

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *J. Mater. Chem. A*, 2020, **8**, 18173Chemical and photoelectrochemical instability of amorphous TiO₂ layers quantified by spectroscopic ellipsometry†H. Kriegel,^a J. Kollmann,^a R. Raudsepp,^a T. Klassen^{ab} and M. Schieda^{*,a}

Amorphous TiO₂ films deposited by Atomic Layer Deposition (ALD) are recently being employed as corrosion protection coatings for photoanodes and photocathodes. However, these protective films are not immune to degradation. We have applied spectroscopic ellipsometry to quantify the loss of material, and the corresponding etch rates, for amorphous TiO₂ in photoelectrochemical cells under a wide range of conditions, at open- and short-circuit, in darkness and under illumination with different wavelength ranges. In 0.5 M sulfuric acid, we found corrosion to proceed *via* two routes: a chemical process, with an activation energy of 57 kJ mol⁻¹, and photoelectrochemically, once excess generated charge carriers are extracted. Compared to the dark condition, the combined degradation rate is more than doubled under illumination at 100 mW cm⁻² and at pH 0.3–2. The observed etch processes are highly dependent on the acidic pH values and largely suppressed at pH > 4. This has important implications for the selection of operating conditions for practical photoelectrochemical cells using amorphous TiO₂ protective coatings.

Received 11th May 2020

Accepted 27th July 2020

DOI: 10.1039/d0ta04878j

rsc.li/materials-a

Introduction

In the pursuit of efficient and stable photoelectrodes for solar water splitting, protective layers of ALD-grown, amorphous TiO₂ have had a large impact on recent and current research strategies in stabilizing efficient but intrinsically unstable absorber materials.¹ A number of groups have demonstrated the potential of thin layers of TiO₂ to extend the lifetime of photoanodes and photocathodes based on silicon,^{2–4} GaAs,⁵ CdTe,⁶ CZTS⁷ and Cu₂O⁸ under PEC (Photoelectrochemical Cell) conditions. Because TiO₂ is intrinsically a poor catalyst for hydrogen or oxygen evolution, such protected electrodes are commonly decorated with an additional catalyst like platinum or nickel. The metal serves to adjust the Fermi-level of the layered system, extracts and stores charge carriers and facilitates charge transfer for the desired reaction. This also reduces the available charge carriers for the electrochemical corrosion pathways of TiO₂ under reducing or oxidizing conditions. The protected systems have shown remarkable photocurrents over hours and days of measurement but often fail after extended testing due to a loss of catalyst^{8–10} or pinhole enabled etching of the underlying absorber.¹¹

Crystalline TiO₂ in the anatase phase has been debated to be either beneficial¹² or detrimental¹⁰ for the electrode functionality, but is accepted to be more stable than amorphous coatings.^{3,10,13} Studies on RF magnetron sputtered TiO₂ on the other hand highlighted the etchability of amorphous titania^{14,15} in warm sulphuric acid or phosphoric acid mixed with hydrogen peroxide. Once crystallized, the etch rate was shown to be more than 20 times lower at 95 °C.¹⁴ Furthermore, recent thermodynamic investigations have determined for amorphous TiO₂ in acidic conditions at 25 °C a solubility three orders of magnitude higher than for the crystalline phases.^{16,17}

Even though decoration with electrocatalysts limits their degradation,^{18,19} in the long term the stability of amorphous TiO₂ protective layers in PEC devices will be determined by their intrinsic behaviour in the corresponding chemical and photoelectrochemical environments. Nevertheless, there seem to be no published systematic studies of etch rates or stability windows under operating conditions for these coatings.

In this work, we have evaluated the corrosion of amorphous ALD-grown TiO₂ layers on highly doped silicon in a PEC device, in the dark and under illumination, with focus on 0.5 M sulfuric acid (H₂SO₄) as the electrolyte. Etch rates were determined from residual layer thicknesses modelled by ellipsometry data measured *ex situ*. The impact of the resulting cell temperature, as well as the influence of the incident illumination wavelength on the etch rate was studied by employing optical filters during the corrosion experiments. In addition, the contribution of photoelectrochemical corrosion could be differentiated from chemical corrosion by careful selection of experimental parameters.

