The effects of antimony doping on the surface structure of rutile TiO$_2$(110)

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Abstract
Titanium dioxide represents a very important wide bandgap photocatalyst that is known to be sensitized to visible light by transition metal doping. Antimony doping has been demonstrated to provide photocatalytic activity when codoped with chromium at an optimum dopant ratio [Sb]/[Cr] of about 1.5. Here, the role of antimony doping on the surface structure of rutile TiO$_2$(110) is studied using non-contact atomic force microscopy (NC-AFM) under ultra-high vacuum conditions. At first glance, the surface structure of antimony-doped TiO$_2$(110) resembles the structure of pristine TiO$_2$(110). However, in contrast to what is found in pristine TiO$_2$(110), a dense layer of protruding features is observed upon antimony doping, which is tentatively ascribed to antimony-rich clusters. Moreover, homogeneously distributed holes are found on the surface, which differ in depth and shape depending on the preparation conditions. Holes with depths ranging from a few up to more than a hundred monatomic steps are observed. These holes are explained by surface segregation of antimony during annealing, as the ionic radius of Sb$^{3+}$ is considerably larger than the ionic radius of Ti$^{4+}$. Our finding provides an indication of why an antimony concentration larger than the optimum ratio results in decreased photocatalytic activity. Moreover, controlling annealing temperature seems to constitute a promising strategy for creating nanosized holes on TiO$_2$ surfaces.

(Some figures in this article are in colour only in the electronic version)

1. Introduction
Titanium dioxide constitutes a very prominent and widely used example of a wide bandgap photocatalyst [1, 2], which is of high importance for the degradation of toxic compounds [3] and for the production of hydrogen [4]. It has been demonstrated that TiO$_2$ can be successfully sensitized to visible light by doping with foreign elements, providing an additional donor or acceptor level in the bandgap of TiO$_2$ [5]. In particular, doping with transition metals has been reported to result in an intense absorption of visible light [6, 7]. Among those transition metals, chromium has also been studied, revealing high visible-light absorption in reflection spectroscopy measurements [8]. However, despite the absorption in the visible-light region, chromium-doped TiO$_2$ has been found to be photocatalytically inactive [8]. This has been explained by the creation of additional oxygen vacancies upon chromium doping [8], as vacancies might constitute recombination sites for electron–hole pairs. Recent non-contact atomic force microscopy (NC-AFM) measurements have provided experimental evidence for vacancy creation upon chromium doping, further supporting this picture [9].

Interestingly, upon codoping with chromium and antimony, rutile titania particles have shown both visible-light absorption as well as photocatalytic activity. The rate of the photocatalytic O$_2$ production has been found to be optimized at a ratio of about 1.5, i.e. in excess of antimony [10, 8]. Even larger antimony excess has been observed to again result in reduced photocatalytic activity. The excess antimony most likely exists in equal amounts of antimony ions with the oxidation states Sb$^{3+}$ and Sb$^{5+}$, replacing Ti$^{4+}$ in the lattice. Consequently, they auto-compensate for charging. NC-AFM images of chromium- and antimony-codoped TiO$_2$ have revealed two important aspects, namely the absence of additional oxygen
vacancies in the case of codoping and a firm integration of the dopant atoms into the crystal lattice [11].

Here, we address the role of excess antimony in chromium- and antimony-codoped titania. The existence of antimony trivalent cations Sb\(^{3+}\) with a comparatively large ionic radius of 76.0 pm compared to 60.0 pm for Sb\(^{5+}\) and 60.5 pm for the replaced titanium ions Ti\(^{4+}\) [12] has been suggested to result in a deformed TiO\(_2\) lattice [8]. In the present work, we study the influence of antimony on the surface structure of rutile TiO\(_2\)(110) using high-resolution NC-AFM. Our results reveal that antimony-doped TiO\(_2\), at first glance, resembles the structure of pristine TiO\(_2\)(110). However, a dense layer of protruding features is observed, which is ascribed to antimony-rich clusters. Besides these clusters, different types of holes are observed on the surface. Shape and depth of these holes appear to depend on the annealing parameters. A possible scenario is given, explaining the formation of these holes based on the large ionic radius of Sb\(^{3+}\).

