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A robust ALD-protected silicon-based hybrid photoelectrode for hydrogen evolution under aqueous conditions†

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Hydrogen production through direct sunlight-driven water splitting in photo-electrochemical cells (PECs) is a promising solution for energy sourcing. PECs need to fulfill three criteria: sustainability, cost-effectiveness and stability. Here we report an efficient and stable photocathode platform for H₂ evolution based on Earth-abundant elements. A p-type silicon surface was protected by atomic layer deposition (ALD) with a 15 nm TiO₂ layer, on top of which a 300 nm mesoporous TiO₂ layer was spin-coated. The cobalt diimine–dioxime molecular catalyst was covalently grafted onto TiO₂ through phosphonate anchors and an additional 0.2 nm ALD-TiO₂ layer was applied for stabilization. This assembly catalyzes water reduction into H₂ in phosphate buffer (pH 7) with an onset potential of +0.47 V vs. RHE. The resulting current density is -1.3 ± 0.1 mA cm⁻² at 0 V vs. RHE under AM 1.5 solar irradiation, corresponding to a turnover number of 260 per hour of operation and a turnover frequency of 0.071 s⁻¹.

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Introduction

The production of solar fuels is one of the most promising strategies to shift towards sustainable energy sourcing as this is the only solution allowing massive amounts of renewable energy to be stored in a durable way. In that context, photo-electrochemical water splitting produces hydrogen (H₂) as a suitable energy vector.¹ Indeed, the mature fuel-cell technology allows its energy content to be turned back into electrical power on demand, with a good efficiency and in a closed cycle.^{2,3} The major challenge for the development of photoelectrochemical cells (PECs) lies in the cost effective fabrication of photo-electrodes with high solar energy conversion efficiencies and high durability under normal utilization.⁴ Fulfilling these requirements raises important scientific questions regarding mainly the elaboration and the combination of the best performing materials able to harvest light and catalyze H₂ evolution.

Silicon is highly abundant and can be cost-effectively processed for the industrial construction of photovoltaic panels into crystalline silicon, which has a conduction band (CB) energy level of ~ -0.5 V vs. NHE (pH = 0) and a narrow band-gap of 1.12 eV.⁵ These features make silicon the material of choice among other semiconductors^{6–9} for the fabrication of H₂-evolving photoelectrodes able to harvest solar irradiation over the whole visible spectrum.^{10,11} However, the major drawback of using silicon as a photoelectrode material is its inherent instability under aqueous or aerobic conditions. This issue can nevertheless be overcome by appropriate surface passivation techniques such as pinhole-free oxide coatings achieved by vacuum deposition,¹² sputtering¹³ or atomic layer deposition (ALD) techniques.¹⁴

Even though a few materials based on Earth-abundant elements already compete with platinum for H₂ evolution over a wide range of pH values,¹⁵ the best catalyst does not necessarily yield the best photoelectrode when it is interfaced with a visible-light-harvesting semiconductor.¹⁶ Indeed, it has been shown that the chemical potential of the catalyst needs to electrochemically equilibrate with the quasi Fermi level of electrons in the irradiated semiconductor,¹⁷ which strongly depends on the way the connection between the semiconductor and the catalyst is achieved. A ground-breaking study has set the use of porous catalyst materials that can adapt their potential through charging/discharging ions from the electrolyte as a way to achieve good efficiency.¹⁸ Such a feature is present in electrodeposited amorphous catalytic materials but it was hypothesized that the same feature can be reproduced in hybrid

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systems¹⁹ through the grafting of molecular catalysts onto a porous conducting transparent support, as a mesoscopic film of TiO₂.^{20,21} Such catalysts could be cobaloximes and diimine-dioxime cobalt complexes since their immobilization on electrode supports yields hybrid materials efficiently producing H₂ from aqueous electrolytes.^{6,10,22–25} Some planar and nano-structured silicon photocathodes interfaced with metal or metal sulphide catalysts can produce current densities > 5 mA cm^{−2} (Table S1†). In contrast, for molecular catalysts/hydrogenase enzymes, only the use of high performance semiconductors such as p-GaP and p-InGaP₂ produces current densities > 1 mA cm^{−2} in photocathode constructs (Table S2†).

We report here a strategy to fabricate a molecular-engineered silicon photocathode (Fig. 1) alleviating the previously observed drawbacks, producing current densities > 1 mA cm^{−2} and achieving sustained photoelectrochemical H₂ evolution in neutral pH under AM 1.5 solar irradiation. To that aim, we combined for the first time ALD protection of a p-Si photocathode and the use of a mesoporous catalytic layer based on TiO₂ interfaced with a molecular diimine dioxime cobalt catalyst.

