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# Black Silicon Photoanodes Entirely Prepared with Abundant Materials by Low-Cost Wet Methods

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KEYWORDS. solar energy conversion, oxygen evolution reaction, nickel, silicon, iron

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3 ABSTRACT We report a new design for water-splitting photoanodes, based on a highly absorbing  
4 black silicon (BSi) substrate modified with catalytic metal nanoparticles (NPs). The overall  
5 fabrication strategy is cost-efficient as it only requires the use of abundant materials and simple  
6 wet procedures such as electrochemical etching and electrodeposition and does not involve the use  
7 of buried homojunction and protection layer. Importantly, these results demonstrate that  
8 electrodeposited transition metal NPs can stabilize structured Si photoelectrodes without the need  
9 for a protection layer.  
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23 The depletion of fossil fuels and the increasing environmental pollution imply an urgent  
24 development of alternative technologies to foster the energy transition.<sup>1</sup> In this frame, the large-  
25 scale use of solar energy is particularly promising because this is the most abundant renewable  
26 energy on Earth.<sup>2</sup> Solar energy research went through a frantic period after the oil crisis in the  
27 early seventies and, in this period, water-splitting photoelectrochemical cells (PECs) started to  
28 gain attention.<sup>3</sup> These devices are based on light-absorbing semiconductor (SC) materials  
29 interfaced with a liquid phase. Under illumination, the charge carriers generated in the SC phase  
30 are driven to the solid-liquid interface where they react to generate products in solution.<sup>4</sup> In the  
31 case of water-splitting PECs, photogenerated holes ( $h^+$ ) and electrons ( $e^-$ ) respectively oxidize  
32 and reduce water to generate  $O_2$  and  $H_2$ , the latter being a highly promising clean energy carrier.<sup>1</sup>  
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34 An effective utilization of sunlight is required for PEC systems, which can be ideally done by  
35 employing small band gap and/or structured SCs.<sup>5</sup> Si is a highly abundant low band gap SC ( $E_g =$   
36 1.1 eV) that would be an excellent candidate for manufacturing PECs, especially because this  
37 material can be readily processed at the industrial level.<sup>6</sup> However, Si is difficult to be employed  
38 as a photoelectrode due to its notorious instability. This issue is even more pronounced when Si  
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3 is used as a photoanode in alkaline media because of: *i*) photocorrosion which induces fast  
4 electrochemical deactivation by generating an insulating SiO<sub>x</sub> layer<sup>7</sup> and *ii*) alkaline etching (AE)  
5 which causes the irreversible dissolution of the material.<sup>8</sup> The main strategy reported so far for  
6 manufacturing stable Si-based oxygen evolution reaction (OER) photoanodes consisted in first  
7 modifying the Si surface with a transparent and conformal protection layer such as TiO<sub>2</sub>,<sup>9,10</sup>  
8 Al<sub>2</sub>O<sub>3</sub>,<sup>11</sup> SiO<sub>2</sub><sup>12</sup> and then applying a co-catalytic (*cocat*) coating in order to improve oxygen  
9 evolution reaction OER kinetics. Recently, unexpected reports from our group<sup>13,14</sup> and  
10 others<sup>15,16,17</sup> have revealed that heterogeneous nanoscale pinched-off *n*-Si/metal (metal = Ni or  
11 Co) junctions, prepared only by a simple aqueous-phase electrodeposition process, can be  
12 employed to manufacture Si photoanodes exhibiting high photocurrent densities and stability  
13 without a protection layer. In addition, we have shown lately that these photoanodes can be  
14 implemented in complete monolithic water-splitting PEC to produce H<sub>2</sub> under simulated  
15 sunlight.<sup>18</sup> The transposition of this concept to structured *n*-Si surfaces has been reported only  
16 once,<sup>19</sup> however, with a Si structuration process based on lithography and vacuum-based  
17 physical methods.<sup>19</sup> In addition, there has been recently a strong interest in producing  
18 photoelectrode based on black silicon (BSi) (that is, highly absorbing structured Si).<sup>20,21</sup> These  
19 systems required, so far, the use of a TiO<sub>2</sub> protection layer deposited by atomic layer deposition  
20 (ALD).<sup>22</sup> In this communication, we report the first example of protection layer-free BSi  
21 photoanode, prepared by a simple and cost-efficient electrodeposition strategy.

