Evidence for two fluids in the ground state of cuprates

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We report charge density measurements, using NMR, in the superconducting compound (Ca$_x$La$_{1-x}$)(Ba$_{1.75-x}$La$_{0.25+x}$)$_2$Cu$_3$O$_y$, which has two independent variables $x$ (family) and $y$ (oxygen). For underdoped samples we find the rate at which holes are introduced into the plane upon oxygenation to be family-independent. In contrast, not all carriers contribute to either antiferromagnetic or superconducting order parameters. This result is consistent with a two fluid phenomenology or intrinsic mesoscopic inhomogeneities in the bulk. We also discuss the impact of weak-chemical-disorder on $T_c$.

Proper counting of holes in the cuprates is essential for understanding their properties. Since they have a small ratio of coherence length to mean free path, they are considered clean superconductors where the carrier density $n$ is the same as the superconducting carrier density $n_s$ in the zero temperature limit. In contrast, modern experiments in the superconducting part of the phase diagram consistently find strong inhomogeneities in these materials. Moreover, several theories of cuprate superconductivity are based on two fluids comprised of: hole-poor and hole-rich regions, bosons and fermions, hot and cooled electrons, etc. This contradiction leads to the question: do all holes participate in the superconducting order parameter? Addressing this question requires a simultaneous measurement of $n$ and $n_s$. The in-plane $^{63}$Cu(2) nuclear quadrupole resonance (NQR) parameter $\nu_Q$ is a direct measure of $n$. $n_s$ can be extracted from the penetration depth. In this work we compare the two numbers for different superconducting families. We find that $n_s$ is not a universal fraction of $n$, and provide a simple relation between $n_s$ and $y$, from parent to overdoped samples. The data support two fluids-based theories for the entire phase diagram, but constrain the possible division between the fluids.

Our study is done on the (Ca$_x$La$_{1-x}$)(Ba$_{1.75-x}$La$_{0.25+x}$)$_2$Cu$_3$O$_y$ (CLBLCO) compound where each value of $x$ is a superconducting family with its own maximum $T_c$ ($T_{c}^{\max}$) ranging between 58 and 80 K, as shown in Fig. 1(a). By choosing a sample with a particular $x$ and oxygen level $y$ one can control $n$ and $n_s$ independently. This allows determination of the rates at which oxygenation produces carriers and carriers turn superconducting. We evaluate the average $\nu_Q$ form $^{63}$Cu(2) nuclear magnetic resonance (NMR) measurements. In passing, we estimate the chemical disorder from the width, $\Delta \nu_Q$, of the $\nu_Q$ distribution. $n_s$ is obtained from previous muon spin rotation ($\mu$SR) measurements.

The ability of NMR to determine carrier density is based on the fact that $^{63}$Cu, with its spin 3/2 nuclei, is directly coupled to charge degrees of freedom via the electric field gradient (EFG), and $\nu_Q$ is a measure of this coupling. $\nu_Q$, in turn, depends linearly on the hole density $\delta$ according to

$$\nu_Q = A n_s + \nu_Q^0,$$

where $A$ and $\nu_Q^0$ are doping-independent, but, in principle, could be family-dependent. This linear dependence was demonstrated for various compounds such as Y123, La$_2$-Sr$_x$CuO$_4$ (La214), and HaBa$_2$CuO$_{4+\delta}$ (Ha124). Therefore, Eq. (1) and the ability of NMR to detect the in-plane copper $[\nu_Q(2)]$ $\nu_Q$ selectively will allow us to determine the evolution of the in-plane carrier concentration and the width of its distribution.

The measurements were done on powder samples fully enriched with $^{63}$Cu. Their preparation is described in Ref. [12]. The oxygen content was measured by double iodometric titration. The accuracy of this method in the enriched CLBLCO is about 0.01. We measured between five and seven different samples for each $x$ in the normal state at 100 K. The most overdoped sample is a non superconducting $x = 0.1$ compound. The NMR measurements were done by sweeping the field in a constant applied frequency $f_{\text{app}} = 77.95$ MHz, using a $\pi/2 - \pi$ echo sequence. The echo signal was averaged 100,000 times and its area evaluated as a function of field. The full spectrum of the optimally doped $x = 0.4$ sample ($y = 7.156$) is shown in the inset of Fig. 1.

