

Temperature dependence of lower critical field $H_{c1}(T)$ shows nodeless superconductivity in FeSeM. Abdel-Hafiez,^{1,*} J. Ge,² A. N. Vasiliev,^{3,4} D. A. Chareev,⁵ J. Van de Vondel,² V. V. Moshchalkov,² and A. V. Silhanek¹¹*Département de Physique, Université de Liège, B-4000 Sart Tilman, Belgium*²*INPAC—Institute for Nanoscale Physics and Chemistry, Nanoscale Superconductivity and Magnetism Group, K.U. Leuven, Celestijnenlaan 200D, B-3001 Leuven, Belgium*³*Low Temperature Physics and Superconductivity Department, M.V. Lomonosov Moscow State University, 119991 Moscow, Russia*⁴*Theoretical Physics and Applied Mathematics Department, Ural Federal University, 620002 Ekaterinburg, Russia*⁵*Institute of Experimental Mineralogy, Russian Academy of Sciences, 142432 Chernogolovka, Moscow District, Russia*

(Received 30 August 2013; revised manuscript received 28 October 2013; published 18 November 2013)

We investigate the temperature dependence of the lower critical field $H_{c1}(T)$ of a high-quality FeSe single crystal under static magnetic fields H parallel to the c axis. The temperature dependence of the first vortex penetration field has been experimentally obtained by two independent methods and the corresponding $H_{c1}(T)$ was deduced by taking into account demagnetization factors. A pronounced change in the $H_{c1}(T)$ curvature is observed, which is attributed to anisotropic s -wave or multiband superconductivity. The London penetration depth $\lambda_{ab}(T)$ calculated from the lower critical field does not follow an exponential behavior at low temperatures, as it would be expected for a fully gapped clean s -wave superconductor. Using either a two-band model with s -wave-like gaps of magnitudes $\Delta_1 = 0.41 \pm 0.1$ meV and $\Delta_2 = 3.33 \pm 0.25$ meV or a single anisotropic s -wave order parameter, the temperature dependence of the lower critical field $H_{c1}(T)$ can be well described. These observations clearly show that the superconducting energy gap in FeSe is nodeless.

DOI: [10.1103/PhysRevB.88.174512](https://doi.org/10.1103/PhysRevB.88.174512)

PACS number(s): 74.20.Rp, 74.25.Dw, 74.25.Jb, 74.70.Dd

I. INTRODUCTION

Superconductivity in Fe-based superconductors (FeSCs) has been studied intensively due to the relatively large transition temperatures T_c up to 55 K,^{1–4} a high upper critical field, and a layered structure similar to the cuprates. Superconductivity emerges when the magnetic order is suppressed by charge doping or by applying an external pressure, and takes place within FeAs, FeP, or FeSe crystallographic planes. These planes are separated by layers of other elements serving as charge reservoirs. Among the very few members of this FeSCs class becoming superconducting at ambient pressure and without doping, we find the iron selenide, FeSe. This compound has the simplest crystal structure and stoichiometry while keeping a moderate superconducting critical temperature of about 8 K for polycrystalline samples.⁵ Furthermore, high-quality single crystals with rather large dimensions can be grown,^{6–8} which are necessary for an accurate determination of bulk physical properties. Interestingly, if the tetragonal FeSe system is submitted to a pressure of 8.9 GPa, a huge enhancement of the T_c up to 36.7 K is obtained.^{5,9} This system also stands out due to the absence of nesting between the hole and electron pockets of the Fermi surface.⁵ In addition, density functional calculations of the electronic structure indicate that the electron-phonon coupling cannot explain superconductivity at such a high transition temperature.¹⁰ Therefore, FeSe falls into the category of unconventional superconductivity and appears as an ideal candidate to study the fundamental properties of superconductivity in clean iron-based superconductors.

One of the crucial issues to elucidate the mechanism leading to high-temperature superconductivity is the nature of pairing, e.g., the symmetry and structure of the superconducting order parameter. Up to now, there have been several investigations on the pairing symmetry of FeSe superconductors. Thermal conductivity measurements¹¹ show the absence of nodes in

the superconducting gap. Furthermore, recent upper critical field studies of β -FeSe crystals have revealed that two-band effects dominate $H_{c2}(T)$, with the possible influence of a spin paramagnetic effect.¹² The presence of both an isotropic s -wave and extended s -wave order parameters coexisting in a superconducting single-crystal FeSe has also been proposed based on specific-heat measurements.¹³ Very recently, multiple Andreev reflection spectroscopy pointed to the existence of two-gap superconductivity.⁷ In addition, muon-spin rotation studies of the penetration depth $\lambda_{ab}^{-2}(T)$ in FeSe_{0.85} were consistent with either two-gap ($s + s$) or anisotropic s -wave order parameter symmetries, thus implying that the superconducting energy gap contains no nodes.¹⁴ In contrast to that, scanning tunneling spectroscopy experiments in the stoichiometric FeSe provided clear evidence for nodal superconductivity.¹⁵ The observed gap function was attributed to an extended s -wave pairing structure with a mixture of $s_{x^2+y^2}$ and $s_{x^2-y^2}$ pairing symmetries.¹⁶ Clearly, to date, there is no general consensus on the origin of the superconducting pairing mechanism in FeSe compounds and further measurements to elucidate this issue are necessary.

The lower critical field $H_{c1}(T)$, i.e., the thermodynamic field at which the presence of vortices into the sample becomes energetically favorable, and the magnetic penetration depth $\lambda(T)$ are very useful parameters, providing key information regarding bulk thermodynamic properties. Indeed, the gap properties of different families of FeSCs have been investigated by tracking the $H_{c1}(T)$ and the magnetic penetration depth.^{17–23} The gap properties of these compounds display single to double gaps and even the presence of nodes. This variety of gap properties appears to be related to the nature and the level of doping. It should be mentioned that this quest for new multiband superconductivity is a timely subject due to the possible emergence of nonmonotonic vortex-vortex interactions, a fingerprint of so-called type 1.5 superconductivity.^{24–26}

