Magnetic Dispersion of the Diagonal Incommensurate Phase in Lightly Doped La_{2-x}Sr_xCuO₄

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We present inelastic neutron scattering experiments on a single-domain crystal of lightly doped $La_{1.96}Sr_{0.04}CuO_4$. We find that the magnetic excitation spectrum in this insulating phase with a diagonal incommensurate spin modulation is remarkably similar to that in the superconducting regime, where the spin modulation is bond parallel. In particular, we find that the dispersion slope at low energy is essentially independent of doping and temperature over a significant range. The energy at which the excitations cross the commensurate antiferromagnetic wave vector increases roughly linearly with doping through the underdoped regime.

DOI: 10.1103/PhysRevLett.101.197001 PACS numbers: 74.72.Dn, 75.40.Gb

Superconductivity in the cuprates is obtained by doping holes into a parent antiferromagnetic insulator. How the transition occurs from antiferromagnetic to superconducting ground states as a function of doping remains a controversial issue. In the case of La_{2-r}Sr_rCuO₄ (LSCO), the commensurate antiferromagnetic order disappears at x =0.02, while superconductivity appears for $x \ge 0.055$; the region in between is often called the "spin-glass" phase [1]. Neutron scattering studies have shown that, at low temperatures ($T < T_{\rm el} \sim 20$ K), magnetic ordering is indicated by the appearance of incommensurate elastic peaks split about the antiferromagnetic wave vector, \mathbf{Q}_{AF} [2,3]. The corresponding spin modulation occurs within the CuO₂ planes along a direction at 45° to the Cu-O bonds; in fact, it has a unique orientation with respect to the orthorhombic axes [4]. The magnetic incommensurability rotates to parallel to the Cu-O bonds with the onset of superconductivity at $x \sim 0.055$. While there have been recent studies of the high-energy magnetic dispersion [5] and the scaling behavior of the low-energy spin fluctuations [6] in LSCO with x = 0.05, the overall nature of the dispersion, and its relation to that in the superconducting phase, has not yet been clarified. This is an important issue, especially in light of recent evidence for a vortex-liquid state extending from the superconducting regime down to $x \approx 0.03$ [7].

Here we present an inelastic neutron scattering study of the spin dynamics in LSCO with x=0.04, in the middle of the spin-glass regime. Despite the rotated incommensurability, we find that the spectrum is consistent with the "hour glass" dispersion observed in superconducting LSCO [8–10] (and also in YBa₂Cu₃O_{6+x} [11–15]). Even at $T\gg T_{\rm el}$, there is a strong anisotropy of the low-energy fluctuations, consistent with the nematiclike response recently reported for YBa₂Cu₃O_{6.45} [16,17]. Furthermore, the effective velocity of the low-energy excitations is ap-

proximately independent of energy, so that the energy, $E_{\rm cross}$, at which the low-energy excitations cross ${\bf Q}_{\rm AF}$ scales linearly with doping, following the elastic limit of the magnetic incommensurability. The lack of variation of the spin-fluctuation velocity is reminiscent of the doping independence of the Fermi velocity observed in photoemission studies [18]; however, we can rule out a spin-densitywave instability as the source of the low-energy magnetic correlations, because the magnitude of $2k_{\rm F}$ (twice the Fermi wave vector) measured by photoemission [19] is inconsistent with the magnetic wave vector. At energies above E_{cross} , the magnetic excitations are consistent with the effective antiferromagnetic dispersion previously observed in LSCO with x = 0.05 [5]. This supports the concept that the magnetic excitations evolve directly from the antiferromagnetic insulator.

Besides La_{1.96}Sr_{0.04}CuO₄, we will also show some results for $La_{1.95}Sr_{0.05}Cu_{0.97}Zn_{0.03}O_4$ (LSCZO); previous work has shown that the Zn increases the spin correlation length while slightly reducing the incommensurability [20]. Both crystals were grown by the traveling solvent floating zone method and have dimensions of $\sim 6\phi \times$ 25 mm³. Each crystal contains essentially a single domain of the low-temperature orthorhombic structure, with lattice parameters a = 5.34 Å and b = 5.42 Å for the pure LSCO crystal at low temperature. The a and b axes are at 45° to the Cu-O bonds. In this coordinate system, $\mathbf{Q}_{AF} = (100)$, (010), in reciprocal lattice units (r.l.u.), $(2\pi/a, 2\pi/b,$ $2\pi/c$). The incommensurate magnetic peaks are split uniquely along \mathbf{b}^* by an amount $\boldsymbol{\epsilon}$, so that the magnetic peaks are observed at $(2m, 2n + 1 \pm \epsilon, 0)$ and (2m + 1, $2n \pm \epsilon$, 0) for integer m, n. The LSCO sample yielded incommensurate peaks with $\epsilon = 0.0513(7)$ r.l.u. for $T \lesssim$ 20 K, with a squared Lorentzian line width of $\kappa =$ 0.04 Å^{-1} , consistent with previous results [21]. For the LSCZO crystal, $\epsilon = 0.0543$ r.l.u. [20].

