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Perspectives on Oxide Heterostructures – The Curious Case of γ-Al₂O₃/SrTiO₃

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Abstract

The heterostructure formed by depositing nanoscale thin films of spinel γ -Al₂O₃ on perovskite SrTiO₃ exhibits a range of exciting properties including room temperature epitaxial growth, high electron mobility, strain-tunable magnetic order, and a symmetry-related reordering of the conduction bands. In comparison to the benchmark LaAlO₃/SrTiO₃ heterostructure, the γ -Al₂O₃/SrTiO₃ heterostructure has been more sparsely investigated, which leaves plenty of room for scientific and technological discoveries. In this perspective article, I describe the key findings of the γ -Al₂O₃/SrTiO₃ heterostructure and propose five directions for future research: 1) an exploration of novel phenomena emerging when relaxing the conventional epitaxial constraint of matching crystal structures across the interface, 2) a dynamic switching of a strong polarization through nanoscale electromigration of aluminum vacancies, 3) autonomous and forced enhancement of the electron mobility via oxygen vacancy diffusion, 4) writing and erasing of magnetic and conducting nanolines using ferroelastic domain walls, and 5) a multiferroic state formed by combining ferroelectricity, ferromagnetism, and ferroelasticity. The proposed research directions may shed light on both fundamental aspects of the heterostructure and pave the way for applications within green energy devices and nanoelectronics.



The strong market pull for high-speed and power-saving electronics has fueled the tremendous development in realizing faster, smaller, and more energy-efficient nanoelectronic devices. Two routes are used to satisfy this market pull: (i) Improving existing devices or (ii) designing devices with new functionalities. Appealing functionalities may be realized by using materials beyond the semiconducting materials that currently constitute the backbone of state-of-the-art electronic devices. A key example is the emergence of neuromorphic computation where non-volatile changes in the resistance observed on the nanoscale in, e.g., memristive oxides can be used to form a computational framework inspired by the human brain, which features both improved energy efficiency, learning capabilities and suitability for deep learning tasks [1]. A material system particularly rich in novel functionalities is the oxide heterostructure formed by depositing a nanoscale thin film of LaAlO₃ epitaxially on SrTiO₃ [2]. Despite both oxides being considered non-magnetic and insulating, conductivity and magnetism emerge at the interface along with numerous other functionalities [3]. In the wake of the large interest in the $LaAlO_3/SrTiO_3$ heterostructure, a new artificial system was obtained by replacing perovskite LaAlO₃ with spinel γ -Al₂O₃ [4]. The γ -Al₂O₃/SrTiO₃ heterostructure has shown to be an astonishing material platform in many ways, yet, the heterostructure has been much less studied compared to the benchmark LaAlO₃/SrTiO₃, which leaves plenty of room for future research adventures. Nonetheless, the heterostructure has already proven to possess several exciting properties not observed in the LaAlO₃/SrTiO₃ counterpart or in other SrTiO₃based heterostructures. In this perspective article, I will first highlight some of the key findings that make the y-Al₂O₃/SrTiO₃ heterostructure an exciting material platform to study, and secondly discuss several key opportunities for future research within nanoscale tailoring of this heterostructure.

Key findings:

