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ABSTRACT
Functional oxide superlattices offer new and exciting possibilities for the exploration of emergent properties at the nanoscale. While the behavior of \( \text{La}_{1-x}\text{Sr}_x\text{FeO}_3 \) films has been extensively investigated at low temperatures, few studies have been carried out at high temperatures, particularly for \( \text{LaFeO}_3/\text{SrFeO}_3 \) superlattice systems. Here, we investigate the transport behavior and optical properties of \( (\text{LaFeO}_3)_n/(\text{SrFeO}_3)_1 \) superlattices at 373 K and above. Using optical spectroscopy, we observe a low energy excitation at \( \sim 1 \text{ eV} \), attributable to charge transfer between the \( \text{O} 2p \) and \( \text{Fe} 3d \) states of the \( \delta \)-doped single \( \text{SrFeO}_3 \) layer. From in-plane conductivity measurements on the superlattices, we determine activation energies that are much lower than those of alloyed samples and vary with the total number of \( \text{SrFeO}_3 \) layers. This suggests that polaronic transport is confined near the \( \text{SrFeO}_3 \) regions, permitting mobilities significantly enhanced over those in alloyed thin films.

La\(_{1-x}\)Sr\(_x\)FeO\(_3\) (LSFO) is a strongly correlated oxide known to display a variety of unique material properties. While it exhibits many interesting electronic and magnetic transitions near or below room temperature,\(^{11}\) at higher temperatures its mixed ionic-electronic properties make it useful for sensors, catalysts, solid oxide fuel cells, and a host of other energy-related applications.\(^{4-11}\) Researchers can now directly examine the fundamental mechanisms governing transport at these elevated temperatures, utilizing model systems and state-of-the-art characterization probes to help distinguish between the effects of microstructure and intrinsic/extrinsic contributors to electrical conduction.\(^{12-15}\) Here we exploit advances in oxide thin film synthesis to investigate the properties of modulation doped (001)-oriented epitaxial heterostructures with SrO layers periodically inserted into the LaFeO\(_3\) matrix. Such atomically controlled, layered structures allow the study of cation ordering effects on electronic behavior,\(^{16-19}\) a topic of increasing interest in materials design\(^{20,21}\) and the one that may lead to the development of novel materials for energy.\(^{22,23}\) The substitution of Sr\(^{2+}\) for La\(^{3+}\) in the charge-transfer insulator LaFeO\(_3\) leads to increased p-type conductivity up to \( x = 0.5 \). Electronic transport is known to occur by small polaron conduction,\(^{24}\) involving thermally activated hops of hole polarons between
states of primarily oxygen character, as the Fe ions have been found to remain in the $3d^9$ configuration up to $x \approx 0.5$. At the limit of $x = 1$, perovskite SrFeO$_x$ is a well-known metal, and the conductivity of LSFO increases with higher Sr concentration. Previous transport studies, however, did not address the effect of geometric confinement in the LSFO films.

Here, we focus on $p$-type electronic transport in (001)-oriented (LaFeO)$_n$/(SrFeO)$_1$ or (LFO)$_n$/(SFO) superlattices at elevated temperatures (373–673 K) and high oxygen pressures (1300 Pa), where a supercell consists of $n$ consecutive layers of LaFeO$_3$ and a single layer of SrFeO$_3$. The superlattices were grown by ozone-assisted molecular beam epitaxy (MBE) on TiO$_2$-terminated SrTiO$_3$ (001) substrates. Throughout deposition, the ozone pressure in the chamber was fixed at $2.7 \times 10^{-6}$ Pa and the sample temperature kept at 923 K. Fluxes from the La, Sr, and Fe effusion cells were calibrated prior to growth. For deposition of the artificially ordered layers, we employed the sequential deposition method, alternately opening and closing each cation species for a time calibrated to result in the deposition of one monolayer. In situ reflection high-energy electron diffraction (RHEED) was used to monitor the growth process, and we observed pronounced intensity oscillations of the specular reflection throughout deposition. Superlattices with $n = 1, 2, 4, 7, 16$ were grown, always starting and terminating with a LaFeO$_3$ layer, leading to approximate Sr concentrations of $x = 0.50, 0.33, 0.20, 0.125,$ and 0.059, respectively. A pure LaFeO$_3$ film and a random alloy of La$_{0.875}$Sr$_{0.125}$FeO$_3$ grown by codeposition of La and Sr were used for comparison.

