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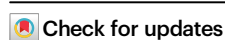


Local inversion-symmetry breaking in a bismuthate high- T_c superconductor

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The doped perovskite BaBiO_3 exhibits a maximum superconducting transition temperature (T_c) of 34 K and was the first high- T_c oxide to be discovered, yet pivotal questions regarding the nature of both the metallic and superconducting states remain unresolved. Although it is generally thought that superconductivity in the bismuthates is of the conventional s -wave type, the pairing mechanism is still debated, with strong electron-phonon coupling and bismuth valence or bond disproportionation possibly playing a role. Here we use diffuse x-ray scattering and Monte Carlo modeling to study the local structure of $\text{Ba}_{1-x}\text{K}_x\text{BiO}_3$ across its insulator-metal boundary. We find no evidence for either long- or short-range disproportionation, which resolves a major conundrum, as disproportionation and the related polaronic effects are likely not relevant for the metallic and superconducting states. Instead, we uncover nanoscale structural correlations that break inversion symmetry, with far-reaching implications for the electronic physics. This unexpected finding furthermore establishes that the bismuthates belong to the broader classes of materials with hidden spin-orbit coupling and a tendency towards inversion-breaking displacements.

Similar to other perovskites and perovskite-derived superconducting oxides such as strontium titanate, strontium ruthenate, and the high- T_c cuprates, the basic structural unit in the (nearly) cubic bismuthates is a metal-oxygen octahedron^{1,2} (Fig. 1). The parent compound BaBiO_3 is an insulator with commensurate charge-density-wave (CDW) order that is thought to be caused by either a periodic modulation of the effective bismuth valence² or a Bi-O charge-transfer mechanism with charge localization on the oxygen orbitals^{3,4}. With hole doping, this modulated phase quickly disappears, and a superconducting metal appears at sufficiently high hole concentrations. Due to strong electron-lattice coupling, the CDW in the parent compound involves three-dimensional oxygen-breathing distortions of the octahedra (Fig. 1d). One of the prevailing hypotheses regarding the doped bismuthates is that short-range CDW correlations survive at high doping levels^{3,5-7} and that these can be viewed as polarons-complex bound states of charge carriers and oxygen breathing distortions⁸. The

metallic phase would then amount to a polaronic liquid, with the polarons binding into bipolarons that, in turn, condense to form a superfluid at T_c ^{3,9}. On the other hand, it has been argued that superconductivity arises through a conventional electron-phonon pairing mechanism^{10,11}, from a homogeneous metallic state well-described by Fermi-liquid theory. Microscopically, the two pictures are radically different, yet conclusive experimental evidence for either scenario is lacking.

Here we use diffuse x-ray scattering and Monte Carlo modeling to study short-range structural correlations in $\text{Ba}_{1-x}\text{K}_x\text{BiO}_3$ (BKBO). We find that static and low-energy breathing distortions are absent in samples with a sufficiently high potassium (and, hence, hole) concentration, on either side of the insulator-metal boundary. This places strong constraints on any local CDW scenario and the associated polaronic effects. However, we uncover correlations of a different nature that involve anticorrelated bismuth and oxygen shifts and break

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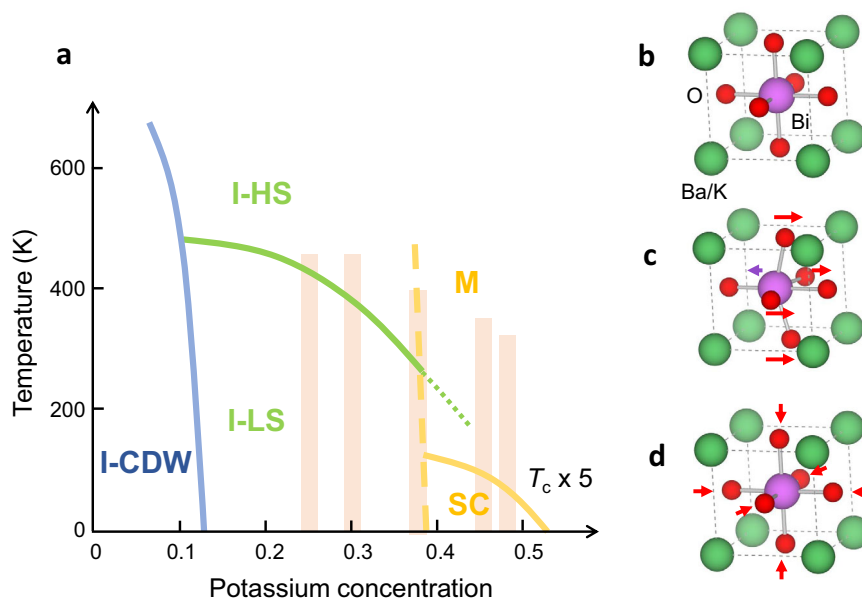


Fig. 1 | Phase diagram and unit-cell distortions in $\text{Ba}_{1-x}\text{K}_x\text{BiO}_3$. **a** Schematic phase diagram based on refs. ^{2,7,20}. In the insulating (I) phase, there exists a commensurate charge-density wave (I-CDW) up to $x \sim 10\text{--}15\%$. Metallic (M) and superconducting (SC) phases appear above $x \sim 35\%$. At intermediate K concentrations, long-range CDW correlations are absent⁷; the high-temperature structure (I-HS) is cubic, whereas long-range rigid octahedral tilts lead to tetragonal and orthorhombic distortions at low temperatures (I-LS; see the “Methods” section and

refs. ^{7,20} for details). Weak long-range tilts persist in the metallic phase (green dotted line). Shaded bands indicate estimated potassium concentrations for the samples studied in this work. **b** Cubic unit cell, that corresponds to the average structure at high doping/temperature. **c** Schematic inversion-breaking distortion; two of the six oxygen atoms do not change position. **d** Schematic oxygen-breathing distortion. The long-range CDW state (I-CDW) involves a three-dimensional alternating pattern of positive and negative breathing distortions.

inversion symmetry (Fig. 1c). We show that these distortions are the result of a complex energy balance that involves screened ionic interactions, precipitated by random potassium–barium substitution. The short-range structural symmetry is therefore considerably lower than the average symmetry, implying that BKBO is a locally non-centrosymmetric superconductor. This finding raises the possibility of unanticipated channels for electron–lattice interactions, with similarities to materials with hidden spin–orbit coupling^{12,13} and systems without global inversion symmetry¹⁴ or close to an inversion-breaking instability^{15–17}.

