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# Prussian blue-derived synthesis of uniform nanoflakes-assembled NiS<sub>2</sub> hierarchical microspheres as highly efficient electrocatalysts in dye-sensitized solar cells†

Shoushuang Huang,<sup>a</sup> Haitao Wang,<sup>a</sup> Yang Zhang,<sup>b</sup> Shangdai Wang,<sup>a</sup> Zhiwen Chen,<sup>a\*</sup> Zhangjun Hu<sup>a</sup> and Xuefeng Qian<sup>b\*</sup>

It's urgent and challenging to explore cost-effective and robust electrocatalyst in the development of large-scaled dye-sensitized solar cells (DSSCs). In this work, we develop a novel strategy to prepare 3D hierarchical NiS<sub>2</sub> microspheres constructed by nanoflakes through a facile chemical etching/anion exchange reaction. Nickel-cobalt Prussian blue analogous (PBA) nanocubes and (NH<sub>4</sub>)<sub>2</sub>S are employed to initially produce uniform  $\gamma$ -NiOOH/NiS<sub>x</sub> hierarchical microspheres, which were then converted to uniform 3D hierarchical NiS<sub>2</sub> microspheres by a controlled annealing treatment. Due to their favorable structural features, the as-obtained NiS<sub>2</sub> hierarchical microspheres possess large surface areas, high structural void porosity and accessible inner surface. All of these advantages facilitate the mass diffusion and charge transport between electrolyte and counter electrode material. As a result, the titled NiS<sub>2</sub> hierarchical microspheres exhibit excellent electrocatalytic activity toward the reduction of I<sub>3</sub><sup>−</sup> ions in DSSCs. A typical DSSC with NiS<sub>2</sub> achieves an impressive power conversion efficiency of 8.46% under AM1.5G illumination (100 mW cm<sup>−2</sup>), higher than that of pyrolysis Pt electrodes (8.04%). Moreover, the fast activity onset and relatively long stability further demonstrate that the NiS<sub>2</sub> hierarchical microspheres are promising alternatives to Pt in DSSCs.

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## 1. Introduction

Dye-sensitized solar cells (DSSCs) have received extensive research interest of becoming the third generation solar cells due to their facile fabrication processes, low-cost, and relatively high photoelectric conversion efficiency.<sup>1–3</sup> A typical DSSC consists of a dye-sensitized TiO<sub>2</sub> photoanode, a redox electrolyte containing I<sub>3</sub><sup>−</sup>/I<sup>−</sup> redox couples, and a counter electrode (CE). As one of the most pivotal components, the CE is responsible for collecting electrons from external circuits and then reducing the I<sub>3</sub><sup>−</sup> to I<sup>−</sup> ions.<sup>4</sup> Recently, noble metal platinum (Pt) is the most preferred CE electrocatalyst due to its high conductivity and outstanding electrocatalytic activity. However, Pt is of high cost and scarcity and thereby seriously limits its practical application. Additionally, Pt could be easily corroded by iodine-based electrolyte,<sup>5</sup> which would inevitably deteriorate the photovoltaic performance and the long-term stability of DSSCs.

Therefore, the development of earth-abundant and active electrocatalyst with high stability is urgently needed to realize the commercial application of DSSCs.

Recently, considerable progress has been achieved on the developments of alternative materials to Pt CE, such as carbon materials,<sup>6–9</sup> conductive polymers,<sup>10,11</sup> metal alloys,<sup>12,13</sup> and transitional-metal compounds.<sup>14,15</sup> Among them, nickel sulfide with a variety of chemical formulas is found to be promising CE material for the I<sub>3</sub><sup>−</sup> reduction reaction in DSSCs.<sup>16–19</sup> For example, the hierarchal Ni<sub>3</sub>S<sub>4</sub> microspheres reported by Yang *et al.* displayed a power conversion efficiency (PCE) of 6.81% under AM1.5G illumination, comparable to that of Pt-based DSSC (6.85%).<sup>16</sup> Additionally, no obvious degradation of photovoltaic performance was observed in 504 hours, indicating the high stability of nickel sulfide in iodine-based electrolyte. Zheng *et al.* found that both of octahedral and cubic NiS<sub>2</sub> nanocrystals exhibited excellent catalytic activity toward the reduction of I<sub>3</sub><sup>−</sup>. The corresponding DSSCs with octahedral NiS<sub>2</sub> and cubic NiS<sub>2</sub> achieved a high PCE of 5.98% and 5.43%, respectively.<sup>20</sup> The well-defined NiS/Ni<sub>3</sub>S<sub>2</sub> nanorod composite array prepared by Pan's group also manifested a high catalytic activity for the regeneration of I<sub>3</sub><sup>−</sup> ions, and attained comparable photovoltaic performances with Pt CE.<sup>21</sup> These extraordinary electrochemical performance of DSSCs depends on the