The main motivation of the present paper is to tackle a longstanding question concerning whether the superconducting properties of these materials can or cannot be accounted for a nodal order parameter. To that end, we reliably determine the temperature dependence of the H_{c1} from magnetization measurements. We then compare the most popular approach of determining the first vortex penetration as the point of deviation from a linear $M(H)$ response, to the value obtained from the onset of the trapped magnetic moment (M_t). Although this latter approach to determine $H_{c1}(T)$ has been performed in detail in high- T_c superconductors, i.e., $\text{YBa}_2\text{Cu}_3\text{O}_{0.69}$ as reported in Ref. 27, its application to Fe-based superconductors still needs to be presented. In particular, our results show that the method of the trapped magnetization onset is more sensitive than the method determined from the point of deviation from a linear $M(H)$ response. In addition, a kink around 7 K is obtained on the $H_{c1}(T)$ curve, which can be accounted for the anisotropic s -wave or multiband superconductivity in this system. Our analysis further shows that the superconducting gaps determined through fittings to the in-plane London penetration depth cannot be described with the single-band weak-coupling BCS scheme; rather, it implies the presence of either two s -wave-like gaps with different magnitudes and contributions or a single anisotropic s -wave gap. Our London penetration depth results are contrasted to values obtained through the more sophisticated technique of muon-spin rotation.¹⁴

II. EXPERIMENT

We investigated a selected platelike FeSe single crystal grown in an evacuated quartz ampoule using the AlCl_3/KCl flux technique with a constant temperature gradient of $5^\circ\text{C}/\text{cm}$ along the ampoule length (the temperature of the hot end was kept at 427°C , and the temperature of the cold end was about 380°C). The phase purity of the resulting crystal was checked with x-ray diffraction.⁷ The sample has lateral dimensions $a \times b \times c = 1.05 \pm 0.08 \times 1.25 \pm 0.1 \times 0.02 \pm 0.01 \text{ mm}^3$ with a mass of 1.2 mg. Magnetization measurements were performed using a superconducting quantum interference device magnetometer (MPMS-XL5) from Quantum Design. The good quality of the crystals is confirmed from a sharp specific-heat jump (9.45 mJ/mol K^2 with zero residual specific heat γ_r),¹³ indicative of a complete superconducting volume. The temperature dependence of resistance $R(T)$ demonstrates a metallic behavior with $T_c = 9.4 \text{ K}$.⁷

III. RESULTS AND DISCUSSIONS

A. Irreversible magnetization

Figure 1(a) presents the field dependence of the isothermal magnetization M at certain selected temperatures up to 40 kOe for $H \parallel c$ (main panel) and $H \parallel ab$ [see the inset of Fig. 1(a)]. For $H \parallel c$, the magnetic irreversibility presents a second peak, whereas no second peak is observed for $H \parallel ab$. This significant anisotropic behavior in the appearance of the second peak has been reported previously in other Fe-based superconductors, e.g., $\text{Ba}(\text{Fe}_{0.93}\text{Co}_{0.07})_2\text{As}_2$ (Ref. 28) and LiFeAs (Ref. 29) single crystals, and is typically associated with the nature of pinning.³⁰ As reported in the 122 and 111

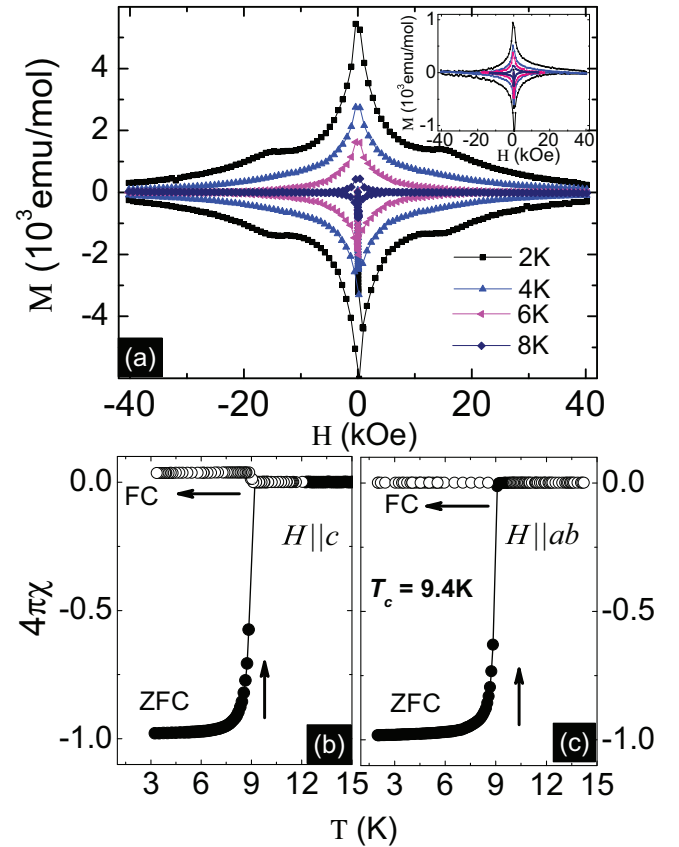


FIG. 1. (Color online) (a) Magnetic field dependence of the isothermal magnetization M vs H loops measured at different temperatures ranging from 2 to 8 K up to 40 kOe with the field parallel to both the c axis and ab plane as an inset. (b) and (c) present the temperature dependence of the magnetic susceptibility χ after demagnetization correction in an external field of 1 Oe applied along c and ab , respectively. The χ has been deduced from the dc magnetization measured with $H \parallel c$ and $H \parallel ab$ following ZFC and FC protocols for FeSe single crystals.

systems, the second peak is regarded as the crossover from plastic to elastic pinning.^{28,29}

Figures 1(b) and 1(c) show the temperature dependence of the magnetic susceptibility of the FeSe single crystal measured by following the zero-field-cooled (ZFC) and field-cooled (FC) procedures in an external field of 1 Oe applied along the c and ab axis, respectively. The ZFC data for both orientations show a sharp diamagnetic signal, thus confirming bulk superconductivity in our investigated system. The magnetic susceptibility exhibits a superconducting transition with an onset transition temperature T_c^{χ} of 9.4 K for both orientations. The clear irreversibility between FC and ZFC measurements is a consequence of a strong vortex trapping mechanism, either by surface barriers or bulk pinning. Notice that the magnetic moment in field-cooled conditions for the $H \parallel c$ axis becomes positive for $T < T_c$ [see Fig. 1(b)]. A similar behavior has been observed in conventional as well as Fe-based superconductors.³¹

The fact that the hysteresis loops for both orientations are symmetric around $M = 0$ points to the relatively weak surface barriers and is indicative of strong bulk pinning.

This consideration holds for all studied temperatures, even close to T_c , and guarantees that vortex penetration occurs at a field close to H_{c1} . In contrast to that, if surface barriers were predominant, the first vortex entrance could take place at much higher field ($\sim H_c$). This is a very important point in order to obtain reliable estimations of the thermodynamic lower critical field, as we will discuss below. It is worth noting that the superconducting $M(H)$ exhibits a very weak magnetic background. This indicates that the sample contains negligible amounts of magnetic impurities.