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51 The fabrication strategy is described in **Scheme 1**. First, *n*-type Si(100) wafers were structured  
52 by employing a rapid (<10 min) two-step method previously introduced by our group to fabricate  
53 a ~10 μm-thick porous Si layer (**Figure 1a**) composed of vertically-aligned macropores bearing  
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3 spikes at their extremities (Figure 1b). As previously reported,<sup>23</sup> these structures efficiently trap  
4 the incident light, as shown by their pitch-black visual appearance and their low reflectance (*vide*  
5 *infra*), we refer herein to them as BSi. Next, we modified the BSi surfaces with a catalytic and  
6 stabilizing coating. Toward this goal, we first electroplated Ni<sup>0</sup> on the freshly-hydrogenated BSi  
7 using the same method as the one previously employed on planar Si.<sup>13,14</sup> This electrodeposition  
8 leads to the homogeneous decoration of the porous BSi layer with randomly-dispersed Ni  
9 nanoparticles (NPs). The role of these NPs role is to generate an array of robust nanoscale  
10 Schottky junctions, promoting an efficient hole transfer across the Si/Ni and Ni/electrolyte  
11 interface as well as a high photovoltage for sunlight-assisted OER, caused by the pinch-off  
12 effect.<sup>24-26</sup> In addition, considering that NiFe alloys are currently among the best OER  
13 electrocatalysts,<sup>27</sup> and that Fe incorporation improves OER kinetics in Si/Ni planar  
14 photoanodes,<sup>14</sup> we have decided to coat the Ni NPs with a NiFe layer by a second aqueous  
15 electrodeposition step. To do so, the BSi surfaces modified by the Ni NPs were first thermally  
16 oxidized after the first electrodeposition step by placing them into an oven at 90 °C for 1 h, in  
17 order to generate a SiO<sub>x</sub> layer onto the unprotected Si surface (red color in Scheme 1). The  
18 presence of this electrically-passivating layer ensures the electrodeposition of the NiFe coating  
19 only on the Ni NPs by preventing the deposition on the BSi, as shown in Scheme 1 (note that the  
20 electrodeposition was performed employing an aqueous electrolyte containing Ni and Fe salts at  
21 a 10/1 molar ratio, see SI and Figure S4 for more details). Scanning and transmission electron  
22 microscopy (SEM and TEM) analyses were performed before and after electrodeposition in  
23 order to characterize the coatings. Cross-section SEM (Figure 1c), confirmed the presence of  
24 randomly-dispersed Ni/NiFe NPs on the walls of the BSi pores and spikes. In addition, TEM  
25 allowed a clear morphological characterization of the Si/Ni nanoscale Schottky junctions (Figure  
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3 1d) as well as the metal NPs (inset of Figure 1d). As shown in these figures, the metal particles  