<sup>a</sup>School of Environmental and Chemical Engineering, Shanghai University, Shanghai 200444, China. E-mail: zwchen@shu.edu.cn

<sup>b</sup>Shanghai Electrochemical Energy Devices Research Center, School of Chemistry and Chemical Engineering, State Key Laboratory of Metal Matrix Composites, Shanghai Jiao Tong University, Shanghai 200240, China

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fascinating morphologies, rich redox reactions, and diverse photoelectric properties of nickel sulfide CEs. Therefore, taking into consideration the high earth abundance, the superior catalytic activity and the excellent chemical stability, nickel sulfide is a promising CE candidate for the large-scale manufacture of low-cost DSSCs.

It's well known that the architecture is one of the key factors in the determination of electrochemical performance of electrode materials. Recently, two-dimensional (2D) nanomaterial has been considered as one kind of ideal morphological foundation for the surface-dependent electrochemical reactions due to the high percentages of surface atoms, abundant active sites and high exposures of low-energy facets.<sup>22,23</sup> However, the major problem associated with the application of 2D materials in catalysis lies in their insufficient number of internal spaces caused by their direct and free accumulation on devices.<sup>24</sup> In contrast, 3D hierarchical architectures provide the potential of combining the merits of low-dimensional material in a single structure, such as higher surface/volume ratio, larger porosity and accessible interior space.<sup>25</sup> These characteristics can facilitate the filling and diffusion of electrolyte, shorten the diffusion distance as well as increase the catalytic active sites, which are important to boost the photovoltaic performances of DSSCs. On the basis of the above investigations, it seems logical to envision that 3D hierarchical NiS<sub>2</sub> architectures constructed by 2D building blocks can be favorable for constructing high performance of DSSCs. Up to now, typical strategy for fabricating NiS<sub>2</sub> superstructures is template-engaged synthesis.<sup>26–28</sup> Therefore, the fabrication of 3D hierarchical NiS<sub>2</sub> superstructures using simple wet chemical methods under mild conditions is still relatively challenging.

Recently, chemical etching has been regarded as an efficient method to prepare complex nanostructures with unique physicochemical properties, and a series of high-quality metal oxides/sulfides complex nanostructure have been successfully fabricated by selective etching of Prussian blue analogues (PBA).<sup>29–31</sup> Inspired by them, in the present work, nanoflakes-assembled NiS<sub>2</sub> hierarchical microspheres were synthesized through a facile chemical etching/anion exchange reaction. Nickel–cobalt Prussian blue analogous (PBA) nanocubes and (NH<sub>4</sub>)<sub>2</sub>S are employed to initially produce uniform  $\gamma$ -NiOOH/NiS<sub>x</sub> hierarchical microspheres, which are then converted to uniform 3D hierarchical NiS<sub>2</sub> microspheres *via* a controlled annealing treatment. The as-obtained NiS<sub>2</sub> hierarchical microspheres combine the merits of 2D and 3D materials when employed as CE catalysts in DSSCs: (1) the 2D nanoflakes provide larger electrochemical active sites and shorter electron diffusion path to catalyze the reduction of I<sub>3</sub><sup>−</sup>. (2) The 3D hierarchical architectures offer a higher surface/volume ratio, larger porosity and accessible inner surface to facilitate the diffusion of electrolyte. (3) The 3D hierarchical structures provide wide spaces for acetylene black filling in and make the microspheres contact more efficiently to conductive substrate, which helps to reduce the resistance of devices. As a result, the NiS<sub>2</sub> hierarchical microspheres exhibited low charge transfer resistance at the electrolyte–electrode interface, high electrocatalytic activity and fast reaction kinetics for the I<sup>−</sup>/I<sub>3</sub><sup>−</sup> redox reaction.

