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Partial breakdown of translation symmetry at a structural quantum critical point associated with a ferroelectric soft mode

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We report for the first time that complete suppression of a phonon-driven structural phase transition causes partial breakdown of a three-dimensional translation symmetry in a well-defined sublattice. This state is revealed for a dielectric compound, $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$, that comprises an AlO_4 network incorporated into a hexagonal $\text{Ba}(\text{Sr})$ sublattice. Pair distribution function analyses and inelastic neutron scattering experiments provide clear-cut evidence of the AlO_4 network forming a continuum of Al-O short-range correlations similar to glasses, whereas the $\text{Ba}(\text{Sr})$ sublattice preserves the original translational symmetry. This glassy network significantly dampens the phonon spectrum and transforms it into the broad one resembling those typically observed in glass materials.

I. INTRODUCTION

Symmetry breaking in crystals has attracted broad interest in condensed matter physics. Particularly, exotic quantum phases in which spatial- or time-reversal symmetries are broken have been extensively studied over the last 20 years [1–7]. As a result, various fruitful research fields, such as topological insulators/superconductors, multiferroics, spintronics, and exotic superconductors, have been developed. Just as a point symmetry, a translation symmetry is also an essential component of crystals. However, few studies focus on a partial breaking of translation symmetry as a counterpart of point symmetry.

The aim of this study is to demonstrate that a three-dimensional translation symmetry is partially broken in a well-defined crystal by complete suppression of a soft-mode driven structural phase transition. The target material is BaAl_2O_4 , which comprises an AlO_4 tetrahedral network with six-membered cavities occupied by Ba atoms. This crystal structure is described as a combination of a Ba sublattice and an AlO_4 network (Figs. 1 (a) and (b)). It experiences a structural phase transition [8] at $T_C = 450$ K from the high-temperature phase (space group $P6_322$) to the low-temperature phase ($P6_3$) with an enlarged cell of $2a \times 2b \times c$ [9]. Simultaneously, the Ba atoms slightly shift along the c axis, which gives a spontaneous polarization. This structural phase transition is driven by the acoustic soft mode (irreducible representation M_2), whose vibration pattern is characterized as

tilting of AlO_4 accompanied by in-plane rotation of O1 atoms that connect the tetrahedra along the c axis [9]. Generally, in a soft-mode driven structural phase transition, the resultant low-temperature structure is closely related to the vibration pattern of the soft mode [10].

This acoustic soft mode is tunable by the chemical composition. As shown in the structural phase diagram (Fig. 2), T_C is rapidly suppressed by substituting Sr for Ba atoms and disappears near $x = 0.1$ [11, 12]. For the composition window of $0.1 \leq x \leq 0.5$, characteristic honeycomb-type thermal diffuse scattering has been observed in electron diffraction patterns over a wide temperature range of at least 100 K to 800 K. This fact means that the soft mode exists at very low frequencies or that a static structural fluctuation lies there. A further increase in x results in another structural phase transition to the $P2_1$ phase at $x = 0.55$. Details of the structural phase diagram of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ is reviewed in the Supplemental Material [13] together with the structural relationship between the parent $P6_322$ phase and the low-temperature phases.

In this paper, we investigate the local structure and phonon spectra of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ in the middle composition window. We show that complete suppression of the phonon-driven structural phase transition realizes a sublattice glass state in which the three-dimensional translation symmetry is partially broken in the well-defined crystal.

II. EXPERIMENTAL

Powder samples of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ were synthesized using a conventional solid-state reaction. Powder X-