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5 were hemispherical with diameters below 100 nm.  
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9 The chemical composition of the modified surfaces was determined by employing several  
10 methods. First, scanning transmission electron microscopy coupled with energy-dispersive X-ray  
11 spectroscopy (STEM-EDS) mapping (**Figure 2a-d**), performed on a BSi area containing 2  
12 isolated NPs (Figure 2a), confirmed the presence of Si in the substrate (red signal in Figure 2b)  
13 and Ni in the NPs (blue signal in Figure 2c). However, Fe mapping shown in Figure 2d was not  
14 fully conclusive: whereas its presence could be discerned for the largest NP, it was not possible  
15 to observe it for the smallest NP. X-ray photoelectron spectroscopy (XPS, survey spectra are  
16 shown in Figure S1) confirmed the presence of the two metals after surface modification, as  
17 revealed by the appearance of two Ni 2p<sub>3/2</sub> peaks at 851.9 and 855.8 eV (Figure 2e),  
18 corresponding respectively to Ni<sup>0</sup> and oxidized Ni (generated during thermal annealing before  
19 NiFe deposition), as well as the Fe 2p<sub>3/2</sub> peak (Figure 2f) at 710.8 eV. The analysis of these two  
20 regions allowed to estimate a Ni/Fe ratio of 13 ± 5, a stoichiometry that fits well with those  
21 reported for highly-efficient OER-active Ni<sub>1-x</sub>Fe<sub>x</sub>OOH materials. This oxyhydroxide species is  
22 the known OER-active phase for NiFe mixed oxides, that is electrochemically generated at the  
23 OER potential.<sup>28</sup> The successful electrodeposition of NiFe on the surface was further confirmed  
24 by X-ray absorption near edge structure (XANES, Figure 2g), which showed no signal at the Fe  
25 K-edge after the first electrodeposition step but clearly exhibited an edge at E<sub>0</sub> = 7133 eV after  
26 the second electrodeposition step. These results confirm that our preparation method is effective  
27 to modify BSi surface by Ni/NiFe NPs. Next, we study the optical and photoelectrochemical  
28 properties of these surfaces.  
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3 The total reflectance of several samples was measured using an integrating sphere in order to  
4 quantify their light-trapping ability (**Figure 3a**). The comparison between the reflectance  
5 spectrum obtained for planar Si (purple curve) and that measured for BSi (green curve) shows a  
6 considerable reflectance decrease caused by the multiple internal reflections of the incident light  
7 within the porous structure. In addition, it is interesting to note that the modification with the  
8 Ni/NiFe NPs (red curve) did not affect significantly the reflectance of the BSi, that remained  
