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## Hidden Fermi-liquid Charge Transport in the Antiferromagnetic Phase of the Electron-Doped Cuprate Superconductors

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Systematic analysis of the planar resistivity, Hall effect, and cotangent of the Hall angle for the electron-doped cuprates reveals underlying Fermi-liquid behavior even deep in the antiferromagnetic part of the phase diagram. The transport scattering rate exhibits a quadratic temperature dependence, and is nearly independent of doping and compound and carrier type (electrons versus holes), and hence is universal. Our analysis moreover indicates that the material-specific resistivity upturn at low temperatures and low doping has the same origin in both electron- and hole-doped cuprates.

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The cuprates feature a complex phase diagram that is asymmetric upon electron-versus-hole doping [1] and plagued by compound-specific features associated with different types of disorder and crystal structures [2], often rendering it difficult to discern universal properties. What is known for certain is that the parent compounds are antiferromagnetic (AF) insulators, that AF correlations are more robust against doping with electrons than with holes [3,4], and that pseudogap (PG) phenomena, seemingly unusual charge transport behavior, and *d*-wave superconductivity appear upon doping the quintessential CuO<sub>2</sub> planes [1]. The nature of the metallic state that emerges upon doping the insulating parent compounds has remained a central open question. Moreover, below a compound specific doping level, the low-temperature resistivity for both types of cuprates develops a logarithmic upturn that appears to be related to disorder, yet whose microscopic origin has remained unknown [1,5–7]. In contrast, at high dopant concentrations, the cuprates are good metals with well-defined Fermi surfaces and clear evidence for Fermi-liquid (FL) behavior [8–14].

In a new development, the hole-doped cuprates were found to exhibit FL properties in an extended temperature range below the characteristic temperature  $T^{**}$  ( $T^{**} < T^*$ ;  $T^*$  is the PG temperature): (i) the resistivity per CuO<sub>2</sub> sheet exhibits a universal, quadratic temperature dependence, and is inversely proportional to the doped carrier density  $p$ ,  $\rho \propto T^2/p$  [15]; (ii) Kohler’s rule for the magnetoresistivity, the characteristic of a conventional metal with a single relaxation rate, is obeyed, with a FL scattering rate  $1/\tau \propto T^2$  [16]; (iii) the optical scattering rate exhibits the quadratic frequency dependence and the temperature-frequency scaling expected for a Fermi liquid [17]. In this part of the phase diagram, the Hall coefficient is known to be approximately independent of temperature and to take on a value that corresponds to  $p$ ,  $R_H \propto 1/p$  [18].

In order to explore the possible connection among the different regions of the phase diagram, an important quantity to consider is the cotangent of the Hall angle,  $\cot(\theta_H) = \rho/(HR_H)$ . For simple metals, this quantity is proportional to the transport scattering rate,  $\cot(\theta_H) \propto m^*/\tau$  ( $H$  is the magnetic field, and  $m^*$  is the effective mass). It has long been known that  $\cot(\theta_H) \propto T^2$  in the “strange-metal” regime ( $T > T^*$ ) of the hole-doped cuprates [19], where  $\rho \propto T$  [1], which has been interpreted as the result of distinct longitudinal and transversal scattering rates [20] or due to an anisotropic scattering rate [21,22]. Remarkably, for the model compound HgBa<sub>2</sub>CuO<sub>4+ $\delta$</sub>  (Hg1201) it was recently found that  $\cot(\theta_H) \propto T^2$  is independent of doping and does not exhibit a noticeable change upon crossing the characteristic temperatures  $T^*$  and  $T^{**}$ , thus providing a direct link between the strange-metal and PG-FL regimes [23]. Upon combining this result with those for other hole-doped cuprates, it was furthermore shown that the transport scattering rate is approximately compound independent, and hence that the scattering mechanism characteristic of the FL at high doping levels ( $p \approx 0.3$ ) prevails even at  $p = 0.01$ , very close to the Mott-insulating state [23].

Here, we consider the electron-doped half of the phase diagram. Unlike for the hole-doped cuprates [23], we find it necessary to explicitly consider the low- $T$  logarithmic resistivity upturn,  $\Delta\rho(T) \propto -\log(T)$ . The magnitude of this upturn is nonuniversal, can vary from sample to sample for the same compound and doping level, and is particularly large deep in the AF state of the electron-doped compounds. This analysis reveals underlying FL behavior in the AF state. Moreover, the scattering rate is nearly the same as for the hole-doped cuprates. These new insights allow us to extend the prior analysis of hole-doped compounds to lower temperature and to demonstrate that the resistivity upturn must have the same physical origin in both electron- and hole-doped cuprates.

