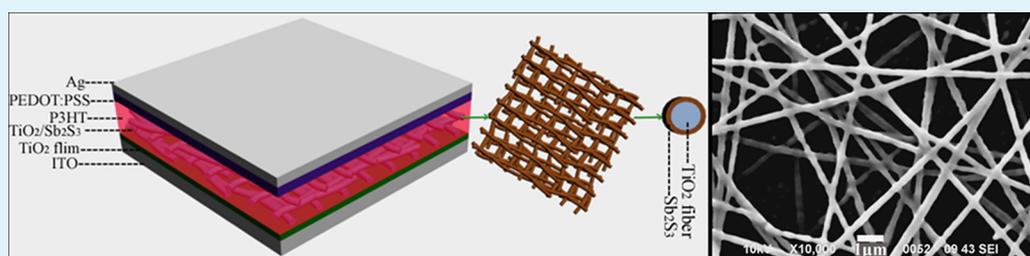


# High Efficiency Solar Cells As Fabricated by $\text{Sb}_2\text{S}_3$ -Modified $\text{TiO}_2$ Nanofibrous Networks

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**ABSTRACT:** High-efficiency hybrid solar cells (HSCs) based on electrospun titanium dioxide ( $\text{TiO}_2$ ) nanofibers plus poly(3-hexylthiophene) (P3HT) are fabricated by means of both the pretreatment using tetrahydrofuran (THF) vapor and the surface modification using n-type antimony chalcogenide ( $\text{Sb}_2\text{S}_3$ ) on the  $\text{TiO}_2$  nanofibrous networks. It is revealed that the THF pretreatment not only reinforces the interfacial physical contact but also suppresses the interfacial recombination. The  $\text{Sb}_2\text{S}_3$  modification improves the light absorption and charge transfer. Given that the active layer of the HSCs is as thin as 300 nm, it is demonstrated that the power conversion efficiency (PCE) is enhanced over 175%, exhibiting a PCE of 2.32%.

**KEYWORDS:** hybrid solar cells, electrospun  $\text{TiO}_2$  nanofibrous networks, THF pretreatment,  $\text{Sb}_2\text{S}_3$  modification, P3HT

## 1. INTRODUCTION

Recently, photovoltaic devices based on composites of n-type inorganic nanomaterials and p-type organic conjugated polymers have attracted much attentions because of their relatively high electron mobility and affinity, high dielectric constant, and potentials for large-area solar cells production at low cost.<sup>1–5</sup> Inorganic nanomaterials also feature high physical and chemical stability. One of the major problems in such kind of hybrid solar cells (HSCs) is the low power conversion efficiency (PCE) because the electron transfer is hindered by both the poor contact at inorganic–organic interface and the broken transport paths in the active layers. A well proven and widely adopted roadmap to improve electron transport both at the interface and in the active layers is to use nanocrystalline structures such as template porous structures,<sup>6</sup> vertically aligned nanorods,<sup>7,8</sup> and electrospun nanofibers.<sup>2–4</sup> Among these options, electrospinning technique is more promising, as it provides a simple, cost-effective, and large-area approach to prepare various materials with one-dimensional (1D) nanostructures. The as-prepared electrospun nanofibers have already been used in various solar cells. Unfortunately, the efficiency obtained so far is still low (<1.1%) compared with dye-sensitized solar cells.<sup>2</sup>

$\text{TiO}_2$  as an n-type semiconductor is a good candidate for photovoltaic material because of its good air stability, high electron mobility, and appropriate band alignment. So far, the HSCs based on  $\text{TiO}_2$  nanostructures yet have low PCE. One

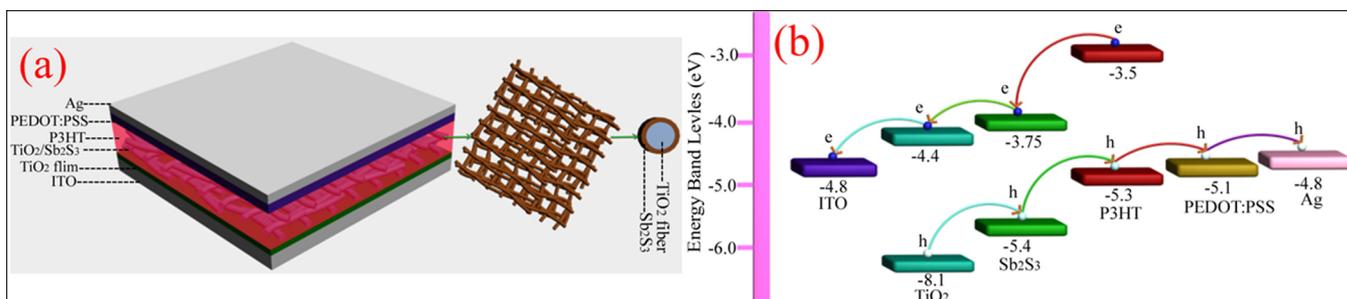
reason is that  $\text{TiO}_2$  can only absorb the UV region radiation due to its wide band gap. To enhance the light harvesting, we can anchor sensitizers such as organic dye or inorganic semiconductor materials on the surface of  $\text{TiO}_2$ .<sup>9–11</sup> Inorganic semiconductors offer many advantages such as high extinction coefficient and capability of optical absorption tailoring over a wider wavelength range.<sup>7,8,12–14</sup>  $\text{Sb}_2\text{S}_3$  as one of metal chalcogenides is an attractive one among these sensitizers. It has high optical absorption coefficient ( $7.3 \times 10^4 \text{ cm}^{-1}$  at 600 nm) and suitable energy band gap of 1.7–1.9 eV. These allow a strong and broad absorption in the visible region, which in turn contributes to more efficient photon harvesting.<sup>7,12,13</sup>

