Structure of a Superhydrophilic Surface: Wet Chemically Prepared Rutile-TiO$_2$(110)(1 × 1)

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Supporting Information

ABSTRACT: Surface X-ray diffraction has been employed to quantitatively determine the geometric structure of an X-ray-induced superhydrophilic rutile-TiO$_2$(110)(1 × 1) surface. A scatterer, assumed to be oxygen, is found at a distance of 1.90 ± 0.02 Å above the five-fold-coordinated surface Ti atom, indicating surface hydroxylation. Two more oxygen atoms, situated further from the substrate, are also included to achieve the optimal agreement between experimental and simulated diffraction data. It is concluded that these latter scatterers are from water molecules, surface-localized through hydrogen bonding. Comparing this interfacial structure with previous studies suggests that the superhydrophilicity of titania is most likely to be a result of the depletion of surface carbon contamination coupled to extensive surface hydroxylation.

INTRODUCTION

Ever since Wang et al.’s discovery that UV irradiation of titania results in a superhydrophilic surface, there has been a great deal of effort to both exploit and understand this novel phenomenon. Significant success has been achieved in the former of these two goals, with applications including self-cleaning windows and antifogging mirrors. In contrast, uncertainty still remains as to the origin of the superhydrophilicity. Currently, there are a number of potential explanations to be found in the literature, but none are supported by compelling experimental evidence. For instance, it is proposed that the superhydrophilicity is simply a result of the removal of surface carbon contamination. Other researchers suggest that modification of the surface chemistry/chemistry of the titania substrate (e.g., surface hydroxylation) underpins this macroscopic property. Longer range structural changes are also purported to be important, including the formation of nanoscale hydrophobic and hydrophilic domains. Here we directly address this topic, employing surface X-ray diffraction (SXRD) to quantitatively determine the structure of a model titania surface, rutile-TiO$_2$(110), that exhibits superhydrophilicity induced through X-ray exposure.

Previously, Shirasawa et al. (SEL) have undertaken SXRD measurements from rutile-TiO$_2$(110) to identify changes in surface structure associated with the UV-induced hydrophobic-to-hydrophilic transition. They report that the application of a wet chemical preparation (WCP) recipe to the substrate resulted in a hydrophobic (1 × 1) surface termination, which became hydrophilic upon UV irradiation. It is suggested that this transition is associated with the presence of surface hydroxyls (OH$_x$), as surface five-fold-coordinated titanium atoms (Ti$_{5c}$) and bridging oxygens (O$_b$) become hydroxylated following the exposure to UV light. Figure 1 illustrates the changes in interface geometry concluded in ref 13.

In this Article, we revisit the structure of the superhydrophilic rutile-TiO$_2$(110)(1 × 1) surface. A WCP method...
Concerning sample preparation, a previously published WCP recipe was again employed for sample preparation, but with X-rays being used to induce superhydrophilicity. Similar to ref 13, the surface is found to be extensively hydroxylated, including OH bound to Ti ions. The diffraction data acquired in this study, however, resemble much more closely those acquired from the hydrophobic termination in ref 13. As argued in detail below, this somewhat unexpected finding suggests that the analysis and interpretation of SEL require revision.

EXPERIMENTAL METHODS

Concerning sample preparation, a previously published WCP recipe was applied to the rutile-TiO$_2$(110) substrate (10 × 10 × 1 mm sample from PI-KEM), which is known to produce a well-ordered (1 × 1) surface termination. In brief, this procedure involves sequential sonication in acetone, ethanol, and deionized water and then annealing in air at ∼973 K for ~90 min. Subsequently, the sample is immersed in aqua regia solution (3:1 by volume ratio of concentrated HCl and HNO$_3$) at room temperature for ~45 min. The final UV-ozone treatment step described in ref 14 was not undertaken in this study. Please note that the TiO$_2$ sample remained transparent following the application of this WCP recipe, indicating that there was no bulk reduction.