Synthesis and electrical conductivity of γ -Al₂O₃/SrTiO₃: The first remarkable characteristic of the γ -Al₂O₃/SrTiO₃ heterostructure is the possibility to grow the spinel γ -Al₂O₃ epitaxially on cubic perovskite SrTiO₃ using e.g. pulsed laser deposition and atomic layer deposition [5–7]. Typically, thin films of γ -Al₂O₃ with a thickness below 10 nm are deposited [5– 7]. Despite the materials having different crystal structures, the oxygen sub-lattices are highly compatible, which enables the non-isomorphic epitaxial growth, even when depositing γ -Al₂O₃ at room temperature with pulsed laser deposition [4]. The matching oxygen sublattices are displayed in Figure 1a, which depicts SrTiO₃ capped with a generic spinel structure AB₂O₄ where aluminum occupies both A and B sites in the case of γ - Al_2O_3 . The symmetry breaking at the spinel/perovskite interface is central to understanding the interface properties. For instance, when electrically insulating γ -Al₂O₃ is deposited on the insulating SrTiO₃, the resulting y-Al₂O₃/SrTiO₃ heterostructure becomes conducting, and the itinerant electrons are confined at the interface by virtue of the broken symmetry [4,8]. The itinerant electrons confined to the interface live in a band structure that differs significantly from other SrTiO3-based heterostructures due to the broken lattice symmetry [9,10]. From annealing in oxygen [11], high-temperature equilibrium conductance experiments [12], and numerical modeling [13] it is deduced that the itinerant electrons originate from oxygen vacancy donors formed in SrTiO₃ during the deposition. Hence, the origin of the itinerant electrons shares similarities to other SrTiO₃-based systems where amorphous or metal top layers such as Al are deposited on SrTiO₃ [14–16]. This origin raises an interesting paradox: The high-temperature equilibrium conductance measurements attribute the interface conductivity in LaAlO₃/SrTiO₃ to the LaAlO₃ polarity, but this is not the case for γ -Al₂O₃/SrTiO₃ although γ -Al₂O₃ has been considered polar [12]. Revisiting the atomic structure of γ -Al₂O₃, it was deduced that γ -Al₂O₃ can be non-polar for particular distributions of the aluminum vacancies that are intrinsically present in γ -Al₂O₃ [17].

Tunable electronic properties of γ-Al₂O₃/SrTiO₃:

Using the understanding of how the conductivity arises in γ -Al₂O₃/SrTiO₃, the carrier density can be tuned by controlling the oxygen vacancy concentration by varying the γ -Al₂O₃ deposition parameter [18] or performing post-annealing in oxygen at temperatures below 300 °C [11]. A similar tunability through defect engineering was also found in the heterostructure



Figure 1 – Properties of the γ -Al₂O₃/SrTiO₃ **heterostructure:** (a) Schematics of the crystal structure of the spinel/perovskite heterostructure where both A and B sites are occupied by aluminum with $2^2/_3$ aluminum vacancies pr. unit cell in the case of γ -Al₂O₃. Despite the different crystal structures, γ -Al₂O₃ can grow epitaxially on SrTiO₃ due to the compatible oxygen sublattice as depicted on the right-hand side. (b) Electron mobility (μ) as a function of temperature (T) including a fit using three different scattering contributions. (c) Scanning SQUID maps of the stray magnetic field above the γ -Al₂O₃/SrTiO₃ surface arising from magnetic order in the heterostructure as the probe is scanned in non-contact mode (lower 4 panels) and in contact mode with increasing pressure (upper 3 panels). The figures 1b and 1c are adapted from Ref. [21] and [21] with permission ₂ from American Physical Society and Springer Nature, respectively.



Figure 2 – Self-enhancing electron mobility: (a) Simulated depth-distribution of oxygen vacancy donors and background impurities (top) as well as itinerant electrons (middle) in a γ -Al₂O₃/SrTiO₃ heterostructure stored at room temperature for different periods after deposition. Here, x denotes the distance from the γ -Al₂O₃/SrTiO₃ interface. The electrostatic potential (bottom) associated with the oxygen vacancy distribution, which governs the distribution of itinerant electrons. (b) Simulated and measured electron mobility at 2 K ($\mu_{T=2K}$) for various storage times (t) after deposition of the y-Al₂O₃/SrTiO₃ heterostructure. The figure are adapted from Ref. [25] with permission from American Physical Society.

composed of amorphous LaAlO₃ deposited on SrTiO₃ where the electrons also originate from oxygen vacancies [11]. Electrons can also be induced in a non-volatile fashion using external stimuli on the macroscopic scale through conventional electrostatic gating [18], ionic liquid gating [19] or light exposure [18,20], and on the nanoscale using a biased atomic force microscopy tip [11]. The latter allows for a versatile patterning of circuits with nanoscale resolution [11].