To improve the efficiency of  $\text{TiO}_2$  nanomaterial-based HSCs, other aspects include consideration of geometric shapes and material processing. High-surface-area 1D nanofibrous networks can provide large-area interface for charge separation and offer continuous transport paths for photogenerated charges. Moreover, compared with  $\text{TiO}_2$ /P3HT nanoparticles hybrids, the anisotropic  $\text{TiO}_2$  nanofibers in P3HT favor increased vertical stacking of P3HT crystallites, thus providing more effective hole transport perpendicular to the substrate.<sup>15</sup> Another problem associated with P3HT is that oxygen can induce traps in P3HT and deteriorate the performance of

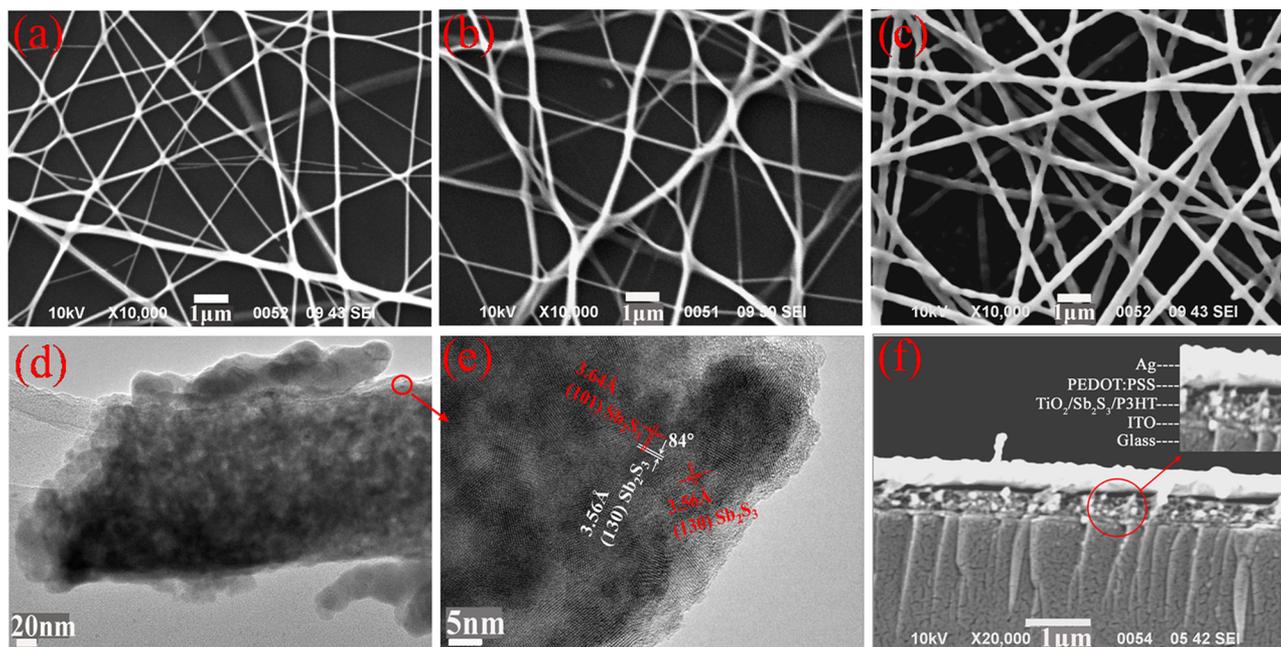
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**Figure 1.** (a) Schematic diagram of the HSC device based on  $\text{Sb}_2\text{S}_3$ -modified  $\text{TiO}_2$  nanofibrous networks and P3HT. (b) Corresponding energy level diagram of the corresponding materials used in our devices.



**Figure 2.** SEM images of (a)  $\text{TiO}_2(\text{UT})$ , (b)  $\text{TiO}_2(\text{T})$ , and (c)  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3$  nanofibers. (d) TEM image of an electrospun  $\text{TiO}_2$  nanofiber modified with  $\text{Sb}_2\text{S}_3$  (growth time: 3h). (e) High resolution TEM image of  $\text{Sb}_2\text{S}_3$  deposited on a  $\text{TiO}_2$  nanofiber. (f) Cross-sectional SEM view of the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure. UT, untreated  $\text{TiO}_2$  nanofibers; T,  $\text{TiO}_2$  nanofibers pretreated with THF vapor.

HSCs.<sup>16–18</sup> Therefore, almost all HSCs based on  $\text{TiO}_2/\text{Sb}_2\text{S}_3$  and P3HT ( $\text{TiO}_2/\text{Sb}_2\text{S}_3/\text{P3HT}$ ) structures are fabricated in inert atmosphere or vacuum. A method that can solve this oxygen-relevant problem and allow device fabrication in air is highly desirable, and will be both cost- and technology-competitive.

Taking care of these considerations, in this work HSCs based on P3HT and electrospun  $\text{TiO}_2$  nanofibrous network is fabricated in ambient air. In order to both reinforce the physical contact and increase the interfacial area to facilitate rapid charge separation and transport, we select THF vapor to pretreat the  $\text{TiO}_2$  nanofiber networks, then modify the nanofibers by depositing a  $\text{Sb}_2\text{S}_3$  layer on  $\text{TiO}_2$  nanofiber surface. The durations of the THF vapor pretreatment and the  $\text{Sb}_2\text{S}_3$  layer growth are optimized. Subsequently, we investigate the performance of the HSCs based on the  $\text{TiO}_2/\text{Sb}_2\text{S}_3/\text{P3HT}$  structures. The recombination in the as-prepared heterojunction is suppressed dramatically upon the THF pretreatment. The light absorption in the visible region is enhanced significantly by  $\text{Sb}_2\text{S}_3$  modification. Eventually, we demonstrate an improvement of 175% of the PCE with respect to reference cell.

