



Surface Science Letters

Stoichiometry-driven switching between surface reconstructions on SrTiO₃(001)[☆]Stefan Gerhold^{*}, Zhiming Wang, Michael Schmid, Ulrike Diebold¹

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ABSTRACT

Controlling the surface structure on the atomic scale is a major difficulty for most transition metal oxides; this is especially true for the ternary perovskites. The influence of surface stoichiometry on the atomic structure of the SrTiO₃(001) surface was examined with scanning tunneling microscopy, low-energy electron diffraction, low-energy He⁺ ion scattering (LEIS), and X-ray photoelectron spectroscopy (XPS). Vapor deposition of 0.8 monolayer (ML) strontium and 0.3 ML titanium, with subsequent annealing to 850 °C in 4×10^{-6} mbar O₂, reversibly switches the surface between $c(4 \times 2)$ and (2×2) reconstructions, respectively. The combination of LEIS and XPS shows a different stoichiometry that is confined to the top layer. Geometric models for these reconstructions need to take into account these different surface compositions.

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1. Introduction

Strontium titanate (SrTiO₃, STO) belongs to the class of cubic perovskite oxides and it is well known for its useful bulk and surface properties, e.g., a high dielectric constant at low temperatures [1], photocatalytic water-splitting [2], lattice matching for growth of high-T_c superconductors [3], and the formation of two-dimensional electron gases at its surfaces [4,5], and at interfaces with other perovskites [6]. For most applications, a control of the surface structure at the atomic scale is of central importance.

The most important surface, STO(001), exhibits a large variety of reconstructions. Typical preparation procedures consist of sputtering with Ar⁺ ions of different energies (~1 keV) and fluences, and annealing to high temperatures (~1000 °C) in various oxygen pressures (atmospheric to $<10^{-10}$ mbar O₂). Often the outcome also depends on the sample preparation history. Different groups report numerous STO(001) surface reconstructions, including (1×1) [7–13], (2×1) [9–12,14,15], (2×2) [7,9,10,16], $c(4 \times 2)$ [12,14,16,17], $c(4 \times 4)$ [12,16], (4×4) [16], $c(6 \times 2)$ [11,15,17–19], $(\sqrt{5} \times \sqrt{5}) - R26.6^\circ$ [13,16,20,21], and $(\sqrt{13} \times \sqrt{13}) - R33.7^\circ$ [11,16]. Sometimes different reconstructions are reported for very similar preparation conditions.

At the STO(110) surface, a study by Wang et al. [22] showed that a series of reconstructions can be created and converted into each other

reversibly by controlling the surface stoichiometry, i.e., by depositing Sr or Ti onto the surface and annealing in 10^{-6} mbar O₂. This was recently also demonstrated at the STO(111) surface by Feng et al. [23]. Following this approach, the present Letter reports on the response of STO(001) surface reconstructions to a similar procedure. Using low-energy electron diffraction (LEED), scanning tunneling microscopy (STM), X-ray photoelectron spectroscopy (XPS), and low-energy He⁺ ion scattering (LEIS) it is concluded that the transition between the (2×2) and the $c(4 \times 2)$ STO(001) surface reconstructions is driven by surface stoichiometry. The implications for models proposed in the literature are discussed.

2. Experimental methods

Nb-doped (0.5 wt.%) SrTiO₃(001) single crystals were purchased from MaTeck Company, Germany. After ultrasonic cleaning in acetone, the samples were introduced into a two-chamber UHV system. One chamber is equipped with evaporators (Sr, Ti), a sputter gun, a home-built quartz crystal microbalance (QCM) and a leak valve for admitting O₂ into the chamber. Its base pressure was below 10^{-9} mbar. The second chamber, with a base pressure below 10^{-10} mbar, was used for analysis with LEED (SpectraLEED, Omicron), STM (Aarhus 150, SPECS), XPS (non-monochromatized dual-anode), and a scanning ion gun for LEIS. XPS spectra were acquired using Mg K α radiation. For LEIS, 1 keV He⁺ ions were used. Backscattered ions (scattering angle $\vartheta = 137^\circ$) and photoelectrons (emission normal to sample surface) were detected with a SPECS Phoibos 100 hemispherical analyzer with 5-channel detection. The XPS peaks were fitted after subtracting a Shirley background and the ISS peak intensity was summed up over the peak area after subtraction of a linear background.

