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Physically driven enhancement of the stability of Bi₂O₃-based ionic conductors via grain boundary engineering

Incheol Jeong¹, Seung Jin Jeong², Byung-Hyun Yun³, Jong-Won Lee³, Chan-Woo Lee⁴, WooChul Jung^{2,5} and Kang Taek Lee^{1,5}

Abstract

Fast oxygen-ion conductors for use as electrolyte materials have been sought for energy conversion and storage. Bi₂O₃-based ionic conductors that exhibit the highest known oxygen-ion conductivities have received attention for use in next-generation solid electrolytes. However, at intermediate temperatures below ~600 °C, their conductivities degrade rapidly owing to a cubic-to-rhombohedral phase transformation. Here, we demonstrate that physical manipulation of the grain structure can be used to preserve the superior ionic conductivity of Bi₂O₃. To investigate the effects of microstructural control on stability, epitaxial and nanopolycrystalline model films of Er_{0.25}Bi_{0.75}O_{1.5} were fabricated by pulsed laser deposition. Interestingly, in situ impedance and ex situ XRD analyses showed that the grain boundary-free epitaxial film significantly improved the stability of the cubic phase, while severe degradation was observed in the conductivity of its polycrystalline counterpart. Consistently, the cation interdiffusion coefficient measured by the Boltzmann–Matano method was much lower for the epitaxial thin film compared to the polycrystalline thin film. Furthermore, first-principles calculations revealed that the presence of grain boundaries triggered the structural resemblance between cubic and rhombohedral phases, as evidenced by radial distribution functions. Additionally, phase transition energetics predicted that the thermodynamic stability of the cubic phase with respect to the rhombohedral counterpart is reduced near grain boundaries. Thus, these findings provide novel insights into the development of highly durable superionic conductors via microstructural engineering.

Introduction

Extensive research has been conducted on the potential of oxygen-ion conductors for use as solid electrolytes in electrochemical devices such as oxygen sensors^{1,2}, solid oxide fuel cells^{3–5}, and oxygen pumps⁶. To date, most studies of oxygen-ion electrolytes have been focused on oxide materials with cubic fluorite structures, such as doped ZrO₂⁷, CeO₂^{8–10}, and Bi₂O₃^{11–13}. Among these, the

δ-Bi₂O₃ polymorph, stands out as a promising electrolyte because of its high oxygen ionic conductivity ($\approx 1 \text{ S cm}^{-1}$ at 800 °C); this stems from the high concentration of disordered oxygen vacancies, which account for 25% of the anion sublattice¹⁴.

Despite the superior ionic conductivity of δ-Bi₂O₃, the thermodynamic stability of the δ phase is confined within a narrow temperature range (729–824 °C); outside of this temperature range, the structure undergoes phase transitions to structures with reduced symmetries, such as the tetragonal β or monoclinic α phases, which exhibit insufficient ionic conductivities at lower temperatures¹⁵. Such phase transitions severely limit the application of δ-Bi₂O₃ and have inspired multiple attempts to stabilize the cubic phase across a broader temperature range by doping

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with rare earth oxides^{16–20}. In particular, Er_2O_3 is frequently selected because it allows fast ionic conduction ($\approx 0.35 \text{ S}\cdot\text{cm}^{-1}$ at 700°C). Notably, an Er_2O_3 concentration of $\sim 15\%$ is sufficient to stabilize the cubic phase, while associations among the dopant and oxygen vacancies are lower than those of alternative dopants²¹.

However, it has been reported that when erbia-stabilized Bi_2O_3 (ESB) is exposed to reduced temperatures (below $\sim 600^\circ\text{C}$), the conductivity deteriorates dramatically due to a phase transformation to a rhombohedral structure that involves breaking the high symmetry of the cubic phase²². Therefore, studies have been conducted to sustain the cubic structure and fast ionic conductivity of stabilized Bi_2O_3 ^{22–24}. A recent study reported that the cubic phase was maintained by doping aliovalent cations and may enhance the long-term behavior²⁵. Specifically, it was demonstrated that the phase transition kinetics were highly associated with diffusion of cations by investigating the time-dependent ionic conductivities of acceptor- and donor-doped systems. Consequently, it was suggested that the control of cation diffusion enhanced the stability of Bi_2O_3 .

In parallel, for many ionic-conducting polycrystalline oxides, grain boundaries (GBs) often interfere with rapid ionic diffusion, particularly during the transfer of ionic carriers²⁶. This is the case for fluorite-based electrolytes such as doped zirconia and ceria, which has led to many attempts to improve the conductivity by increasing the grain size and thereby reducing the GB density^{27,28}. However, unlike other fluorites, the GBs of stabilized Bi_2O_3 are remarkably conductive and, therefore, have little effect on the overall ionic conductivity²⁹. Partly due to these unique interfacial properties of Bi_2O_3 , there have not been many attempts to control the microstructure of the Bi_2O_3 electrolyte and manipulate the density of the GBs. Considering the influence of GBs on phenomena involving atomic movement and recrystallization^{30–32}, such as grain growth, precipitation, and phase transitions, a systematic study elucidating the correlations between GBs and the phase stability of stabilized Bi_2O_3 would suggest new insights.

