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BaFe₂As₂/Fe bilayers with [001]-tilt grain boundary on MgO and SrTiO₃ bicrystal substrates

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Abstract

Co-doped BaFe₂As₂ (Ba-122) can be realized on both MgO and SrTiO₃ bicrystal substrates with [001]-tilt grain boundary by employing Fe buffer layers. However, an additional spinel (i.e. MgAl₂O₄) buffer between Fe and SrTiO₃ is necessary since an epitaxial, smooth surface of Fe layer can not be grown on bare SrTiO₃. Both types of bicrystal films show good crystalline quality.

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1. Introduction

In high- T_c cuprates, [001]-tilt grain boundaries (GBs) in MgO and SrTiO₃ bicrystal substrates have been commonly used for realizing Josephson junctions as well as investigating transport properties of GBs. After the discovery of Fe-based pnictide superconductors,[1] the community became immediately interested in exploring grain boundaries in this new class of superconductors.

Despite the recent success of realizing Co-doped BaFe₂As₂ (Ba-122) on [001]-tilt GB MgO and SrTiO₃ bicrystal substrates,[2-3] these substrates may not be suitable for several reasons. Firstly, a large misfit between Ba-122 and MgO of around 6 % leads to low crystalline quality or even non-epitaxial growth under our deposition condition.[4] Secondly, the SrTiO₃ substrate becomes electrically conductive under ultra-high vacuum (UHV) conditions, which may compromise transport measurements particularly near the normal-superconducting transition. However, implementation of Fe for MgO and Fe+MgAl₂O₄ for SrTiO₃ as buffer layers can solve those problems. An Fe layer itself is electrically conductive, however, the recalculation of the superconducting transition temperature (T_c) and the critical current density (J_c) is possible due to its thin layer thickness (20 nm).[5] In this report, the fabrication of Co-doped Ba-122 thin films on both [001]-tilt bicrystal MgO and SrTiO₃ substrates and their transport properties are presented in detail. For the aim of bicrystal Josephson junctions, substrates with a large [001]-tilt angle of over 20° have been used in this investigation.

2. Experimental procedure

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2.1. Ba-122 on [001]-tilt MgO bicrystal

A 20 nm thick Fe buffer layer was deposited by means of pulse laser deposition (PLD) on a [001]-tilt MgO bicrystal substrate with misorientation angle of $\theta_{GB}=36.8^\circ$ (nominal value) at room temperature, followed by a high-temperature annealing at 700°C . This process is very effective for obtaining epitaxial Fe layers with flat surface. After the Fe buffer preparation, Co-doped Ba-122 ($\text{BaFe}_{1.8}\text{Co}_{0.2}\text{As}_2$) with 100 nm thickness was deposited at 700°C . All deposition processes have been conducted under UHV condition. A detailed description of both Fe buffer and Co-doped Ba-122 deposition procedure can be found in Ref. [6]. After the structural characterization described in sub-section 2.3, Au layers were deposited on the films by PLD at room temperature followed by ion beam etching to fabricate bridges of 0.5 mm width and 1 mm length for transport measurements.

2.2. Ba-122 on [001]-tilt SrTiO_3 bicrystal

A MgAl_2O_4 buffer was deposited on a [001]-tilt GB SrTiO_3 bicrystal substrate with $\theta_{GB} = 30^\circ$ at 650°C by PLD in UHV condition (pulse number 4000 at 40 mJ). In a pilot experiment, we have confirmed that MgAl_2O_4 can be grown epitaxially on SrTiO_3 (100) substrate. After the primary buffer deposition, the MgAl_2O_4 -buffered SrTiO_3 substrate was cooled to room temperature for the secondary, Fe buffer deposition. A 20 nm thick Fe buffer was prepared by the same procedure as described in sub-section 2.1 except for the annealing temperature. Here, a slightly higher temperature of 750°C was employed. After the Fe buffer preparation, a 100 nm thick Co-doped Ba-122 ($\text{BaFe}_{1.84}\text{Co}_{0.16}\text{As}_2$) layer was deposited. *In-situ* gold layer deposition at room temperature was conducted after the Ba-122 deposition for the aim of protection from possible damage during the structuring process.

