Superconducting state coexisting with a phase-separated static magnetic order in (Ba,K)Fe$_2$As$_2$, (Sr,Na)Fe$_2$As$_2$, and CaFe$_2$As$_2$

AUTHOR(S):
Goko, T.; Aczel, A. A.; Baggio-Saitovitch, E.; Bud'ko, S. L.; Canfield, P. C.; Carlo, J. P.; Chen, G. F.; ... Yamamoto, T.; Yu, W.; Uemura, Y. J.

CITATION:
Goko, T. ...[et al]. Superconducting state coexisting with a phase-separated static magnetic order in (Ba,K)Fe$_2$As$_2$, (Sr,Na)Fe$_2$As$_2$, and CaFe$_2$As$_2$. PHYSICAL REVIEW B 2009, 80(2): 024508.

ISSUE DATE:
2009-07

URL:
http://hdl.handle.net/2433/109861

RIGHT:
© 2009 The American Physical Society
Superconducting state coexisting with a phase-separated static magnetic order in (Ba,K)Fe$_2$As$_2$, (Sr,Na)Fe$_2$As$_2$, and CaFe$_2$As$_2$

T. Goko,1,2,3 A. A. Aczel,3 E. Baggio-Saitovitch,4 S. L. Bud’ko,5 P. C. Canfield,5 J. P. Carlo,2 G. F. Chen,6 Pengcheng Dai,7 A. C. Hamann,8 W. Z. Hu,6 H. Kageyama,9 G. M. Luke,9 J. L. Luo,6 B. Nachumi,2 N. Ni,5 D. Reznik,8 D. R. Sanchez-Candela,4 A. T. Savici,10 K. J. Sikes,2 N. L. Wang,6 C. R. Wiebe,11 T. J. Williams,3 T. Yamamoto,9 W. Yu,3 and Y. J. Uemura2,6

1TRIUMF, 4004 Wesbrook Mall, Vancouver, British Columbia, Canada V6T 2A3
2Department of Physics, Columbia University, New York, New York 10027, USA
3Department of Physics and Astronomy, McMaster University, Hamilton, Ontario, Canada L8S 4M1
4Centro Brasileiro de Pesquisas Físicas, Rua Xavier Sigaud 150 Urca, CEP 22290-180 Rio de Janeiro, RJ, Brazil
5Department of Physics and Astronomy and Ames Laboratory, Iowa State University, Ames, Iowa 50011, USA
6Beijing National Laboratory for Condensed Matter Physics, Institute of Physics, Chinese Academy of Sciences, Beijing 100080, People’s Republic of China
7Department of Physics and Astronomy, University of Tennessee, Knoxville, Tennessee 37996, USA
8Forschungszentrum Karlsruhe, Institut für Festkörperphysik, Postfach 3640, D-76021 Karlsruhe, Germany
9Department of Chemistry, Kyoto University, Kyoto 606-8502, Japan
10Department of Physics and Astronomy, Johns Hopkins University, Baltimore, Maryland 21218, USA
11Department of Physics, Florida State University, Tallahassee, Florida 32310, USA

Received 20 August 2008; revised manuscript received 10 June 2009; published 14 July 2009

By muon spin-relaxation measurements on single-crystal specimens, we show that superconductivity in the Fe$_2$As$_2$ (A=Ca,Ba,Sr) systems, in both the cases of composition and pressure tunings, coexists with a strong static magnetic order in a partial volume fraction. The superfluid response from the remaining paramagnetic volume fraction of (Ba$_{0.55}$K$_{0.45}$)Fe$_2$As$_2$ exhibits a nearly linear variation in $T$ at low temperatures, suggesting an anisotropic energy gap with line nodes and/or multigap effects.