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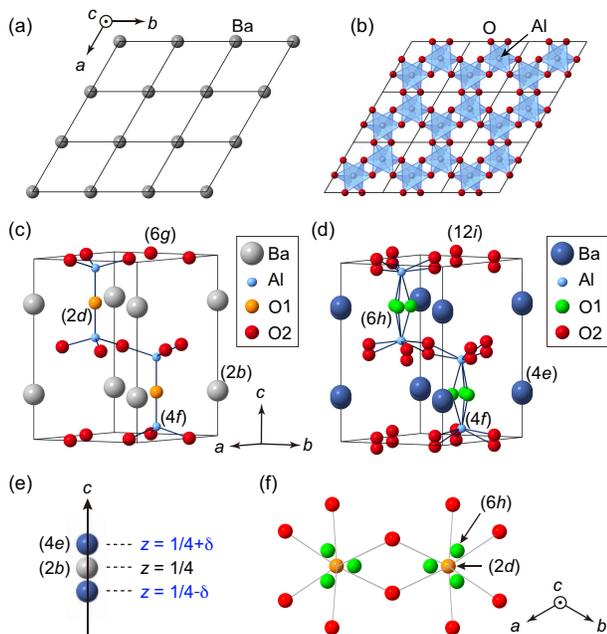


FIG. 1. Relationship between the (a) Ba sublattice and (b) AlO_4 network of the high-temperature phase of BaAl_2O_4 (space group $P6_322$). Solid boxes indicate the multiple unit cells. (c) Ideal crystal structure of the high-temperature phase. The cell parameters determined by using laboratory X-ray diffraction are $a = 5.2326(3)$ Å and $c = 8.8174(7)$ Å at 500 K. (d) Split atom model used as an average structure model for $x \geq 0.05$. In this model, the Ba, O1, and O2 sites are split into several seats with lowered symmetries and partial occupancies. Panels (e) and (f) describe the relationship between the ideal and split positions of Ba and O1 atoms, respectively. (e) Ba atoms are accommodated at the $2b$ site at $z = 1/4$ in the ideal $P6_322$ structure. When the Ba atoms have a small displacement ($\pm\delta$) along the c axis from the ideal site, they can be accommodated at the $4e$ site with lowered symmetry. (f) O1 atoms are accommodated at the $2d$ site in the ideal structure. Al atoms are not shown here. Because the $2d$ site is on the threefold axis along the c axis, the three seats of the $6h$ site are generated in the split atom model.

ray diffraction was carried out at the BL02B2 beamline (SPring-8). The X-ray energy was set at 25 keV. X-ray total scattering measurements were carried out at room temperature at the BL04B2 beamline (SPring-8) with the X-ray energy of 113.2 keV. Multi-incident energy time-of-flight INS measurements were performed at AMATERAS [14] installed at BL14 in the Materials and Life Science Experimental Facility (MLF) in Japan Proton Accelerator Research Complex (J-PARC). The incident neutron energies were set at $E_i = 52.45, 17.255,$ and 8.485 meV. For the inelastic neutron scattering experiment, the powder samples of $x = 0, 0.03, 0.07, 0.2, 0.3$ (mass ≈ 5 g each) were separately enclosed in thin-walled aluminum cans with He exchange gas. A top-loading closed cycle refrigerator was used for temperature control. Obtained data were visualized using Mslice [15] after data reduction [16].

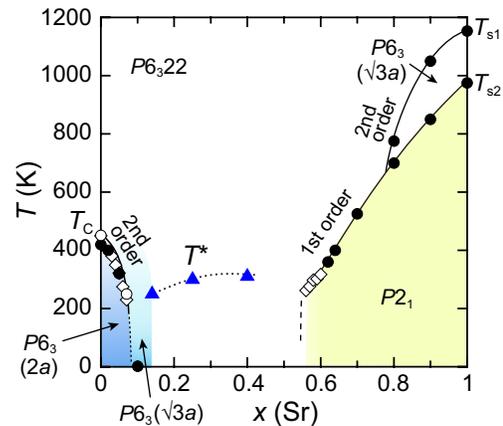


FIG. 2. Full phase diagram of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ [9, 11, 12, 22–24]. The phase boundaries are determined using powder X-ray diffraction (filled circles), synchrotron X-ray diffraction experiments for single crystals (open circles), and dielectric measurements (diamonds). T^* denotes the temperature at which thermal diffuse scattering caused by the M_2 and K_2 soft modes shows a maximum intensity. The K_2 mode is another dominant soft mode in the high-temperature phase of BaAl_2O_4 , and the $P6_3(\sqrt{3}a)$ structure is the condensed state of the K_2 mode [24]. The Supplemental Material provides further details of the phase diagram of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ [13].

