



Effect of template post-annealing on Y(Dy)BaCuO nucleation on CeO₂ buffered metallic tapes



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ABSTRACT

Substrate engineering is very significant in the synthesis of the high-temperature superconductor (HTS) coated conductor. Here we design and synthesize several distinct and stable Cerium oxide (CeO₂) surface reconstructions which are used to grow epitaxial films of the HTS YBa₂Cu₃O_{7-δ} (YBCO). To identify the influence of annealing and post-annealing surroundings on the nature of nucleation centers, including Ar/5%H₂, humid Ar/5%H₂ and O₂ in high temperature annealing process, we study the well-controlled structure, surface morphology, crystal constants and surface redox processes of the ceria buffers by using X-ray diffraction (XRD), X-ray photoelectron spectroscopy (XPS) and field-emission scanning electronic microscopy (FE-SEM), respectively. The ceria film post-annealed under humid Ar/5%H₂ gas shows the best buffer layer properties. Furthermore, the film absorbs more oxygen ions, which appears to contribute to oxygenation of superconductor film. The film is well-suited for ceria model studies as well as a perfect substitute for CeO₂ bulk material.

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1. Introduction

The second-generation (2G) high-temperature superconductor REBa₂Cu₃O_{7-δ} (REBCO, RE=Y and Gd, Dy etc rare earths) coated conductors (CCs) wires have now entered into the marketplace reproducibility. However, the current density of long-length 2G wire production still needs to be better optimized [1–3]. In order to improve the critical current density of YBCO, various techniques have been developed, including improving grain connectivity, introducing high density nano-size defects, and increasing effective superconducting thickness. Besides, superconducting properties of REBCO are significantly affected by surface morphology and crystallinity of buffer layer. It has been found that CeO₂ is a most important material in heterogeneous catalysis due to its high oxygen storage capacity (OSC) based on the oxidation and reduction of cerium ions [4,5]. It is reported that CeO₂ has large oxygen storage capacity because oxygen vacancy defects (CeO_{2-x}) can be rapidly formed and eliminated [6]. And the cubic fluorite structure of CeO₂ is also great interesting due to its outstanding properties such as wide band gap, high thermal and chemical stability, high dielec-

tric constant and unique optical properties, making CeO₂ films a promising material for various applications, such as buffer layers gate dielectric, fuel cells [7–9], sunscreen cosmetics [10], catalysis [11] and ultraviolet blocking filter [12–16]. CeO₂ has been successfully used as cap buffer layer for preventing the interdiffusion and enhancing the compatibility between REBCO and the metallic substrate, for its small lattice mismatch with REBCO, chemical tenability, self-assembling crystalline structure. Usually the structure and nuclei density of YBCO are influenced by the well defined substrate characteristics and processing conditions. Many methods have been used to optimize CeO₂ cap layers, such as UV light irradiation, better synthesize parameters, rare earth doping and post-annealing. Post-annealing is a simple and promising route to optimize the quality of CeO₂. However, there are few reports on post-annealing of CeO₂ under various atmosphere. In this work, we mainly investigate the influence of post-annealing under different processing atmospheres on CeO₂ buffers as well as the subsequent superconducting Y(Dy)BCO layers prepared by metal organic deposition.

2. Experimental

Nano-crystalline CeO₂ thin films around 40 nm are grown on long-length IBAD tape: un-textured Hastelloy alloy metal tape

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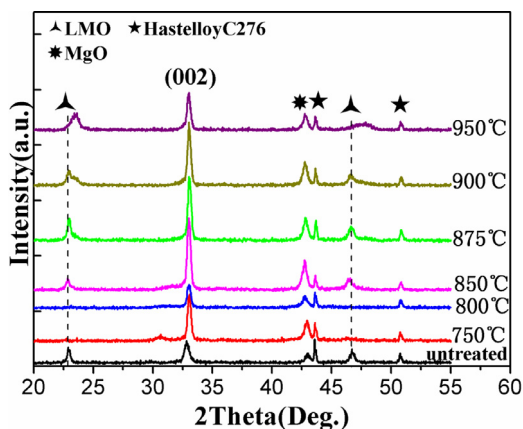


Fig. 1. X-ray θ – 2θ diffraction patterns of post-annealed CeO₂ films under different temperatures.

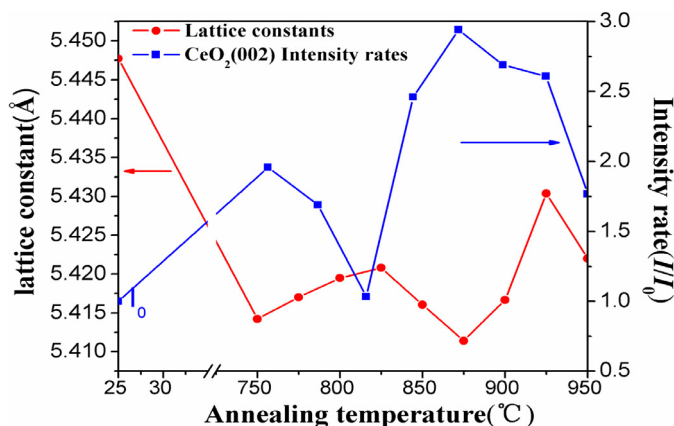


Fig. 2. The dependence of CeO₂ lattice parameters and CeO₂(002) peak intensity with different temperatures.

buffered with Al₂O₃/ Y₂O₃/ IBAD-MgO/ Epi-MgO/ LaMnO₃. Cerium oxide template buffers, which annealed at 800 °C is approximately 40 nm thick, were deposited using a large-area magnetron sputtering deposition system on LaMnO₃ layers. After deposition, CeO₂ buffers are post-annealed ex-situ at different temperatures. It is found that the crystalline quality of CeO₂ is sensitive to annealing temperature from Fig. 1.