Results

Figure 2 shows representative diffuse scattering data for two BKBO samples on either side of the metal–insulator transition. Qualitatively, the insulating samples show remarkably little diffuse scattering overall, indicative of nearly absent short-range correlations. In contrast, distinct features are immediately obvious in the metallic samples, with a notable asymmetry of the diffuse intensities around the Bragg peaks—the scattering is always stronger on the low-wavevector side of the peaks. This is a signature of distortions induced by local lattice contraction or expansion, known generically as size-effect scattering¹⁸. Yet in our case this is not simply due to Ba–K substitutional disorder, since the ionic radii of Ba and K are nearly the same. In fact, the K^+ ion is slightly larger than Ba^{2+} while having a smaller x-ray form factor, which should lead to size-effect scattering that is stronger on the high-wavevector side¹⁸. Moreover, the asymmetric scattering shows a nontrivial doping dependence, with a strong increase in the metallic phase, and weak temperature dependence in the studied range (Fig. 2e). In contrast to Pb-doped BaBiO_3 , which displays metastable structures and structural phase separation on the 10–20 nm scale^{2,19}, BKBO is thought to be single-phase in the metallic/superconducting state^{2,20}. We note that the intensity maximum around $x \approx 45\%$ could be partially due to experimental uncertainties (see the “Methods” section) and that the $x \approx 38\%$ sample, which lies close to the insulator–metal boundary, might have

a nonzero insulating volume fraction, which would decrease the integrated intensity.

Importantly, no diffuse signal is present around $(\frac{1}{2} \frac{1}{2} \frac{1}{2})$ and equivalent positions in any of the studied samples (Supplementary Fig. S1). This demonstrates, within our experimental sensitivity, the absence of short-range breathing distortions that double the quasi-cubic unit cell. Instead, in insulating samples, we observe sharp, weak superstructure peaks that originate from a qualitatively different structural distortion—small long-range octahedral tilts—known from previous work²⁰. Diffuse x-ray scattering is an energy-integrating probe and, in principle, sensitive to distortions at any energy scale. Yet the corresponding structure factors decay fairly quickly with the energy of lattice excitations²¹, and dynamic distortions thus become difficult to observe above 10–20 meV. We therefore cannot rule out the presence of breathing distortions at higher energies, and breathing phonon modes are certainly present (between 70 and 80 meV¹⁰). However, the absence of low-energy distortions still places stringent constraints on polaronic theories of the metallic state, especially for small polarons with high binding energies and low mobilities, where significant distortions would be expected⁸. Moreover, our result for the insulating phase is certainly not consistent with a frozen polaronic state.

We can achieve a more refined understanding of the local structure by generating a three-dimensional vector pair distribution function (3D- Δ PDF) using established punch-and-fill methods to eliminate contributions from the average structure (see the “Methods” section). The 3D- Δ PDF therefore only shows short-range pair correlations that deviate from the average structure and enable model-free insight into the real-space local structure^{22,23}. For a given distance between a pair of atoms, the 3D- Δ PDF is positive if the probability to find the atoms at that distance is higher than for uncorrelated, random displacements from the average structure, and negative if this probability is lower. BKBO is particularly well-suited to deduce the local structure from the 3D- Δ PDF due to its simple average structure and a small number of atoms per unit cell. Two-dimensional cuts through the 3D- Δ PDF in two representative planes are shown in Fig. 2 for the $x \approx 45\%$ sample, with

