Skin Layer of BiFeO$_3$ Single Crystals

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A surface layer ("skin") different from the bulk was found in single crystals of BiFeO$_3$. Impedance analysis and grazing incidence x-ray diffraction reveal a phase transition at $T^* \sim 275 \pm 5$ °C that is confined within the surface of BiFeO$_3$. X-ray photoelectron spectroscopy and refraction-corrected x-ray diffraction as a function of incidence angle and photon wavelength indicate a reduced electron density and an elongated out-of-plane lattice parameter within a few nanometers of the surface. The skin will affect samples with large surface to volume ratios, as well as devices that rely on interfacial coupling such as exchange bias.

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Bismuth ferrite, BiFeO$_3$, has become a cornerstone of magnetoelectric research, owing to its high ferroelectric and magnetic ordering temperatures, large ferroelectric polarization, and electrically controllable magnetic easy plane [1–3]. Yet the phase diagram of this archetypal material still remains unresolved, with a growing number of structural and/or functional anomalies reported as a function of temperature that may or may not signal the existence of true phase transitions (see Ref. [3] for a critical review of some of them). Another unaddressed issue concerns the existence of a surface layer (a “skin”) in BiFeO$_3$. Notice that several important perovskites [e.g., SrTiO$_3$ [4,5], BaTiO$_3$ [6,7], or relaxor Pb(Mg$_{1/3}$Nb$_{2/3}$)$_3$O$_3$ [8]] are known to have surface layers that are structurally different from the interior of the crystal. As we shall see, resolving the phase diagram of BiFeO$_3$ and unravelling its surface structure are in fact linked problems: the surface has its own symmetry and undergoes its own phase transitions which interfere with measurements of bulk BiFeO$_3$. Understanding the BiFeO$_3$ skin is also important because most devices proposed for this material rely on interfacial interactions such as exchange bias [2,9–11].

Here we have investigated the electronic and structural properties of the skin layer of BiFeO$_3$ single crystals as a function of temperature by techniques allowing a tunable information depth. We have found a phase transition at $T^* \sim 275 \pm 5$ °C confined within the surface layer of the crystal. Meanwhile, at room temperature, refraction-corrected grazing incidence x-ray diffraction reveals that the skin of BiFeO$_3$ has an elongated out-of-plane lattice parameter (tensile strain of 0.7%) in the topmost few nanometers’ region. Impedance spectroscopy and x-ray photoelectron spectroscopy show that the surface layer of BiFeO$_3$ is charge-depleted.

The crystal used for x-ray investigations was rosettelike with (001) habitus [12], grown using the method proposed by Kubel and Schmid [13]. It was first optically polished using sandpaper and diamond paste; subsequently, in order to minimize the mechanically damaged surface layer while preserving smoothness, a 30 min chemical-mechanical polishing using silica slurry (Syton SF1; Logitech) was employed. The pressure applied during the last stage was less than 100 g/cm$^2$ so as to minimize polishing stress at the surface. A photograph of the crystal is shown in Fig. 1 (inset). Ferroelectric or ferroelastic domains are optically observable using standard birefringence. The quadrant arrangement is consistent with the polarization pointing alternately along the diagonals of the unit cell, although closure cannot be confirmed by birefringence alone. Precedents of ferroelectric flux closure have so far been limited to nanoscopic domains [14]. The local roughness was 0.8 nm, determined by atomic force microscopy, and in agreement with the x-ray reflectivity (Fig. 1) value of (1.3 ± 0.1) nm.

(110)-oriented dendritic crystals were also made as described in Refs. [15–18]. Their ac impedance [Figs. 2(a) and 2(b)] was measured using an Agilent impedance analyzer, model 4294A. The electrodes were sputtered gold, with Pt wires attached to them using silver paste. The impedance shows classic Maxwell-Wagner behavior [19]: a giant and frequency-dependent steplike increase in capacitance, typical of a material with two lossy dielectric
components in series. The bulk of the crystal becomes conducting as temperature increases (hence the large losses), so that the voltage is then mostly dropped on a charge-depleted interfacial barrier layer (contact capacitance); because the interface is thin, the apparent capacitance becomes large [20,21]. At low frequency (where interfacial capacitance dominates), there is a peak whose temperature \(T' = 272 \text{°C}\) is frequency-independent thus signaling a true phase transition. On the other hand, the peak’s absence from the high-frequency curves (where bulk capacitance dominates) implies that the inside of the crystal is not undergoing any change [22]. The \(T'\) peak must therefore correspond to a phase transition confined within the interface. The anomaly is observed also in as-grown unpolished samples (Fig. 2(c)) and therefore the surface transition is not caused by polishing.