## 2. EXPERIMENTAL DETAILS

**Materials and methods.** Figure 1a shows a schematic diagram of the hybrid solar cell (HSC) device to be fabricated in this work. For each sample, cleaned ITO substrate is first coated with a compact  $\text{TiO}_2$  film of about 40 nm in thickness by spin-coating sol–gel method. The  $\text{TiO}_2$  nanofibrous networks are then electrospun at a flow rate of 0.3 mL/h onto the  $\text{TiO}_2$  film surface from a precursor gel containing tetrabutyl titanate (5%) and poly (vinylpyrrolidone) (PVP) (5%) in the solution of isopropanol and 2-methoxyethanol (2:1 in volume) in ambient air. For this electrospinning, a voltage of 9.0 kV between a metal orifice and a grounded rotating collector with a distance of 8 cm is applied for 45 min.

Subsequently, several of the as-prepared samples are chosen as the reference samples and the other samples are submitted to next processing. For the THF vapor pretreatment, the samples are kept for different durations in an anhydrous THF chamber without direct contact with THF. Samples are untreated and chosen as the reference. After the pretreatment, these samples together with the reference samples are annealed at 500 °C for 15 min by rapid thermal processing (RTP) to remove organic components and allow nucleation and growth of  $\text{TiO}_2$  grains inside the fibers. Then, the  $\text{TiO}_2$  nanofibers besides the reference samples are covered with a  $\text{Sb}_2\text{S}_3$  layer using a chemical bath deposition method in an aqueous solution of  $\text{SbCl}_3$  and  $\text{Na}_2\text{S}_2\text{O}_3$  in ambient air.<sup>7,19–21</sup> The thickness of the  $\text{Sb}_2\text{S}_3$  layer is controlled by the growth time.

After the  $\text{Sb}_2\text{S}_3$  deposition, the as-prepared samples including the reference samples are rinsed thoroughly with deionized water and dried in air, and then annealed in nitrogen ( $\text{N}_2$ ) ambient at  $350\text{ }^\circ\text{C}$  for 20 min. Here, it is found that the ambient for the subsequent cooling immediately after the annealing is critical for the sample performance. Because the  $\text{Sb}_2\text{S}_3$  can be partially oxidized to  $\text{Sb}_2\text{O}_3$  during the cooling process in ambient air and the  $\text{Sb}_2\text{O}_3$  oxide presumably acts as a passivation layer benefiting to the reduction of the recombination between electrons in the  $\text{Sb}_2\text{S}_3$  layer and holes in the P3HT.<sup>22</sup> That is why better performance is achieved with the samples cooled in air rather than in  $\text{N}_2$ .

To fabricate the HSCs stemming from the above samples, a P3HT layer is infiltrated into the  $\text{TiO}_2$  nanofibrous networks by spin-coating a solution of P3HT (Reike, regioregularity is  $\sim 93\%$ , 20 mg/mL dissolved in toluene) at 1500 rpm for 30 s in ambient air. Our earlier results show that the deposition of P3HT in ambient air can obtain a performance comparable to that of the cells fabricated in an inert environment.<sup>23</sup> So the deposition of P3HT is carried out in ambient air. Subsequently, a layer of poly(3,4-ethylenedioxythiophene):poly(4-styrene sulfonate) (PEDOT:PSS, Bayron P, Bayer A.G.) is spun on the top of P3HT layer in ambient air, followed by thermal annealing at  $120\text{ }^\circ\text{C}$  for 20 min in a glovebox filled with high purity  $\text{N}_2$ . Finally, an Ag electrode is evaporated on the sample surface through a shadow mask under a vacuum of  $1 \times 10^{-4}$  Pa and these samples are postannealed at  $100\text{ }^\circ\text{C}$  for 20 min. To this stage, the HSCs are fabricated with the standard in-plane size of  $2\text{ mm} \times 2\text{ mm}$ . The morphology, dimensions, and crystallinity of the  $\text{TiO}_2$  nanofibers covered with the  $\text{Sb}_2\text{S}_3$  layer are characterized by scanning electron microscopy (SEM, JEOL 5700, Japan), transmission electron microscopy (TEM, F2010, Japan), and X-ray diffraction (XRD, X'Pert PRO, Cu K $\alpha$  radiation), respectively.

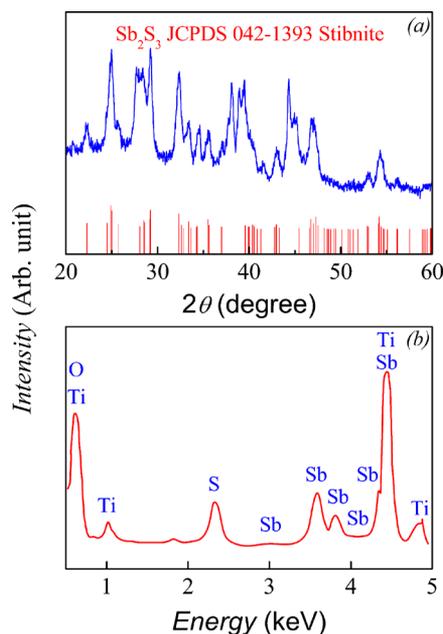
**Device Characterizations.** The photovoltaic performance of these HSCs is characterized using a Keithley 2400 source meter under an illumination of  $100\text{ mW cm}^{-2}$  (Newport 91160, 150W solar simulator equipped with an AM1.5G filter). The radiation intensity is calibrated by a standard silicon solar cell (certified by NREL) as the reference. The external quantum efficiency (EQE) and the UV–vis absorption spectra are measured using a standard EQE system (Newport 66902). The electrochemical impedance spectroscopy (EIS) measurements are performed on the Zahner Zennium electrochemical workstation in the dark. A 10 mV ac sinusoidal signal source is employed over the constant bias with the frequency ranging from 1 Hz to 4 MHz.