Results and discussion

The native surface oxide layer of commercial boron-doped p-type silicon (100) wafers (725 μm-thick, resistivity 1–50 Ω cm) was removed using hydrofluoric acid (HF) (see the Experimental part for the detailed etching process). Then the p-Si electrodes were rapidly transferred to the chamber of an ALD system. The electrodes were coated with a first layer of TiO₂ through 710 ALD cycles performed in continuous mode at 255 °C. Fig. 2A shows the top view scanning electron microscopy (SEM) image of this p-Si|ALD-TiO₂ electrode, revealing a TiO₂ layer for the protection of p-Si against corrosion in aqueous electrolytes. The thickness of the ALD-TiO₂ layer measured by ellipsometry was ~15 nm. A second layer of mesoporous TiO₂ was then spin-coated (SC) on top of the first layer to increase the surface area of the electrode and thus afford higher catalyst loading. Fig. 2B and C show the top view and cross-sectional SEM images of p-Si|ALD-TiO₂|SC-TiO₂ electrodes, revealing a ~300 nm thickness for the SC-TiO₂ layer. We selected the diimine-dioxime cobalt complex (Co_{C11P}) derivative containing a pendant phosphonic acid anchor as the H₂-evolving molecular catalyst.²⁶ Soaking the

p-Si|ALD-TiO₂|SC-TiO₂ electrodes in a methanolic solution of Co_{C11P} results in the straightforward immobilization of the complex. The grafting occurs through the formation of a mixed organic-inorganic phosphonate linkage,²⁷ as evidenced by electrochemistry, X-ray photoelectron spectroscopy (XPS) and infrared spectroscopy (see below). To further stabilize the catalyst anchoring against hydrolysis during long term operation in aqueous electrolytes, we specifically undertook a second ALD process, inspired by the “Mummy” strategy.²⁸ In this process, an ultra-thin overlayer of TiO₂ is deposited at the mesoporous Co_{C11P}-modified SC-TiO₂ interface to shield the catalyst-surface linkage.²⁸ We thus performed 10 ALD cycles in a 5 s exposure mode at 150 °C. The low deposition temperature was used to avoid potential degradation of the molecular structure of the catalyst while the selected exposure mode ensured that the ~0.2 nm thick overlayer is uniformly deposited within the whole mesoporous layer of Co_{C11P}-modified TiO₂.

Fig. 3 shows the XPS spectra of the Co 2p core region of p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P} (black trace) and p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ (red trace) electrodes. The peaks observed at 795.4 and 780.2 eV correspond to Co 2p_{1/2} and 2p_{3/2} core levels, respectively. The binding energies are similar to those previously reported for the same catalytic moiety bound to carbon nanotubes, NiO or ITO.^{22,25,26} The fact that the binding energies of the Co 2p peaks are not modified upon coating strongly suggests that this overlayer did not degrade the molecular structure of the Co_{C11P} catalyst in p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrodes.

A Fourier transform infrared (FTIR) spectrum was measured on the Co_{C11P}-modified TiO₂ material scratched from a p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrode and sampled as a KBr pellet. Fig. 4 compares the FTIR spectrum of this material (red trace) with that of the Co_{C11P} molecular catalyst (black trace). Signals corresponding to the stretching frequencies of the oxime, imine and C–C bond in the imine-oxime moieties are found almost unchanged in both spectra at 1140/1146, 1519/1514 and 1251 cm^{−1}, respectively (assignments made by comparison with the related cobaloxime compounds following ref. 29). The broad peak at ~1625 cm^{−1} observed for the p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrode (red trace) corresponds to the bending vibration band of the hydrogen-bonded adsorbed water on unmodified TiO₂ (blue

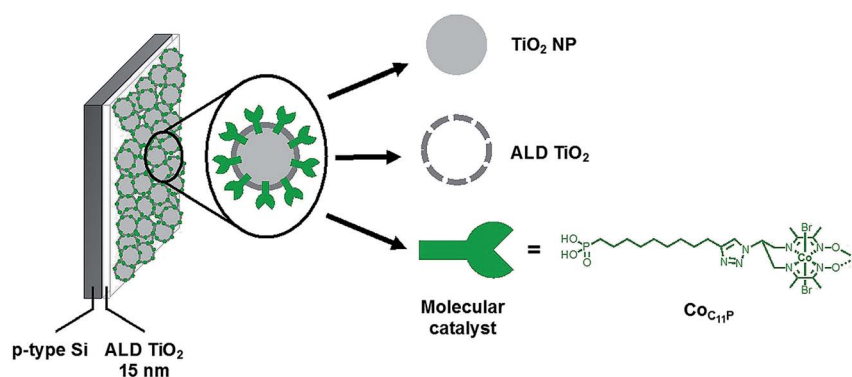


Fig. 1 Molecular-engineered silicon photocathode for H₂-evolution.