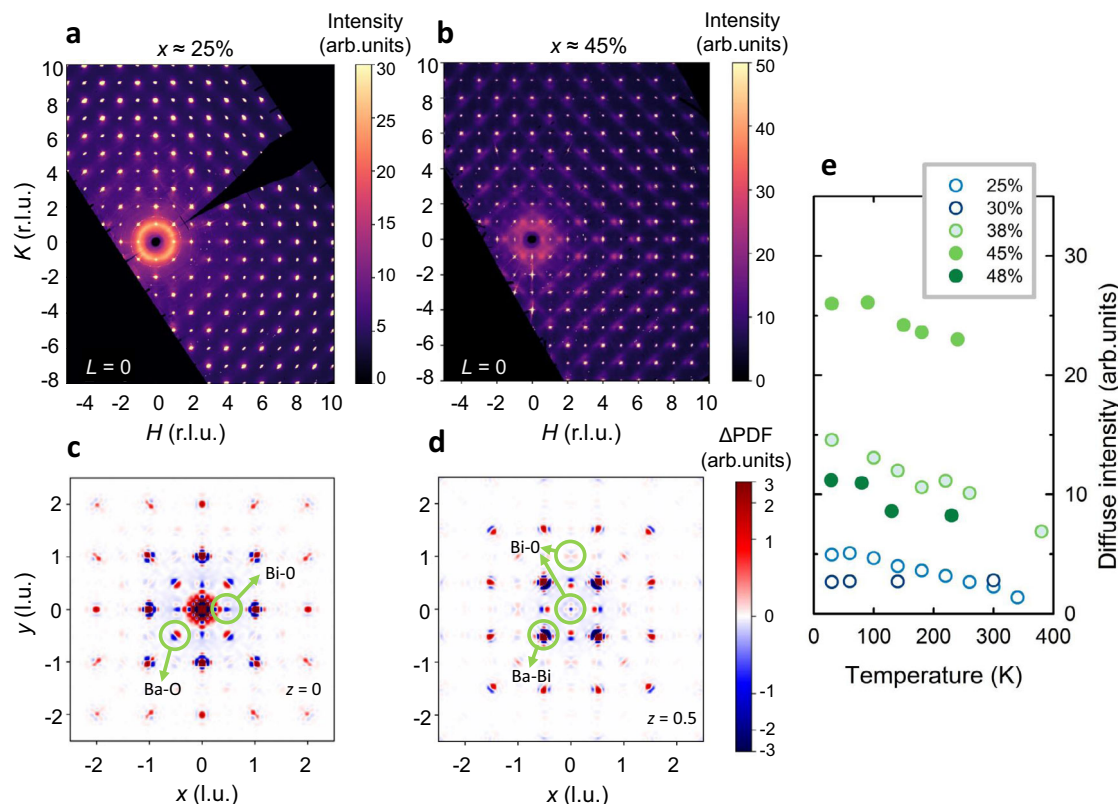


Fig. 2 | Local structure of $\text{Ba}_{1-x}\text{K}_x\text{BiO}_3$. **a, b** Normalized diffuse x-ray intensity in the $L = 0$ plane at 30 K for (a) $x \approx 0.25$, in the insulating state, and (b) $x \approx 0.45$, in the metallic state; r.l.u. refers to reciprocal lattice units. Structured diffuse scattering that is asymmetric around the Bragg peaks is clearly seen in the metallic sample. **c, d** Vector pair distribution functions (3D- Δ PDF) obtained for the sample with $x \approx 0.45$ (same as in **b**), shown in two representative planes ($z = 0$ in **c** and $z = 0.5$ in **d**), with a number of Ba-O, Ba-Bi, and Bi-O correlation peaks annotated; l.u. refers to lattice units. Correlations involving K are suppressed compared to Ba because of the significantly smaller x-ray form factor of K. Red and blue indicate higher and lower probabilities of finding the respective pair of atoms at a given distance, respectively, compared to the distances that correspond to the average structure. Three main features are clearly seen in the 3D- Δ PDF: dipole-like Ba-O and Ba-Bi

peaks, which imply that Ba and O atoms tend to be closer together, and Ba and Bi atoms further apart than in the average structure; and Bi-O peaks that suggest that the oxygen displacements are perpendicular to the Bi-Bi bond. An inversion-breaking local distortion can explain all these features. **e** Temperature dependence of the asymmetric component of the diffuse scattering across the insulator-metal boundary (one-dimensional cuts for the $x \approx 0.45$ sample are shown in Supplementary Fig. S3). Open and full symbols indicate insulating and metallic compositions, respectively; the $x \approx 0.38$ sample is close to the phase boundary. The intensities in **a, b**, and **e** are normalized to nearby Bragg peaks; although it is difficult to extract absolute intensities from x-ray data (see the “Methods” section), a strong increase in relative intensity is clearly seen in the metallic phase.

distinct pair correlations annotated. It is immediately clear that the characteristic length of the correlated displacements is short: the signals decay significantly within 2–3 unit-cell distances, i.e., about 1–2 nm (see also Supplementary Fig. S2). It is straightforward to identify most of the features in the 3D- Δ PDF, especially at non-integer positions where there is no multiplicity. In the $z = 0$ plane, Ba-O correlation peaks are clearly visible, and the dipolar shape of the features shows that Ba and O atoms are more likely to be closer to each other than in the average structure. Note that K-O correlations would appear at the same positions, but the Ba-O signal dominates due to the much larger form factor of Ba. In contrast, Bi-Ba correlations (visible in the $z = 1/2$ plane) show that Bi and Ba are more likely to be further apart. Finally, Bi-O correlations do not involve appreciable changes in the interatomic distance, but rather a displacement of oxygen perpendicular to the average Bi-Bi line. This is seen as a four-fold symmetric pattern around $(0\ 0\ 1/2)$, and also quite clearly in the next-nearest-neighbor Bi-O correlations around $(0\ 1\ 1/2)$ and equivalent positions.

Two important conclusions can be drawn from the 3D- Δ PDF. First, features that would correspond to breathing distortions of the Bi-O bonds are absent, as already clear from the reciprocal space data (Supplementary Fig. S1). Second, the only way to reconcile the Bi-O correlations with the opposite signs of the Ba-O and Ba-Bi correlations is a displacement pattern that breaks inversion symmetry. In local

environments where Ba and K are not symmetrically arranged, the inversion-breaking Bi-O distortion of Fig. 1c occurs, with the oxygen atoms shifting toward the Ba-rich side of the unit cell, and the Bi atoms shifting away from it (but with a much smaller amplitude). The Ba ions then adjust their positions to be further away from Bi, but the oxygen shift is larger than the Ba shift, so the Ba-O distance is still smaller than in the average structure. Importantly, purely rigid BiO_6 octahedra tilts would not lead to pronounced Ba-O size-effect features (that are even visible, e.g., in the next-nearest-neighbor Ba-O peaks around $(1\ 1/2\ 1/2)$), but would yield a symmetric Mexican-hat feature centered at the origin in the 3D- Δ PDF²². Yet it is possible that a local rigid tilt component is superposed with the inversion-breaking displacement.