From the magnetization hysteresis loops $M(H)$, we calculated the critical current density J_c by using the critical state model with the assumption of field-independent J_c .^{32,33}

$$J_c = \frac{20\Delta M}{\left[a\left(1 - \frac{a}{3b}\right)\right]}, \quad (1)$$

where $\Delta M = M_{dn} - M_{up}$, M_{dn} and M_{up} are the magnetizations measured with decreasing and increasing applied field, respectively, and a (cm) and b (cm) are sample widths ($a < b$). The unit of ΔM is in electromagnetic unit per cubic centimeter and the calculated J_c is in ampere per square centimeter. We obtain $J_c(2\text{ K}) \sim 1.34 \times 10^4$ A/cm² for $H \parallel c$ and $J_c(2\text{ K}) \sim 1.8 \times 10^4$ A/cm² for $H \parallel ab$. These values are lower than those reported in Ba-122, 1111, 11, and the 111 systems^{29,34–36} and higher than those observed in $\text{K}_{0.64}\text{Fe}_{1.44}\text{Se}_2$.³⁷

B. Experimental determination of the lower critical field

Determining the lower critical field from magnetization measurements has never been an easy task, particularly since H_{c1} is an equilibrium thermodynamic field, whereas the magnetization curve is highly irreversible as a consequence of metastable vortex states far from equilibrium. The most popular method to estimate H_{c1} (here tagged as method A) consists of detecting the transition from a Meissner-like linear $M(H)$ regime to a nonlinear $M(H)$ response, once the vortices penetrate into the sample and build up a critical state. This transition is not abrupt, therefore bearing a substantial error bar.

These sorts of measurements are obtained by tracking the virgin $M(H)$ curve at low fields at several temperatures, as shown in the upper panel of Fig. 2 for $H \parallel c$. We have adopted a rigorous procedure (i.e., with a user-independent outcome) to determine the transition from linear to nonlinear $M(H)$, which consists of calculating the regression coefficient R of a linear fit to the data points collected between 0 and H , as a function of H . Then, H_{c1} is taken as the point where the function $R(H)$ departs from 1. This procedure is illustrated for a particular temperature $T = 3\text{ K}$ in the inset of the upper panel of Fig. 2.

An alternative and seemingly more reliable way to determine the lower critical field (here tagged as method B) can be obtained by measuring the onset of the trapped moment M_t as described in Refs. 27 and 38. In contrast to method A, where a heavy data postprocessing is needed, now a careful measurement protocol needs to be followed with little data analysis. Indeed, the trapped flux moment M_t is obtained by (i) warming the sample up to temperatures above T_c ($T = 20\text{ K}$), then (ii) cooling the sample at zero field down to the chosen temperature; subsequently (iii) the external magnetic field is increased to a certain maximum value H_m and

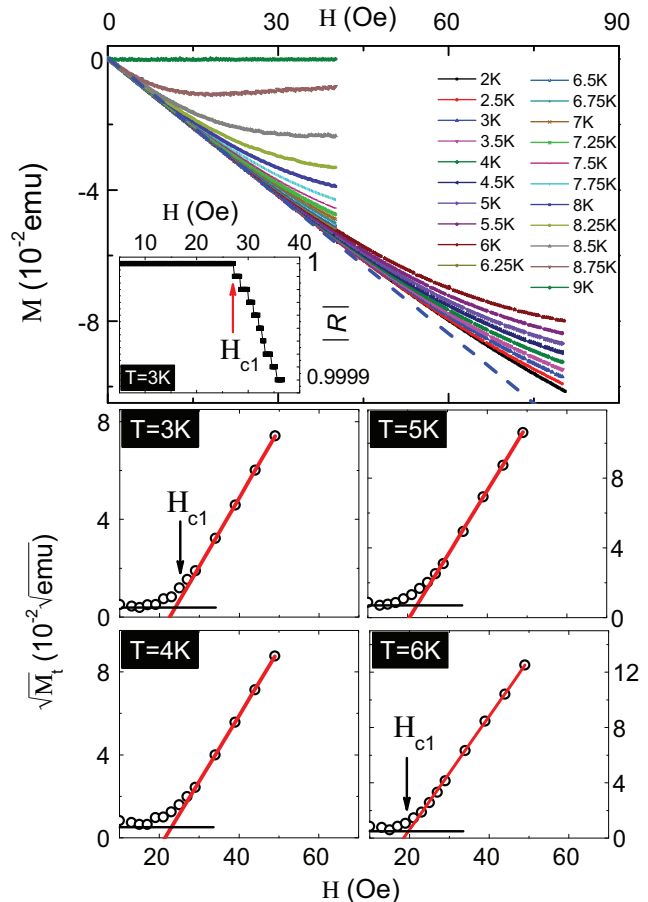


FIG. 2. (Color online) The upper panel shows the superconducting initial part of the measured magnetization curves of β -FeSe single crystals at various temperatures for $H \parallel c$. The dashed line depicts the Meissner line (linear fits between 0 and 15 Oe). The inset depicts an example used to determine the H_{c1} value using the regression factor R at $T = 3\text{ K}$ (see text). The lower panels present the field dependence of the typical plot of $\sqrt{M_t}$ vs H at various temperatures. The solid lines are a linear fit to the high-field data of $\sqrt{M_t}$ vs H . H_{c1} values are determined by extrapolating the linear fit to $\sqrt{M_t} = 0$.

(iv) measurements are made of the remanent magnetization M_t after the applied field has been switched off. The field H_m at which M_t deviates from zero determines the H_{c1} value at the desired temperature. It is important to notice that this method furnishes us with a rather independent determination of H_{c1} , weakly linked to the first procedure (method A) described above.

When taking into account the reversible magnetization, the trapped magnetic moment is $M_t \propto (H - H_{c1})^2$.²⁷ Then the extrapolation $\sqrt{M_t} \rightarrow 0$ determines the exact value of the H_{c1} . The lower panel of Fig. 2 presents the typical plot of $\sqrt{M_t}$ versus the applied field H for our FeSe single crystal. The solid line is a linear fit to the high-field data of $\sqrt{M_t}$ vs H . H_{c1} is determined by extrapolating the linear fit to $\sqrt{M_t} = 0$.

Once the values of H_{c1} have been experimentally determined, we need to correct them to account for the demagnetization effects. Indeed, the deflection of field lines around the sample leads to a more pronounced Meissner slope given by $M/H_a = -1/(1 - N)$, where N is the demagnetization factor.