DOI: 10.1103/PhysRevB.80.024508 PACS numbers: 74.25.Dw, 74.25.Nf, 75.25.−z, 76.75.+i

The announcement of superconductivity in La(O,F)FeAs ($T_c=26$ K) (Ref. 1) triggered an unprecedented burst of research activities in FeAs-based superconductors and their parent compounds. By now, superconductivity has been reported in systems with four different crystal structures, including the “1111” systems RE(O,F)FeAs with rare earth=La, Nd, Ce, etc.,2 and the “122” systems AFe$_2$As$_2$ (A=Ba, Sr, Ca).3−5 Extensive measurements by neutron scattering,6−8 Mössbauer effect,9−11 and muon spin relaxation ($\mu$SR) (Refs. 11−15) have revealed collinear antiferromagnetic order in undoped parent compounds,6,16,17 hyperfine splitting of $^{57}$Fe Mössbauer spectra, and $\mu$SR frequencies indicative of a static moment size ranging between 0.3 and 0.8 Bohr magnetons per Fe,9,10,15 and nearly linear scaling between $T_c$ and the superfluid density in the 1111 systems following the trend found in cuprate and other exotic superconductors.18

In studies of magnetic phase diagrams of the 1111 systems, as a function of increasing (O,F) substitution, La(O,F)FeAs shows19 an abrupt and first-order-like evolution from an antiferromagnetic to superconducting state, Ce(O,F)FeAs shows nearly second-order-like evolution,7 and Sm(O,F)FeAs (Ref. 20) exhibits phase-separated coexistence of static magnetism and superconductivity in a small concentration region around the phase boundary. Despite these differences, superconductivity appears mostly in the region without static magnetic order in the 1111 systems, similar to the case of the cuprates. In contrast, very little has been reported on the phase diagrams of the 122 systems. Recent powder neutron measurements on (Ba,K)Fe$_2$As$_2$ (Ref. 8) found a phase diagram similar to the one for Sm(O,F)FeAs with coexisting long-range magnetic order and superconductivity near the phase boundary, without providing information on the volume fraction of the magnetically ordered region. We have also reported $\mu$SR measurements on a single crystal of (Ba$_{0.55}$K$_{0.45}$)Fe$_2$As$_2$ (Ref. 15) which found the coexistence of phase-separated static magnetic order and superconductivity. The superfluid density of this crystal was much lower than that in the corresponding 1111 systems with comparable $T_c$’s, which is suggestive of insufficient carrier doping. In the 122 systems, more definitive studies of magnetic phase diagrams can be expected due to the availability of large single crystals,21−23 improvement of the growth method, and applicability of pressure tuning free from randomness due to substitution.

In this paper, we report $\mu$SR measurements of superconducting single crystals of (Ba$_{0.55}$K$_{0.45}$)Fe$_2$As$_2$ ($T_c \sim 37$ K) and (Sr$_{0.5}$Na$_{0.5}$)Fe$_2$As$_2$ ($T_c \sim 35$ K) in ambient pressure, and of CaFe$_2$As$_2$ in ambient and applied pressure $p$ up to $p = 10$ kbar, performed at TRIUMF in Vancouver, Canada. The former two crystals were prepared at the Institute of Physics in Beijing using the FeAs flux method23 and weighed $\sim 100$ and 40 mg, respectively. As shown in Fig. 1, sharp superconducting transitions were observed in the magnetic susceptibility of both crystals, which suggests that they are of good quality. No anomaly due to a structural transition can be seen in the resistivity results. These crystals were mounted with their $ab$ planes perpendicular to the muon beam at the M20 channel. In our $\mu$SR measurements in transverse external fields applied perpendicular to the $ab$ plane, both of these crystals exhibited a superfluid response.
FIG. 1. (Color online) Temperature dependences of the resistivity and the magnetic susceptibility of our specimens of (a) (Ba$_{0.8}$K$_{0.2}$)Fe$_2$As$_2$ ($T_c \sim 37$ K) and (b) (Sr$_{0.5}$Na$_{0.5}$)Fe$_2$As$_2$ ($T_c \sim 35$ K). Due to the irregular shape of the specimen which prevents accurate estimate of the demagnetizing factor, we put the results of the latter system on a relative (arbitrary) scale. The inset figure of (b) shows the magnetic susceptibility obtained in the field-cooling and zero-field-cooling procedures.

Over 1 g of CaFe$_2$As$_2$ crystals (in more than 100 pieces) prepared in Ames Laboratory using the Sn flux method were mounted in a pressure cell having a sample space of 7 mm in diameter and 12 mm long. The cell was pressurized at room temperature before being mounted in the cryostat. Daphne oil was used as the pressure mediator. This was chosen because it is known not to solidify at room temperature up to $10$ kbar and so generates hydrostatic pressure over a wider pressure range than Fluorinert, which solidifies above $15$ kbar at room temperature. The crystals were aligned with their $ab$ planes perpendicular to the muon beam at the M9B channel, where the initial muon spin polarization is tuned to be perpendicular to the beam direction.