Mechanically tunable magnetism:

Surprisingly, γ -Al₂O₃/SrTiO₃ was found to possess a mechanically tunable magnetic state [21]. The magnetic state manifests itself at T < 40 K both via the anomalous Hall effect and a magnetoresistance that deviates from Kohler's rule. Scanning SQUID was further used to detect a magnetic order extending hundreds of micrometers with modulations occurring along the tetragonal domain walls of SrTiO₃ (see Figure 1c). Applying force on the γ -Al₂O₃/SrTiO₃ surface using the scanning SQUID changes the magnetic order drastically, resulting in much stronger modulations (see Figure 1c). It was speculated that the applied force enhanced the magnetization by shifting the balance between the competition of a ferromagnetic order and a nonmagnetic order. The ferromagnetic order is likely to arise from localized magnetic moments on oxygen vacancies that exchange couple to the itinerant electrons. The non-magnetic order may originate from, e.g., a singlet electron pairing as observed in a non-superconducting state for other SrTiO₃-based heterostructures [22-24].

Electron mobility:

annealing at ~100 °C or during prolonged room temperature storage [29]. The finite mobility of oxygen vacancies - acting both as electron donors and scattering centers - was attributed to being the cause of this self-enhancing mobility [13]. Driftdiffusion calculations were performed to describe the oxygen vacancy movement (see Figure 2). Here, part of the oxygen remained fixed at the γ -Al₂O₃/SrTiO₃ interface due to the broken lattice symmetry, whereas other oxygen vacancies formed a diffusion front that moved gradually deeper into the bulk of SrTiO₃ due to electrostatic repulsion (Figure 2a). The simulations further show that the resulting electrostatic potential from the charged oxygen vacancies caused a significant fraction of the electrons to be distributed between the interface and the diffusion front, thereby forming a high-way for high-mobility conduction in between the regions of high oxygen donor concentrations. A resulting self-enhancing electron mobility was predicted, which matched the experimental data well (Figure 2b). The high carrier density of $\sim 5 \cdot 10^{14}$ cm⁻² is therefore predicted to be accompanied by a nanometric electron depth distribution after deposition to extending a micrometer or more into the SrTiO₃ substrate after prolonged aging for several months. This can be compared to experimental depth distributions extending 0.9 nm and 7.5 nm into SrTiO₃ as deduced from angle-dependent x-ray photoemission spectroscopy at room temperature and cryogenic infrared ellipsometry performed on samples without prolonged aging [4,8].

The mobility in γ -Al₂O₃/SrTiO₃ can be enhanced by gentle post-

Future perspectives:

At room temperature the electron mobility in γ -Al₂O₃/SrTiO₃ is generally found to be limited to 12 cm²/Vs by phonon scattering [25], but with a few reports stating values exceeding 30 cm²/Vs [26,27]. At 2 K the mobility exceeds 100,000 cm²/Vs (see Figure 1b) at a high carrier density of ~5·10¹⁴ cm⁻² [4,25], which differs fundamentally from bulk conducting SrTiO₃ and most other SrTiO₃-based heterostructures where high mobility is found at low carrier densities [28]. This mobility value at low temperatures is amongst the highest obtained for oxides [28]. In contrast to the prototypical perovskite/perovskite LaAlO₃/ SrTiO₃ interface, the spinel/perovskite γ -Al₂O₃/SrTiO₃ interface has been investigated more sparsely, which leaves plenty of room for exploring how the altered crystal structure across the interface may lead to unexpected phenomena. Below, I outline five key opportunities for future research inspired by the current knowledge of the γ -Al₂O₃/SrTiO₃ heterostructure:



Figure 4 – Polarity in the γ -Al₂O₃/SrTiO₃ heterostructure: (a) Schematics of the nominal polarity in LaAlO₃ deposited epitaxially on SrTiO₃ arising from polar ionic layers. (b-f) Schematics of the nominal polarity in γ -Al₂O₃ deposited epitaxially on SrTiO₃ if aluminum vacancies are (b) not present in the structure (charged structure), (c) present solely on octahedral sites, (d) present solely on tetrahedral sites, (e-f) distributed inhomogeneously to form overall non-polar states. (g) The electrostatic potential build-up due to the polarity as calculated by an electrostatic capacitor model. The figure is reproduced from Ref. [13] with permission from Elsevier.