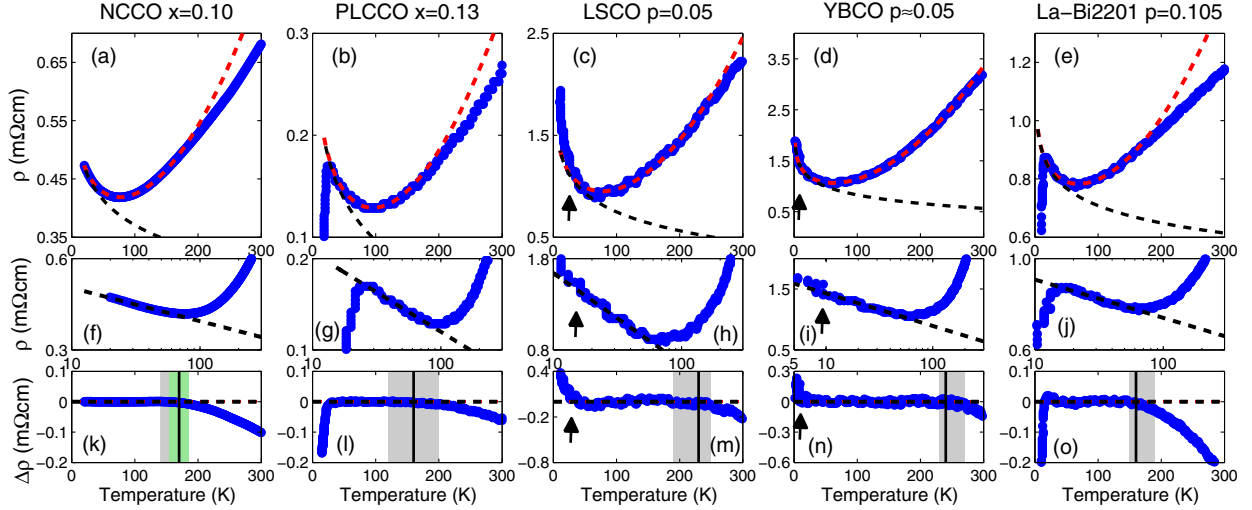


FIG. 1. *ab*-plane resistivity of various cuprate materials. (a)–(e) Raw data (blue circles) and fit to Eq. (2b) (red dashed curves). The estimated contributions  $A_0 - A_{\log} \log(T/1K)$  are shown as black dashed lines. (f)–(j) Semilog plots of the resistivity. The dashed lines indicate the logarithmic contribution. (k)–(o) Differences between the raw data and fits. The horizontal black dashed lines indicate zero difference and are guides to the eye. The gray shaded bands indicate the temperature range in which the underlying quadratic temperature dependence of the planar resistivity breaks down [38,41]. The Néel temperature ( $T_N \approx 165$  K) of NCCO ( $x = 0.10$ ) is shown as a green shaded band [4]. The arrows indicate low-temperature deviations from logarithmic behavior in lightly doped LSCO and YBCO. Except for the new NCCO data, which are consistent with prior work [26], the data are adapted from Refs. [36,38,40].

It is instructive to recall the systematic study of initially very clean, hole-doped  $\text{YBa}_2\text{Cu}_3\text{O}_{6+\delta}$  (YBCO) samples, with intrinsic resistivity  $\rho_i(T) \propto T^2$  in the PG regime [15], that were subsequently exposed to electron-beam irradiation [7]. Upon increasing the radiation dose, and hence the density of point defects, the resistivity was found to be enhanced by a  $T$ -independent contribution ( $\rho_0$ ) and a low- $T$  upturn [ $\Delta\rho(T)$ ]. This suggests that the resistivity can be decomposed into three terms:

$$\rho = \rho_0 + \Delta\rho(T) + \rho_i(T). \quad (1)$$

Except at very low doping levels and temperatures, the nonuniversal upturn is known to exhibit a logarithmic temperature dependence [6,7,24,25].