μSR measurements were performed in zero field (ZF) and weak transverse field (WTF) of $30–100$ G to study magnetic ordering, and in transverse field (TF) of $300–500$ G to study superfluid density. A recent study on (Ca,Sr)RuO$_3$ and MnSi in applied pressure has demonstrated μSR’s unique capability of determining volume fractions of regions with and without static magnetic order in systems having phase separation. Details of the μSR methods can be found in Refs. 24 and 25.

Figure 2 shows (a) the muon spin precession frequencies observed in ZF μSR and (b) the paramagnetic volume fraction determined from WTF-μSR measurements of single-crystal specimens of (Ba$_{0.8}$K$_{0.2}$)Fe$_2$As$_2$ ($T_c \sim 37$ K) and (Sr$_{0.5}$Na$_{0.5}$)Fe$_2$As$_2$ ($T_c \sim 35$ K). The results from the present work (solid and open symbols) are compared with those of the undoped parent compounds BaFe$_2$As$_2$ (Ref. 15) and SrFe$_2$As$_2$ (Ref. 10) (solid and broken lines).

In TF μSR, the precession signal from the paramagnetic volume fraction exhibits damping below $T_c$ due to an inhomogeneous field distribution in the flux vortex lattice. The relaxation rate $\sigma$, obtained by fitting the spectra to a Gaussian function $\exp(-\alpha^2I^2/2)$, is given by $\sigma \propto 1/\lambda - n_s/m^*$, where $\lambda$ is the penetration depth, $n_s$ is the superconducting carrier density, and $m^*$ is the effective mass. An increase in $\sigma$ was observed in both the (Ba,K) and (Sr,Na) crystals below the superconducting $T_c$'s. Since the statistical accuracy of the data is much better for the former system with the larger paramagnetic volume fraction, here we present the results of (Ba$_{0.5}$K$_{0.5}$)Fe$_2$As$_2$ in TF=500 G applied parallel to...
the $c$ axis in Figs. 3(a) and 3(b). The temperature dependence of $\sigma$ in (a) is nearly linear with $T$, as demonstrated by the good agreement with the scaled data from a YBa$_2$Cu$_{3-y}$O$_{6.95}$ (YBCO) system. The observed behavior is distinctly different from the case for an isotropic energy gap by the broken line representing a calculation for BCS $s$-wave coupling. The observed temperature dependence may be attributed to (1) line nodes in an anisotropic energy gap or (2) widely different magnitudes of multiple isotropic gaps as seen in calculations based on multiple bands. An angle resolved photoemission spectroscopy (ARPES) measurement on (Ba,K)Fe$_2$As$_2$ reported evidence for isotropic multiple gaps, consistent with theories based on an extended $s$-wave pairing.

The absolute value of $\sigma(T=0)$ is about a factor of 3 larger than that observed in (Ba$_{0.55}$K$_{0.45}$)Fe$_2$As$_2$ in our previous measurements. Given that the $H_{c2}$ anisotropy is relatively low near $T_c$, varying between 3.5 and 2.5 for $H<14 \, T$ (Ref. 21) and decreasing for higher fields, we plot the present results in the $\sigma(T=0)$ versus $T_c$ plot in Fig. 3(b) without single crystal to polycrystalline conversion corrections. The resulting point from the present (Ba,K) crystal (red solid square symbol) indicates that the present system has a superfluid density close to those in the 1111 systems with comparable $T_c$'s and that a sufficiently doped 122 FeAs system follows the nearly linear relationship between $T_c$ and $\eta/\mu^*$ found in the cuprates and 1111 systems.