In this study, we used pulsed laser deposition (PLD) techniques to investigate how the long-term phase stabilities of ESBs change depending on rigorously controlled grain boundary conditions. Both epitaxial and polycrystalline thin films of $\text{Er}_{0.25}\text{Bi}_{0.75}\text{O}_{1.5}$ (25ESB) were prepared on two different single crystal substrates (SiO_2 (001) and C-plane Al_2O_3 (0001)). These polycrystalline films with nanosized columnar grains are expected to clarify the contribution of GBs to cubic phase stability when compared to the GB-free epitaxial counterpart. The surface morphologies and crystallographic structures of the fabricated model samples were characterized through atomic force microscopy (AFM) and X-ray diffraction (XRD), respectively. Next, the time-dependent behavior of the ionic

conductivity was analyzed in situ using electrochemical spectroscopy (EIS). To elucidate the effects of GBs on the phase transition kinetics, the cation interdiffusion coefficient was measured by the Boltzmann–Matano method. An atomic-level understanding of the stabilities of ESBs arising from the presence of GBs was obtained by density functional theory (DFT) calculations with a focus on local atomic geometries, electronic structures of the GB region, and phase transition energetics. To the best of our knowledge, the current work provides the first indication of how the grain boundary condition affects the long-term behaviors of ESBs with a thin-film model study.

Materials and methods

Sample preparation

The ESB thin films and CeO_2 interlayer were prepared using PLD from oxide targets of the respective materials. The ESB target was prepared via a conventional solid-state reaction in which a mixture containing stoichiometric amounts of Bi_2O_3 (99.99% pure, YeeYoung Cerachem) and Er_2O_3 (99.99% pure, Alfa Aesar) was ball-milled for 24 h with zirconia media in a HDPE bottle. The powder mixture was calcined at 800°C for 16 h before the calcined powder was crushed finely with a mortar and pestle and sieved with a $45 \mu\text{m}$ mesh. The CeO_2 target was prepared using micron powder (99.9%, Alfa Aesar). Finally, the ESB and CeO_2 powders were uniaxially pressed under a pressure of 50 MPa to obtain the disk-shaped PLD target with a diameter of 2.54 cm (1 in.) and a thickness of 1 cm. The final ESB and CeO_2 targets were obtained after sintering at 890°C for 16 h and 1400°C for 6 h, respectively.

Polycrystalline thin films were grown on a single-crystal SiO_2 (001) substrate with a CeO_2 interlayer to prevent diffusion of Si. The effect of the interlayer is described in the Supporting Information and Fig. S1. Epitaxial thin films were grown on C-plane Al_2O_3 (0001) substrates. The PLD system was operated using a KrF excimer laser with an emission wavelength of 248 nm (Coherent COMPex 205) at a pulsed laser energy of 270 mJ and a laser repetition rate of 5 Hz for a selected duration. During the laser ablation process, the substrate temperature and O_2 working pressure were kept at 700°C and 10 mTorr, respectively. After the deposition process was completed, the chamber was kept at 700°C under an O_2 working pressure of 1 Torr for 30 min to facilitate oxidation of the films. It was confirmed in Fig. S2, which shows XRD results obtained before and after heat treatment, that the ESB films were prepared well and without any oxygen deficiency.

Characterization

To investigate the crystal structures of the deposited ESB films, XRD measurements were conducted using a Bragg–Brentano diffractometer (Rigaku Ultima IV,

Tokyo, Japan, Cu K α wavelength of 1.541 Å). The in-plane reflections recorded in the high-resolution XRD measurements were utilized to perform a precise analysis of the growth structures and crystallinities of the deposited films using a diffractometer (X'Pert-PRO MRD, PANalytical) operated at 45 kV and 40 mA with a fixed Cu anode. An AFM (Park Systems XE-150) was used to study the surface morphologies of the specimens.

The electrical conductivities of the ESB thin films were evaluated using two-probe electrochemical impedance spectroscopy (EIS, Biologic VMP-300). The details of the experimental setup and analytical method for the in-plane conductivity measurements are given in the Supporting Information and Fig. S3. The impedance of each specimen was measured under ambient air conditions (balanced by mixing O₂ and Ar gas flows) over a frequency range of 5 MHz to 0.05 Hz and an amplitude range of 100–400 mV. In addition, the measurements were carried out based on a carefully controlled heating schedule, considering the degradation occurring at elevated temperatures. The details are included in the Supporting Information and Fig. S4.

A focused ion beam (FIB, Hitachi NB 5000) was used to prepare trenches for the thin films, and a composition analysis was performed at each location using transmission electron microscopy combined with energy-dispersive X-ray spectroscopy (TEM EDS, Hitachi HF-3300). The impurity diffusion profiles of the polycrystalline ESB thin film samples were analyzed by time-of-flight secondary ion mass spectrometry (ToF-SIMS; TOF-SIMS5, Germany) after the annealing process. Here, 30 keV Bi⁺ clusters with a raster of 256 × 256 measured points scanned over an area of 100 × 100 μm² were used to generate negatively charged secondary ions, and the secondary ion intensity levels of all relevant isotopes and clusters were analyzed simultaneously. For depth profiling, 1 keV Cs⁺ ions (300 × 300 μm²) were used in noninterlaced mode (analyze 1 frame, sputter 3 s, pause 4 s) for sequential ablation of the surface between measurements of the mass spectra.

Computational details

DFT calculations were performed using VASP^{33,34}. The generalized gradient approximation (GGA-PBE) was chosen for the exchange-correlation functionals. Projector-augmented-wave (PAW) potentials with valence configurations of 4f²5d¹⁰6s²6p³ and 2s²2p⁴ were applied to describe Bi and O, respectively. Plane waves with an energy cutoff of 450 eV based on the convergence test within 0.02 eV/f.u. were used to expand the electronic wave functions. We used 1 × 1 × 1 and 2 × 2 × 2 *k*-meshes³⁵ within the Monkhorst–Pack scheme for the GB and bulk models, respectively. The *k*-point settings were optimized by energy convergence tests based on a 1 × 1 × 1 Bi₂O₃ unit cell. The convergence threshold for the electronic self-consistent iterations was set to 10⁻⁴ eV/cell.