9 below 10% in the visible range. The surface was tested for OER in 1 M NaOH (measured pH =  
10 13.6), as shown by the cyclic voltammograms (CVs) of Figure 3b. In the dark, no photoresponse  
11 could be obtained (black curve). In contrast, high photocurrents were recorded under  
12 illumination with simulated sunlight (AM 1.5G, 100 mW cm<sup>-2</sup>, red curve). TEM experiments,  
13 performed on a photoanode that was employed for OER during 50 CV cycles, demonstrated the  
14 good adhesion of the NPs on the BSi surface (Figure 1d). The photoelectrochemical  
15 measurements obtained for 10 independently-prepared electrodes (Table S1), allowed to  
16 calculate benchmark values of maximum photocurrent  $j_{max} = 23.8 \pm 0.8$  mA cm<sup>-2</sup> and  
17 photocurrent density at the standard potential of the O<sub>2</sub>/H<sub>2</sub>O couple,  $j_{1.23V} = 4.3 \pm 0.8$  mA cm<sup>-2</sup>. In  
18 contrast, non-modified BSi surfaces failed to promote OER, as shown by the CV obtained under  
19 illumination with such a surface (green curve in Figure 3b). Photocurrent spectroscopy, recorded  
20 with a modified BSi surface at +2 V, revealed that the incident photon-to-current efficiency  
21 (IPCE, red curve in Figure 3b) is comprised between 50 and 68% in the visible range. The  
22 integrated value of the photocurrent spectrum (purple curve in Figure 3b) obtained by using the  
23 IPCE and the AM 1.5G solar spectrum is in good agreement (22.8 mA cm<sup>-2</sup>) with the value  
24 obtained by cyclic voltammetry at the same potential, confirming the validity of our  
25 measurements. An optical detection system was employed to measure the quantity of produced  
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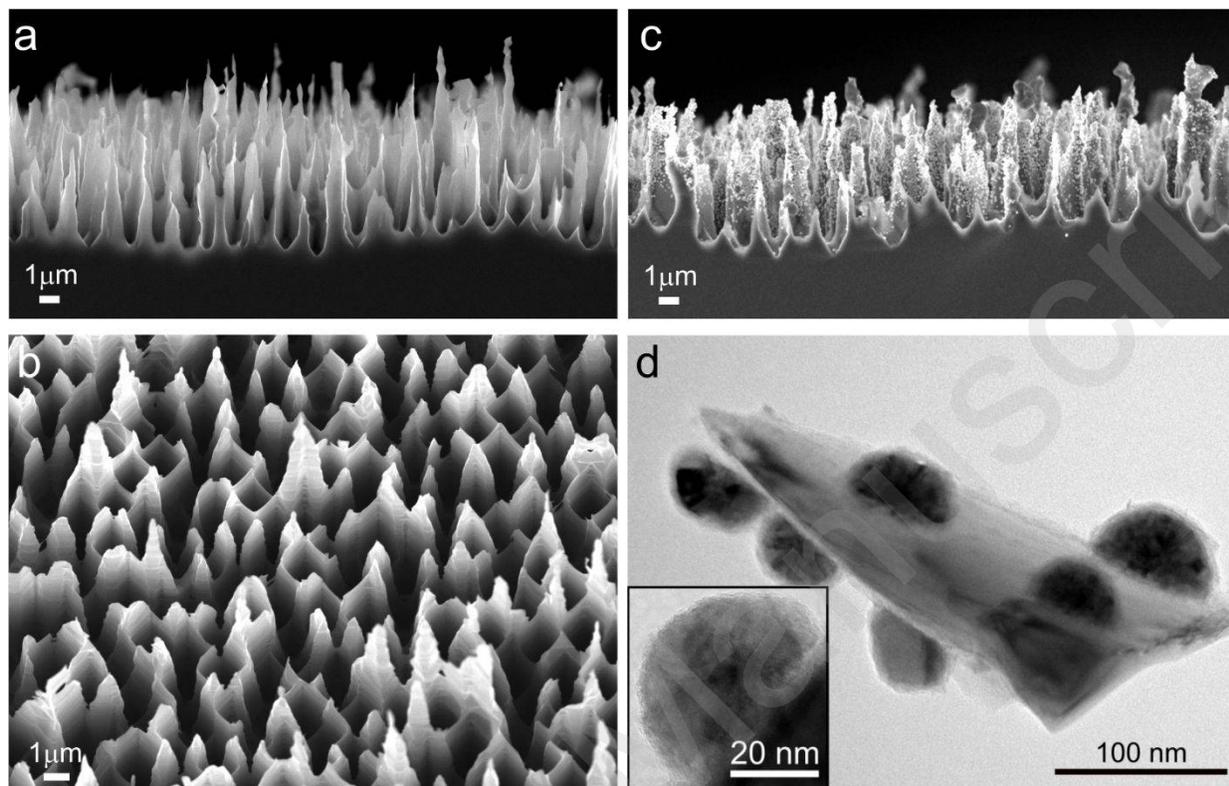
O<sub>2</sub> during a 30 min-long preparative electrolysis (red curve in Figure 3d). This measurement, which was performed under illumination, confirmed that O<sub>2</sub> was produced when the modified photoanode was biased at +2 V ( $t = 15$  min), which ceased immediately when the polarization was stopped ( $t = 45$  min). The calculated production rate is  $220.6 \mu\text{mol h}^{-1} \text{cm}^{-2}$ . Based on the electrical charge consumed during this experiment (the chronoamperogram (CA) is shown in Figure S2), we calculated the theoretical amount of O<sub>2</sub>, that is plotted as purple disks in Figure 3d. Taking into account the experimental and the theoretical numbers of mole of O<sub>2</sub>, we calculated a Faradaic efficiency ( $\eta$ ) of 96%. The stability of a photoanode was tested by a prolonged electrolysis test at +2 V in 1 M NaOH (Figure 3e), which revealed that OER could be performed over 16 h with a rather slow degradation of the electrode, as shown by the fact that it maintained a photocurrent higher than 75% of its initial value at the end of the test.

