Magnetic orientation and magnetic anisotropy in paramagnetic layered oxides containing rare-earth ions

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Abstract
The magnetic anisotropies and easy axes of magnetization at room temperature were determined, and the effects of rare-earth (RE) ions were clarified for RE-based cuprates, RE-doped bismuth-based cuprates and RE-doped Bi-based cobaltite regarding the grain orientation by magnetic field. The easy axis, determined from the powder orientation in a static field of 10 T, depended qualitatively on the type of RE ion for all three systems. On the other hand, the magnetization measurement of the \(c\)-axis oriented powders, aligned in static or rotating fields, revealed that the type of RE ion strongly affected not only the directions of the easy axis but also the absolute value of magnetic anisotropy, and an appropriate choice of RE ion is required to minimize the magnetic field used for grain orientation. We also studied the possibility of triaxial grain orientation in high-critical-temperature superconductors by a modulated oval magnetic field. In particular, triaxial orientation was attempted in a high-oxygen-pressure phase of orthorhombic RE-based cuprates \(Y_2Ba_4Cu_7O_y\). Although the experiment was performed in epoxy resin, which is not practical, in-plane alignment within 3\(^\circ\) was achieved.

Keywords: rare-earth element, magnetic alignment, magnetic anisotropy, layered oxides

(Some figures in this article are in colour only in the electronic version)

1. Introduction
Functional compounds with anisotropic crystal structures, such as layered structures, generally exhibit anisotropic physical properties and possess functionally superior crystallographic structure. In particular, the effect of the anisotropy on electronic transport properties is a major factor determining the functionality of thermoelectric and superconducting materials. For applications of these layered functional compounds as bulk materials, a grain orientation process is intrinsically indispensable. Moreover, in the case of superconducting materials based on copper oxides, which are high-critical-temperature (\(T_c\)) cuprate superconductors, not only the \(c\)-axis grain orientation of superconducting \(CuO_2\) planes extending in the \(ab\)-planes but also the alignment of crystallographic axes in the \(CuO_2\) planes, that is, triaxial or biaxial grain orientation, is required for practical use.

Magnetic orientation is one of the most useful methods of grain orientation. The recent development of superconducting magnets has realized the 10 T class of high static fields in a room-temperature bore without using liquid helium, and these magnetic fields can provide a kinetic energy comparable to gravity and thermal energy at room temperature, even for feeble magnetic materials, paramagnetic and diamagnetic...
materials. Therefore, magnetic orientation using cryogen-free superconducting magnets is one of the most useful techniques for layered compounds with feebly magnetic properties at room temperature. The materials requirements for the achievement of magnetic orientation are sufficiently large magnetic anisotropy at room temperature and the desired easy axis of magnetization. Although understanding these magnetic properties at room temperature is important for grain orientation, the factors determining magnetic anisotropy and the easy axis in functional layered oxides and methods for controlling these properties have not been clarified yet.

In this study, we focus on three different layered oxide systems, REBa$_2$Cu$_3$O$_y$ (RE123), Bi$_2$Sr$_2$CaCu$_2$O$_y$ (Bi2212) and Bi$_2$Ca$_2$CO$_2$O$_y$ (BiCa222), the former two of which are high-$T_c$ cuprate superconductors with a CuO$_2$ layer and the latter is a candidate thermoelectric oxide with a CoO$_2$ layer. Figure 1 shows the crystal structures of these three compounds, all of which possess a layered structure with alternately stacked blocking layers and two-dimensional $TMO_2$ ($T$ = Cu and Co) layers. To clarify the possibility of control of the easy axis of magnetization and of the magnetic anisotropy between the in-plane and out-of-plane directions by RE ions, the magnetic roles of RE in RE123, RE-doped Bi2212 and RE-doped BiCa222 were investigated from the viewpoint of the magnetic susceptibility of magnetically oriented powders. Furthermore, we also discuss the possibility of formation of a triaxially oriented microstructure in high-$T_c$ cuprates by magnetic field. We focus on Y$_2$Ba$_2$Cu$_3$O$_y$ (Y247, $y \approx 15$), which is a superconducting compound related to Y123 and does not possess a twin microstructure in its grains. We also attempted to fabricate a triaxially oriented powder in epoxy resin using a modulated oval magnetic field [1]. This work is very important from the viewpoint of the practical use of RE-based cuprates because the symmetry of superconducting quasi-particles is $d$-wave, and because a change in the superconducting current depending on the misorientation angle at a single grain boundary emerges even for the in-plane direction [2].