### 3. RESULTS AND DISCUSSION

The energy level diagram of the  $\text{TiO}_2/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure is shown in Figure 1b, where the energy levels across the  $\text{TiO}_2$ ,  $\text{Sb}_2\text{S}_3$ , and P3HT are aligned so that the excitation dissociation and charge transfer at the interfaces are energetically favorable. Moreover, successful sensitization of  $\text{TiO}_2$  by  $\text{Sb}_2\text{S}_3$  has been demonstrated, indicating also that the energy levels of  $\text{Sb}_2\text{S}_3$  and  $\text{TiO}_2$  match with each other.<sup>7,12,13,19–21</sup> Figure 2a–c show the SEM images of the only calcined  $\text{TiO}_2$  nanofibers (untreated,  $\text{TiO}_2(\text{UT})$ ),  $\text{TiO}_2$  nanofibers pretreated with THF vapor ( $\text{TiO}_2(\text{T})$ ), and  $\text{TiO}_2$  nanofibers pretreated with THF vapor and modified with  $\text{Sb}_2\text{S}_3$  (i.e., covered with a  $\text{Sb}_2\text{S}_3$  layer) ( $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3$ ), respectively. All these  $\text{TiO}_2$  nanofibers for SEM observations are coated on the underlying  $\text{TiO}_2$  films, respectively. It is seen that these nanofibers have a diameter distribution over 100–200 nm and they cross over one and another. Such fiber crossing can improve the effective interfacial contact, benefiting to the charge transfer. The THF pretreatment (Figure 2b) induces etched features on the  $\text{TiO}_2$  fiber surface, benefiting to the interjunction points and electrical contact between the ITO substrate and  $\text{TiO}_2$  fibrous networks.<sup>5</sup>

In comparison with the  $\text{TiO}_2(\text{UT})$ ,  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3$  fibers shown in Figure 2c become much coarser and indicate good

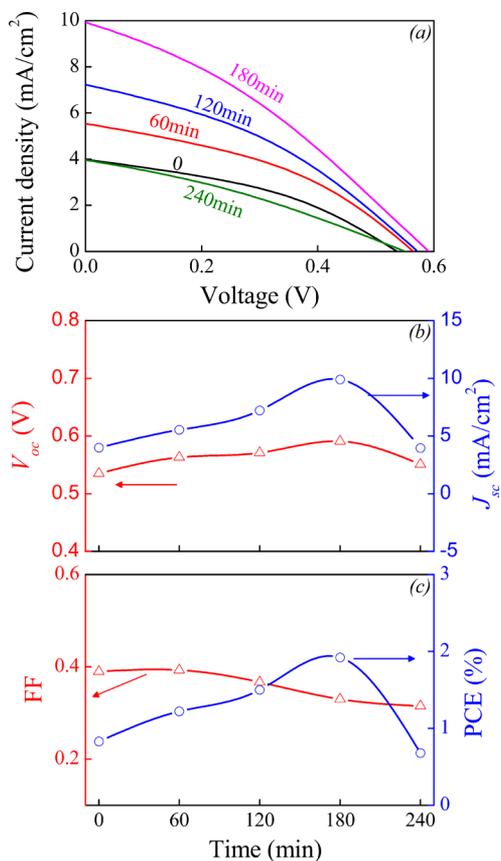
interfacial contact between them. And the interpore distance between nanofibrous channels is retained, preserving the effective surface area for P3HT penetration. Figure 2f exhibits the cross-sectional view of the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure, and it is seen that the space in-between the  $\text{TiO}_2/\text{Sb}_2\text{S}_3$  nanofibers is completely filled with P3HT. The thickness of the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  active layer is  $\sim 300\text{ nm}$ . Figure 2d, e show the TEM images of a  $\text{TiO}_2/\text{Sb}_2\text{S}_3$  nanofiber at different resolutions. It can be seen that the  $\text{Sb}_2\text{S}_3$  nanoparticles on the  $\text{TiO}_2$  surface have a broad size distribution, and are well-crystallized with lattice constant  $c$  of  $3.64\text{ \AA}$ , consistent with earlier reports.<sup>24</sup> Figure 3a shows the XRD patterns of a dark-



**Figure 3.** (a) XRD spectrum for  $\text{Sb}_2\text{S}_3$  thin film deposited on glass substrate in chemical bath for 3h and then annealed at  $350\text{ }^\circ\text{C}$  for 20 min in  $\text{N}_2$  atmosphere. The vertical lines are the characteristic XRD peaks of stibnite crystal structure (JCPDS 042–1393). (b) Energy-dispersive X-ray spectrum of the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3$  nanofibers.

brown  $\text{Sb}_2\text{S}_3$  layer on a slide glass substrate and the prepared condition is identical to that for the  $\text{Sb}_2\text{S}_3$  layer in the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure. We can see that the peaks can be well indexed by the stibnite crystal structure (JCPDS 042–1393). Figure 3b shows the energy dispersive X-ray spectrum of a  $\text{TiO}_2/\text{Sb}_2\text{S}_3$  nanofiber, indicating that the fiber contains elements Ti, O, Sb, and S.

