange interaction.), in which we set L=0 because the spin waves of BaFe₂As₂ are two-dimensional, especially for the high-energy spin excitations probed in our RIXS measurements. We find $E_0 > \gamma/2$ at all the momenta measured, which indicates that the spin waves are underdamped and far from being critically damped. The intrinsic spin waves of the clearly resolved two different branches $S_h(q)$ and $S_k(q)$

around Γ are consistent with the anisotropic Heisenberg model, in which dispersive spin waves can only arise from the AF wave vector \mathbf{Q}_1 (S_1). The minor deviation of the branch $S_k(q)$ from the anisotropic Heisenberg model can be attributed to the failure of the anisotropic Heisenberg model in describing the small anisotropy of magnetic excitations at high energy, due to the emergence of the spin excitations around $\mathbf{Q}_2 = (0, 1)$ (S_2) at $E \geq 100$ meV observed by neutron scattering 50,51 . Figure 3d shows the energy dispersion of FeSe obtained from the fitting of the magnetic excitations shown in Fig. 2a,b. The filled symbols mark the bare energy dispersion (E_0) without damping effect, and the open symbols represent the energy dispersion for the intensity maxima E_m .

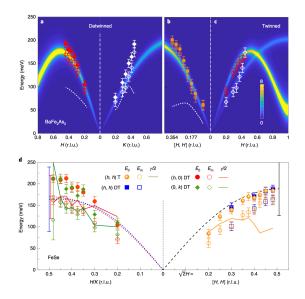


Fig. 3 Energy dispersions and damping factors for the magnetic excitations of FeSe and BaFe₂As₂. a-c, Comparison between dispersions of the magnetic excitations measured with RIXS and a simulation with a Heisenberg model for detwinned (a) and twinned (b,c) BaFe₂As₂ (ref.⁵²). The filled symbols, open symbols and white dashed curves in a and b are undamped energies (E_0), energies for the intensity maxima (E_m) and the damping factor (γ /2) fitted from the spectra shown in Fig. 2c,d with equation (1). The open symbols in c show E_m . d, Spin-excitation energy dispersions of FeSe obtained from the fitting of the RIXS spectra in Fig. 2a,b with equation (1). The filled (open) red circles, green diamonds, orange circles, and blue squares mark the undamped energies E_0 (energies for intensity maxima E_m) of the dispersions along H (detwinned, DT), K (detwinned), [H, H] (twinned, T) and [H, H] (detwinned) directions, respectively. The red, green, and orange lines mark the momentum-dependent damping factor γ /2 along the high-symmetry directions. The black, red, and blue dashed curves are the calculated flavor-wave dispersions in the AFQ phase along the [H, H], and H/K directions, respectively. The error bars for the data points are estimated according to the confidence interval of the fittings and the error in determining the E = 0 position. The error bars mark the damping factor γ /2 along the corresponding directions.

The momentum-dependent damping factors $\gamma/2$ are overall larger than that for BaFe₂As₂, but still in the underdamped regime for most of the excitations. Moreover, the damping factor for S_{hh} (along the [H, H] direction) is smaller than for $S_{h/k}$ in both FeSe and BaFe₂As₂, suggesting a common anisotropic damping effect in FeSC. The (anisotropic) underdamped nature of the magnetic excitations, not inferable in previous neutron scattering and RIXS studies on twinned samples^{28,43}, suggests that the excitations should arise from interacting local moments associated with strong electron correlations⁵³.

Because the largest momentum (0.482, 0) along H is close to the zone boundary (0.5, 0), the excitation energy scale of FeSe ($E_0 \approx 200$ meV and $E_m \approx 160$ meV) at (0.482, 0) and (0, 0.482) (consistent with that in ref.⁴³) reveals a much higher band top in the first Brillouin zone than that ($E \approx 100$ meV)

120–150 meV) observed by neutron scattering in the Brillouin zone around $(1, 0)^{28}$. This is in stark contrast to the case of BaFe₂As₂, where the dispersions measured by RIXS and neutron scattering can be consistently described with one model (Fig. 3)^{39,50}. We attribute this difference to the absence of stripe AF order in FeSe. Because of the translational symmetry of the stripe AF order ($\mathbf{k} = (1, 0)$) in BaFe₂As₂, the magnetic excitations in the first Brillouin zone can be deemed as a replica of the ones in the Brillouin zone centring at Q = (1, 0). However, long-range AF order is not established in spite of the strong in-plane magnetic correlations in FeSe. Accordingly, the dispersions around Γ and (1, 0) do not have to be identical. However, how to quantitatively reconcile the neutron scattering and RIXS results is still an open question.