2. Experimental details

Powders of RE-doped BiCa222, RE-doped Bi2212 and RE123 with an average grain size of 1–3 μm were synthesized by a solid-state reaction. Details of the synthesis conditions are as follows: Bi$_2$(Ca$_{0.5}$RE$_{0.5}$)$_2$CO$_3$O$_y$ [Bi(Ca,RE)222] powders were synthesized from Bi$_2$O$_3$, CaCO$_3$, RE$_2$O$_3$ and CoO$_2$ starting powders. After calcination at 750°C for 12 h in air and thorough grinding, mixtures with an appropriate cationic ratio were sintered at 820°C for 24 h in air. Ball-milling of the sintered powders was performed to prepare fine powders. Bi$_2$Sr$_2$(Ca$_{0.5}$RE$_{0.5}$)$_2$Cu$_2$O$_y$ powders were obtained from Bi$_2$O$_3$, SrCO$_3$, CaCO$_3$, RE$_2$O$_3$ and CuO starting powders. After calcination at 790°C for 12 h in air and thorough grinding, mixtures with an appropriate cationic ratio were sintered at 820°C for 24 h in air. RE123 powders were synthesized from RE$_2$O$_3$, BaCO$_3$ and CuO powders. Mixtures with an appropriate cationic ratio were sintered at 850–900°C for 36 h in flowing gas mixture 1%O$_2$ + 99%Ar for RE = La to Gd and Yb and at 880–900°C for 18 h in air for RE = Dy – Tm. The oxygen contents of the RE123 powders were controlled to $y \sim 6.9$ by post-annealing at 400°C in flowing oxygen.

For all the powders, magnetically aligned (MA) powder samples were fabricated in epoxy resin, which was cured for more than 12 h at room temperature under a static magnetic field ($H_s^{\text{static}}$) of 10 T. The weight ratio of the powder and epoxy resin was 1 to 10. The easy axis of magnetization for each compound was determined from the x-ray diffraction (XRD) pattern of the surface perpendicular to $H_s^{\text{static}}$ using each MA powder sample. For RE123 compounds with the easy axis in the ab-plane, c-axis oriented powder samples were prepared in a rotated magnetic field ($H_s^{\text{rotated}}$) of 12 T at room temperature. In this case, the powders in epoxy resin were horizontally rotated at 30 rpm in a transverse magnetic field of 12 T during the curing of the epoxy resin. The difference in magnetic susceptibility parallel to the c-axis ($\chi_c$) and in the ab-plane ($\chi_{ab}$) at room temperature ($\Delta \chi = \chi_c - \chi_{ab}$)
shows the XRD patterns of the MA samples. They were 4–20 times larger than that of Y123, and therefore the calculated \( \Delta \chi \) was determined using a SQUID magnetometer for the c-axis oriented powder samples of RE123. The mass of RE123 powders in epoxy resin was determined by the Archimedes method.

3. Results and discussion

3.1. Effects of RE-doping on magnetic anisotropy and easy axis of magnetization

To study the magnetic effects of paramagnetic RE ions at room temperature, the easy axes of magnetization and \( \Delta \chi \) values were determined. XRD patterns for the MA powder samples aligned in a static field reveal that RE123 compounds with RE = Y, La, Pr, Nd, Sm, Dy and Ho exhibited an easy axis parallel to the c-axis and those with RE = Eu, Gd, Er, Tm and Yb had an easy axis in the ab-plane. In particular, RE123 compounds with the easy axis in the ab-plane are not desirable for the evaluation of magnetic anisotropy between the in-plane and out-of-plane directions. This is because a certain level of \( \Delta \chi \) includes a component of the magnetic susceptibility normal to \( H^\text{mag} \) in these MA samples. Therefore, c-axis oriented powders are required. We attempted c-axis orientation using a rotated magnetic field, under which the hard axis of magnetization is aligned perpendicular to the \( H^\text{rotated} \) plane [3, 4]. As a result of such grain orientation, c-axis oriented powder samples were successfully fabricated for RE = Eu, Er, Tm and Yb, and it was found that the rotated field was effective for the c-axis orientation of RE123 compounds with \( \chi_c < \chi_{ab} \).

Figure 2 shows the relation between the number of 4f electrons and \( \Delta \chi \) for the RE123 compounds, which was determined from magnetization measurements in directions parallel to the c-axis and ab-plane using the c-axis aligned powders samples. Clearly, \( \Delta \chi \) largely depended on the type of RE, and \( |\Delta \chi| \) in RE123 with a heavy RE was larger than that in RE123 with a light RE. The absolute values of \( \Delta \chi \), \( |\Delta \chi| \), were on the order of \( 10^{-4} \) for RE = Dy, Ho, Er and Tm. They were 4–20 times larger than that of Y123, and therefore the calculated \( H^\text{a} \) for magnetic grain orientation in the case of Dy123 is approximately 5 times lower than that for Y123. This behavior has also been observed qualitatively in oxygen-reduced RE123 compounds with \( y < 6.5 \) [5], indicating that the roles of RE ions are basically preserved upon a change in the local crystal structure around the RE ions.

