

Electronic inhomogeneity and competing phases in electron-doped superconducting $\text{Pr}_{0.88}\text{LaCe}_{0.12}\text{CuO}_{4-\delta}$

Pengcheng Dai,^{1,2,*} H. J. Kang,¹ H. A. Mook,² M. Matsuura,² J. W. Lynn,³ Y. Kurita,⁴ Seiki Komiya,⁴ and Yoichi Ando⁴

¹*Department of Physics and Astronomy, The University of Tennessee, Knoxville, Tennessee 37996-1200, USA*

²*Condensed Matter Sciences Division, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831-6393, USA*

³*NIST Center for Neutron Research, National Institute of Standards and Technology, Gaithersburg, Maryland 20899-8562, USA*

⁴*Central Research Institute of Electric Power Industry, Komae, Tokyo 201-8511, Japan*

(Received 15 May 2004; published 21 March 2005)

We use neutron scattering to demonstrate that electron-doped superconducting $\text{Pr}_{0.88}\text{LaCe}_{0.12}\text{CuO}_{4-\delta}$ in the underdoped regime is electronically phase separated in the ground state, showing the coexistence of a superconducting phase with a three-dimensional antiferromagnetically ordered phase and a quasi-two-dimensional spin-density wave modulation. The Néel temperature of both antiferromagnetic phases decreases linearly with increasing superconducting transition temperature (T_c) and vanishes when optimal superconductivity is achieved. These results indicate that the electron-doped copper oxides are close to a quantum critical point, where the delicate energetic balance between different competing states leads to microscopic heterogeneity.

DOI: 10.1103/PhysRevB.71.100502

PACS number(s): 74.72.Jt, 61.12.Ld, 75.25.+z, 75.50.Ee

One of the most important unresolved problems in high-transition temperature (high- T_c) superconductors is the determination of the microscopic state when charge carriers, either holes or electrons, are introduced to the CuO_2 planes of their insulating long-range antiferromagnetically ordered parent compounds.¹ One school of thought suggests that the doped charge carriers segregate into inhomogeneous patterns, such as stripes, to allow the antiferromagnetic (AF) regions to survive.² In this picture, the observed quasi-two-dimensional (2D) incommensurate spin-density wave (SDW) in hole-doped high- T_c superconductors, such as $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$ (Ref. 3) and $\text{La}_2\text{CuO}_{4+\delta}$ (Ref. 4), is due to remnants of the AF insulating phase that compete with superconductivity (SC).^{5,6} Alternatively, carrier doping into the CuO_2 planes may evolve the system through a quantum phase transition (QPT) between competing ordered and disordered phases.⁷ While much is unknown about these competing phases and their relationship to SC, the presence of such a QPT has been reported in both hole-doped^{8–11} and electron-doped¹² materials.

In this paper, we use neutron scattering to demonstrate that the AF to SC transition in electron-doped $\text{Pr}_{0.88}\text{LaCe}_{0.12}\text{CuO}_{4-\delta}$ (PLCCO) is close to a QPT (Ref. 13) and the proximity to such a QPT leads to an inhomogeneous electronic state due to competing order parameters.¹⁴ Specifically, we find that when SC first appears in PLCCO with increasing doping, a quasi-2D commensurate SDW modulation is also induced, and both coexist with the three-dimensional (3D) AF state. The Néel temperature (T_N) of both the 2D and 3D AF phases decreases linearly with increasing T_c and vanishes when optimal SC is achieved [Fig. 1(f)]. Combined with earlier reports,^{3–6,15–17} our discovery establishes that the quasi-2D SDW modulation and its associated electronic inhomogeneity are general features of charge-carrier-doped copper oxides. Such microscopic heterogeneity naturally results from the competition of different order parameters near a QPT.