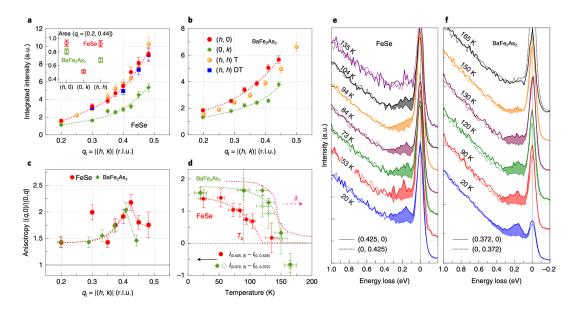


Fig. 4 Anisotropic magnetic excitations in detwinned FeSe and BaFe₂As₂. a,b, Momentum-dependent energy-integrated intensity of magnetic excitations for FeSe (a) and BaFe₂As₂ (b). The inset shows the momentum-integrated intensity of $S_{h/k/hh}$ for FeSe (red circles) and BaFe₂As₂ (green squares) in the range of $q_{\parallel} = [0.2, 0.44]$. c, Spin excitation anisotropy between S_h and S_k , defined as the ratio between the integrated intensity for (q, 0) and (0, q) as shown in a and b. d, Temperature dependence of the spin-excitation difference between $S_h(q_{\parallel})$ and $S_k(q_{\parallel})$, in which the data points are the integrated intensity of $S_h(q_{\parallel}) - S_k(q_{\parallel})$ in the energy range of [0.08, 0.4] eV, as shown in e and f. The dashed red and green lines in c and d are guides to the eye. The pink dashed curve shows the lattice distortion $\delta = (a-b)/(a+b)$ of BaFe₂As₂ under a uniaxial pressure of P~20 MPa, which reaches 0.36% below $T\sim100$ K. The vertical black dashed line marks the $T_s = 90$ K for FeSe. The vertical error bars in a-d are estimated from the data statistics and the confidence interval of the fittings, which determine the error bars in c. The horizontal bars in d mark the temperature range where data were collected. e,f, Temperature-dependent RIXS spectra for FeSe and BaFe₂As₂ measured at $q_{\parallel} = 0.425$ and 0.372, respectively. The shaded areas in e and f mark the intensity difference between $S_h(q_{\parallel})$ and $S_k(q_{\parallel})$.

Figure 4a,b shows the momentum-dependent energy-integrated intensity of the spin excitations S_h , S_k and S_{hh} for FeSe and BaFe₂As₂, respectively. With increasing q_{\parallel} , the integrated intensities along all three high-symmetry directions increase monotonically. The spin-excitation intensity for FeSe is slightly higher than for BaFe₂As₂, qualitatively consistent with previous neutron scattering results²⁸. Note that the momentum-direction dependence of the amplitude for $S_{h/k/hh}$ is modulated by the anisotropic damping factor inherent to $S_{1,2}$, which requires $S_{1,2} = 0$ at Γ (ref. ⁵²). To quantify the spin-excitation anisotropy, we plotted the ratio $I(q, 0)/I(0, q) = S_h/S_k$ for FeSe and BaFe₂As₂ in the same panel (Fig. 4c). It is surprising that the spin-excitation anisotropy of FeSe (reflecting the nematic spin

correlations) is rather similar to the spin-wave anisotropy of BaFe₂As₂ in both amplitude and energy scale. Furthermore, in the high-energy/momentum region, the anisotropy of FeSe is even larger than for BaFe₂As₂. This prominent spin-excitation anisotropy signifies strong electronic nematicity with large energy scale ($E \approx 200 \text{ meV}$) and magnitude in FeSe. Moreover, the temperature dependence of the difference between S_h and S_k for FeSe at selected $q_{\parallel} = 0.425$ decreases with increasing temperature, persists to a temperature (104 K) 20% higher than T_s and finally drastically reduces at a temperature (135 K) well above T_s , indicating a gradual suppression of nematic spin correlations above T_s (Fig. 4d,e). Because FeSe is under uniaxial strain applied from the BaFe₂As₂ substrate (pink dashed curve, Fig. 4d), T_s is no longer well defined. For this reason, it is not surprising that the spin-excitation anisotropy disappears at a temperature above the zero-pressure T_s . This temperature dependence is similar to that for BaFe₂As₂, as shown in Fig. 4d,f, where the anisotropy at $q_{\parallel} = 0.372$ persists at a temperature slightly higher than T_N but vanishes at T = 165 K.