The superconducting state can also be obtained by applying pressure (using a liquid pressure medium) to the undoped parent compounds of the 122 systems. In particular, the CaFe$_2$As$_2$ system shows superconductivity below $T\sim 10 \, K$ at relatively low pressures $p$ of 3–8 kbar, which is attainable using the available $\mu$SR piston-cylinder pressure cell. We studied static magnetic order of CaFe$_2$As$_2$ at ambient pressure and at $p=3.9, 6.2,$ and 9.9 kbar by performing WFT $\mu$SR with WFT=50 G. Solid symbols in Fig. 4(a) show the low-temperature ($T\rightarrow 0$) values of the volume fraction of the magnetically ordered region (from WFT data) as well as the $\mu$SR precession frequency in ZF $\mu$SR, which is proportional to the size of the ordered Fe moment. We present the resulting pressure-temperature phase diagram of CaFe$_2$As$_2$ in Fig. 4(c). The superconducting phase boundary in this figure is based on the reported resistivity results. Figure 4(c) indicates that a rather strong magnetism exists in a substantial volume fraction below $T=50–100 \, K$, which is well above the superconducting $T_c$ as in the cases of the (Ba,K) and (Sr,Na) crystals at ambient pressure.

Resistivity and neutron measurements in applied pressure, the former (the latter) using Fluorinert (He gas) as the pressure medium, have been reported on CaFe$_2$As$_2$ single crystals prepared by an identical method to that used for the present specimen.

In the resistivity studies, a sharp jump was observed at $T=170 \, K$ at ambient pressure, corresponding to the first-order tetragonal-to-orthorhombic structural phase transition below which magnetic order was detected both by neutrons and muons. With increasing pressure this feature broadens and ordering moves toward lower temperatures, which is qualitatively consistent with the present results in Fig. 4(c). The resistivity anomaly disappears above $p=4 \, kbar$, and the neutron magnetic Bragg-peak intensity at $T=50 \, K$ becomes nearly equal to the background level at $p=6.3 \, kbar$ [Fig. 1(c) in Ref. 33], while $\mu$SR detected magnetic order continuing to exist at $p=6.2 \, kbar$, albeit in a partial volume fraction.

Quite recently, additional neutron measurements under pressure, using Fluorinert (and not He gas) as the pressure medium, were performed to examine CaFe$_2$As$_2$ specimens...
FIG. 4. (Color online) (a) The volume fraction of regions without static magnetic order in CaFe$_2$As$_2$, as a function of temperature and pressure, determined by WTF-μSR measurements with WTF $\sim 50$ G. The points with closed (open) symbols were obtained in measurements with (without) a pressure cell. (b) Pressure dependence of the volume fraction of the magnetically ordered region (purple closed circles; left axis) from WTF μSR and the muon spin precession frequency (blue triangles; right axis) from ZF μSR at $T \sim 2$ K in CaFe$_2$As$_2$. (c) The phase diagram as a function of pressure and temperature in CaFe$_2$As$_2$. The $T_N$ values are taken from the reported resistivity results (Ref. 4). Upper, middle, and lower temperatures attached to the closed circle symbols for $T_N$ represent temperatures at which the volume fraction with static magnetic order becomes 30%, 50%, and 70% of the value at $T \rightarrow 0$, respectively.

grown by the same group$^{22}$ as those discussed in the present work. These neutron measurements revealed the coexistence of two structural phases (antiferromagnetic orthorhombic and nonmagnetic collapsed tetragonal phases) at low temperatures, which is also consistent with our μSR results. This behavior probably arises from the applied pressure not actually being truly hydrostatic, as CaFe$_2$As$_2$ seems to be strongly affected by any slight deviation from hydrostaticity. More specifically, superconductivity was not observed in resistivity and susceptibility measurements$^{35}$ of CaFe$_2$As$_2$ under He gas pressure, which is capable of providing true hydrostatic pressure over a much wider pressure temperature range than either Fluorinert or Daphne oil. The crystals investigated by Yu et al. were also grown by the same group$^{22}$ as those studied in the present work. Superconductivity, the coexistence of magnetically ordered and nonmagnetic fractions and the coexistence of the orthorhombic and collapsed tetragonal structures appear only under pressure using a liquid pressure medium. The magnetic order in a partial volume fraction may play an important role in the emergence of superconductivity in CaFe$_2$As$_2$.