^aHelmholtz-Zentrum Geesthacht, Centre for Materials and Coastal Research GmbH, Max-Planck-Str. 1, D-21502 Geesthacht, Germany. E-mail: mauricio.schieda@hzg.de^bUniversity of the Federal Armed Forces, Helmut-Schmidt-University, Holstenhofweg 85, D-22043 Hamburg, Germany

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ta04878j

Experimental

Sample preparation

Amorphous TiO₂ films were grown on highly doped silicon wafers (5 mohm cm) by plasma assisted Atomic Layer Deposition (ALD). Substrates were cleaned in buffered oxide etch (1 : 6 HF : NH₄F) prior to loading in the ALD reactor (Oxford Instruments FlexAL). The depositions were performed at 100 °C, using tetrakis(dimethylamido)titanium (TDMAT) and oxygen plasma. The steps consisted of 0.8 s precursor dose, 6 s Ar purge, 1 s pre-plasma gas stabilisation, 3 s plasma with RF power of 300 W and 4 s post-plasma purge. 900 cycles were used to obtain 68 nm TiO₂ film thickness at a growth rate of 0.77 Å per cycle. For comparison, crystalline films were grown with the same parameters at 300 °C, resulting in 60 nm films of TiO₂ in anatase phase.

Characterization

Raman spectra of as-grown films were measured with a WITec confocal Raman-microscope (alpha300 R) using a 633 nm laser. X-ray diffractograms were acquired with a Bruker D8 diffractometer under grazing incidence angle of 1.0° using a Cu K-α source.

The (photoelectro-)chemical corrosion experiments were conducted in a commercial PEC cell (Zahner-Elektrik) with a quartz window for illumination and the exposed sample area defined by a 5 mm O-ring. Illumination was supplied by a 300 W xenon lamp (LOT-Oriel) fitted with an AM1.5 global filter (Fig. S1†). Additional adjustment of the wavelength range was achieved using optical filters (Thorlabs). Experiments were performed under open-circuit conditions or in “zero resistance ammeter” (ZRA) mode with a Versastat 4 potentiostat (PAR/Ametek). Unstirred 0.5 M sulphuric acid was used as the electrolyte if not stated otherwise.

The residual layer thickness was determined by spectroscopic ellipsometry on a J.A. Woollam M-2000 ellipsometer with wavelengths in the range of 245 to 995 nm, at 75° angle of incidence. The thickness of the remaining TiO₂ layer was modelled with the Cauchy²⁰ relation for wavelengths >400 nm and with dispersions of Cody–Lorentz^{21,22} and Tauc–Lorentz.²³ In general, the results of fitting ellipsometry data to the various models were in close agreement. In all cases with 15 nm or more remaining, the residual film thickness can be determined to within ±0.3%. The homogeneity of the residual TiO₂ layer thickness in the area of the sample exposed to the electrolyte was verified by ellipsometric mapping, on a J. A. Woollam RC2 ellipsometer fitted with focusing probes, at 60°, 65° and 70°, in a wavelength range of 210 to 2500 nm. For the full experimental and fitted ellipsometry data see ESI, Fig. S4–S7.† Tabulated optical data for the silicon substrate and a SiO₂ interlayer were taken from Herzinger *et al.*²⁴

Results and discussion

Film properties

The as-grown TiO₂ layers on silicon were characterized by grazing incidence X-ray diffraction (GIXRD), Raman

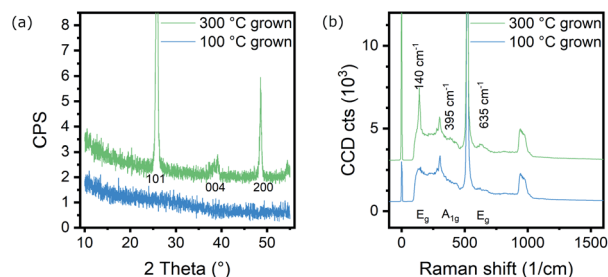


Fig. 1 (a) X-ray diffractogram and (b) Raman spectrum of TiO₂ films grown at 100 °C and 300 °C to yield amorphous or crystalline layers respectively. For the amorphous films, even the strongest signals indicating anatase at 25.3° and 140 cm⁻¹ cannot be distinguished from the noise or fluorescent background.

spectroscopy, atomic force microscopy (AFM) and spectroscopic ellipsometry (SE). Fig. 1 shows the diffractogram and Raman spectrum of the samples studied in this work, compared to a 300 °C grown film, highlighting the absence of any significant amount of crystalline TiO₂ in the 100 °C deposition. AFM topography (Fig. S2, S3 and Table S1†) shows sub-nm surface roughness for the films and is bare of the typical hillocks observed for (partially) crystalline coatings.^{25,26} Spectroscopic ellipsometry was used to determine the film thickness at 68.1 ± 0.5 nm with a roughness of approximately 2 nm. Modelling the SE data with a Tauc–Lorentz dispersion for the TiO₂ layer yields an optical band gap of 3.30–3.45 eV (details in ESI, Fig. S4–S6†), which is in agreement with the amorphous nature of the material.

Corrosion at open-circuit in the dark and under different illumination conditions

Fig. 2 shows the residual film thicknesses after exposing the samples to a strongly acidic electrolyte (0.5 M H₂SO₄) for 2–60 h and the respective average etch rates, both in the dark and under illumination with 100 mW cm⁻². It shows the steady decrease in film thickness over time, which is enhanced when the samples are illuminated. The average etch rates are 0.39–0.55 nm h⁻¹ in the dark and up to 2.1 nm h⁻¹ under illumination (Fig. 2b).