$S(Q, E)$ were simulated using OCLIMAX [17] based on the phonon calculation. Phonon calculations were performed by using Phonopy [18] in conjunction with DFT calculations using Vienna ab initio package (VASP) code [19–21]. Thermal conductivity was measured for the densified polycrystalline samples of $x = 0$ –1 using the steady-state method in a physical property measurement system (PPMS, Quantum Design). Further experimental details are included in the Supplemental Material [13].

III. RESULTS

The synchrotron powder X-ray diffraction profiles at 15 K shown in Fig. 3 (a) verify that the $1/2 \ 3/2 \ 1$ superlattice reflection arising from the low-temperature phase is entirely suppressed in the composition window of $0.06 < x < 0.1$. The full width at half maximum (FWHM) is analyzed and converted to the correlation length (ξ) in Figs. 3 (b) and (c), respectively. ξ decreases with x and reaches ≈ 20 nm at $x = 0.06$. This variation reflects the microstructure being divided into smaller domains as x increases [23]. That is, the long-range order of the low-temperature phase is suppressed as x increases. Each domain is separated by anti-phase boundaries in the ab plane [23, 25]. The reported domain size is typically ≈ 100 nm at $x = 0$, decreases with increasing x , and reaches ≈ 20 nm at $x = 0.07$ [23]. These values are consistent with the value of ξ shown in Fig. 3 (c). The FWHM of the fundamental reflections are independent of

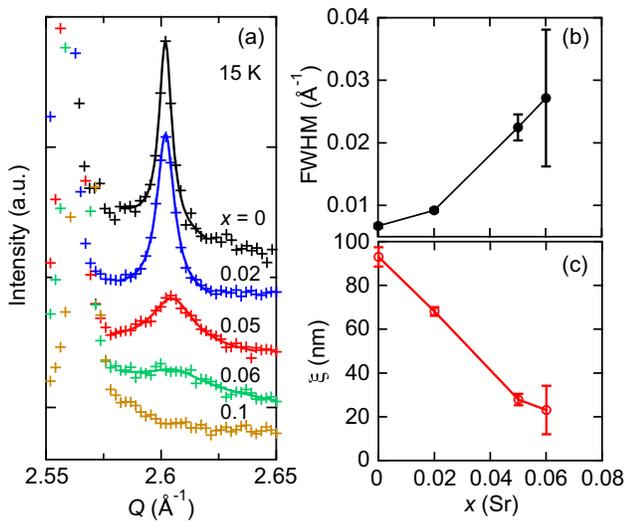


FIG. 3. (a) Synchrotron powder X-ray diffraction profiles in the range of $Q=2.55\text{--}2.65\text{ \AA}^{-1}$ for $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ ($x=0\text{--}0.1$) samples at 15 K. The $1/2\ 3/2\ 1$ superlattice reflection observed at $Q \approx 2.6\text{ \AA}^{-1}$ disappears between $x = 0.06$ and 0.1 . (b) FWHM as a function of Sr composition (x). (c) Correlation length (ξ) converted from the FWHM value. The value of ξ decreases with x and reaches ≈ 20 nm at $x = 0.06$. This length corresponds to only ≈ 20 unit cells of the low-temperature phase.

the Sr composition and exhibit almost a constant value, as revealed by various diffraction methods [11, 12].

The effect that changes the long-range order to short-range order should also affect the local structure of this material. Fig. 4 (a) displays the results of local structure analyses for $x = 0\text{--}0.3$ samples using the PDFs ($g(r)$) measured at 300 K. The profiles computed using the average structure models obtained with powder X-ray structural refinements [12] are also shown. The standard program package PDFgui [26] was employed for the computation. Because of the structural phase transition, the structure model of the low-temperature phase ($P6_3, 2a$) is employed as the average structure for the profile computation at $x < 0.05$. In contrast, the computation for $x \geq 0.05$ employs the split atom model of $P6_322$ (see Fig. 1 (d)) as the average structure instead of the ideal structure (see Fig. 1(c)) because of the large atomic fluctuation. As shown in Fig. 4 (a), the observed PDF profile for $x = 0$ is well reproduced by the average structure model. Several characteristic peaks indicated by triangles are assigned based on the partial $g(r)$ plotted at the bottom of Fig. 4 (a); the peak at $r \approx 1.75\text{ \AA}$ comes from the Al-O1 and Al-O2 pairs in the AlO_4 tetrahedra. The peaks observed at $r \approx 3.45$ and 6.80 \AA correspond to the Ba-Al and Ba-Ba pairs, respectively. These pairs of atoms are indicated by double-headed arrows in the inset of Fig. 4 (a).