Previous measurements of the spin dynamics in detwinned FeSe were constrained to low energies $(E \leq 10 \text{ meV})^{35}$ and were unable to discriminate between the different scenarios for the nematicity in FeSe. Our present measurements over a large energy window enable the discovery that the detwinned FeSe harbours high-energy (up to 200 meV) anisotropic spin excitations that are dispersive and underdamped. This surprising finding appears to be at variance with the itinerant mechanism for the nematicity in FeSe. In the itinerant picture, the leading contribution to the spin excitation spectrum comes from a two-particle process, that is, a convolution of single-electron and single-hole excitations. Because of this particle-hole nature of the spin excitations, damping already appears in the leading-order contribution to the spin excitations is generically already overdamped; higher-order processes further enhance the damping. An itinerant model calculation indeed shows that the spin excitations at such pertinent (high) energies are highly overdamped⁵⁴. Our work is expected to motivate further studies on this issue within the itinerant description.

Instead, the dispersive and underdamped nature of the high-energy spin excitations point to a local-moment starting point to describe the nematicity. Taking together that the energy scale of the spin-excitation anisotropy is far beyond the d_{xz}/d_{yz} orbital splitting ($E \approx 50$ meV), and the nematic spin correlations decrease and finally disappear at a temperature above T_s , we conclude that the nematicity is probably spin-driven.

This has led us to consider a generalized bilinear-biquadratic model on a square lattice with local moments³¹. Here, the local moments capture the majority of the spin degrees of freedom, especially at high energies. Indeed, the importance of electron correlations in iron-based superconductors has been clearly demonstrated in recent years. Although several specific theoretical approaches have been taken to address the effect of electron correlations, including Hund's metal⁵⁵, proximity to an orbital-selective Mott phase^{6,56} and a Mott–Hund's picture⁵⁷, they share the common feature that the single-electron states have a mixture of incoherent and coherent states (which dominate at high and low energies, respectively). Indeed, for FeSe, ARPES has directly observed the high-energy Hubbard bands^{58,59}, which mediate effective exchange interactions between quasi-localized moments⁶⁰.

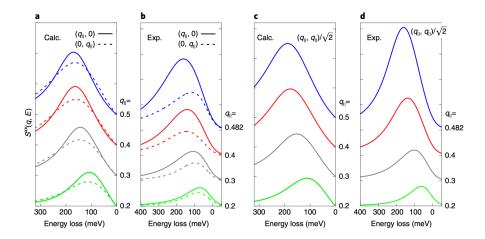


Fig. 5 Calculated spin excitation spectra of the AFQ phase and their comparison with the fitting curve of the experimental $S(q_{\parallel})$ from RIXS. a,c, Calculated spectra along the H/K directions (a) and the [H, H] direction (c). b,d, Fitting curves of the experimental $S(q_{\parallel})$ along the H/K directions (b) and [H, H] direction (d).

In the generalized bilinear-biquadratic model, the leading-order contributions to the spin excitation spectrum comprise a one-particle process. Because only sub-leading contributions that involve more-than-one-particle processes contribute to damping, the high-energy spin excitations are underdamped. More specifically, we have calculated the spin dynamics in the proposed picture for the nematicity based on a (1, 0) antiferroquadrupolar state using the flavor-wave method³¹. This approach treats the spin dipolar and quadrupolar excitations on an equal footing, as the bosonic flavor waves. Consistent with the physical picture, the spin-excitation spectrum is dispersive and underdamped (for details see the Supplementary Information). In Fig. 5 we show that the calculated spin dynamics provide a good understanding of the experimental data. Thus, our results provide evidence for a local-moment-based picture for the nematicity of FeSe.

The rich properties of the Fe-based superconductors in general and nematic FeSe in particular have been addressed from both weak^{5,37,61} and strong^{6,30,31} coupling perspectives, with each having had successes. Our work brings about a hitherto unknown and remarkable feature of universality across Febased superconductors. For BaFe₂As₂ with collinear AF order, the anisotropy can be readily interpreted in the picture of intrinsic spin waves^{50,51}. FeSe, by contrast, lacks long-range magnetic order even though it is nematic. The local-moment-based understanding, as dictated by the qualitative similarities in the high-energy spin excitations of FeSe and BaFe₂As₂, suggests that the largest spin spectral weight in both the iron chalcogenides and pnictides is associated with the incoherent electronic excitations induced by the underlying electron correlations. As such, our work not only allows for discriminating the proposed mechanisms for the nematicity of FeSe, but also points to a unified understanding of the correlation physics ⁴² across the seemingly distinct classes of Fe-based superconductors.