Traditionally, the transition from AF to SC in copper ox-

ides is achieved by chemical doping, which also introduces additional complications such as structural and/or chemical disorder that may mask the nature of the transition.¹³ We choose to study this transition in electron-doped PLCCO because, as grown, these materials are nonsuperconducting (NSC) AF insulators and SC can be achieved by only annealing the samples with minor oxygen modification that changes the charge-carrier density.^{18,19} While recent neutron-scattering experiments on superconducting $\text{Nd}_{1.85}\text{Ce}_{0.15}\text{CuO}_4$ (NCCO) show a drastic suppression of the static 3D AF order, how SC is established is unknown in part because the magnetic nature of the Nd^{3+} in NCCO complicates the interpretation of the data.²⁰ A systematic investigation on single crystals of PLCCO, where the Pr^{3+} has a nonmagnetic singlet ground state,^{16,17,21} should shed light on our understanding of the transition from the AF to SC.

We grew single crystals of PLCCO (mosaicity $<1^\circ$, cylindrical rods $\phi 5 \times \sim 20$ mm) using the traveling solvent floating zone technique.²² As grown, the PLCCO crystals are not superconducting and exhibit the noncollinear AF structure of undoped Pr_2CuO_4 [Fig. 1(a)] due to the weak pseudo-dipolar exchange interaction between Cu and Pr (Refs. 23–25). By annealing the samples at different temperatures in pure Ar, we obtain bulk superconducting PLCCO with different T_c 's and therefore control the transition from the AF state to the superconducting state (Ref. 18). Magnetic-susceptibility measurements on our PLCCO crystals show the onset of bulk SC, which corresponds to zero resistivity, at $T_c=24, 21,$ and 16 K, respectively [Fig. 1(c)].

We used elastic neutron scattering to probe AF order and label the momentum transfer $Q=(q_x, q_y, q_z)$ as $(H, K, L) = (q_x a/2\pi, q_y a/2\pi, q_z c/2\pi)$ in the reciprocal lattice units (rlu) appropriate for the tetragonal unit cell of PLCCO (space group $I4/mmm$, $a=3.98$, and $c=12.27$ Å). Experiments were performed in $[H, K, 0]$ and $[H, H, L]$ zones using triple-axis spectroscopy. In the latter geometry we also probed the 2D magnetic scattering using a two-axis energy-integration mode.²⁶