To study the impact of incident wavelength, optical filters were used for experiments of 15 h etching under illumination. Using a longpass UV-filter with a 400 nm cut-on (Thorlabs FGL400) and considering that the band gap of TiO₂ is >3 eV, exclusion of UV light should minimize photochemical or photoelectrochemical etching (any carriers generated by absorption in the degenerately doped silicon substrate are rapidly lost to recombination). The resulting etch rate was 1.4 nm h⁻¹, close to the rate at full illumination. Using other filters, *i.e.* an UV-vis bandpass filter (Thorlabs FGS900, 900 nm cut-off) and an UV-passing filter (Thorlabs FGUV, transmission 325–385 nm & 700–900 nm) showed etch rates of 1.0 and 0.7 nm h⁻¹, respectively, with no apparent influence of the chosen wavelength. Fig. 3a summarizes the average rates with several of the employed filters for experiments of 15 h etching duration at open-circuit (black frames).



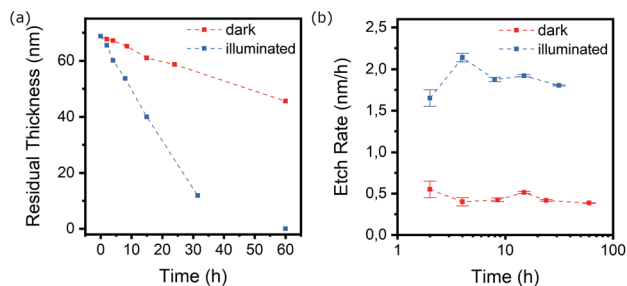


Fig. 2 (a) Residual thickness and (b) etch rate of TiO_2 in 0.5 M H_2SO_4 over time in the dark (red) and under 100 mW cm^{-2} illumination (blue).

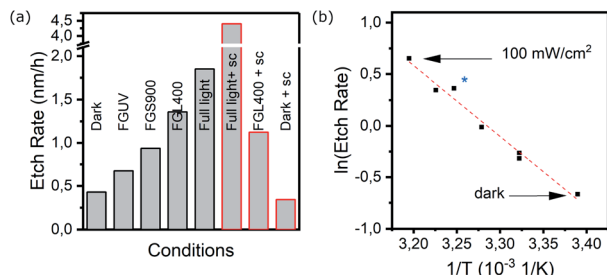


Fig. 3 (a) Average etch rate over 15 h under full or partial illumination in 0.5 M H_2SO_4 . FGL400, FGS900 and FGUV denote various filters to remove UV- and IR contributions from incoming light (L400/S900) or limit excitation to UV only (FGUV). "sc" indicates that the cell was wired directly to the Pt counter electrode to allow free transfer of photo-generated charge carriers. Due to higher etch rate, data for fully illuminated, short-circuited sample was determined after 4 hours of etching. (b) Arrhenius plot of TiO_2 etch rate based on warming up of the PEC cell under illumination using various filters. The blue star indicates heating up of the cell, sample and electrolyte externally but in the dark (see text for details). All temperatures in the graph were measured within the electrolyte, close to the sample. With a slope of $-6.873 \times 10^3 \text{ K}^{-1}$, resulting in $E_a = 57 \text{ kJ mol}^{-1}$.

A warming up of the PEC cell under constant illumination was noted and the electrolyte temperature in the cell was measured with a Pt-100 resistance thermometer, both in close proximity to the sample, with the sensor tip illuminated, and more remotely with the sensor not exposed to light. Without any filters, the measured temperatures rose to 38°C and 41°C with the sensor shadowed and exposed respectively (Fig. S8†). The surface temperature of the sample may even exceed those values but could not be determined directly.

The observed increased etch rates under illumination with and without various filters, indicated a temperature dependent etching rate, which was confirmed by performing experiments in the dark with a heated cell (approx. 38°C electrolyte). In this case, the observed rate of 1.5 nm h^{-1} was comparable to the previously described experiment with the UV filter, for which a similar temperature was measured.

To verify the influence of temperature increase and stabilization during the early stage of continuous illumination, the TiO_2 thickness was determined in a series of repeated 1 h intervals under illumination, showing an etch rate of 1.10 nm h^{-1} (Fig. S9†). A second series of measurements was performed at

intervals of 2.8 h, each time allowing the cell to approach its saturation temperature, resulting in an etch rate of 1.63 nm h^{-1} . As the reproducibility of the etch rates over 24 hours and 27 nm for the onset illumination conditions further shows, the slow initial corrosion rates are not linked to differences in the TiO_2 film surface related to deposition conditions.