By comparing the influence of different annealing temperatures on lattice parameters and peak intensity of CeO₂, we attempt to manipulate the stabilization process for optimization of annealing temperature region in Fig. 2.

The crystallite size of the films, D, was calculated by Scherrer's formula as: $D = K\lambda / (\beta \cos \theta)$, Where $k = 0.9$ is the Scherrer constant, $\lambda = 1.5406 \text{ \AA}$ is the wavelength of X-ray diffraction. β is a full width at half maxima of (002) peak, and θ corresponds to the Bragg angle.

Considering both peak shift of LaMnO₃(LMO) and lattice constants of CeO₂, we optimized different post-annealing temperatures. The dependence of lattice parameters of CeO₂ peak intensity ratio of CeO₂ on post-annealing temperatures are shown in Fig. 2. As shown in Figs. 1 and 2, no obvious peak shift of LaMnO₃(LMO) and excellent crystallinity of CeO₂ is observed for the specimen post-annealed at 875 °C. It was reported that high oxygen partial pressure can decrease the particle energy, posing a hazard for film qualities, but other results argue the variation in oxygen partial pressure has no effect on the crystalline orientation of CeO₂ layers [17]. So we did not pay more attention to it. A significant improvement of CeO₂ crystallinity is obtained for annealing temperature

at 875 °C. Thus we fixed post-annealing temperature at 875 °C and turned to varying post-annealing atmospheres. The Cerium oxide are post-annealed for 60 min at 875 °C in various flowing gas [18] as follows: (a) Ar-5%H₂, (b) humid Ar-5%H₂, (c) O₂, with the corresponding samples termed S1,S2,S3 respectively and the specimen without post-annealing is named S0.

Finally the Y(Dy)BCO layers about 1 μm are deposited on the above ceria buffers via a low fluorine metal organic deposition (MOD) process. The synthesize and heat-treatment process for low-fluorine Y(Dy)BCO films have been reported in our previous work [19–21].

The structure and the texture of the grown films of CeO₂ with and without post-annealing are characterized by glancing incidence X-ray diffraction θ – 2θ scans and ω -scans (Philips X'Pert PRO, CuK α radiation operated at 40 mA and 40 kV). The morphology, valence state, microstructure and size of CeO₂ are evaluated by scanning electron microscopy(SEM), atomic force microscopy(AFM) and X-rays photoelectron spectroscopy(XPS). The spectroscopic measurements of samples are carried out in a ultra-high-vacuum system (5×10^{-8} mbar) with an analysis chamber. And the measurements (here Al K α irradiation used at an excitation voltage of 14.5 KV and an emission current of 18.15 mA, scan area is $500 \mu\text{m} \times 500 \mu\text{m}$) scan deeper layer < 10 nm without annealing in XPS chamber. The surface of Y(Dy)BCO profile is tested by metallographic microscope. The J_c of Y(Dy)BCO films were measured at 77 K by an inductive measurement system from Theva. The critical current (I_c) of the coated conductors were obtained via multiplying critical current density (J_c) by the corresponding cross-sectional area (S).

3. Results and discussion

Comparing with the conventional evaluation system about cap layer, the present mechanism manifests the surface quality and the lattice mismatch with the substrate, or the cap layer in the case of multilayered structures influencing the interfacial nucleation phenomena of Y(Dy)BCO grains strongly [22–26].

In order to distinguish the lattice mismatch with Y(Dy)BCO and surface quality of the post-annealing CeO₂, the post-annealed samples are analyzed by different characterization methods.

Fig. 3 (a–d) show the high-resolution reciprocal space mapping(HR-RSM) of the CeO₂ (002) deposited at different ambient flowing gas. After post-annealed, the surface of buffer layer occurs adsorption and separation reaction with the atoms of different gases. The gas atoms and atoms from surface buffer layer exchange electrons, forming chemical bond. Therefore, the out-of-plane texture of ceria oxide undergoes different changes. The HR-RSM results show that the CeO₂ films post-annealed under different atmosphere exhibit different growth morphology. Fig. 3b and d display broader RSMs range in the 2Theta direction, revealing poorer crystallization of the film post-annealed under such atmosphere. When the CeO₂ layer was post-annealed in the surrounding of humid Ar-5%H₂ gas, the RSMs show a considerably converged pattern in the 2Theta and omega directions, demonstrating humid Ar/5%H₂ gas is suitable for the regrowth of CeO₂ layer on the LMO substrate. Based on the above analysis, the structural optimization of CeO₂ film post-processed in wet Ar/5%H₂ flowing gas overwhelms that of other films. Furthermore, the HR-RSM area of the CeO₂ post-annealed in humid Ar/5%H₂ atmosphere exhibits smallest overall c-axis tilts.

The ϕ -scans are carried out to evaluate the in-plane texture of CeO₂ with and without being post-annealed. Four symmetric peaks average full width at half-maximum (FWHM) values about the ϕ -scan patterns are shown in Table 1, indicating all the CeO₂ films have strong biaxial texture. Obviously the structure of S2 is slightly better than other samples and more suitable for subse-

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