Interplay of structural and electronic phase separation in single crystalline La$_2$CuO$_{4.05}$ studied by neutron and Raman scattering

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(Dated: November 7, 2018)

We report a neutron and Raman scattering study of a single-crystal of La$_2$CuO$_{4.05}$ prepared by high temperature electrochemical oxidation. Elastic neutron scattering measurements show the presence of two phases, corresponding to the two edges of the first miscibility gap, all the way up to 300 K. An additional oxygen redistribution, driven by electronic energies, is identified at 250 K in Raman scattering (RS) experiments by the simultaneous onset of two-phonon and two-magnon scattering, which are fingerprints of the insulating phase. Elastic neutron scattering measurements show directly an antiferromagnetic ordering below a Néel temperature of $T_N = 210$ K. The opening of the superconducting gap manifests itself as a redistribution of electronic Raman scattering below the superconducting transition temperature, $T_c = 24$ K. A pronounced temperature-dependent suppression of the intensity of the (100) magnetic Bragg peak has been detected below $T_c$. We ascribe this phenomenon to a change of relative volume fraction of superconducting and antiferromagnetic phases with decreasing temperature caused by a form of a superconducting proximity effect.

I. INTRODUCTION

The development of different kinds of charge inhomogeneities at low temperature is a well known intrinsic property of many strongly-correlated-electron systems. Recent studies of Matsuda et al. on very lightly doped La$_{2-x}$Sr$_x$CuO$_4$ show that even at $x < 0.02$ the system demonstrates an electronic phase separation in which regions of hole-rich and hole-free phases are formed. This observation indicates the possibility for charges to create nonuniform phases in spite of low doping and an expected uniform distribution of Sr-dopants.

Oxygen-doped La$_2$CuO$_{4+\delta}$ presents a system in which charge inhomogeneities exist on the background of phase-separated oxygen interstitials with an intimate interrelation of structural (excess oxygen) and charge subsystems. The discovery of high-$T_c$ superconductivity in samples of nominally pure La$_2$CuO$_4$ led to the first evidence of charge disproportionation in this material. The connection between superconductivity and the hole-rich phase with a finite concentration of interstitial oxygen was directly proved by annealing in high pressure oxygen. It was reported that after annealing, the superconducting volume fraction had been increased. Although the average concentration of charge carriers, roughly estimated as $p \sim 26$, was moderate, the relatively high temperature of the superconducting phase transition, $T_c = 32$ K, was achieved even at $\delta = 0.05$.

Many investigations of La$_2$CuO$_{4+\delta}$ have focused on the structural aspects of the phase separation phenomena (for reviews, see Refs. [17]). The powder neutron diffraction measurements by Jorgensen et al. showed that at low temperature, samples contained two very similar orthorhombic phases, a primary one with $\delta \approx 0$, and a second, oxygen-rich phase that was superconducting. At high temperatures, only a single phase was present, with a reversible oxygen phase separation occurring (in one sample) at 320 K. The development of electrochemical-oxidation techniques has made it possible to determine the $T-\delta$ phase diagram of La$_2$CuO$_{4+\delta}$, the most prominent feature of which is a miscibility gap in the region $0.01 < \delta < 0.05$. Elastic neutron scattering from large electrochemically-oxygenated single crystals revealed a new kind of periodicity in oxygen-rich regions—an ordering of the interstitially-occupied oxygen layers along the $c$-axis, commonly referred to as staging behavior. These results indicated that the phase diagram of La$_2$CuO$_{4+\delta}$ at higher oxidation levels includes new miscibility gaps separating regions of staged phases.

It is generally presumed that the first miscibility gap is driven by a tendency for holes doped into antiferromagnetic Cu$_2$O$_2$ planes to phase separate. The phase separation is screened by the mobile oxygen interstitials. The existence of hole-poor and hole-rich phases at lower temperatures is demonstrated by the observations of, respectively, antiferromagnetic order and superconductivity, however, the ordering temperatures are lower than the structural phase transition temperature for the interstitials. One motivation for the present work is to test more directly the connection between the electronic and interstitial phase separation transitions through the use of Raman scattering, which is sensitive to the presence of a correlated-insulator phase.
through two-magnon and two-phonon scattering.