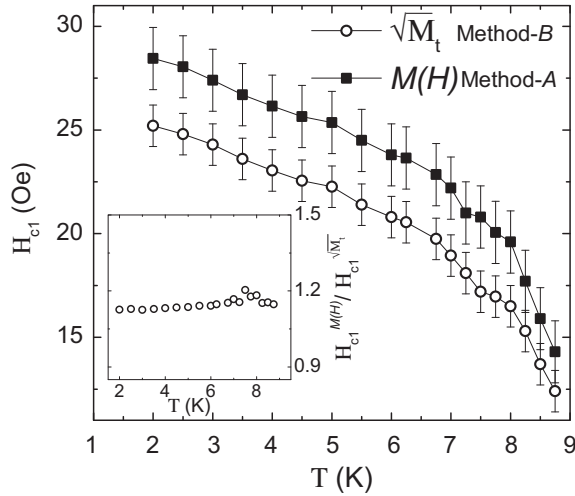


FIG. 3. The lower critical fields of the FeSe single crystal for the field applied parallel to the c axis. H_{c1} has been estimated by two different methods—from the extrapolation of $\sqrt{M_t} \rightarrow 0$ (see the lower panels of Fig. 2) and from the regression factor (see the inset of the upper panel in Fig. 2). The bars show the uncertainty estimated by the deviating point of the regression fits and the linear fit of $\sqrt{M_t}$. The inset shows the ratio of $H_{c1}(T)$ values obtained by both methods.

Taking into account these effects, the absolute value of H_{c1} can be estimated by using the relation proposed by Brandt:³⁹

$$q_{\text{disk}} = \frac{4}{3\pi} + \frac{2}{3\pi} \tanh \left[1.27 \frac{c}{a} \ln \left(1 + \frac{a}{c} \right) \right], \quad (2)$$

where $q_{\text{disk}} \equiv (|M/H_a| - 1)(c/a)$, and a and c are the dimensions perpendicular to the field and thickness of our investigated sample, respectively. For our sample we find $N \approx 0.9623$. In addition, an alternative way to determine the demagnetization factor is from a rectangular prism approximation based on the dimensions of the crystal, giving us $N \approx 0.9688$.⁴⁰

The corrected values of H_{c1} obtained by following the two methods described above are illustrated in Fig. 3 for $H \parallel c$. Even though both procedures yield different values of H_{c1} , the ratio of both methods is just a constant factor with no change in the shape or the dependence (see the inset of Fig. 3). This fact shows that both methods can provide a qualitative estimation of the temperature dependence of H_{c1} . However, method B shows lower H_{c1} values than method A, which means that the former method is much more sensitive than the latter method, although to obtain a more quantitative result we should compare these methods to high resolution imaging techniques such as magneto-optical imaging, Bitter decoration, or scanning Hall probe microscopy.

C. Theoretical fitting of the lower critical field

Irrespective of the method used to obtain H_{c1} , a pronounced change of the curvature is observed around 7 K. This may be attributed to the anisotropic s -wave or multiband nature of superconductivity in our system. This behavior is reminiscent of that reported for the two-band superconductors MgB_2 (Ref. 41) and Fe-based superconductors,^{17,19} in which similar $H_{c1}(T)$ curves were well fitted by a two-gap weak-coupling BCS model.

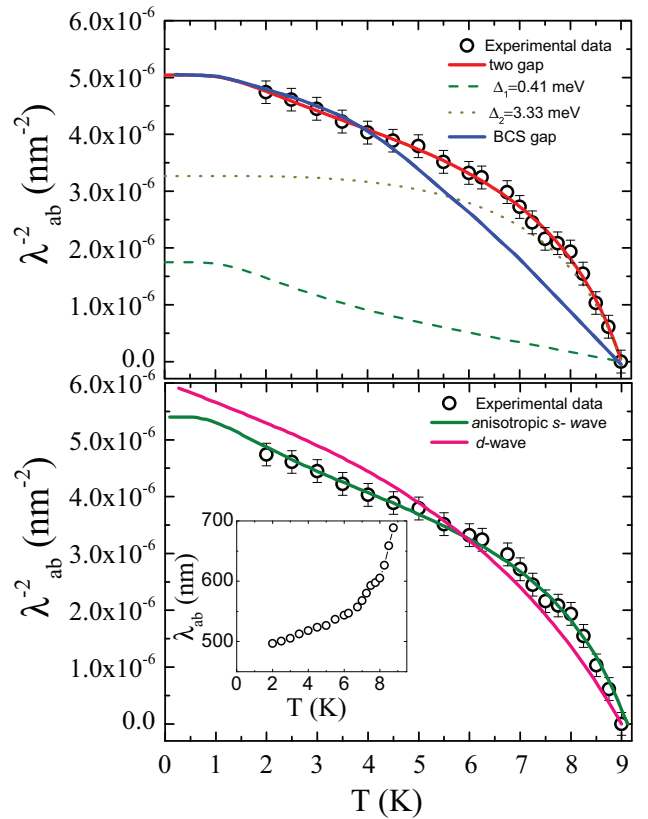


FIG. 4. (Color online) Upper panel: The temperature dependence of the London penetration depth $\lambda_{ab}^{-2}(T)$ for FeSe. The solid red line is the fitting curve using the two-gap model. The dashed and dashed dotted lines show the contributions in the two-band model of the big gap and small gap, respectively (see text). The blue line corresponds to a single-gap BCS curve. The saturation yields $\lambda_{ab}(0) = 445(15)$ nm. Lower panel: The fitting curves (solid lines) were obtained within the following anisotropic s -wave and d -wave models of the gap symmetries (see text). The inset presents the temperature dependence of the magnetic penetration depths $\lambda_{ab}(T)$ of FeSe.

Alternatively, in order to shed light on the pairing symmetry in our system, we determined the temperature dependence of the magnetic London penetration depth (λ_{ab}) applied along the c axis by using the following formula (taking the demagnetization effect into account): $\mu_0 H_{c1}^{\parallel c} = (\phi_0/4\pi\lambda_{ab}^2) \ln \kappa_c$, where ϕ_0 is the magnetic-flux quantum and $\phi_0 = h/e^* = 2.07 \times 10^{-7}$ Oe cm², and $\kappa_c = \lambda_{ab}/\xi_{ab} = 72.3$ (Ref. 42) is the Ginzburg-Landau parameter. The results of these calculations are shown in the inset of Fig. 4. Since we believe that method B is more accurate in determining H_{c1} , we have calculated λ_{ab} only for this method. The penetration depth of our FeSe shows similar behavior to the penetration depths as reported in LiFeAs .¹⁹ At low temperatures, $\lambda_{ab}(T)$ does not show the typical exponential behavior expected for a fully gapped clean s -wave superconductor.

