Magnetic Resonant Mode in the Low-Energy Spin-Excitation Spectrum of Superconducting Rb₂Fe₄Se₅ Single Crystals

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We have studied the low-energy spin-excitation spectrum of the single-crystalline Rb₂Fe₄Se₅ superconductor ($T_c = 32$ K) by means of inelastic neutron scattering. In the superconducting state, we observe a magnetic resonant mode centered at an energy of $\hbar \omega_{res} = 14$ meV and at the (0.5 0.25 0.5) wave vector (unfolded Fe-sublattice notation), which differs from the ones characterizing magnetic resonant modes in other iron-based superconductors. Our finding suggests that the 245-iron selenides are unconventional superconductors with a sign-changing order parameter, in which bulk superconductivity coexists with the $\sqrt{5} \times \sqrt{5}$ magnetic superstructure. The estimated ratios of $\hbar \omega_{\rm res}/k_B T_c \approx 5.1 \pm 0.4$ and $\hbar \omega_{\rm res}/2\Delta \approx$ 0.7 ± 0.1 , where Δ is the superconducting gap, indicate moderate pairing strength in this compound, similar to that in optimally doped 1111 and 122 pnictides.

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Soon after the discovery of arsenic-free iron-selenide superconductors A_2 Fe₄Se₅ (A = K, Rb, Cs), also known as 245 compounds [1], their unprecedented physical properties came to light, such as the coexistence of high- T_c superconductivity with strong antiferromagnetism [2,3]. The pairing mechanism and the symmetry of the superconducting (SC) order parameter in this family of compounds still remain among the major open questions. In the majority of other Fe-based superconductors, it is widely accepted that the strong nesting between the holelike Fermi surface (FS) at the Brilliouin zone (BZ) center and electronlike FS at the BZ boundary leads to the sign-changing *s*-wave (s_+ -wave) pairing symmetry [4]. This scenario has been supported by different experimental methods, such as angle-resolved photoemission spectroscopy (ARPES) [5], quasiparticle interference [6], and inelastic neutron scattering (INS) [7,8]. On the other hand, recent theoretical calculations [9] and ARPES experiments [10,11] on the 245 system revealed the absence of holelike FSs at the BZ center in the electronic structure, implying that the nesting between the hole- and electronlike FS sheets is no longer present. Hence, several theoretical studies proposed alternative pairing instabilities, such as *d*-wave or another type of s_+ -wave symmetry with a sign change in the order parameter between bonding and antibonding states [12–14]. As a hallmark of sign-changing SC order parameter, several authors theoretically predicted a resonant mode in the magnetic excitation spectrum below the SC transition; yet, its precise position in momentum space still remains controversial [12,13].

A major complication in treating the 245 compounds theoretically arises from the presence of a crystallographic superstructure of Fe vacancies that has been consistently reported both from x-ray and neutron diffraction experiments [15]. This $\sqrt{5} \times \sqrt{5}$ superstructure is closely related to the static antiferromagnetic (AFM) order persisting up to the Néel temperature, $T_N \approx 540$ K [16]. Although most of the existing band structure calculations have so far neglected the superstructure, several others have pointed out that it may have a strong influence on the FS shape [17,18]. However, these pronounced FS reconstruction effects have not been experimentally confirmed so far [3,10,11]. Such an uncertainty in the FS geometry and its nesting properties makes it hard to predict the exact



FIG. 1 (color online). The dc magnetic susceptibility measurements on three representative single-crystalline Rb₂Fe₄Se₅ samples from the same batch. A sharp diamagnetic response is observed in the ZFC measurement right below 32 K, indicating $\sim 100\%$ exclusion of the external magnetic field.



FIG. 2 (color online). (a) The in-plane projection of twinned magnetic and nuclear Bragg peak positions arising from the $\sqrt{5} \times \sqrt{5}$ Fe-vacancy superstructure. The two sets of dots and the corresponding dashed lines denote magnetic superstructure Bragg reflections and the magnetic BZ boundaries for the right and left twin domains, respectively. Black dotted lines represent the Fe1 unfolded BZ boundary. The arrow shows the trajectory of elastic momentum scans. (b) Elastic scans along the K direction [arrow in (a) and equivalent scans] measured at 1.5 K. The almost identical neutron intensities of two symmetric magnetic Bragg peaks indicate the nearly equal population of the two twin domains in the sample. (c) INS intensity at 11.5 meV in the vicinity of the magnetic ordering wave vector $(1.3 \ 0.1 \ 0.5)_{\text{Fe}_1}$. The intense spin-wave excitation peak is consistent with recent time-of-flight INS measurements on an insulating $Rb_{2+\delta}Fe_4Se_5$ compound [26].