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Methods

Sample preparation. The high-quality BaFe₂As₂ and FeSe single crystals used in the present study were grown using self-flux and the chemical vapour transport method, respectively. The BaFe₂As₂ single crystals were oriented using a Laue camera and cut along the tetragonal [110] and [1-10] directions using a high-precision wire saw (WS-25). The directions of the self-cleaved edges of the FeSe single crystals were also determined using a Laue camera. The well-cut BaFe₂As₂ crystals, with typical dimensions of 5 mm × 4.3 mm × 0.5 mm, were pre-cleaved before the final preparation. For RIXS measurements of BaFe₂As₂, we placed a ceramic top post onto the upper surface of BaFe₂As₂ for in situ cleaving. For RIXS measurements of FeSe, we glued thin FeSe crystals onto the upper surface of BaFe₂As₂ along the same direction using epoxy Stycast 1266 and a small ceramic top post onto the surface of the FeSe. The prepared crystals with the posts were inserted into the slot of the uniaxial-pressure devices, which were mounted on a modified copper sample holder of the RIXS spectrometer (Fig. 1b)⁶².

Experimental set-ups. The RIXS and XAS measurements were performed with the RIXS spectrometer at the ADRESS beamline of the Swiss Light Source at the Paul Scherrer Institut^{63,64}. The beam size at the sample position was $4 \times 55 \mu m^2$. All the measurements shown in the main text were collected using linear horizontal (LH) polarization (electric field vector of the incident photons lying within the horizontal scattering plane), denoted as π polarization. The RIXS spectra were collected with a grazing-incidence configuration, as shown in Fig. 1d. The scattering angle was set to $2\theta_s$ = 130°, with which a substantial area of the first Brillouin zone is accessible (Fig. 1d, grey circle). The measurements were performed along high-symmetry directions H/K and [H, H] in orthorhombic notation. The total energy resolution for the RIXS measurements was set to 80 meV. The in-plane momentum q_{\parallel} could be tuned continuously by rotating the sample and thereby changing the angle δ . Before the measurements, the sample holder was inserted into the manipulator head and the top post was removed by cleaving at low temperature (~20 K) and in ultrahigh vacuum (<10⁻¹⁰ mbar). We defined the wave vector \mathbf{Q} in reciprocal space as $\mathbf{Q} = H\mathbf{a}^* + K\mathbf{b}^* + L\mathbf{c}^*$, where H, K, L are Miller indices and $\mathbf{a}^* = \mathbf{\hat{a}} 2\pi/a_0$, $\mathbf{b}^* = \mathbf{\hat{a}} 2\pi/b_0$, and $\mathbf{c}^* = 2\pi/c$ are reciprocal lattice unit vectors with $a_0 \approx 5.334$ Å, $b_0 \approx 5.308$ Å and $c \approx 5.486$ Å. The FeSe tri-layer height o was $d \approx 5.5$ Å. The orthorhombic lattice s distortion of FeSe was $\delta = (a_o - b_o)/(a_o + b_o) \approx 0.27\%$ at a temperature well below T_s .

Data availability

All data that support the plots in this paper are available from the corresponding author upon reasonable request. Source data are provided with this paper. The data can also be found at Figshare public repository⁶⁵.

Code availability

All relevant source code is available from the corresponding author upon reasonable request.

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Author contributions

X.L. conceived this project and developed the detwinning strategy. X.L. and T.S. wrote the beamtime proposals and coordinated the experiments as well as all other project phases. X.L., W.Z., Y.T., E.P., R.L., Z.T. and T.S. carried out the RIXS experiments with the support of V.N.S. X.L. analysed the data with assistance from Y.S. P.L., R.L. and Z.T. prepared the BaFe2 As 2 single crystals. T.C. and P.D. provided the FeSe single crystals. R.Y. and Q.S. carried out theoretical and computational analyses. X.L., P.D. and T.S. wrote the manuscript with input from R.Y. and Q.S. All authors made comments.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information The supplementary material is attached at the end of this manuscript.

