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PROCESS PARAMETERS, ORIENTATION, AND FUNCTIONAL PROPERTIES OF MELT-PROCESSED BULK Y-Ba-Cu-O SUPERCONDUCTORS*

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ABSTRACT

This study compared the microstructure, texturing, and functional properties (critical currents) of YBa₂Cu₃O_{7-x}-based bulk pellets that were prepared by the quench-melt-growth-process (QMGP), melt-textured growth (MTG), and conventional solid-state reaction (SSR) approaches. Using two X-ray diffraction (XRD) methods, θ -2 θ , and rocking curves, we found that the individual grains of two melt-processed pellets exhibited remarkable preferred orientational alignment (best rocking curve width = 3.2°). However, the direction of the preferred orientation among the grains was random. Among the three types of bulk materials studied, the QMGP sample was found to have the best J_c values, ≈4500 A/cm² at 77 K in a field of 2 kG, as determined from SQUID magnetic data.

INTRODUCTION

Since its discovery in 1987, $YBa_2Cu_3O_{7-x}$ has probably been the most extensively studied superconductor [1]. $YBa_2Cu_3O_{7-x}$ single crystals exhibit

better homogeneity. transition temperature (T_c). critical current density (J_c), and preferred orientation than bulk samples. Usually, in the bulk ceramic superconductors, J_c is orders of magnitude lower (150–600 A/cm² at 77 K) than it is in a single crystal (5 x 10⁶ A/cm² at 77 K) [2,3]. Weak links are typical for bulk superconductors and include dislocations, impurities, and high–angle grain boundaries that often hinder the possibilities of obtaining higher J_c values. Generally, the melt–textured–growth (MTG) approach is to heat bulk YBa₂Cu₃O_{7-x} (123) above its peritectic temperature (1010°C) for a few minutes and then rapidly cool the pellets to different temperature plateaus, followed by slow cooling to ≈950°C. The quench–melt–growth process (QMGP) is very similar to the MTG approach, except for the high–temperature (1300°C) quenching step that is included at the beginning to ensure that the Y₂BaCuO₅ (211) phase is finely distributed.

The sequence of events during premelting and subsequent solidification have been extensively studied by Rodriguez et al. [4] and Golden et al. [5]. The high-temperature reactions of 123 showed the development of a phase, tentatively identified as 211, along with the 123 phase at temperatures between 700 and 940°C. Golden et al. [5], found that even materials quenched from temperatures as low as 950°C contained a phase mixture. This indicated the presence of a metastable liquid at temperatures of 950°C and above. The end product of MTG usually contains mixed phases of 211, BaCuO₂, and CuO. Some evidence indicates that the 123 phase does not nucleate at 211 particles in Y-deficient liquid [6].

A very dense and preferentially aligned microstructure obtained by melt processing can give rise to dramatically improved J_c , and significantly reduced

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field dependence. In the MTG polycrystalline 123, J_c is usually $\approx 5000 \text{ A/cm}^2$, at H = 0 and T = 77 K.

In this detailed investigation, we have studied, for the first time, the preferred orientation of each grain in melt-processed 123 samples. We show that the melt-processed samples exhibit preferred orientations comparable to those in single crystals.

EXPERIMENTAL METHOD

The powders Y_2O_3 , BaCO₃, and CuO were mixed, ground, and heated in a tube furnace with flowing oxygen gas. The total oxygen pressure inside the furnace was maintained at 3.3 mm Hg with a vacuum pump. During cooling, the pressure inside the furnace was raised to ambient pressure at 850°C, and further cooling was continued in the flowing oxygen at ambient pressure [7].

The first of the three pellets was made by QMGP. Approximately 20 g of the above-described powder was quenched from 1350°C. The powder was then pressed into \approx 3-cm-diameter pellets under 50 MPa pressure. It was then melt processed in air according to the following scheme: the pellet was ramped from 25°C at the rate of 200°C/h to 1040°C. Then it was ramped at a rate of 60°C/h to 1060°C, and held for 1 h. It was cooled at 500°C/h to 1000°C, at 3°C/h to 900°C, and finally, at 60°C/h to 25°C. The pellet was then oxygenated at \approx 550°C for 24 h in 100% oxygen gas and slowly cooled to room temperature.