From the respective electrolyte temperatures determined under (partial) illumination and the resulting etch rates, an Arrhenius plot was constructed (Fig. 3b) which yields an activation energy of 57 kJ mol^{-1} for the dissolution of amorphous TiO_2 , comparable to the values reported for anatase-rutile particles in HF-HCl mixtures.²⁷

Corrosion under short-circuit conditions

With the working electrode at open-circuit, electrochemical corrosion is expected to occur only if significant inhomogeneities are present. Such inhomogeneities, *e.g.* in the form of pinholes to the substrate or considerable phase differences, could present a difference in electrochemical potentials and divide the surface locally into microscopic anodic and cathodic regions that facilitate oxidation and reduction reactions. In our experiments, however, degradation proceeds isotropically, as evidenced by atomic force microscopy and spectroscopic ellipsometry mapping (Fig. S2 and S7†). On such a homogenous surfaces, the electrochemical anodic or cathodic corrosion would lead to a build-up of charges that result in an electrochemical potential that opposes further corrosion. Connecting the sample in short-circuit to a counter electrode or using a potentiostat in zero resistance mode, allows these charges to be extracted and the study of the influence of (photo-)electrochemical corrosion without applied bias.

Results from the additional short-circuit experiments are included in Fig. 3a and highlighted by red frames. Under dark conditions but with the TiO_2 electrode connected externally to the Pt counter electrode, the etch rate over 15 hours was largely unaffected and remained $<0.5 \text{ nm h}^{-1}$, while under full illumination, a corrosion of $>4.0 \text{ nm h}^{-1}$ was determined. Consequently, under short-circuit conditions and illumination, the etch process is accelerated more than 2-fold with the free flow of excess charge carriers to the counter electrode.

The importance of wavelength can herein be highlighted, as the use of the 400 nm cut-on filter during short-circuit experiments resulted in an etch rate of 1.2 nm h^{-1} , comparable to the open-circuit conditions, and no apparent enhancement from extracting excess charge.

Considering the mechanism of photoinduced corrosion, with light dependent generation of reactive holes in the valence band that provide potential either for water oxidation or oxidation of the semiconductor, it is apparent that the increase in film removal under full illumination is related to excitation by photons of UV wavelength and energy greater than the band gap of the amorphous TiO_2 . The anodic decomposition potential of TiO_2 is less positive than the potential at its valence band maximum (VBM), and more positive but close the redox potential of the oxygen evolution reaction (OER).^{28,29} With an intrinsically low activity towards water oxidation and immense



overpotentials for oxygen evolution, the kinetic stability against photocorrosion, as was pointed out by Gerischer,³⁰ and Bard,³¹ is overcome for these amorphous films.

Corrosion at different pH values

Studies for PEC absorber materials, catalysts and protection schemes are most commonly conducted in either very alkaline or very acidic electrolyte, due to the high conductivity of these solutions and often superior activity of the catalysts at the edges of the pH range.³²

In our experiments we have similarly observed significantly less degradation of the films at pH = 7 and even pH = 4, with a change in film thickness of <1 nm over 15 h of corrosion at open-circuit or 4 h allowing charge extraction. Instability in amorphous TiO₂ films has also been reported for alkaline environments.³³ In 1 M KOH or NaOH, the degradation of our samples proceeds inhomogeneously and is not easily determined through ellipsometry. It is uncertain at this point, if this is based on non-uniform degradation of the TiO₂ layer or related to a pinhole induced photocorrosion process of the underlying substrate, as proposed by Didden *et al.* for thin films of TiO₂ on CdS.¹¹

In the range of pH = 0.3 to pH = 4, we determined the corrosion rates at OCP in the dark and under illumination and in zero resistance ammeter mode under illumination. Fig. 4 shows the resulting rates and highlights the pH influence for all measurement conditions. In the dark, corrosion rates drop from 0.51 nm h⁻¹ at pH = 0.3 to 0.05 nm h⁻¹ at pH = 2–3 and from 1.92 to 0.13 nm h⁻¹ under illumination. With charge extraction, the corrosion rates remain higher by a factor of 2.5 to 4 compared to open-circuit but decrease rapidly with increasing pH value.