The performance of our photoanodes is compared with that of other reported systems based on structured *n*-type Si in Table S2. Note that in this table, we have deliberately omitted the electrodes based on a Si homojunction (*e.g.*  $np^+$ -Si), that require additional fabrication steps and fall outside the scope of the present article. From that table, it can be observed that our method is the only one that does not imply the use of a conformal protection layer (such as ALD TiO<sub>2</sub>) and that is entirely based on wet methods. Interestingly, we can also notice that our operation time is the highest that has been reported so far for protection-free structured photoanodes and that the performance we obtained in terms of  $j_{1.23V}$  and operation time is higher than that reported for BSi using an ALD-deposited TiO<sub>2</sub> protection layer.<sup>22</sup>

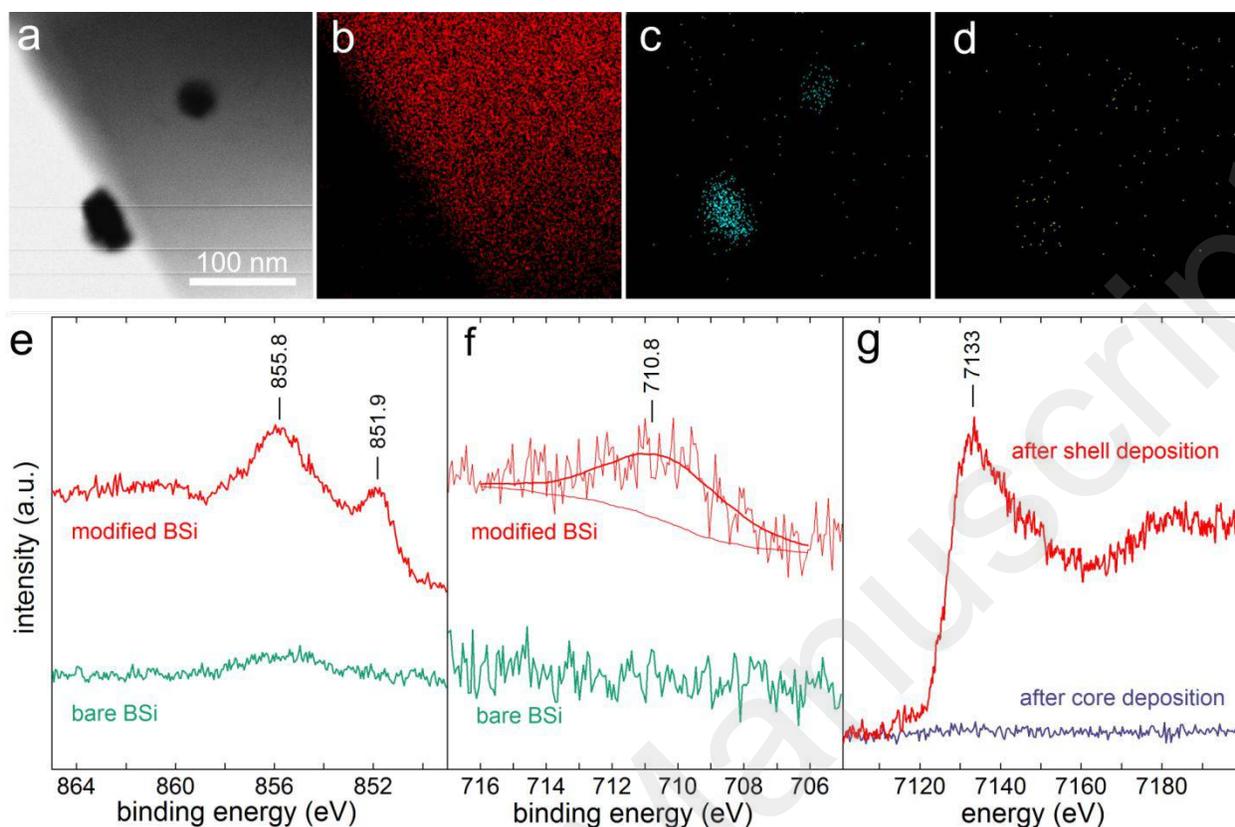
In conclusion, we have reported a new strategy based on simple, low-cost wet methods to manufacture BSi-based photoanodes. The light-trapping *n*-type BSi substrate is prepared by a