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Supplementary information: Spin excitation anisotropy in the nematic state of detwinned FeSe

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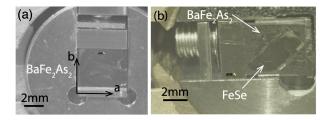


Fig. S1. Mechanical device for detwinning (a) BaFe₂As₂ and (b) FeSe installed on a RIXS sample holder on SAXES spectrometer at ADRESS beamline, SLS. Uniaxial pressure is applied via tuning the screw on one end. The pressure can be held at low temperature by a Belleville spring washer. A large BaFe₂As₂ crystal ($5 \times 4.3 \times 0.4$ mm³) was inserted into the device in (a) and a FeSe/BaFe₂As₂ sample in (b). Both samples were cleaved at ~ 20 K in high vacuum ($< 1 \times 10^{-8}$ torr).

Spin excitation anisotropy and nematic spin correlations

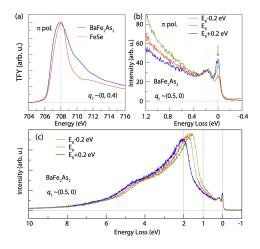


Fig. S2. (a) Comparison of the total fluorescence yield (TFY) XAS spectrum between BaFe₂As₂ (blue curve) and FeSe (red curve). The Fe-L₃ resonance energy (E_R) of FeSe is slightly (~ 0.2 eV) higher than that for BaFe₂As₂. The data were collected at $\mathbf{q}_{\parallel} \sim (0, 0.4)$ and T=20 K with π polarization. (b), (c) Brief energy detuning RIXS measurements of detwinned BaFe₂As₂ near the Fe-L₃ edge. The measurements were performed at T=20 K with $\mathbf{q}_{\parallel}=(0.5, 0)$. The vertical arrow in (b) marks the E_i -independent spin excitations near the Fe-L₃ edge, indicative of its Raman-mode nature. As a comparison, the fluorescence peak (at ~ 1.5 - 2 eV) in (c) shifts to higher energy with increasing E_i .

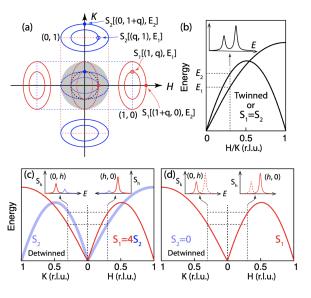


FIG. S3. (a) Reciprocal space and Brillouin zone for typical FeSC systems such as BaFe₂As₂ and FeSe. The red (blue) dashed rectangle are first BZ centering at Γ point associated with $\mathbf{Q}_1 = (1, 0)$ ($\mathbf{Q}_2 = (0, 1)$). The inner (outer) ellipses mark the constant-energy excitations with an energy transfer of E_1 (E_2). (b)-(d) Schematic dispersion for (b) a twinned sample or a sample with no nematic spin correlations ($S_1 = S_2$), (c) a detwinned sample with nematic spin correlations $S_1 > S_2$, and (d) a detwinned sample with S_1 only. The insets in (b)-(d) show the corresponding spectra S_h and S_k .

Figure 3 shows how we can connect the spin-excitation anisotropy measured with RIXS to the intrinsic nematic spin correlations. As we have described in the main text, we define the spin-excitation anisotropy as $\phi(q_{\parallel}) = \frac{\int S_h(q_{\parallel},E)dE}{\int S_k(q_{\parallel},E)dE} = S_h(q_{\parallel})/S_k(q_{\parallel})$, in which $S_h(q_{\parallel})$ and $S_k(q_{\parallel})$ are energy integrated spin excitations measured at $(q_{\parallel},0)$ and $(0,q_{\parallel})$, respectively. In our previous neutron scattering study, we have defined the nematic spin correlations as $\psi(E) = \frac{S_1(E) - S_2(E)}{S_1(E) + S_2(E)}$ (or $\psi(q,E) = \frac{S_1(q,E) - S_2(q,E)}{S_1(q,E) + S_2(q,E)}$), where $S_{1,2}(E) = \frac{\int S_{1,2}(q,E)dq}{\int dq}$, and the integral $\int dq$ runs over the whole Brillouin zone (BZ). Note q is a reduced momentum within a BZ.