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After loading the samples into the system, they were sputtered with 1 keV Ar^+ ions ($\sim 8 \times 10^{13} \text{ Ar}^+/\text{cm}^2\text{s}$) for typically 10 min and annealed at $\sim 850^\circ\text{C}$ in 10^{-6} mbar O_2 for 40 min. Titanium was evaporated from an electron beam evaporator (Omicron) and strontium was evaporated using a low-temperature effusion cell (CreaTec). The flux was monitored by a QCM. The temperature was measured with an optical pyrometer using an emissivity of 1. While this measurement method is not very accurate for oxides, the structures presented here are stable within a wide range of temperatures ($700\text{--}900^\circ\text{C}$), thus ambiguities in temperature measurement should not be a major problem.

All samples were treated with the following sequence. In order to get a clean, well-defined surface structure, samples introduced to the UHV system were sputtered (at least once) and then annealed. A $c(4 \times 2)$ structure (determined by LEED or STM) was observed after this procedure. Sr was evaporated onto this surface and the sample was annealed after deposition. Sr was deposited until a change of surface reconstruction to the (2×2) structure was observed (by LEED). For reversing the reconstruction, Ti was deposited until the structure changed back to the original surface reconstruction [$c(4 \times 2)$]. A full switch of reconstructions was obtained by depositing a nominal thickness (determined by the QCM) of $\sim 3 \text{ \AA}$ of Sr or $\sim 0.35 \text{ \AA}$ of Ti. Taking the densities of Ti and Sr and the size of the $\text{STO}(001)$ unit cell into account, this corresponds to 0.3 and 0.8 monolayers (MLs) for Ti and Sr, respectively. After deposition, the samples were annealed for 15 min (850°C in 10^{-6} mbar O_2).

3. Results

A two-domain $c(4 \times 2)$ reconstructed surface was obtained through multiple sputtering and annealing cycles; no other superstructures were detected by STM and LEED. Fig. 1(a) shows a large-scale STM image of this surface. The inset shows the associated LEED pattern. This structure was identified as the two-domain $c(4 \times 2)$ reconstruction using the software LEEDpat30 [24]. The terraces are $\sim 30 \text{ nm}$ wide and the step-height is equivalent to one unit cell of STO (3.9 \AA). An atomically resolved image of the two-domain $c(4 \times 2)$ structure and its corresponding fast Fourier transformation is shown in Fig. 1(a) with unit cells indicated.

One prominent feature of this surface is the straight step edges that run along the $[010]$ and $[100]$ directions. On the $c(4 \times 2)$ reconstructed surface, step edges aligned with the $\langle 100 \rangle$ directions have also been

reported by Jiang and Zegenhagen [17] and Castell [12]. Deposition of 0.8 ML (3 \AA QCM reading) of Sr onto this surface and annealing to 850°C in 10^{-6} mbar O_2 changes the surface (Fig. 1b). The LEED pattern reveals a (2×2) structure (inset). Again the terraces are $\sim 30 \text{ nm}$ wide with one unit cell step-height. The sharp LEED pattern indicates that the structure covers the surface uniformly and with good order for the most part. However, faint intermediate spots in the LEED image indicate the presence of a minority phase with different periodicity. Note the different appearance of the step edges; for the (2×2) structure the step edges are wavy, as was also found by Silly et al. [25]. Because it was not possible to image the (2×2) structure atomically, UHV flash-annealing ($\sim 850^\circ\text{C}$, 5 min) was applied (to increase the conductivity). However, the surface transformed into a structure with $c(4 \times 4)$ symmetry (not shown). This transition was already reported by Silly et al. [25] and the structural appearance in STM exhibited also the same $c(4 \times 4)$ structure, as was found in their work.