The structure of the superlattices was studied with x-ray reflectivity (XRR) using Cu $K_α$ radiation (Bruker D8 Discover), atomic force microscopy (AFM, Asylum Research MFP-3D), and cross-sectional scanning transmission electron microscopy (STEM) using a NION UltraSTEM100 operated at 100 kV. Since LSFO is a known mixed conductor, we performed both in-plane direct and alternating current (DC and AC) electrical measurements using a Keithley 2400 Sourcemeter and a CH Instruments 760D potentiostat, respectively. The electrical measurements were conducted in an environmental probe station capable of a wide variety of temperatures (300–1200 K) and gas environments. Optical properties of superlattices were measured by spectroscopic ellipsometry (VASE Ellipsometer, J. A. Woollam Co., Inc.) at room temperature. Since the optical wavelengths were much longer than the superlattice periodicities, each sample was approximated as a single effective medium for analysis.

A representative AFM image of the (LFO)$_n$/(SFO) surface after growth is shown in Fig. 1(a). As seen, the ~0.4 nm step height from the SrTiO$_3$ substrate was preserved during the growth process. This, along with the observed RHEED oscillations during growth, indicates that the superlattices were grown in a layer-by-layer fashion, resulting in atomically smooth surfaces and little to no excess material. The XRR data shown in Fig. 1(b) indicate that the superlattice structures were maintained after growth at high temperature, with interfacial roughnesses ranging from 0.1 to 0.3 nm, replicating the surface roughness of the substrate. Evidence for sharp interfaces can also be seen in the cross-sectional high-angle annular dark-field (HAADF) STEM image for one of the superlattice systems, (LFO)$_2$/(SFO), as shown in Fig. 1(c). Long-range periodicity is observed along the growth direction, consisting of two bright layers alternating with one dark layer, due to the higher atomic number of La compared with Sr. Chemical mapping with electron energy-loss spectroscopy (EELS) [Fig. 1(d)] shows that the atom distribution in the superlattice follows the intended ordering sequence, with little to no La signal in the darker layers and uniform Fe distribution.

The average out-of-plane lattice constant, $c$, as measured from high x-ray angle diffraction (Fig. S1) decreases from 0.393(1) nm to

![FIG. 1. Structural results of the (LFO)$_n$/(SFO) superlattices. (a) Representative AFM image of the film surface [from the (LFO)$_n$/(SFO) superlattice]. (b) Low-angle X-ray reflectivity measurements from the (LFO)$_n$/(SFO) superlattices. The first order superlattice X-ray reflections are indicated by the asterisks. (c) Cross-sectional HAADF STEM image of the (LFO)$_2$/(SFO) superlattice film grown on TiO$_2$-terminated SrTiO$_3$ (001). As depicted, the first deposited layer is LaO. The alternating bright and dark layers stem from the different atomic numbers of La and Sr. (d) Chemical maps obtained from the EELS spectra for the area marked with a green rectangle in (c) showing Fe (green; integrated Fe L$_{2,3}$ edge intensity) and La (red; integrated La M$_{4,5}$) atomic column distributions. The composite image was obtained from the two maps after intensity normalization. Note that vertical distortions in the maps are artifacts due to specimen drift during the acquisition.](image-url)
0.388(1) nm as \( x \) increases from 0.059 to 0.5, respectively. These values are in good agreement with the pseudocubic lattice parameters of air-cooled, bulk LSFO,\(^{15,32,35–39}\) as well as fully-oxidized epitaxial thin films.\(^{15,33–35}\) The \( c/vs x \) behavior is as expected based on the small misfit strain with respect to the substrate, which ranges from \( -0.6\% \) to \( +0.5\% \) as \( x \) is varied from 0 to 0.5.