The atomic positions and cell parameters were relaxed until the residual force reached 10⁻³ eV/Å. We obtained a lattice constant of $x = y = z = 5.7$ Å for the 1 × 1 × 1 unit cell after fitting the calculated total energy as a function of the unit cell volume to the Birch–Murnaghan equation of state³⁶. This yielded good agreement (~3%) with the experimental result³⁷ extrapolated to 0 K (5.553 Å), which was within the typical overestimation calculated by GGA functionals. Based on the lattice constant of the grain condition, a tilt Σ3[110]/(111) GB model was generated because it was reported based on previous calculations to have a lower GB formation energy compared to other models for fluorite systems^{38–40}. A slab model with a vacuum region was developed to depict the atomic structure of the GB region with a sufficient bulk region. The GB model was constructed by mirroring and shifting the (111) plane, which featured 240 atoms and had dimensions of 8.06 × 8.06 × 98.98 Å³. We used slabs sandwiched by a vacuum region of approximately 20 Å along the *z*-direction to simulate the GBs. To minimize the effects of surface electronic redistribution on the GB area and to describe the bulk region adequately, three terminal atomic layers were fixed (one stoichiometric Bi₂O₃ (111) trilayer). In addition, two oxygen layers were fixed at the center of the GB model to prevent the structure from being fully relaxed along the GB region because the GB blurs and dissipates owing to facile relaxation of oxygen ions originating from the disordered nature of the anion sublattice in Bi₂O₃. To approximate the disorder of oxygen vacancies, the special quasi-random structure (SQS) method from the Alloy Theoretical Automated Toolkit (ATAT) was used. In the SQS approach, oxygen ions are arranged on the anion sublattice so that their most relevant radial correlation functions imitate those of the reference structure, which is perfectly random⁴¹. Doublet and triplet clusters are considered as proposed from previous studies^{42,43}, and the generated structures showed energy convergence to within 0.1 eV/f.u. However, additional consideration of quadruplets did not lead to improvements. Furthermore, the maximum cutoff radius within the supercell was used to ensure that all possible ranges of clusters were covered. The effective coordination number (ECN) was calculated by weighting the individual bond lengths between the central ion and the nearby ions in comparison to the minimum bond length of the central ion⁴⁴. The ECN was calculated by summing the weights as follows:

$$\text{ECN} = \sum_i \exp\left(1 - \left(\frac{l_i}{l_{\min}}\right)^6\right) \quad (1)$$

where l_i denotes the bond lengths between the central and nearby i ions, while l_{\min} denotes the minimum bond lengths from the central ion.

Results and discussion

Grain boundary manipulation via thin-film deposition

Two types of thin films with different microstructures were used as model samples to confirm the effect of GBs on phase stability and ion conductivity. Figure 1a shows a schematic drawing of the epitaxial and columnar growth of the ESB thin films. Figure 1b, c present AFM images of the as-deposited surface morphologies of the epitaxial ESB thin film (Epi-ESB) and polycrystalline ESB thin film (Poly-ESB), respectively. Epi-ESB had a smooth surface ($R_q = 0.686$ nm) without GBs, whereas Poly-ESB exhibited widespread surface irregularities ($R_q = 2.737$ nm) corresponding to GBs over an area of $1 \mu\text{m} \times 1 \mu\text{m}$. Figure 2a shows the out-of-plane XRD patterns (ω - 2θ) for Epi-ESB and Poly-ESB. The crystalline cubic fluorite structure (δ -phase) was observed for both films without any secondary phase formation. Indeed, Epi-ESB showed only the (111) peak, which, as reported elsewhere, was indicative of coherent film growth. The in-plane diffraction patterns shown in Fig. 2b provided further evidence for epitaxial

growth of Epi-ESB. In contrast to Poly-ESB, for which no peaks were observed in the in-plane ϕ -scan results, Epi-ESB showed sharp in-plane peaks with a six-fold rotational symmetry of 60° , thus confirming the strong alignment registry between the film and the substrate. These crystallographic features of epitaxially grown ESB films on C- Al_2O_3 (0001) substrates were repeatedly confirmed in previous studies^{45,46}. Alternatively, several orientations, namely, (111), (200), (220), and (331), were observed for Poly-ESB, indicating the polycrystalline growth of ESB. The possibility of a phase change for SiO_2 at high temperature was reported in a previous study⁴⁷. However, as seen from the high-temperature real-time XRD results of the z-cut SiO_2 substrate in Fig. S5, the phase change of SiO_2 was negligible and certainly had no effect on the growth of Poly-ESB films.

Furthermore, additional characterization of the Epi-ESB film was performed by means of high-resolution TEM (HR-TEM), as shown in Figs. S6, 7. First, the cross-sectional HR-TEM image (Fig. S6d, e) of the ESB film