These findings agree well with Pourbaix diagrams that account for hydrated species of TiO₂ and allow for TiO²⁺ or TiO₂⁺ species at low pH.^{34,35} The common transitions here are the reduction to Ti³⁺ at potentials more favourable than proton reduction and oxidation towards a TiO₂²⁺ species approximately 700 mV more positive than the thermodynamic potential of

oxygen evolution. Similarly, Knauss *et al.* have shown the hydrolysis of TiO₂ even in rutile phase at elevated temperature to proceed through Ti(OH)₄ to Ti(OH)₃⁺ and Ti(OH)₂²⁺ in acidic media or to Ti(OH)₅⁻ in alkaline environment.³⁶

Short comparison to other TiO₂ thin films

The ALD growth of crystalline TiO₂ proceeds through an initial amorphous thin film until achieving a critical thickness upon which fully crystalline films are generally observed.^{37,38} In the early growth phase, *e.g.* up to tens of nm – dependent on growth parameters – the amorphous layer hosts the seeds of crystallites, which progressively grow as anatase throughout the film both by continuous epitaxial growth at the exposed surface and crystallization through the film, consuming the amorphous TiO₂.²⁶

The electrodes of the multi-crystalline films referenced for comparison in Fig. 1, that were grown at 300 °C and consist predominantly of anatase phase, exhibit no apparent corrosion even during extensive electrochemical testing. Other films, grown also at 300 °C but below the thickness threshold for full crystallisation show the degradation of the amorphous material only, while the embedded crystallites become successively more exposed³⁹ which is in agreement with findings by other groups regarding the greater intrinsic stability of crystalline grown films.⁸

The solubility of Ti generally increases at pH values further from the point of zero charge (PZC) but the results show that this parameter is not an adequate indicator for comparing different phases.⁴⁰ The PZC of amorphous TiO₂ is commonly found to be more acidic (4.0–4.6)^{41,42} than either that for anatase (5.2–6.2)^{41,43} or rutile (5.4–6.0)^{43,44} and to change during aging and crystallisation.⁴¹

Structurally, amorphous TiO₂ is more similar to anatase than rutile but the disorder in the chains of octahedra and the broad dispersion of bond lengths give rise to weak links in form of elongated Ti–O bonds with 1/2 to 1/3 of the bond-strength of the crystalline counterpart.⁴⁵ Stability is additionally decreased by the lower average coordination numbers (CN) of oxygen and titanium (avg. CN_{Ti} = 5.49) in the amorphous phase compared to the crystalline phases (CN_{Ti} = 6).⁴⁵

Sputtering profiles of XPS measurements (Fig. S10†) reveal low percentage impurity levels of carbon and nitrogen throughout the films originating from incompletely removed precursor ligands during the plasma-assisted deposition process. All films were exposed to atmosphere and show high contamination at the surface. The low temperature deposition at 100 °C results in higher levels of Nitrogen and Carbon contamination around 4 and 5 at%, in comparison to elevated deposition temperatures of 300 °C with impurity levels <1 at%.

The similar corrosion behaviour in 100 °C and 300 °C grown amorphous material points to a more dominant influence from the phase of TiO₂, compared to the incorporation of N or C. Photoelectrochemically, our electrodes containing amorphous TiO₂ commonly exhibit a reduction feature at 0.15 V *vs.* SHE (Fig. 5). This is only present after anodic polarization beyond a specific potential under illumination and has similar

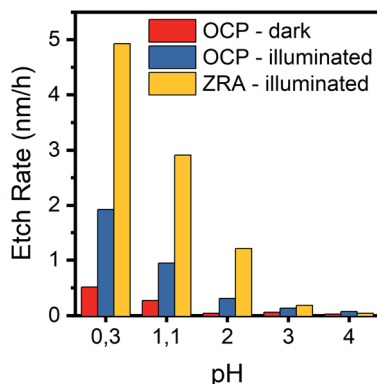


Fig. 4 Etch rate dependence on the pH value of electrolyte (H₂SO₄ + K₂SO₄, 0.5 M SO₄²⁻ total) at open-circuit conditions (OCP) and in zero resistance ammeter mode (ZRA). Data for OCP experiments all after 15 h, ZRA data for 4 or 8 h experiments due to higher etch rates.



characteristics to a peak observed by Bertoluzzi *et al.* for Fe_2O_3 .⁴⁶ Their study attributes these peaks to the forced surface oxidation and the formation of traps, with further arguments towards a slow detrapping mechanism. For the TiO_2 electrodes in this work, the appearance of reduction peaks coincides with the amorphous phase and instability of the material.

Preliminary insights from EIS

The study of electrochemical impedance spectroscopy (EIS) can shed further light on bias dependent capacitances and resistances in the electrochemical system. Models treating the charge transfer either directly from the bands or *via* surface states^{47,48} may help to identify bottlenecks, but were a poor approach to fitting the experimental data for our (partially) corroding electrodes. Instead, we focussed on simple equivalent circuits (Fig. 6a) for a blocking layer (dark) or single Randles circuit both containing one constant phase element (CPE) to account for inhomogeneity or mixed processes that cannot at this stage be fully unravelled.