Starting with new data for a sample of the archetypal electron-doped cuprate  $\text{Nd}_{2-x}\text{Ce}_x\text{CuO}_{4+\delta}$  (NCCO) that exhibits robust AF order ( $x = 0.10$ ; superconductivity in NCCO appears for  $x \approx 0.13$  [4]), we follow the evolution of these three contributions as a function of temperature and doping for a large number of compounds: electron-doped NCCO [26–30],  $\text{La}_{2-x}\text{Y}_x\text{CuO}_{4+\delta}$  (LYCO) [31],  $\text{Pr}_{2-x}\text{Ce}_x\text{CuO}_{4+\delta}$  (PCCO) [32–35],  $\text{Pr}_{1.3-x}\text{La}_{0.7}\text{Ce}_x\text{CuO}_{4+\delta}$  (PLCCO) [36], and  $\text{La}_{2-x}\text{Ce}_x\text{CuO}_{4+\delta}$  (LCCO) [37], and hole-doped  $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$  (LSCO) [38], YBCO [7,38,39] and  $\text{Bi}_2\text{Sr}_{2-x}\text{La}_x\text{Cu}_2\text{O}_{8+\delta}$  (La-Bi2201) [24,40]. Representative resistivity data are shown in Fig. 1 (for a summary of sample characteristics, see Ref. [41]). In all cases, the logarithmic contribution is apparent.

Equation (1) can be written in two identical forms:

$$\rho = \rho_{\text{res}} - A_{\log} \log(T/T_{\log}) + A_2 T^2, \quad (2a)$$

$$\rho = A_0 - A_{\log} \log(T/1K) + A_2 T^2, \quad (2b)$$

where  $\rho_{\text{res}}$  is the residual ( $T = 0$ ) resistivity and  $A_0 = \rho_{\text{res}} + A_{\log} \log(T_{\log}/1K)$ . We fit the data (Fig. 1) to the second form, as it contains three rather than four parameters. This procedure resembles that in Ref. [7], with  $A_0 = \rho_0$ , except that we allow all three parameters to vary; i.e., we do not use the high- $T$  data to fix  $A_2$  and  $\rho_0$ ; the difference between the two approaches, which lead to very similar conclusions, is further analyzed in Ref. [41].

For the hole-doped cuprates, it was demonstrated that the sheet resistance coefficient  $A_{2\Box}$  is universal [15]. Thus, to compare  $A_{\log}$  and  $A_2$  for different compounds, we convert to sheet resistance units. Figure 2(a) shows that  $A_{2\Box}$  for the electron-doped materials is approximately inversely proportional to the cerium concentration, and hence to the nominal electron concentration. For comparison, Fig. 2(b) shows  $A_{2\Box} \propto 1/p$  obtained from fits to Eq. (2b) for the hole-doped cuprates. Remarkably, the absolute values of  $A_{2\Box}$  for hole- and electron-doped materials at the same nominal doping level are very similar.

In contrast, at the same nominal doping level,  $A_{\log\Box}$  and  $A_{0\Box}$  are nearly an order of magnitude larger for the electron-doped compounds; similar to  $A_{2\Box}$ , these coefficients exhibit power-law doping dependences [41]:

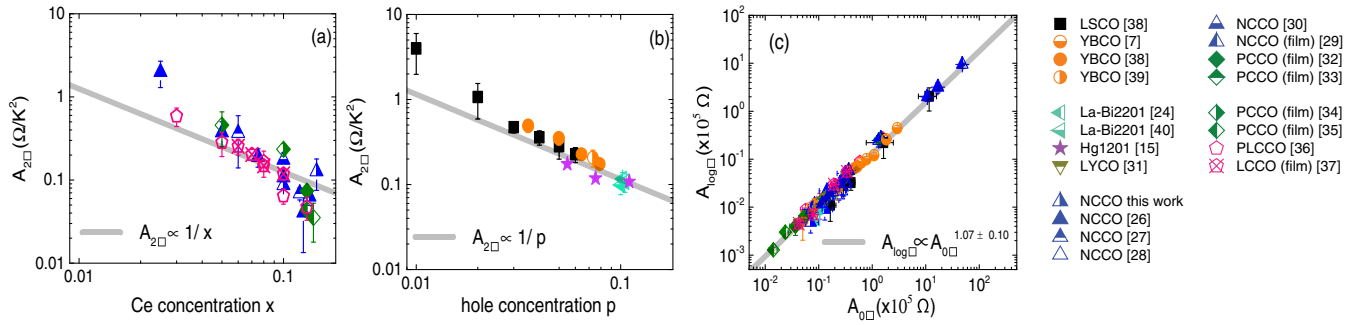


FIG. 2. Doping dependence of  $A_{2\Omega}$  for (a) electron- and (b) hole-doped materials, as obtained from fits to Eq. (2b). Note that in Ref. [15]  $A_{2\Omega}$  was estimated for the hole-doped cuprates without considering the logarithmic contribution; see also Ref. [41]. The gray lines are guides to the eye and indicate  $1/x$  and  $1/p$  dependences, respectively. (c) Scaling relation between  $A_{\log\Omega}$  and  $A_{0\Omega}$ . Data are adapted from Refs. [7,15,24,26–40]. The error bars are 1 standard deviation, estimated from fits with various temperature ranges. At low doping, the low-temperature upturn can no longer be well described by a logarithmic contribution (see Fig. 1 and Ref. [41]), and the obtained coefficients thus depend on the fit temperature range and have large error bars.

$A_{\log\Omega} \propto x^{-3.6 \pm 0.3}$  and  $A_{0\Omega} \propto x^{-3.4 \pm 0.3}$  (electron doping) and  $A_{\log\Omega} \propto p^{-3.0 \pm 0.2}$  and  $A_{0\Omega} \propto p^{-2.7 \pm 0.3}$  (hole doping). Within error, the exponents are the same on each side of the phase diagram. Figure 2(c), which treats the doping level as an implicit parameter, highlights this point by revealing an approximately linear relationship between  $A_{0\Omega}$  and  $A_{\log\Omega}$ , which holds over many orders of magnitude. This observation carries several important messages. First, it implies that the dominant contribution to  $A_0$  is unrelated to residual impurity scattering [recall that  $A_0 = \rho_{\text{res}} + A_{\log} \log(T_{\log}/1\text{K})$ ]. Second,  $T_{\log}$  should not vary considerably. Indeed,  $T_{\log}$  obtained from fits to Eq. (2a) is on the order of 100 K and exhibits a similar monotonic doping dependence for all materials [41]. Finally, it points to a universal mechanism responsible

for the resistivity upturns in both electron- and hole-doped cuprates, which seems to be related to disorder [7] rather than a reconstruction of the Fermi surface [43].

Motivated by these insights and by the recent finding of a universal scattering rate throughout the phase diagram of the hole-doped cuprates [23] [ $\cot(\theta_H) = C_0 + C_2 T^2$  holds with the universal value of  $C_2 = 0.0175(20)$  and with a compound, doping, disorder dependent  $C_0$ ], we take a closer look at prior comprehensive data for the AF phase of NCCO [44]. Figure 3 shows the procedure to disentangle the resistivity contributions and to obtain  $\cot(\theta_H)$  for NCCO ( $x = 0.075$  and  $0.10$ ). The underlying FL scattering rate is only revealed upon considering the logarithmic upturn. [The same procedure was applied for  $x = 0.05$  and  $0.125$  [26,44]; see Ref. [41]]. Once we subtract the nonuniversal contribution, we find