The elastic magnetic scattering was characterized on the cold triple-axis spectrometer LTAS at the Japanese Research Reactor (JRR-3) of the Japan Atomic Energy Agency (JAEA). Neutrons with an energy of 5 meV were used, together with a horizontal collimator sequence of guide-80'-S-80'-80'; contamination from higher-order beams was effectively eliminated using Be filters.

The inelastic scattering measurements on LSCO were performed on the thermal triple-axis spectrometers TAS-1 and TAS-2 at JRR-3 with a fixed final neutron energy, E_f , of 13.7 meV. The horizontal collimator sequences were open-80'-S-80'-80' on TAS-1 and guide-80'-S-80'-open on TAS-2. For LSCZO, the measurements were performed on the thermal triple-axis spectrometer HB-1 at the High Flux Isotope Reactor (HFIR), Oak Ridge National Laboratory. The measurement conditions were $E_f = 14.7$ meV, horizontal collimations of 48'-60'-S-80'-240'. Pyrolytic graphite (PG) filters were used to suppress higher harmonics. The single crystals were oriented in the (HK0) scattering plane and were mounted in a closed-cycle He gas refrigerator.

Figure 1 shows the typical neutron inelastic scattering spectra between 2.5 meV and 30 meV in LSCO (x = 0.04) measured at low temperature along the modulation direction. At energies of 4 meV and below, the two peak structure is clear. With increasing energy, the peak separation becomes smaller and the peak width also becomes

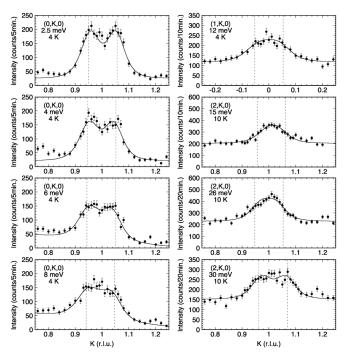


FIG. 1. Neutron inelastic scans along (0, K, 0), (1, K, 0), or (2, K, 0) in La_{1.96}Sr_{0.04}CuO₄. The solid lines are the results of fits as discussed in the text. The broken lines represent the peak positions observed at $\hbar\omega = 0$ meV. The measurements for $\hbar\omega \le 12$ meV were performed on TAS-2, and for $\hbar\omega \ge 15$ meV on TAS-1 (where the effective flux is $\sim 3 \times$ higher).

broadened. We have confirmed that the peak broadening corresponds to the resolution effect. The solid lines in Fig. 1 are the results of fits of a convolution of the resolution function with two two-dimensional (2D) squared Lorentzians. In the fitting, the linewidth (κ) of each peak, which corresponds to the effective inverse correlation length, is assumed to be isotropic in the a-b plane and fixed at the value determined from the elastic scattering (0.04 Å^{-1}) . This model reproduces the observed spectra reasonably well for all measured energies $(2.5 \le \hbar\omega \le$ 30 meV). The analysis indicates that the peak separation (2ϵ) decreases towards zero as $\hbar\omega$ is raised to 15 meV. Strong, but weakly dispersing, phonon scattering creates a background that varies slowly with \mathbf{Q} near \mathbf{Q}_{AF} ; however, its strength hampers analysis in the range $15 < \hbar\omega <$ 25 meV. For $\hbar\omega > 25$ meV, the peak separation appears to grow. The Q-integrated intensity decreases monotonically with increasing energy. This behavior is consistent with that observed in LSCO with x = 0.05 [5]; however, it is distinct from the reported behavior in LSCO with x =0.16 [9] and in La_{1.875}Ba_{0.125}CuO₄ [8], where the intensity peaks at finite energy.