The Cu spin Hamiltonian can be written as:

$$\mathcal{H}/\hbar = -\nu_I \mathbf{I} \cdot (1 + \mathbf{K}) \cdot \hat{\mathbf{H}} + \frac{\nu_Q}{6} [3I_x^2 - I_y^2 + \eta(I_x^2 - I_y^2)],$$

where $\nu_I = (63\gamma/2\pi)H$, $\mathbf{H}$ is a unit vector in the direction of the field, $\mathbf{K}$ is the shift tensor, and $\eta$ is the asymmetry parameter of the EFG. In the absence of magnetic field, there is only one transition frequency given by $f = \nu_Q \sqrt{1 + \eta/3}$, so $\nu_Q$ cannot be separated from $\eta$, and the use of the magnetic field is essential. This field, applied in the direction $\theta$ and $\phi$ with respect to the principal axis of the EFG, lifts this degeneracy, and the transition frequencies $\nu_m(H, \theta, \phi)$ are expected: a center line which corresponds to the $1/2 \rightarrow -1/2$ transition ($m = 0$), and two satellites which correspond to the $3/2 \rightarrow 1/2$ ($m = 1$) and $-1/2 \rightarrow -3/2$ ($m = -1$) transitions. Expressions for $\nu_m(H, \theta, \phi)$ up to second order perturbation theory in $\nu_Q$ for completely asymmetric
of strong chemical disorder. This means that the evolution of the main peaks for \( x = 0.4 \) as a function of oxygen level \( y \) for the \( x = 0.1 \) and \( 0.4 \) CLBLCO samples.

A zoom on the main features of the Cu(2) signal of all four optimally doped samples is depicted in Fig. 2 (note the three axis breakers). The evolution of the main peaks as \( x \) increases is highlighted by the dotted lines. It is clear that as \( x \) decreases the peaks move away from each other. This means that \( \nu_Q \) at optimal doping is a decreasing function of \( x \). A more interesting observation is the fact that there is no change in the width of the peaks, at least not one that can easily be spotted by the naked eye. This means that the distribution of \( \nu_Q \) is \( x \)-independent and that there is no difference in the weak-chemical-disorder (WCD) between the optimally doped samples of the different families. By WCD we mean disorder that is not strong enough to “wipe out” the contribution of the Cu(2) in its vicinity from the spectrum. Thus, WCD is not relevant to the variation of \( T_c^\text{max} \) between the different families. As we shall see, this conclusion is supported by more rigorous analysis.

Although we limit our conclusion to WCD, it is important to place a few limitations on the possible existence of strong chemical disorder. The difference in ionic size between La and Ba in the Ba layer creates local lattice distortions. These impact the apical oxygen O(4) position, which contributes to \( \nu_q \). However a simple calculation based on Eqs. 4 and 5 in Ref. 15 shown that the probability of finding an O(4) out of its ideal place increases with \( x \). Therefore, if the O(4) displacement were important for chemical disorder we would have expected the \( x = 0.4 \) family to have a lower \( T_c \) than the \( x = 0.1 \), in contrast to observation. In addition, magic angle spinning (MAS) Ca NMR found only one line. This ensures the existence of only one Ca site. Also, since Ca and Ba have the same valance, and the total amount of La is fixed, doping is done only by the chain layer oxygen, which is relatively remote from the CuO\(_2\) layer. Therefore, chemical disorder is expected to be minimal. Overall, we cannot find evidence for the existence or importance of strong chemical disorder.

The evolution of the main peaks for \( x = 0.4 \) as a func-

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**FIG. 1:** (a) The superconducting part of the \((\text{Ca}_x\text{La}_{1-x})(\text{Ba}_{1.75-\delta}\text{La}_0.25+\delta)\text{Cu}_3\text{O}_9\) phase diagram. (b) Normalized critical temperatures plotted as a function of \( K(x)\Delta y \), including the glass \( T_r \) temperature, and corrected Néel temperature \( T_r^{\text{corr}} \) (see text). Inset, the normalized muon spin relaxation rates taken from Ref. 2 as a function of oxygen level \( y \) for the \( x = 0.1 \) and \( 0.4 \) CLBLCO samples.

**FIG. 2:** NMR spectra of \( ^{63}\text{Cu} \) at \( T = 100 \) K in optimally doped CLBLCO samples with varying \( x \). The inset shows the full spectrum of the \( x = 0.4 \) compound including contributions from Cu(1) and Cu(2). The main figure zooms in on the Cu(2) contribution (note the three axis breakers). The position of the Cu(2) peaks are shown by dotted lines.
FIG. 3: NMR spectra of $^{63}$Cu(2) at $T = 100$ K in CLBLCO samples with $x = 0.4$ and varying $y$. The positions of the peaks are shown by dotted lines.

The fit is not perfect, mostly because the spectrum is not symmetric. Such an asymmetric spectrum is a result of correlations between different a priori random parameters in the Hamiltonian [21]. For example, if the shift tensor $\mathbf{K}$ is also non uniform in the sample but its values are correlated with the values of $\nu_Q$, the spectrum could be non symmetric. Attempts to take this kind of effect into account failed due to the enormous increase in computing time. We continue the discussion based on the best fit we could practically achieve. For a discussion on the error bars evaluation see Ref. [17]. We could not determine, $K_z$, $K_x$ and $K_y$ very accurately, and found very small $\eta$ for all the samples, ranging from 0 to 0.1, in agreement with various estimates for Y123 [22].