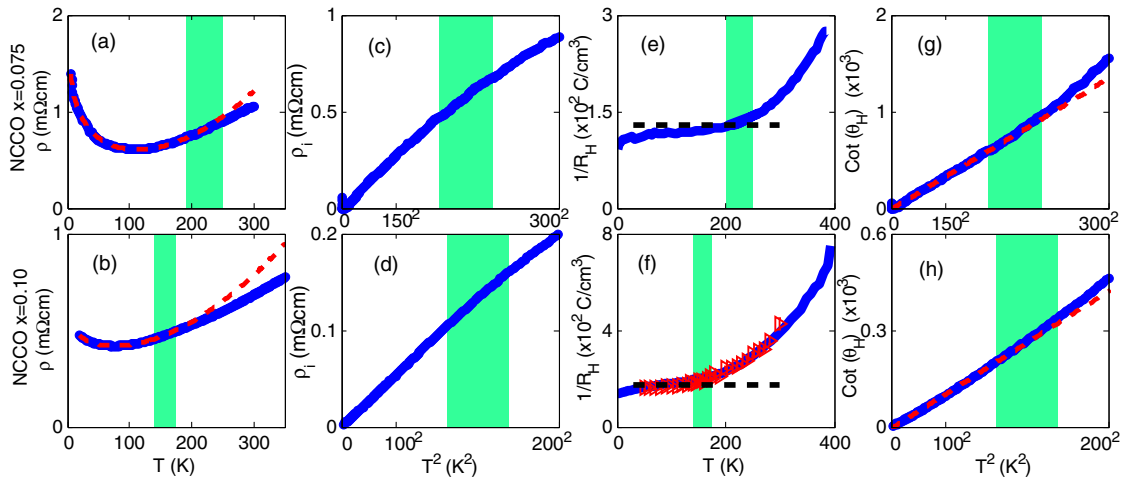


FIG. 3. Resistivity data and fit for NCCO with (a)  $x = 0.075$  [26] and (b)  $x = 0.10$  [present work, same data as in Fig. 1(a)]. (c),(d) Resistivity after subtraction of fitted logarithmic contribution. (e),(f)  $1/R_H$  adapted from Ref. [44]; the  $1/R_H$  data obtained for our NCCO ( $x = 0.10$ ) sample (red triangles) agree with the prior work [44]. The black horizontal dashed lines indicate the magnitude of  $1/R_H$  estimated from the chemical dopant concentrations  $x = 0.075$  and  $0.10$ . (d),(g) Cotangent of the Hall angle. The red dashed lines indicate  $\cot(\theta_H) \propto C_2 T^2$ . As in Fig. 1(k), the green bands indicate the estimated Néel temperatures [4].

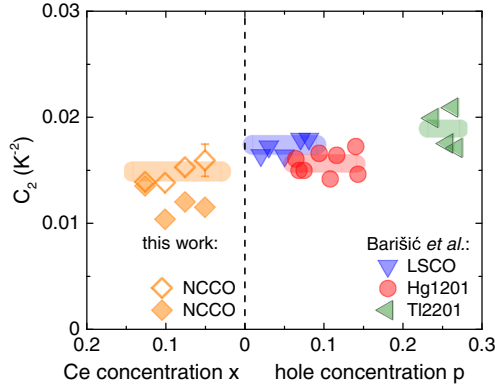


FIG. 4.  $C_2$  for various single-layer cuprates. LSCO, Hg1201, and Tl2201 results adapted from Ref. [23].  $C_2$  for NCCO is determined both by method 1 (open symbols) and method 2 (solid symbols). See also Ref. [41].

$$\cot(\theta_H) = \rho_i / (HR_H) = C_2 T^2, \quad (3)$$

where  $\rho_i = \rho - (A_0 - A_{\log} \log(T/1K))$ .

Above the Néel temperature,  $\cot(\theta_H)$  deviates from this simple quadratic behavior. This is consistent with Fig. 1(k) and appears to be the result of Fermi-surface reconstruction [26]: upon increasing the temperature above  $T_N$ , the Fermi surface evolves from simple electron pockets to a more complex shape, and the Hall coefficient ceases to be a good measure of the carrier density [23,44]. In order to address the properties of the high- $T$  regime, a more elaborate analysis is required, which is beyond the scope of the present Letter.

As seen from Fig. 4, our analysis (“method 1”) of the dc resistivity and Hall coefficient for NCCO yields values of  $C_2$  that are nearly identical to those found previously for the hole-doped cuprates. This suggests that the umklapp process is the scattering mechanism in both cases. We test the robustness of this result with regard to the fit procedure by determining the slope of the quadratic term ( $A_2$ ) simply from the high- $T$  part of the resistivity, as suggested in Refs. [7,23], neglecting the low- $T$  upturns (“method 2”). In essence, method 2 yields a lower bound for  $A_2$ , since the logarithmic contribution is neglected [41]. As summarized in Fig. 4, we find that  $C_2$  changes no more than 30% for NCCO in the studied doping range, confirming the robustness of our analysis.

Hole-doped LSCO exhibits a considerable resistivity upturn at moderate and low doping. We have analyzed LSCO data ( $p = 0.05, 0.07, 0.08$ ) [38] with method 1 and find hardly any change in  $C_2$  compared to the prior result based on method 2 [23]. However, as seen from Fig. 1(c), for  $p = 0.05$  at the lowest temperatures, the resistivity upturn is stronger than logarithmic. Method 1 no longer gives an accurate description for  $p \leq 0.03$  [41].

The relatively small difference (about a factor of 2) in the value of  $C_2$  between electron- and hole-doped cuprates can

be attributed to a difference in the effective mass. We are not aware of reliable measurements of  $m^*$  for the electron-doped cuprates in the relevant doping range, but band-structure calculations indicate a smaller value than for the hole-doped compounds [45].