Another aspect that we explore is the effect of quenched disorder. In samples of La$_2$CuO$_{4+\delta}$ prepared by electrochemical intercalation of oxygen in an aqueous solution, a reversible phase separation is generally observed.\cite{12,13} In the oxygen-rich phase, the oxygen interstitials exhibit staging order, indicating substantial mobility of the oxygens. Modest reductions in the superconducting transition temperature, $T_c$, can be achieved by rapid cooling.\cite{25} In contrast, crystals intercalated electrochemically in a molten salt at relatively high temperature can exhibit a large distribution of lattice parameters at room temperature\cite{26} indicative of quenched disorder. Previous studies\cite{10,20} of such crystals have indicated low oxygen mobility and depressed $T_c$s of 15 to 25 K.

The sample that we study here has, as we will demonstrate, a fine-grained mixture of oxygen-poor and oxygen-rich domains, with substantial strain, at 300 K and below. Neutron diffraction indicates the gradual onset of antiferromagnetic order below 210 K, while $T_c$ is measured to be 24 K. These low transition temperatures, together with the absence of staging order of the interstitials in the oxygen-rich phase, indicate that quenched disorder has pinned the interstitial distributions within the first miscibility gap.\cite{24}

The first surprise that we observe is that the Raman signature of an insulating, antiferromagnetically-correlated phase does not appear until the sample is cooled below $T_I \approx 250$ K, even though a significant degree of oxygen phase separation is already evident. As we will discuss, this result suggests that, contrary to expectations, the miscibility gap is driven largely by lattice, rather than electronic, energies. Such behavior is consistent with a study of miscibility gaps in the related system La$_{2-x}$Sr$_x$NiO$_{4+\delta}$, where the structural phase diagram is found to depend only on the concentration of interstitials and not on the net hole concentration.\cite{21}

The second surprise is a reduction of the antiferromagnetic Bragg peak intensity on cooling through $T_c = 24$ K. A somewhat similar reduction in intensity has been observed in association with an electronic phase-separation phenomenon in La$_{2-x}$Sr$_x$CuO$_4$ with $x \sim 0.015$.\cite{25} In the latter case, the lost intensity reappears as diffuse incommensurate scattering; however, we do not see such a response in our oxygen-doped sample. Instead, we suggest that the measurement indicates a reduction in the net volume of antiferromagnetic phase due to a superconducting proximity effect. Such a proximity effect may be necessary for the superconducting phase to propagate through the finely mixed insulating and metallic phases.

Behavior of this type, associated with quenched structural disorder, has been discussed recently by Dagotto and collaborators.\cite{21,22} Their model calculations show some general features of competition between two ordered phases placed on the background of intrinsic inhomogeneities. Extended to manganites, this model predicts a colossal magnetoresistance effect. In the case of underdoped cuprates at low temperatures with competing superconductivity and antiferromagnetism, the authors claim that one can expect a “colossal superconducting proximity effect.”\cite{22} Previously, such a phenomenon has been observed in underdoped YBa$_2$Cu$_3$O$_{6+\delta}$.\cite{14}

The rest of the paper is organized as follows. A description of experimental details is given in the next section. The results are presented and discussed in Sec. III. Our conclusions are summarized in Sec. IV.

II. EXPERIMENTAL DETAILS

As is now well known, the way of preparing La$_2$CuO$_{4+\delta}$ plays an important role with regard to the physical properties of this material and more especially its superconducting properties. Electrochemical oxidation has been shown\cite{8} to be a reliable and controlled method to induce an oxygen surplus. Using this process, oxygen atoms can be inserted into La$_2$CuO$_4$ at interstitial positions of the La$_2$O$_2$ layers leading to the formation of mobile holes of concentration $p \approx 2\delta$ in the CuO$_2$ planes. Our crystal was prepared differently from the usual electrochemical-oxidation method in which the intercalation process occurs in an aqueous solution at a temperature of 100°C and lower.\cite{8,9,10,12,13} The oxygenation of our sample was performed electrochemically in a NaOH melt at $T = 330$°C, with an electrical current density of 10 mA/cm$^2$ for ~47 minutes. These are the same conditions used for crystal L2 in Ref.\cite{24}. The oxygen excess $\delta$ was estimated to ~0.05.

In-plane resistivity measurements on our sample, Fig. 1 show semiconducting behavior. If this were a single-phase sample, then given the large magnitude of the resistivity at $T > T_c$, one would not expect it to go superconducting.\cite{27,28} The crystal was also characterized by SQUID magnetometry. The superconducting transition temperature was found to be $T_c = 24$ K (see inset in Fig. 1).