## 2. Experiment section

### 2.1 Synthesis of NiCo PBA nanocubes

The NiCo PBA nanocubes were synthesized through a previously reported method with slight modification.<sup>32</sup> In brief, 3.0 mmol of Ni(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O and 4.5 mmol of Na<sub>3</sub>C<sub>6</sub>H<sub>5</sub>O<sub>7</sub>·2H<sub>2</sub>O were dissolved in 100 mL of deionized water under vigorous stirring to form a clear solution. Then, this green solution was rapidly injected into 100 mL of aqueous solution containing 2.0 mmol of K<sub>3</sub>[Co(CN)<sub>6</sub>]. The obtained mixed solution was allowed to age for 24 h at room temperature. Finally, the product was collected by centrifugation and washed with water and ethanol for several times, and dried at 80 °C in a vacuum oven for 12 h.

### 2.2 Synthesis of 3D hierarchical NiS<sub>2</sub> microspheres

In a typical synthesis, 60 mg of the above as-prepared NiCo PBA nanocubes were dispersed into 60 mL of ethanol to form a homogeneous suspension with the assistance of ultrasonication for 30 min. At the meantime, 3.0 mL (NH<sub>4</sub>)<sub>2</sub>S (≥8.0%) aqueous solution was dissolved into 30 mL of deionized water. The (NH<sub>4</sub>)<sub>2</sub>S solution was then poured into the above NiCo PBA suspension under magnetic stirring for 30 min. Finally, the resulting mixture was transferred into a Teflon-lined stainless-steel autoclave and kept at 100 °C for 10 hours in an electric oven. The final products were collected by centrifugation, and washed with water and ethanol before drying at 80 °C overnight. Then, 30 mg of the as-obtained sample was placed at the downstream side of crucible, and 120 mg of sublimed sulfur powder was placed at the upstream side of the crucible. The crucible was subsequently placed into the tube furnace and calcined under nitrogen atmosphere at 350 °C for 2 hours with a ramp rate of 2 °C min<sup>−1</sup>. After cooling down to room temperature, 3D NiS<sub>2</sub> hierarchical microspheres were obtained.

### 2.3 Preparation of counter electrodes

CEs were prepared by a slurry coating procedure on the cleaned FTO substrates. The slurry was prepared by mixing 80 mg of NiS<sub>2</sub> catalyst, 10 mg of acetylene black and 10 mg of polyvinylidene fluoride into 1.0 mL *N*-methyl-2-pyrrolidone followed by sonication for 30 min to form a homogeneous ink. The resulting slurry was then coated onto the FTO substrate using a doctor blade method. The area of active layer was masked by the transparent adhesive tape layers tape with an aperture area of 0.5 × 0.5 cm<sup>2</sup>, and the thickness of the CE film was strictly controlled to be 4 μm. Finally, the electrodes were dried at 60 °C for 12 hours in a vacuum oven. For comparison, pyrolytic Pt CE was prepared by drop-casting 50 μL of H<sub>2</sub>PtCl<sub>6</sub> in ethanol (5 mM) on a 1.5 × 1.5 cm<sup>2</sup> FTO glass followed by sintering at 400 °C for 30 min.

### 2.4 Fabrication of DSSCs

The commercial TiO<sub>2</sub> photoanodes (Ying kou Opvtech New Energy Co., Ltd) were first annealed at 500 °C for 1 hour. After being cooled down to 80 °C, the TiO<sub>2</sub> photoanodes were immersed in a 0.5 mM ethanol solution of N719 dye (Solaronix



SA, Switzerland) for 20 h. DSSCs were fabricated by assembling the N719-sensitized TiO<sub>2</sub> photoanodes and CE with a hot-melt film as spacer (Surlyn 1702; DuPont, 60 μm in thickness). The cell internal space was filled with electrolyte using a vacuum pump followed by a sealing procedure with a Surlyn film and a cover glass under heating. The liquid electrolyte was composed of 0.60 M 1-butyl-3-methylimidazolium iodide, 0.03 M I<sub>2</sub>, 0.50 M 4-*tert*-butyl pyridine, 0.06 M lithium iodide and 0.10 M guanidinium thiocyanate of anhydrous acetonitrile. The sealed DSSCs were used for the photocurrent–voltage test with an active area of 0.16 cm<sup>2</sup>.