In principle, there are two distinct ways to understand the simultaneous FL and logarithmic transport behaviors captured by Eq. (2). The first is via Matthiessen’s rule, which assumes that scattering rates for different scattering processes simply add up ( $1/\tau_{\text{total}} = 1/\tau_1 + 1/\tau_2 + \dots$ ). The second possibility is that Eq. (2) describes a serial-resistor network, which would imply the existence of metallic and nonmetallic regions [46,47]. We can distinguish between these two possibilities by considering a recent result obtained for the hole-doped cuprates [16], namely, that Kohler scaling for the magnetoresistance holds for compounds or samples that exhibit negligible residual resistivity ( $\rho_{\text{res}} \approx 0$ ) and FL behavior [ $(\rho - \rho_{H=0})/\rho_{H=0} \propto H^2/\rho_{H=0}^2$ , where  $\rho_{H=0}$  is the zero field resistivity] below  $T^{**}$ . This scaling follows directly from the Boltzmann transport equation and unmistakably demonstrates the FL character of the pseudogap phase. For LSCO, which exhibits a large resistivity, it was found that Kohler’s rule is obeyed only if  $\rho_{H=0}$  is replaced by  $\rho_{H=0} - A_0$ . This result is incompatible with Matthiessen’s rule for a homogeneous system. However, it is compatible with a serial-resistor network [47] in which only metallic patches contribute to the magnetoresistivity and nonmetallic regions characterized by logarithmic behavior have negligible influence. This conclusion also provides an explanation for the heuristic logarithmic term and for our finding (from the approximately linear scaling between  $A_0$  and  $A_{\log}$ ) that  $\rho_{\text{res}}$  is negligible. Such upturns naturally appear at sufficiently large temperatures in systems with strongly coupled metallic grains separated by an insulating matrix [48].

We have established that the planar charge transport of the electron-doped cuprates exhibits hidden FL behavior in the AF phase (for an early discussion on FL behavior outside of the AF phase, see Ref. [49]). This fact had previously gone unnoticed because the FL transport is masked by a particularly large nonmetallic contribution to the resistivity. The FL behavior is remarkably robust and universal. The sheet resistance coefficients  $A_{2\Box}$  are very similar for electron and hole-doped compounds at the same nominal doping level. Moreover, by considering the Hall effect, we demonstrate that the scattering rate is nearly the same on both sides of the phase diagram. We also find that the nonuniversal additive logarithmic contribution is characterized by a temperature scale that has the same magnitude for electron- and hole-doped compounds and exhibits a weak doping dependence. This fact is exemplified by the approximate (linear) scaling between  $A_{\log\Box}$  and  $A_{0\Box}$  that is found to hold over 4 orders of magnitude. We therefore conclude that the nonmetallic contribution must have the same physical origin in all compounds. Most likely, it is

associated with charge transport involving metallic and (nonuniversal) nonmetallic regions of the material. The metallic regions continue to follow simple FL behavior to the lowest temperatures and doping levels studied here. Overall, these insights shed new light on the physics of the quintessential copper-oxygen planes and they demonstrate that important aspects of the charge transport of electron- and hole-doped cuprates are universal.