To investigate the universality of the magnetic roles of RE ions, RE doping was attempted for Bi2212 and BiCa222 as another superconducting cuprate and another functional layered oxide, respectively. Their easy axes of magnetization were revealed from the XRD measurement of these MA samples. Figure 3 shows the XRD patterns of the MA samples of Bi(Ca,RE)222, together with the patterns of the MA sample of RE-free BiCa222. In the case of RE-free BiCa222, only (200) and (020) peaks were enhanced by the static magnetic field. That is, the easy axis for RE-free BiCa222 lies in the \( ab \)-plane, parallel to the thermoelectric CoO2 layers. The easy axis is derived from the magnetic anisotropy of the CoO2 layers because cations other than Co ions, that strongly affect the magnetic properties, are not present in the system.

For the doping of RE = Y − Gd, the XRD patterns were roughly coincident with that of the RE-free sample. Therefore, the easy axis was preserved in the \( ab \)-plane, and 5 mol% of doping by RE did not affect the magnetic anisotropy and easy axis of magnetization in the case of doping of a light RE. However, for the doping of RE = Tb − Ho, the (200) and (020) peaks disappeared and only (001) peaks were seen. Even at a low doping level of RE, the easy axis was converted completely from the \( ab \)-plane to the c-axis direction. Moreover, in the case of RE = Er − Yb, clear enhancement of the (200) peak was observed, indicating that the magnetic anisotropy of RE123 was also enhanced, with the easy axis in the \( ab \)-plane (\( ab \)-plane magnetic anisotropy). These results mean that heavy RE ions affect the magnetic anisotropy and easy axis more than light RE ions. This clear difference in the magnetic roles of light and heavy RE ions is qualitatively consistent with the results for RE123 obtained from figure 2.

We investigated the effects of doping RE = Eu, Er, Tm and Yb, which had the ability to enhance the \( ab \)-plane magnetic anisotropy in BiCa222, on the easy axes for Bi2212 with the easy axis parallel to the c-axis. Figure 4 shows XRD patterns of the MA samples for RE-doped Bi2212 (RE = Eu, Er, Tm and Yb), together with the patterns for the MA sample of RE-free Bi2212. As mentioned above, the easy axis for RE-free Bi2212 is the \( c \)-axis. In the case of RE = Er and Tm, although RE doping induced a Bi2Sr2CuOy (Bi2201) phase, the (001) peaks disappeared and the (200) and (020) peaks were markedly enhanced. Namely, the easy axis was converted from the \( ab \)-plane to the c-axis by the doping of Er and Tm. However, in the case of RE = Eu and Yb, the 50% substitution of RE is insufficient for the complete conversion of the easy axis into the \( ab \)-plane. This is probably because the \( ab \)-plane magnetic anisotropies generated by the 50% doping of these two RE ions were smaller than or comparable to the intrinsic c-axis magnetic anisotropy of the CuO2 plane in RE-free Bi2212. This result also demonstrates that the ability to control the easy axis and magnetic anisotropy depended on the type of RE.
The easy axes of the three functional layered oxides obtained in this study are summarized in Table 1, also with the sign of the second-order Stevens factor, $\alpha_J$, [6, 7] for each RE ion. From the comparison between the directions of the easy axis in RE123 and the sign of $\alpha_J$, the easy axis was clearly correlated with the sign of $\alpha_J$ and can be explained in terms of the single-ion anisotropy of $\text{RE}^{3+}$. However, the absolute values of $\Delta \chi$ are significantly different between light and heavy RE ions, which is in contrast to the magnetic anisotropy of RE metals [8]. Note that the calculated magnetic anisotropy constants for the heavy and light RE were almost the same. This is presumably due to differences in the structural symmetry around the RE and the measurement temperature, meaning that it is important to experimentally evaluate the roles of RE ions for each material. The easy axes for RE-doped Bi2212 corresponded well with the result for RE123, whereas those for RE-doped BiCa222 were different, particularly in the case of doping of a light RE. This is due to the intrinsic difference in the easy axes and magnetic anisotropy among these RE-free compounds. As suggested from the RE-free, Y and La-doped samples, the easy axes for the CuO$_2$ and CoO$_2$ planes were the $c$-axis and an axis in the $ab$-plane, respectively. Therefore, for the doping of a light RE with small magnetic anisotropy, such as Pr or Nd, these $c$-axis magnetic anisotropies are insufficient to convert the easy axis of BiCa222 to the $c$-axis. On the other hand, in the case of a heavy RE, such as Tb, Dy or Ho, with $c$-axis magnetic anisotropy much larger than those of the light RE, even 5 mol% RE substitution can convert the easy axis effectively.