Elastic neutron scattering measurements were performed on the BT-2 triple-axis spectrometer at the NIST...
Center for Neutron Research. The sample was approximately 0.2 cm$^3$ in size. After orienting the crystal on a holder, it was put into an Al can with He exchange gas, and then mounted on the cold finger of a closed-cycle He refrigerator. All of the diffraction measurements were performed using an incident neutron energy of 13.7 meV, with either one or two pyrolytic graphite filters in the incident beam to suppress neutrons at harmonic wavelengths. The resolution was varied by changing the horizontal collimators, as necessary.

Raman scattering measurements were carried out in quasi-backscattering geometry using the 514.5 nm argon-laser line. A piece of the same sample used for the neutron measurements was mounted on the holder of a He-gas-flow cryostat. The incident laser beam of 10 mW power was focused onto a spot of 0.1 mm diameter on the $3 \times 3$ mm$^2$ sample surface area. The scattered light was analyzed with a DILOR XY triple spectrometer combined with a nitrogen-cooled CCD detector.

To check the influence of the sample surface condition, the Raman measurements were performed both on the as-grown surface and on the mirror-like surface obtained by polishing and chemically etching in isopropanole and acetone. As no significant differences were observed, we report here the measurements performed on the polished surface. We note that the penetration depth of the laser light is about 2000 $\mu$m. As is usually done in Raman studies of La-cuprates.

The crystal structure of stoichiometric La$_2$CuO$_4$ is orthorhombic ($D_{18h}$) at room temperature and tetragonal ($D_{4h}$) above about 515K. The orthorhombic distortion is small and this allows us to assume tetragonal symmetry as is usually done in Raman studies of La-cuprates. The $x$- and $y$-axes were taken to be along the CuO bonds, with $z$ perpendicular to the CuO$_2$ plane. Within the tetragonal point group $D_{4h}$, the $zz$ geometry couples to excitations of $A_{1g}$ symmetry, $xy$ to $B_{2g}$ symmetry, and $xx$ to the combination of $A_{1g}$ and $B_{1g}$ symmetries. If $x'$ and $y'$ denote axes rotated by 45° from $x$ and $y$, then $x'y'$ geometry allows coupling to excitations with $B_{1g}$ symmetry and $x'x'$ geometry to a combination of $A_{1g}$ and $B_{2g}$ components. The spectra reported here were measured in the $xx$, $xy$, $x'y'$, and $x'y'$ scattering configurations.

III. RESULTS AND DISCUSSIONS

A. Structural data

For single crystals prepared electrochemically in an aqueous solution of NaOH, an ordering of the oxygen interstitials in oxygen-rich phases has been observed. The interstitials separate into domains such that, in a given domain, interstitials are present only in every $n^{th}$ layer. This “staged” structure is easily identified by the presence of specific superlattice peaks. We expected that it should be possible to identify the oxygen content of the intercalated phase by characterizing the superlattice peaks. Surprisingly, neutron diffraction measurements on the present crystal revealed an absence of such superlattice peaks at all temperatures. On the other hand, measurements of fundamental Bragg peaks indicate the presence of two phases, with lattice parameters consistent with oxygen-rich and oxygen-poor phases.

The presence of two phases is most clearly indicated by (00l) reflections, and we have focused on the (006). Although two distinct peaks cannot be resolved, the observed peak is quite broad and asymmetric. Fitting with two gaussian peaks yields the $c$ lattice parameters shown in Fig. 2(a). These results are consistent with previous studies of phase-separated samples. The phase with the larger $c$ lattice parameter should be the oxygen-rich phase. The difference between lattice parameters changes little with temperature, suggesting that the oxygen phase separation temperature, $T_{ps}$, is well above 300 K.

The relative intensities of the two fitted (006) peaks are shown as a function of temperature in Fig. 3. Assuming similar structure factors, there are comparable amounts of the two phases, with slightly more of the oxygen-rich phase. The temperature dependence of the relative intensities might indicate changes in the volume fractions; however, it is also possible that the changes are due to relief from extinction associated with strain. For the oxygen-rich phase, the widths of the (002), (006), and (008) Bragg peaks increase with momentum trans-
fer, indicating a substantial amount of strain. This strain is expected to have an impact on the phase separation and oxygen ordering dynamics; it indicates a significant amount of quenched disorder. If we assume that the fitted peak widths obtained at (002) are due purely to particle-size broadening (ignoring resolution and strain), then we get a lower limit of 600 Å for the diameters of the oxygen-poor and oxygen-rich domains.