Furthermore, if we compare our data to the single-gap BCS theory (i.e., a weak-coupling approach), we find that a single BCS gap cannot be reconciled with our experimental data (see the blue dashed line in Fig. 4). Indeed, the single BCS gap leads to a rather different trend and shows a systematic deviation from the data in the whole temperature range below T_c . In addition, it largely misses the kink around 7 K.

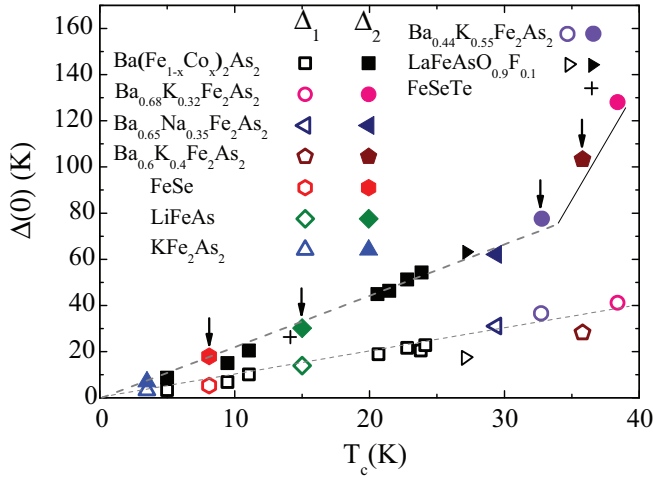


FIG. 5. (Color online) The superconducting-gap evolution of FeSe (this study), together with $\text{Ba}_{0.65}\text{Na}_{0.35}\text{Fe}_2\text{As}_2$ (Ref. 31), $\text{Ba}(\text{Fe}_{1-x}\text{Co}_x)_2\text{As}_2$ (Ref. 48) for $0.05 \leq x \leq 0.146$, $\text{Ba}_{0.68}\text{K}_{0.32}\text{Fe}_2\text{As}_2$ (Ref. 49), $\text{Ba}_{0.6}\text{K}_{0.4}\text{Fe}_2\text{As}_2$ (Ref. 17), KFe_2As_2 (Ref. 50), LiFeAs (Ref. 19), $\text{Ba}_{0.45}\text{K}_{0.55}\text{Fe}_2\text{As}_2$ (Ref. 51), $\text{LaFeAsO}_{0.9}\text{F}_{0.1}$ (Ref. 52), and FeSeTe (Ref. 53). Lines are guides to the eye. The arrows show that both gap values are determined using the lower critical field $H_{c1}(T)$ studies.

Knowing that (i) a single isotropic gap scenario cannot describe our data and (ii) the presence of two superconducting gaps is observed in a variety of different pnictides (see Fig. 5), we applied a phenomenological two-gap model reported by Carrington and Manzano.⁴³ Within this model, the temperature dependence of each energy gap can be approximated as⁴³ $\Delta_i(T) = \Delta_i(0) \tanh\{1.82[1.018(\frac{T_{ci}}{T} - 1)]^{0.51}\}$. In the Carrington and Manzano approach the one-band expression is straightforwardly generalized to the two-band case. The obtained experimental temperature dependence of $\lambda^{-2}(T)$ is fitted using a model of two BCS superconducting bands within the clean limit approach for a London superconductor with different gaps.⁴⁴ According to Ref. 45, for each band, $\lambda_i^{-2}(T)$ is given by

$$\lambda_i^{-2}(T) = \frac{\Delta_i(T) \tanh\left(\frac{\Delta_i(T)}{2k_B T}\right)}{\lambda_i^2(0)\Delta_i(0)}, \quad (3)$$

where $\lambda_i(0)$ is the residual penetration depth for each band, and k_B is the Boltzmann constant. Considering different contributions of each band to the whole $\lambda^{-2}(T)$, the following expression was used: $\lambda^{-2}(T) = r\lambda_1^{-2}(T) + (1-r)\lambda_2^{-2}(T)$, with r being the weighting factor that indicates the contribution of the small gap.

TABLE I. The superconducting transition temperature T_c (K), the d -wave approach ($\Delta_0 = \text{meV}$), the anisotropic s -wave approach ($\Delta_0 = \text{meV}$), two s -wave gaps (meV), and the London penetration depth $\lambda_{ab}(0)$ extracted from the the temperature dependence of the London penetration depth for FeSe and $\text{FeSe}_{0.85}$ (Ref. 14).

System	T_c	d wave	Anisotropic s wave	Two s waves	λ (nm)	Ref.
FeSe	9.4	2.045(5)	1.663(5)	0.41 and 3.33	445(15)	This work
$\text{FeSe}_{0.85}$	8.26	1.8(5)	2.2(3)	0.38 and 1.6	405(7)	14

To calculate the theoretical curves, the parameters $\Delta_1(0)$, $\Delta_2(0)$, and their respective ratios are adjusted. The results of the calculation using the above equations are shown in Fig. 4. The best description of the experimental data is obtained using values of $\Delta_1(0) = 0.41 \pm 0.1$ meV, $\Delta_2(0) = 3.33 \pm 0.25$ meV, and $r = 0.2$. The calculated penetration depth data are represented by the solid red line in the upper panel of Fig. 4.

The temperature dependence of the magnetic penetration depth of the anisotropic s -wave and d -wave gap calculations were performed using the following functional form:^{14,46}

$$\frac{\lambda_{ab}^{-2}(T)}{\lambda_{ab}^{-2}(0)} = 1 + \frac{1}{\pi} \int_0^{2\pi} \int_{\Delta(T,\varphi)}^{\infty} \left(\frac{\partial f}{\partial E} \right) \frac{E dE d\varphi}{\sqrt{E^2 - \Delta(T,\varphi)^2}}, \quad (4)$$

$$f = \frac{1}{[1 + \exp(E/k_B T)]}, \quad (5)$$

where f is the Fermi function, φ is the angle along the Fermi surface, and $\Delta(T,\varphi) = \Delta_0 \delta(T/T_c) g(\varphi)$ (Δ_0 is the maximum gap value at $T = 0$). The function $g(\varphi)$ is given by $g^d(\varphi) = |\cos(2\varphi)|$ for the d -wave gap, while $g^s(\varphi) = (1 + a \cos 4\varphi)/(1 + a)$ for the anisotropic s -wave gap.⁴⁷ The results of the analysis are presented in the lower panel of Fig. 4 by solid lines. The best description of the experimental data for the anisotropic s wave is obtained using values of $\Delta_0 = 1.663(5)$ meV, $a = 4.772$, and $\lambda_{ab}(0) = 430(15)$ nm. For the d -wave case, we get $\Delta_0 = 2.045(5)$ meV. It is obvious that the d -wave case cannot describe the penetration depth data. On the other hand, the experimental data are well described for both anisotropic s - and two-gap s -wave models.