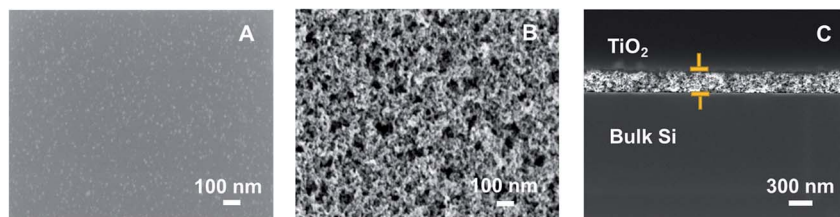


Fig. 2 SEM images of TiO₂ layers on p-Si: ALD-TiO₂ (top view) (A) and ALD-TiO₂/SC-TiO₂ top view (B) and cross-sectional view (C).

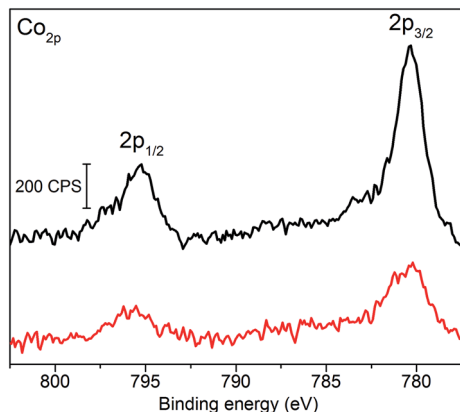


Fig. 3 X-ray photoelectron spectra of the Co 2p core region of the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P electrode (black trace) and p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode (red trace).

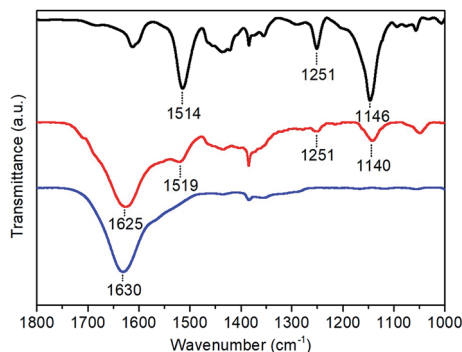


Fig. 4 FTIR spectra of the CoC₁₁P molecular catalyst as synthesized (black trace), p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode (red trace) and unmodified TiO₂ (blue trace).

trace, 1630 cm⁻¹).^{30,31} This analysis further confirmed that the final 10 ALD cycles for TiO₂ deposition at 150 °C did not degrade the molecular structure of the CoC₁₁P catalyst.

CoC₁₁P loading on the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode was quantified by inductively coupled plasma atomic emission spectrometry (ICP-AES) after digestion of the ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ layer in sulfuric acid. ICP-AES analysis indicated a Co loading of 67.4 ± 0.42 nmol cm⁻². We, however, note that only a fraction of the grafted catalysts may be electrochemically addressable, as reported previously for similar catalyst-modified mesoporous conducting metal oxide electrodes.^{9,26}

The photo-electrocatalytic properties of the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode for H₂ evolution were assessed in degassed 1 M phosphate buffer (pH 7) under solar AM 1.5 irradiation in a three-electrode system. Negligible current densities were observed during linear sweep voltammetry (LSV, 10 mV s⁻¹) under dark conditions for all the fabricated photocathodes (Fig. 5). Under illumination (1 sun AM 1.5), the LSV of the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode (Fig. 5, red trace) displays a current density of -1.25 mA cm⁻² at 0 V vs. RHE and an onset photocurrent potential of +0.47 V vs. RHE (fill factor of 0.17). In comparison, the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P electrode without the ALD-TiO₂ overlayer displays much lower performances, with a threefold lower (-0.5 mA cm⁻²) current density at 0 V vs. RHE and an onset photocurrent potential of +0.09 V vs. RHE (Fig. 5, black trace). In both cases, the photovoltage provided at the p-Si/TiO₂ results in catalysis, occurring from potentials significantly more positive than the redox potential of the Co^{II}/Co^I couple observed at ~0 V vs. RHE for CoC₁₁P-modified TiO₂ electrodes deposited onto FTO (see Fig. S1†). The higher current density displayed by the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode compared to the same construction incorporating decylphosphonic acid C₁₀P, as a surrogate reproducing surface modification through phosphonate binding, clearly evidences the role of CoC₁₁P in catalyzing H₂ evolution (Fig. S2†). The current density displayed by the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ photocathode is one order of magnitude higher than that of a similar hybrid