The peak positions (r) observed for $x = 0$ at $r \approx 1.75$, 3.45 , and 6.80 \AA are tracked and plotted as a function of

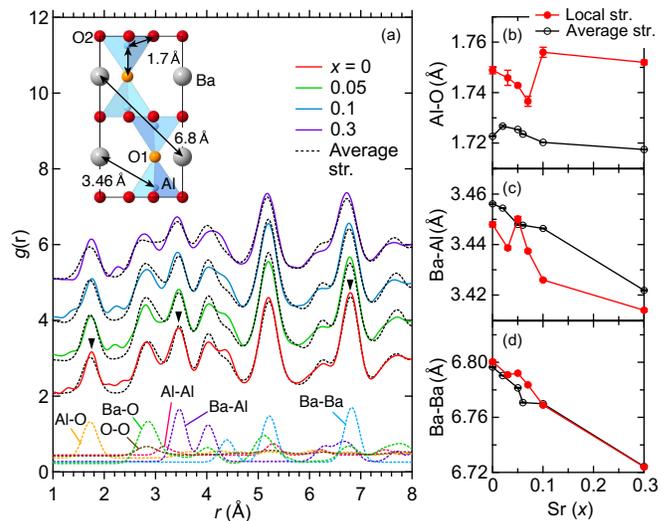


FIG. 4. (a) Experimentally derived PDFs ($g(r)$) at 300 K of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ with $x = 0\text{--}0.3$ (solid curves). The $g(r)$ curves for different compositions are plotted with a relative offset. Model calculations (broken curves) are also performed for each composition by using the average structures obtained with powder X-ray structural refinements [12]. Partial $g(r)$ curves for the average structure of $x = 0$ are shown at the bottom. The peak assigned as Al-O comes from the Al-O1 and Al-O2 correlations in the AlO_4 tetrahedra. (b), (c), and (d) Peak positions of the Al-O, Ba-Al, and Ba-Ba correlations, respectively.

x in Figs. 4 (b)–(d) together with the values calculated from the average structure models. The experimentally observed peak positions reflect the local structure of the compound. In the computed profile shown in Fig. 4 (a), the O-O and Ba-O peaks are superimposed on each other at $r \approx 2.9\text{ \AA}$. In addition, the computed peak at $r \approx 2.9\text{ \AA}$ is significantly broadened when x increases because of the Debye-Waller (DW) factors of Ba(Sr) and O atoms [12]. For these reasons, the position of the peak at $r \approx 2.9\text{ \AA}$ is not analyzed.

As shown in Fig. 4 (d), the calculated peak positions of the Ba(Sr)-Ba(Sr) pair almost linearly decrease with increasing x , reflecting the variation in the lattice constant [11]. The experimentally observed peak positions of the Ba(Sr)-Ba(Sr) correlation are entirely consistent with those of the average structure models, indicating that the Ba(Sr) atoms are at the average position. A similar trend is also observed for the Ba(Sr)-Al pair, as shown in Fig. 4 (c), although a small deviation is observed between the experimental and computational values. In contrast, the calculated values of the Al-O pair are almost independent of the Sr concentration, as shown in Fig. 4 (b). In addition, the experimentally observed peak position of the Al-O pair exhibits a clear upshift at $x = 0.1$. This Al-O peak is well separated from the other peaks, including the Ba-O peak at $r \approx 2.9\text{ \AA}$. Therefore, the results shown in Figs. 4 (b)–(d) indicate that