The present μSR results do not provide direct evidence for distinguishing whether superconductivity lives uniformly over the entire volume or exclusively in the paramagnetic volume fraction. Evidence for the latter case has been reported in ARPES studies on a lightly doped single crystal of (Ba$_x$K)$_2$Fe$_2$As$_2$, which found an intensity ratio of an ungapped magnetic response to a gapped superconducting response comparable to the ratio of volumes with and without static magnetic order found by μSR. This situation is similar to the case of (La$_x$Eu,Sr)$_2$CuO$_4$ where the static volume fraction and superfluid density in μSR exhibit a tradeoff.$^{37}$

The present study does not provide an estimate of the length scale of the phase-separated regions. In La$_{2-x}$CuO$_4$, we estimated this length to be a few nanometers.

Phase separation at the border of magnetic and superconducting states has been found also in the organic (BEDT-TTF)$_2$X superconductors$^{39}$ and CeCu$_2$Si$_2$. First-order phase transitions, similar to those in the La$_{1111}$FeAs system, have been found in the A$_x$C$_{60}$, Ce(Co,Cd)In$_5$, and CeRhIn$_5$ (Ref. 42) systems. Phase separation was also discovered at the border of a collinear antiferromagnetic state and a nonmagnetic spin-gap state in an insulating $J_1$–$J_2$ spin system Cu(Cl,Br)La(Nb,Ta)$_2$O$_7$,$^{43}$ and at the border of an itinerant heliferromagnetic state and paramagnetic state in MnSi and (Sr,Ca)RuO$_3$. Further exploration of these behaviors will lead to a better understanding of superconductivity and magnetism in correlated-electron systems.

Regarding the pairing symmetry, available experimental results on the 1111 and 122 systems are divided between those favoring an isotropic nodeless gap$^{44}$ and those supporting line nodes and/or multigap features.$^{26,27,45}$ The (Ba$_{0.5}$K$_{0.5}$)$_2$Fe$_2$As$_2$ result in Fig. 3(a) has established at least one definite case which does not agree with a single isotropic energy gap. The different temperature dependences of the superfluid density between the two (Ba$_x$K)$_2$Fe$_2$As$_2$ crystals in Fig. 3(a) may be due to a doping dependence of multiple gap magnitudes, as proposed in Ref. 30. The nearly linear relationship between $T_N$ and the superfluid density [Fig. 3(b)] followed by cuprates, 1111 FeAs, 122 FeAs, and A$_x$C$_{60}$ sys-
SUPERCONDUCTING STATE COEXISTING WITH A...

*Author to whom correspondence should be addressed: tomo@lorentz.phys.columbia.edu