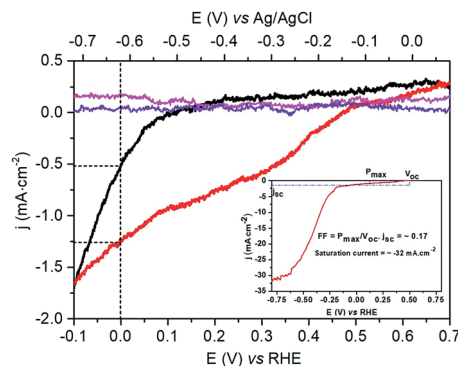


Fig. 5 LSV (10 mV s⁻¹) of the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P electrode (magenta, resp. black trace) and p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode (violet, resp. red trace) in 1 M phosphate buffer (pH 7) under dark conditions, respectively, one sun irradiation. The inset shows the p-Si|ALD-TiO₂|SC-TiO₂|CoC₁₁P|ALD-TiO₂ electrode (red trace) at more negative potentials and the calculated fill factor (FF).



construction without ALD-TiO₂ layers (p-Si|mesoTiO₂|NiP).¹⁰ At the same time the onset potential is shifted 350 mV more positive. The significantly improved performance of the electrode functionalized with the 10 cycle ALD-TiO₂ overlayer confirms that such an overlayer enables the sequestering of the Co_{C11P} molecular catalyst within the mesoporous TiO₂ layer where it can mediate H₂ evolution.

The superior behavior fostered by the ALD-TiO₂ overlayer was further supported by chronoamperometric measurements performed for 1 hour at 0 V vs. RHE under similar conditions (Fig. 6).³² An initial decay of current, also observed in catalyst-free electrodes (Fig. S3†), might originate from the charging behavior at the silicon/TiO₂ interface and/or within the n-type TiO₂ semiconductor caused by the sudden illumination. Then, both electrodes exhibited stable current densities of the same magnitude as that measured during LSV experiments, *i.e.* −0.5 and −1.3 mA cm^{−2} for p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P} and the overcoated p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrodes, respectively (Fig. 6, black and red traces, respectively). The chronoamperometric trace of the p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrode shows staggered variations due to the intermittent release of H₂ bubbles from the electrode surface. Gas chromatography analysis quantified the production of 17.3 μmol cm^{−2} of H₂, which evolved concomitantly with the passage of 3.96 C cm^{−2} through the p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrode, yielding a faradaic efficiency of 84%. With these data and the catalyst loading determined from ICP-AES, a turnover number (TON) of 260 and a turnover frequency (TOF) of 0.071 s^{−1} were calculated.

We note that higher current densities are observed at more negative potentials (see the inset in Fig. 5) with a second photoinduced process starting at −0.25 V vs. RHE. This second process likely corresponds to H₂ evolution directly mediated at the surface of TiO₂ as it is also observed for the p-Si|ALD-TiO₂|SC-TiO₂ electrode at higher negative potential (Fig. S4†). Possibly the construction described here does not allow for sufficient catalyst loading (based on the intrinsic turnover frequency of each molecular catalyst) to sustain such high

current densities. Alternatively, the electronic coupling between the Si wafer and the TiO₂ layers may not be optimal to allow the catalyst layer to equilibrate with the conduction band of Si.

We also investigated the behaviour of the p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrodes in 0.1 M NaOH (pH 13) (Fig. S5†). From LSV measurements, the current density and onset photocurrent potential were measured as −1 mA cm^{−2} at 0 V vs. RHE and +0.53 V vs. RHE under solar AM 1.5 illumination, respectively. These figures indicate good H₂-evolution properties of the Co_{C11P} catalyst in basic aqueous electrolyte, in line with Turner's report on the catalytic activity of cobaloximes under similar conditions.⁶ Current densities similar to those observed in LSV were sustained during chronoamperometric measurements (0 V vs. RHE) at the p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrode in 0.1 M NaOH (pH 13) for a period of one hour under AM 1.5 solar irradiation (Fig. S6†), yielding a faradaic efficiency of 83%. TON and TOF were calculated to be 215 and 0.06 s^{−1}, respectively, based on the catalytic loading determined through ICP-AES. XPS analysis of the Co 2p core region (Fig. S7†) carried out at the end of this chronoamperometry experiment supports the presence of the retained Co_{C11P} molecular catalyst on the p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}|ALD-TiO₂ electrode, with a Co 2p_{3/2} peak at 780.2 eV, similar to that in Fig. 3, whereas alteration of the Co complex would result in a shoulder peak around 782 eV.^{6,33} The noisiness of this feature indicates that the complex is present at the very surface of the mesoporous TiO₂ layer with a decreased concentration compared to the as-prepared sample. By contrast, a system based on a similar diimine-dioxime catalyst attached on TiO₂ shows full disappearance of the XPS Co 2p_{3/2} signature after activity.¹⁰ Retention of the Co 2p_{3/2} XPS binding energy after activity is another indication that the ALD-strategy is effective in stabilizing the grafting of the molecular catalyst during turn-over. These data show for the first time the versatility and stability of the Co_{C11P} molecular catalyst and phosphonate anchor, respectively, which is operative over a wide range of pH values, paralleling a report on cobaloximes with carboxylate anchors by Turner and coworkers.⁶ Furthermore, working of photocathodes in basic aqueous electrolytes is of great interest since that would help to tether them with TiO₂-based photoanodes in a tandem cell configuration for unbiased water splitting.