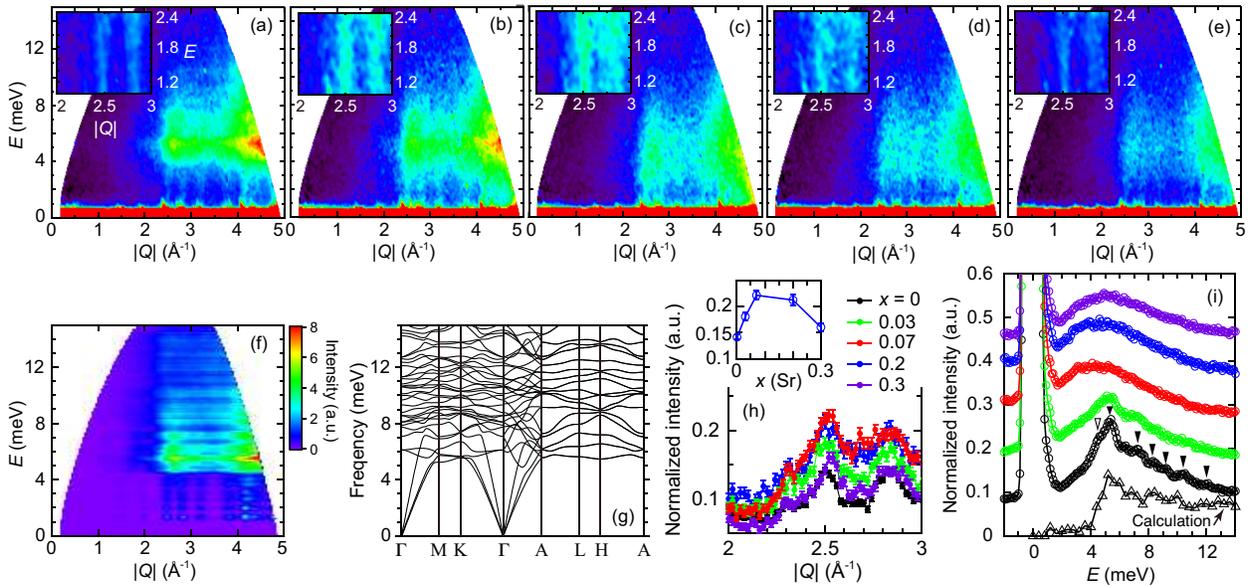


FIG. 5. Dynamical structure factor, $S(Q, E)$, for $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ powder samples of (a) $x = 0$, (b) 0.03, (c) 0.07, (d) 0.2, and (e) 0.3 measured at 100 K with $E_i = 17.255$ meV. The insets show an enlarged view in the range of $E=1.0\text{--}2.4$ meV and $|Q| = 2\text{--}3 \text{ \AA}^{-1}$ obtained with $E_i = 8.485$ meV. The Al sample can background has been subtracted from each scan. (f) $S(Q, E)$ calculated using the OCLIMAX program [17] based on the phonon calculation for BaAl_2O_4 . (g) Phonon-dispersion relations calculated on the ferroelectric $P6_3(2a)$ structure of BaAl_2O_4 . The calculation uses a $2 \times 2 \times 3$ supercell of the unit cell. (h) Inelastic spectra energy-integrated over the energy transfer $E=1\text{--}2.3$ meV with $E_i = 8.485$ meV. The inset presents the maximum intensity of the peak at $Q \approx 2.5 \text{ \AA}^{-1}$. (i) Inelastic spectra Q -integrated over $|Q|=2\text{--}4 \text{ \AA}^{-1}$ with $E_i = 17.255$ meV. Profiles are offset by 0.09 for clarity. The intensities in panels (h) and (i) are normalized using a coherent cross-section of the 111 nuclear peak whose structure factor is independent of the Ba/Sr ratio.

the local structure of AlO_4 tetrahedra deviates from the average structure at $x \geq 0.1$.