Upon the completion of surface preparation, surface hydrophilicity was evaluated by delivering a small droplet of deionized water to the TiO$_2$(110) surface using a syringe/hypodermic needle. Once the contact angle of the deposited droplet had been determined by visual inspection, the sample was blown dry with high-purity nitrogen. The sample was then transferred to the diffractometer located in EH1 of beamline I07 at the Diamond Light Source synchrotron facility for SXRD measurements. It was mounted in an X-ray transparent (cylindrical Kapton window) environmental cell. Once the cell was closed up, high-purity helium was flowed through it for the duration of the SXRD measurements. It should be noted that it was not possible to monitor either oxygen or water vapor concentration within the cell.

SXRD data were collected at an incidence angle of 1° with the substrate at room temperature using a photon energy of $hv = 17.7$ keV and a 2D Pilatus photon detector. A systematic series of X-ray reflections was acquired from the sample; that is, for a given $(h,k)$ integer, data were measured as a function of $l$ to facilitate the generation of so-called crystal truncation rods (CTRs). $h$, $k$, and $l$ are the reciprocal lattice vectors. They are defined with reference to the real-space (1 × 1) unit cell of the (110) surface, described by lattice vectors $(a_1, a_2, a_3)$ which are parallel, to the [110], [001], and [110] directions, respectively; $a_1 = a_2 = a\sqrt{2}$, and $a_3 = c$ where $a = 4.593$ Å and $c = 2.958$ Å are the lattice constants of the tetragonal rutile crystal structure. A surface-sensitive reflection (i.e., one that is located well away from any bulk Bragg peak), namely, $(-1, 0, 0.9)$, was recorded at regular intervals to monitor the surface integrity.

To facilitate fully quantitative structure determination, the raw 2D diffraction images were integrated, including background removal, and corrected to enable plots of structure factor versus perpendicular momentum transfer for each CTR to be compiled. This procedure generated a total of 1068 nonequivalent reflections from eight distinct CTRs. The ROD software was employed to simulate these data, with structural and nonstructural parameters being refined to achieve the overall best fit between experiment and theory. Reduced $\chi^2$ was used to evaluate the goodness of fit, which is defined as follows:

$$\chi^2 = \frac{1}{N-P} \sum_{i=1}^{N} \left( \frac{|F_{\text{exp}}(hkl)| - |F_{\text{th}}(hkl)|}{\sigma_{\text{th}}(hkl)} \right)^2$$

$N$ is the number of measured structure factors, $P$ is the number of parameters optimized during fitting, and $F_{\text{exp}}(hkl)$ and $F_{\text{th}}(hkl)$ are the experimental and theoretically calculated structure factors, respectively. $\sigma_{\text{th}}(hkl)$ is the uncertainty associated with $F_{\text{th}}(hkl)$. $\chi^2$ behaves such that a value of 1 indicates that experiment and theory are essentially coincident, with agreement decreasing with increasing $\chi^2$. The quoted precision of each fitted parameter is determined by systematically varying the parameter about its optimal value and for each step optimizing all other parameters, until $\chi^2$ has increased by 1/($N-P$) from its minimum value.

RESULTS AND DISCUSSION

The application of our WCP recipe to the rutile-TiO$_2$(110) sample resulted in a deionized water contact angle of ~80°. This value is consistent with that reported in ref 14 for a (1 × 1) surface termination subsequent to immersion in aqua regia but not exposed to UV-ozone treatment. Following exposure to 107°'s photon beam, the contact angle was found to fall to essentially 0°; that is, a superhydrophilic transition was induced by X-ray exposure. All diffraction measurements were undertaken with the rutile-TiO$_2$(110)(1 × 1) surface in this state; a contact-angle measurement at the end of data collection indicated that a value of 0° was maintained throughout this period. Data from the (-1, 0, 0.9) reflection also revealed no substantive surface degradation.

Figure 2 shows four of the experimental CTRs acquired in the current study (black markers with error bars), together with equivalent data collected by SEL from rutile-TiO$_2$(110) following UV exposure (blue markers with error bars). A priori, as both data sets were recorded from superhydrophilic
surfaces, it was expected that they would be very similar. However, there are significant differences. For example, the local maxima in our data at (0, 1, ∼3) and (1, 0, ∼2), are not replicated in those from SEL. In contrast, our CTR profiles are much more comparable to those reported by SEL for their pre-UV exposure (hydrophobic) surface. These data are also shown in Figure 2 as red markers with error bars. We note that on an adsorbate-free rutile-TiO2(110)(1 × 1) surface, prepared in ultrahigh vacuum (UHV), the aforementioned local maxima are associated with significant displacements of surface atoms away from their bulk positions; that is, they are not a direct signature of surface hydrophilicity. An absence of such features may be a result of either a more bulk-like termination or surface roughening.