2.3. Structural characterization and transport measurements

Phase purity and out-of-plane texture were investigated by means of X-ray diffraction in Bragg-Brentano geometry with Co-K α radiation. In-plane orientation of both Fe and Co-doped Ba-122 were investigated by using the 110 and 103 poles, respectively, in a texture goniometer operating with Cu-K α radiation. The respective measured reflection of the MgO and SrTiO_3 bicrystal substrates for in-plane texture measurements are 220 and 110.

Superconducting properties were measured in a Physical Property Measurement System (PPMS, Quantum Design) by a standard four-probe method with a criterion of $1 \times 10^{-4} \text{ V cm}^{-1}$ for evaluating J_c .

3. Results and discussion

XRD patterns in logarithmic scale (fig. 1 (a)) clearly show that the film has been grown with high phase purity and *c*-axis texture, i.e. with [001] perpendicular to the substrate surface. Fig. 1 (b) shows the ϕ -scans of the 103 Ba-122, the 110 Fe and the 220 MgO. It is evident from fig. 1 (b) that two grains with $\theta_{GB} \sim 37.1^\circ$, which is slightly larger than the nominal value of 36.8° , are grown due to the perfect transfer of the grain orientation from the substrate via Fe buffer to the Ba-122 layer. Here the respective average $\Delta\phi_{\text{Ba-122}}$ and $\Delta\phi_{\text{Fe}}$ are 0.84° and 0.80° (not corrected for instrumental broadening), indicative of good in-plane alignment for both layers. The GB is symmetric, i.e. it is situated with $\theta_{GB}/2$ to (100) in both grains.

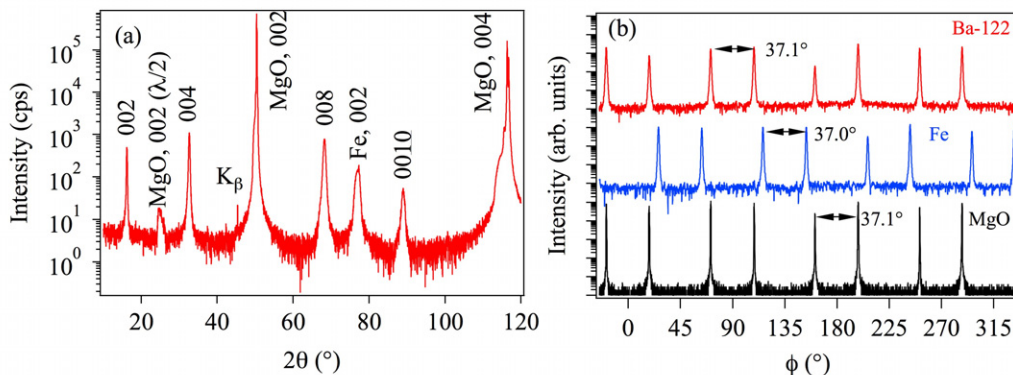


Fig. 1. (a) The $\theta/2\theta$ scan of Ba-122/Fe bilayer on [001]-tilt GB MgO bicrystal substrate in Bragg-Brentano geometry using Co-K α radiation. (b) The ϕ -scans of the 103 Ba-122, the 110 Fe and the 220 MgO. Average misorientation angles are 37.1° , 37.0° , and 37.1° for Ba-122, Fe and MgO, respectively.

R - T curves in fig. 2 (a) show that both the inter- and intra-grain bridge have the same onset T_c as well as zero resistance, indicating the high quality of the bicrystal films. The field dependence of $J_c(B||c)$ for both the inter- and intra-grain bridge at various temperatures is displayed in fig. 2 (b). Clearly, the intra-grain J_c (J_c^{intra}) is higher than the inter-grain J_c (J_c^{inter}) at low magnetic fields regime due to the large θ_{GB} . The difference between J_c^{intra} and J_c^{inter} is getting smaller with increasing applied field and finally both curves overlap at the irreversibility field for $\theta_{GB} \sim 37^\circ$. On the assumption that the weak-link behavior is empirically described by $J_c^{\text{inter}} = J_c^{\text{intra}} \exp(-\theta_{GB}/\theta_0)$ in low field regime, the characteristic angle (θ_0) is estimated to around 7.9° for our Co-doped Ba-122 bicrystal films (fig.2 (c)), which is almost twice as large as that of $\text{YBa}_2\text{Cu}_3\text{O}_7$ [7].