We obtain further microscopic insight through classical Monte Carlo modeling of the local structure. First, we explore a toy model of (static) short-range CDW correlations using an effective Ising-type Hamiltonian based on Bi valence disproportionation. The bismuth valences +3 and +5 play the role of Ising spins, and K substitution disrupts the CDW since it locally favors the +5 state. Octahedral breathing distortions are then introduced by using a valence-dependent interaction and relaxing the Bi-O distances using Monte Carlo importance sampling (see the “Methods” section). For the sake of computational efficiency, the model neglects the possibility of partial charge transfer between different BiO_6 octahedra and operates

only with discrete values of charge perfectly localized on Bi atoms. Though likely unrealistic, this assumption still serves our purpose, since the relevant diffuse features will mostly be determined by the range of CDW correlations and the octahedral breathing amplitude, and not by the exact Bi oxidation states. The charge should thus be viewed as an effective variable. The free parameters of the model (see the “Methods” section) can be adjusted so that long-range CDW order disappears at low K concentrations, as expected, but short-range correlations survive up to higher K concentrations. The CDW correlations are extremely short-ranged in this regime, with correlation lengths of ~ 2 unit cells. Yet we invariably find diffuse peaks at $(\frac{1}{2}\frac{1}{2}\frac{1}{2})$ and equivalent reciprocal space positions (Supplementary Fig. S1), i.e., a signature of local CDW correlations that is clearly absent in our data.

In order to capture the observed diffuse scattering features, we then construct a minimal model that only includes Bi and Ba/K displacements, under the assumption that all Bi atoms are equivalent. Furthermore, it is assumed that local lattice distortions are caused by an incomplete screening of the local ionic charges, which leads to electrostatic repulsion that is stronger for Bi–Ba pairs than for Bi–K pairs. This is achieved with a screened Coulomb (i.e., Yukawa) potential to describe ionic interactions (see the “Methods” section). In the relaxed structure, Ba and Bi then prefer to be further apart than K and Bi (Fig. 3b). Since the scattering is dominated by the heavy atoms, the main reciprocal space features are well reproduced (Fig. 3a), with stronger diffuse scattering on the low-wavevector side of the Bragg peaks. We note that we have not attempted a detailed quantitative comparison of the data and modeling, since the simple minimal model should not be expected to completely reproduce the measured diffuse scattering. In particular, different forms of the interaction potential lead to slightly different diffuse patterns. A full quantitative refinement that takes this into account would be difficult to perform reliably and is therefore left for future work.

We further investigate the role of oxygen by extending the minimal model to include oxygen displacements perpendicular to the Bi–Bi direction. Using the bond valence sum method to estimate the oxidation states of Bi, Ba, and K (see the “Methods” section), we find Bi and K oxidation states in reasonable agreement with their nominal values, whereas Ba is significantly underbonded. At low K

concentrations, the system compensates for this through symmetry-lowering structural transitions⁷, as permitted by the known size mismatch of Ba to the surrounding oxygen cage (Supplementary Fig. S4). On the other hand, at sufficiently high K concentrations, long-range tilt formation is disrupted by randomly distributed K atoms. Inversion-breaking oxygen displacements then become an alternative way to compensate for the underbonding of Ba in unit cells with an asymmetric arrangement of Ba and K. We include this in the Monte Carlo modeling by employing the squared bond valence mismatch to penalize chemically unstable coordination environments (see the “Methods” section). The obtained local structure contains a mixture of rigid tilts and inversion-breaking displacements that decrease the Ba–O distances (Fig. 3c), in agreement with the 3D- Δ PDF analysis. The inversion-breaking distortions are found to increase with increasing K concentration (Supplementary Fig. S6). We emphasize that the random K/Ba substitution and short-range electrostatic interactions are crucial in generating the distortions.

Discussion

Our results establish that bismuth valence disproportionation and oxygen breathing-mode polarons are likely not relevant to superconducting BKBO. However, other electron–phonon composite states could be present, particularly states that involve the inversion-breaking Bi–O phonon mode. Ab initio studies suggest that the coupling of electrons to the longitudinal Bi–O mode around 20 meV is in fact stronger than the coupling to high-energy oxygen breathing modes¹⁰, which may lead to polaronic physics. The appearance of a broad infrared peak in the optical conductivity²⁴ is consistent with electron–phonon bound states. However, it is also possible that the local inversion-breaking distortions are predominantly precipitated by structural properties, with electron–phonon coupling playing a secondary role and a more conventional metallic state. This is supported by the increase of inversion-breaking correlations with doping in our simulations, which do not explicitly include electron–phonon coupling. A related, important question is the interplay between the inversion-breaking displacements and rigid octahedral tilts—the latter are caused by atomic-size mismatch and decrease with doping, whereas the former clearly increase substantially in the metallic phase