It is understood that the thickness of the  $\text{Sb}_2\text{S}_3$  layer on the  $\text{TiO}_2$  nanofiber surface is crucial for the device performance, and it should be optimized to keep a balance between optical absorbance and charge transfer. Figure 4a shows the current density–voltage ( $J$ – $V$ ) characteristics of a series of  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structures with different  $\text{Sb}_2\text{S}_3$  growth time. Figure 4b, c show the detailed photovoltaic parameters including the short-circuit current density ( $J_{sc}$ ), open-circuit voltage ( $V_{oc}$ ), fill factor ( $FF$ ) and PCE as functions of the growth time. In the initial stage,  $J_{sc}$ ,  $V_{oc}$  and PCE all increase with the time, while  $FF$  decreases. The reduced  $FF$  may be due to the rough nature of the  $\text{Sb}_2\text{S}_3$  surface, caused by small crystallites developed during the annealing process, leading to small shunt resistance.<sup>8</sup> The duration time for the optimized parameters  $J_{sc} \sim 9.9\text{ mA/cm}^2$ ,  $V_{oc} \sim 0.591\text{ V}$ ,  $FF \sim 0.33$ , and

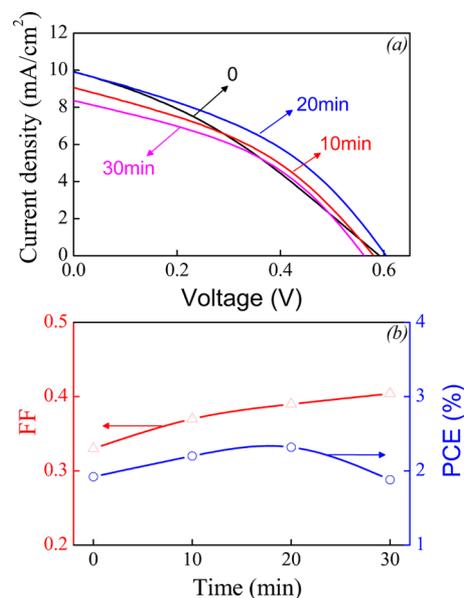


**Figure 4.** (a) Current–voltage ( $J$ – $V$ ) curves of  $\text{TiO}_2$  (UT)/ $\text{Sb}_2\text{S}_3$ /P3HT structure with different growth times for  $\text{Sb}_2\text{S}_3$  layer. (b) Measured  $V_{oc}$  and  $J_{sc}$  and (c) FF and PCE as a function of the growth time for  $\text{Sb}_2\text{S}_3$  layer in the  $\text{TiO}_2$  (UT)/ $\text{Sb}_2\text{S}_3$ /P3HT structure.

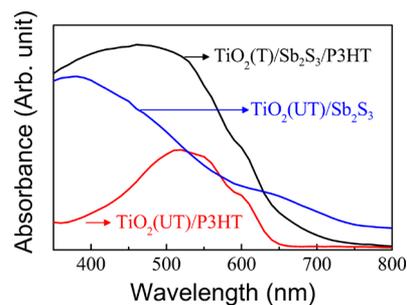
PCE  $\sim 1.92\%$ , is 3 h. This optimization can be confirmed by the absorbance spectrum and impedance data to be presented below. The thicker-than-optimal  $\text{Sb}_2\text{S}_3$  layer yields the lower PCE. This effect is mainly attributed to the decrease in  $J_{sc}$  and  $V_{oc}$ , because of the difficulty of deep penetration of P3HT, resulting in insufficient charge transport.<sup>12</sup>

The effect of the THF pretreatment on the charge transfer promotion is demonstrated by the  $J$ – $V$  curves obtained for different THF duration time, as shown in Figure 5a. Figure 5b demonstrates the dependence of parameters FF and PCE on the time. The PCE increases in the first 20 min and then decreases due to the decay of  $V_{oc}$  and  $J_{sc}$ . The optimal pretreatment time is 20 min, with the obtained parameters:  $V_{oc} \sim 0.603$  V,  $J_{sc} \sim 9.88$  mA/cm<sup>2</sup>, FF  $\sim 0.39$ , and PCE  $\sim 2.32\%$ .

It is believed that the optimized performance of the HSCs is attributed to three major factors: the increased absorption, accelerated charge transfer due to the interfacial contact reinforcement, and good band alignment at the interface between  $\text{Sb}_2\text{S}_3$  and P3HT. To clarify this issue, we present in Figure 6 the absorption spectra of the  $\text{TiO}_2$ (UT)/P3HT,  $\text{TiO}_2$ (UT)/ $\text{Sb}_2\text{S}_3$ , and  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT films, respectively. Obviously, the best performance is obtained for the  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT structure in terms of the high and broad absorption. Because  $\text{Sb}_2\text{S}_3$  offers high absorption coefficient and proper band gap of 1.7–1.9 eV,<sup>13</sup> the significant increase of the absorption benefits to the increase of  $J_{sc}$  due to the  $\text{Sb}_2\text{S}_3$  coverage and THF vapor pretreatment.



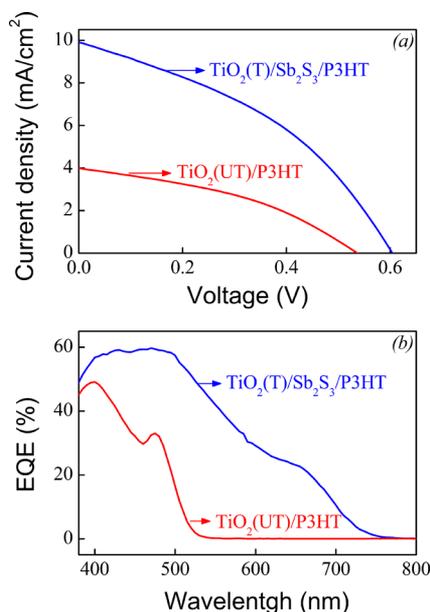
**Figure 5.** (a)  $J$ – $V$  curves, and (b) FF and PCE as a function of THF pretreatment time in the  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT structure.