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3 fast (<10 min) double-step technique, and then covered by Ni/NiFe NPs using aqueous  
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5 electrodeposition. In this system, the Ni cores generate an effective pinched-off heterogeneous  
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7 junction with *n*-Si<sup>24</sup> and the NiFe layer promote efficient OER catalysis. These photoanodes can  
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9 operate for 16 h, producing O<sub>2</sub> at high rates with a quasi-quantitative Faradaic efficiency. To  
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11 date, this is the first example reported of buried homojunction-free and protection layer-free BSi  
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13 only based on simple wet methods. In addition, these results demonstrate that the stabilization of  
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15 *n*-Si by electroplated Ni NPs, which was reported for planar Si(100),<sup>13,14</sup> is also effective on  
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17 structured *n*-Si. We are confident that these results will inspire other groups to develop original  
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19 designs for photoanodes that will be employed for cost-effective water splitting PECs.  
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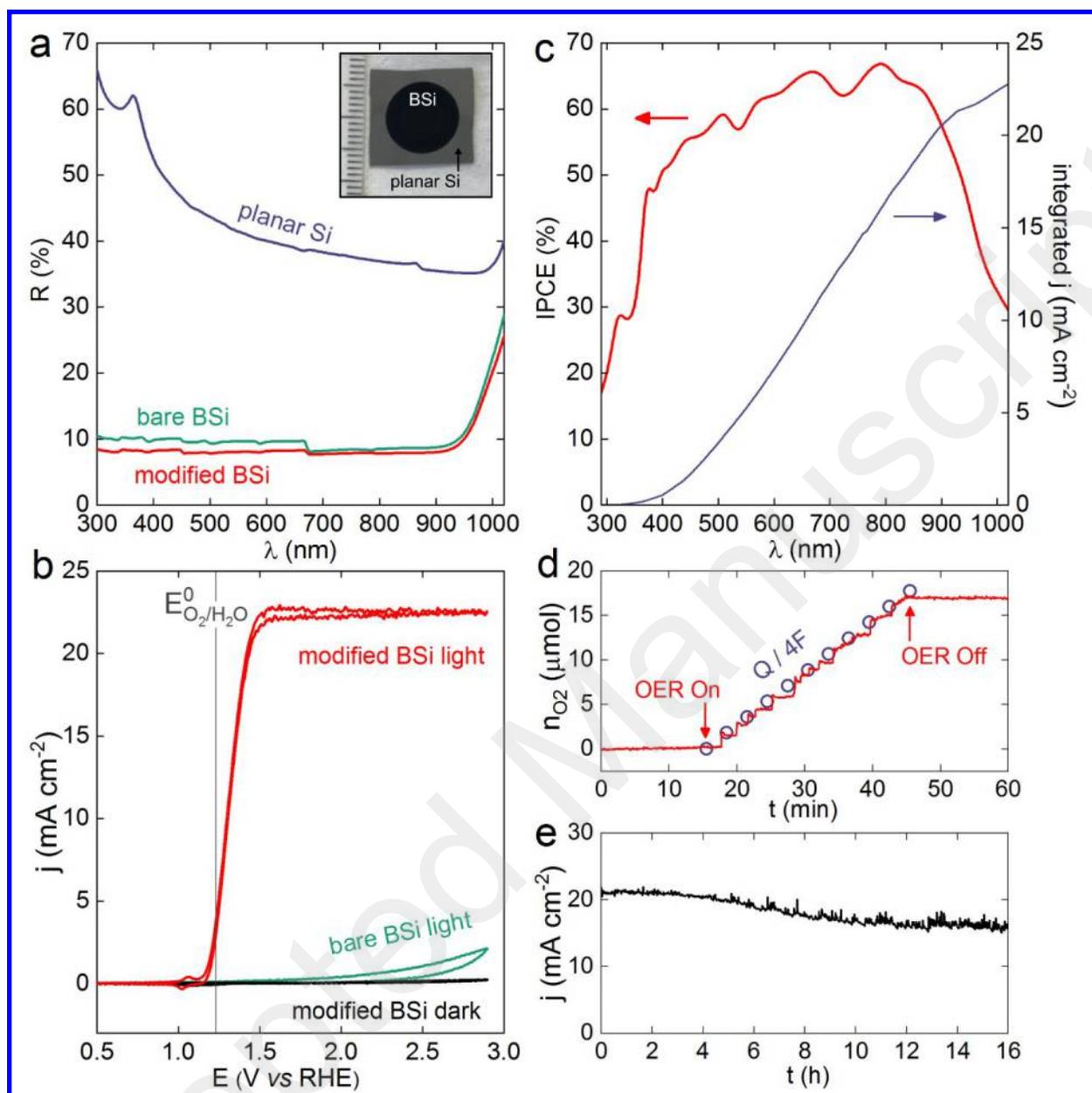
## FIGURES