A scan through (200)/(020) shows two peaks. (These reflections are seen along the same direction because of twinning.) One should expect to see three peaks due to the difference in orthorhombicity found previously for the oxygen rich and poor phases. The observed peaks are slightly asymmetric at low temperature, but a fit with more than two peaks is not stable over a broad temperature range. The lattice parameters obtained from the 2-peaks fits are shown in Fig. 2(b). The error bars reflect the distribution of lattice parameters determined from the peak widths after correcting for the calculated resolution. The width of the b distribution is comparable to the splitting in b for the two phases found by Radaelli et al. The temperature dependence of structural superlattice peaks at (012) appears to be consistent with a very gradual decrease in orthorhombic splitting with increasing temperature.

B. Onset of insulating state and features of magnetic order

Raman spectra of La$_2$CuO$_{4.05}$ at different temperatures in the $x'x'$ and $x'y'$ polarization configurations are shown in Figs. 1 and 3, respectively. For both configurations, one can see very strong two-phonon scattering at low temperature, but very little at 295 K. (Note that there are no single phonon excitations above $\sim 700$ cm$^{-1}$.) In addition, one can see strong two-magnon scattering of $A_{1g}$ symmetry at $\sim 3000$ cm$^{-1}$ in the $x'x'$ configuration with two-phonon scattering.

In Raman studies of La$_{2-x}$Sr$_x$CuO$_4$, Sugai and coworkers have shown that strong two-magnon and two-phonon scattering peaks appear together in the antiferromagnetic-insulator phase for $x \approx 0$, but both are dramatically reduced in doped phases with $x \gtrsim 0.06$. Thus, it appears that the two-phonon signal, along with the two-magnon scattering, is a useful indicator of the insulating phase. To test the temperature dependence of

FIG. 3: Ratio of integrated intensities of the two (006) peaks vs. temperature.

FIG. 4: Raman spectra of La$_2$CuO$_{4.05}$ at 5, 200, and 295 K in $x'x'$ polarization configuration. The inset shows the magnetic susceptibility $\chi$ of the sample.

FIG. 5: Raman spectra of La$_2$CuO$_{4.05}$ at 5 and 295 K in $x'y'$ polarization configuration. The inset shows the Raman response function $\chi''(\omega)$ obtained by dividing the original spectra by the Bose-Einstein thermal factor $[1 - \exp(-\hbar\omega/k_BT)]^{-1}$. The downturn at low frequency is a signature of the superconducting state.
this signature, we performed measurements on the insulator La$_{1.69}$Nd$_{0.31}$CuO$_4$. The temperature-dependent intensities of several two-phonon peaks, relative to the intensity of the single-phonon peak at 428 cm$^{-1}$, are plotted in Fig. 4. We see that there is only a modest decrease in the intensity ratio between 5 K and 295 K.

The two-phonon intensity ratios for the La$_2$CuO$_{4.05}$ sample are also plotted in Fig. 6. Compared to the insulator, the relative two-phonon signal decreases more rapidly with increasing temperature, dropping rapidly toward zero near 250 K. We interpret these results as indicating the absence of an insulating phase for $T > T_r \approx 250$ K, and the rapid appearance of the insulating phase for $T < T_r$.

To detect the onset of antiferromagnetic order, we rely on neutron scattering measurements. Figure 6 shows a scan along $Q = (1, 0, l)$ through the magnetic Bragg peak at $l = 0$, measured at $T = 9$ K. Besides the Bragg peak, there is also broad diffuse scattering along the antiferromagnetic rod. The temperature dependence of the (100) magnetic peak is shown in Fig. 8. The Néel temperature appears to be approximately 210 K, which is significantly lower than $T_r$. This magnetic ordering temperature is also lower than the more typical value of $\sim 250$ K found for the hole-poor phase in two-phase samples, indicating that the quenched disorder causes some frustration of the phase separation.

**C. Analysis of the phase separation transitions**

The neutron diffraction measurements indicate the presence of two phases near the two edges of the first miscibility gap, all the way up to 300 K. In previous studies, the oxygen-poor phase has been identified as an antiferromagnetic insulator, based on detection of antiferromagnetic order at lower temperatures; however, in our case the Raman spectra indicate that an antiferromagnetic insulating phase does not appear until the temperature drops below $T_r$. How can we reconcile these observations?