**Figure 6.** Absorbance spectra of  $\text{TiO}_2$ (UT)/P3HT,  $\text{TiO}_2$ (UT)/ $\text{Sb}_2\text{S}_3$ , and  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT films.

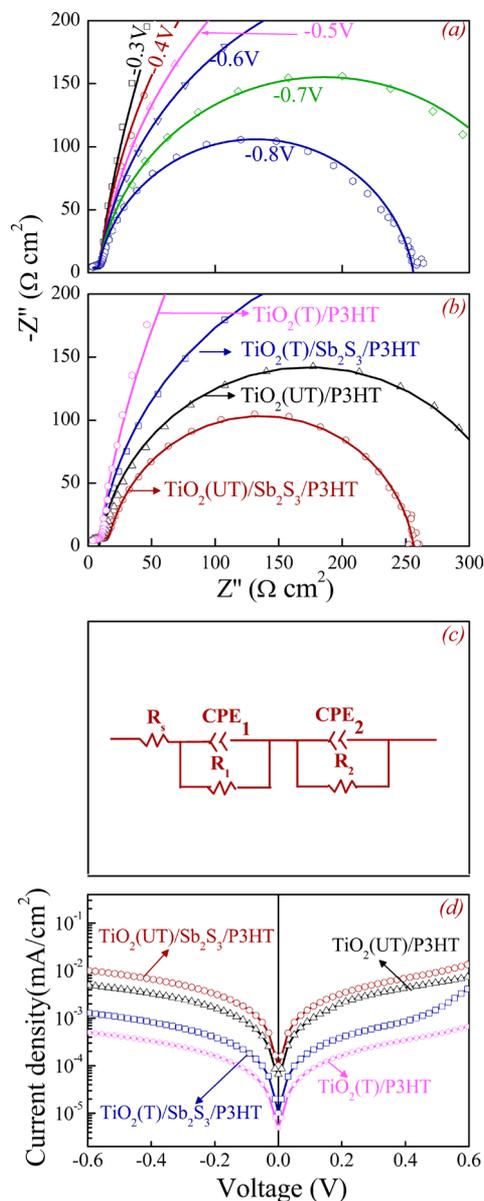
Figure 7a shows the  $J$ – $V$  curves for the  $\text{TiO}_2$ (UT)/P3HT and  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT structures prepared under the optimal conditions. For the fresh  $\text{TiO}_2$ (UT)/P3HT structure, one obtains  $V_{oc} \sim 0.535$  V,  $J_{sc} \sim 3.98$  mA/cm<sup>2</sup>, FF  $\sim 0.39$ , and PCE  $\sim 0.83\%$ . The  $\text{Sb}_2\text{S}_3$  coverage and THF pretreatment allow enhancement of the four parameters up to 0.603 V, 9.88 mA/cm<sup>2</sup>, 0.39, and 2.32%, respectively. This implies an improvement of PCE over 175% compared with the fresh structure. Figure 7b presents the measured EQE data for the  $\text{TiO}_2$ (UT)/P3HT and  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT structures, indicating the dramatic enhancement in the visible range. The EQE spectrum is consistent with the absorption enhancement induced by the  $\text{Sb}_2\text{S}_3$ . The high EQE in the  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT structure suggests that our device architecture performs well because of the good band alignment and an intimate contact between the  $\text{Sb}_2\text{S}_3$  and P3HT, as seen in the cross-sectional SEM of Figure 2(f).

For a better understanding of the microscopic mechanisms for the observed enhancement in the performance upon the THF pretreatment and  $\text{Sb}_2\text{S}_3$  coverage, the EIS and dark  $J$ – $V$  measurements are carried out. The EIS is used to evaluate the charge transfer and recombination at the interfaces. In our measurement, two semicircles in the Nyquist plots for each device are observed. As an example, Figure 8(a) shows the Nyquist plots of the EIS data for the  $\text{TiO}_2$ (T)/ $\text{Sb}_2\text{S}_3$ /P3HT



**Figure 7.** (a)  $J$ - $V$  curves, and (b) External quantum efficiency (EQE) of  $\text{TiO}_2(\text{UT})/\text{P3HT}$  and  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structures.

structure as a function of bias voltage. The plot shows two major features: one in the high frequency range and the other in the low frequency range. The arc observed in the high frequency range is independent of the applied bias, implying that it does not depend on the Fermi level position and can be ascribed to the parallel association of the capacitance  $\text{CPE}_{\text{ct}}$  of the cells and the charge transfer resistance  $R_{\text{ct}}$  at the P3HT/PEDOT:PSS/Ag interface.<sup>19</sup> It is noted that this resistance is expected to vary very slowly with voltage. The second is the large semicircle in the low frequency range which is ascribed to the parallel combination of the recombination resistance  $R_{\text{rec}}$  of electrons in the  $\text{TiO}_2$  with that of holes in P3HT or accepting states in  $\text{Sb}_2\text{S}_3$ , and the chemical capacitance  $\text{CPE}_{\text{rec}}$  of electrons in  $\text{TiO}_2$ .<sup>20</sup> The chemical capacitance is related to the electron Fermi level in  $\text{TiO}_2$  caused by the variation of the electron density as a function of bias voltage.<sup>25</sup> The size of the arc in the low-frequency range decreases as the applied bias voltage increases, as shown in Figure 8a. This is due to the decrease in  $R_{\text{rec}}$  with the increase of electron density. Subsequently, we compare the Nyquist plots for the four structures:  $\text{TiO}_2(\text{UT})/\text{P3HT}$ ,  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$ ,  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  and  $\text{TiO}_2(\text{T})/\text{P3HT}$  at fixed bias voltage of  $-0.6$  V (close to  $V_{\text{oc}}$ ) and the results are presented in Figure 8b. The solid lines in panels a and b in Figure 8 are the fittings of the experimental data using the model in Figure 8c. For more accurate fitting, CPE is used instead of ideal capacitances, C. It can be seen that the measured Nyquist plots can be fitted well. As shown in Figure 8b, compared with the fresh  $\text{TiO}_2(\text{UT})/\text{P3HT}$  structure, the  $\text{TiO}_2(\text{T})/\text{P3HT}$  structure has the highest  $R_{\text{rec}}$ , whereas the  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$  shows the lowest  $R_{\text{rec}}$ .<sup>26–30</sup> This indicates that the recombination rate increases upon the  $\text{Sb}_2\text{S}_3$  coating, and decreases upon the THF pretreatment because the recombination rate is inversely proportional to  $R_{\text{rec}}$ .<sup>25</sup> Because the effect of THF pretreatment on the recombination rate is greater than that of  $\text{Sb}_2\text{S}_3$  coating, the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure has lower recombination rate than that of  $\text{TiO}_2(\text{UT})/\text{P3HT}$  structure. Although the  $\text{Sb}_2\text{S}_3$  coating enhances the recombination rate, it significantly increases the light absorption, leading to the