The MTG pellet was produced with the same powder and procedure used for the QMGP pellet, except that the powder was not quenched from 1350°C. All melting and cooling experiments were carried out in air under ambient conditions. Oxygenation was performed in 100% oxygen gas flow.

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-3-

The third, and conventional, pellet was made by solid-state reaction (SSR) from the same $YBa_2Cu_3O_7$ precursor powder used for the others. The pellets were sintered at 945°C for 10 h, oxygenated at ~550°C, and slow cooled in 100% oxygen.

RESULTS AND DISCUSSION

The specimens for X-ray and magnetic characterization were cut in the form of cubes or identical parallelepipeds from three bulk samples, identified as MTG, QMGP, and SSR. X-ray diffraction measurements were carried out on all of the faces of the specimens as shown in Figs. 1 and 2. The rather narrow peaks in the X-ray diffraction patterns (Fig. 1), and their high intensities, suggest that rather large grains are present in the MTG pellet. The degree of orientation in the QMGP pellet has been investigated in detail with rocking curve measurements similar to the ones shown in Fig. 2 for MTG.

The MTG sample exhibited excellent preferred orientation, as indicated by a thorough analysis of the data obtained at each face of the cut sample (Figs. 1 and 2). The best preferred orientation was determined to be on Side C of the rectangular piece and this is shown in Fig. 2 by the rocking curve distribution width, full width half maximum (FWHM), of 3.2°. This outstanding preferred grain orientation is comparable to that which is usually seen only in YBa₂Cu₃O_{7-x} single crystals with FWHM ~8–10° [8]. Bulk YBa₂Cu₃O_{7-x} commonly exhibits a preferred orientation in the range of 20–30° [8]. The θ –2 θ X–ray diffraction pattern for MTG shows that the c–axis preferred orientation is in the 001 direction. However, this preferred orientation differs in its direction on each side of this small piece with dimensions 3.0 x 3.2 x 5.1 mm.

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On side A, which is parallel to side C, the preferred orientation is in two directions, corresponding to the 110 and 220 planes (Fig. 2). The similar preferred orientation occurs on parallel sides D and B. On side D, the θ -2 θ and rocking curve XRDs show that the preferred orientation in the 200 plane has a narrow range of \approx 6.4° (Figs. 1 and 2). On side B, the θ -2 θ and rocking curves XRDs show 200– and 103-plane orientation in the range of 3.1° and 3.0°, respectively.

We found that our samples had different preferred orientations within 1 mm, but with high intensities, suggesting that the individual large grains are well oriented, but with deviations among them. This result is similar to that from single crystals. One important difference is that, in our study, we looked at all the faces of the specimen, which is not usually the case for studies reported in literature [9,10]. It may be that differences, on a micrometer (μ m) scale, do exist in single crystals as well and what has been reported is simply an average FWHM.

We do observe some inhomogeneity in the samples, which may be caused by other phases that are present. This is common for melt–processed 123 materials, because they are cooled from above the peritectic temperature. The domain, grain, and defect phase dimensions were measured by optical microscopy (Fig. 3). The domains of the MTG sample were of \approx 500 µm, and the 211 phase (in the shape of rods) in the samples was unoriented, randomly distributed, and \approx 10–15 µm long and 5 µm wide (Fig. 3a). However, CuO inclusions were quite rare (<1%) and predominately found at domain edges, near defects and/or impurities. The size of the CuO inclusions depended greatly on the crack spacing (some cracks and pores were present in most of these materials, whereas SSR samples were usually somewhat porous, as

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indicated by a density that was lower than the expected ideal one), but most were $\approx 5 \ \mu m$.