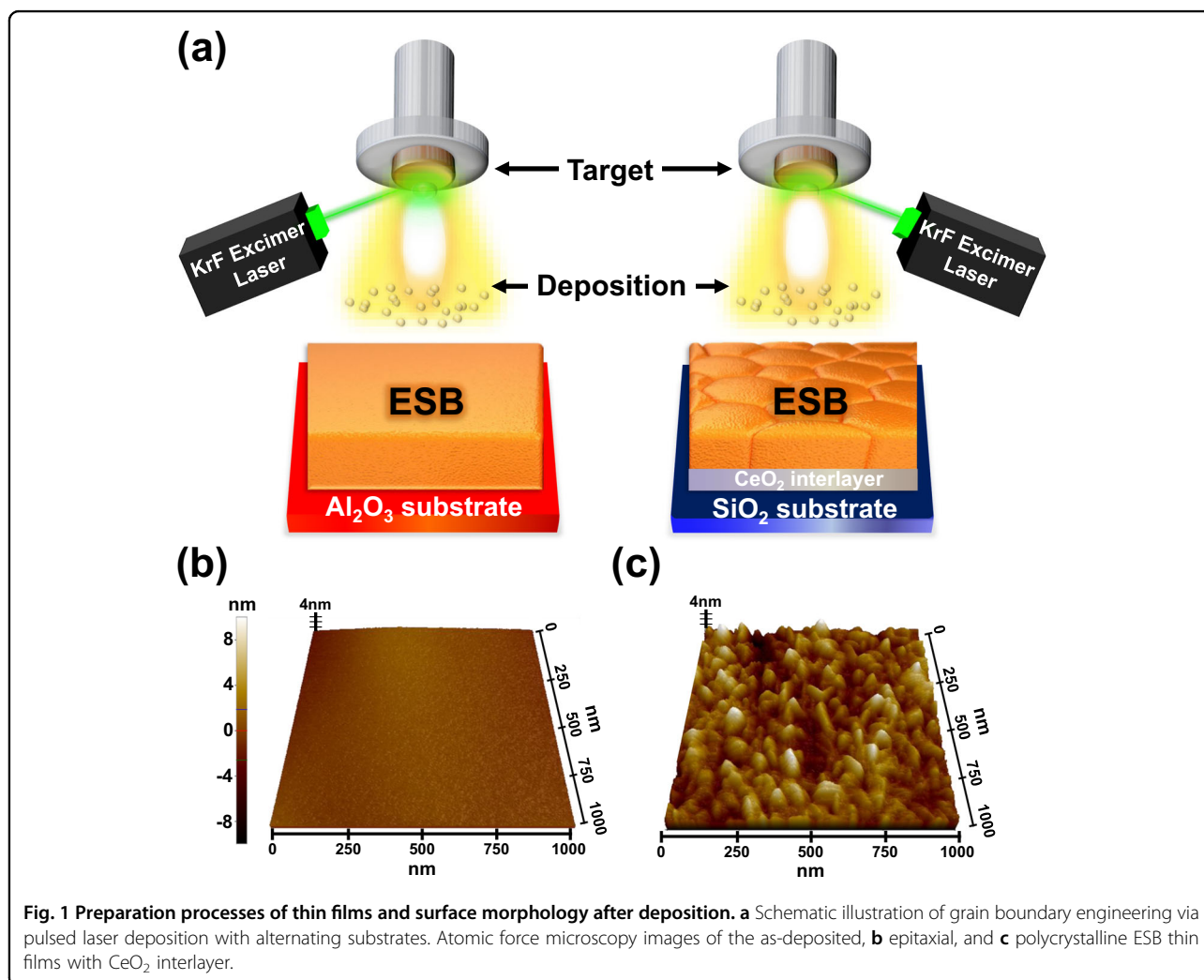
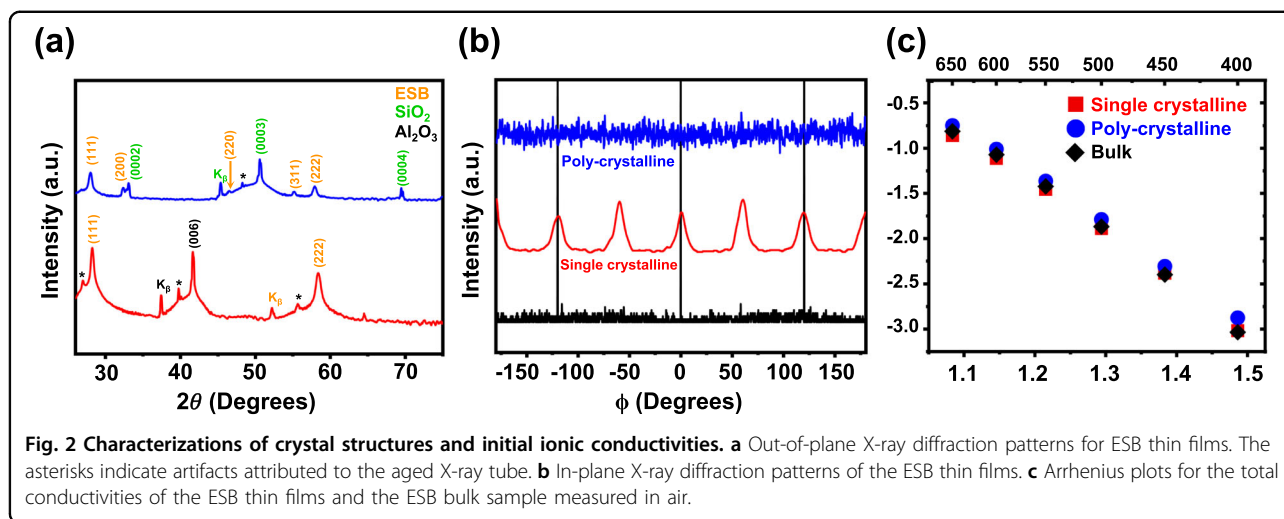


Fig. 1 Preparation processes of thin films and surface morphology after deposition. **a** Schematic illustration of grain boundary engineering via pulsed laser deposition with alternating substrates. Atomic force microscopy images of the as-deposited, **b** epitaxial, and **c** polycrystalline ESB thin films with CeO_2 interlayer.