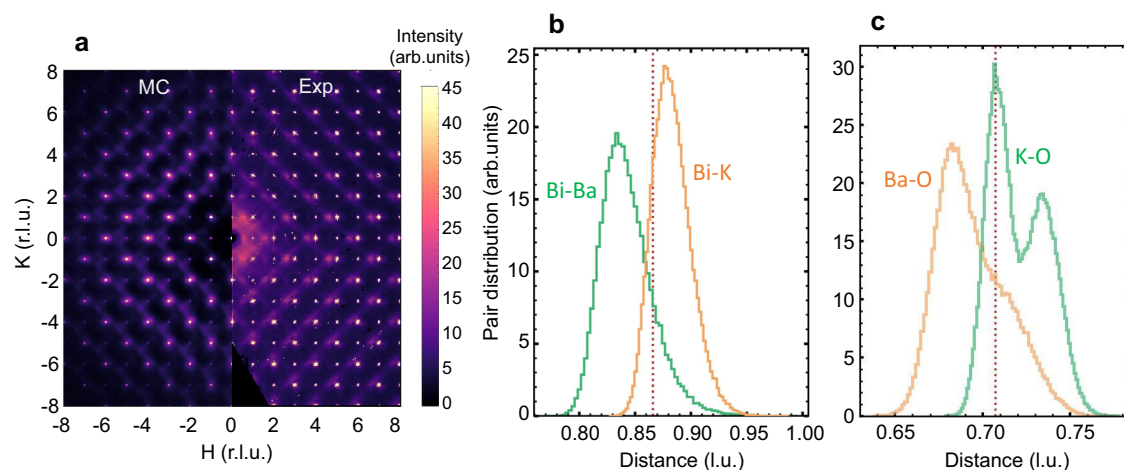


Fig. 3 | Monte Carlo modeling of the local structure. **a** Calculated reciprocal-space image (left half) from a simple model (see the “Methods” section) that only includes negative Bi–Ba and positive Bi–K correlations; r.l.u. refers to reciprocal lattice units. The model captures the main features observed in the experiment, in particular the diagonal rods of scattering and the asymmetry of the diffuse intensity around Bragg peaks (right half). **b** Distribution of Bi–Ba and Bi–K distances within the model, showing a significant difference, which is the microscopic origin of the observed asymmetric diffuse features. Vertical dashed lines indicate the distances

in the average structure. **c** Distribution of nearest-neighbor Ba–O and K–O distances in an expanded model that includes oxygen displacements (see the “Methods” section). Due to inversion-breaking distortions, the Ba–O distances are shorter than average, in agreement with the experimental 3D- Δ PDF result. The calculated O–O distance distribution, which is not directly observable in the experiment due to small oxygen x-ray form factors, is shown in Supplementary Fig. S5.

(Fig. 2e). It is, therefore, possible that the rigid tilts compete with the polar displacements, which might then only proliferate below a critical tilt value, with a direct relation to the insulator–metal transition. An analogous interplay between octahedral tilts and electronic properties has been observed in, e.g., certain cuprates²⁵. Targeted uniaxial-strain experiments to influence the tilts, as recently demonstrated in rare-earth titanates²⁶, might provide more information about this aspect of bismuthate physics, especially given the possible role of tilts in enhancing electron–phonon coupling²⁷. Another interesting possibility is that screening by mobile carriers is essential to obtain local distortions. Our calculations use the screened Yukawa potential for electrostatic interactions, which severely limits their spatial range and permits inhomogeneity on the scale of the screening length. This is appropriate in the metallic phase, but not in the insulating phase, where long-range Coulomb interactions likely favor configurations with significantly less short-range charge disorder, which might explain the enhancement of diffuse scattering for metallic compositions.

The existence of local correlated inhomogeneity has profound implications for both the metallic and the superconducting states. Similar to other prominent metallic perovskites, such as titanates²⁸, nickelates²⁹, and vanadates³⁰, the resistivity of the bismuthates is approximately proportional to the temperature squared, consistent with Fermi-liquid physics, but with a magnitude that is unrealistically large for electron–electron scattering³¹. It has been proposed that this might generically be understood through the interaction of small polarons with low-lying phonon modes³⁰. The phonon spectrum of the bismuthates contains several optical modes in the 10–20 meV range¹⁰ that could lead to such behavior, but the polarons would probably involve the inversion-breaking displacements instead of breathing distortions. Moreover, local inversion-symmetry breaking can cause electron–lattice interactions that are forbidden in the average high-symmetry structure; electron-phonon coupling in a conventional Fermi liquid can also enhance T^2 resistivity³². Importantly, the Thomas–Fermi screening length in metallic BKBO is about 7 Å and likely sets the structural correlation length, implying that the dipole moment associated with the inversion-breaking distortions is not completely screened. A Rashba-type electron–phonon coupling that is linear in phonon amplitudes might be important for both normal-state properties and superconducting pairing, as proposed for materials close to a polar instability, such as strontium titanate^{15,33}. Moreover, since the inversion-symmetry breaking appears at length scales comparable to the superconducting coherence length² of 30–60 Å, the bismuthates can be viewed as locally non-centrosymmetric superconductors. This indicates that the pairing might be more exotic than just simple *s*-wave, with admixtures of odd-momentum gap functions allowed by symmetry¹⁴ and the appearance of local time-reversal symmetry breaking. Finally, it will be interesting to see if similar effects are present in BaPb_{1-x}Bi_xO₃ and the newly discovered isostructural antimony-based superconductor³⁴ (Ba,K)SbO₃ and in other substitutionally doped oxides.

Methods

Samples

Ba_{1-x}K_xBiO₃ crystals were grown using a low-temperature electrocrystallization method similar to previous work^{35,36}. The growth precursor consisted of 140 g KOH, which acts as flux, 10.5 g Bi₂O₃, and 3.5–20 g Ba(OH)₂·8H₂O. The precursor materials were combined in a PTFE crucible, melted at 400 °C, and homogenized by a magnetic stirrer at 60 rpm in a humid nitrogen atmosphere. A small amount (2–4 g) of KNO₃ was added to counteract the degradation of the crucible³⁶. Platinum electrodes were inserted into the molten solution and a constant current between 0.1 and 1 mA was supplied for 18 h. BKBO crystals deposit on the anode, and excess bismuth deposits on the cathode. The amount of Bi₂O₃ far exceeds the solubility limit of

KOH at the beginning of growth, enabling the Bi₂O₃ concentration to remain constant for the duration of the growth. The ratio of Ba(OH)₂·8H₂O to KOH controls the doping level of the deposited crystal. Using this method, crystals with potassium content between $x = 0.25$ and 0.51 were grown.

Sharp superconducting transitions from dc magnetization and ac susceptibility (including nonlinear susceptibility³⁷ measurements were used as an indicator to select metallic samples. For both metallic and insulating samples, high crystalline quality was established from sharp, well-defined Bragg peaks in the x-ray data. The potassium concentration was estimated from lattice parameter values obtained during the refinement of the diffuse x-ray scattering data, and by employing the known linear relation between doping and unit cell size⁷ at 300 K. In the case of metallic samples, these values were found to be consistent with those determined from the well-defined relationship between T_c and doping⁷.