Due to long-range Coulomb interactions, it is clear that some degree of electronic phase separation must occur concomitantly with the development of two phases with different interstitial concentrations. At the same time, the antiferromagnetic phase of lightly-doped La$_2$CuO$_4$ is quite sensitive to hole content, with $T_N$ dropping rapidly to zero at $p = 0.02$ and the transport properties varying rapidly with $p$ in this regime. Hence, the oxygen-poor phase at $T > T_r$ may correspond to $p \gtrsim 0.02$. On cooling...
through $T_c$, the electronic energies favoring expulsion of holes from the antiferromagnetic phase become important and drive a slight increase in the degree of phase separation, resulting in the onset of detectability of signatures of the antiferromagnetic phase. We presume that the slight adjustment of interstitial segregation at this point was masked in the diffraction experiments by the large strain effects.

The idea that the initial, high-temperature phase separation is driven largely by lattice energies rather than electronic energies is consistent with a recent study of room-temperature structural phase diagrams in the related system La$_{2-x}$Sr$_x$NiO$_{4+\delta}$. With $x = 0$, pure La$_2$NiO$_{4+\delta}$ is known to have a complicated phase diagram with multiple miscibility gaps. If these phase separations were driven dominantly by electronic energies, then on co-doping with Sr, one would expect the phase boundaries to scale with the net hole content, $p = x + 2\delta$. To the contrary, diffraction measurements on numerous samples clearly show that the phase boundaries depend almost exclusively on $\delta$ (at least for $x < 0.08$). Thus, it appears that the lattice strain energies associated with interstitial oxygens must play an important role in the phase separation phenomena.

### D. Competition between superconductivity and antiferromagnetism

The variation of the (100) magnetic peak intensity at low temperature is shown in the inset of Fig. 3. On cooling, the intensity appears to hit a plateau at about 35 K, and then below $T_c$ it decreases. This effect looks similar to that observed in antiferromagnetic La$_{2-x}$Sr$_x$CuO$_4$ with 0.01 $< x < 0.02$, and in oxygenated La$_2$CuO$_3$ with extremely low Néel temperature ($T_N \sim 90$ K). However, in contrast to those observations, a corresponding increase in the diffuse magnetic scattering was not observed in our case.

The drop in the (100) magnetic peak intensity at low temperature is quite remarkable. We propose that the observed effect may involve a change in the relative volume fraction of metallic and insulating phases for temperatures below the superconducting phase transition. At such low temperatures the mobility of excess oxygen is negligible, so the changes should be largely electronic in nature. It appears that a form of superconducting proximity effect may be involved. The model calculations of Burgy et al. indicate that such a phenomenon may occur when a system with quenched disorder is below a normal-state percolative threshold. We note that the in-plane resistivity of our crystal (Fig. 1) looks qualitatively quite similar to calculated curves in Fig. 4(b) of Ref. [24].

### E. Features of the superconducting state

Since the discovery of high-temperature superconductivity, many experimental techniques, such as quasiparticle tunneling, angle-resolved photoemission, microwave absorption, IR and Raman spectroscopy, etc., have been used with the aim to clear up the nature of the superconducting gap (SCG). Raman scattering established itself as a powerful method to study this problem. At room temperature all cuprates exhibit a broad and rather flat background that is usually attributed to electronic excitations. As the temperature decreases below $T_c$, the frequency distribution of the electronic background changes reflecting the opening of the SCG. This renormalization consists of the formation of a broad peak associated with the pair-breaking process at the energy $2\Delta$ together with the decrease of the scattering intensity at energies lower than $2\Delta$. Unlike conventional superconductors with an isotropic gap, where no scattering is expected for frequencies $\omega \leq 2\Delta$, the low frequency depletion in high-temperature superconductors (HTSC’s) is not complete. Moreover, the peak frequency positions and renormalization of the scattering intensity below $T_c$ is different for $A_{1g}$, $B_{1g}$, and $B_{2g}$ symmetry components. An appropriate choice of the polarizations of incident and scattered light in the RS experiment can reveal the SCG anisotropy and the symmetry of the order parameter.