The results for fitted $\nu_Q$ and $\Delta\nu_Q$ are shown in Fig. 4 (a) and (b), respectively. From Fig. 4(a) it is clear that $\nu_Q$ grows linearly with doping in the underdoped side of the phase diagram, in agreement with other compounds [9, 10, 11], and Eq. 1. It is also clear that $\nu_Q$ in this equation is $x$-dependent, but this could be attributed to NQR base line properties. In contrast, the behavior in the overdoped side of the $x = 0.1$ is surprising since $\nu_Q$ saturates. As pointed out before in Ref. [17], the added holes on the overdoped side no longer go into the planes.

The most interesting finding is that within the experimental error, the slope of $\nu_Q(x, y)$ in the underdoped side is $x$-independent, as demonstrated by the parallel solid lines, and $\partial\nu_Q(x, y)/\partial y = 7.2(1.4)$ MHz/Oxygen. This means that the rate at which holes are introduced into the CuO$_2$ planes, $\partial n/\partial y$, is a constant independent of $x$ or $y$ in the underdoped region. Using further the ubiquitous assumption that the optimal hole density, $n^{\text{opt}}$, at optimal oxygennation, $y^{\text{opt}}$, is universal, we conclude that $n(x, y)$ is a function only of $\Delta y = y - y^{\text{opt}}$. This result was predicted previously [13] based on bond valence calculations.

An implication of this conclusion is that the Presland et al. formula $T_c/T_c^{\text{max}} = 1 - 82.6(n - 0.16)^2$ could not be correct for CLBLCO. For example, the $x = 0.1$ and $x = 0.4$ samples with $T_c = 0$ have different $\Delta y$, and according to our experiment different $n$. However, the Presland formula predicts the same $n$.

If the conversion from oxygen to holes is universal, but different families begin to superconduct at different $y$ values (see Fig. 1), it means that the rate at which holes turn into superconducting carriers is family-dependent. To demonstrate this concept further we focus on the region of the CLBLCO phase diagram where $T_c$ grows linearly
with doping, as emphasized by the solid lines in Fig. 4(a). This behavior is unique to CLBLCO, which does not show a 1/8 dip or plateau in $T_c$. We determine $n_s$ from $\mu$SR measurement at $T \rightarrow 0$. The $\mu$SR relaxation rate $\sigma$ is a measure of the density of superconducting carriers, and $\sigma/\sigma_{opt} = n_s/n_{s, opt}$. In the inset of Fig. 4(b) we depict $\sigma/\sigma_{opt}$, taken from Ref. [2], as a function of $y$ for two extreme families with $x = 0.1$ and 0.4. In this particular region $\partial n_s/\partial y$ is a constant, which is family-dependent and denoted hereafter by $K(x)$. Using the universality of $\partial n/\partial y$ we find that $\partial n_s/\partial n = K(x)(\partial n/\partial y)^{-1}$, which varies between families. This, again, rules out the possibility that $n_s = n$ in all CLBLCO samples.

In light of previous work a stronger conclusion could in fact be extended to the entire phase diagram of CLBLCO. In Fig. 4(b), taken from Ref. [2], we depict $T_{N, opt}^c, T_g$ and $T_c$, normalized by $T_c^{max}$ of each family. $T_{N, opt}^c$ is the Néel temperature after the contribution from anisotropies have been divided out. The scaled variable $K(x)\Delta y$, with $K = 0.77, 0.67, 0.54, 0.47$, for the $x = 0.1$ to 0.4 respectively, collapsed the entire phase diagram into a single function. However, no interpretation was given for $K(x)$. The present experiment suggests that $\partial n_s/\partial y = K(x)$ over the entire phase diagram, and that $n_s$ should be considered as carriers participating in both superconducting and antiferromagnetic order parameters.

As for the line width, examination of Fig. 4(b) reassures us of our previous intuition that $\Delta \nu_Q$ is not changing between the different families, especially for the optimally doped samples. This becomes obvious when considering $\Delta \nu_Q$ of YBCO at $T = 100$ K, which also has $\nu_Q = 31$ MHz, but $\Delta \nu_Q = 0.5$ MHz [23]. The changes in $\Delta \nu_Q$ between $x = 0.4$ and $x = 0.1$ are minute compared to the changes between $x = 0.1$ and YBCO. Yet $x = 0.4$ has a $T_c^{max}$ very similar to YBCO. Thus, if YBCO is considered disorder free, the difference in WCD between $x = 0.4$ and $x = 0.1$ either does not exist or is not relevant. The same conclusion was reached by MAS Ca NMR [10].