Diffuse x-ray scattering

The measurements were performed on beamline 6-ID-D of the Advanced Photon Source, Argonne National Laboratory, USA. The beamline uses a superconducting magnet undulator insertion device and a double-crystal monochromator to produce a monochromatic high-energy x-ray beam. We used a photon energy of 87 keV, slightly below the Bi K-edge at 90.526 keV, which likely suppresses the relative intensities of the diffuse scattering/PDF peaks that involve Bi. A Pilatus 2 M CdTe detector with a sensor layer optimized for high energies was used to collect frames with 0.1 s exposure time, while samples were continuously rotated at 1° s⁻¹ about a horizontal axis. Three sets of rotations were performed in each measurement, between which (i) the detector was translated by 5 mm in both the horizontal and vertical directions to cover gaps between detector chips, and (ii) the sample rotation axis was offset by ±15° from perpendicular to the beam to allow masking of artifacts caused by scattering in the sensor layer. The overall counting time at each temperature setting was 20 min. See Supplementary Material in ref. ²³ for more details. Data covering a range of about ±15 Å⁻¹ in all directions were collected down to 30 K using an Oxford N-Helix helium cryocooler, and transformed to S(**Q**) using the software package CCTW (<https://sourceforge.net/projects/cctw/>).

3D-ΔPDF

The 3D-ΔPDF shown in Fig. 2 was generated similarly to ref. ²³ by symmetrizing the $x = 45\%$ data with all operations included in the $m\bar{3}m$ point group, “punching” out spheres of reciprocal space of radius 0.2 r.l.u. around each Bragg position, and “filling” the removed spheres by interpolating with a three-dimensional Gaussian kernel. A Tukey window ($\alpha = 0.5$) was applied to the “filled” data to reduce leakage artifacts in the Fourier transform, and a discrete Fourier transform was then applied to the data. The 3D-ΔPDF shown is the real part of this Fourier transform.

The real-space correlation length of the displacement was obtained from a representative linecut ($0.5 \leq x \leq 6$, $y = 0$, and $z = 0$) of the 3D-ΔPDF. The absolute values of the signal peaks along the linecut were fit to an exponential function (Supplementary Fig. S2). The decay constant was found to be 1.1 unit cells, which indicates a significant decay within about 2 unit cells (about 1 nm).

Temperature and doping dependences of diffuse scattering

The temperature and doping dependences of the asymmetric component of the diffuse signal were obtained as follows. First, a Bragg peak free of any artifacts such as streaking (due to Compton scattering) or powder rings was chosen. For a comparison of relative scattering intensities among different samples it is essential that the same Bragg peak and Brillouin zone be used, as there exists considerable intensity variation across different zones. Because BKBO is nearly cubic at the measured temperatures, there are 8 equivalent sites for

each hkl position in reciprocal space, and the positions ($\pm 4 \pm 6$) were chosen. A one-dimensional (1D) intensity profile along [100] was obtained by summing between ± 0.04 r.l.u. along [010] and ± 0.08 r.l.u. along [001]. The result in Fig. 2e was obtained from fast-Fourier transformations of these 1D datasets, centered on the respective Bragg peak, via

$$y(k) = \sum_{n=0}^{N-1} e^{-2\pi i \frac{kn}{N}} x(n), \quad (1)$$

where N is the length of the 1D array, x is the intensity at index n , and $0 \leq k \leq 41$ (this is well into the range of k where the contribution of each higher value becomes vanishingly small). This transformation returns both real values, which correspond to the amplitudes of the symmetric parts of the signal, and imaginary values, which correspond to the asymmetric scattering intensities. For each sample at each temperature, the asymmetric intensities were summed and normalized by the maximum value in the 1D dataset to obtain a ‘temperature factor’. This normalization functioned quite well for a comparison of datasets for the same sample; however, a second normalization was required to compare results for different samples and to account for differences in illuminated sample volumes and background levels. This normalization consisted of finding the Bragg peak intensity at the same temperature for each sample and scaling each sample’s temperature factors by the ratio of its Bragg peak intensity to that of the sample with the strongest Bragg peak intensity. Notably, this procedure involved uncertainties that were difficult to quantify, most importantly possible partial saturation (nonlinearity) of the detector at the Bragg peaks for the largest samples, which might affect the comparison in Fig. 2e. The relative intensities should therefore be taken as semi-quantitative, with the main, robust result that the diffuse intensity increases in the metallic phase.

Monte Carlo modeling

We used classical Monte Carlo (MC) to simulate short-range ordered structures³⁸, with K randomly substituted for Ba. As a first step, a clear diffuse scattering signature of short-range CDW correlations was established by constructing a simple, yet robust Ising-type model with the Hamiltonian

$$H = J_1 \sum_{\langle ij \rangle} s_i s_j + J_2 \sum_{\langle ij \rangle} \sigma_i s_j, \quad (2)$$

where bismuth oxidation states +3 and +5 are mapped to the respective values +1 and -1 of the spin variable s , the chemical species of nearest-neighbor A-site cations are encoded as $\sigma = +1$ (K) and 0 (Ba), the indices i and j label different unit cells, and the sums are restricted to the nearest-neighbour pairs. Since we aim for a phenomenological model of short-range CDW formation, the coupling constants $J_{1,2}$ can be estimated from the requirement that long-range order disappears for K concentrations above^{15,20} -0.1, while short-range CDW correlations persist at higher doping. Long-range CDW order corresponds to a nonzero expectation value of the staggered magnetization

$$m = \frac{1}{N} \left\langle \sum_{ijk} (-1)^{i+j+k} s_{ijk} \right\rangle \quad (3)$$

where N is the total number of unit cells, and the brackets denote the ensemble obtained via MC importance sampling. Similarly, short-range order is captured by the chemical correlation coefficient³⁹