1 Y. Kamihara, T. Watanabe, M. Hirano, and H. Hosono, J. Am. Chem. Soc. 130, 3296 (2008).
2 X. H. Chen, T. Wu, G. Wu, R. H. Liu, H. Chen, and D. F. Fang, Nature (London) 453, 761 (2008); G. F. Chen, Z. Li, D. Wu, G. Li, W. Z. Hu, J. Dong, P. Zheng, J. L. Luo, and N. L. Wang, Phys. Rev. Lett. 100, 247002 (2008); Z.-A. Ren, Guang-Can Che, Xiao-Li Dong, Jie Yang, Wei Lu, Wei Yi, Xiao-Li Shen, Zheng-Cai Li, Li-Ling Sun, Fang Zhou, and Zhong-Xian Zhao, EPL 83, 17002 (2008).
3 M. Rotter, M. Tagel, and D. Johrendt, Phys. Rev. Lett. 101, 107006 (2008).
4 M. S. Torikachvili, S. L. Bud’ko, N. Ni, and P. C. Canfield, Phys. Rev. Lett. 101, 057006 (2008).
5 P. L. Alireza, Y. T. Chris Ko, J. Gillett, C. M. Petrone, J. M. Cole, S. E. Sebastian, and G. G. Lonzarich, J. Phys.: Condens. Matter 21, 012208 (2009).
6 C. de la Cruz, Q. Huang, J. W. Lynn, Jiying Li, W. Ratcliff II, J. W. Lynn, and Pengcheng Dai, Nature (London) 453, 899 (2008).
7 J. Zhao, Q. Huang, Clarina de la Cruz, Shiliang Li, J. W. Lynn, Y. Chen, M. A. Green, G. F. Chen, G. Li, Z. Li, J. L. Luo, N. L. Wang, and Pengcheng Dai, Nature Mater. 7, 953 (2008).
8 H. Chen, Y. Ren, Y. Qiu, Wei Bao, R. H. Liu, G. Wu, T. Wu, Y. L. Xie, X. F. Wang, Q. Huang, and X. H. Chen, EPL 85, 17006 (2009).
9 M. Rotter, M. Tagel, and D. Johrendt, I. Schellenberg, W. Hermes, R. Pottgen, Phys. Rev. B 78, 020503(R) (2008).
10 A. Jeshie, N. Caro-Canales, H. Rosner, H. Borrman, A. Ormechi, D. Kasinathan, H. H. Klauss, H. Luetskens, R. Khasanov, A. Amato, A. Hosier, K. Kaneko, C. Krellner, and C. Geibel, Phys. Rev. B 78, 180504(R) (2008).
11 H.-H. Klauss, H. Luetskens, R. Klingeler, C. Hess, F. J. Litterst, M. Kraken, M. K. Korshunov, I. Eremeev, S.-L. Drechsler, R. Khasanov, A. Amato, J. Hamann-Borrero, N. Leps, A. Konradt, G. Behr, J. Werner, and B. Büncher, Phys. Rev. Lett. 101, 077005 (2008).
12 H. Luetskens, H.-H. Klauss, R. Khasanov, A. Amato, R. Klingeler, I. Hellmann, N. Leps, A. Konradt, C. Hess, A. Köhler, G. Behr, J. Werner, and B. Büchner, Phys. Rev. Lett. 101, 097009 (2008); R. Khasanov, H. Luetskens, A. Amato, H.-H. Klauss, Z.-A. Ren, J. Yang, W. Lu, and Z.-X. Zhao, Phys. Rev. B 78, 092506 (2008).
13 A. J. Drew, F. L. Pratt, T. Lancaster, S. J. Blundell, P. J. Baker, R. H. Liu, G. Wu, X. H. Chen, I. Watanabe, V. K. Malik, A. Dubroka, K. W. Kim, M. Rössle, and C. Bernhard, Phys. Rev. Lett. 101, 097010 (2008).
14 J. P. Carlo, Y. J. Uemura, T. Goko, G. J. MacDougall, J. A. Rodriguez, W. Yu, G. M. Luke, Pengcheng Dai, N. Shannon, S. Miyasaka, S. Suzuki, S. Tajima, G. F. Chen, W. Z. Hu, J. L. Luo, and N. L. Wang, Phys. Rev. Lett. 102, 087001 (2009).
15 A. A. Azcel, E. Baggio-Saitovitch, S. L. Budko, P. C. Canfield, J. P. Carlo, G. F. Chen, Pengcheng Dai, T. Goko, W. Z. Hu, G. M. Luke, J. L. Luo, N. Ni, R. Sanchez-Cardela, F. Tafiri, N. L. Wang, T. J. Williams, W. Yu, and Y. J. Uemura, Phys. Rev. B 78, 214503 (2008).
16 Q. Huang, Y. Qiu, W. Bao, M. A. Green, J. W. Lynn, Y. C. Gasparovic, T. Wu, G. Wu, and X. H. Chen, Phys. Rev. Lett. 101, 257003 (2008).
17 A. I. Goldman, D. N. Argyriou, B. Ouladdiaf, T. Chatterji, A. Kreyssig, S. Nandi, N. Ni, S. L. Bud’ko, P. C. Canfield, and R. J. McQueeney, Phys. Rev. B 78, 100506(R) (2008).
18 Y. J. Uemura, G. M. Luke, B. J. Sternlieb, J. H. Brewer, J. F. Carolan, W. N. Hardy, R. Kadono, J. R. Kempton, R. F. Kiefl, S. R. Kreitzman, P. Mulhern, T. M. Riseman, D. Li, Williams, B. X. Yang, S. Uchida, H. Takagi, J. Gopalakrishnan, A. W. Sleight, M. A. Subramanian, C. L. Chien, M. Z. Cieplak, Gang Xiao, V. Y. Lee, B. W. Statt, C. E. Stronach, W. J. Kossler, and X. H. Yu, Phys. Rev. Lett. 62, 2317 (1989); Y. J. Uemura, L. P. Le, G. M. Luke, B. J. Sternlieb, W. D. Wu, J. H. Brewer, T. M. Riseman, C. L. Seaman, M. B. Maple, M. Ishikawa, D. G. Hinks, J. D. Jorgensen, G. Saito, and H. Yamochi, ibid. 66, 2665 (1991); Y. J. Uemura, Physica B 374-375, 1 (2006).
19 H. Luetskens, H.-H. Klauss, M. Kraken, F. J. Litterst, T. Dellmann, R. Klingeler, C. Hess, R. Khasanov, A. Amato, C. Baines, M. Kosmala, O. J. Schumann, M. Braden, J. Hamann-Borrero, N. Leps, A. Konradt, G. Behr, J. Werner, and B. Büchner, Nature Mater. 8, 305 (2009).
20 A. J. Drew, Ch. Niehammer, P. J. Baker, F. L. Pratt, S. J. Blundell, T. Lancaster, R. H. Liu, G. Wu, X. H. Chen, I. Watanabe, V. K. Malik, A. Dubroka, M. Rössle, K. W. Kim, C. Baines, and C. Bernhard, Nature Mater. 8, 310 (2009).
21 N. Ni, S. L. Bud’ko, A. Kreyssig, S. Nandi, G. E. Rustan, A. I. Goldman, S. Gupta, J. D. Corbett, A. Kracher, and P. C. Canfield, Phys. Rev. B 78, 014507 (2008).
22 N. Ni, S. Nandi, A. Kreyssig, A. I. Goldman, E. D. Mun, S. L. Bud’ko, and P. C. Canfield, Phys. Rev. B 78, 014523 (2008).
23 G. F. Chen, Z. Li, J. Dong, G. Li, W. Z. Hu, X. D. Zhang, X. H. Song, P. Zheng, N. L. Wang, and J. L. Luo, Phys. Rev. B 78, 224512 (2008).
24 Y. J. Uemura, T. Goko, I. M. Gat-Malureanu, J. P. Carlo, P. L. Russo, A. T. Savici, A. Azcel, G. J. MacDougall, J. A. Rodriguez, G. M. Luke, S. R. Dunsiger, A. McCollam, J. Arafi, Ch. Pflieger, P. Böni, K. Yoshimura, E. Baggio-Saitovitch, M. B. Fontes, J. Larrea, Y. V. Sushko, and J. Sereni, Nat. Phys. 3, 29