Broken lattice symmetry:

Conventional epitaxy relies on a high degree of structural coherency across the interface. Matching the crystal structure and lattice constant enables the growth of high-quality thin films using the substrate as a growth template. Broken symmetry is typically introduced by altering the composition of the constituent materials while still maintaining a reasonable match of the crystal structure across the heterointerface. Whereas high-quality homo- and heterostructures can be formed using this route, it imposes strict limitations on the choice of materials that can be combined. The γ -Al₂O₃/SrTiO₃ heterostructure partially relaxes these limitations by introducing broken symmetry from changing both the composition, lattice constant and crystal structure across the interface while still maintaining a high-quality epitaxial growth (see Figure 1a). A key example of how the broken symmetry at the perovskite/spinel interface influences the functional properties is the altered band structure. In (001)-oriented LaAlO₃/SrTiO₃, the bare SrTiO₃ surface as well as SrTiO₃ capped with AI metal, the lowest-lying band is of d_{xy} character, however, the presence of positively charged aluminum arranged in a spinel structure induces an increase in the d_{xy} energy so it shifts up above the level of the d_{xz} and d_{yz} bands in γ -Al₂O₃/SrTiO₃ [9,10,30–32].

It remains, however, largely unclear how other properties are influenced by transitioning from perovskite/perovskite to spinel/perovskite symmetries. In particular, the interesting electronic and magnetic properties originate from an interfacenear region, making them susceptible to the lattice symmetry across the interface. The broken symmetry also influences the structural relaxation at the interface, which may form an interfacial polarization, emergent pyroelectricity and oxygen vacancy stabilization. In addition, it is of high interest to explore how the limitations of epitaxy can be further overcome not only by the non-isomorph epitaxial growth of different crystal structures but also by exploiting the recent progress within freestanding oxide thin films [33,34]. Using the latter approach, crystalline oxide thin films can be detached from the growth substrate and reunited with other films or substrates to form new interfaces not obtainable by conventional epitaxy (see Figure 4) [35]. Here, the high-quality epitaxial growth of γ -Al₂O₃ on perovskites may enable y-Al₂O₃ to be used either as a sacrificial layer etchable with, e.g, NaOH [36] (blue layer in Figure 4) or as the active layer to be released from the substrate (yellow or red layer). This may enable high-quality spinel crystal structures with potentially sub-unit cell thicknesses to be released and subsequently recombined with other spinel or perovskite structures. The potential sub-unit cell release and recombination of 0.2 nm membranes (1/4 of a unit cell) allow for a larger surface-to-volume ratio as well as smaller building blocks



Figure 3 – Stacking and twisting of freestanding oxides: An epitaxial heterostructure composed of a substrate (dark blue), sacrificial layer (light blue) and active layer (red and yellow) subjected to etching of the sacrificial layer can be used to detach the top layers from the growth substrate. This enables stacking of layers with different material classes (perovskites, spinels etc.) with a flexibility not allowed by conventional epitaxy. The figure is adapted from Ref. [29] with permission from Wiley.

for the construction of stacked heterostructures. Of particular interest is the formation of conducting interfaces in heterostructures formed by stacking individual membranes. As the mechanism for forming conductivity in epitaxially grown γ -Al₂O₃/SrTiO₃ relies on the movement of oxygen across the interface, kinetic limitations may prevent conducting interfaces to be formed by room temperature stacking of non-conducting γ -Al₂O₃ and SrTiO₃ membranes. In contrast, conductivity may be formed by high-temperature reduction of SrTiO₃, which forms oxygen vacancy donors that can be kinetically trapped inside SrTiO₃ by a protective layer of γ -Al₂O₃. The new γ -Al₂O₃/SrTiO₃ interface may also serve to localize and thermodynamically stabilize oxygen vacancies at the interface.