3.2. Triaxial orientation of Y247 by a modulated oval magnetic field

Polycrystalline Y247 was synthesized by a solid-state reaction under high oxygen pressure. Details of the synthesis process are described elsewhere [9]. Y247 powder with particles of mean diameter 6 $\mu$m was mixed with epoxy resin, and this mixture was cured for 8 h at room temperature in a modulated oval magnetic field. The weight ratio of the Y247 powder and epoxy resin is 1:10. Triaxial orientation by the modulated oval field was performed by the repetition of static and rotation
modes for 8 h. Details are as follows; in the rotation mode, the mixture was horizontally rotated at a rotation speed of 50 rpm for 15 min in a transverse field (\(H_a\)) of 12 T. Then, the rotation was stopped for 15 min and the sample stage was located such that the direction normal to the \(\alpha\)-plane (see inset of figure 5) overlapped with the \(H_a\) direction in each static mode.

Figure 5 shows XRD patterns of the \(\alpha\)-, \(\beta\)- and \(\gamma\)-planes (see inset) for the Y247 powders magnetically oriented using the modulated oval field, together with the pattern taken for the \(\delta\)-plane, which is perpendicular to the \(\alpha\)-plane and tilted at 45° relative to the \(\beta\)- and \(\gamma\)-planes. For the \(\alpha\)-plane, only the (001) peaks were observed, and the easy axis of magnetization was the \(c\)-axis. This is consistent with the result obtained for magnetic grain orientation using a static field of 12 T. When one focuses on the XRD patterns of the \(\beta\)- and \(\gamma\)-planes, the positions of reflection peaks, which are based on (001) or (020), were very close for both planes. However, a slight difference of approximately 0.3° was observed between the (200) and (020) peak positions. The difference coincided well with that obtained from a simulation by the Rietveld method [10]. That is, the main peaks at the \(\beta\)- and \(\gamma\)-planes were derived from the (020) and (200) reflections, respectively. The magnetic susceptibilities for the three crystallographic axes were related as \(\chi_c > \chi_a > \chi_b\). Even in the Y247 system without the twin microstructure and with small paramagnetic anisotropy, the modulated oval field aligned all the crystallographic axes, namely, triaxial grain orientation was achieved.

To evaluate the degree of in-plane orientation of thus obtained Y247, a pole graph was recorded. Figure 6 shows a contour mapping of the pole figure of the (0113) peak. Twofold symmetric reflection spots were clearly observed, confirming the achievement of triaxial orientation. The full-width at half-maximum of these spots was 2.4° for the \(b^*\)-direction, indicating that the degree of in-plane orientation was 2.4°. From a practical point of view, a degree of in-plane orientation of within 3° is superior to those for the \(a\)-axis growth region of the melt-solidified bulks of RE123 (5–10°) [11] and RE123 films grown on metal tape with several buffer layers (∼4°) [12]. Therefore, one can recognize that this magnetic orientation technique using the modulated oval field is a novel triaxial orientation process for the production of high-\(T_c\) superconductor materials.

A serious problem in the above grain-orientation technique at the current stage is that the excellent triaxial orientation was achieved in epoxy resin with high viscosity. Therefore, to improve practicality of this process, the fabrication of quasi-single-crystalline bulk samples and thick films is required. Also it is indispensable to use a colloidal solution [13–15] containing Y247 powder, in which the viscosity is much smaller. This is a serious practical problem to overcome, and, in practice, the use of other modulated
oval field techniques, the choice of appropriate RE ions [5] in RE247 and the optimization of grain orientation conditions will be required. Furthermore, if the grain orientation technique using the modulated oval field can be applied to the RE123 phase, there is a strong possibility that this magneto-scientific technique will become a practical production method for bulk magnets and superconducting cables. Of course, one should take into account that a twin microstructure is formed in the general sintering of RE123 and development a method suppressing the disappearance of the in-plane magnetic anisotropy induced by the twin microstructure.

4. Conclusions

The easy axes of magnetization and magnetic anisotropies at room temperature of the magnetically oriented powders of RE123, RE-doped Bi2212 and RE-doped BiCa222 depend on the type of RE ion. The magnetic role of each RE ion was qualitatively almost the same for the three compounds, and the magnetic anisotropies generated by heavy RE ions were much larger than those generated by light RE ions. The appropriate choice of RE dopant leads to not only a change in the direction of the easy axis but also a minimization of the magnetic field required for grain orientation. Furthermore, we demonstrated the triaxial orientation of Y247 without the twin microstructure in epoxy resin using a modulated oval field. According to the pole figure of the (0113) peak, a high degree of in-plane orientation of less than 3° was achieved, which was smaller than that of coated RE123 conductors. Therefore, there is a strong possibility that a magnetic triaxial orientation technique will be used in the production of superconducting bulks and thick films if a grain orientation technique based on a colloidal process is developed.

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