The potential dependent analysis of the separate elements (Fig. 6b and c) reveals several features that can be linked to the CV scans (Fig. 5):

- (1) The pure capacitance $C(1)$ encountered in the model in the dark exhibits a dip at the exact potential of the reductive peak in the CV (0.15 V), or a local maximum at 0.25 V (Fig. 6b).
- (2) The CPE based capacitance and resistance sharply change in parallel with the onset of reductive current (<0.2 V).
- (3) Under illumination, the dominant capacitance (derived from the CPE), though generally an order of magnitude larger, has a minor peak at 0.15 V and a maximum of resistance at 0.25 V *vs.* SHE (Fig. 6c).

In line with any modelling of complex data, the analysis of electrochemical impedance spectroscopy offers numerous pitfalls and is susceptible to assumptions in selecting an appropriate electrochemical equivalent circuit. In order to justify a more comprehensive model, systematic variations of the layer studied and the overall electrochemical system are required to attribute physical significance to any selected element. In a comprehensive study currently ongoing, we expect to shed light on the contributing factors to the chemical and photoelectrochemical corrosion, as well

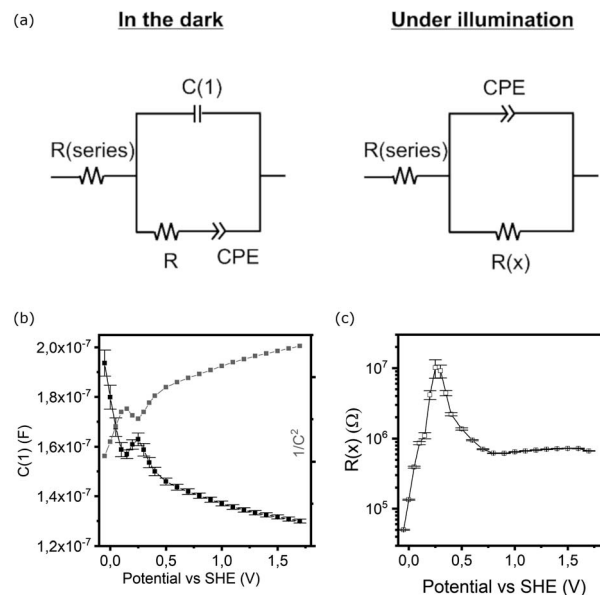


Fig. 6 (a) Electrochemical equivalent circuits used for modelling EIS data, (b) capacitive element in parallel ($C(1)$) in the dark and (c) main resistance element under illumination.

as a link to the intricate interplay of deposition parameters in film growth by ALD. This could help identify the optimal deposition pathway to stable protective coatings for photoanodes and cathodes, as well as provide means to identify weak points in electrochemically exposed layers.

Conclusions

The corrosion of ALD-grown, amorphous TiO_2 films in 0.5 M sulfuric acid ($\text{pH} = 0.3$) was studied by spectroscopic ellipsometry. At open-circuit potential the etch rate in the dark is 0.5 nm h^{-1} . Under 100 mW cm^{-2} illumination, the etch rate increases to 2.1 nm h^{-1} , which can be attributed to the temperature change of the photoelectrochemical cell under continuous illumination from room temperature to 41°C . By using cut-on and cut-off filters to remove different fractions of the spectrum (*e.g.* UV or IR), it could be shown that excitation wavelength is not responsible for an increase in etch rates but rather the different cell temperatures resulting from illumination. The corresponding activation energy of the TiO_2 dissolution is 57 kJ mol^{-1} .

Controlling the flow of excess generated charge carriers in short-circuit configuration, as opposed to open-circuit conditions, furthermore revealed a photoelectrochemical corrosion contribution dependent on UV excitation, increasing etch rates more than twofold. In agreement with recently published Pourbaix diagrams that account for hydrated TiO_2 , the degradation is strongly pH dependent and becomes negligible at $\text{pH} = 4$ in the time frames investigated. Consequently, in order to ensure long term stability, for PEC devices operating in acidic electrolyte, and incorporating amorphous TiO_2 coatings which are at least partially in contact with the electrolyte, the operational conditions should be limited to $\text{pH} > 4$.

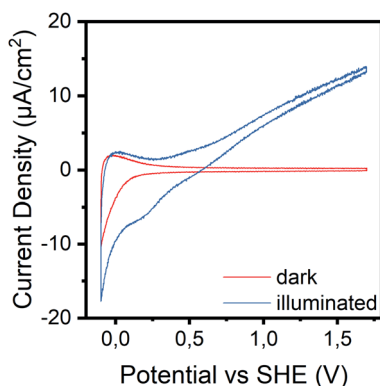


Fig. 5 Cyclic voltammogram (CV) of amorphous TiO_2 on $n^{++}\text{Si}$ in the dark and under 100 mW cm^{-2} illumination.