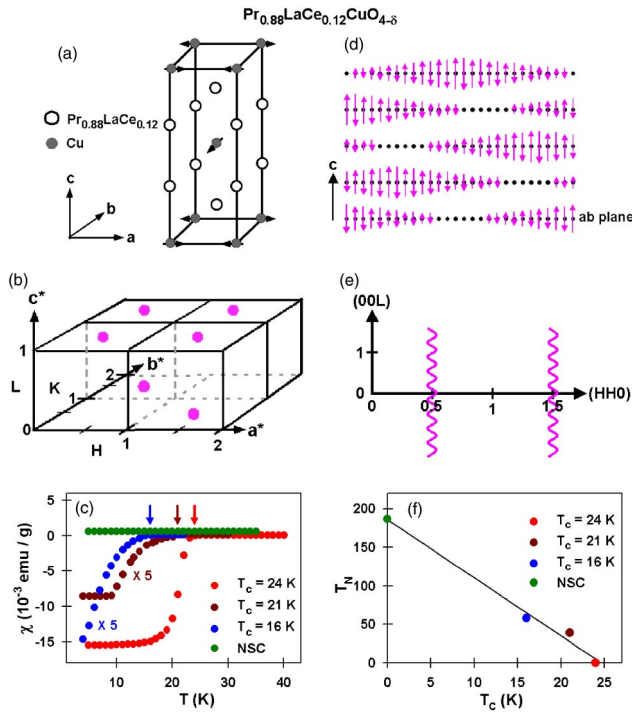


FIG. 1. Spin structure models, magnetic susceptibilities, and summary of results in the neutron experiments. For magnetic susceptibility, we used a 7-T SQUID magnetometer. Our neutron-scattering experiments were performed on HB-1, HB-1A, and HB-3 triple-axis spectrometers at the High-Flux Isotope Reactor (HFIR), Oak Ridge National Laboratory. Typical collimations were, proceeding from the reactor to the detector, $48'-40'-40'-120'$ (full width at half maximum), and the final neutron energy was fixed at $E_f=14.78$ meV with ~ 1 meV energy resolution. The monochromator, analyzer; and filters were all pyrolytic graphite. (a) The non-collinear type-I/III spin structure of the 3D AF order in PLCCO (Refs. 23 and 24). (b) Expected Bragg peaks from the AF structure of (a). (c) T dependence of the magnetic susceptibility for the four crystals investigated. The $T_c=24$ - and 21-K samples were obtained by annealing as-grown single crystals in pure Ar at 970 and 940 °C for 24 h, respectively. To obtain the $T_c=16$ -K crystal, the as-grown sample was annealed in pure Ar at 915 °C for 1 week. (d) Simplified model for AF order in the CuO_2 plane, but weakly correlated along the c axis. For clarity we plot the moment direction along the c axis, whereas the actual spin direction is in the a - b plane. (e) Expected scattering in reciprocal space from (d). (f) The relationship between T_c and T_N for samples investigated.

The magnetic Bragg peaks can be used to determine the size and temperature dependence of the ordered Cu (and Pr, if they are significant) moments, and Fig. 2 summarizes the experiments in the $[H,K,0]$ plane for the four samples investigated. Because of the c -axis spin-spin coupling, the magnetic scattering of the noncollinear AF structure [Fig. 1(a)] is nonzero at $(0.5,1.5,0)$, but vanishes at $(0.5,0.5,0)$ [Fig. 1(b)] (Refs. 23–25). The annealing process necessary for producing SC also induces a small ($\sim 0.9\%$) epitaxial secondary impurity phase, cubic $(\text{Nd,Ce})_2\text{O}_3$ in the case of $\text{Nd}_{2-x}\text{Ce}_x\text{CuO}_4$ (Refs. 19 and 27) and $(\text{Pr,L a,Ce})_2\text{O}_3$ for PLCCO (Ref. 28), which has Bragg peaks at $(0.5,0.5,0)$ and $(0.5,1.5,0)$. For the optimally doped ($T_c=24$ K) sample, nei-

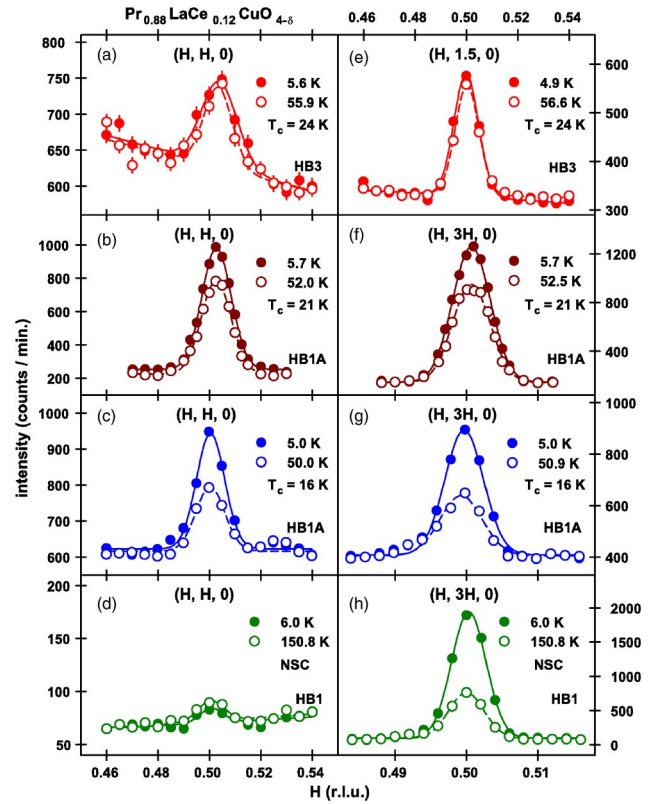


FIG. 2. Wave-vector scans in the $[H,K,0]$ zone around $(0.5,0.5,0)$ and $(0.5,1.5,0)$ at different temperatures for the four PLCCO samples. The in-plane and out-of-plane (vertical) resolutions of the spectrometers at $(0.5,0.5,0)$ are ~ 0.025 and ~ 0.1 – 0.2 \AA^{-1} , respectively. (a-d) Scans along the $[H,H,0]$ direction around $(0.5,0.5,0)$ at different temperatures. (e-h) Similar scans around $(0.5,1.5,0)$. Solid lines are Gaussian fits to the peaks on sloping backgrounds. The peaks are resolution limited and give a minimum coherence length of ~ 200 \AA .

