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EVIDENCE FOR ELECTRONIC LOCALIZATION IN YBa2-yLayCu3Ox

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Electrical resistivity, thermoelectric power, magnetic susceptibility and neutron scattering experiments on $YBa_{2-1}La_1Cu_3O_3$, with y=1, show that this compound is neither superconductive nor magnetically ordered but are consistent with it being an electronically localized system

In efforts to understand the physics of the RBa₂Cu₃O_x high temperature superconductors, substitutional studies have concentrated on interchanging rare earth R atoms [1] and replacing some of the Cu atoms with various transition elements [2]. Although these studies have proven useful in revealing aspects important for an understanding of high temperature superconductivity, surprisingly little efford [3] has been given to investigating the role that the Ba sites play in these materials. Because trivalent La is nearly the same size as divalent Ba, it is possible to prepare $RBa_{2-\nu}La_{\nu}Cu_{3}O_{\kappa}$ with y at least equal to 1. With this substitution the nominal valence of Cu is reduced from 2.3 (for y=0) to 2.0 (for y=1), assuming that the oxygen content $x \approx 7$ in both cases. Here we report characteristics of YBaLaCu₃O $_x$ that are distinctively different than those of YBa2Cu3Ox and suggest a substitutionally-induced transition to a localized state.

Polycrystalline samples of YBaLaCu₃O_x were prepared using standard ceramic techniques with starting materials of Y₂O₃, BaCO₃, La₂O₃, and CuO. The starting materials were thoroughly ground together, calcined in air, reground, fired at 1080° C in oxygen and slowly cooled to 400° C before being removed from the furnace. X-ray analysis of the resulting pellets showed a dominant ($\leq 90\%$) orthorhombic phase, characteristic of RBa₂Cu₃O_x materials, with lattice parameters a=3.840 Å, b=3.856 Å and c=11.55 Å. These values, compared to a=3.8203 Å, b=3.8855 Å and c=11.6835 Å for YBa₂Cu₃O_{6 9} [4],

indicate that La addition reduces the orthorhombiticity. The presence of a minority ($\gtrsim 5\%$) phase that could be indexed to Y2Cu2O5 and at least one additional unidentified phase (≤5%) also was detected. The oxygen content of these samples was not determined, although a similar preparation procedure is known to give $x \approx 7$ in YBa₂Cu₃O_x. In this regard, however, a crude analysis of the c-lattice parameter change with La substitution in terms of observations by Segre et al. [3] for LaBa_{2-x}La_xCu₃O_{7+o} suggests that our sample may be slightly oxygen deficient. Sections from the pellets were used for electrical resistivity, thermoelectric power and magnetic susceptibility measurements. Several large pellets, combined weight ~50 g, were powdered for magnetic neutron scattering experiments at the National Bureau of Standards reactor. The susceptibility of random pieces from the neutron samples all showed the same temperature dependence.

The inverse magnetic susceptibility $1/\chi$ is shown in fig. 1 as a function of temperature. The Curie-Weiss behavior and maximum in χ at 12 K (see inset) suggest the possibility of antiferromagnetic order. Neutron diffraction at low temperatures failed to reveal any evidence for an antiferromagnetic transition in the majority phase YBaLaCu₃O_x. However, clear evidence for antiferromagnetic order at 12 K was found in the second phase Y₂Cu₂O₅. Recent susceptibility measurements on this compound by Troc et al. [5] show a maximum in χ at 13 K and at higher temperatures $\chi(T) \propto \mu_{\rm eff}^2/(T-38)$ where

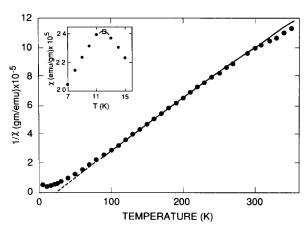


Fig. 1. Inverse magnetic susceptibility $1/\chi$ versus temperature of a sample of nominal composition "YBaLaCu₃O_x". The solid line is that expected if 8.5% of the sample weight were Y₂Cu₂O₅. The inset shows the presence of a maximum in χ at 12 K that is also consistent with 8.5% Y₂Cu₂O₅ (open square symbol).