In both cases, relatively high photocurrent densities measured at 0 V vs. RHE result from the combination of an ALD-coated compact TiO₂ underlayer protecting p-Si from corrosion, a mesoporous TiO₂ layer allowing for high catalytic loading, and a second ALD-coated compact ultrathin TiO₂ overlayer stabilizing the anchorage of the molecular catalyst onto the mesoporous layer. Our control experiments (Fig. S3†) demonstrate that all three components are crucial to achieve efficiency and stability: (i) the presence of a thin TiO₂ protective layer allows generation of photocurrent in the mA cm^{−2} range for photoelectrodes stored in ambient atmosphere for days, (ii) the SC mesoporous TiO₂ allows a 10-fold increase of the current density compared to a flat ALD-TiO₂ surface (compare red and orange curves in Fig. S3b†), and (iii) the presence of an ALD-TiO₂ overlayer results in a sustained photocurrent at its

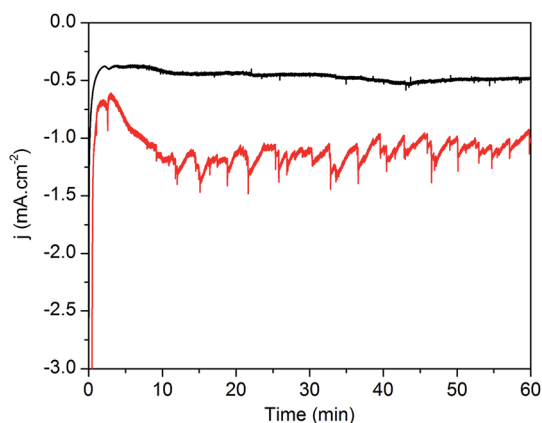


Fig. 6 Chronoamperometric profiles of the p-Si|ALD-TiO₂|SC-TiO₂-|Co_{C11P} electrode (black trace) and p-Si|ALD-TiO₂|SC-TiO₂|Co_{C11P}-|ALD-TiO₂ electrode (red trace) in 1 M phosphate buffer (pH 7) recorded at 0 V vs. RHE under one sun irradiation.



nominal value measured during LSV. This result is particularly remarkable since the non-stabilized grafting of a similar diimine–dioxime cobalt catalyst led to rapid leaching of the catalyst from the mesoporous TiO_2 layer (30 min, <2 TONs achieved),¹⁰ while the ALD-coated TiO_2 overlayer of this study allowed a sustained TOF (260 TONs achieved within 1 h with stable current density) for H_2 evolution by the molecular catalyst. With such an optimized platform we offer the possibility of using cost-effective silicon wafers for the preparation of hybrid photocathodes displaying performances previously restricted to expensive III–V semiconductors such as $\text{GaP}^{7,8}$ or GaInP_2 .⁶

Conclusion

We successfully designed a stable photocathode platform that unlocks the exploitation of cost-effective and technology-ready silicon wafers for the construction of robust photocathodes exclusively based on Earth-abundant elements, as an asset for sustainability. ALD and sol–gel coating techniques, which we used in our strategy, are currently deployed for various industrial applications, en route for the future scaling up of the production of such photoelectrodes. The assets of this photocathode architecture are resistance to corrosion, high catalyst loading and retention of the molecular structure of the catalyst on the electrode surface. These assets result in a platform suitable to anchor various molecular catalysts onto inorganic semiconductors in future research on (photoelectro)catalysis. Reaching the current density displayed by high-performing semiconductors by improving the loading densities in molecular catalysts at hybrid electrodes is currently being investigated in our laboratory and could benefit from the hierarchical inverse opal TiO_2 structures recently developed for photoelectrochemical biological systems.³⁴

Experimental

Chemicals

Hydrofluoric acid (HF, 48%), solvents and starting materials were purchased from Sigma Aldrich. Ultrapure nitric acid (65%) for ICP-AES analysis was purchased from Chem-Lab. Co_{C11P} was prepared as previously described.²⁶ Decylphosphonic acid ($\text{C}_{10\text{P}}$) was purchased from Sigma-Aldrich.