As shown in Fig. S3(a), the RIXS spectra $S_h(q)$ and $S_k(q)$ measured at (q, 0) (red dot) and (0, q) (blue dot) in the first BZ, can be written as:

$$S_h(q) = S_1[(q,0), E_2] + S_2[(q,0), E_1]$$
(1)

$$S_k(q) = S_1[(0,q), E_1] + S_2[(0,q), E_2]$$
(2)

As $S_h(q) > S_k(q)$ at all the momenta measured, the lower limit for $\phi(q)$ is 1, which can be realized in a twinned sample, or a system without nematic spin correlations, as shown in Fig. S3(b). The upper limit for $\phi(q)$ is $S_1[(q,0),E_2]/S_1[(0,q),E_1]$, which can be achieved in a system where S_2 vanishes, such as the Heisenberg model calculation of the intrinsic spin waves in BaFe₂As₂ [Fig. S3(d)]. Note that

the upper limit could be different in BaFe₂As₂ and FeSe. This could undermine our comparison of the spin-excitation anisotropy between BaFe₂As₂ and FeSe. For a system with nematic state, the spin-excitation anisotropy $\phi(q)$ can reflect the amplitude and energy scale of the nematic spin correlations [Fig. S3(c)].

Fitting of the RIXS spectra

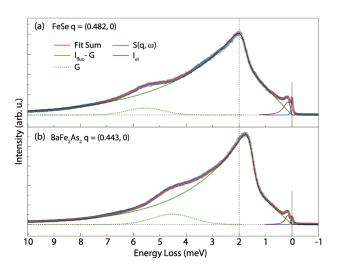


FIG. S4. Fitting of the full RIXS spectra measured at (a) $\mathbf{q}_{\parallel} = (0.482, 0)$ for FeSe, and (b) $\mathbf{q}_{\parallel} = (0.443, 0)$ for BaFe₂As₂.

To extract the energy dispersion E(q), damping rate, and intensity of the magnetic excitations, we need to use an efficient model to account for the fluorescence (I_{fluo}) , (para)magnon (S(q, E)) and elastic peak (I_{el}) of RIXS spectra. RIXS spectrum at q can be expressed as:

$$I(E) = I_{\text{fluo}} + S(q, E) + I_{\text{el}}$$
(3)

where E is energy transfer (loss), and $I_{\rm el}$ can be described with a resolution-limited gaussian peak. The RIXS spectral weight of iron-based superconductors (FeSC) are usually dominated by strong fluorescence signal at $E \gtrsim 0.5$ eV. Accurate description of fluorescence is important for extracting the magnetic scattering. Previous RIXS studies of FeSC have shown that the fluorescence at Fe $-L_3$ edge can be well described by the formula:

$$I_{\text{fluo}} = (bE^2 + aE) \cdot (1 - g_{\text{v}}) + I_0 \exp(-\alpha E) \cdot g_{\text{v}} + G \tag{4}$$

with

$$g_{\gamma} = \left(\exp\left(\frac{E + E^*}{\Gamma}\right) + 1\right)^{-1} \tag{5}$$

where $G = A_0 \exp\left(\frac{(E+E_S)^2}{2\sigma^2}\right)$ is a gaussian function. g_{γ} generates a smooth crossover from the quasilinear region with $E \lesssim 1 \, \text{eV}$ (the first term of Eq. (2)) to the exponential region (the second term of Eq. (4)) [1-3]. As shown in Fig. S4, the overall line-shape of the fluorescence can be well fitted by

Eq. (4). Because the magnetic excitations exist only at low energy ($E \leq 0.8 \text{eV}$), while the exponential decay dominates the range with much higher energy, we neglect g_{γ} and describe the low-energy fluorescence using $bE^2 + aE$ in fitting the magnetic excitations.

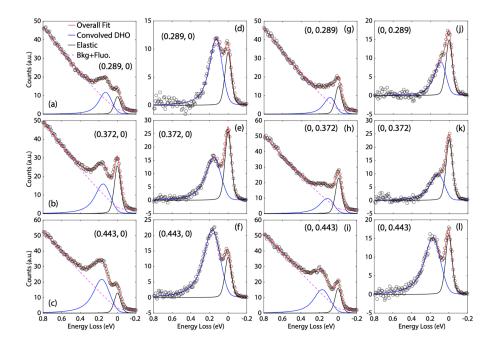


FIG. S5. Selected RIXS spectra along [H, 0] and [0, K] directions for detwinned BaFe₂As₂ and their fittings using the model as described in Eq. (4). Black circles in (a)-(c), (g)-(i) are raw RIXS spectra, and in (d)-(f), (j)-(l) are spectra with fluorescence and background subtracted. Blue curves are a damped harmonic oscillator (DHO) model convolved with an energy-resolution limited gaussian. Black curves are gaussian fittings of the elastic scattering peaks. Pink dashed curves are fluorescence and background. Red curves are the overall fit of the spectra.