It is interesting to compare the extracted values with those obtained previously on the off-stoichiometry compound $\text{FeSe}_{0.85}$ (see Table I). It is worth mentioning here that the two-gap model describes the in-plane penetration depth data on $\text{FeSe}_{0.85}$ with gap values of 1.60 and 0.38 meV, substantially different from the values we report here for FeSe.¹⁴ This might not be surprising since it has been well established that changing the Se content not only leads to a different T_c but, as shown by McQueen *et al.*,⁵⁴ slight changes from the ideal 1:1 ratio in FeSe lead to severe changes in the superconducting properties. For instance, the low-field magnetization data of various $\text{FeSe}_{1\pm\delta}$ samples showed that the strongest superconducting signal occurs for the most stoichiometric sample, whereas it has been shown that for the $\text{FeSe}_{0.82}$ case there is no superconducting signal.⁵⁵

The extracted gap values for the two-gap s -wave model are also different from $\Delta_{1,2}(0) = 2.5$ and 5.1 meV, reported for $\text{FeTe}_{0.55}\text{Se}_{0.45}$,⁵⁶ but comparable to the two-band s -wave fit for the multiple Andreev reflection spectroscopy⁵⁷ [$\Delta_{1,2}(0) = 0.8 \pm 0.2$ and 2.75 ± 0.3 meV]. Such a multigap nature seems to be a common scenario for Fe-based superconductors. It

should be noted that both gap values are not far from those reported for LiFeAs single crystals,¹⁹ but are much smaller than those reported in Ba_{0.6}K_{0.4}Fe₂As₂ and Ba_{0.45}K_{0.55}Fe₂As₂.^{17,51} The lower gap in LiFeAs, Ba_{0.6}K_{0.4}FeAs, and in our system is smaller but significantly affects the zero-temperature penetration depth. It should be pointed out that such a small gap is also in line with specific-heat data on a similar FeSe crystal.¹³ The contribution to the in-plane penetration depth data from each band is also shown in Fig. 4 by dashed and dotted lines, respectively. In fully gapped superconductors, the penetration depth data should show a flat behavior at low temperatures. However, using the two-band model, we can get an expected saturating behavior below 0.4 K, indicating a full-gap superconducting state. This saturation yields $\lambda_{ab}(0) = 445(15)$ nm, which is somewhat smaller than [560(20) nm] in Fe(Te, Se).⁵⁸ Our estimated $\lambda_{ab}(0)$ value is indeed comparable to the $\lambda_{ab}(0)$ value [405(7) nm] derived from muon-spin rotation studies.¹⁴ Our results provide another strong evidence that FeSe is not a simple single gap.

Finally, for the sake of comparison, the gap amplitudes as a function of T_c of FeSe single crystals are shown in Fig. 5, together with Ba(Fe_{1-x}Co_x)₂As₂ (Ref. 48) for $0.05 \leq x \leq 0.146$, Ba_{0.68}K_{0.32}Fe₂As₂,⁴⁹ Ba_{0.65}Na_{0.35}Fe₂As₂,³¹ KFe₂As₂,⁵⁰ LiFeAs,¹⁹ Ba_{0.45}K_{0.55}Fe₂As₂,⁵¹ LaFeAsO_{0.9}F_{0.1},⁵² and FeSeTe.⁵³ As it can be seen, the gap values differ for different compounds within the 122 family and also for 11, 111, and 1111 compounds. Furthermore, Ponomarev *et al.*⁵⁷ have proven that the small and larger gaps increase linearly with T_c . It is also clear that the larger gap increases stronger than linear with T_c for $T_c \geq 30$ K. In addition, the values of the underdoped, optimally, and overdoped K-doped data fit onto the same curves of $\Delta_1(0)$, $\Delta_2(0)$. On the other hand, for the large gap values, a tendency for strong-coupling

effects, e.g., the compounds with the highest T_c , as in K- and Na-doped superconductors, is given. Very recently the evolution of the electronic structure of the single-layer FeSe film during the annealing process illustrates that the superconductivity is in the strong-coupling regime. In addition, both the superconducting gap and the transition temperature increase with the annealing process.⁵⁹ We do not yet have a good understanding for such large gap behavior, especially above 30 K. However, it is worth mentioning that the contribution of both gap values fits well with the other hole- and electron-doped 122 systems as well as with the 111 and 1111 compounds.

D. Summary

In conclusion, we have determined the temperature dependence of the lower critical field $H_{c1}(T)$ of FeSe by the onset of either the trapped moment or nonlinear $M(H)$ response. Assuming either two s -wave-like gaps with magnitudes $\Delta_1 = 0.41 \pm 0.1$ meV and $\Delta_2 = 3.33 \pm 0.25$ meV, or an anisotropic s -wave using values of $\Delta_0 = 1.663$ meV, we account for the temperature dependence of the lower critical field $H_{c1}(T)$. These observations clearly show that there are no nodes in the superconducting energy gap of FeSe. The London penetration depth $\lambda_{ab}(T)$ is calculated from the lower critical field and yields $\lambda_{ab}(0) = 445(15)$ nm.

ACKNOWLEDGMENTS

The authors thank Paulo de Sousa Pereira, Jun Li, and Kelly Houben for fruitful discussions. This work is supported by the FNRS projects, “crédit de démarrage U.Lg.,” the MP1201 COST Action, and by the Methusalem Funding of the Flemish Government. The work of A.N.V. and D.A.C. was supported by RFBR Grants No. 13-02-00174 and No. 12-02-90405. J.V.d.V. acknowledges support from FWO-VI.

*m.mohamed@ulg.ac.be

¹Z.-A. Ren, W. Lu, J. Yang, W. Yi, X.-L. Shen, Z.-C. Li, G.-C. Che, X.-L. Dong, L.-L. Sun, F. Zhou, and Z.-X. Zhao, *Chin. Phys. Lett.* **25**, 2215 (2008).

²H. Kito, H. Eisaki, and A. Iyo, *J. Phys. Soc. Jpn.* **77**, 063707 (2008).