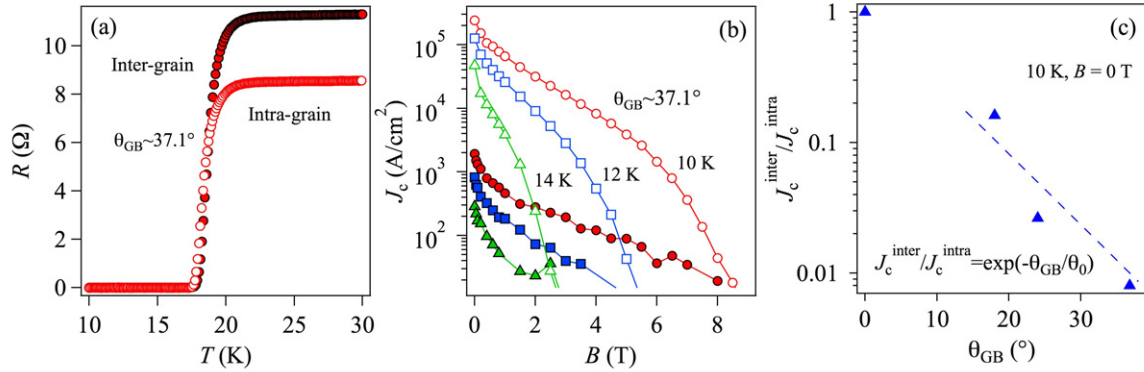


Fig. 2. (a) R - T curves of inter- and intra-grains. Both curves show the same T_c of around 20 K. (b) J_c - B performances ($B||c$) of the corresponding grains at different temperatures. Open symbols represent the intra-grain and the solid symbols show the inter-grain curves. (c) Normalized critical current density ($J_c^{\text{inter}}/J_c^{\text{intra}}$) as a function of θ_{GB} at 10 K.

The growth of epitaxial Fe with smooth surface on SrTiO_3 substrate is not straightforward, which is presumably due to the difference in crystal structure rather than to the lattice mismatch. As a result, epitaxial growth of Ba-122 is hardly achieved on Fe-buffered SrTiO_3 . Indeed, fiber textured Fe is observed (fig. 3 (a)), when it is prepared directly on SrTiO_3 substrate with the same procedure as on MgO substrates. In order to avoid this problem, an additional buffer layer of MgAl_2O_4 between Fe and SrTiO_3 substrate has been used. MgAl_2O_4 fits better to accommodate the lattice mismatch to Fe rather than MgO as shown in Table 1. Here, the lattice mismatch is defined as $(a_f - a_s)/a_f$, where a_f and a_s are the lattice parameters of Fe and substrates, respectively. Additionally, a small lattice mismatch of 3.4 % between MgAl_2O_4 and SrTiO_3 may lead to good epitaxial growth of the MgAl_2O_4 buffer.

Fig. 3 (b) shows the 110 pole figure of Fe on MgAl_2O_4 -buffered SrTiO_3 fabricated by the same procedure described in sub-section 2.1. It is clear from fig. 3 (b) that a perfect textured growth of Fe is realized (average $\Delta\phi_{Fe} = 0.61^\circ$). During heating of Fe, evolution of the Fe-layer texture was observed through reflection high energy electron diffraction (RHEED), which is a similar observation of Fe on single crystalline MgO substrates.[8] The RHEED images of the Fe layer on MgAl_2O_4 -buffered SrTiO_3 after the high temperature annealing showed only streak patterns, indicative of surface smoothing. This architecture also prevents the current-shunting effect between SrTiO_3 and Co-doped Ba-122. This buffer architecture offers the opportunity to grow smooth epitaxial Fe layers on various perovskite substrate such as $(\text{La},\text{Sr})(\text{Al},\text{Ta})\text{O}_3(100)$ substrate (not shown in this paper).