Further evidence for an electronically different surface layer comes from x-ray photoelectron spectroscopy (XPS). The Bi \(4f\) core level results are shown in Figs. 2(d) and 2(e). Spectra were acquired at angles of 90° and 30° with respect to the sample surface using small spot Al \(K\alpha\) radiation. Two components are observed in the Bi \(4f\) spectrum indicating two different electronic environments for Bi\(^{3+}\). The higher binding energy component is more intense at glancing angle detection [Fig. 2(d)] than at normal incidence [Fig. 2(e)], showing that it should be associated with the surface or near-surface region. No carbonate species were observed in the C \(1s\) spectra (not shown); thus, the high binding energy component is not due to adsorbates. Higher binding energy is consistent with lower screening and therefore with lower electron density (charge depletion), as also observed in the skin of other ferroelectrics [6]. The Fe \(2p\) spectrum was checked and showed a spin orbit splitting (13.45 eV) and \(2p_{3/2}\) binding energy (710.8 eV), fully consistent with trivalent Fe. The XPS evidence thus points to a distinct electronic environment while maintaining the same chemistry as the bulk BiFeO\(_3\).

Grazing incidence x-ray diffraction allows tracking surface structural changes, with tunable information depth from a few unit cells to several hundreds of microns [23]. The sample in Fig. 1 has been investigated by grazing incidence x-ray diffraction on the six-circle diffractometer at SpLine beam line, ESRF [25]. The sample was placed on a heating stage (25°C-400°C) covered with an airtight Kapton housing filled with 1 bar of pure oxygen to minimize oxygen vacancies.

In order to tune the penetration depth we have devised a dual approach, changing both the incidence angle and the incident photon energy. The justification is that, while either parameter alone permits probing different depths [Fig. 3(b); [23,24]], the parameters also yield refraction-induced peak shifts [26], so that both real (structural) shifts...
and artificial (refractive) ones are convolved. In order to separate them we note that refraction behaves differently as a function of angle and energy [Fig. 3(b)], especially near the critical angle and/or the absorption edge (Bi L\textsubscript{1} at ~13 keV). The penetration depth $\delta$ and refractive shift $\Delta L$ plotted in Figs. 3(b) and 3(c) have been calculated using [23]

$$\delta = 1/\text{Im}(q_z), \quad \Delta L = a_z\text{Re}(q_z - Q_z)/(2\pi),$$

where $a_z$ is the vertical lattice parameter, and $Q_z$ and $q_z$ are the vertical components of the scattering vector in vacuum and in the material, respectively:

$$Q_z = K[\sin(\alpha_i) + \sin(\alpha_f)],$$

$$q_z = K[\sqrt{\sin^2(\alpha_i) - 2(1 - n)} + \sqrt{\sin^2(\alpha_f) - 2(1 - n)}],$$

with $\alpha_i, \alpha_f$ the incidence and exit angles, $n$ the complex refraction index, and $K = 2\pi/\lambda$ the wave vector length in vacuum. Figures 3(b) and 3(c) show that, at an incidence angle of 0.22°, the refraction shift is exactly the same for 10 keV and 15 keV. This set of angles and energies is therefore convenient because it allows discriminating penetration depths between a few angstroms and several tens of nanometers keeping a constant refraction.

Diffraeted intensity was mapped around the (102) reflection. We used a pseudocubic unit cell with perpendicular reciprocal lattice vectors $H$, $K$, and $L$, with $L$ being perpendicular to the crystal surface. In Fig. 3(a) we show two perpendicular sections (tomographs) of the three-dimensional mesh. Because of the ferroelastic twinning, two peaks corresponding to the (102) and (−102) reflections are observed. On top of this, there is broadening of the maxima due to mosaicity [27]. We have therefore tracked the modulus of the momentum transfer $Q = \sqrt{H^2 + K^2 + L^2}$, in analogy with powder diffraction, and used its changes to identify structural distortions.