Thickness-dependent dynamic polarity switching:

Beyond the potential polarity formed at the interface in y- $Al_2O_3/SrTiO_3$, polarity may also form within the y-Al_2O_3 layer. To obtain charge neutrality in y-Al₂O₃, each spinel unit cell intrinsically contains $2^2/_3$ aluminum vacancies (Al_{211/2}O₃₂) located on lattice sites with tetragonal and/or octahedral symmetries (see Figure 1a). Depending on the film thickness of γ -Al₂O₃ as well as the distribution of aluminum vacancies in y-Al₂O₃, the film may either be highly polar or overall non-polar (Figure 3). By distributing the aluminum vacancies freely on tetragonal or octahedral sites, it was shown that no distributions of aluminum vacancies exist that give a non-polar film for thicknesses below one unit cell (0.8 nm corresponding to 8 sublayers) [4]. This should be compared to RHEED-monitored growth with sub-unit cell control of the thickness as 4 RHEED oscillations correspond to a single unit cell [4]. For a γ -Al₂O₃ thickness of 1 unit cell, the aluminum vacancies can either distribute in ways that produce a strong polarity (Figure 3c-d) or no overall polarity (Figure 3e-f) with a large predicted impact on the associated electrostatic potential build-up in the $\gamma\text{-}Al_2O_3$ layer (Figure 3g). A strong polarization of 75 μ C/cm² is expected for aluminum vacancies distributed according to Figure 3c, which exceeds that found in single crystals of the strongly polar material BaTiO₃ (~25 μ C/cm² [37]) as well as the spontaneous polarization at the surface of single-crystalline SrTiO₃ substrates (~5 μ C/cm² [38]). A yet untested hypothesis exists suggesting that the polarity in γ -Al₂O₃ might be tunable by electric fields due to the electromigration of aluminum ions [17]. If disregarding the internal electrostatic build-up associated with the various cation configurations, the movement of aluminum vacancies is expected to have an activation barrier on the order of 0.6 eV for moving aluminum vacancies between lattice sites within y-Al₂O₃ [39]. This is on the same order of magnitude as the barrier for oxygen vacancy diffusion in SrTiO₃ where electromigration is commonly observed [40,41]. This would offer switchable functionalities with similarities to conventional ferroelectrics, but with the switching caused by large-scale defect migration rather than small polar lattice distortions within a unit cell. In addition, the polarity may also be sensitive to applied stress and temperature, resulting in piezoelectric and pyroelectric properties.



Figure 5 – Ferroelastic domain wall memristor: Schematics of a ferroelastic domain wall memristor where external stimuli such as stress or electric fields may be used to control the landscape of ferroelastic domain walls and hence the resulting electrical resistivity in a non-volatile fashion.

Self-enhancing electron mobility:

Oxygen vacancies in γ-Al₂O₃/SrTiO₃ serve key roles as electron dopants, scattering sites limiting the low-temperature mobility, and a likely source for magnetism. The diffusivity of oxygen vacancies in SrTiO₃ with an energy barrier on the order of 0.6 eV makes it possible for these key defects to move in absence of high temperatures. This stands in stark contrast to conventional electronic systems where donors are typically static, and it, therefore, opens up opportunities to tune functional properties through post-deposition treatment [11,42]. As mentioned above, it was suggested that the autonomous rearrangement of oxygen vacancies at room temperature resulted in a depth distribution, which favors high-mobility carriers [13]. One may expect that the diffusion can be changed from autonomous to controlled on-demand by adding an electric field from electrostatic gates to control the electromigration of oxygen vacancies. In this way, one can envision a dynamic control of both the location of electron donors as well as the resulting confinement potential and electron distribution in a non-volatile fashion. Back-gates are expected to primarily modulate the depth distribution of donors and vacancies whereas side gates or local gates from probes may control the lateral distribution as well. This is expected to have a strong impact on the electron mobility as well as other functional properties such as the magnetism observed in y-Al₂O₃/SrTiO₃. The y-Al₂O₃/SrTiO₃ interface here is expected to play the particular role of partially trapping the oxygen vacancies near the broken interface, which may be enhanced or diminished by replacing the γ -Al₂O₃ layer with other top films such as amorphous top layers [14], spinel layers with cation substitutions [43] or metal layers [16] to form a set of oxygen-deficient SrTiO₃-based heterostructures.