In comparison to the stable, crystalline counterpart anatase, amorphous TiO₂ shows a photoinduced reduction peak near 0.2 V vs. SHE that can be linked to features in the EIS analysis. Ongoing comprehensive studies in this direction may be helpful to identify the exact origin of the instability in TiO₂ films, their dependence on synthesis parameters and electrochemical contribution in the overall system.

Altogether, the results show that for PEC systems a study of the electrodes without added catalyst or enabled photoreaction can reveal underlying stability limitations.

Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript and declare no competing interests.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors are grateful to A. Schwartzberg, D. Olynick, and S. Cabrini for their insight and support for the preparation and coating of the samples by atomic layer deposition at The Molecular Foundry of the Lawrence Berkeley National Laboratory. The authors furthermore gratefully acknowledge the financial support by the Federal Ministry of Education and Research of Germany in the framework of the project "FocusH2" (no. 03SF0479A). Work at the Molecular Foundry was supported by the Office of Science, Office of Basic Energy Sciences, of the U.S. Department of Energy under contract no. DE-AC02-05CH11231.

Notes and references

- 1 S. Hu, N. S. Lewis, J. W. Ager, J. Yang, J. R. McKone and N. C. Strandwitz, *J. Phys. Chem. C*, 2015, **119**, 24201–24228.
- 2 Y. W. Chen, J. D. Prange, S. Dühnen, Y. Park, M. Gunji, C. E. D. Chidsey and P. C. McIntyre, *Nat. Mater.*, 2011, **10**, 539–544.
- 3 B. Seger, T. Pedersen, A. B. Laursen, P. C. K. Vesborg, O. Hansen and I. Chorkendorff, *J. Am. Chem. Soc.*, 2013, **135**, 1057–1064.
- 4 S. Hu, M. R. Shaner, J. A. Beardslee, M. Lichterman, B. S. Brunshwig and N. S. Lewis, *Science*, 2014, **344**, 1005–1009.
- 5 J. Qiu, G. Zeng, M.-A. Ha, B. Hou, M. Mecklenburg, H. Shi, A. N. Alexandrova and S. B. Cronin, *Chem. Mater.*, 2015, **27**, 7977–7981.
- 6 M. F. Lichterman, A. I. Carim, M. T. McDowell, S. Hu, H. B. Gray, B. S. Brunshwig and N. S. Lewis, *Energy Environ. Sci.*, 2014, **7**, 3334–3337.
- 7 L. Rovelli, S. D. Tilley and K. Sivula, *ACS Appl. Mater. Interfaces*, 2013, **5**, 8018–8024.
- 8 A. Paracchino, N. Mathews, T. Hisatomi, M. Stefl, S. David Tilley and M. Grätzel, *Energy Environ. Sci.*, 2012, **5**, 8673–8681.
- 9 B. Seger, S. David Tilley, T. Pedersen, P. C. K. Vesborg, O. Hansen, M. Grätzel and I. Chorkendorff, *J. Mater. Chem. A*, 2013, **1**, 15089–15094.
- 10 B. Seger, D. S. Tilley, T. Pedersen, P. C. K. Vesborg, O. Hansen, M. Grätzel and I. Chorkendorff, *RSC Adv.*, 2013, **3**, 25902–25907.
- 11 A. Didden, P. Hillebrand, B. Dam and R. van de Krol, *Int. J. Photoenergy*, 2015, **2015**, 1–8.
- 12 C. Ros, T. Andreu, M. D. Hernández-Alonso, G. Penelas-Pérez, J. Arbiol and J. R. Morante, *ACS Appl. Mater. Interfaces*, 2017, **9**, 17932–17941.
- 13 M. Hannula, H. Ali-Löytty, K. Lahtonen, E. Sarlin, J. Saari and M. Valden, *Chem. Mater.*, 2018, **30**, 1199–1208.
- 14 J. Ohkubo, Y. Hirose, E. Sakai, S. Nakao, T. Hitosugi and T. Hasegawa, *Jpn. J. Appl. Phys.*, 2011, **50**, 018002.
- 15 S. Okazaki, T. Ohhashi, S. Nakao, Y. Hirose, T. Hitosugi and T. Hasegawa, *Jpn. J. Appl. Phys.*, 2013, **52**, 098002.
- 16 S. M. Kramer, I. G. Gorichev, Yu. A. Lainer, I. V. Artamonova and M. V. Terekhova, *Russian Metallurgy*, 2014, **2014**, 704–707.
- 17 E. V. Shkol'nikov, *Russ. J. Phys. Chem. A*, 2016, **90**, 567–571.
- 18 F. Nandjou and S. Haussener, *J. Phys. D: Appl. Phys.*, 2017, **50**, 124002.
- 19 F. Nandjou and S. Haussener, *ChemSusChem*, 2019, **12**, 1984–1994.
- 20 E. Langereis, S. B. S. Heil, H. C. M. Knoop, W. Keuning, M. C. M. van de Sanden and W. M. M. Kessels, *J. Phys. D: Appl. Phys.*, 2009, **42**, 073001.
- 21 A. S. Ferlauto, G. M. Ferreira, J. M. Pearce, C. R. Wronski, R. W. Collins, X. Deng and G. Ganguly, *J. Appl. Phys.*, 2002, **92**, 2424–2436.
- 22 J. Price, P. Y. Hung, T. Rhoad, B. Foran and A. C. Diebold, *Appl. Phys. Lett.*, 2004, **85**, 1701–1703.
- 23 G. E. Jellison and F. A. Modine, *Appl. Phys. Lett.*, 1996, **69**, 371–373.
- 24 C. M. Herzinger, B. Johs, W. A. McGahan, J. A. Woollam and W. Paulson, *J. Appl. Phys.*, 1998, **83**, 3323–3336.
- 25 W.-J. Lee and M.-H. Hon, *J. Phys. Chem. C*, 2010, **114**, 6917–6921.
- 26 G. Luka, B. S. Witkowski, L. Wachnicki, M. Andrzejczuk, M. Lewandowska and M. Godlewski, *CrystEngComm*, 2013, **15**, 9949–9954.
- 27 E. Bright and D. W. Readey, *J. Am. Ceram. Soc.*, 1987, **70**, 900–906.
- 28 H. Gerischer, *J. Vac. Sci. Technol.*, 1978, **15**, 1422–1428.
- 29 S. Chen and L.-W. Wang, *Chem. Mater.*, 2012, **24**, 3659–3666.
- 30 H. Gerischer, *J. Electroanal. Chem. Interfacial Electrochem.*, 1977, **82**, 133–143.
- 31 A. J. Bard and M. S. Wrighton, *J. Electrochem. Soc.*, 1977, **124**, 1706.
- 32 Z. Chen, H. Dinh and E. Miller, *Photoelectrochemical Water Splitting: Standards, Experimental Methods, and Protocols*, Springer-Verlag, New York, 2013.