Considering the qualitative comparison outlined above, it was expected that fitting of our experimental SXRD data would result in the hydrophobic structure determined by SEL, where molecular H2O is bound atop TiO (see Figure 1). Figure 3 shows the best fit (blue line) achieved using SEL’s hydrophobic structure as a starting point and simply allowing the displacement of both atomic positions and nonstructural parameter values. As indicated by \( \chi^2 = 2.60 \), as well as visual inspection, the experiment–theory agreement is far from perfect, suggesting that the correct structural solution had not been found. On this basis, we explored other potential surface terminations, including those consistent with the presence of surface hydroxyls. The resulting overall best fit to the experimental CTRs is shown in Figure 3 (red line). To achieve this fit, 78 parameters were optimized, that is, 51 atomic coordinates, 21 Debye–Waller (DW) factors, a scale factor, surface roughness (\( \beta \)), three fractional occupancies, and surface fraction. The corresponding \( \chi^2 \) is 1.05; that is, there is an excellent level of agreement between the experimental and simulated data.

The optimum geometry of the first few atomic layers emerging from the best fit to the experimental CTR profiles is depicted in Figure 4. Selected corresponding interatomic distances are listed in Table 1. A ball-and-stick model showing all atoms displaced during fitting is shown in Figure S1, along with a complete list of the optimized coordinates, DW factors,

Figure 2. Comparison of the (0, 1, 0), (1,0, 0), (1, 1, 0), and (2, 0, 0) experimental CTRs acquired from X-ray-induced superhydrophilic rutile-TiO2(110) in the current study with data from SEL. Current study: black markers with error bars; pre-UV exposure from SEL: red markers with error bars; post-UV exposure from SEL: blue markers with error bars. Profiles are systematically offset for clarity.

Figure 3. Comparison of experimental CTR data (black markers with error bars), acquired from X-ray-induced superhydrophilic rutile-TiO2(110), and theoretical best-fit simulations. Solid blue line indicates the best fit achieved following relaxation of the hydrophobic structure reported by SEL. Solid red line indicates the overall best fit, with \( \beta = 0.24 \) and a surface fraction of 0.96.

Figure 4. Ball-and-stick models of the X-ray-induced superhydrophilic rutile-TiO2(110) surface structure determined from SXRD data. Perspective (top) and plane (bottom) views are shown. Red spheres are Ti atoms, and darker (lighter) blue spheres are substrate (adsorbate) oxygen atoms. H atoms (pink spheres) are depicted, although they were not explicitly included in the structure determination. Possible hydrogen bonds are indicated by dashed lines. The numerical labeling of the atoms is employed in Table 1 and Table S1 for identification purposes. Symmetry-paired atoms are denoted by *.

Table 1. Selected Interatomic Distances Derived from Atomic Coordinates (Table S1) of Optimized Superhydrophilic TiO2(110)(1 × 1) Structure

| atoms     | interatomic distance (Å) |
|-----------|--------------------------|
| O(3′)−O(1′) | 2.70 ± 0.06             |
| O(3′)−O(2′) | 2.65 ± 0.05             |
| O(2′)−O(1′) | 2.68 ± 0.03             |
| O(1′)−Ti(2) | 1.90 ± 0.02             |
| O(3)−Ti(1)  | 1.83 ± 0.02             |
| O(2)−Ti(1)  | 1.98 ± 0.02             |
| O(2)−Ti(2)  | 1.95 ± 0.02             |
| O(3)−Ti(1)  | 1.94 ± 0.01             |
| O(4)−Ti(2)  | 1.94 ± 0.01             |
molecules, which are localized through hydrogen bonding. Oxygen atoms O(2) at a distance of 2.65 Å from Ti(2) are located atop at a distance of 1.90 ± 0.02 Å, which is consistent with the presence of a bound terminal hydroxyl (OHt). The oxygen atoms O(3) are at somewhat greater distances from the topmost substrate atoms. (labeled O(1)), with O(3)′ nearest neighbor being O(1)′ at a distance of 2.65 ± 0.05 Å. These interatomic separations suggest that oxygen atoms O(2)′ and O(3)′ arise from water molecules, which are localized through hydrogen bonding.