³J. Yang, Z.-C. Li, W. Lu, W. Yi, X.-L. Shen, Z.-A. Ren, G.-C. Che, X.-L. Dong, L.-L. Sun, F. Zhou, and Z.-X. Zhao, *Supercond. Sci. Technol.* **21**, 082001 (2008).

⁴C. Wang, L. Li, S. Chi, Z. Zhu, Z. Ren, Y. Li, Y. Wang, X. Lin, Y. Luo, S. Jiang, X. Xu, G. Cao, and Z. Xu, *Europhys. Lett.* **83**, 67006 (2008).

⁵Y. Mizuguchi, F. Tomioka, S. Tsuda, T. Yamaguchi, and Y. Takano, *Appl. Phys. Lett.* **93**, 152505 (2008).

⁶A. E. Böhrner, F. Hardy, F. Eilers, D. Ernst, P. Adelman, P. Schweiss, T. Wolf, and C. Meingast, *Phys. Rev. B* **87**, 180505(R) (2013).

⁷D. Chareev, E. Osadchii, T. Kuzmichev, J.-Y. Lin, S. Kuzmichev, O. Volkova, and A. Vasiliev, *CrystEngComm* **15**, 1989 (2013).

⁸R. Hu, H. Lei, M. Abeykoon, E. S. Bozin, S. J. L. Billinge, J. B. Warren, T. Siegrist, and C. Petrovic, *Phys. Rev. B* **83**, 224502 (2011).

⁹S. Medvedev, T. M. McQueen, I. A. Troyan, T. Palasyuk, M. I. Eremets, R. J. Cava, S. Naghavi, F. Casper, V. Ksenofontov, G. Wortmann, and C. Felser, *Nat. Mater.* **8**, 630 (2009).

¹⁰A. Subedi, L. Zhang, D. J. Singh, and M. H. Du, *Phys. Rev. B* **78**, 134514 (2008).

¹¹J. K. Dong, T. Y. Guan, S. Y. Zhou, X. Qiu, L. Ding, C. Zhang, U. Patel, Z. L. Xiao, and S. Y. Li, *Phys. Rev. B* **80**, 024518 (2009).

¹²H. Lei, D. Graf, R. Hu, Hyejin Ryu, E. S. Choi, S. W. Tozer, and C. Petrovic, *Phys. Rev. B* **85**, 094515 (2012).

¹³J.-Y. Lin, Y. S. Hsieh, D. A. Chareev, A. N. Vasiliev, Y. Parsons, and H. D. Yang, *Phys. Rev. B* **84**, 220507(R) (2011).

¹⁴R. Khasanov, K. Conder, E. Pomjakushina, A. Amato, C. Baines, Z. Bukowski, J. Karpinski, S. Katrych, H.-H. Klauss, H. Luetkens, A. Shengelaya, and N. D. Zhigadlo, *Phys. Rev. B* **78**, 220510(R) (2008).

¹⁵C.-L. Song, Y.-L. Wang, P. Cheng, Y.-P. Jiang, W. Li, T. Zhang, Z. Li, K. He, L. Wang, J.-F. Jia, H.-H. Hung, C. Wu, X. Ma, X. Chen, and Q.-K. Xue, *Science* **332**, 1410 (2011).

¹⁶H.-H. Hung, C.-L. Song, X. Chen, X. Ma, Q.-k. Xue, and C. Wu, *Phys. Rev. B* **85**, 104510 (2012).