ther peak shows any temperature dependence, suggesting no magnetic component. Scans along $(0.5,0.5,L)$ through $L=1,3$ where magnetic peaks are expected, also found no evidence of static AF order [Figs. 3(a) and 3(b)], consistent with the phase diagram of $\text{Pr}_{1-x}\text{LaCe}_x\text{CuO}_{4-\delta}$ (Ref. 18).

For the underdoped superconducting samples, on cooling to 5 K the intensity increases for the $(0.5,1.5,0)$ magnetic Bragg peak [Figs. 2(f) and 2(g)] as expected. Assuming that the residual AF structure in superconducting PLCCO is identical to that of the as-grown PLCCO, we estimate that the effective static low-temperature Cu ordered moment (M_{Cu}) reduces from 0.06 to 0.04 μ_B (with no appreciable induced Pr moment) when T_c increases from 16 to 21 K [Figs. 3(d) and 3(f)]. Concomitantly, T_N of the system decreases from 60 K [Fig. 3(f)] to 40 K [Fig. 3(d)]. For comparison, we cut a piece off the $T_c=16$ -K crystal and quenched SC by annealing it in air at 900 °C for 24 h [Fig. 1(c)]. In this NSC PLCCO sample, the T_N becomes 186 K and M_{Cu} is 0.12 μ_B at 97 K [Fig. 3(g)].

In the underdoped samples, we also observed magnetic intensity at the $(0.5,0.5,0)$ peak [Figs. 2(b) and 2(c)]. The temperature dependence of the scattering at $(0.5,0.5,0)$ and $(0.5,1.5,0)$ for the $T_c=21$ - and 16-K crystals is quite similar