Figure 5. Ball-and-stick models of the optimum interfacial structures determined from SXRD for X-ray-induced superhydrophilic rutile-TiO2(110) (current study), rutile-TiO2(110) subsequent to dipping in H2O(l), and rutile-TiO2(110) submerged in H2O(l). Side (top) and plane (bottom) views are shown. Red spheres are Ti atoms, and darker (lighter) blue spheres are substrate (adsorbate) oxygen atoms. Selected interatomic distances are annotated. Fractional occupancies of adsorbate oxygen atoms are indicated by the values inscribed on the lighter blue spheres.

and fractional occupancies in Table S1. Neglecting the details of atomic relaxation, the surface mimics the stoichiometry and geometry of bulk-terminated rutile-TiO2(110)(1 × 1) but is decorated with oxygen species. Focusing on Ti5c (labeled Ti(2)), an adsorbed oxygen atom (labeled O(1)′) is located atop at a distance of 1.90 ± 0.02 Å, which is consistent with the presence of a bound terminal hydroxyl (OHt). The experimental distance from ref. 20 is 1.95 ± 0.03 Å, with a moderately longer distance of 2.07 Å being obtained from molecular dynamics calculations. Two additional non-substrate oxygen atoms (labeled O(2′) and O(3′)) are at somewhat greater distances from the topmost substrate atoms. O(2′) is 2.68 ± 0.03 Å above the bridging oxygen atom (labeled O(1)), with O(3)′ nearest neighbor being O(1)′ at a distance of 2.65 ± 0.05 Å. These interatomic separations suggest that oxygen atoms O(2′) and O(3′) arise from water molecules, which are localized through hydrogen bonding.

For illustrative purposes, we have included H atoms in Figure 4 but stress that these species were not explicitly included during the generation of simulated SXRD data due to their negligible X-ray scattering.

Given the optimized structure displayed in Figure 4, it is interesting to compare this result with other pertinent studies. Focusing initially on SEL’s work, the present diffraction data are very similar to those acquired from their hydrophobic surface, as demonstrated in Figure 2. Because our surface is superhydrophilic, as a result of X-ray exposure, this agreement presents a conundrum. One plausible explanation, arising from discussion with SEL, is that the Ti5c footprint of the X-ray beam employed for their SXRD measurements induced superhydrophilicity only in this region. Hence the water contact-angle measurement, where the droplet employed covered a much larger surface area, did not reveal this local X-ray-induced superhydrophilicity. It should be noted that on 107 almost the entire sample surface would have been exposed to the X-ray beam during alignment and measurement.

On the basis that SEL’s pre-UV-exposure surface is superhydrophilic in the area probed by the X-ray beam, then one further issue requires resolution. Specifically, despite the similar experimental data, the discrepancy between our optimized structure and SEL’s needs to be understood, for example, the variation in the Ti(2)–O(1)′ distance (1.90 ± 0.02 versus 2.09 ± 0.03 Å). To this end, our experimental data set (eight CTRs) was reduced to match that of SEL (six CTRs), and fitting was undertaken. Under these conditions, we were able to effectively model the data with SEL’s hydrophobic structure. On this basis, it is evident that fewer experimental CTRs leads to a local χ2 minimum, resulting in a significantly different surface structure.

One other matter emerging from the preceding discussion is the origin of the UV-induced change in CTR profiles observed by SEL. Assuming that their pre-UV data is acquired from a superhydrophilic area of the rutile-TiO2(110) surface, then the observed changes cannot be accounted for by a hydrophobic–hydrophilic transition. This deduction implies that UV-irradiation leads to additional interfacial modification; that is, a unique surface structure is formed upon exposure to UV light. Currently, this suggestion is essentially conjecture, but is worthy of further investigation.