**Figure 8.** (Color online) (a) Example of Nyquist plots of  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure as a function of bias voltages in the dark. (b) Nyquist plots of  $\text{TiO}_2(\text{UT})/\text{P3HT}$ ,  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$ ,  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  and  $\text{TiO}_2(\text{T})/\text{P3HT}$  structures measured at the bias voltage of  $-0.6$  V (close to  $V_{\text{oc}}$ ) in the dark. (c) Simplified equivalent circuit employed to fit the Nyquist plots. Solid lines in (a) and (b) are the fittings of the experimental data using the model in panel (c). (d) Dark  $J$ - $V$  curves of  $\text{TiO}_2(\text{UT})/\text{P3HT}$ ,  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$ ,  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  and  $\text{TiO}_2(\text{T})/\text{P3HT}$  structures.

enhancement of  $J_{\text{sc}}$ . And the THF pretreatment improves the interjunction and the electrical contact between the ITO substrate and  $\text{TiO}_2$  fibrous networks, which will benefit to the effective interfacial contact and charge transfer. Therefore, the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure shows the best performance. Compared with the  $\text{TiO}_2(\text{UT})/\text{P3HT}$  structure, the better performance in the  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure can be attributed to the intimate contact between the  $\text{Sb}_2\text{S}_3$  and P3HT, high and broad light absorption, and the proper band alignment at the interface.<sup>12</sup> Figure 8c shows the equivalent circuit, where a series resistance ( $R_s$ ) is included because of the contacts and wires.

For further investigation, the dark  $J$ - $V$  measurements are conducted for these samples too. Figure 8(d) shows the dark  $J$ - $V$  curves of the  $\text{TiO}_2(\text{UT})/\text{P3HT}$ ,  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$ ,  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  and  $\text{TiO}_2(\text{T})/\text{P3HT}$  structures. It is known that the dark current primarily results from the recombination and electron leakage current at the interface of the heterojunctions.<sup>31</sup> Compared with the fresh  $\text{TiO}_2(\text{UT})/\text{P3HT}$  structure, the dark current density in the  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure is higher, but in the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  and  $\text{TiO}_2(\text{T})/\text{P3HT}$  structures they are lower upon the forward bias. For the reverse bias, the  $\text{TiO}_2(\text{UT})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure exhibits a higher leakage current compared to the  $\text{TiO}_2(\text{UT})/\text{P3HT}$  structure, suggesting that more recombination events occur because of the presence of the  $\text{Sb}_2\text{S}_3$  layer. The  $\text{TiO}_2(\text{T})/\text{P3HT}$  structure exhibits the lowest leakage current, indicating that less recombination events occur because of the THF pretreatment. In addition, the lower dark current density in the forward bias indicates less recombination sites and interfacial traps in the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure. These results allow a conclusion that the  $\text{TiO}_2(\text{T})/\text{Sb}_2\text{S}_3/\text{P3HT}$  structure not only exhibits efficient light harvesting but also lower recombination rate and thus the best PCE.

#### 4. CONCLUSION

A series of HSCs based on electrospun  $\text{TiO}_2$  nanofibrous network and P3HT have been fabricated. We have demonstrated that the THF vapor pretreatment and  $\text{Sb}_2\text{S}_3$  modification of the  $\text{TiO}_2$  nanofibrous networks can improve significantly the performance of the as-prepared HSC. The underlying mechanism is intrinsically related to the increased light absorption and reduced recombination rate at the organic/inorganic heterojunctions. The measured PCE value of the as-prepared HSCs reaches up to 2.32 from 0.83% for the reference HSC without the THF pretreatment and  $\text{Sb}_2\text{S}_3$  modification.

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##### Notes

The authors declare no competing financial interest.