To investigate the nature of phonons, inelastic neutron scattering was performed on the powder samples. Fig. 5 (a) presents the contour plot of the dynamical structure factor ($S(Q, E)$) for $x = 0$ obtained at 100 K with incident energy $E_i = 17.255$ meV. The calculated phonon dispersion for the $P6_3$ low-temperature structure of $x = 0$ is shown in Fig. 5 (g). Fig. 5 (f) shows the simulated $S(Q, E)$ based on the phonon calculation results. Overall, the simulation well reproduces the experimental result for $x = 0$. The prominent scattering intensity observed at ≈ 5 meV in Fig. 5 (a) mainly comes from the optical phonons with a flat dispersion at the corresponding energies in Fig. 5 (g). The weak streaks observed at $2.5, 2.8, 3.2, 3.8,$ and 4.2 \AA^{-1} are ascribed to the dispersive acoustic modes at low energies.

Interestingly, as shown in Figs. 5 (b)–(e), the energy distribution of the INS intensity deforms and shifts toward the lower-energy region as x increases. The insets of Figs. 5 (a)–(e) show the spectra in the low-energy region measured with $E_i = 8.485$ meV. The intensity in the low-energy region exhibits increasing trend toward $x = 0.07$. This trend is more explicitly observed in Fig. 5 (h), where each spectrum is integrated over the energy transfer $E = 1\text{--}2.3$ meV. To discuss the intensities quantitatively, they are normalized using a coherent cross-section of the 111

nuclear peak. Two peaks at $Q = 2.5$ and 2.8 \AA^{-1} arise from the acoustic modes, and the intensities strongly depend on x and show the largest value near $x = 0.07$. In the inset, the intensity at $Q \approx 2.5 \text{ \AA}^{-1}$ is plotted as a function of x . After showing a maximum near $x = 0.07$, it decreases with increasing x .

Fig. 5 (i) shows the $S(E)$ with $E_i = 17.255$ meV integrated over $|Q| = 2\text{--}4 \text{ \AA}^{-1}$. The calculation well reproduces the experimental result for $x = 0$. Several peaks, indicated by filled triangles, arise from the optical modes. In addition, the most prominent peak has a shoulder, as indicated by an open triangle, which is ascribed to the acoustic modes. At $x = 0.03$, the sharp peaks arising from the optical modes are significantly smeared out. They entirely disappear at $x = 0.07$, and the spectrum shows a broad character.

The glassy nature observed in the INS spectra should also appear in the thermal conductivity, κ . Figs. 6 (a) and (b) show the temperature dependence of κ . Generally in crystals, it behaves as $\kappa \propto T^\alpha$ ($\alpha = 3$) at the lowest temperatures, followed by $\kappa \propto \exp(E/T)$ at several 10 K, and accordingly shows a peak at ≈ 20 K. This characteristic peak is clearly observed for the end members, as indicated by filled triangles in Figs. 6 (a) and (b), both of which have the long-range structures of $P6_3$ and $P2_1$, respectively (see also Fig. 2 and Fig. S2 in the Supplemental Material [13]). Values of 2.3 and 2.6 are

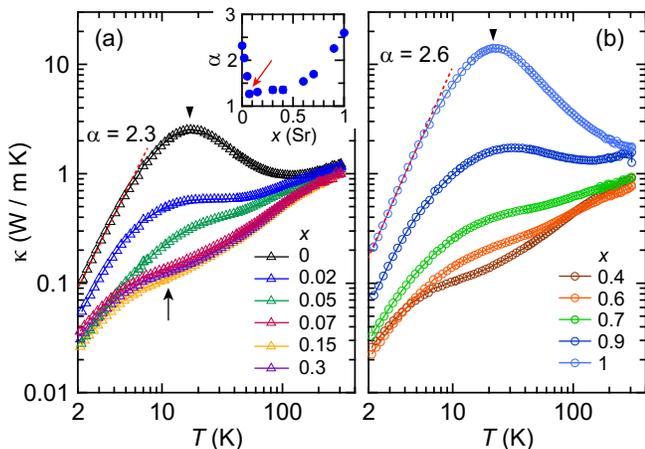


FIG. 6. Log-log plots of the thermal conductivity as a function of temperature for $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ of (a) $x = 0-0.3$ and (b) $x = 0.4-1$. Error bars are smaller than the symbol size. Broken lines show the power fit using $\kappa \propto T^\alpha$ for the experimental data of $x = 0$ and $x = 1$. A plateau is observed for $x = 0.07-0.4$ near 10 K, as indicated by an arrow. The inset shows the power exponent α plotted against x .