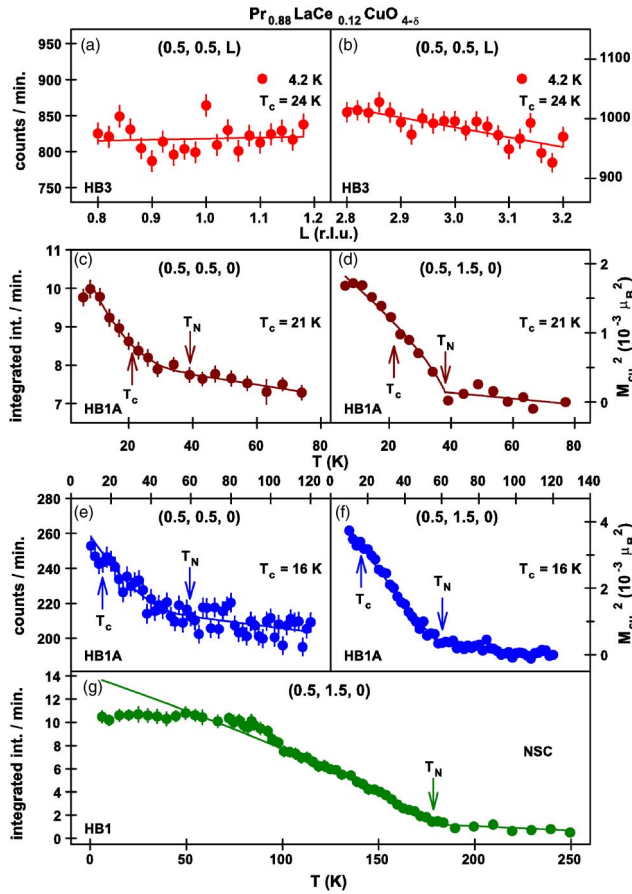


FIG. 3. (a,b) Wave-vector scans through $(0.5, 0.5, 1)$ and $(0.5, 0.5, 3)$ for $T_c=24$ K PLCCO at 4.2 K. T dependence of the scattering at $(0.5, 0.5, 0)$ and $(0.5, 1.5, 0)$ in the $[H, K, 0]$ zone for (c,d) $T_c=21$ K, (e,f) $T_c=16$ K, and (g) NSC PLCCO samples used to extract their T_N 's. The solid lines in (d,f) and (g) are power-law fits describing the contribution of Cu spins (Ref. 23). The ordered Cu moments (M_{Cu}) in (d,f) are estimated by normalizing the magnetic intensity to the weak $(1, 1, 0)$ nuclear Bragg peaks without considering absorption or extinction effects. At 97 K, $M_{Cu}=0.12 \mu_B$ in (g). No measurable Pr moment was induced in (d,f) while an induced Pr moment is clearly evident below 95 K in (g) (see Ref. 24).

[Figs. 3(c)–3(f)], and thus suggests at first glance that they are both related to the onset of 3D AF order. However, equivalent scans in the $[H, H, L]$ scattering plane indicate that the two types of peaks must have a different origin. In particular, while both $(0.5, 0.5, 0)$ and $(0.5, 1.5, 0)$ peaks are resolution limited in the CuO_2 plane [Figs. 2(b), 2(c), 2(f), and 2(g)], a scan along the L direction of $(0.5, 0.5, 0)$ in the $[H, H, L]$ zone at 6 K shows only slightly enhanced “background” scattering compared to the 80-K data on the $T_c=16$ -K crystal [Fig. 4(a)]. Scans along the $(0.5, 0.5, L)$ direction find no appreciable differences between 6 and 60 K except at $(0.5, 0.5, 1)$, where the long-range 3D AF order develops [Figs. 4(b) and 4(c)]. Measurements at the $(0.5, 0.5, 1)$, $(0.5, 0.5, 2)$, $(0.5, 0.5, 3)$, and $(0.5, 0.5, 5)$ positions (not shown), on the other hand, confirm quantitatively that the magnetic structure is indeed that of Fig. 1(a) with a $T_N \approx 60$ K and negligible induced Pr moment.