The process of transforming the surface structure can be reversed; switching between $c(4 \times 2)$ and (2×2) reconstructions was reproduced more than ten times. As indicated by the arrow in the bottom of Fig. 1, deposition of Ti onto the (2×2) structure and annealing at $\sim 850^\circ\text{C}$ in 10^{-6} mbar O_2 leads back to the two-domain $c(4 \times 2)$ structure. The amount of Ti necessary to switch between the two structures was measured to be 0.3 ML ($\sim 0.35 \text{ \AA}$ QCM reading). This procedure yields sharp LEED patterns, no degradation of the surface quality (as judged by LEED) was observed for multiple switchings.

To determine the chemical composition of the surface, XPS and LEIS were performed. Fig. 2 shows the XPS and LEIS spectra for the two-domain $c(4 \times 2)$ and the (2×2) reconstructed surface, respectively. Well-defined $\text{O}1s$, $\text{Ti}2p$, and $\text{Sr}3d$ core-level spectra were observed. The $\text{Ti}2p$ spectrum shows the single doublet feature of the Ti^{4+} state as in bulk STO. Furthermore, there is no detectable feature related to hydroxyls in $\text{O}1s$ spectra, indicating that those samples were fully oxidized and not affected by, e.g., adsorbed water. All peaks of the $c(4 \times 2)$ reconstructed surface have a binding energy that is 0.11 eV higher than those of the (2×2) reconstructed surface. This indicates downward band bending on the $c(4 \times 2)$ surface, i.e., a surface that is more positively charged. The ratio of the intensities of the $\text{Sr}3d_{5/2}$ to $\text{Ti}2p_{3/2}$ peaks was determined in order to reveal compositional differences. Within the statistical error, the XPS signal (in normal emission) did not show any difference (see Table 1). Note that the escape depth of photoelectrons

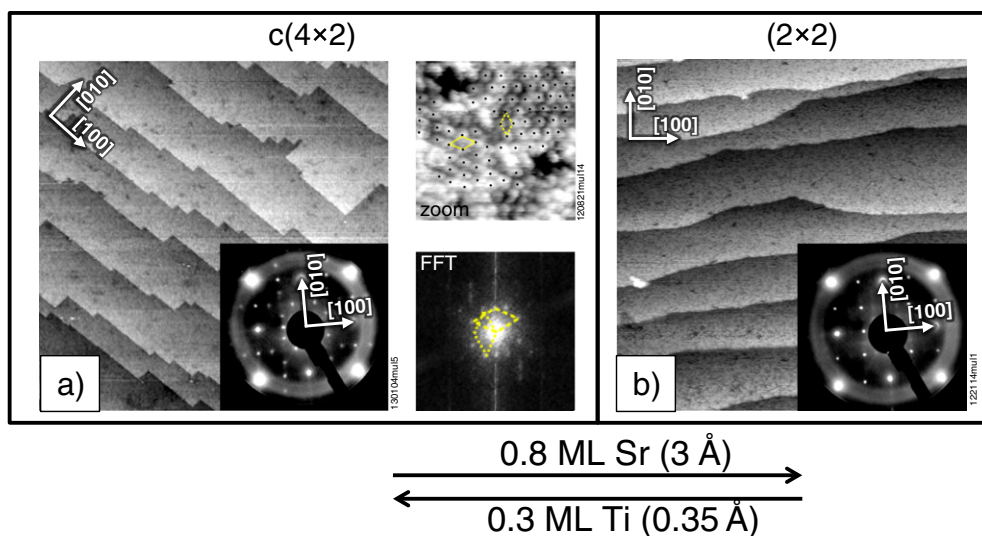


Fig. 1. Large-scale STM images ($200 \times 200 \text{ nm}^2$) of differently reconstructed $\text{STO}(001)$ surfaces. The insets show the LEED patterns at an electron energy of 69 eV. (a) Multiple sputtering and annealing cycles produce the two-domain $c(4 \times 2)$ surface (tunneling parameters: $V_s = +3.5 \text{ V}$, $I_t = 0.48 \text{ nA}$). An atomically resolved STM image and the corresponding FFT are shown at the side of (a). (b) Deposition of Sr with subsequent annealing changes the reconstruction to a (2×2) structure (tunneling parameters: $V_s = +2.3 \text{ V}$, $I_t = 0.26 \text{ nA}$). Ti deposition and annealing revert the surface to the $c(4 \times 2)$ structure. Note that the step edges are different in (a) and (b).

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