SUPERCONDUCTING STATE COEXISTING WITH A...
(2007).

25 J. E. Sonier, J. H. Brewer, and R. F. Kiefl, Rev. Mod. Phys. 72, 769 (2000).

26 L. Benfatto, M. Capone, S. Caprara, C. Castellani, and C. Di Castro, Phys. Rev. B 78, 140502(R) (2008).

27 C. Ren, Z.-S. Wang, H.-Q. Luo, H. Yang, L. Shan, and H.-H. Wen, Phys. Rev. Lett. 101, 257006 (2008).

28 H. Ding, P. Richard, K. Nakayama, K. Sugawara, T. Arakane, Y. Sekiba, A. Takayama, S. Souma, T. Sato, T. Takahashi, Z. Wang, X. Dai, Z. Fang, G. F. Chen, J. L. Luo, and N. L. Wang, EPL 83, 47001 (2008).

29 I. I. Mazin, D. J. Singh, M. D. Johannes, and M. H. Du, Phys. Rev. Lett. 101, 057003 (2008); K. Kuroki, S. Onari, R. Arita, H. Usui, Y. Tanaka, H. Kontani, and H. Aoki, ibid. 101, 087004 (2008).

30 M. M. Parish, J. Hu, and B. A. Bernevig, Phys. Rev. B 78, 144514 (2008).

31 M. M. Altarawneh, K. Collar, C. H. Mielke, N. Ni, S. L. Bud’ko, and P. C. Canfield, Phys. Rev. B 78, 220505(R) (2008); H. Q. Yuan, J. Singleton, F. F. Balakirev, S. A. Baily, G. F. Chen, J. L. Luo, and N. L. Wang, Nature (London) 457, 565 (2009).

32 P. L. Russo,, C. R. Wiebe, Y. J. Uemura, A. T. Savici, G. J. MacDougall, J. Rodriguez, G. M. Luke, N. Kaneko, H. Eisaki, M. Greven, O. P. Vajk, S. Ono, Yoichi Ando, K. Fujita, K. M. Kojima, and S. Uchida, Phys. Rev. B 75, 054511 (2007).