The QMGP sample texture was similar to that of the MTG sample; however, the domains were a bit larger (600 μ m). For the QMGP sample, slightly fewer 211–phase particles were observed and they were smaller, i.e., on the order of 5–10 μ m long and 3 μ m wide (Fig. 3b). These results are similar to those reported by Murakami et al. [11], who observed that the average size of a 211 particle was ≈2 μ m in diameter. In the QMGP sample, the CuO particles were rare and found in the same places (cracks) where they were observed in the MTG sample. From these results, we can suggest that, in the MTG samples, the 211 phase solidified first in the 123 matrix and then the liquid CuO moved into the thermal-gradient cracks to solidify later.

Both melt-processed samples exhibited a highly continuous matrix of 123 phase, whereas, in contrast, the conventionally prepared sample exhibited a somewhat porous matrix of 123 phase. The size of the 123 grains was on the order of \approx 5–6 µm (Fig. 3c).

SQUID magnetic measurements have been used to determine superconducting properties such as T_c and J_c . The T_c values of the QMGP, MTG, and conventional pellets were 92, 91.5, and 91 K, respectively, as derived from magnetization data measured during the warming runs after cooling the samples in a zero field. The J_c values at 77 K were determined from the width of the hysteresis loops at an applied field of 2 kG. Figure 4 shows the SQUID hysteresis loop measured at 77 K for the QMGP sample.

For the melt-textured bulk material, J_c values in the range of 4000–17,000 A/cm² at H = 0, and 77 K have been reported [8,12]. The QMGP sample in our

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study exhibited J_c values of $\approx 4500 \text{ A/cm}^2$ at H = 2000 G, and 77 K. The J_c of the MTG sample was $\approx 1100 \text{ A/cm}^2$ at 77 K (H = 2000 G). The slightly higher J_c of the QMGP sample was probably due to fewer and smaller 211 particles embedded in the 123 matrix.

For all three pellets, the J_c values were calculated from hysteresis loops at 77 K, with a modified version of Bean's J_c model [13], and the widely accepted formula [11] for a square sample of side–length d given by

$$J_c = 30 \frac{\Delta M}{d}$$
 (Gauss).

The high J_c of the melt-textured samples is thought to be caused by single-crystallike properties such as large grain size with excellent preferred orientation within each large grain.

SUMMARY

In this paper, we compared the orientation and functional properties of QMGP, MTG, and conventionally processed (SSR) bulk 123 superconductors. The melt-processed samples exhibited preferred orientation characteristics in each grain, with a narrow distribution of orientation (rocking curve XRD $\approx 3.2^{\circ}$ wide) comparable to that in single crystals (8–10°). The preferred orientation of each grain is excellent in all directions even though the direction of the preferred orientation changes slightly from grain to grain. Such a detailed study of the preferred orientational fluctuations in all directions has not been reported in the past for QMGP samples [8,11,12]. The XRD peaks are narrow and intense because, in our samples, the domains are large ($\approx 500 \ \mu m$). The QMGP has the highest J_c of the three bulk samples and it is $\approx 4500 \ A/cm^2$, at

H = 2000 G and T = 77 K, most probably because of fewer and smaller particles of 211 phase in the 123 matrix, as compared to the MTG sample.

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Figure captions

- Fig. 1. Four largest θ -2 θ XRD spectrums of the MTG sample. Each side has different large peaks, suggesting different preferred orientations.
- Fig. 2. Rocking curve XRD spectrums of three sides of MTG YBa₂Cu₃O_{7-x} sample. Each side has narrower (FWHM ≅3.2–6.4°) curves than are found in most single crystals (8–10°).
- Fig. 3. Optical photomicrographs of (a) MTG, (b) QMGP, and (c) solid-state sintered YBa₂Cu₃O_{7-x} bulk pellets. Note that fewer and smaller 211
 (light colored) particles are present in the QMGP sample.
- Fig. 4. SQUID hysteresis loop at 77 K for QMGP YBa₂Cu₃O_{7-x} sample.







Figure 3.



Figure 4.