ALD of TiO_2 on the silicon surface (p-Si|ALD- TiO_2)

Boron doped p-type silicon with a resistivity of 1–50 ohm cm, orientation 100, thickness 725 μm and diameter 200 mm was bought from MEMC. The silicon was cut into 0.5×0.5 cm pieces and etched with 1 : 1 HF : ethanol mixture for 10 min. It was then stored in an argon degassed heptane solution and dried under argon before transferring into the ALD chamber. TiO_2 was deposited using an ALD system (Savannah 100 ALD system, Cambridge Nano Tech Inc.) with a precursor temperature at 95 °C, pump line temperature at 150 °C and sample chamber temperature at 255 °C in continuous mode for 710 cycles with a nitrogen flow of 5 sccm. The precursors used for TiO_2 deposition were titanium(IV) isopropoxide (TTIP) purchased from

Strem Chemicals Inc. and water as sources for titanium and oxygen, respectively. A cycle typically comprised a 0.5 s TTIP pulse and a 0.02 s water pulse with a 5 s purging time in between.

SC of mesoporous TiO_2 on the ALD- TiO_2 deposited silicon surface (p-Si|ALD- TiO_2 |SC- TiO_2)

A suspension of TiO_2 anatase nanoparticles of size 15–20 nm (Ti-Nanoxide T300/SC, Solaronix) was spin-coated on ALD- TiO_2 deposited on silicon (polished side). The spin-coating was carried out at 5000 rpm for 30 s, with an acceleration of 2000 rpm s^{-1} . The coated electrodes were then thermally treated at 475 °C for 30 min in a tubular furnace under an argon flow.

Immobilization of the Co_{C11P} molecular catalyst and $\text{C}_{10\text{P}}$ molecular anchor on the TiO_2 coated silicon surface (p-Si|ALD- TiO_2 |SC- TiO_2 | Co_{C11P} and p-Si|ALD- TiO_2 |SC- TiO_2 | $\text{C}_{10\text{P}}$)

A saturated solution of Co_{C11P} molecular catalyst²⁶ was prepared with methanol. Silicon substrates coated with ALD- TiO_2 and mesoporous SC- TiO_2 were soaked overnight in the catalyst solution. The substrates were then washed with methanol before further modification. A similar procedure was followed to sensitize the electrode with the $\text{C}_{10\text{P}}$ molecular anchors.

ALD- TiO_2 deposition on the Co_{C11P} molecular catalyst immobilized TiO_2 coated silicon surface (p-Si|ALD- TiO_2 |SC- TiO_2 | Co_{C11P} |ALD- TiO_2)

Co_{C11P} -functionalized TiO_2 -modified silicon electrodes were coated with 10 cycles of ALD- TiO_2 . ALD was performed in exposure mode with an exposure time of 5 s. During the exposure, pumping on the chamber is stopped in order to allow time for the precursors to diffuse in the mesoporous TiO_2 layer. Except for the temperature of the deposition chamber which was decreased from 255 °C to 150 °C to avoid degradation of the catalysts, the other parameters (temperature, nitrogen flow, pulse duration and purging time) were kept the same as in continuous mode.

Preparation of TiO_2 - Co_{C11P} electrodes

Screen-printed TiO_2 substrates were purchased from Solaronix. The electrodes were sintered under atmospheric conditions in a Harry Gestigkeit flat furnace operated with a Dettlef Gestigkeit-Programmer PR5 control board programmed with the following temperature ramp: R.T. to 450 °C (30 min), 30 min at 450 °C. The thickness of the TiO_2 layer is ca. 600 nm. The substrates were cooled to ca. 90 °C and introduced in PTFE boxes containing 10 mL of a 0.5 mM methanolic solution with Co_{C11P} (5 μmol). The boxes were closed and the content was stirred for 24 h on an orbital stirring plate. The electrodes were removed from the solution, washed by dipping them in pristine methanol for 10 min under orbital stirring and dried with N_2 , to yield the decorated $\text{TiO}_2/\text{Co}_{\text{C11P}}$ electrodes.

Instrumental characterization techniques

A Zeiss Ultra+ instrument was used for SEM imaging of ALD- TiO_2 and mesoporous SC- TiO_2 in In-Lens mode at accelerating voltages of 20 kV and 2 kV, respectively.



An ellipsometer (M 2000 model) from J. A. Woollam Co., In. was used to measure the thickness of the ALD-TiO₂ substrate, using the CompleteEASE software (V4.72). It was used in the “off-sample” configuration with an incident angle of attack of 75° with respect to the film plane normal, and the light wavelength was scanned between 200 nm and 1700 nm. The obtained data were fitted with the CompleteEASE software (V4.72). A typical example of the recorded spectrum and its fit are displayed in Fig. S8.†

FTIR spectra were recorded on a Perkin-Elmer Spectrum 100 spectrometer. The spectra were collected over 64 scans, with a resolution of 1 cm⁻¹ and background corrected with an ambient atmosphere spectrum. Spectra of the samples were recorded and analyzed using SPECTRUM version 10.5.2 software, in the range of 1000–2000 cm⁻¹. The powdered samples were mixed in KBr and pressed to form pellets. Typically, the pellets were made of the scratched overlayer of the electrode (4 cm²) mixed with 100 mg of KBr.