### 2. Experimental section

The scanning probe technique used in this study is NC-AFM [13] operated in the frequency modulation mode [14]. For this technique, a sharp nanoscopic tip mounted at the end of an oscillating cantilever is scanned in close proximity over the sample surface while keeping the oscillation amplitude constant. The forces acting between tip and surface atoms shift the cantilever’s eigenfrequency. This frequency shift can be directly used as an imaging signal when scanning the cantilever at a fixed tip–surface distance (constant height mode). Alternatively, the frequency shift can be kept constant to measure the height profile of equal frequency shift (referred to as topography). Measurements are performed at room temperature in an ultra-high vacuum (UHV) chamber (base pressure below 10\(^{-10}\) mbar) equipped with a VT AFM 25 (Omicron, Taunusstein, Germany) and an easy-PLL Plus phase-locked loop controller (Nanosurf, Liestal, Switzerland) for oscillation excitation and signal demodulation. We use silicon cantilevers (type PPP-NCH) from NanoWorld (Neuchâtel, Switzerland) with an eigenfrequency of about 300 kHz operated at an amplitude of 10 nm. For these experiments, the cantilevers are sputtered with Ar\(^+\) ions after insertion into the chamber. The TiO\(_2\) samples are crystals of the highest quality available (MTI, Richmond, MD, USA). The crystals were doped with antimony by calcining them together with Sb\(_2\)O\(_3\) powder in a crucible at 1420 K for 10 h in air [8]. The total amount of antimony in the doped crystal is difficult to quantify. However, in the upper few micrometers the antimony concentration is known to be of the order of 3 at.\% [10]. This order of magnitude was confirmed by x-ray photoelectron spectroscopy (XPS) measurements of the freshly calcined wafer. After insertion into the UHV chamber, the surface was cleaned by cycles of 15 min Ar\(^+\) ion sputtering at 1 keV and subsequent annealing. The sample was either annealed for 15 min at 1100 K or for 1 h at 930 K. The preparation parameters were chosen in accordance with our experience on pristine TiO\(_2\)(110). In order to minimize long-range electrostatic interactions, a bias voltage in the range of −0.6 V to −1.4 V was applied to the tip.

### 3. Results and discussion

Here, the investigation of a TiO\(_2\)(110) sample doped with an antimony concentration of about 3 at.\% is presented. We first describe the surface structure at the atomic scale and compare it to the surface structure of pristine titania. Second, the surface structure at larger frame size is presented, which is dominated by the formation of holes of different shape and depth. In the third part we highlight the influence of the annealing procedure on the hole formation.

Figure 1. NC-AFM topography images obtained on antimony-doped TiO\(_2\)(110) after annealing at 1100 K. The surface is found to be covered by a dense layer of protruding features, which is tentatively ascribed to antimony-rich clusters. The dark spot in the center of each cluster is a scan artifact, which can be explained by slow distance feedback loop settings as described in [17]. Besides the clusters, a row pattern resembling the TiO\(_2\)(110) surface can be identified in both images. The periodicity perpendicular to [001] was measured to be (660 ± 30) pm.
slightly bluish after UHV treatment, as shown in figure 2(b). The crystal is found to be transparent after calcination and only tied to a dark blue color of the entire crystal. In contrast, the sample would be heavily reduced. Such a reduced sample the features observed here might be of a different nature.

However, the features observed here appear to have a much larger diameter than typical hydroxyl defects, indicating that the features observed here might be of a different nature. Moreover, if the observed features would be hydroxyl defects, the sample would be heavily reduced. Such a reduced sample is tied to a dark blue color of the entire crystal. In contrast, the crystal is found to be transparent after calcination and only slightly bluish after UHV treatment, as shown in figure 2(b).

We cannot rule out that antimony doping results in surface rather than bulk reduction due to external strain [16], which would not necessarily be accompanied by a dark blue color. However, as outlined below, we favor the assignment of the protruding features being antimony-rich clusters rather than hydroxyl defects.

Besides the clusters, the typical row pattern of the rutile surface is also visible in the images in figure 1. The periodicity perpendicular to the [001] direction was measured to be (660 ± 30) pm, which is within the error identical to the surface unit cell length of the pristine titania surface.