yielded for the low-temperature exponent α for $x = 0$ and 1, respectively. The smaller values than the ideal $\alpha = 3$ are explained by the isotope effect [27]. The peak observed for $x = 0$ is suppressed toward $x = 0.07$ and entirely disappears above this composition. In the composition window of $x = 0.07-0.4$, a plateau-like region appears, as indicated by an arrow, which is the typical characteristic observed in amorphous solids [28, 29]. The large peak resurges as x increases further. That is, this plateau-like behavior is only observed in the middle composition range where the low-temperature structures are suppressed.

IV. DISCUSSION

The transition point where the structural phase transition occurs at $T = 0$ has been known as the structural quantum critical point (sQCP) [30–32]. The complete suppression of the superlattice peak observed in Fig. 3 (a) indicates that the sQCP lies in the composition window of $x = 0.06$ and 0.1. The absence of the structural phase transition has also been confirmed in dielectric measurements down to 2 K [23]. The inset of Fig. 6 (a) summarizes the temperature exponent α of the low-temperature κ . It shows a dip near $x = 0.07$, as indicated by an arrow in the inset. In addition, the low-temperature specific heat in our previous work considerably increases and exhibits a maximum near $x = 0.1$, and the Debye temperature (Θ_D) shows a dip [13, 24].

The structural quantum material $(\text{Sr}_{1-x}\text{Ca}_x)_3\text{Rh}_4\text{Sn}_{13}$ has also been reported to show a dip in α and Θ_D near the sQCP at $x \approx 0.9$ [30, 33]. In addition, the

low-temperature specific heat of this system considerably increases toward the sQCP [34]. These characteristics are entirely consistent with those observed in the $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$ system. Similar behavior has also been reported for the Θ_D of $\text{LaCu}_{6-x}\text{Au}_x$ near the sQCP at $x \approx 0.3$ [31]. Thus, the observed behavior of the thermal parameters and the complete suppression of the superlattice reflection testify to the existence of the sQCP in $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$. The INS results shown in the inset of Fig. 5 (h) indicate that the abundance of low-energy phonons should be the essential effect of the sQCP.

In the energy dependence of the INS spectrum for $x = 0$ shown in Fig. 5 (a), several peaks are observed at specific energies due to strong scattering by optical phonons. This peak structure disappears at $x \geq 0.07$ and deforms into broad spectra resembling a boson peak commonly observed in amorphous solids [35–37]. That is, the broad spectra observed at $x \geq 0.07$ indicate that glasslike static disorder is realized along with the long-range periodic structure.

Because the PDF analyses reveal that the Ba(Sr) atoms are at the average position, the glasslike disorder is ascribed to the AlO_4 network in which the Al-O bond lengths deviate from the average value. This fact means that while Ba(Sr) atoms retain a rigid framework adapting to the $P6_322$ symmetry, the AlO_4 network forms a continuum of short-range correlations of AlO_4 tetrahedra and partially breaks the translational symmetry at higher Sr concentrations than the sQCP. Because of this glassy Al-O network, the thermal conductivity for samples with $x = 0.07-0.4$ behaves as if they were amorphous solids. As described in the Introduction, the $P2_1$ phase evolves at $x > 0.5$. As a result, the glassy network disappears, and thereby, the peak in the thermal conductivity reappears.

Generally, in crystalline solids, the phonon mean free path is so large at low temperatures that the value of κ varies with the number of phonons, which depends on temperature as $\propto T^3$. In other words, at low temperatures, more phonons should carry more heat, resulting in a larger value of the thermal conductivity. However, the thermal conductivity decreases toward the sQCP composition despite the increased number of low-energy phonons, as shown in Fig. 5 (h). This contradiction indicates that the phonon mean free path decreases near the sQCP because of the glassy Al-O correlation, which likely suppresses phonon propagation.