## 2.5 Characterization

The phase and crystal structure of the as-obtained products was characterized by powder X-ray diffraction (XRD) (Shimadzu XRD-6000) with Cu Kα radiation. The morphologies and crystal lattice of the samples were characterized by field-emission scanning electron microscopy (FESEM, JEOL, JSM-6700F) and high-resolution transmission electron microscopy (HRTEM, JEOL, JEM-2100F), respectively. The X-ray photoelectron spectroscopy (XPS) analysis was performed on a VG Scientific ESCLAB 220iXL X-ray photoelectron spectrometer with Al Kα (1486.6 eV) as the X-ray source. The surface area and porosity of the materials was analyzed by using a NOVA 2200e instrument. The photocurrent–voltage tests of DSSCs were performed under AM 1.5 illumination (100 mW cm<sup>−2</sup>) in ambient condition on a 94023A Oriel sol3A solar simulator (Newport) with 450 W xenon lamp as light source. Electrochemical impedance spectroscopy (EIS) experiments and Tafel-polarization curves were measured with dummy cells in the dark by using a Zahner electrochemical workstation (Zahner Zennium CIMPS-1, Germany). The frequency range of EIS experiments was from 100 mHz to 1 MHz with an AC modulation signal of 10 mV and bias DC voltage of 0 V. The resultant impedance spectra were analyzed *via* the Z-view software. Tafel-polarization measurements were carried out with a scan rate of 5 mV s<sup>−1</sup>. Cyclic voltammetry was conducted in a three-electrode system in an acetonitrile solution of 0.1 mmol LiClO<sub>4</sub>, 10 mmol LiI and 1 mmol I<sub>2</sub> at a scan rate of 50 mV s<sup>−1</sup>. Platinum nets electrode served as a CE, and the Ag/AgCl couple was used as a reference electrode.

## 3. Results and discussion

Field emission scanning electron microscopy (FESEM) images showed that the as-prepared Ni<sub>3</sub>[Co(CN)<sub>6</sub>]<sub>2</sub> PBA nanocubes had a highly uniform and well-defined cubic structure with an average size of 242 nm (Fig. 1a and b), and the corresponding transmission electron microscopy (TEM) image indicated these PBA nanocubes presented a solid and dense texture (Fig. 1c). The NiCo PBA nanocubes were then employed as nickel source to react with (NH<sub>4</sub>)<sub>2</sub>S aqueous solution at 100 °C for 10 hours. FESEM images clearly revealed that the as-obtained product was composed of uniform and monodispersed microspheres with an average diameter of 2.0 μm (Fig. 1d). An enlarged SEM image indicated that each microsphere was constructed by 2D

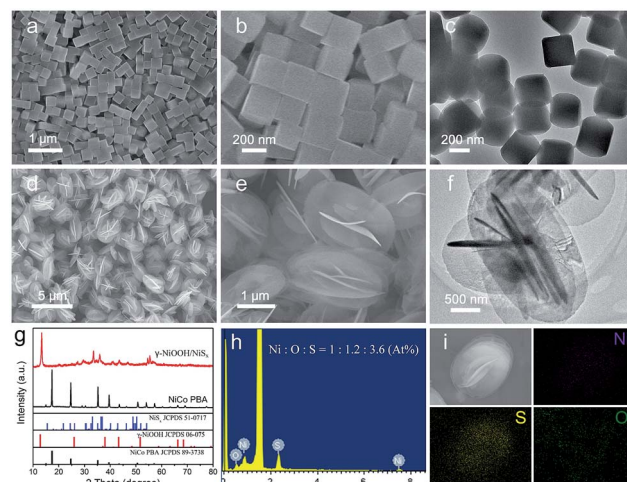


Fig. 1 SEM (a, b, d, e) and TEM (c, f) images of the as-prepared NiCo PBA nanocubes (a–c) and γ-NiOOH/NiS<sub>x</sub> (d–f) microspheres; XRD patterns (g), EDX spectrum (h) and element mapping (i) of the γ-NiOOH/NiS<sub>x</sub> hierarchical microspheres.