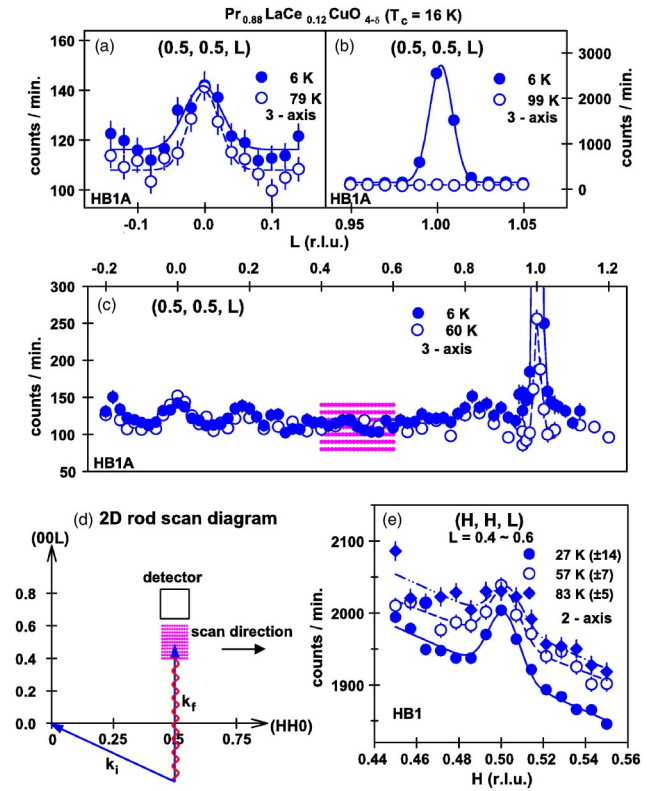


FIG. 4. Measurements in the $[H, H, L]$ zone on the $T_c=16$ -K crystal. Scans along the L direction around (a) $(0.5, 0.5, 0)$ and (b) $(0.5, 0.5, 1)$ at different temperatures. The magnetic peak in (b) is resolution limited and gives a c axis (ξ_c) AF coherence length of 790 \AA . (c) Scans along the $[0.5, 0.5, L]$ direction at 6 K (closed circles) and 60 K (open circles). The peaks at $L \sim 0.2, 0.8$ are non-magnetic. The magenta areas in (c) and (d) mark the L region probed in the two-axis integration mode. (d) The experimental geometry in the two-axis mode in the $[H, H, L]$ zone (Ref. 26). (e) Scans along the $[H, H]$ direction around $(0.5, 0.5)$ with a range of L at different temperatures.

The scattering associated with the $(0.5, 0.5, 0)$ peak therefore appears to be diffuse and quasi-2D along the c -axis direction.²⁹ To test if this is indeed the case, we carried out measurements in the two-axis energy-integrated mode by aligning the outgoing wave vector k_f parallel to the 2D rod direction.²⁶ In this geometry, we measure the wave-vector-dependent susceptibility that includes the elastic scattering around $(0.5, 0.5, L)$ with $0.4 < L < 0.6$ [see magenta areas in Figs. 4(c) and 4(d)]. The data reveal an unambiguous peak centered at $(0.5, 0.5, L)$ whose intensity decreases with increasing temperature [Fig. 4(e)]. This, and measurements around $(0.5, 0.5, 0)$ (Fig. 2), suggest that $(0.5, 0.5, L)$ is weakly L dependent. Therefore, underdoped PLCCO has a quasi-2D SDW modulation [Figs. 1(d) and 1(e)] that coexists with SC and 3D AF order.

The relation between T_c 's and T_N 's of our PLCCO crystals is summarized in Fig. 1(f), which demonstrates that T_N decreases linearly with increasing T_c and vanishes when optimal superconductivity is established. While these results suggest the presence of a QPT to SC when T_N approaches zero, how do we understand the coexistence of quasi-2D SDW and

long-range 3D AF order and their relationship to SC?

One concern might be that the annealing process necessary for producing SC in PLCCO causes macroscopic oxygen inhomogeneities, giving rise to separate nonmagnetic SC and AF NSC phases. In this scenario one would expect to observe both a sharp T_c and a robust T_N weakly depending on oxygen doping as in the case of oxygen-doped La_2CuO_4 (Ref. 30). However, the drastic decrease in 3D transition to long-range magnetic order (T_N decreasing from 186 K for NSC to 40 K for the $T_c=21$ -K sample) combined with relatively sharp magnetic transition [as in hole-doped La_2CuO_4 (Ref. 31)] demonstrates that this is not the case. The relatively broad SC transition is then a consequence of an intrinsic electronic inhomogeneity occurring on a length scale longer than the (short) SC coherence length. While neutrons are a bulk probe and cannot determine how SC coexists with the AF phases at zero field, magnetic-field experiments show

that the AF phase competes directly with SC (Refs. 32 and 33). In this case, the observed commensurate quasi-2D SDW in PLCCO is analogous to the incommensurate 2D SDW in hole-doped $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$ and $\text{La}_2\text{CuO}_{4+\delta}$ (Refs. 3 and 4). In the electron-doped case, the commensurate SDW may arise from the in-phase domains of “stripes,”²² as compared to the antiphase domains of “stripes” in hole-doped materials.² While a detailed picture of PLCCO may be obtained from future spin dynamical measurements, it appears that microscopic heterogeneity and the resulting competition of different order parameters are general features of doped copper oxides near a QPT [Fig. 1(f)].

We would like to thank Y. S. Lee and K. Yamada for helpful discussions. This work was supported by the US NSF-DMR-0139882 and DOE under Contract No. DE-AC-00OR22725 with UT/Battelle, LLC.