XPS analyses were carried out with a Versa Probe II spectrometer (ULVAC-PHI) equipped with a monochromated Al K α source ($h\nu = 1486.6$ eV). The core level peaks were recorded with a constant pass energy of 23.3 eV. The XPS spectra were fitted with CasaXPS 2.3.15 software using Shirley background and a combination of Gaussian (70%) and Lorentzian (30%) distributions. Binding energies are referenced with respect to the adventitious carbon (C 1s BE = 284.6 eV).

ICP-AES samples were analyzed using a Shimadzu ICPE-9000 instrument with a mini plasma torch in axial reading mode. The sample preparation involves the digestion of the p-Si|ALD-TiO₂-[SC-TiO₂][Co_{C11P}]ALD-TiO₂ electrode (22.5 cm² area) in 10 mL of 98% sulfuric acid at 60 °C for one hour under sonication. The solution was then diluted to 30 mL with 1% ultrapure nitric acid solution prepared with MilliQ water and sonicated for 5 hours. Further, 1.67 mL of this solution was diluted to 10 mL with 1% ultrapure nitric acid prepared with MilliQ water. This final solution (10 mL) was analyzed for cobalt concentration by ICP-AES. The samples were prepared in triplicate.

Photocurrent and microGC measurements

Electrochemical analysis was performed using a BioLogic SP300 bipotentiostat and a representative three-electrode setup. A platinum wire was used as the auxiliary electrode and Ag/AgCl, KCl (3 M) (denoted as Ag/AgCl) as the reference electrode bridged to the electrolyte through a Vycor frit. The unpolished side of silicon was cotton swabbed with an In–Ga eutectic mixture and superposed with a copper plate. The electrical connection was made through the copper plate. Photoelectrodes were front-illuminated with a 300 W ozone-free xenon lamp (Newport) operated at 280 W coupled to an AM 1.5 filter. Irradiance at the substrate surface was measured to be 100 mW cm⁻² using a Newport PM1918-R power meter.

Calibration of the reference electrode was realized with [K₄Fe(CN)₆] in 0.1 M potassium phosphate buffer at pH 7 (ref. 35) and conversion of potentials against the normal hydrogen electrode (NHE) and the reversible hydrogen electrode (RHE) was performed as previously described.³⁶

Fill factor calculation:³⁷

$$FF = \frac{P_{\max}}{V_{\text{oc}}j_{\text{sc}}}$$

where P_{\max} is the product of voltage (V) and current (A) at maximum power, V_{oc} (V) is the open circuit voltage and j_{sc} (A) is the short circuit current. We normalized the voltage (V) vs. RHE.

A MicroGC S3000 (SRA Instruments) with a diamond LV Ms5A 14m module was operated via a Soprane chrome interface. Throughout the chronoamperometric run, argon gas was purged into the electrolyte at a flow rate of 5 mL min⁻¹ controlled with a mass flow controller, and the outlet gas mixture was analyzed using a microGC.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 M. G. Walter, E. L. Warren, J. R. McKone, S. W. Boettcher, Q. Mi, E. A. Santori and N. S. Lewis, *Chem. Rev.*, 2010, **110**, 6446–6473.
- 2 N. Armaroli and V. Balzani, *ChemSusChem*, 2011, **4**, 21–36.
- 3 N. Armaroli and V. Balzani, *Energy Environ. Sci.*, 2011, **4**, 3193–3222.
- 4 J. R. McKone, N. S. Lewis and H. B. Gray, *Chem. Mater.*, 2014, **26**, 407–414.
- 5 D. Zhang, J. Shi, W. Zi, P. Wang and S. Liu, *ChemSusChem*, 2017, **10**, 4324–4341.
- 6 J. Gu, Y. Yan, J. L. Young, K. X. Steirer, N. R. Neale and J. A. Turner, *Nat. Mater.*, 2015, **15**, 456.
- 7 A. M. Beiler, D. Khusnutdinova, S. I. Jacob and G. F. Moore, *Ind. Eng. Chem. Res.*, 2016, **55**, 5306–5314.
- 8 D. Khusnutdinova, A. M. Beiler, B. L. Wadsworth, S. I. Jacob and G. F. Moore, *Chem. Sci.*, 2017, **8**, 253–259.
- 9 B. L. Wadsworth, A. M. Beiler, D. Khusnutdinova, S. I. Jacob and G. F. Moore, *ACS Catal.*, 2016, **6**, 8048–8057.
- 10 J. J. Leung, J. Warnan, D. H. Nam, J. Z. Zhang, J. Willkomm and E. Reisner, *Chem. Sci.*, 2017, **8**, 5172–5180.
- 11 S. Chandrasekaran, T. Nann and N. H. Voelcker, *Nano Energy*, 2015, **17**, 308–322.
- 12 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.
- 13 B. Seger, T. Pedersen, A. B. Laursen, P. C. K. Vesborg, O. Hansen and I. Chorkendorff, *J. Am. Chem. Soc.*, 2013, **135**, 1057–1064.
- 14 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.