Writing functionalities with structural domain walls:

The magnetism and current distribution in γ -Al₂O₃/SrTiO₃ have stripy modulations at cryogenic temperatures aligned along the [100], [010], and [110] crystal directions [21,25]. The stripy features are attributed to the ferroelastic domains arising when SrTiO₃ transitions from cubic to tetragonal around 105 K. The transition forms three different domains where the elongation of the tetragonal unit cell is along the x-, y-, and z-direction, respectively. This leads to a distribution of domains separated by domain walls where the spatial differences in the lattice here form modulations of the electronic and magnetic properties [21,25]. The size, number, and spatial distribution of domains and domain walls can be controlled using external stimuli such as mechanical stress and electric fields on both the macroscopic and microscopic scales [21,44]. This opens up for dynamically writing and erasing nanoscale lines with modified magnetic and conducting properties. As these lines are coupled to the underlying lattice structure, the changes are expected to be nonvolatile, which may open up for designing spintronic devices and ferroelastic memristors (see Figure 5).

Multiferroicity:

As described above, SrTiO₃ turns ferroelastic below approximately 105 K. In addition, SrTiO₃ is a quantum paraelectric at cryogenic temperatures, which is characterized by the ferroelectric polarity of the unit cell being suppressed by quantum tunneling between two polar states, hence on average producing a non-polar state. As SrTiO3 is on the verge of becoming a ferroelectric material, ferroelectricity can be induced by lattice modification such as applying stress [45], substituting Sr for Ca [46], or replacing ¹⁶O with ¹⁸O isotopes [47]. In addition, a ferromagnetic state appears to be formed as well in the y-Al₂O₃/SrTiO₃ heterostructure [21]. This suggests that a multiferroic state awaits to be discovered in y-Al₂O₃/SrTiO₃, which combines both ferroelastic, ferroelectric, and ferromagnetic properties in a single material system. Of particular interest would be to study to what extent the ferroelastic, ferroelectric, and ferromagnetic degrees of freedom couple together to form, e.g., magnetoelectric effects. Although this multiferroic state has yet to be observed in y-Al₂O₃/SrTiO₃, several studies point to an intimate coupling between the spin, lattice and charge degrees of freedom as illustrated in Figure 6. Here, the magnetization is dynamically coupled to the lattice through its strain-tunable property as well as through its modulation on ferroelastic domain walls [21]. Similarly, the charge flow is also modulated by the ferroelastic domain walls [25]. Through an electron-phonon coupling, the charge further distorts the lattice to form a polaron with an enhanced effective mass as the charge flows in the structure [10]. Lastly, the spin and charge degrees of freedom are also coupled through the anomalous Hall effect and exchange coupling of spins with the itinerant charge carriers [21]. Whereas the ways these degrees of freedom couple together are not exclusive to γ -Al₂O₃/SrTiO₃ but rather occur in other SrTiO₃based material systems as well [21,44,48-53], all coupled degrees of freedom have been observed in a single y-Al₂O₃/SrTiO₃ sample. In addition, the coupling between charge and spin degrees of freedom through the anomalous Hall effect manifests itself in γ -Al₂O₃/SrTiO₃ as a strongly non-linear Hall resistance below a threshold temperature, which coincides with

the emergence of a stripy modulation of the magnetic signal picked up by scanning magnetometry [21].

Conclusion:

The γ -Al₂O₃/SrTiO₃ heterostructure has already proven to be an exciting and multifunctional material system, which is less explored compared to the archetypical LaAlO₃/SrTiO₃ interface. Nonetheless, replacing the LaAlO₃ film with y-Al₂O₃ results in, e.g., a severely broken lattice symmetry, reordering of the conduction band, and high electron mobility. I here propose to further expand the range of properties by 1) investigation of heterointerfaces not bound by conventional epitaxial constraints, 2) dynamic and thickness-dependent polarity switching, 3) electromigrative control of electron donors, 4) writing and erasing of conducting and magnetic nanolines, and 5) realization of co-existing ferroelasticity, ferromagnetism, and ferroelectricity in γ -Al₂O₃/SrTiO₃. This renders the material property an interesting material platform to explore for scientific purposes as well as for applications utilizing ferroelectricity, high-mobility electronics, and multiferroicity.

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Figure 6 – Coupling of spin, lattice and charge degrees of freedom: The wide range of properties observed in γ -Al₂O₃/SrTiO₃ arises due to the coupling of spin, lattice and charge degrees of freedom. The figure is adapted from Ref. [17] with permission from Springer Nature.

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