33 A. Kreyssig, M. A. Green, Y. Lee, G. D. Samolyuk, P. Zajdel, J. W. Lynn, S. L. Bud’ko, M. S. Torikachvili, N. Ni, S. Nandi, J. B. Leão, S. J. Poulton, D. N. Argyriou, B. N. Harmon, R. J. McQueeney, P. C. Canfield, and A. I. Goldman, Phys. Rev. B 78, 184517 (2008).

34 A. I. Goldman, A. Kreyssig, K. Prokeš, D. K. Pratt, D. N. Argyriou, J. W. Lynn, S. Nandi, S. A. J. Kimber, Y. Chen, Y. B. Lee, G. Samolyuk, J. B. Leão, S. J. Poulton, S. L. Bud’ko, N. Ni, P. C. Canfield, B. N. Harmon, and R. J. McQueeney, Phys. Rev. B 79, 024513 (2009).

35 W. Yu, A. A. Aczel, T. J. Williams, S. L. Bud’ko, N. Ni, P. C. Canfield, and G. M. Luke, Phys. Rev. B 79, 020511(R) (2009).

36 D. V. Evstushinsky, D. S. Inosov, V. B. Zabolotnyy, A. Koitzsch, M. Knupfer, B. Büchner, M. S. Viazovska, G. L. Sun, V. Hinkov, A. V. Boris, C. T. Lin, B. Keimer, A. Varykhalov, A. A. Kordyuk, and S. V. Borisenko, Phys. Rev. B 79, 054517 (2009).

37 K. M. Kojima, S. Uchida, Y. Fudamoto, I. M. Gat, M. I. Larkin, Y. J. Uemura, and G. M. Luke, Physica B 326, 316 (2003).

38 A. T. Savici, Y. Fudamoto, I. M. Gat, T. Ito, M. I. Larkin, Y. J. Uemura, G. M. Luke, K. M. Kojima, Y. S. Lee, M. A. Kastner, R. J. Birgeneau, and K. Yamada, Phys. Rev. B 66, 014524 (2002).

39 T. Sasaki, N. Yoneyama, and N. Kobayashi, Phys. Rev. B 77, 054505 (2008).

40 G. M. Luke, A. Keren, K. Kojima, L. P. Le, B. J. Sternlieb, W. D. Wu, Y. J. Uemura, Y. Onuki, and T. Komatsubara, Phys. Rev. Lett. 73, 1853 (1994).

41 J. Arvanitidis, K. Papagelis, Y. Takabayashi, T. Takenobu, Y. Iwasa, M. J. Rosseinsky, and K. Prassides, J. Phys.: Condens. Matter 19, 386235 (2007).

42 T. Park, F. Ronning, H. Q. Yuan, M. B. Salamon, R. Movshovich, I. L. Sarrao, and J. D. Thompson, Nature (London) 440, 65 (2006).

43 Y. J. Uemura, A. Aczel, Y. Ajiro, J. Carlo, T. Goko, D. Goldfeld, A. Kitada, G. Luke, G. MacDougall, I. Mihăilescu, J. Rodriguez, P. Russo, Y. Tsujimoto, C. Wiebe, T. Williams, T. Yamamoto, K. Yoshimura, and H. Kageyama, arXiv:0806.2021 (unpublished).

44 K. Hashimoto, T. Shibuchi, T. Kato, K. Ikada, R. Okazaki, H. Shishido, M. Ishikado, H. Kito, A. Iyo, H. Eisaki, S. Shamoto, and Y. Matsuda, Phys. Rev. Lett. 102, 017002 (2009).

45 L. Shan, Y. Wang, X. Zhu, G. Mu, L. Fang, C. Ren, and H.-H. Wen, EPL 83, 57004 (2008); H.-J. Graf, D. Paar, G. Lang, N. J. Curro, G. Behr, J. Werner, J. Hamann-Borger, C. Hess, N. Leps, R. Klingeler, and B. Büchner, Phys. Rev. Lett. 101, 047003 (2008).

46 Y. J. Uemura, arXiv:0811.1546 (unpublished).