location of itinerant spin fluctuations in reciprocal space. Moreover, band structure calculations in the vacancyordered magnetic state result in insulating solutions for the stoichiometric 245 compound [17]. A possible way to reconcile these apparently contradictory observations is to assume a nanoscale phase separation of (i) insulating vacancy-ordered magnetic domains and (ii) metallic nonmagnetic phase domains with effective electron doping that could host superconductivity at low temperatures. Such a kind of electronic phase segregation, resembling the situation in hole-doped 122 pnictides [19], found recent support from ARPES [3], scanning nanofocus singlecrystal x-ray diffraction imaging [20], scanning-tunneling microscopy [21], and optical spectroscopy [22]. In this Letter, we provide experimental insight by using INS to directly probe the elementary magnetic excitations in superconducting Rb₂Fe₄Se₅ (RFS).

For the present study, we used a mosaic of RFS single crystals with a total mass of ~ 1 g, grown by the Bridgman method [23]. The nearly stoichiometric and homogeneous composition with Rb:Fe:Se = 0.796:1.596:2.000(1.99:3.99:5) has been determined by wavelength dispersive x-ray electron-probe microanalysis using a Camebax SX50 analyzer with an accuracy of 0.5% for Fe and up to 1% for Se. The SC properties of the sample were characterized by magnetometry, where $\sim 100\%$ flux exclusion was observed in the zero-field-cooled (ZFC) measurement for temperatures up to $T_c = 32$ K (Fig. 1). The INS experiment was performed at the thermal-neutron tripleaxis spectrometer IN8 (ILL, Grenoble), with the sample mosaic mounted in the (HH0)/(00L) or (H00)/(00L)scattering planes. The wave vectors $\mathbf{Q} = (0.5 \ 0.5 \ L)_{\text{Fe}_1}$ and $(0.5 \ 0 \ L)_{\text{Fe}_1}$ were directly accessible in our scattering planes, and the spectrometer further allowed us to tilt the sample in order to access **Q** vectors in a certain range out of the scattering planes. Here and henceforth, we are using unfolded reciprocal-space notation corresponding to the Fe sublattice, which we denote as Fe1 because of its simplicity and the natural correspondence to the symmetry of the observed signal [24]. We quote the wave vector $\mathbf{Q} =$ (HKL) in reciprocal-lattice units (r.l.u.), i.e., in units of the reciprocal-lattice vectors \mathbf{a}^* , \mathbf{b}^* , and \mathbf{c}^* of the Fe sublattice ($a^* = 2\pi/a$, etc.). Here, a = b = 2.76 Å is the room-temperature distance between the nearestneighbor Fe atoms, and c = 7.25 Å is the distance between Fe layers. All INS measurements were done in the fixed- k_f $(k_f = 2.662 \text{ Å}^{-1}) \text{ mode, using a double-focused PG}(002)$ monochromator and analyzer. A 5-cm-thick oriented PG filter was installed before the analyzer to eliminate higherorder neutron contamination, and no collimation was applied to maximize the intensity.

We start with the magnetic Bragg peak patterns arising from the $\sqrt{5} \times \sqrt{5}$ Fe-vacancy superstructure. Panel (a) in Fig. 2 is a sketch of magnetic and nuclear superstructure Bragg reflections, projected onto the two-dimensional $\mathbf{Q}_{\parallel} = (HK)$ plane. The black dots and the large dashed rectangle correspond to the center and boundaries of the unfolded Fe₁ BZ, respectively. The solid dots represent magnetic Bragg reflections from two twin domains, and the corresponding dashed lines, rotated clockwise and counterclockwise with respect to the Fe₁ BZ, are magnetic zone boundaries of the two twin domains. Forbidden magnetic Bragg peaks, which coincide with the nuclear Fe-vacancy superstructure reflections seen by x-ray diffraction [15], are shown by empty circles. Figure 2(b) shows elastic scans crossing two magnetic Bragg peaks at $(0.7 \pm 0.1 \ 0.5)_{\text{Fe}_1}$, as shown by the arrow in (a), and along equivalent reciprocalspace directions in higher BZs. Note that the two magnetic Bragg peaks at $K = \pm 0.1$ originate from different twin domains, so that their similar intensity indicates almost equal population of both twins in our sample. The magnetic Bragg peak intensity decreases more rapidly

when moving to a higher BZ along the out-of-plane direction than in-plane, indicating that the magnetic moment is oriented predominantly along the *L* direction in this system. This is consistent with the reported spin configuration in the magnetically ordered phase, in which spins are alternatively pointing up and down along the *c* axis [25]. Figure 2(c) shows inelastic magnetic intensity in the vicinity of the AFM ordering wave vector $\mathbf{Q} =$ $(1.3 \ 0.1 \ 0.5)_{\text{Fe}_1}$ at 11.5 meV, measured at a low temperature, T = 1.5 K. The intense spin-wave peak is consistent with recent time-of-flight INS measurements on an insulating $\text{Rb}_{2+\delta}\text{Fe}_4\text{Se}_5$ compound [26].

