THERMOELECTRIC POWER IN THE NORMAL STATE OF HIGH-$T_c$ SUPERCONDUCTORS RBA$_x$Cu$_3$O$_y$ (R = YTTRIUM OR RARE EARTH)

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Received 31 July 1987

We report measurements of the thermoelectric power $S$ and resistivity $\rho$ of high-$T_c$ superconductors RBA$_x$Cu$_3$O$_y$. For $R = Y$ or a rare earth and $x = 7$, $S$ is positive and nearly temperature independent but its magnitude can vary depending on $R$. We have systematically varied the oxygen stoichiometry ($6 < x < 7$) and find that changes in $x$ have a far more dramatic effect on $S$ and $\rho$ than does $R$. When $x$ approaches 6.5, there is a cross-over to new behavior with $S$ approximately linear in temperature. This metallic-like behavior does not appear in the resistivity. The thermopower results suggest that RBA$_x$Cu$_3$O$_y$ compounds have strong on-site Coulomb correlations.

Among recent measurements on high-$T_c$ copper-oxide superconductors, to the best of our knowledge, no systematic study of the thermopower has been reported. We have performed thermopower measurements on RBA$_x$Cu$_3$O$_y$ (where $R = Y$, Eu, Gd, Ho) and studied the effect on resistivity as well as thermopower due to oxygen deficiency. This study was motivated in part by the large discrepancies between several groups reporting measurements on the normal state properties of nominally comparable materials [1]. We believe that several factors complicate the issue: chemical stoichiometry, the density of polycrystalline material, and expected anisotropy of the transport properties. Hall effect experiments [2] indicate that the carrier concentration changes with oxygen stoichiometry suggesting that other transport properties may be affected similarly.

All samples were prepared by conventional ceramic powder techniques. To vary the oxygen stoichiometry, we heated samples of the same batch to 300–600°C in various oxygen pressures for 2 to 24 hours, then allowed the sample to cool slowly to room temperature. Assuming the weight loss incurred by heating resulted solely from the removal of oxygen provided a semiquantitative estimate of the oxygen content.

The electrical resistivity was measured by a standard dc 4-probe method and the Seebeck effect was detected through copper reference wires. The absolute thermopower of the sample was obtained by correcting the measured Seebeck coefficient with the absolute thermopower of copper at various temperatures. We used a differential Au–0.07%Fe/Cromel–P thermocouple to detect the temperature gradient of $\approx 0.5$ K across the sample. Our apparatus was calibrated with constantan wires and the measured Seebeck coefficient agreed with NBS values to within 5% accuracy.

The thermopower $S$ of RBA$_x$Cu$_3$O$_y$ samples is shown in fig. 1. That $S$ is positive suggests conduction is dominated by holes, which is confirmed by Hall effect measurements [2, 3]. The room temperature values of $S$ are similar to that predicted by the free electron approximation if one allows the carriers to be positively charged. ($S_{d} = \pi^{2}k^{2}T/ee_{F} \approx 10 \mu$V/K with $e_{F} \approx 2$ eV). However, this is not consistent with the temperature independence of $S$ above $T_c$, which for a metal is expected to increase linearly with temperature. Within experimental uncertainty, $S$ goes to zero below $T_{c}$, as expected.

A first glance at fig. 1 might suggest that $S$
increases as the mass of R increases. However, the oxygen stoichiometry of these compounds has far more influence on the thermopower and resistivity because the values can vary significantly from one batch to another. We have systematically varied the oxygen stoichiometry of YBa$_2$Cu$_x$O$_y$ and EuBa$_2$Cu$_x$O$_y$ for 6 < x ≤ 7. The results for both are similar and we discuss our observations in the Eu compound below.

The thermopower and resistivity are plotted in fig. 2 for the starting material EuBa$_2$Cu$_x$O$_y$ (where x ≈ 7.0 as determined by chemical analysis). As x decreases (see fig. 3), the resistivity increases and the mid-point $T_c$ decreases.

For $x < 6.45$, the resistivity increases drastically as a power law of decreasing temperature. The mid-point $T_c$ and resistivity at 250 K are shown in the inset of fig. 3. It is clear that the scattering mechanism changes near $x = 6.45$ which is about the stoichiometry where an orthorhombic–tetragonal structural phase transition occurs [4].

The thermopower for EuBa$_2$Cu$_x$O$_y$ (where 6 < x ≤ 7) is plotted in fig. 4 with different scales. As $x$ decreases, $S$ increases and shows a slight temperature dependence above $T_c$ for $x > 6.45$ [5]. For 6.1 < $x$ < 6.45, the thermopower shows a metallic-like behavior, i.e. $S = AT + BT^3$ for $T < 40$ K, while the resistivity behaves otherwise. The values of $S$ at room temperature have a similar dependence on oxygen stoichiometry as the Hall coefficient [2]. At $x = 6.1$, the thermopower is semiconductor-like with a large negative value at room temperature. The sign change at $x = 6.1$ suggests that conduction is now dominated by electron-like carriers for $T > 100$ K.

It has been proposed [3] that the Hubbard model well describes the transport properties of these compounds. We plot $S$ at 250 K in the inset of fig. 4 along with the Heikes expression [6] for a system with strong on-site electron–electron repulsion, i.e. $S = (k/e) \ln[2(1 - \rho)/\rho]$, where $\rho$ is the number of free electrons per site. Here, $\rho = (13 - 2x)/3$ per Cu-site for $x < 6.5$ and $\rho =$
1 - (2x - 13)/3 per Cu-site for x > 6.5. We see that Heikes expression qualitatively accounts for our observations, including the sign change in $S$.

In conclusion, the resistivity and thermopower depend strongly on the oxygen stoichiometry and little on various rare earth substitutions. A simple two-band model seems inadequate to explain the temperature independence of $S$ together with the observed effects of oxygen stoichiometry. Further, thermopower measurements suggest that these compounds have strong on-site Coulomb correlations.

Acknowledgement

Work at Los Alamos was performed under the auspices of the US Department of Energy.

References

[5] Qualitatively similar behavior is reported on YBa$_2$Cu$_3$O$_{6.5}$ by S. Uchida et al. (preprint).