

Superconductivity in a New YBaCuO Compound at 105 K.

I. KIRSCHNER(*), J. BÁNKUTI(*), M. GÁL(**), K. TORKOS(**), K. G. SÓLYMOS(***)
and G. HORVÁTH(*)

(*) *Department for Low Temperature Physics, Roland Eötvös University
1088 Budapest, Puskin u. 5-7, Hungary*

(**) *Department of General and Inorganic Chemistry, Roland Eötvös University -
1088 Budapest, Muzeum krt. 6-8, Hungary*

(***) *Department of Petrology and Geochemistry, Roland Eötvös University
1088 Budapest, Muzeum krt. 4/a, Hungary*

(received 27 April 1987; accepted in final form 5 June 1987)

PACS. 74.70. - Superconducting materials.

PACS. 74.70N - Dirty superconductors.

Abstract. - A superconducting transition has been detected in a $(Y_{0.8}Ba_{0.2})(CuO_{4-\delta})_2$ compound by electrical and magnetic measurements. The resistivity begins to decrease at 173 K and the zero-resistivity state sets in at 105 K.

The experiences of investigation of high- T_c superconductors [1-7] led us to search the possible highest- T_c superconductors among Y-Ba-Cu-O compounds. The optimum value of the ionic radii, of the substitution of lanthanides (L) with alkaline earths (AE) and that of the ratio of L + AE to Cu and the redox processes of Cu were taken into account during our experiments [8].

A new kind of Y-Ba-Cu-O system was developed with nominal composition of $(Y_{0.8}Ba_{0.2})(CuO_{4-\delta})_2$, which was prepared by solid-state reaction from pure chemicals of Y_2O_3 , $BaCO_3$ and CuO.

During previous experiments we found that the lower molar ratio of (Y + Ba):Cu improves the superconducting parameters, so it was chosen as 1:2 in this case. The pulverized and homogenized mixture was heated gradually over 3 hours to 925 °C, found to be the optimum temperature for the solid-state reaction by thermogravimetry. After 18 hours the sample was quenched within 15 minutes. Powder X-ray patterns (Siemens D-500) on reacted and pulverized mixture show a microcrystalline structure. The powder was then pressed into a pellet on 20 kbar and sintered once more. Photographs taken by a scanning electron microscope (Jeol JXA-50A) show that the pellet consists of a medley of different crystalline shapes.

Samples of dimension $(0.7 \times 1 \times 8)mm^3$ were cut out of the pellet. A circular diamond saw was used and when the temperature rose to somewhat above 100 °C the sample was cooled with water. We mention this detail since it may be relevant to the sample's behaviour. The

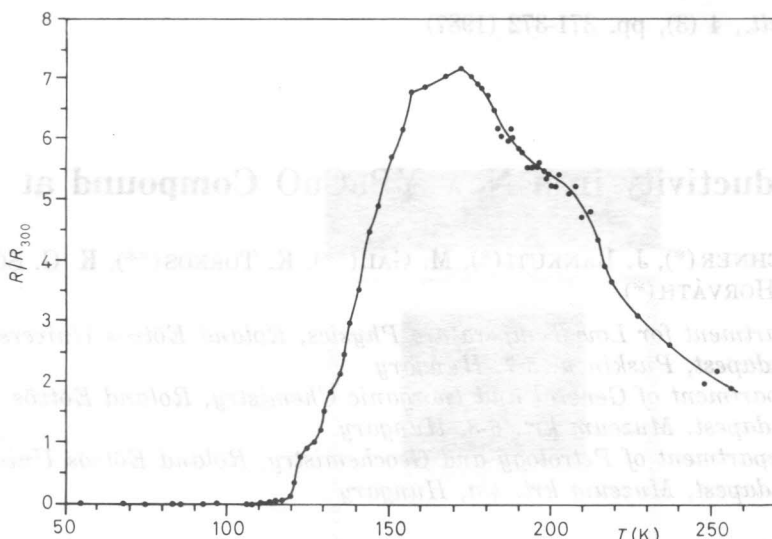


Fig. 1. - Dependence on temperature of the resistance of a specimen $0.7 \times 1 \times 8$ mm. $R_{300} = 39.52 \Omega$, $\rho_{300} = 350$ m Ω cm.

experiments were done over a 48 h period, a week after the preparation of the sample. About thirty cyclings between room temperature and 77 K left the behaviour unchanged.

The dependence of the resistivity R on temperature T was measured at $10 \mu\text{A}$, $100 \mu\text{A}$ and 1 mA. The data obtained are independent of the current in this range. The smallest measurable voltage signal was 10 nV and so the sensitivity of the measurement of specific resistivity ρ is $8.8 \cdot 10^{-8} \Omega$ cm.

The temperature dependence of the resistance is shown in fig. 1. There is a kink in the curve at 173 K, the sharp drop starts at 156 K and the zero-resistance state appears at 105 K. The measurements were repeated 4 days later: the onset temperature of the rapid drop increased to 159 K and zero resistance appeared at 101 K. After 3 weeks the sample showed no superconductivity: the resistance increased monotonously with falling temperature. Details of the sample preparation will be further studied.

Magnetic measurement carried out at 77 K show a partial penetration of the external field into the sample and so it hints at a type-II superconductivity. Placing the sample in a low magnetic field of 762 Oe, about 21 vol.% is in Meissner's state at this temperature.

REFERENCES

- [1] BEDNORZ J. G. and MÜLLER K. A., *Z. Phys. B*, **64** (1986) 189.
- [2] BEDNORZ J. G., TAKASHI M. and MÜLLER K. A., *Europhys. Lett.*, **3** (1987) 379.
- [3] CHU C. W., HOR P. H., MENG R. L., GAO L., HUANG Z. J. and WANG Y. Q., *Phys. Rev. Lett.*, **58** (1987) 405.
- [4] CAVA R. J., VAN DOVER R. B., BATLOGG B. and RIETMAN E. A., *Phys. Rev. Lett.*, **58** (1987) 408.
- [5] WU M. K., ASHBURN J. R., TORNG C. J., HOR P. H., MENG L., GAO L., HUANG Z. J., WANG Y. Q. and CHU C. W., *Phys. Rev. Lett.*, **58** (1987) 908.
- [6] HOR P. H., GAO L., MENG R. L., HUANG Z. J., WANG Y. Q., FORSTER K., VASSILIOU J., CHU C. W., WU M. K., ASHBURN J. R. and TORNG C. J., *Phys. Rev. Lett.*, **58** (1987) 911.
- [7] UCHIDA S., TAKAGI H., KITAZAWA K. and TANAKA S., *Jpn. J. Appl. Phys.*, **26** (1987) L-1.
- [8] KIRSCHNER I., BÁNKUTI J., GÁL M., TORKOS K., SÓLYMOS K. G. and HORVÁTH G., *Europhys. Lett.*, **3** (1987) 1309.