Figure 2(a) shows 300 K data for the real part of the optical conductivity spectra, \( \sigma(\omega) \). The transition at \( \sim 3.2 \text{ eV} \) for pure LaFeO\(_3\) stems from the electronic band gap. Using well-established procedures, \( \) we determine an optical bandgap of 2.68(2) eV for LaFeO\(_3\), which is in good agreement with the prior result of 2.64 eV from Scafetta et al.\(^{15,32}\) Similar to the results for alloyed LSFO films,\(^{15,32}\) Sr doping induces a red shift in this transition. We also observe the appearance of a transition at \( \sim 1.2 \text{ eV} \), attributed to states formed within the bandgap and from which we can extract a new, smaller bandgap as a function of Sr concentration [Fig. 2(a), inset].

With an increasing amount of Sr, we observe the following systematic changes in the electronic structure of the superlattices:

1. The appearance and development of absorption in the midgap region at \( \sim 1 \text{ eV} \); 2. increasing spectral weight (SW) at the \( \sim 3 \text{ eV} \) photon energy range; 3. decreasing spectral weight above 4 eV. The prominent amount of absorption at around 1 eV and 2–3 eV can be attributed to charge transfer transitions between O 2p and various Fe 3d orbital states (O 2p to Fe 3d \( t_{2g} \) and \( e_g \), respectively).\(^{15,35–39}\)

The nature of the \( \sim 1 \text{ eV} \) peak is the key to understanding transport properties of the ferrite system. To obtain further insights into this low energy excitation, we plot the bandgap of the absorption and its spectral weight \( \text{[SW = } \int \sigma(\omega) \ d\omega \text{]} \) as a function of Sr doping \( (x) \), as shown in Fig. 2(b). Due to the wavelength limitation of the spectroscopic ellipsometer (1, 700 nm), we also measured the reflectance spectra of the superlattices down to the mid-infrared range (0.07 eV) and confirmed the low-photon-energy tail based on the fitting of reflectance spectra, shown as dotted lines in the inset of Fig. 2(a). As seen, the SW of the \( \sim 1 \text{ eV} \) absorption peak almost increases linearly with \( x \), possibly due to the increased hole concentration with Sr doping within the superlattice. Therefore, the peak can be attributed to the newly introduced electronic states from the SrFeO\(_3\) layers in the superlattice.\(^{15} \) The optical bandgap determined at lower absorption energies also shows a systematic trend. Interestingly, these values are consistent with the activation energies obtained from the previous high temperature transport measurements on alloyed LSFO samples.

To address the effect of geometric confinement within the (LFO)\(_{x}\)/[SFO] superlattices on the transport properties, electrical measurements were performed using platinum interdigitated electrodes with 5 \( \mu \text{m} \) spacing. AC measurements performed between \( 10^{-1} \text{ Hz} \) and \( 10^{7} \text{ Hz} \) showed no evidence of frequency dispersion up to 673 K resulting in conductance values nearly identical to those from the DC experiments, suggesting that ionic transport is not a major contributor to the electrical conductivity in this temperature range. Consequently, all of the values reported here were obtained from DC measurements conducted after an initial 60 min oxygen anneal at a fixed oxygen partial pressure of 1300 Pa, with a maximum temperature of 673 K to minimize the degree of interdiffusion.\(^{40}\) The samples were given at least 2 h to equilibrate after changing temperatures and reach time-independent conductance values. Room temperature XRR measurements taken from a sample before and after annealing at both 473 K and 673 K are shown in Fig. S2 of the supplementary material. As seen, anneals at these temperatures had a negligible effect on the cation-ordered structure.