Now we turn to the INS measurements across T_c near a few candidate **Q** vectors, where the magnetic resonant mode could be expected. Figures 3(a) and 3(b) display raw energy-scan spectra recorded above and below T_c at **Q** = $(0.5 \ 0.3125 \ 0.5)_{\text{Fe}_1}$, where the resonance has been theoretically predicted [12], and at $(0.5 \ 0 \ 0.5)_{\text{Fe}_1}$, where it is usually found in other Fe-based superconductors [7,8]. In the absence of any resonant enhancement, the intensity is expected to be higher in the normal state due to the influence of the Bose factor at low energies. Already in the raw data, one can see that this is the case for all data points except a narrow energy region around 14 meV at **Q** = $(0.5 \ 0.3125 \ 0.5)_{\text{Fe}_1}$.

To emphasize this effect and to eliminate the energydependent background, we plot temperature differences of the same data sets in Figs. 3(c) and 3(d). Also shown are the difference spectra for $\mathbf{Q} = (0.5 \ 0.5 \ 0.5)_{\text{Fe}_1}$, $(0.5 \ 0.25 \ 0.5)_{\text{Fe}_1}$, and $(0.5 \ 0 \ 0)_{\text{Fe}_1}$. As seen in Fig. 3(c), a prominent peak (shaded region) is found at $\hbar\omega_{\rm res} \approx$ 14 meV for $\mathbf{Q} = (0.5 \ 0.3125 \ 0.5)_{\text{Fe}_1}$ $\mathbf{Q} =$ and $(0.5 \ 0.25 \ 0.5)_{\text{Fe}_1}$, which we attribute to the magnetic resonant mode. However, no such peak is observed at Q = $(0.5 \ 0.5 \ 0.5)_{\text{Fe}_1}$, in contrast to some alternative predictions based on the *d*-wave pairing symmetry [13]. Figure 3(d)also demonstrates the absence of any resonant mode at $\mathbf{Q} = (0.5 \ 0 \ 0.5)_{\text{Fe}_1}$ and $(0.5 \ 0 \ 0)_{\text{Fe}_1}$, where it is usually found in iron pnictides [7,8]. At these wave vectors, the data simply follow the solid line, which is the Bose-factor difference between 1.5 and 35 K.

To verify whether the observed redistribution of spectral weight at low temperatures is related to the SC transition, we have measured the temperature dependence of the resonance intensity at $\mathbf{Q} = (0.5 \ 0.3125 \ 0.5)_{\text{Fe}_1}$, which is shown in Fig. 3(e). Indeed, an order-parameter-like increase of intensity below T_c is found, which is accepted as the hallmark of the magnetic resonant mode.

To pin down the exact location of the resonance in \mathbf{Q} space, we have measured momentum scans along the BZ boundary at both temperatures. Their difference is presented in Fig. 3(f) and suggests a maximum at the commensurate nesting wave vector $\mathbf{Q}_{res} = (0.5 \ 0.25 \ 0.5)_{Fe_1}$ shown by the star symbols in Fig. 2(a), close to the



FIG. 3 (color online). (a),(b) Raw energy scans measured in the SC (1.5 K) and normal (35 K) states at $\mathbf{Q} =$ $(0.5 \ 0.3125 \ 0.5)_{Fe_1}$ and $(0.5 \ 0 \ 0.5)_{Fe_1}$, respectively. The inset in (a) shows the zoomed-in part of the resonant peak in the raw data. (c) Intensity difference between the SC state and the normal state at three **Q** vectors: $(0.5 \ 0.25 \ 0.5)_{\text{Fe}_1}$, $(0.5 \ 0.3125 \ 0.5)_{Fe_1}$, and $(0.5 \ 0.5 \ 0.5)_{Fe_1}$. While there is no positive intensity at (0.5 0.5 0.5), a clear resonance peak (shaded region) is observed around 14 meV both at $(0.5 \ 0.25 \ 0.5)_{Fe_1}$ and $(0.5 \ 0.3125 \ 0.5)_{Fe_1}$. (d) The same plot as in (c) but for $\mathbf{Q} =$ $(0.5\ 0\ 0.5)_{Fe_1}$ and $(0.5\ 0\ 0)_{Fe_1}$, where the magnetic resonant mode has been found in other Fe-based superconductors but is absent here. The base line in (c),(d) is the difference of the Bose factors. (e) Temperature dependence of the raw INS intensity at 14 meV and $\mathbf{Q} = (0.5 \ 0.3125 \ 0.5)_{\text{Fe}_1}$ that demonstrates an order-parameter-like behavior with an onset at T_c . (f) Intensity difference of momentum scans along the BZ boundary, measured below and above T_c , with a maximum at the commensurate wave vector $\mathbf{Q}_{res} = (0.5 \ 0.25 \ 0.5)_{Fe_1}$. The solid line is a Gaussian fit with a linear background. Different symbols represent identical momentum scans measured in different experiments, rescaled to the (002) nuclear Bragg peak intensity. The position of the resonant mode predicted by Maier et al. [12] is shown by the arrow.