**Figure 1. Structural characterization of the BSi photoanodes.** a) Cross-section SEM picture showing the BSi layer before electrodeposition. b) Tilted SEM picture showing the top of the BSi surface. c) Cross-section SEM picture showing the BSi layer after electrodeposition of the Ni/NiFe NPs. d) TEM picture showing a modified BSi spike detached from the layer (in this case the surface has been employed for OER, *vide infra*). Inset: high magnification TEM picture showing a single Ni/NiFe NP after electrodeposition of NiFe.



**Figure 2. Chemical characterization of the BSi photoanodes.** a) Bright field STEM picture image of Ni/NiFe NPs deposited on BSi and corresponding b) Si, c) Ni and d) Fe EDS maps. e,f) XPS spectra showing the Ni 2p<sub>3/2</sub> and the Fe 2p<sub>3/2</sub> regions. g) XANES spectra recorded at the Fe K-edge. In e-g), the green spectra were recorded before Ni electrodeposition, the purple spectrum was recorded after Ni electrodeposition and the red spectra were recorded after NiFe electrodeposition.



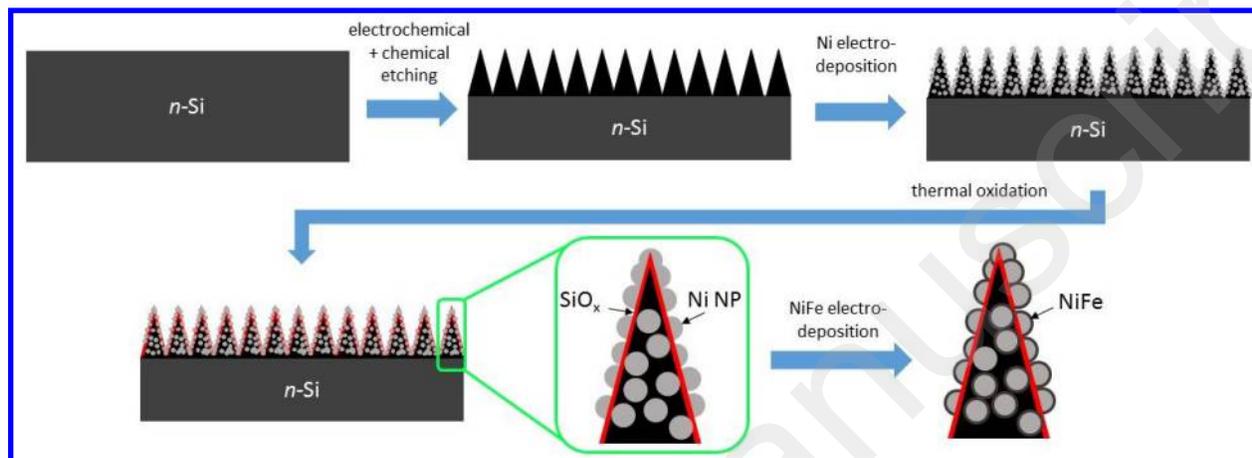
**Figure 3. Optical and photoelectrochemical characterization of the BSi photoanodes.** a) Total reflectance spectra. Inset: photograph showing the visual appearance of a BSi disk prepared on a flat Si square. b) Cyclic voltammograms recorded in the dark (black curve) or under simulated sunlight (colored curves) at 100 mV s<sup>-1</sup> in 1 M NaOH (not corrected from the Ohmic drop). c) IPCE spectrum (red curve) and corresponding integrated photocurrent (purple curve) recorded at +2 V vs RHE. d) Experimentally measured (red curve) and theoretical (purple disks) O<sub>2</sub> detection

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3 curves, obtained during an electrolysis at +2 V vs RHE under illumination with simulated sunlight.

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5 e) Chronoamperogram showing the photocurrent density as a function of time during a preparative  
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7 electrolysis at +2 V vs RHE under illumination with simulated sunlight. In a,b) the purple curve  
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9 was recorded on planar Si, the green curves were recorded on bare BSi and the red curves were  
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11 recorded on Ni/NiFe-modified BSi.  
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## SCHEME

**Scheme 1.** Scheme showing the fabrication steps that were employed to prepare the BSi photoanodes.



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3 ASSOCIATED CONTENT  
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6 **Supporting Information.** Experimental section, XPS survey spectra, supplementary figures, and  
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8 tables are provided in supporting information. The following files are available free of charge.  
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14 AUTHOR INFORMATION  
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23 **Author Contributions**  
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25 The manuscript was written through contributions of all authors. All authors have given approval  
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27 to the final version of the manuscript.  
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## SYNOPSIS