- ¹⁷C. Ren, Z.-S. Wang, H.-Q. Luo, H. Yang, L. Shan, and H.-H. Wen, *Phys. Rev. Lett.* **101**, 257006 (2008).
- ¹⁸K. Sasmal, B. Lv, Z. Tang, F. Y. Wei, Y. Y. Xue, A. M. Guloy, and C. W. Chu, *Phys. Rev. B* **81**, 144512 (2010).
- ¹⁹Y. Song, J. Ghim, J. Yoon, K. Lee, M. Jung, H. Ji, J. Shim, and Y. Kwon, *Europhys. Lett.* **94**, 57008 (2011).
- ²⁰X. L. Wang, S. X. Dou, Z.-A. Ren, W. Yi, Z.-C. Li, Z.-X. Zhao, and S.-I. Lee, *J. Phys.: Condens. Matter* **21**, 205701 (2009).
- ²¹C. Ren, Z.-S. Wang, H. Yang, X. Zhu, L. Fang, G. Mu, L. Shan, and H.-H. Wen, arXiv:0804.1726.
- ²²C. Martin, R. T. Gordon, M. A. Tanatar, H. Kim, N. Ni, S. L. Budko, P. C. Canfield, H. Luo, H. H. Wen, Z. Wang, A. B. Vorontsov, V. G. Kogan, and R. Prozorov, *Phys. Rev. B* **80**, 020501(R) (2009).
- ²³R. T. Gordon, N. Ni, C. Martin, M. A. Tanatar, M. D. Vannette, H. Kim, G. D. Samolyuk, J. Schmalian, S. Nandi, A. Kreyssig, A. I. Goldman, J. Q. Yan, S. L. Budko, P. C. Canfield, and R. Prozorov, *Phys. Rev. Lett.* **102**, 127004 (2009).
- ²⁴V. V. Moshchalkov, M. Menghini, T. Nishio, Q. H. Chen, A. V. Silhanek, V. H. Dao, L. F. Chibotaru, N. D. Zhigadlo, and J. Karpinski, *Phys. Rev. Lett.* **102**, 117001 (2009).
- ²⁵N. V. Orlova, A. A. Shanenko, M. V. Milosevic, F. M. Peeters, A. V. Vagov, and V. M. Axt, *Phys. Rev. B* **87**, 134510 (2013).
- ²⁶J. Carlström, J. Garaud, and E. Babaev, *Phys. Rev. B* **84**, 134518 (2011).
- ²⁷V. V. Moshchalkov, J. Y. Henry, C. Marin, J. Rossat-Mignod, and J. F. Jacquot, *Physica C* **175**, 407 (1991).
- ²⁸R. Prozorov, N. Ni, M. A. Tanatar, V. G. Kogan, R. T. Gordon, C. Martin, E. C. Blomberg, P. Prommapan, J. Q. Yan, S. L. Budko, and P. C. Canfield, *Phys. Rev. B* **78**, 224506 (2008).
- ²⁹A. K. Pramanik, L. Harnagea, C. Nacke, A. U. B. Wolter, S. Wurmehl, V. Kataev, and B. Büchner, *Phys. Rev. B* **83**, 094502 (2011).
- ³⁰D. Giller, A. Shaulov, R. Prozorov, Y. Abulafia, Y. Wolfus, L. Burlachkov, Y. Yeshurun, E. Zeldov, V. M. Vinokur, J. L. Peng, and R. L. Greene, *Phys. Rev. Lett.* **79**, 2542 (1997).
- ³¹A. K. Pramanik, M. Abdel-Hafiez, S. Aswartham, A. U. B. Wolter, S. Wurmehl, V. Kataev, and B. Büchner, *Phys. Rev. B* **84**, 064525 (2011).
- ³²B. B. Goodman and M. Wertheimer, *Phys. Lett.* **18**, 236 (1965).
- ³³S. H. Goedmoed, C. Van Kolmeschate, J. W. Metselaar, and D. De Kier, *Physica* **31**, 573 (1965).
- ³⁴B. Shen, P. Cheng, Z. Wang, L. Fang, C. Ren, L. Shan, and H.-H. Wen, *Phys. Rev. B* **81**, 014503 (2010).
- ³⁵D. Bhoi, P. Mandal, P. Choudhury, S. Dash, and A. Banerjee, *Physica C* **471**, 258 (2011).
- ³⁶R. W. Hu, E. S. Bozin, J. B. Warren, and C. Petrovic, *Phys. Rev. B* **80**, 214514 (2009).
- ³⁷H. Lei and C. Petrovic, *Phys. Rev. B* **83**, 184504 (2011).
- ³⁸M. Angst, R. Puzniak, A. Wisniewski, J. Jun, S. M. Kazakov, J. Karpinski, J. Roos, and H. Keller, *Phys. Rev. Lett.* **88**, 167004 (2002).
- ³⁹E. H. Brandt, *Phys. Rev. B* **60**, 11939 (1999).
- ⁴⁰E. Pardo, D.-X. Chen, and A. Sanchez, *J. Appl. Phys.* **96**, 5365 (2004).
- ⁴¹K. Papagelis, J. Arvanitidis, K. Prassides, A. Schenck, T. Takenobu, and Y. Iwasa, *Europhys. Lett.* **61**, 254 (2003).
- ⁴²H. Lei, R. Hu, and C. Petrovic, *Phys. Rev. B* **84**, 014520 (2011).
- ⁴³A. Carrington and F. Manzano, *Physica C* **385**, 205 (2003).
- ⁴⁴A. Brinkman, A. A. Golubov, H. Rogalla, O. V. Dolgov, J. Kortus, Y. Kong, O. Jepsen, and O. K. Andersen, *Phys. Rev. B* **65**, 180517(R) (2002).
- ⁴⁵V. A. Gasparov, N. S. Sidorov, and I. I. Zver'kova, *Phys. Rev. B* **73**, 094510 (2006).
- ⁴⁶R. Khasanov, A. Shengelaya, A. Maisuradze, F. La Mattina, A. Bussmann-Holder, H. Keller, and K. A. Müller, *Phys. Rev. Lett.* **98**, 057007 (2007).
- ⁴⁷L. Shan, Y. Huang, H. Gao, Y. Wang, S. L. Li, P. C. Dai, F. Zhou, J. W. Xiong, W. X. Ti, and H. H. Wen, *Phys. Rev. B* **72**, 144506 (2005).
- ⁴⁸F. Hardy, P. Burger, T. Wolf, R. A. Fisher, P. Schweiss, P. Adelman, R. Heid, R. Fromknecht, R. Eder, D. Ernst, H. v. Löhneysen, and C. Meingast, *Europhys. Lett.* **91**, 47008 (2010).
- ⁴⁹P. Popovich, A. V. Boris, O. V. Dolgov, A. A. Golubov, D. L. Sun, C. T. Lin, R. K. Kremer, and B. Keimer, *Phys. Rev. Lett.* **105**, 027003 (2010).
- ⁵⁰M. Abdel-Hafiez, S. Aswartham, S. Wurmehl, V. Grinenko, C. Hess, S.-L. Drechsler, S. Johnston, A. U. B. Wolter, B. Büchner, H. Rosner, and L. Boeri, *Phys. Rev. B* **85**, 134533 (2012).
- ⁵¹T. Shibauchi, K. Hashimoto, R. Okazaki, and Y. Matsuda, *Physica C* **469**, 590 (2009).
- ⁵²Ya. G. Ponomarev, S. A. Kuzmichev, M. G. Mikheev, M. V. Sudakova, S. N. Tchesnokov, O. S. Volkova, A. N. Vasiliev, T. Hänke, C. Hess, G. Behr, R. Klingeler, and B. Büchner, *Phys. Rev. B* **79**, 224517 (2009).
- ⁵³T. Kato, Y. Mizuguchi, H. Nakamura, T. Machida, H. Sakata, and Y. Takano, *Phys. Rev. B* **80**, 180507(R) (2009).
- ⁵⁴T. M. McQueen, Q. Huang, V. Ksenofontov, C. Felser, Q. Xu, H. Zandbergen, Y. S. Hor, J. Allred, A. J. Williams, D. Qu, J. Checkelsky, N. P. Ong, and R. J. Cava, *Phys. Rev. B* **79**, 014522 (2009).
- ⁵⁵A. J. Williams, T. M. McQueen, and R. J. Cava, *Solid State Commun.* **149**, 1507 (2009).
- ⁵⁶C. C. Homes, A. Akrap, J. S. Wen, Z. J. Xu, Z. W. Lin, Q. Li, and G. D. Gu, *Phys. Rev. B* **81**, 180508(R) (2010).
- ⁵⁷Ya. G. Ponomarev, S. A. Kuzmichev, M. G. Mikheev, M. V. Sudakova, S. N. Tchesnokov, T. E. Shanygina, O. S. Volkova, A. N. Vasiliev, and Th. Wolf, *J. Exp. Theor. Phys.* **113**, 459 (2011).
- ⁵⁸H. Kim, C. Martin, R. T. Gordon, M. A. Tanatar, J. Hu, B. Qian, Z. Q. Mao, R. W. Hu, C. Petrovic, N. Salovich, R. Giannetta, and R. Prozorov, *Phys. Rev. B* **81**, 180503 (2010).
- ⁵⁹S. He, J. He, W. Zhang, L. Zhao, D. Liu, X. Liu, D. Mou, Y.-B. Ou, Q. YanWang, Z. Li, L. Wang, Y. Peng, Y. Liu, C. Chen, L. Yu, G. Liu, X. Dong, J. Zhang, C. Chen, Z. Xu, X. Chen, X. Ma, Q. Xue, and X. J. Zhou, *Nat. Mater.* **12**, 605 (2013).