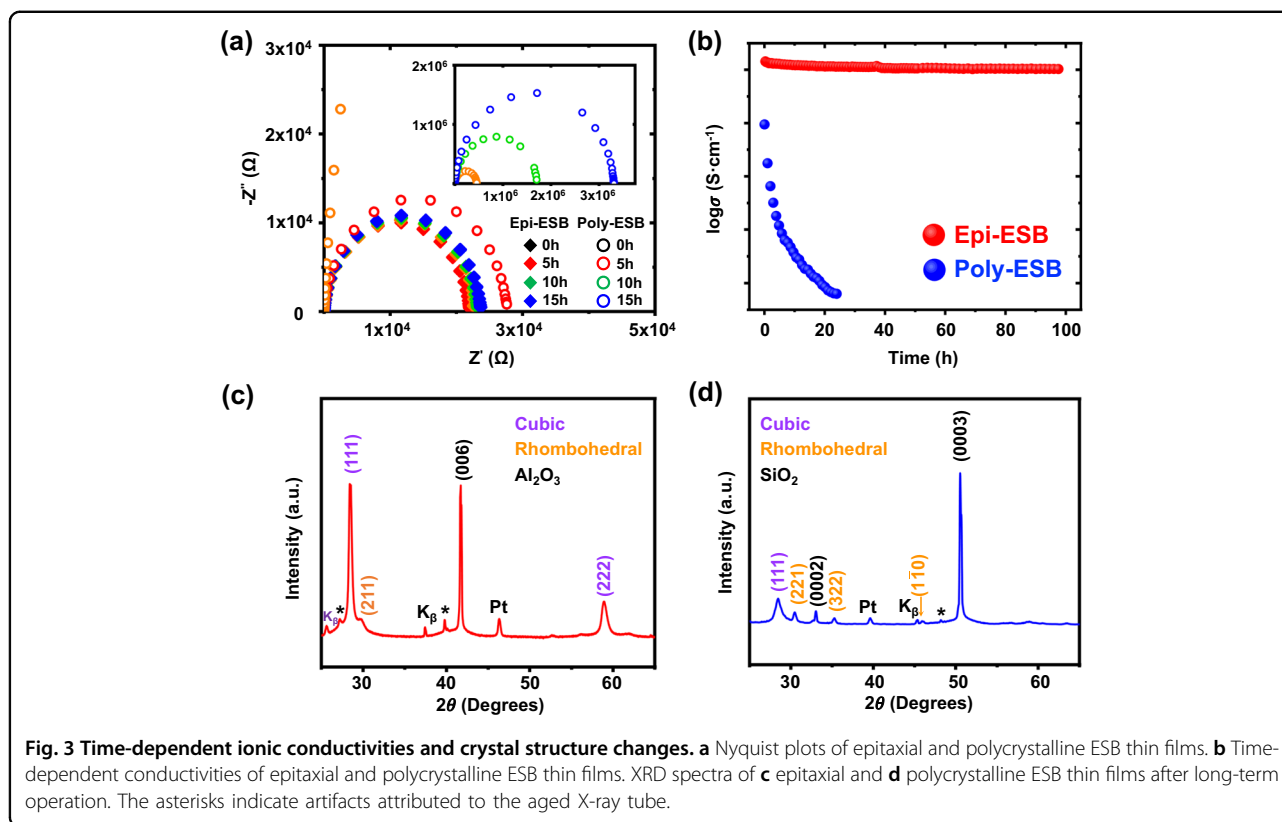
region and ESB|Substrate interface region revealed that the dense ESB thin film was well grown and without any grain boundaries. Figure S6b, c shows the selected area electron diffraction (SAED) patterns for the film area and interface region, respectively, as marked in Fig. S6a. The heteroepitaxial relationship between the ESB thin film and the C-Al₂O₃ substrate is clearly shown, and the highly oriented and crystalline growth feature of Epi-ESB was confirmed. The strong spots are the Bragg reflections of fluorite-like structures, while the other spots are satellite reflections, as reported elsewhere^{26,48}.

Figure 2c presents the ionic conductivities of Epi-ESB and Poly-ESB alongside that of bulk ESB in the Arrhenius plots. The conductivity values were derived by investigating the origin of the resulting impedance spectra, and the detailed process is included in the Supporting Information and Figs. S3 and S8–S10. Initially, measured ionic conductivities and calculated activation energies of bulk (1.1 eV), Epi- (1.1 eV), Poly-ESB (1.0 eV) matched each other well and agreed with the results of our previous work, which demonstrated that GBs in ESB serve neither as preferential paths for oxygen diffusion nor as obstacles to oxygen conduction^{29,49}. In summary, these results emphasize that the thin films fabricated herein were well defined, thereby maintaining the inherent transport properties of ESB.

Time-dependent conductivity behavior

Figure 3 shows how the ionic conductivity and crystal structure of ESB change over time at 600 °C. Specifically, Fig. 3a shows the Nyquist plots for Epi-ESB and Poly-ESB annealed for 20 h. Surprisingly, the ohmic resistance was maintained in Epi-ESB, whereas a dramatic increase in the resistance was observed for Poly-ESB. Figure 3b presents the time-dependent conductivity behaviors of ESB thin films heated at 600 °C for 100 h: the conductivity of Poly-