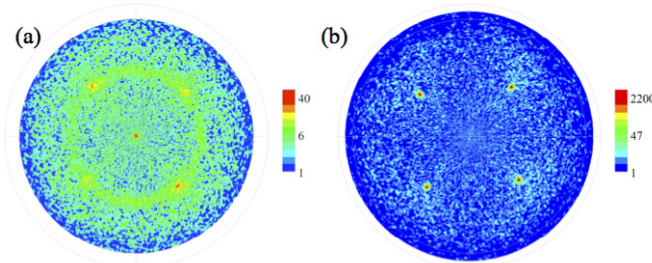


Fig. 3. (a) The 110 pole figure measurement of Fe on bare SrTiO_3 and (b) MgAl_2O_4 -buffered SrTiO_3 .

Table 1. The crystal structure, lattice parameter and the lattice mismatch to Fe of MgO and MgAl₂O₄.

Materials	Crystal structure	Lattice parameter (nm)	Lattice mismatch (%)
MgO	Rock salt	0.421	3.7
MgAl ₂ O ₄	Spinel	0.8083 ($a/2=0.4042$)	-0.4

Structural characterization of Ba-122/Fe/MgAl₂O₄ on [001]-tilt GB SrTiO₃ substrate by means of X-ray diffraction is summarized in figs. 4. As stated earlier, *in-situ* Au layer deposition was conducted after the Ba-122 deposition, which explains the 111 reflection of Au in fig. 4 (a). It is further clear from fig. 4 (a) that only 00/ reflections of Ba-122 with a small amount of the 110 reflection are observed. Fig. 4 (b) displays the ϕ -scans of the 103 Ba-122, the 110 Fe and the 110 SrTiO₃. Clear 8 peaks from two grains with a rotational angle of 30° are apparent. Accordingly, an epitaxial Ba-122 film is formed on [001]-tilt SrTiO₃ bicrystal substrate. The onset $T_c = 24$ K of the film is almost identical to that of Ba-122/Fe bilayers on single crystal substrates. (fig. 4 (c)) This value is higher than the intra-grain T_c deposited on MgO bicrystal presented in fig. 2 (a) due to a higher deposition temperature as well as different PLD targets (i.e. different Co concentration). The resultant bicrystal junctions show clear Josephson effects and the detailed studies can be found in Ref. [9].

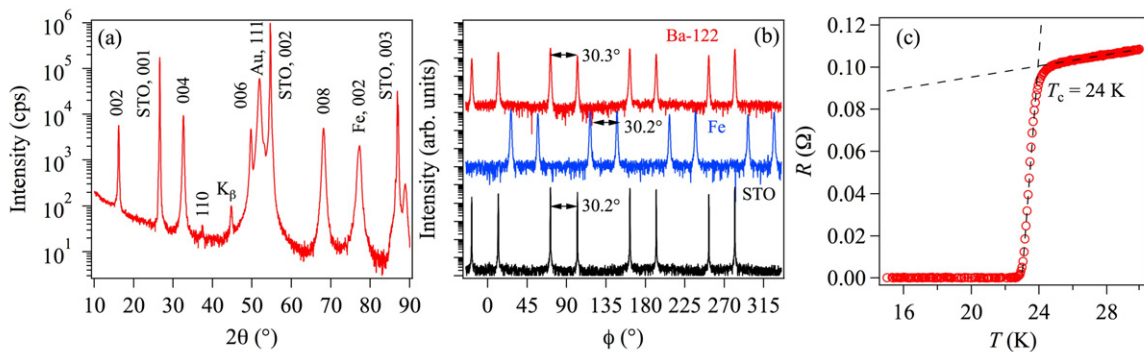


Fig. 4. (a) The $\theta/2\theta$ scan of Ba-122/Fe/MgAl₂O₄ trilayer on [001]-tilt SrTiO₃ (STO) bicrystal substrate in Bragg- Brentano geometry using Co-K α radiation. (b) The ϕ -scans of the 103 Ba-122, the 110 Fe, and 110 STO. Average misorientation angles are 30.3°, 30.2°, and 30.2° for Ba-122, Fe and STO, respectively. (c) Intra-grain T_c was recorded as 24 K.

4. Summary

High-quality Co-doped Ba-122 bicrystal films can be realized on both [001]-tilt MgO and SrTiO₃ substrates by employing Fe buffer layers via pulsed laser deposition. The additional MgAl₂O₄ buffer between Fe and SrTiO₃ is necessary for realizing epitaxial Ba-122. The characteristic angle of Co-doped Ba-122 in low field regime is about 8°, which is almost twice as large as that of YBa₂Cu₃O₇.

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