In summary, we find that the rate at which holes are doped into the planes when oxygen is added, $\partial n/\partial y$, is identical in all families in the superconducting underdoped region. In part of this region, the rate at which holes contribute to the condensate $\partial n_s/\partial n \propto K(x)$ is a family-dependent constant. Based on this and previous findings, we conclude that in CLBLCO not all oxygens contribute holes to the superconducting and antiferromagnetic order parameters. We also show that weak-disorder, as determined from $\Delta \nu_Q$, is not playing a role in the $T_c^{max}$ variations. This reinforces previous conclusions that the changes in $T_c^{max}$ between different CLBLCO families is caused only by variation in the in-plane coupling constant $J$ [24]. Finally, since $\nu_Q$ is not changing in the overdoped side as the oxygen level increases, in this side holes are not added to the planes. The open question in this study is which physical or chemical property of CLBLCO sets $K(x)$.

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[1] J. L. Tallon, J. R. Cooper, S. H. Naqib, and J. W. Loram, Phys. Rev. B 73, 180504(R) (2006).
[2] A. Keren, A. Kanigel, J. S. Lord, A. Amato, Solid State Commun. 126, 39 (2003). A. Kanigel, A. Keren, A. Knizhnik, O. Shafir, Phys. Rev. B 71, 224511 (2005).
[3] J. M. Tranquada et al, Nature 375, 561 (1995); Ch. Niedermayer, C. Bernhard, T. Blasius, A. Golnik, A. Moodenbaugh, and J. I. Budnick, Phys. Rev. Lett. 80, 3843
(1998); C. Howald, P. Fournier, and A. Kapitulnik, Phys. Rev. B 64, 100504(R) (2001); S. H. Pan et al., Nature 413, 282 (2001); M. Vershinin et al., Science 303, 1995 (2004).
[4] V. J. Emery and S. A. Kivelson, Physica C 209, 597 (1993);
[5] E. Altman and A. Auerbach Phys. Rev. B 65, 104508 (2002); J. Ranninger, J.M. Robin, and M. Eschrig, Phys. Rev. Lett. 74, 4027 (1995); R. Friedberg and T.D. Lee, Phys. Rev. B 40, 6745 (1989).
[6] D. Pines, cond-mat/0404151
[7] R. Ofer et al., cond-mat/0606695
[8] K. Asayama et al., Prog. Nuc. Mag. Reso. Spec. 28, 221 (1996).
[9] H. Yasuoka, in Spectroscopy of Mott Insulator and Correlated Metals, edited by A. Fujimori and Y. Tokura, Solid State Sciences, Vol. 119 (Springer-Verlag, Berlin, 1995), p. 213.
[10] G.-q. Zheng et al., J. Phys. Soc. Jpn. 64, 2524 (1995).
[11] A. A. Gippius et al., Physica C 276, 57 (1997).
[12] D. Goldschmidt, A. Knizhnik, Y. Direktovitch, G. M. Reisner, and Y. Eckstein, Phys. Rev. B 49, 15928 (1994).
[13] C. P. Slichter, Principles of Magnetic Resonance, Harper and Row, New York, (1963).
[14] J. F. Baugher, P.C. Taylor, T. Oja and P.J. Bray, J. of Chem. Phys. 50, 4914 (1969).
[15] A. Keren et al., Phys. Rev. B 57, 10745 (1998).
[16] S. Marchand, Ph. D. thesis, Universite Paris 6.
[17] A. Kanigel and A. Keren, cond-mat/0606475
[18] O. Chmaissem, Y. Eckstein, C. G. Kuper, Phys. Rev. B 63, 174510 (2001).
[19] M. R. Preslan et al., Physica C 176, 95 (1991).
[20] W. H. Press et al. numerical Recipes in C, Cambridge University Press (1988).
[21] J. Haase, C. P. Slichter, R. Stern, C. T. Milling and D. G. Hinks, J. Supercond. 13, 723 (2000).
[22] C. H. Pennington, D. J. Durand, D. B. Zax, C. P. Slichter, J. P. Rice, and D. M. Ginsberg, Phys. Rev. B 37, R7944 (1988); T. Shimizu et al., Journal of the Physical Society of Japan, 57, 2494 (1988).
[23] A. Kanigel et al., Phys. Rev. Lett. 88, 137003 (2002).
[24] R. Ofer, S. Levy, A. Kanigel, and A. Keren, Phys. Rev. B 73, 012503, (2006).
[25] E. P. Stoll and P. F. Meier, Private Communication.