For the small, nonadiabatic polaron hopping mechanism, the conductivity is predicted to have the following temperature dependence:\(^{11,42}\)

\[
\sigma = \frac{\sigma_0}{T^{3/2}} e^{-E_a/k_B T}
\]

where \( \sigma_0 \) is a pre-exponential factor, \( E_a \) is the activation energy, and \( k_B \) is the Boltzmann constant. A plot of \( \sigma T^{3/2} \) vs 1000/T is shown in Fig. 3(a) for both the cation-ordered superlattices and the random alloy film. The conductivities follow the Arrhenius behavior predicted in Eq. (1), in accordance with the small polaron model, and increase with the Sr concentration, \( x \), as expected based on carrier doping. Interestingly, all of the superlattices exhibit higher
This shows that the change in conductance is mainly dependent on the number of SrFeO₃ layers and not the LFO:SFO ratio, which is markedly different from the behavior of alloyed LSFO samples. This discrepancy between the alloy samples and our superlattices can be understood if the polarons are geometrically confined to the SrFeO₃ layers. Note that both the formation energy of oxygen vacancies in SrFeO₃ (about 0.4 eV) and the energy for ionic conduction in bulk SrFeO₃ (about 0.5 eV) are larger than the activation energies observed here.

As seen in Fig. 3, the cation-ordered structures exhibit activation energies smaller than those for the random alloy samples, by roughly 0.2 eV (except at x = 0.5). Since the hole concentrations of the superlattice samples in Fig. 2(b) scale with the degree of Sr doping, it appears that the reduced activation energy of the superlattices originates from an enhancement in mobility related to cation ordering. The decrease in activation energy with x also indicates that hole transport is not perturbed by cation disorder and may be confined to the SrFeO₃ layers, as shown in Fig. 3(c). If increasing x implies a progression toward more metallic perovskite SrFeO₃ rather than more insulating brownmillerite SrFeO₂.₅, one can reasonably explain the decrease in bandgap and the increase in conductivity of the superlattices. In fact, the ~1 eV low energy absorption peak in Fig. 2(b) is a signature of the perovskite SrFeO₃ structure.

To confirm confined, in-plane polaronic hopping along the SrFeO₃ layers, we fabricated (LFO)ₓ/(SFO) and (LFO)ₓ/(SFO) superlattices, each with a total of seven SFO layers (63 and 35 unit cells thick, respectively). The conductance of (LFO)ₓ/(SFO) is nearly the same as that of the (LFO)ₓ/(SFO) superlattice, as shown in Fig. 3(d). The conductance therefore appears to be independent of both the total thickness and the number of LaFeO₃ unit cells. This shows that the change in conductance is mainly dependent on the number of SrFeO₃ layers and not the LFO:SFO ratio, which is markedly different from the behavior of alloyed LSFO samples. This discrepancy between the alloy samples and our superlattices can be understood if the polarons are geometrically confined to the SrFeO₃ layers. Note that both the formation energy of oxygen vacancies in SrFeO₃ (about 0.4 eV) and the energy for ionic conduction in bulk SrFeO₃ (about 0.5 eV) are larger than the activation energies observed here.

In summary, we report the observation of confined transport behavior in (LFO)ₓ/(SFO) superlattices at high temperatures. The results of optical spectroscopy indicate enhanced absorption at around 1 eV, originating from charge transfer between the O 2p and Fe 3d 12g states. Changes in the corresponding bandgap are consistent with changes in the activation energies of the (LFO)ₓ/(SFO) superlattices as a function of the Sr concentration. The lower activation energies of (LFO)ₓ/(SFO) superlattices as compared to those of alloyed LSFO samples may be understood in terms of confined polaronic transport. While spectroscopic measurements of the high temperature electronic structure would be worthwhile, this initial study provides important insights into unique properties of digitally synthesized materials.

See supplementary material for high angle, θ − 2θ x-ray diffraction scans (using Cu Ka radiation) from all of the superlattices and low angle reflectivity scans from one of the superlattices, illustrating the effect of annealing on superlattice structures.

Work by S. K. Kim, Y.-M. Kim, Y. Dong, C. M. Folkman, D. W. Jeong, A. Y. Borisevich, J. A. Eastman, A. Bhattacharya, and D. D. Fong was supported by the U.S. Department of Energy (DOE), Office of Science, Basic Energy Sciences (BES), Materials Sciences and Engineering Division. Work by S. H. Chang was supported by the Chung-Ang University Research Grants in 2017.
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