Figure 3 reveals that the clusters cover all terraces and islands of the surface. Interestingly, the clusters seem not to adsorb in close proximity to the step edges. In a thin stripe at the edges of the islands (indicated by red arrow), the line pattern of the surface is visible. From a Fourier transformation of this image (not shown), the lines are measured to be separated by (670 ± 30) pm, similar to what was obtained from figure 1. The findings at the atomic scale are independent of the applied annealing recipes (i.e. 15 min at 1100 K or 1 h at 930 K). In particular, the density of clusters was found to be virtually identical after each preparation cycle.

3.2. Holes

A typical image of large frame size obtained on the antimony-doped titania surface is presented in figure 4. The terraces are found to be formed similar to what we typically observe on pristine TiO$_2$(110). The monatomic step height was measured to be identical to what we found on pristine titania.

Furthermore, two deep holes are visible in figure 4(a) with a diameter of about 50 nm. The depth of these holes cannot be obtained from NC-AFM measurements as explained in the following. The line profiles presented in figure 4 show a horizontal scan line in the forward direction of the topography (c), the detuning (d) and the amplitude (e) image. At the beginning of the hole the detuning drops to zero, indicating a too large tip–sample distance. The topography signal shows that the scanner approaches the sample at the hole site by more than 10 nm, but the detuning remains zero. Apparently, the base of the hole was not reached by the tip. At the end of the hole, the tip strongly interacts with the backside of the hole, which is indicated by large negative and even positive detuning values. This coincides with a decrease in amplitude of about 10%, indicating a highly dissipative interaction. This behavior was observed at all hole sites, even with the fastest distance feedback loop settings available. From various attempts to scan such holes at small frame size (not shown), the typical depth can be assumed to be definitely larger than 30 nm. Assuming an aspect ratio of the tip of typically 2 at the highest, which corresponds to a cone angle of the tip apex of at least 90°, the depth of a 50 nm wide hole is—as a matter of principle—not accessible if the depth is larger than 25 nm.

The holes are found to be rather homogeneously distributed at the surface as indicated by figure 5. The mean distance between neighboring holes was measured to be about 1 μm. Interestingly, all holes are found to be surrounded by small islands as depicted by the detailed images in figure 5, resembling a volcano-like crater.
Figure 4. NC-AFM topography (a) and detuning (b) image obtained on antimony-doped TiO$_2$(110). Two holes can be identified with a diameter of about 50 nm. Line profiles depict a horizontal scan line in forward direction of the topography (c), the detuning (d) and the amplitude (e) signal at a hole site (indicated by the black line). The line profiles indicate that the hole must be deeper than 12 nm as the detuning remains zero within the hole while the scanner approaches the surface. At the backside of the hole, the tip strongly interacts with the sample as the scanner cannot retract the tip fast enough. This is indicated by even positive detuning values and a decrease in amplitude.

Figure 5. NC-AFM topographic images of antimony-doped TiO$_2$(110) obtained after annealing at 1100 K. In total, 13 holes are visible with a mean distance of about 1 µm. The cut-outs extracted from (a) reveal that all holes are surrounded by small islands. This results in an appearance that resembles a volcano-like shape. The step edges can be found to run either parallel to the [001] direction (a) or enclose an angle with the [001] direction (b) as indicated by the red and black arrows. Cone-shaped troughs located at the hole sites and aligned in the [001] direction are, therefore, only observed in the case of step edges that are not parallel to [001]. The holes (black circles) and cone-shaped troughs (gray triangles) identified in (b) are sketched in (c) for illustration.
Figure 6. NC-AFM topographic images of antimony-doped TiO$_2$(110) obtained after annealing at 930 K. The terraces are perforated and numerous small islands exist, similar to the pristine surface when annealed at the same temperature. Besides a few large and deep holes, which appear identical as observed after annealing at 1100 K, a large number of small pits are present here. These pits are typically less than 2 nm deep and less than 10 nm wide. All holes in figure 5(b) appear at the sharp end of a cone-shaped trough located at each hole. For illustration, these troughs are sketched in figure 5(c). The holes in figure 5(a) have no troughs. This difference can be explained by differently oriented step edges in the two images, as explained in the following. In figure 5 two typical images are presented, either showing step edges running parallel to the [001] direction (a) or step edges that enclose an angle with the [001] direction (b), as indicated by the red and black arrows. The cone-shaped troughs are also aligned in the [001] direction. Assuming that the terraces are growing in the [001] direction during annealing, the observed troughs might be explained by shadowing at hole sites.