Figures 3(d) and 3(e) show the radial distribution of intensities integrated for each shell of constant $Q$ (i.e., the histogram of the moduli of $Q$) for the three-dimensional $HKL$ maps. Fig. 3(d) shows data at identical incidence angle (0.22°) but different photon energies (10 and 15 keV); in 3(e) the photon energy is constant (15 keV) and the incidence angle is varied (0.1° and 5°). For each panel, the lighter (red) curve denotes the surface of the crystal, while the bulk is in the dark (blue) curve. We have normalized the intensities to allow a comparison between light (red) and dark (blue) curves, and corrected the peaks for refraction shift [26]. In all cases the data show an unambiguous difference between surface and bulk. Such a difference cannot be explained by refraction as the shift of the histograms was identical for all data in 3(d) and, moreover, the corrections in 3(e) lead to the same results. The conclusion is that there is an expansion of the (102) interplanar distance located at the topmost nanometers of the crystal. Since the skin must be in-plane coherent with the bulk, the peak shift translates into an out-of-plane lattice parameter elongation of 0.7% at the surface.

The temperature dependence of the out-of-plane lattice parameter has been measured and is shown in Fig. 4 for the bulk and for the surface ($\alpha_i = 0.1^\circ$, penetration depth of 1 nm). The difference between the two curves is striking: whereas the bulk shows a featureless linear thermal expansion, the skin shows a marked anomaly, with negative thermal expansion setting in around 260 °C, followed by a sharp expansion of almost 0.2% at 280 °C. This temperature agrees with $T^*$ from impedance analysis. Its detection only at the subcritical angle and absence from supercritical incidence confirms that the phase transition is confined within a skin layer that is only a few unit cells thick; though its exact thickness cannot be measured, the complete absence from x-ray measurements with a penetration depth of 50 nm places an upper bound and suggests an order of magnitude in the region of 10 nm or less.

Surface-sensitive probes such as backscattering Raman [28] have shown an onset of phonon softening at $T^* = 270$ °C and culminating at $T_N$. It is not clear that it is the same transition, but the fact that $T^*$ is the same is unlikely to be a coincidence. Skin effects may also account for other

FIG. 3 (color online). (a) Reciprocal space tomograph around the (102) reflection. Calculated penetration depth (b) and refractive shift (c) as a function of incidence angle for two different beam energies, showing that for an angle of 0.22°, changing the energy allows tuning the penetration depth without changing the refractive shift. The Q-histogram analysis of the HKL measured by changing the photon energy (d) and the incidence angle (e), both of which confirm that the lattice parameter is expanded at the surface.
also note that the (100) planes of films are comparable to the thickness of the skin. Likewise, in thinning of the Bi sublattice for a radius smaller than 9 nm [35], structurally different from large grained ones, with a melt-to-volume ratio. Fine grained ceramics, for example, are charge depletion. be considered, and may in fact contribute to the observed as the surface relaxation energy of perovskites [33,34]. We confined transition would be facilitated by these, because the small energy barrier between them is of the same order as the surface relaxation energy of perovskites [33,34]. We also note that the (100) planes of BiFeO₃ are not neutral, so the electrostatic cost of the surface termination should also be considered, and may in fact contribute to the observed charge depletion.

The skin will impact on samples with a large surface to volume ratio. Fine grained ceramics, for example, are structurally different from large grained ones, with a melting of the Bi sublattice for a radius smaller than 9 nm [35], comparable to the thickness of the skin. Likewise, in thin films a T' anomaly has been reported [36]. The distinct properties of the surface are also likely to affect the performance of interfacial coupling devices such as those based on exchange bias [11]. Given its repercussions, then, the surface of BiFeO₃ deserves close scrutiny from the multiferroic community.

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FIG. 4 (color online). Comparison between the reciprocal thermal expansion at the skin (empty symbols) and inside the crystal (solid symbols) evidencing the local phase transition at T' in the surface of BiFeO₃.

unexplained anomalies of BiFeO₃ [3]. In particular, singularities at 200 K and 140 K have been observed by surface-sensitive probes [29,30] but not by bulk-sensitive ones such as transmission neutron diffraction and magnetometry [31], which suggests surface effects. As for the origin of the skin transition, the calculations of Dieguez et al. [32] predict a plethora of metastable phases within just 100 meV of the ground state of bulk BiFeO₃. A surface-confined transition would be facilitated by these, because the small energy barrier between them is of the same order as the surface relaxation energy of perovskites [33,34]. We also note that the (100) planes of BiFeO₃ are not neutral, so the electrostatic cost of the surface termination should also be considered, and may in fact contribute to the observed charge depletion.

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[27] From the angular broadening of the diffraction maxima we can estimate also the mean misorientation of the mosaic blocks: ∆ω = (1.3 ± 0.3) deg.