$$c_{ij} = \frac{P_{ij} - \theta^2}{\theta(1 - \theta)} \quad (4)$$

where P_{ij} is the joint probability that bismuth sites i and j are occupied by the same oxidation state +3 (+5), and θ is the overall concentration of +3 (+5) atoms. Negative values of c_{ij} correspond to situations where the two sites tend to be occupied by different oxidation states, which is used to quantify remnant CDW correlations. The first term in the Hamiltonian ($J_1 > 0$) models the Coulomb interaction between nearest-neighbor charges localized on Bi atoms, and is thus responsible for CDW formation. The second term ($J_2 < 0$) captures the disruption of the CDW order by the presence of K atoms, which play the role of effectively negatively charged impurities and locally favor the +5 state. A representative statistical ensemble of equilibrium configurations was obtained by simulating many $20 \times 20 \times 20$ supercells with periodic boundary conditions, such that the basic move involves a swap of +3 and +5 atoms in order to preserve the effective charge balance, and by using simulated annealing⁴⁰ to prevent the simulation from getting stuck in local minima. Furthermore, we used the coupling between the local charge and octahedral breathing amplitude^{41,42} that is responsible for stabilizing the charge modulation, through the valence-dependent harmonic interaction between Bi and O atoms

$$U_i(r) = \frac{1}{2} K [r - r_0(s_i)]^2 \quad (5)$$

where K is the effective spring constant and $r_0(s_i)$ the Bi–O equilibrium distance for two different oxidation states i . The value of the spring constant $K \approx 19$ eV/Å² can be estimated by fitting the Raman-active 70 meV breathing mode^{42,43}, and the average values of r_0 (+5) and r_0 (+3) in the parent compound are known from previous studies^{2,7,20} to be -2.11 and -2.29 Å, respectively. Since we aimed to reproduce the local structure that would result from short-range CDW, the above valence-dependent potential was sufficient to reproduce the relevant effects and avoided microscopic details of electron–phonon coupling and the intricacies of stability of polaronic and bipolaronic solutions^{3,5}. Importantly, the calculation allows for independent optimization of each oxygen position within a given octahedron and therefore does not assume a specific local symmetry for the breathing distortions. Consequently, all possible distortions that constrain the oxygen motion along the Bi–O bond are allowed. Using MC sampling with positional displacements as the basic move to relax the Bi–O bonds, and subsequently calculating diffuse scattering⁴⁴, we found diffuse superstructure peaks at specific half-integer positions in reciprocal space, which are absent in experiment (Supplementary Fig. S1). The configuration and diffuse scattering shown in Supplementary Fig. S1 was obtained using $J_2/J_1 = 4$ and J_1 equal to the effective Monte Carlo temperature. For this choice of parameters, no long-range order is present, but short-range correlations persist (Supplementary Fig. S1). The results do not significantly depend on the ratio of J_1 to the temperature if the ratio is smaller than or close to one.

The second part of the analysis involved a minimal model that captures the essential features of the observed diffuse scattering (Fig. 2). First, since we find no evidence of charge modulation in the studied K concentration range, we assumed a homogenous charge distribution on the Bi sublattice, with the oxidation state of Bi fixed by charge neutrality ($V_{\text{Bi}} = 4 + x$). The pseudo-cubic lattice constant obtained from this assignment using bond valence sum rules⁴⁵ for the BiO₆ octahedral complex is consistent with $a = 4.35\text{--}0.17x$ Å, known from structural studies⁷, which indicates that the unit cell size is mainly controlled by the strength of the Bi–O bonds and gives credibility to our assumption. Also, the asymmetry of the observed diffuse features indicates that the local lattice distortion is not simply due to slight differences in the ionic sizes of Ba²⁺ (~1.49 Å) and K⁺ (~1.52 Å) (Shannon–Prewitt ionic radii⁴⁶). Therefore, we assumed that the effect is caused by residual electrostatic interactions between different non-bonded ions, where the static screening in the metallic regime is

approximately accounted for by the familiar Yukawa potential

$$\phi_{ij}(r) = \frac{Q_i Q_j}{r} e^{-k_s r} \quad (6)$$

where Q_i represents the ionic charge of the i -th ion and $k_s \approx 0.49 \text{ \AA}^{-1}$ the screening wavenumber²⁰, which sets the length scale of the local lattice distortion. In the insulating state, the screened potential is likely not appropriate, and long-range Coulomb interactions might drastically change the overall energy balance; this is, however, significantly more difficult to simulate, and has not been attempted. The ionic charges were assumed to be equal to atomic oxidation states, which suffices for the desired semi-quantitative insights that we aimed for. To make the model more realistic, we introduced Bi–O, Ba–O, and K–O covalent bonds via the Morse pair potential

$$U(r) = D \left(1 - e^{-(r-r_0)/b} \right)^2 \quad (7)$$

where the equilibrium distance between atoms r_0 is determined by the size of the unit cell. The constant b determines the “softness” of the bond and was taken from⁴⁷. D was obtained by comparing to the known values of atomic displacement parameters^{7,20} (thermal factors); $D_{\text{Bi-O}} = 10$ and $D_{\text{Ba-O}} = D_{\text{K-O}} = 2$ in units of MC temperature. The structure was relaxed by performing the MC thermalization on an ensemble of $15 \times 15 \times 15$ supercells. We found that the calculated diffuse scattering averaged over many equilibrium configurations and symmetrized in the $m3m$ point group agreed well with the data, with the asymmetry on the low-wavevector side of the Bragg peaks reproduced well (Fig. 3a).