predicted resonance position, $\mathbf{Q} = (0.5 \ 0.3125 \ 0.5)_{Fe_1}$ [12]. Yet, the disagreement is small compared to the \mathbf{Q} width of the peak, which explains the similar INS response at both \mathbf{Q} vectors, as seen from Fig. 3(c). Because the



FIG. 4 (color online). Normalized resonance energy, $\hbar\omega_{\rm res}/k_BT_c$, in Fe-based superconductors for $q_z = 0$ and $q_z = \pi$ [29]. This ratio for RFS is slightly higher than for 122 compounds but comparable to 11-, 111-, and 1111-type superconductors.

position of the nesting vector is strongly doping dependent and the calculations in Ref. [12] were done for the arbitrary doping of 0.1 electrons per Fe, a quantitative agreement with our results is not expected. The observed wave vector suggests an even higher level of the effective electron doping in the metallic phase of the sample, which is difficult to reconcile with the stoichiometric chemical composition unless we assume electronic phase segregation into electron-rich and electron-poor regions of the kind discussed in Refs. [20,21]. Such a high doping level of the metallic regions would also agree qualitatively with the ARPES results [3,10,11].

By comparing the normalized resonance intensity in RFS with that in the nearly optimally doped Ba(Fe_{1-x}Ni_x)₂As₂ (BFNA), measured in a similar experimental configuration at the same spectrometer [24], we find that the intensity at the resonance energy in RFS is approximately a factor of 3 smaller than in BFNA. Because we expect four nonequivalent resonant peaks in the BZ of RFS from symmetry considerations, as opposed to only two such peaks [$\mathbf{Q}_{res} = (0.5 \ 0)_{Fe_1}$ and $(0 \ 0.5)_{Fe_1}$] in the 122 system, the total resonant spectral weight in both compounds turns out to be comparable.

It has been shown that the resonance energy scales linearly with T_c in iron-based superconductors, with a ratio of $\hbar\omega_{res}/k_BT_c$ that slightly varies between different families [8,24,27] but is generally lower than the respective ratio of ~5.3 measured in high- T_c cuprates [28]. In Fig. 4, we compare this ratio in all Fe-based superconductors, in which the resonant mode has been found (see Ref. [29] and references therein). The value for RFS, $\hbar\omega_{res}/k_BT_c \approx$ 5.1 ± 0.4, lies slightly above the nearly universal ratio of 4.3 estimated for 122 compounds (solid line) [24] but is close to that in FeTe_{1-x}Se_x, LiFeAs, La-1111, and cuprate superconductors.

Another important dimensionless parameter that allows an assessment of the pairing strength in unconventional superconductors is the $\hbar\omega_{\rm res}/2\Delta$ ratio, where Δ is the superconducting gap. We have recently shown that in Febased superconductors this ratio varies between the universal value of ~0.64, typical for strong-coupling superconductors with high T_c [30], and unity for low- T_c compounds [29]. In the 245 systems, the SC gap has been measured by ARPES and NMR [11,31], producing the average $2\Delta/k_BT_c$ ratio of ~7.2 ± 0.4 [29]. It corresponds to the $\hbar\omega_{\rm res}/2\Delta$ ratio of ~0.7 ± 0.1 in RFS, slightly above the strong-coupling limit, supporting the general trend found in all Fe-based superconductors [29].

To conclude, we have observed the magnetic resonant mode coexisting with the AFM order in RFS below the SC transition. Our finding suggests unconventional pairing with a sign-changing order parameter in the 245 systems, qualitatively consistent with theoretical predictions made under the assumption of finite electron doping in the metallic phase volume. Although this result points toward an electronic phase separation scenario [20], the resonant spectral weight is comparable to that in optimally doped 122 compounds. It evidences bulk superconductivity in our sample, consistent with the $\sim 100\%$ flux exclusion observed in the ZFC magnetization measurement, and implies an unusual coexistence or microscopic segregation of strong antiferromagnetism with superconductivity, which should be addressed by future theoretical work. The estimated ratios of $\hbar\omega_{\rm res}/k_BT_c \approx 5.1 \pm 0.4$ and $\hbar\omega_{\rm res}/2\Delta \approx$ 0.7 ± 0.1 in this compound indicate moderately strong pairing, similar to other Fe-based superconductors.

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