Given that the magnetic excitations are heavily damped in itinerant FeSC, we use a general damped harmonic oscillator function to capture the features of the magnetic excitations [4-7]:

$$S(q,E) = A \frac{E_0}{1 - e^{-\beta E}} \frac{2\gamma E}{(E^2 - E_0^2)^2 + (E\gamma)^2},$$
(6)

where β is $\frac{1}{k_{\rm B}T}$ ($k_{\rm B}$ is Boltzmann constant). $E_0(q)$ is the undamped energy and $\gamma(q)$ the excitation life time (which represents the damping rate). When the excitations are underdamped ($\gamma/2 < E_0$), we can also describe the spectrum using an antisymmetrized Lorentzian function [5-7],

$$S(q,E) \propto \frac{\gamma/2}{(E - E_a)^2 + (\gamma/2)^2} + \frac{\gamma/2}{(E + E_a)^2 + (\gamma/2)^2},$$
 (7)

generating the fitted pole energies $E_q = [E_0^2 - (\gamma/2)^2]^{-1/2}$. In a general case, the undamped energy E_0 and the excitation life time γ are essential information we need to extract from the fitting of the

spectra.

We show in Fig. S4 the overall fitting of the RIXS spectra (blue open circles) using Eq. (4) (red curves), and in Figs. S5 and S6 the fitting of the low-energy magnetic excitations for BaFe₂As₂ and FeSe using Eq. (6). The spectra are well fitted by the damped harmonic oscillator model. The energy dispersions and integrated intensities for BaFe₂As₂ and FeSe have been shown in the main text.

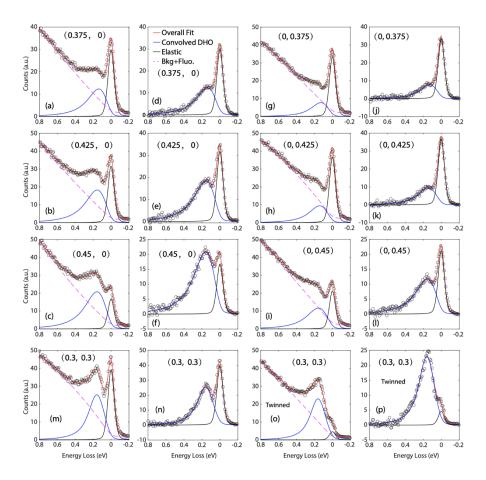


FIG. S6. Fitting of the RIXS spectra of FeSe following the same procedure as that for BaFe₂As₂ in Fig. 3. The spectra in (o) and (p) are measured on twinned sample while the others on detwinned sample.

Theoretical results

Motivated by the dispersive and underdamped feature of the RIXS spectrum in the detwinned FeSe, we study the spin dynamics of an S=1 bilinear-biquadratic Heisenberg model. The Hamiltonian reads

$$H = \frac{1}{2} \sum_{i,\delta_n,\alpha,\beta} \left\{ J_n S_i \cdot S_j + K_n \left(S_i \cdot S_j \right)^2 \right\},\tag{8}$$

where $j = i + \delta_n$, and δ_n connects a site i in a square lattice and its n's nearest neighbor sites with n = 1,2,3. Here J_n , and K_n are respectively the bilinear and biquadratic couplings between the n's

nearest neighbor spins. Unlike BaFe₂As₂, the ground state of FeSe is not magnetically ordered but can hold an antiferroquadrupolar (AFQ) order with wave vector $(\pi, 0)$ [8].

Following this, we study the spin dynamics through Schwinger bosons in an SU(3) representation [10]. At each site, we use $|-1\rangle$, $|0\rangle$, and $|1\rangle$ to denote the three eigenstates of the spin operator S^z . They are used for a time-reversal invariant basis of the SU(3) representation:

$$|x\rangle = \frac{i}{\sqrt{2}}(|1\rangle - |-1\rangle),$$

$$|y\rangle = \frac{1}{\sqrt{2}}(|1\rangle + |-1\rangle),$$

$$|z\rangle = -i|0\rangle.$$
(9)