3.3. Influence of annealing temperature

The images presented in figure 5 were obtained after annealing at 1100 K for 15 min. For comparison similar images obtained after annealing at 930 K for 1 h are shown in figure 6. The terraces appear perforated and covered by small islands similar to what we observed on pristine TiO$_2$(110) annealed at the same temperature. In contrast to the high temperature prepared doped surface, here a large number of small pits exist besides the large, deep holes. These small pits have a typical depth of 1–2 nm and an average nearest-neighbor distance of about 50 nm.

Besides the two already described types of holes, a third type of hole is seldom observed after annealing at 1100 K, which is typically less than 10 nm deep, but extends over several hundred nanometers. Two of this kind are shown in figure 7. A detailed analysis of the flat holes revealed that the typical surface structure with step edges of monatomic height can be found at the bottom of the holes, as indicated by figure 8. Besides these flat, extended holes also deep holes with their cone-shaped trough as introduced in figures 5(b) and (c) exist in the surrounding surface region.

The images presented in figures 5(b) and 7 are obtained after the very same preparation cycle at macroscopically different surface areas. However, the size and number of holes are different. Taking the form and flatness of terraces as the reference for the annealing temperature and assuming large, island-free terraces to result from higher temperature preparation, the situation found in figure 7 might be explained by an annealing temperature slightly lower than 1100 K. Due to different thermal contact to the silicon stripe used for resistive heating of the sample, slightly different annealing temperatures at macroscopically different sample sites are conceivable. Obviously, the number as well as the size of the holes and pits crucially depend on the exact annealing procedure. Thus, it might be possible to control the size and density of the pits by properly tuned annealing parameters.

3.4. Summary

In the following, we outline a possible scenario for explaining the observed structural characteristics. Antimony induces a large lattice stress in the titania crystal as already suggested by the too large ionic radius of Sb$^{3+}$. Actually, an Sb$^{3+}$ ion requires almost double the volume an Ti$^{4+}$ ion occupies when considering the ionic radii. Although Sb$^{3+}$ ions nearly perfectly fit into the crystal, they cannot remain alone as this would lead to charging of the crystal. Consequently, antimony cannot be integrated into the titania lattice at all. Thus, antimony is observed to leave the titania crystal when mobile enough. Upon vacuum annealing, antimony can segregate to the surface, forming segregation channels that appear as nanosized volcanos. At low annealing temperature, only antimony from the upper layers might have enough thermal
energy to reach the surface. Thus, a large number of small, flat pits is created.

At higher annealing temperature the mobility of antimony dopants is increased. Hence, material from deeper layers can also segregate to the surface. Furthermore, more material is segregated on a shorter timescale. The large and deep holes, created upon high temperature annealing, indicate that the sample surface is almost exploding as soon as antimony gets enough thermal energy to leave the bulk. In this case, instead of numerous small pits, only a few large segregation channels are established to reduce the surface energy. These holes are rather homogeneously distributed. Cone-shaped troughs are observed to exist close to hole sites, which might result from shadowing at these holes. A possible explanation is that terraces growing during annealing are distorted at the holes. The erupted material consisting of TiO₂ forms new islands around the holes resulting in the volcano-like appearance. Antimony, erupted through the holes, is adsorbed as clusters on top of the surface.

4. Conclusion
In conclusion, the antimony-doped surface was found to exhibit terraces, steps and a row pattern similar to what is typically observed on pristine titania. However, the
terraces were covered by a dense layer of clusters, which was tentatively explained by antimony-rich clusters. Holes with depths ranging from a few up to more than a hundred monatomic steps were observed. The width and depth of these holes were found to crucially depend on the annealing procedure. The deepest holes were surrounded by small islands, resembling a volcano-like shape.

The different observed structures were explained by a simple model considering the ionic radii of the dopant atoms. Our findings draw a picture of weakly integrated antimony in antimony-doped titania. These results, thus, also indicate why an excess of antimony beyond the optimum ratio of 1.5 is unfavorable for photocatalytic applications, as the additional antimony ions cannot be integrated into the bulk. Moreover, the presented study revealed another interesting phenomenon. It seems to be possible to create nanosized holes in a controlled way by tuning the annealing parameters of antimony-doped TiO$_2$(110). This might be an appropriate strategy for creating nanopatterned surfaces with macroscopic techniques.

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