Most of the RS experiments imply that the SCG in HTSC’s is strongly anisotropic. Such a behavior was examined by different authors in the framework of a $d$-wave $s + id$ or anisotropic $s$-pairing. On the other hand, some experiments performed mainly on single CuO$_2$-layered compounds suggest the presence of an isotropic $s$-wave type gap. Up to now there is no definite consensus how to interpret these conflicting results. Furthermore, it is not clear whether the SCG anisotropy depends on doping level. Raman scattering experiments on Tl$_2$Ba$_2$CuO$_{6+\delta}$ and Bi$_2$Sr$_2$CaCu$_2$O$_{8+\delta}$ showed that overdoped samples exhibit a symmetry independent gap in contrast to the anisotropic SCG in the near optimum doping case. A SCG anisotropy was observed in YBa$_2$Cu$_3$O$_{7-\delta}$ with different doping levels. The anisotropy of the SCG and variation of the maximum in the electronic RS with scattering selection rules have been discussed in detail in a recent paper of M. Opel et al. It was pointed out that $d$-wave symmetry is widely accepted but complications arise both in the underdoped and overdoped ranges of the phase diagram. It is thus of interest to check the redistribution of the electronic continuum below $T_c$ in the metallic phase of the sample that undergoes phase separation.

In our experiments on La$_2$CuO$_{4.05}$ a frequency redistribution of the electronic continua is observed in all scattering geometries investigated when the temperature decreases below $T_c = 24$ K. In order to evaluate the $A_{1g}$ scattering component, we subtracted the $xy$ spec-
La$_2$CuO$_{4.05}$, but it should be noted that we also do not observe a $\omega^4$ law in the $B_{1g}$ scattering component (see Fig. 9 and inset in Fig. 9). The data shown here are very consistent with the experimental results of Ref. 51; however, in our study the situation is complicated due to phase separation. As discussed above and shown in Fig. 1 for temperatures below $T_c$ we detect both the Raman signal characteristic for the insulating as well as the superconducting phases. Moreover, we expect the oxygen concentration of the oxygen-rich regions to vary significantly, consistent with the strain detected by neutron diffraction.

**IV. CONCLUSION**

In summary, we have carried out Raman and elastic neutron scattering experiments on an excess-oxygen-doped La$_2$CuO$_4$ single crystal. The neutron diffraction measurements indicate that there are both oxygen-rich and oxygen-poor phases present in the sample and that the oxygen phase separation temperature is well above 300 K. The absence of detectable staging behavior in oxygen-rich regions allows us to conclude that a significant amount of quenched disorder must be present in the sample. In the Raman scattering experiments, the La$_2$CuO$_{4.05}$ sample displays at room temperature a response similar to cuprates with substantial doping. Below the temperature $T_c \approx 250$ K, the RS spectra change drastically: features characteristic for the insulating state appear in the spectra. We have proposed that electronic energies cause an additional redistribution of interstitial oxygen at $T_c$, making the oxygen-poor phase more insulating and resulting in the appearance of the two-phonon and two-magnon scattering. The Néel temperature $T_N \approx 210$ K was estimated from the neutron scattering measurements. A decrease of the (100) magnetic Bragg peak intensity for $T < T_c$ is attributed to subsequent electron-density redistribution connected with a superconducting proximity effect. Below the superconducting transition temperature, two-magnon scattering, strong two-phonon scattering, and a change of the electronic continua (that occurs as a result of the superconducting gap opening) are observed in the RS spectra, simultaneously. Based on the polarization dependence of the electronic continua, we conclude that the SCG in La$_2$CuO$_{4.05}$ is anisotropic with an anisotropy ratio $\Delta(B_{1g})/\Delta(B_{2g})$ $\approx$ 1.9 for the CuO$_2$ plane. The peculiarities of the electronic RS, i.e., the low-frequency power law dependence of the different scattering components at low frequencies and their different peak positions, are found to be very close to those observed earlier for Li$_{2-x}$Sr$_x$CuO$_2$. 51
Acknowledgments

We thank E. Ya. Sherman for helpful discussions. This work was supported in part by NATO Collaborative Linkage Grant PST.CLG.977766 and INTAS Grant 96-0410. JMT is supported at Brookhaven by the U.S. Department of Energy’s Office of Science under Contract No. DE-AC02-98CH10886. We acknowledge the support of the National Institute of Standards and Technology, U.S. Department of Commerce, in providing the neutron facilities used in this work.

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