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- [1] N. P. Armitage, P. Fournier, and R. L. Greene, *Rev. Mod. Phys.* **82**, 2421 (2010).
- [2] H. Eisaki, N. Kaneko, D. L. Feng, A. Damascelli, P. K. Mang, K. M. Shen, Z. X. Shen, and M. Greven, *Phys. Rev. B* **69**, 064512 (2004).
- [3] B. Keimer, N. Belk, R. J. Birgeneau, A. Cassanho, C. Y. Chen, M. Greven, M. A. Kastner, A. Aharony, Y. Endoh, R. W. Erwin, and G. Shirane, *Phys. Rev. B* **46**, 14034 (1992).
- [4] E. M. Motoyama, G. Yu, I. M. Vishik, O. P. Vajk, P. K. Mang, and M. Greven, *Nature (London)* **445**, 186 (2007).
- [5] V. Radhakrishnan, C. K. Subramaniam, V. Sankaranarayanan, G. V. Subba Rao, and Srinivasan, *Physica (Amsterdam)* **167C**, 53 (1990).
- [6] Y. Ando, G. S. Boebinger, A. Passner, T. Kimura, and K. Kishio, *Phys. Rev. Lett.* **75**, 4662 (1995).
- [7] F. Rullier-Albenque, H. Alloul, F. Balakirev, and C. Proust, *Europhys. Lett.* **81**, 37008 (2008).
- [8] T. Yoshida, X. J. Zhou, M. Nakamura, S. A. Kellar, P. V. Bogdanov, E. D. Lu, A. Lanzara, Z. Hussain, A. Ino, T. Mizokawa, A. Fujimori, H. Eisaki, C. Kim, Z.-X. Shen, T. Kakeshita, and S. Uchida, *Phys. Rev. B* **63**, 220501(R) (2001).
- [9] S. Nakamae, K. Behnia, N. Mangkorntong, M. Nohara, H. Takagi, S. J. C. Yates, and N. E. Hussey, *Phys. Rev. B* **68**, 100502(R) (2003).
- [10] N. E. Hussey, M. Abdel-Jawad, A. Carrington, A. P. Mackenzie, and L. Balicas, *Nature (London)* **425**, 814 (2003).
- [11] B. Vignolle, A. Carrington, R. A. Cooper, M. M. J. French, A. P. Mackenzie, C. Jaudet, D. Vignolles, C. Proust, and N. E. Hussey, *Nature (London)* **455**, 952 (2008).
- [12] S. Nakamae, K. Behnia, N. Mangkorntong, M. Nohara, H. Takagi, S. J. C. Yates, and N. E. Hussey, *Phys. Rev. B* **79**, 219904(E) (2009).
- [13] T. Helm, M. V. Kartsovnik, M. Bartkowiak, N. Bittner, M. Lambacher, A. Erb, J. Wosnitza, and R. Gross, *Phys. Rev. Lett.* **103**, 157002 (2009).
- [14] T. Helm, M. V. Kartsovnik, I. Sheikin, M. Bartkowiak, F. Wolff-Fabris, N. Bittner, W. Biberacher, M. Lambacher, A. Erb, J. Wosnitza, and R. Gross, *Phys. Rev. Lett.* **105**, 247002 (2010).
- [15] N. Barišić, M. K. Chan, Y. Li, G. Yu, X. Zhao, M. Dressel, A. Smontara, and M. Greven, *Proc. Natl. Acad. Sci. U.S.A.* **110**, 12235 (2013).
- [16] M. K. Chan, M. J. Veit, C. J. Dorow, Y. Ge, Y. Li, W. Tabis, Y. Tang, X. Zhao, N. Barišić, and M. Greven, *Phys. Rev. Lett.* **113**, 177005 (2014).
- [17] S. I. Mirzaei, D. Stricker, J. N. Hancock, C. Berthod, A. Georges, E. van Heumen, M. K. Chan, X. Zhao, Y. Li, M. Greven, N. Barišić, and D. van der Marel, *Proc. Natl. Acad. Sci. U.S.A.* **110**, 5774 (2013).
- [18] Y. Ando, Y. Kurita, S. Komiya, S. Ono, and K. Segawa, *Phys. Rev. Lett.* **92**, 197001 (2004).
- [19] T. R. Chien, Z. Z. Wang, and N. P. Ong, *Phys. Rev. Lett.* **67**, 2088 (1991).
- [20] P. W. Anderson, *Phys. Rev. Lett.* **67**, 2092 (1991).
- [21] N. E. Hussey, *J. Phys. Condens. Matter* **20**, 123201 (2008).
- [22] J. M. Buhmann, M. Ossadnik, T. M. Rice, and M. Sigrist, *Phys. Rev. B* **87**, 035129 (2013).
- [23] N. Barišić, M. K. Chan, M. J. Veit, C. Dorow, Y. Ge, Y. Tang, W. Tabis, G. Yu, X. Zhao, and M. Greven, *arXiv:1507.07885*.
- [24] S. Ono, Y. Ando, T. Murayama, F. F. Balakirev, J. B. Betts, and G. S. Boebinger, *Phys. Rev. Lett.* **85**, 638 (2000).
- [25] N. Doiron-Leyraud, M. Sutherland, S. Y. Li, L. Taillefer, R. Liang, D. A. Bonn, and W. N. Hardy, *Phys. Rev. Lett.* **97**, 207001 (2006).
- [26] Y. Onose, Y. Taguchi, K. Ishizaka, and Y. Tokura, *Phys. Rev. B* **69**, 024504 (2004).
- [27] C. H. Wang, G. Y. Wang, T. Wu, Z. Feng, X. G. Luo, and X. H. Chen, *Phys. Rev. B* **72**, 132506 (2005).
- [28] C. H. Wang, G. Y. Wang, H. B. Song, Z. Feng, S. Y. Li, and X. H. Chen, *Supercond. Sci. Technol.* **18**, 763 (2005).
- [29] S. I. Woods, A. S. Katz, M. C. de Andrade, J. Herrmann, M. B. Maple, and R. C. Dynes, *Phys. Rev. B* **58**, 8800 (1998).
- [30] P. L. Bach, S. R. Saha, K. Kirshenbaum, J. Paglione, and R. L. Greene, *Phys. Rev. B* **83**, 212506 (2011).
- [31] W. Yu, B. Liang, P. Li, S. Fujino, T. Murakami, I. Takeuchi, and R. L. Greene, *Phys. Rev. B* **75**, 020503(R) (2007).
- [32] Y. Dagan, M. M. Qazilbash, C. P. Hill, V. N. Kulkarni, and R. L. Greene, *Phys. Rev. Lett.* **92**, 167001 (2004).
- [33] S. Finkelman, M. Sachs, G. Droulers, N. P. Butch, J. Paglione, P. Bach, R. L. Greene, and Y. Dagan, *Phys. Rev. B* **82**, 094508 (2010).
- [34] J. Gauthier, S. Gagné, J. Renaud, M.-E. Gosselin, P. Fournier, and P. Richard, *Phys. Rev. B* **75**, 024424 (2007).
- [35] F. F. Tafti, F. Laliberté, M. Dion, J. Gaudet, P. Fournier, and L. Taillefer, *Phys. Rev. B* **90**, 024519 (2014).
- [36] X. F. Sun, Y. Kurita, T. Suzuki, S. Komiya, and Y. Ando, *Phys. Rev. Lett.* **92**, 047001 (2004).
- [37] K. Jin, X. H. Zhang, P. Bach, and R. L. Greene, *Phys. Rev. B* **80**, 012501 (2009).
- [38] Y. Ando, S. Komiya, K. Segawa, S. Ono, and Y. Kurita, *Phys. Rev. Lett.* **93**, 267001 (2004).