- 15 C. C. L. McCrory, S. Jung, I. M. Ferrer, S. M. Chatman, J. C. Peters and T. F. Jaramillo, *J. Am. Chem. Soc.*, 2015, **137**, 4347–4357.
- 16 M. R. Nellist, F. A. Laskowski, F. Lin, T. J. Mills and S. W. Boettcher, *Acc. Chem. Res.*, 2016, **49**, 733–740.
- 17 H. Gerischer, *J. Electroanal. Chem. Interfacial Electrochem.*, 1977, **82**, 133–143.
- 18 F. D. Lin and S. W. Boettcher, *Nat. Mater.*, 2014, **13**, 81–86.
- 19 S. Hennessey and P. Farras, *Chem. Commun.*, 2018, **54**, 6662–6680.
- 20 N. Coutard, N. Kaeffer and V. Artero, *Chem. Commun.*, 2016, **52**, 13728–13748.
- 21 N. Queyriaux, N. Kaeffer, A. Morozan, M. Chavarot-Kerlidou and V. Artero, *J. Photochem. Photobiol., C*, 2015, **25**, 90–105.
- 22 E. S. Andreiadis, P. A. Jacques, P. D. Tran, A. Leyris, M. Chavarot-Kerlidou, B. Joussetme, M. Matheron, J. Pecaut, S. Palacin, M. Fontecave and V. Artero, *Nat. Chem.*, 2013, **5**, 48–53.
- 23 N. Kaeffer, M. Chavarot-Kerlidou and V. Artero, *Acc. Chem. Res.*, 2015, **48**, 1286–1295.
- 24 S. Donck, J. Fize, E. Gravel, E. Doris and V. Artero, *Chem. Commun.*, 2016, **52**, 11783–11786.
- 25 N. M. Muresan, J. Willkomm, D. Mersch, Y. Vaynzof and E. Reisner, *Angew. Chem., Int. Ed.*, 2012, **51**, 12749–12753.
- 26 N. Kaeffer, C. D. Windle, R. Brisse, C. Gablin, D. Léonard, B. Joussetme, M. Chavarot-Kerlidou and V. Artero, *Chem. Sci.*, 2018, **9**, 6721–6738.
- 27 S. P. Pujari, L. Scheres, A. T. M. Marcelis and H. Zuilhof, *Angew. Chem., Int. Ed.*, 2014, **53**, 6322–6356.
- 28 A. M. Lapidès, B. D. Sherman, M. K. Brennaman, C. J. Dares, K. R. Skinner, J. L. Templeton and T. J. Meyer, *Chem. Sci.*, 2015, **6**, 6398–6406.
- 29 P. E. Rutherford and D. A. Thornton, *Spectrochim. Acta, Part A*, 1979, **35**, 711–714.
- 30 H. Jensen, A. Soloviev, Z. Li and E. G. Søgaard, *Appl. Surf. Sci.*, 2005, **246**, 239–249.
- 31 A. Litke, Y. Su, I. Tranca, T. Weber, E. J. M. Hensen and J. P. Hofmann, *J. Phys. Chem. C*, 2017, **121**, 7514–7524.
- 32 See Fig. S3† for control data on electrodes lacking the Co_{C11P} catalyst or containing decyl phosphonic acid (C_{10P}).
- 33 S. Cobo, J. Heidkamp, P.-A. Jacques, J. Fize, V. Fourmond, L. Guetaz, B. Joussetme, V. Ivanova, H. Dau, S. Palacin, M. Fontecave and V. Artero, *Nat. Mater.*, 2012, **11**, 802–807.
- 34 D. H. Nam, J. Z. Zhang, V. Andrei, N. Kornienko, N. Heidary, A. Wagner, K. Nakanishi, K. P. Sokol, B. Slater, I. Zebger, S. Hofmann, J. C. Fontecilla-Camps, C. B. Park and E. Reisner, *Angew. Chem., Int. Ed.*, 2018, **57**, 10595–10599.
- 35 J. E. O'Reilly, *Biochim. Biophys. Acta*, 1973, **292**, 509–515.
- 36 N. Kaeffer, J. Massin, C. Lebrun, O. Renault, M. Chavarot-Kerlidou and V. Artero, *J. Am. Chem. Soc.*, 2016, **138**, 12308–12311.
- 37 D. Bartesaghi, I. D. Perez, J. Knierpert, S. Roland, M. Turbiez, D. Neher and L. J. A. Koster, *Nat. Commun.*, 2015, **6**, 7083.