The anisotropy of the scattering, measured parallel and perpendicular to the incommensurate peaks, is shown in Fig. 2 for $T=50~{\rm K}\gg T_{\rm el}$, where there is no static order. This is to illustrate that the character of the low-energy spin correlations does not depend on whether there is a static component. As noted in the introduction, this anisotropy looks very similar to that found recently in YBa₂Cu₃O_{6.45} [16] and consistent with the observed anisotropic electronic response [17,22]. (Of course, in both of these cases,

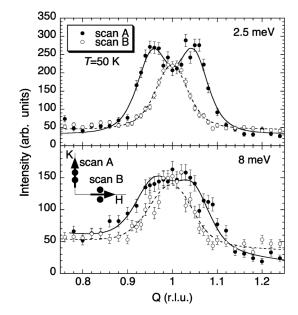


FIG. 2. Scans measured parallel (scan A) and perpendicular (scan B) to the incommensurate wave vector, as indicated in the inset, for energies of 2.5 and 8 meV at T=50 K. The particular positions for the scans were chosen to optimize the resolution.

there is already a reduction from 4-fold symmetry due to the crystal structure.) It is also of interest to consider the 2D distribution of intensity at 25 meV and above; however, resolution effects prevented any definitive conclusions.

Figure 3 summarizes the neutron inelastic results for our samples. For x=0.04, the effective incommensurability decreases towards zero as the energy approaches 15 meV, while it grows for 25 meV and above. The results for the LSCZO sample are almost identical. At higher energies, we expect the dispersion to be similar to that measured in $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$ with x=0.05 [5], and these points are included in the figure. The lower energy results for $\text{La}_{1.875}\text{Ba}_{0.125}\text{CuO}_4$ [8] are also shown for comparison (plotted as a function of $|\mathbf{q}|$, and ignoring the difference in orientation of \mathbf{q}). Overall, the magnetic dispersion has an hour-glass shape similar to that observed in the parallel incommensurate phase.

An intriguing feature is that the slope of the effective low-energy dispersion (which corresponds to a velocity) does not appear to depend strongly on doping. If the velocity is independent of doping, while the incommensurability is linear in x for $x \leq \frac{1}{8}$, then one would expect E_{cross} to be proportional to x [23,24]. In Fig. 4, we plot values for E_{cross} extracted from neutron scattering studies [8–

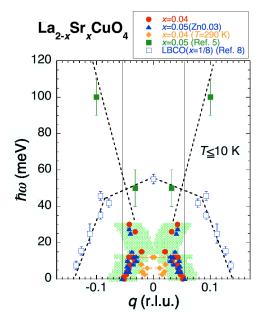


FIG. 3 (color online). Magnetic dispersion relation along q_K in La_{1.96}Sr_{0.04}CuO₄ (filled circles) below 10 K and at 290 K (filled diamonds). For comparison, magnetic dispersion relations in other related compounds are also shown. The filled triangles, filled squares, and open squares are data of La_{1.95}Sr_{0.05}Cu_{0.97}Zn_{0.03}O₄, La_{1.95}Sr_{0.05}CuO₄ (Ref. [5]), and La_{1.875}Ba_{0.125}CuO₄ (Ref. [8]), respectively. It is noted that the peak positions are 45° rotated in La_{1.875}Ba_{0.125}CuO₄. The thick shaded bars represent the full width at half maximum of the excitation peaks in La_{1.96}Sr_{0.04}CuO₄. The broken lines are visual guides.

10,25,26]. Each symbol represents an estimate from interpolating a parabola through the low-energy dispersion, while the error bars indicate the energy range over which constant-energy cuts are consistent with a single peak of minimum width. The results are consistent with $E_{\rm cross} \sim x$ for $x \lesssim \frac{1}{8}$. This trend is opposite to the gradual decrease observed for high-energy, antiferromagneticlike spin excitations [1].

We also investigated the temperature dependence of the magnetic excitation spectra. Figure 5(a) shows scans at $\hbar\omega=2.5$ meV. The lines through the data are similar fits to those in Fig. 1, now with κ taken to be temperature independent. (Even if κ were temperature dependent, we estimate an upper limit at 290 K of ~ 0.05 Å⁻¹, an increase of only 25%.) We conclude that ϵ gradually decreases as the temperature rises, with the intensity also decreasing. The temperature dependence of ϵ evaluated at 2.5, 6, and 12 meV is plotted in Fig. 5(b). In each case, the decrease is roughly linear in temperature. A consequence of this is that the slope of the dispersion does not appear to change.