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#### REFERENCES

- (1) Huynh, W. U.; Dittmer, J. J.; Alivisatos, A. P. *Science* **2002**, *295*, 2425–2427.
- (2) Tai, Q. D.; Yan, F.; Zhao, X. Z. *J. Mater. Chem.* **2010**, *20*, 7366–7371.
- (3) Wu, S. J.; Tai, Q. D.; Yan, F. *J. Phys. Chem. C* **2010**, *114*, 6197–6200.
- (4) Wu, S. J.; Li, J. H.; Lo, S. C.; Tai, Q. D.; Yan, F. *Org. Electron* **2012**, *13*, 1569–1575.
- (5) Sudhagar, P.; Gonzalez-Pedro, V.; Mora-Sero, I.; Fabregat-Santiago, F.; Bisquert, J.; Kang, Y. S. *J. Mater. Chem.* **2012**, *22*, 14228–14235.
- (6) Goh, C.; Coakley, K. M.; McGhee, M. D. *Nano Lett.* **2005**, *5*, 1545–1549.
- (7) Cardoso, J. C.; Grimes, C. A.; Feng, X. J.; Zhang, X. Y.; Komarneni, S.; Zanonid, M. V. B.; Bao, N. Z. *Chem. Commun.* **2012**, *48*, 2818–2820.
- (8) Coakley, K. M.; Liu, Y. X.; Goh, C.; McGhee, M. D. *MRS Bull.* **2005**, *30*, 37–40.
- (9) Lee, Y. L.; Lo, Y. S. *Adv. Funct. Mater.* **2009**, *19*, 604–609.
- (10) Kamat, P. V. *Acc. Chem. Res.* **2012**, *45*, 1906–1915.
- (11) Lee, H. J.; Chen, P.; Moon, S. J.; Sauvage, F.; Sivula, K.; Bessho, T.; Gamelin, D. R.; Comte, P.; Zakeeruddin, S. M.; Seok, S. I.; Gratzel, M.; Nazeeruddin, M. K. *Langmuir* **2009**, *25*, 7602–7608.
- (12) Chang, J. A.; Rhee, J. H.; Im, S. H.; Lee, Y. H.; Kim, H. J.; Seok, S. I.; Nazeeruddin, M. K.; Gratzel, M. *Nano Lett.* **2010**, *10*, 2609–2612.
- (13) Im, S. H.; Lim, C. S.; Chang, J. A.; Lee, Y. H.; Maiti, N.; Kim, H. J.; Nazeeruddin, M. K.; Gratzel, M.; Seok, S. I. *Nano Lett.* **2011**, *11*, 4789–4793.
- (14) Zeng, T. W.; Liu, I. S.; Hsu, F. C.; Huang, K. T.; Liao, H. C.; Su, W. F. *Opt. Express* **2010**, *18*, A357–A365.
- (15) Lin, C. C.; Ho, P. H.; Huang, C. L.; Du, C. H.; Yu, C. C.; Chen, H. L.; Yeh, Y. C.; Li, S. S.; Lee, C. K.; Pao, C. W.; Chu, C. P.; Chang, M. W.; Chen, C. W. *J. Phys. Chem. C* **2012**, *116*, 25081–25088.
- (16) Guerrero, A.; Boix, P. P.; Marchesi, L. F.; Ripolles-Sanchis, T.; Pereira, E. C.; Garcia-Belmonte, G. *Sol. Energy Mater. Sol. Cells* **2012**, *100*, 185–191.
- (17) Bauld, R.; Fleury, L. M.; Walsh, M. V.; Fanchini, G. *Appl. Phys. Lett.* **2012**, *101*, 103306.
- (18) Schafferhans, J.; Baumann, A.; Wagenpfahl, A.; Deibel, C.; Dyakonov, V. *Org. Electron* **2010**, *11*, 1693–1700.
- (19) Boix, P. P.; Lee, Y. H.; Fabregat-Santiago, F.; Im, S. H.; Mora-Sero, I.; Bisquert, J.; Seok, S. I. *ACS Nano* **2012**, *6*, 873–880.
- (20) Boix, P. P.; Larramona, G.; Jacob, A.; Delatouche, B.; Mora-Sero, I.; Bisquert, J. *J. Phys. Chem. C* **2012**, *116*, 1579–1587.
- (21) Lim, C. S.; Im, S. H.; Rhee, J. H.; Lee, Y. H.; Kim, H. J.; Maiti, N.; Kang, Y. K.; Chang, J. A.; Nazeeruddin, M. K.; Gratzel, M.; Seok, S. I. *J. Mater. Chem.* **2012**, *22*, 1107–1111.
- (22) Itzhaik, Y.; Niitsoo, O.; Page, M.; Hodes, G. *J. Phys. Chem. C* **2009**, *113*, 4254–4256.
- (23) Wu, S. J.; Li, J. H.; Tai, Q. D.; Yan, F. *J. Phys. Chem. C* **2010**, *114*, 21873–21877.
- (24) Hu, H. M.; Mo, M. S.; Yang, B. J.; Zhang, X. J.; Li, Q. W.; Yu, W. C.; Qian, Y. T. *J. Cryst. Growth* **2003**, *258*, 106–112.
- (25) Fabregat-Santiago, F.; Garcia-Belmonte, G.; Mora-Sero, I.; Bisquert, J. *Phys. Chem. Chem. Phys.* **2011**, *13*, 9083–9118.
- (26) Franceschetti, D. R.; Macdonald, J. R.; Buck, R. P. *J. Electrochem. Soc.* **1991**, *138*, 1368–1371.
- (27) Adachi, M.; Sakamoto, M.; Jiu, J.; Ogata, Y.; Isoda, S. *J. Phys. Chem. B* **2006**, *110*, 13872–13880.
- (28) Hodes, G. *J. Phys. Chem. C* **2008**, *112*, 17778–17787.
- (29) Mora-Sero, I.; Gimenez, S.; Fabregat-Santiago, F.; Gomez, R.; Shen, Q.; Toyoda, T.; Bisquert, J. *Acc. Chem. Res.* **2009**, *42*, 1848–1857.
- (30) Mora-Sero, I.; Gimenez, S.; Fabregat-Santiago, F.; Azaceta, E.; Tena-Zaera, R.; Bisquert, J. *Phys. Chem. Chem. Phys.* **2011**, *13*, 7162–7169.
- (31) Musselman, K. P.; Marin, A.; Schmidt-Mende, L.; MacManus-Driscoll, J. L. *Adv. Funct. Mater.* **2012**, *22*, 2202–2208.