- 33 J. Zheng, Y. Lyu, R. Wang, C. Xie, H. Zhou, S. P. Jiang and S. Wang, *Nat. Commun.*, 2018, **9**, 1–10.
- 34 M. Pourbaix, *Atlas of electrochemical equilibria in aqueous solutions*, Nat'L Assoc. Of Corrosion, 1974.
- 35 P. Acevedo-Peña, J. Vazquez-Arenas, R. Cabrera-Sierra, L. Lartundo-Rojas and I. González, *J. Electrochem. Soc.*, 2013, **160**, C277–C284.
- 36 K. G. Knauss, M. J. Dibley, W. L. Bourcier and H. F. Shaw, *Appl. Geochem.*, 2001, **16**, 1115–1128.
- 37 D. R. G. Mitchell, D. J. Attard and G. Triani, *Thin Solid Films*, 2003, **441**, 85–95.
- 38 A. P. Alekhin, S. A. Gudkova, A. M. Markeev, A. S. Mitiaev, A. A. Sigarev and V. F. Toknova, *Appl. Surf. Sci.*, 2010, **257**, 186–191.
- 39 R. Raudsepp, H. Kriegel, J. Kollmann, T. Klassen and M. Schieda, manuscript in preparation.
- 40 M. P. Finnegan, H. Zhang and J. F. Banfield, *Chem. Mater.*, 2008, **20**, 3443–3449.
- 41 E. A. Barringer and H. K. Bowen, *Langmuir*, 1985, **1**, 414–420.
- 42 T. Sugimoto and X. Zhou, *J. Colloid Interface Sci.*, 2002, **252**, 347–353.
- 43 K. Bourikas, T. Hiemstra and W. H. Van Riemsdijk, *Langmuir*, 2001, **17**, 749–756.
- 44 M. L. Machesky, D. J. Wesolowski, D. A. Palmer and K. Ichiro-Hayashi, *J. Colloid Interface Sci.*, 1998, **200**, 298–309.
- 45 J. Mavračić, F. C. Mocanu, V. L. Deringer, G. Csányi and S. R. Elliott, *J. Phys. Chem. Lett.*, 2018, **9**, 2985–2990.
- 46 L. Bertoluzzi, L. Badia-Bou, F. Fabregat-Santiago, S. Gimenez and J. Bisquert, *J. Phys. Chem. Lett.*, 2013, **4**, 1334–1339.
- 47 B. Klahr, S. Gimenez, F. Fabregat-Santiago, T. Hamann and J. Bisquert, *J. Am. Chem. Soc.*, 2012, **134**, 4294–4302.
- 48 L. Bertoluzzi, P. Lopez-Varo, J. A. J. Tejada and J. Bisquert, *J. Mater. Chem. A*, 2016, **4**, 2873–2879.