Just as the incommensurability evolves with temperature, so does the electronic conductivity. In particular, the conductivity is poor at room temperature where the magnetic correlations are almost commensurate. The lowfrequency optical conductivity grows with cooling [22], as the spin incommensurability develops. Given the continuous evolution of E_{cross} with doping, providing a connection with La_{1.875}Ba_{0.125}CuO₄ where charge and spin stripe order is known to occur [8], we believe that charge stripes and moment modulation are likely to be an important part of the incommensurate response in LSCO with x = 0.04. We are aware that there has been considerable work on spiral-order models for the spin-glass phase [27– 31]. While we cannot rule out such models on the basis of the observed magnetic spectrum, explaining Fig. 4 may be a challenge for such an approach.

Another possibility is spin-density-wave order due to Fermi-surface nesting [32]. Angle-resolved photoemission measurements on LSCO [19] indicate that $2k_F$ in the nodal direction would yield a diagonal magnetic modulation with

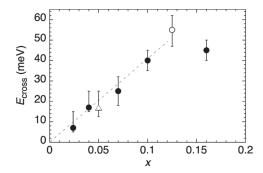


FIG. 4. Plot of $E_{\rm cross}$ vs x in LSCO (filled circles), LSCZO (open triangle), and ${\rm La_{2-x}Ba_xCuO_4}$ (open circle), including results from Refs. [8–10,25,26]. The dashed line is a guide to the eye.

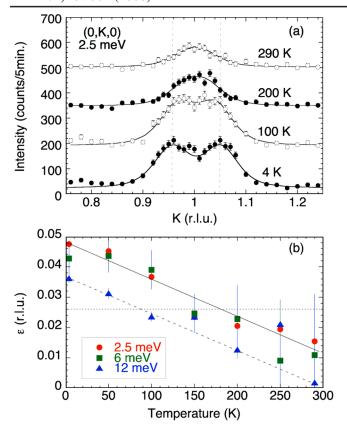


FIG. 5 (color online). (a) Temperature dependence of the magnetic excitation spectra at 2.5 meV. Successive scans have been displaced vertically by 150 counts for clarify. The solid lines are the results of fits of a convolution of the resolution function with two 2D squared Lorentzians. The broken lines represent the peak positions at 4 K. (b) Temperature dependence of the incommensurability (ϵ) at 2.5, 6, and 12 meV in La_{1.96}Sr_{0.04}CuO₄. The solid and broken lines are visual guides for the data. The dotted horizontal line shows the peak width (HWHM). Two peaks can hardly be resolved when ϵ becomes much smaller than this value.

 $\epsilon = 0.15 \text{ Å}^{-1}$ for x = 0.04. This happens to be 2.5 times greater than the experimental value of 0.06 Å⁻¹. Hence, a simple nesting picture appears to be ruled out.

In terms of a stripe picture, the dominant magnetic interaction would be superexchange within locally antiferromagnetic domains. There is still a challenge to understand why the dispersion of the low-energy excitations is not significantly affected by the rotation in stripe orientation. One possibility suggested by Granath [33] is that a diagonal stripe might consist of a staircase pattern of bond-parallel stripes, in which case local interactions would be independent of average stripe orientation. Granath [33] found that such a pattern is necessary in order to obtain consistency with the photoemission experiments [19]. Furthermore, we note that, while spin-wave calculations for a diagonal-stripe model [24,34] provide a good description of the magnetic spectrum observed in insulating

 $\text{La}_{2-x}\text{Sr}_x\text{NiO}_4$ [35,36], they have difficulty in reproducing the hour-glass-like spectrum in the present case. Even in the case of parallel stripes, calculations by Seibold and Lorenzana [37] based on the Hubbard model predict $E_{\text{cross}} > 30 \text{ meV}$ for $x \ge 0.05$, which is well above the experimental trend. Thus, while our results are phenomenologically consistent with stripe correlations, improvements in theoretical models are needed.

We would like to thank T. Tohyama and Y. Koike for stimulating discussions. This study was supported in part by the U.S.-Japan Cooperative Program on Neutron Scattering and by a Grant-in-Aid for Scientific Research from the MEXT of Japan. Work at Oak Ridge National Laboratory and Brookhaven National Laboratory was supported by the U.S. Department of Energy's Office of Science under Contract Nos. DE-AC05-00OR22725A and DE-AC02-98CH10886, respectively.

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