 $\mu_{\rm eff} = 2.81 \, \mu_{\rm B}$ /formula unit. An analysis of the data in fig. 1 in light of their observations leads to the conclusion that 8.5% of the mass of our sample is Y₂Cu₂O₅ and that the remaining fraction has a temperature independent, or at most a weakly temperature dependent, susceptibility of magnitude $\chi_0 \approx 1.6 \times 10^{-7}$ emu/g. Results of this one-adjustableparameter analysis are shown by the solid line in fig. 1 and by the square in the inset, which are in good agreement with the data and with the estimate obtained by X-ray analysis of the fractional amount of $Y_2Cu_2O_5$ present. Assuming that χ_0 is due to a Paulilike contribution from YBaLaCu₃O_x implies that the effective density-of-states at the Fermi energy in YBaLaCu₃O_x is about 1/4 that of the high- T_c superconductor YBa₂Cu₃O_x [6].

The electrical resistance increases monotonically with decreasing temperature and shows no evidence for a superconducting transition above 4 K or for magnetic ordering in $Y_2Cu_2O_5$. In fig. 2 we plot the logarithm of the resistance as a function of $T^{-1/4}$. The linear variation from 9 < T < 125 K covers over three orders-of-magnitude increase in resistance and is highly suggestive of transport by three-dimensional (3D) variable-range hopping [7]. Taking $\ln R \propto (T_0/T)^{1/4}$ yields a characteristic temperature $T_0 \approx 5 \times 10^5$ K, which is comparable in magnitude to values of T_0 found [8] in oxygen-deficient poly-

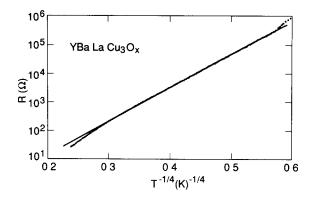


Fig. 2. Resistance of YBaLaCu₃O_x on a logarithmic scale as a function of temperature to the -1/4 power

crystalline samples of La₁₈Sr₀₂CuO_{4-δ}. Such behavior is contradistinctive to the linearly temperature dependent resistance observed in polycrystalline YBa₂Cu₃O₇ and is indicative of the significant role that Ba atoms play in the conduction process. Although we emphasize that the precise temperature dependence may be influenced by unknown anisotropy effects [9], YBaLaCu₃O_x is not unique in displaying localized-like transport: the resistivity in the Cu-O planes of single crystalline La₂CuO₄ is that expected of correlated 2D or 3D variable-range hopping [9]. For non-superconducting RBa₂Cu₃O_x samples in which $x \le 6.2$ or in which 5 at% Zn has been substituted for Cu, one finds respectively variable-range hopping [10] and weak-3D localizationtype conductivities [11].

The thermoelectric power S (fig. 3) of this ma-

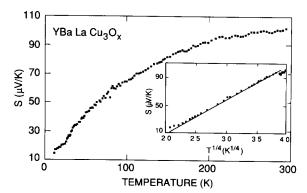


Fig. 3 Thermoelectric power S versus temperature T for YBaLaCu₃O_x. The inset shows that S is a linear function of $T^{1/4}$ for the temperature interval 25 < T < 220 K

terial is large and positive, which in a single-band model implies that entropy transport is dominated by a small number of hole-like carriers. The room temperature value of S is nearly 50 times larger than in RBa₂Cu₃O₇ [6] and comparable to that found in oxygen-deficient samples of this material [10]. Unlike the nearly temperature independent thermopower of polycrystalline high- T_c compounds [6], we find $S \propto T^{1/4}$ for 25 < T < 220 K, roughly the same temperature interval over which $\ln R \propto T^{-1/4}$. For 3D variable-range hopping, S should be proportional to $T^{1/2}$ [7,12], clearly not the temperature dependence observed here. However, as in the case of the electrical resistance, anisotropy effects could distort the intrinsic temperature dependence.

Although conductivity in the high- $T_{\rm c}$ superconductors is generally regarded as being dominated by the Cu-O planes/chains, it is apparant from these results that changes in valence and spatial order of the Ba-site atoms can alter significantly the transport characteristics. In all cases of which we are aware, the approach to a localized-like state (at the expense of superconductivity) is accompanied by a decrease in the state-density at the Fermi level, whether it be produced by oxygen depletion [10], Cu substitution [11] or, in this case, a replacement for Ba, as well as the introduction of disorder on one of these sites. Clearly, these observations should be considered in attempts to raise $T_{\rm c}$ above 100 K.

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