- [39] Y. S. Lee, K. Segawa, Z. Q. Li, W. J. Padilla, M. Dumm, S. V. Dordevic, C. C. Homes, Y. Ando, and D. N. Basov, *Phys. Rev. B* **72**, 054529 (2005).
- [40] Y. Peng, J. Meng, D. Mou, J. He, L. Zhao, Y. Wu, G. Liu, x. Dong, S. He, J. Zhang, X. Wang, Q. Peng, Z. Wang, S. Zhang, F. Yang, C. Chen, Z. Xu, T. K. Lee, and X. J. Zhou, *Nat. Commun.* **4**, 2459 (2013).
- [41] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevLett.117.197001> for sample information and additional analysis, which includes Ref. [42].
- [42] R. Liang, D. A. Bonn, and W. N. Hardy, *Phys. Rev. B* **73**, 180505(R) (2006).
- [43] F. Laliberte, W. Tabis, S. Badoux, B. Vignolle, D. Destraz, N. Momono, T. Kurosawa, K. Yamada, H. Takagi, N. Doiron-Leyraud, C. Proust, and L. Taillefer, [arXiv: 1606.04491](https://arxiv.org/abs/1606.04491).
- [44] Y. Onose, Y. Taguchi, K. Ishizaka, and Y. Tokura, *Phys. Rev. Lett.* **87**, 217001 (2001).
- [45] C. Weber, K. Haule, and G. Kotliar, *Nat. Phys.* **6**, 574 (2010).
- [46] J. C. Phillips, A. Saxena, and A. R. Bishop, *Rep. Prog. Phys.* **66**, 2111 (2003).
- [47] Q. He, C.-H. Yeh, J.-C. Yang, G. Singh-Bhalla, C.-W. Liang, P.-W. Chiu, G. Catalan, L. W. Martin, Y.-H. Chu, J. F. Scott, and R. Ramesh, *Phys. Rev. Lett.* **108**, 067203 (2012).
- [48] I. S. Beloborodov, A. V. Lopatin, V. M. Vinokur, and K. B. Efetov, *Rev. Mod. Phys.* **79**, 469 (2007).
- [49] C. Tsuei, A. Gupta, and G. Koren, *Physica (Amsterdam)* **161C**, 415 (1989).