The structural characteristic of this peculiar sublattice glass state can be understood as follows. Toward the sQCP composition, the microstructure of the ferroelectric phase is divided into small regions [23] with an average size of ≈ 20 nm (Fig. 3 (c)). At higher Sr concentrations than the sQCP, the soft mode presumably incoherently freezes inside the small islands, which has been observed in our previous transmission electron microscopy (TEM) experiments [11]. As shown in Fig. 1 (e), the $P6_322$ space group possesses the $4e$ site above and below the $2b$ site along the c axis. This $4e$ site can

accommodate the Ba(Sr) atoms even though the mode freezes incoherently and the atoms significantly deviate from the ideal position (the $2b$ site). As a result, the local structure of the Ba(Sr) atoms coincides with the average structure described with the split atom model and preserves the periodicity across the crystal.

In contrast, the incoherently frozen O1 atoms cause a large displacement even from the $6h$ site of the split atom model (Fig. 1 (f)) because they are rotating around the $2d$ site in the vibration of the soft mode. O1 atoms have been reported to exhibit an extremely large DW factor even in structural refinement using the split atom model, while O2 atoms exhibit a sufficiently small DW factor [12]. This large displacement from the $6h$ site can reasonably be considered to be detected in the PDF analysis as a deviation from the average structure model. Such a local environment changes the AlO_4 network from a long-range periodic structure to a continuum of Al and O correlations with various Al-O1 bond orientations, giving rise to the sublattice glass state in the AlO_4 network. The atomic displacement of the Al atom is very small in the vibration pattern of the soft mode [9]. Accordingly, the DW factor of Al atoms does not largely increase in the Sr-incorporated samples [12]. This fact is also consistent with the picture of the incoherently frozen mode discussed above.

Figure 5 (i) also indicates that the sublattice glass state significantly changes the nature of phonons of this material. Generally, the damping of phonons is large when the phonon lifetime is reduced, and larger damping gives broader phonon peaks in the phonon spectra. As shown in Fig. 5 (i), all the phonon peaks indicated by triangles are smeared out with increasing x and thereby significantly broadened, indicating phonon damping. That is, the partial breakdown of the three-dimensional translational symmetry in the AlO_4 network dampens all the phonons of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$. Damping of phonons has been theoretically predicted to increase the low-energy vibration states [38, 39], which is entirely consistent with our experimental observations.

Low-energy phonons have recently attracted increasing interest for layered thermoelectric materials with mobile ions at high temperatures [40–42], which exhibit very low thermal conductivity due to the local vibration of the disordered interlayer ions. In the cage compounds, such as filled skutterudites and clathrate compounds [43, 44], the so-called rattling mode is characterized as a localized Einstein mode with a certain vibration frequency of ≈ 5 meV [45], which also contribute to reduce thermal con-

ductivity. In principle, the sublattice glass state revealed in this study is also applicable for other materials, including semiconductors and metals, with a tunable soft mode. Our findings should be a new strategy to explore low- κ materials.

In the stream of studies focusing on the breakdown of spatial- and time-reversal point-symmetry to explore exotic quantum phases, this work reveals for the first time that the translational symmetry can also be partially broken in crystals. Recent theoretical work has predicted that anharmonic phonon damping plays a key role in the superconducting dome at a ferroelectric quantum critical point [46]. The structural nature and the phonon damping observed in this study should pave a new path to pursue unusual quantum phases and phenomena in the future.

V. CONCLUSIONS

By using the PDF analyses and INS measurements, we have demonstrated that short-range correlation with a broken translational symmetry is realized at higher Sr concentrations than the sQCP of $\text{Ba}_{1-x}\text{Sr}_x\text{Al}_2\text{O}_4$. In this state, the AlO_4 network forms a continuum of Al-O short-range correlations, while the original translational symmetry is preserved in the Ba(Sr) sublattice. Incoherent condensation of the soft mode at the sQCP is most likely responsible for this sublattice glass state. This state significantly dampens the phonons in the whole energy range, resulting in glassy thermal properties.

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