Within this representation, three Schwinger bosons are associated with the three states, $b_{\alpha}^{\dagger}|\emptyset\rangle = |\alpha\rangle$, where $\alpha = x, y, z$, and $|\emptyset\rangle$ is the null state of the Schwinger bosons. The bosons satisfy a local constraint at each site:

$$\sum_{\alpha} b_{i\alpha}^{\dagger} b_{i\alpha} = 1 \tag{10}$$

The spin dipolar and quadrupolar operators can be written in terms of the Schwinger boson bilinears as

$$S_i^{\alpha} = -i\epsilon_{\alpha\beta\gamma} \left(b_{i\beta}^{\dagger} b_{i\gamma} - b_{i\gamma}^{\dagger} b_{i\beta} \right), \tag{11}$$

$$Q_i^{\alpha\beta} = -(b_{i\alpha}^{\dagger}b_{i\beta} + b_{i\beta}^{\dagger}b_{i\alpha}),$$

$$Q_i^{x^2 - y^2} = -(b_{ix}^{\dagger} b_{ix} - b_{iy}^{\dagger} b_{iy}), \tag{12}$$

$$Q_i^{r^2-3z^2} = \frac{1}{\sqrt{3}} \left(2b_{iz}^{\dagger} b_{iz} - b_{ix}^{\dagger} b_{ix} - b_{iy}^{\dagger} b_{iy} \right)$$
,

where α , β , and γ run over x, y, and z, and $\epsilon_{\alpha\beta\gamma}$ is the Levi-Civita symbol. The Hamiltonian can be rewritten in terms of these Schwinger bosons. The condensation of the boson(s) describes a quadrupolar order. We focus on condensing the b_x and b_y bosons in alternating columns, which specifies the $(\pi, 0)$ AFQ order of the $Q^{x^2-y^2}$ type.

We study the spin excitations in the AFQ phase by using the flavor-wave theory. Referring the detail of the calculation to Ref. [8, 9], here we outline the main results. The transverse dynamical spin structure factor is given as follows:

$$S_D^{xx}(q,\omega) = \frac{1}{2} \sqrt{\frac{A_q + B_q}{A_q - B_q}} \delta(\omega - \epsilon_q), \tag{13}$$

where the magnetic excitation spectrum is

$$\epsilon_q = \sqrt{A_q^2 - B_q^2},\tag{14}$$

$$A_q = J_1 \cos q_y + J_3 (\cos 2 \, q_x + \cos 2 \, q_y) - K_1 - 2K_3, \tag{15}$$

$$B_q = (K_1 - J_1)\cos q_y + (K_3 - J_3)(\cos 2 q_x + \cos 2 q_y). \tag{16}$$

The dispersion of the magnetic excitations for the parameter set $J_1 = J_2 = 10 \text{ meV}$, $J_3 = -20 \text{ meV}$, $K_1 = -28 \text{ meV}$, $K_2 = 48 \text{ meV}$, and $K_3 = -68 \text{ meV}$ are shown in Fig. 5(a) of the main text. It signals a Goldstone mode at the Brillouin zone center and exhibits anisotropy along the [H, 0] (red line) and [0, H] (blue line) directions. We have made no effort to fine-tune the model parameters, and the theoretical dispersion already provides a good description of the experimental data taking into account the size of the experimental resolution (about 80 meV). Because the dispersion of the transverse excitations does not depend on J_2 and K_2 and also due to the size of the experimental error bar, the model parameters cannot be fully determined by fitting the experimental data.

We further calculate the dynamical spin structure factor along several high symmetric directions of the Brillouin zone. The results along the [H, 0], [0, H], and [H, H] directions are presented in Fig. 5(b) and (c) of the main text. We have adopted a phenomenological damping parameter γ in the calculation, and take $\gamma/2 = 100$ meV, $\gamma/2 = 150$ meV, and $\gamma/2 = 125$ meV along the [H, 0], [0, H], and [H, H] directions, respectively (indicated as the error bars in Fig. 5(a)). By comparing the spectral weights along the [H, 0] and [0, H] directions we find an increasing spectral weight anisotropy with increasing momentum transfer. This anisotropic feature is inherent in the coherence factor of $S_D^{xx}(q, \omega)$ in Eq. (13).

In summary, we have found that both the spectral weight anisotropy and the dispersion of excitations agree well with experimental results. From this, we can infer that well-defined collective magnetic excitations exist up to about 200 meV in FeSe. The calculated magnetic excitations are associated with the $(\pi,0)$ AFQ order, which arise from a model of quasi-localized magnetic moments with frustrated exchange interactions. Our results further highlight the important role of strong electron correlations in FeSe and related iron-based superconductors.

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