*E-mail address: daip@ornl.gov

- ¹M. A. Kastner *et al.*, Rev. Mod. Phys. **70**, 897 (1998).
- ²S. A. Kivelson *et al.*, Rev. Mod. Phys. **75**, 1201 (2003).
- ³H. Kimura *et al.*, Phys. Rev. B **59**, 6517 (1999).
- ⁴Y. S. Lee *et al.*, Phys. Rev. B **60**, 3643 (1999).
- ⁵S. Katano, M. Sato *et al.*, Phys. Rev. B **62**, R14 677 (2000).
- ⁶B. Lake *et al.*, Nature (London) **415**, 299 (2002).
- ⁷S. Sachdev, Rev. Mod. Phys. **75**, 913 (2003).
- ⁸J. L. Tallon and J. W. Loram, Physica C **349**, 53 (2001).
- ⁹F. F. Balakirev *et al.*, Nature (London) **424**, 912 (2003).
- ¹⁰W. Bao *et al.*, Phys. Rev. Lett. **91**, 127005 (2003).
- ¹¹Y. Ando *et al.*, Phys. Rev. Lett. **92**, 247004 (2004).
- ¹²Y. Dagan *et al.*, Phys. Rev. Lett. **92**, 167001 (2004).
- ¹³S. Sachdev and S. C. Zhang, Science **295**, 452 (2002).
- ¹⁴G. Alvarez *et al.*, Phys. Rev. B **71**, 014514 (2005).
- ¹⁵A. T. Savici *et al.*, Phys. Rev. B **66**, 014524 (2002).
- ¹⁶J. A. Sanjurjo *et al.*, Phys. Rev. B **51**, 1185 (1995).
- ¹⁷W. Henggeler *et al.*, Europhys. Lett. **29**, 233 (1995).
- ¹⁸M. Fujita *et al.*, Phys. Rev. B **67**, 014514 (2003).
- ¹⁹K. Kurahashi *et al.*, J. Phys. Soc. Jpn. **71**, 910 (2002).
- ²⁰K. Yamada *et al.*, Phys. Rev. Lett. **90**, 137004 (2003).
- ²¹A. T. Boothroyd *et al.*, Phys. Rev. B **45**, 10 075 (1992).
- ²²X. F. Sun *et al.*, Phys. Rev. Lett. **92**, 047001 (2004).
- ²³I. W. Sumarlin *et al.*, Phys. Rev. B **51**, 5824 (1995).
- ²⁴A. N. Lavrov *et al.*, Phys. Rev. Lett. **92**, 227003 (2004).
- ²⁵D. Petitgrand *et al.*, Phys. Rev. B **59**, 1079 (1999).
- ²⁶R. J. Birgeneau *et al.*, Phys. Rev. B **3**, 1736 (1971).
- ²⁷P. K. Mang *et al.*, Nature (London) **426**, 139 (2003).
- ²⁸H. J. Kang *et al.*, Nature (London) **426**, 140 (2003).
- ²⁹The difference between the $[H, K, 0]$ measurements and the $[H, H, L]$ measurements is then due to the coarse vertical instrumental resolution (0.2 \AA^{-1}), which in the $[H, K, 0]$ geometry integrates over $-0.2 < L < 0.2$, rendering the scattering observable.
- ³⁰A. N. Lavrov *et al.*, Phys. Rev. Lett. **87**, 017007 (2001).
- ³¹O. P. Vajk *et al.*, Science **295**, 1691 (2002).
- ³²H. J. Kang *et al.*, Nature (London) **423**, 522 (2003). Our experiments performed on PLCCO, where the impurity phase $(\text{Pr, La, Ce})_2\text{O}_3$ has no field dependence, find anisotropic field-induced effects at $(0.5, 0.5, 0)$ for both $T_c=16$ - and 21 -K samples and no observable field-induced effect for the $T_c=24$ -K crystal (H. J. Kang *et al.*, unpublished).
- ³³M. Fujita *et al.*, Phys. Rev. Lett. **93**, 147003 (2004).