ESB decreased by 94.6% from 9×10^{-2} to 1×10^{-3} S/cm after approximately 20 h of operation, while the initial conductivity ($\sim 7 \times 10^{-2}$ S/cm) of Epi-ESB remained almost constant during annealing for up to 100 h ($\sim 6 \times 10^{-2}$ S/cm). Furthermore, additional experiments were performed to investigate the effect of lattice strain on conduction characteristics and long-term phase stability. Under identical PLD process conditions with different durations of deposition, films with three different thicknesses (140, 500, and 780 nm) were grown on identical Al₂O₃ substrates. As seen from Fig. S11, all of the films showed epitaxial growth features with highly preferred orientations in the (111) direction. In Fig. S11b, the thinner the film was, the greater the influence received from the substrate, and a larger peak shift was observed compared to that of the fully relaxed ESB25 bulk. Based on the (111) peak shift value, the respective strain level was evaluated as shown in Table S1 by referring to the previous method^{50,51}. Next, conductivity measurements and long-term stability tests at 600 °C were performed on the thin films, as shown in Fig. S12. The conductivity values of the strained films were slightly lower than that of the bulk, but the conductivity value of the 500 nm film was similar to that of the bulk. Furthermore, the results of the long-term stability tests shown in Fig. S12b indicated that all three epitaxial thin films underwent sluggish degradation with rates several orders of magnitude slower than that of the polycrystalline thin film. In this regard, enhancement of the long-term durabilities of epitaxial films did not show a significant correlation with the thicknesses (and lattice strain) of the epitaxial films. Thus, it was demonstrated that, regardless of their thickness, epitaxial ESB films exhibited long-term phase stability much superior to that of polycrystalline ESB film. Figure S13 shows the ToF-SIMS analysis of possible Ce diffusion due to the CeO₂ interlayer in Poly-ESB after the



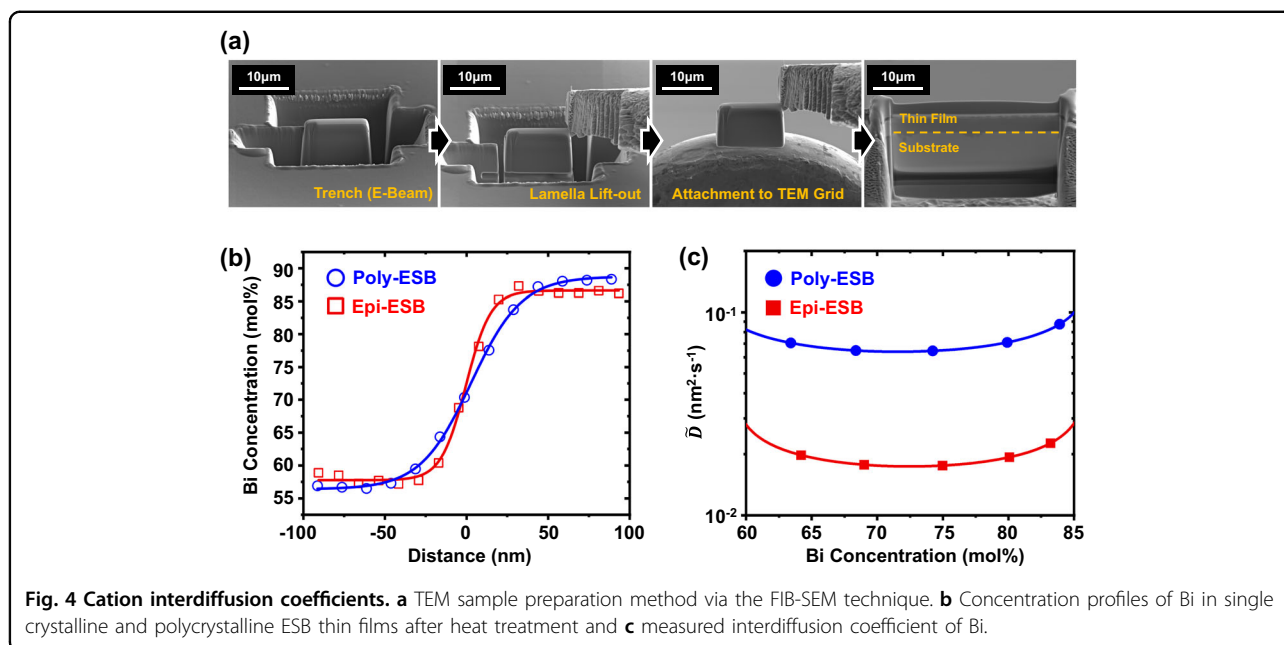
conductivity measurement. However, the level of Ce content in the interior of the ESB thin film was only at the noise level, indicating the absence of chemical reactions or intermixing of chemical species.

After the long-term stability tests, XRD analyses of both samples revealed that the long-term heat treatment led to a clear difference between the films, as shown in Fig. 3c, d. Overall, the cubic phase was maintained for Epi-ESB, except for growth of a negligible rhombohedral peak (Fig. 3c); however, for Poly-ESB, several rhombohedral peaks were formed after 30 h of heat treatment, while minor peaks for the cubic structure vanished, and only the (111) oriented peak remained (Fig. 3d). This result suggested that the degraded conductivities of the films following long-term annealing were mainly due to a transition from the conductive cubic phase to the resistive rhombohedral phase, as reported in our previous work²⁵. In the XRD spectra of Poly-ESB, it appeared that cubic and rhombohedral phases coexisted. Specifically, as a substantial transition occurred, the (111) peak of the cubic phase dropped in intensity and widened, whereas the (211) peak, which is the main peak of the rhombohedral phase, became relatively larger. In the XRD spectra of Epi-ESB, cubic peaks remained, and only a minor rhombohedral peak was observed, indicating significant suppression of the phase transition. Effects related to nanoscale structural evolution, such as amorphous phases and local

dislocations, were not considered in this study, and further studies may be necessary.