Having reconciled the results of this study with those of SEL, a comparison of the geometry of the current superhydrophilic termination with those reported for interfaces formed by the exposure of UHV-prepared TiO2(110)(1 × 1) to liquid water (H2O(l)) is worthwhile. Figure 5 compares the current optimum structure (TiO2(110):Super) that elucidated with SXRD following dipping of TiO2(110)(1 × 1) into H2O(l) and measuring ex situ in UHV (TiO2(110):Dip-H2O(l)) as well as that determined for TiO2(110) submerged in H2O(l) (TiO2(110):Sub-H2O(l)). These three structures are similar but not identical. For example, both TiO2(110):Super and TiO2(110):Sub-H2O(l) exhibit oxygen atoms consistent with hydrogen-bonded H2O molecules, although their configuration differs; such scatterers are not evident in the TiO2(110):Dip-H2O(l) data due to the acquisition in UHV. Turning to Ti5c for each structure displayed in Figure 5, the distance to the atop oxygen atom is consistent with hydroxylation (Ti5c–OH). However, TiO2(110):Super displays a slightly shorter Ti5c–OH distance (1.90 ± 0.02 Å) than either TiO2(110):Dip-H2O(l) or TiO2(110):Sub-H2O(l) (1.95 ± 0.03 Å). This variation may be a result of the former substrate being essentially fully oxidized, whereas the latter two were somewhat reduced as a result of substrate preparation in UHV.
Regarding the origin of the X-ray-induced superhydrophilicity of titania, the current SXRD data rule out the coexistence of hydrophilic and hydrophobic domains, as analysis indicates that almost the entire surface adopts the same geometry; that is, surface fraction is 0.96. It should be emphasized that the present study cannot be used to definitively rule out the existence of such domains on UV-exposed titania. Furthermore, because the diffraction data from TiO$_2$(110):Dip-H$_2$O(l) were acquired in UHV from a surface not irradiated with either UV or X-rays during dipping, the mere presence of OH$^-$ cannot be directly related to photoinduced superhydrophilicity. Given this result, one could suggest that the simple removal of surface carbon most likely underpins this property. It is, however, notable that the fractional occupancy of OH$^-$ (O(1')) for TiO$_2$(110):Super is approximately double that for either TiO$_2$(110):Dip-H$_2$O(l) or TiO$_2$(110):Sub-H$_2$O(l), that is, 1.00 compared to 0.45 and 0.50, respectively, as indicated in Figure 5. Hence, increased surface hydroxylation may play a role in TiO$_2$ superhydrophilicity, coupled to the loss of surface carbon. We remark that in ref 20 ab initio modeling suggests that the presence of OH$^-$ is a result of (near) surface defects driving surface H$_2$O dissociation. Because the substrate in the current study is not expected to possess any significant concentration of defects, the hydroxyl species must arise from elsewhere. Almost certainly, it is photon-induced (or photoexcited electron) chemistry that produces these OH$^-$ adsorbates, which may be the reason that a higher coverage is achieved; we note that this increase in surface hydroxylation is not simply related to carbon removal, as the surfaces in ref 20 are reported to be relatively carbon-free (≤0.1 monolayer).

Finally, we would like to comment on a recent elegant study suggesting that air- or aqueous-solution-exposed rutile-TiO$_2$(110) is commonly decorated by carboxylate species. On the basis that SXRD is not a spectroscopic probe, there is always the potential for misidentification of adsorbates, especially those exhibiting similar X-ray scattering characteristics (e.g., C and O). For the current study, however, we argue that this is not the case. Supporting evidence is two-fold. First, Auger spectra acquired from a superhydrophilic rutile-TiO$_2$(110) surface, prepared following our WCP recipe, show no discernible carbon signal. Second, SXRD data were acquired from a superhydrophilic surface, which is inconsistent with the presence of adsorbed carboxylates.

## CONCLUSIONS

To summarize, SXRD data have been acquired from an X-ray-induced superhydrophilic rutile-TiO$_2$(110) (1 × 1) surface. It is concluded that the five-fold-coordinated surface Ti atom is hydroxylated, as indicated by the presence of an atom, assumed to be O, at a distance of 1.90 ± 0.02 Å. There is also evidence of hydrogen-bonded H$_2$O molecules, which are located somewhat further from the substrate surface. The examination of the current structure, in tandem with previous work, suggests that the X-ray-induced superhydrophilicity of titania is likely to be a result of both the depletion of surface carbon and increased surface hydroxylation.

## ASSOCIATED CONTENT

**Supporting Information**

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jpcc.9b00245.
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