In order to complete the picture, we constructed a model of the local structure of oxygen octahedra, whose features are not obvious in reciprocal-space data, but can be inferred from the 3D- Δ PDF. We used the well-known perovskite (ABO_3) tolerance factor

$$t = \frac{r_A + r_O}{\sqrt{2}(r_B + r_O)} \quad (8)$$

as an indicator of structural stability, where⁴⁶ $r_O = 1.26 \text{ \AA}$ is the ionic radius of O^{2-} . For $r_B + r_O$, we used half the lattice constant, since the size of the unit cell is controlled by the covalent Bi–O bond whose length depends on doping through the effective valence of Bi. Using tabulated values of ionic radii of K^+ and Ba^{2+} in place of r_A we obtained the doping dependences of t_{Ba} and t_{K} (Supplementary Fig. S4), and hence an indication of how well the A-site cations are accommodated by the lattice. t_{K} is in the “ideal” range for all K concentrations, whereas Ba is too small for the ideal cubic lattice, which can be compensated for by the formation of long-range rigid octahedral tilts that lead to the sequence of structural transitions^{7,20} ($Pm3m \rightarrow I4/mcm \rightarrow Ibmm \rightarrow I2_1m$) as the K concentration decreases toward zero. However, for higher K concentrations, the formation of long-range ordered tilts is significantly disrupted by the presence of randomly distributed K atoms. The barium size mismatch and the disorder introduced by potassium doping thus lead to an interplay between short-range tilts and inversion-breaking displacements. This is also reflected in the unusually large anisotropic thermal ellipsoids of oxygen⁷, indicating the propensity for large oxygen displacements perpendicular to the Bi–O bond to accommodate the size discrepancy. Similar conclusions can be drawn if we use the bond valence sum (BVS) method^{45,48} to estimate the empirical valence ν of a given atom as a sum of contributions of each bond that it forms (bond valences s_j) in each bonding environment, with $\nu = \sum_j s_j$ the scalar and $\mathbf{v} = \sum_j s_j \mathbf{e}_j$ the vector bond valence sum, where \mathbf{e}_j are unit vectors along the bonds, and the summation is over bonds. The bond

valences are best understood in the context of (mostly) ionic bonds where bond valence plays the role of electric flux so that the BVS rule above simply becomes Gauss’ law. Using the known correlation between bond valence and bond length

$$s_j = e^{\frac{R_0 - r}{b}} \quad (9)$$

where r is the bond length and R_0 and b are parameters tabulated in ref. ⁴⁷, we calculated the doping dependence of the empirical Ba and K valences in the average structure. In the doping range of interest ($x \sim 0.35$ – 0.45), we found K to be fully bonded (ν_K close to nominal valence +1 of K), and Ba to be significantly underbonded (ν_{Ba} smaller than nominal valence +2 of Ba) (Supplementary Fig. S4). This correlates with our previous geometric analysis but also allowed us to formulate a semi-quantitative model where we use the (squared) valence mismatch, also referred to as the global instability index, defined by⁴⁵

$$g = w_s \sum_i (v_i - V_i)^2 + w_v \sum_i (\mathbf{v}_i^2 - \mathbf{V}_i^2)^2 \quad (10)$$

as a measure of structural instability^{48,49}. The two right-hand side terms correspond to scalar and vector contributions, respectively⁴⁹, with weights $w_s = 2$ and $w_v = 1$. V_i and v_i are the scalar nominal and empirical valences of an atom, respectively, while \mathbf{V}_i and \mathbf{v}_i are the respective valences. The nominal vector valences \mathbf{V}_i are taken to be zero^{45,49}. Using g in the calculations naturally captures multi-body atomic correlations and the tendency of oxygen displacements to compensate for the valence mismatch of Ba. To constrain the model, we implemented soft volume exclusion using the Lennard–Jones repulsive term $A(r_{\text{ion}}/r)^{12}$, where r_{ion} is the tabulated ionic radius of a given atom⁴⁶ and $A = 6$ in units of MC temperature. Furthermore, the interplay of tilts and polar displacements required the inclusion of an effective octahedral rigidity constraint, for which we used a harmonic spring potential between neighboring oxygen atoms as the simplest option:

$$U_R(r) = \frac{1}{2} k (r - r_0)^2, \quad (11)$$

where r is the O–O distance, r_{eq} is their distance in the average structure, and the spring constant $k = 10$ (in MC temperature units) was chosen to be sufficiently large, thus penalizing, but not precluding non-rigid displacement patterns of the oxygen octahedra. Given the qualitative nature of the model, the exact proportion of rigid tilts as opposed to non-rigid polar distortions is hard to estimate, as it heavily depends on the relative weights of g and $U_R(r)$. Still, overall trends in the doping dependence can be discerned either by looking at the distribution of O–O distances (Supplementary Fig. S5) or the local centrosymmetry parameter, defined as⁵⁰

$$p_{\text{CSP}} = \sum_1^{N/2} |\mathbf{r}_i + \mathbf{r}_{i+N/2}|^2 \quad (12)$$

where N is the number of nearest neighbors of a given atom, and \mathbf{r} are vectors from this central atom to a pair of opposite neighbors. This parameter is a convenient measure of deviations from centrosymmetry, and it increases with K concentration (Supplementary Fig. S6).

Data availability

The raw diffuse x-ray scattering data are available as HDF5 files stored in the NeXus format under restricted access because of their large size, and transfer to third parties can be arranged upon request from the corresponding authors.

Code availability

The code used for Monte Carlo simulations and diffuse scattering calculations is available at <https://github.com/mspaic/BKBO-simulations>.

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Author contributions

M.G. and D.P. conceived the work. S.G., J.J., D.Z., and D.P. grew and characterized the samples. S.G., J.J., Z.W.A., M.J.K., D.P., and R.O. performed the x-ray scattering experiments and analyzed the data. M.S. performed the model calculations. D.P., M.G., S.G., and M.S. wrote the manuscript with input from all authors.

Competing interests

The authors declare no competing interests.

Additional information

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