Cation interdiffusion coefficient

An association between cation diffusion and phase transition kinetics has been suggested in previous studies^{22–25}, and we used microstructural control to determine whether there exists a corresponding correlation in this study. The interdiffusion coefficients of Bi cations were measured via the Boltzmann–Matano method. To this end, films with different compositions were deposited sequentially to establish bilayered diffusion couple specimens (40ESB|15ESB) for each of Epi-ESB and Poly-ESB. After subjecting each diffusion couple specimen to heat treatment at 700 °C for 10 h, specimens for transmission electron microscopy (TEM) analysis were fabricated by using an in situ lift-out technique with a FIB. Figure 4a shows the detailed steps of sample preparation. The layer at the diffusion couple interface was obtained by FIB etching and attached to the grid. The Bi concentration as a function of distance was then quantified using the energy-dispersive X-ray spectroscopy (EDS) capability of the TEM system. The concentration profile was drawn with $x = 0$ representing the Matano interface, that is, the interface where the mass is balanced and the diffusion-induced loss on one side equals the corresponding gain on the other side. The concentration profiles after heat



treatment of Epi-ESB and Poly-ESB are shown in Fig. 4b. Upon inspecting the changes in slopes near the Matano interface, the change in the Bi concentration gradient was found to be significantly lower in Epi-ESB than in Poly-ESB. This implied that the diffusion coefficient of Bi in the epitaxial specimen was smaller than that in the polycrystalline specimen. The diffusion coefficient of Bi was quantitatively determined via the Boltzmann–Matano method, as follows:

$$\tilde{D}(C^*) = -\frac{1}{2t} \left(\frac{dx}{dC} \right) \bigg|_{C^*} \int_{C_0}^{C^*} x dC \quad (2)$$

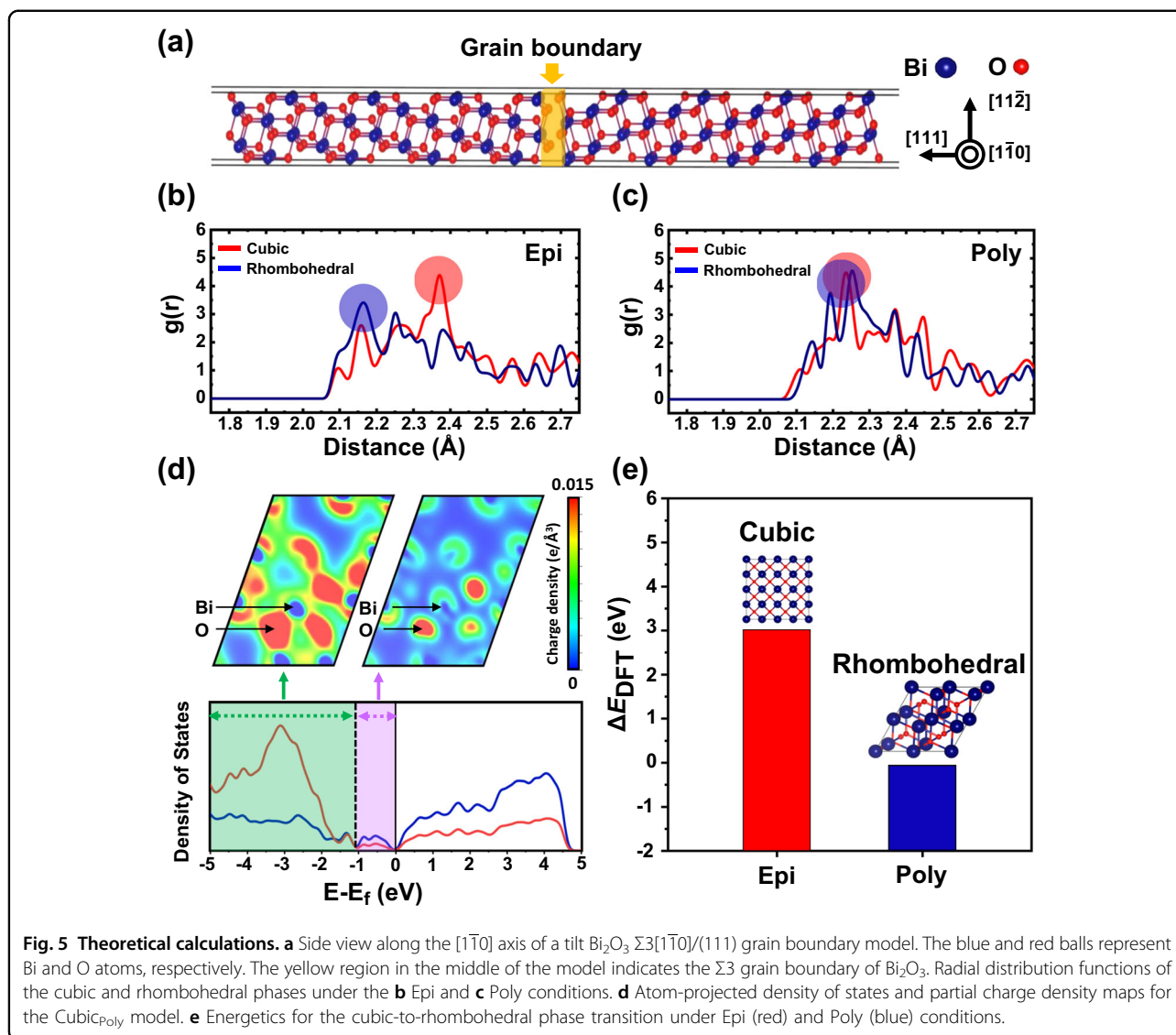
where t denotes a specific time, C_0 is the Bi concentration at the end of the diffusion couple, and C^* represents the Bi content at a specific position. The composition-dependent Bi interdiffusion coefficients were measured for Epi-ESB and Poly-ESB with this method, and they were plotted in Fig. 4c. The measured interdiffusion coefficient of Epi-ESB observed for the 25ESB composition ($0.02 \text{ nm}^2/\text{s}$) was 71% lower than that of Poly-ESB ($0.07 \text{ nm}^2/\text{s}$). This result indicated that cation diffusion can be controlled by microstructural manipulation. Thus, slower diffusion of Bi cations would result in effective suppression of a phase transition for stabilized bismuth oxide.

Role of grain boundaries in phase transitions

To further understand the stability of the cubic phase in the presence of GBs, the atomic-level behavior was simulated using DFT calculations. The GB model was envisaged as alternating O and Bi planes, as shown in

Fig. 5a. To explore how the atomic structure of the grain condition changed due to the presence of GBs, a structural model containing slab layers representing the bulk and GB conditions was developed (Fig. S14). Then, a gradual change due to the presence of GB was observed. The local geometry of the GB model was analyzed from the perspective of Bi cations, and the ECNs⁴⁴ of the Bi ions (ECN_{Bi}) were averaged for each layer of the GB model. The ECN_{Bi} gradually decreased as the Bi ions approached the GB, as shown in Table S2, implying that interactions of Bi with O were weakened near the GB. This might free Bi cations to move from the initial sublattice and thereby speed diffusion of Bi ions, as observed in Poly-ESB, which contained numerous GBs. To understand the changes in stability resulting from reduced interactions between Bi and O upon moving from the bulk condition toward the GB condition, two sets of cubic and rhombohedral structures are prepared with varying degrees of oxygen occupation. One set represented the bulk condition with minimal GB influence (Epi condition), and the other represented the GB condition with maximal GB influence (Poly condition). Cubic and rhombohedral phases for each condition were denoted X_Y ($X = \text{Cubic, or Rhom}; Y = \text{Epi, or Poly}$) for consistency with the experiments. The reproducibilities of the two sets for each condition were verified from the atomic and electronic perspectives, and details are described in the Supporting Information. The schematic processes used for calculations are depicted in Fig. S15.

Figure 5b, c shows the bonding characteristics of the cubic and rhombohedral phases for each condition, as analyzed using radial distribution functions. The



frequencies of the Bi–O bonds were described as a function of distance within the first coordination shell. The most dominant bond lengths were marked with red and blue circles for the cubic and rhombohedral phases, respectively. While there were significant bonding differences between the cubic and rhombohedral phases in the Epi condition, the bonding distributions of the two phases in the Poly condition were more closely aligned, as evidenced by the overlapping red and blue circles. From this structural resemblance between the cubic and rhombohedral phases in the Poly condition, it can be inferred that transformation from the cubic to the rhombohedral phase became easier with less structural deformation near the GBs. Figure 5d shows the density of states and corresponding charge density maps for $\text{Cubic}_{\text{Poly}}$. While the states in the lower energy range (Fig. 5d, green data below -1 eV) showed

results similar to those for $\text{Cubic}_{\text{Epi}}$ (Fig. S17a), the states below the Fermi energy (purple data from -1 to 0 eV) showed antibonding characteristics, which implied instability of the cubic phase under Poly conditions. This behavior can be attributed to the distinctive tendencies observed for atomic relaxation (represented by ECN_{Bi}). As shown in Table S3, the ECN_{Bi} of the rhombohedral phase underwent a greater change on going from Epi to Poly than the ECN_{Bi} of the cubic phase. Specifically, while the ECN_{Bi} of the rhombohedral phase (4.13) was higher than that of the cubic phase (3.99) under the Epi condition, the reverse was true for the Poly condition, where the ECN_{Bi} of the rhombohedral phase (2.87) was lower than that of the cubic phase (3.04). This indicated that rhombohedral Bi_2O_3 responded more readily to changes in the local atomic environment than its cubic counterpart.

In Fig. 5e, the calculated DFT energy, $\Delta E_{\text{DFT}} (= E_{\text{DFT}}^{\text{Rhom}} - E_{\text{DFT}}^{\text{Cubic}})$, predicts the energy cost for transformation from the cubic to the rhombohedral phase under each condition. Under the Epi condition, represented by the red bar, ΔE_{DFT} is positive, signifying that the cubic phase is likely to be maintained. However, under the Poly condition (blue bar), ΔE_{DFT} becomes negative, which indicates that the transformation from the cubic to rhombohedral phase is energetically favorable. In other words, it is predicted that the cubic structure is more stable than the rhombohedral structure in the absence of GB effects; the rhombohedral structure becomes energetically favorable with respect to the cubic structure when there is a grain boundary effect. Even taking the finite-temperature error ($\sim \text{meV/f.u.}$) into account^{52,53}, the difference in energy of $\sim 3 \text{ eV}$ (0.25 eV/f.u.) for the two conditions is distinguishable, indicating that the phase stability is highly sensitive to the grain boundary condition. These predictions indicate that the presence of GBs induces a phase transition for Bi_2O_3 , and these results strongly support our experimental data.

Conclusion

In this study, the long-term stability of the Bi_2O_3 -based superionic conductor was enhanced significantly via precise manipulation of GBs. While the conductivity of the poly-ESB thin film declined rapidly from 9×10^{-2} to $1 \times 10^{-3} \text{ S/cm}$ at 600°C due to the cubic-to-rhombohedral phase transformation, the Epi-ESB thin film maintained its initial conductivity for 100 h without a phase transition. Successful modification of cation diffusion was investigated with the interdiffusion coefficient of Bi, and a proportional relationship was observed between cation diffusion and phase transition kinetics. Furthermore, DFT calculations revealed that under the influence of GBs, a strong structural resemblance was observed between the cubic and rhombohedral phases; antibonding characteristics were observed for the cubic phase, implying high feasibility of transformation from the cubic to rhombohedral phase in ESB. These observations suggest that the superionic conductivity of stabilized Bi_2O_3 can be effectively secured by physically manipulating GB conditions in the bulk, such as by elongating heat treatment conditions, applying a thermomechanical treatment, or adding sintering agents. Thus, our findings provide novel insights for the development of highly conductive and durable solid electrolytes for next-generation energy applications.

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Conflict of interest

The authors declare no competing interests.

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