nanoflakes building blocks in self-assemble style (Fig. 1e). The thickness of the nanoflake was in the range of 17–28 nm determined by TEM image (Fig. 1f). Fig. 1g showed the X-ray diffraction pattern (XRD) of the as-synthesized sample. The crystallographic phase of the product could be identified as γ-NiOOH (JCPDS 06-075) and NiS<sub>x</sub> (JCPDS, 51-0717), respectively. Energy dispersive X-ray spectroscopy (EDX) showed the Ni : O : S atomic ratio to be 1 : 1.2 : 3.6 (Fig. 1h), and the corresponding elemental mapping images indicated that these elements were uniformly distributed in a single microsphere (Fig. 1i). From the results mentioned above, we concluded that the as-obtained product was composed of γ-NiOOH and NiS<sub>x</sub>.

In the present synthesis strategy, the NiCo PBA nanocubes as nickel source was critical for the formation of uniform 3D hierarchical γ-NiOOH/NiS<sub>x</sub> microspheres. Replacing NiCo PBA nanocubes with nickel nitrate or nickel acetate while keeping the other synthesis parameters similar, irregular nano/micro-sized nickel sulfide particles with large size-distribution were obtained (Fig. S1†). This result can be explained by the fact that the excessively release of Ni<sup>2+</sup> cations from metal salts lead to the fast nucleation and growth of nickel sulfides.<sup>33</sup> Further study indicated that the formation of uniform γ-NiOOH/NiS<sub>x</sub> hierarchical microspheres also depended upon reaction temperature. When the reaction temperature was over 140 °C, NiS<sub>2</sub> nanoparticles (NPs) with average size of ~200 nm were obtained (Fig. S2†). In contrast, nearly no product could be obtained when the reaction was carried out at room temperature. The above findings demonstrated that the precise control over reaction rate was crucial for the formation of high-quantity γ-NiOOH/NiS<sub>x</sub> hierarchical microspheres.<sup>29</sup> Therefore, a proper amount of ethanol was introduced into the reaction system to slow down the etching of NiCo PBA nanocubes as well as the nucleation of nickel sulfides (Fig. S3†).

To investigate the structural evolution from NiCo PBA nanocubes to the γ-NiOOH/NiS<sub>x</sub> hierarchical nanostructures, a series of experiments with different reaction times were





conducted. In order to further slow down the reaction rate, the reaction were carried out in 90 mL ethanol and 10 mL H<sub>2</sub>O. The structure and morphologies of the intermediate products collected at different reaction stages were shown in Fig. 2. After a reaction time of 0.5 hour, the edges of the NiCo PBA nanocubes were etched, while their plane surfaces remained unchanged (Fig. 2b and c). This phenomenon indicated the different reaction activity between the edge and plane parts, which is in good agreement with Yu's report.<sup>29,31</sup> When the reaction was carried out for 1.0 hour, the NiCo PBA nanocubes disappeared and thin nanoflakes were observed (Fig. 2d). This nanoflakes were identified mainly to be  $\gamma$ -NiOOH by XRD (Fig. S4†). At a longer reaction time of 10 hours, the nanoflakes were connected with each other and self-assembled into uniform hierarchical microspheres (Fig. 2e).

Based on the experimental results mentioned above, a possible growth mechanism for the  $\gamma$ -NiOOH/NiS<sub>x</sub> hierarchical microspheres was proposed. The double hydrolysis of (NH<sub>4</sub>)<sub>2</sub>S produced NH<sub>3</sub>·H<sub>2</sub>O and H<sub>2</sub>S gas in solution, which were further hydrolyzed to hydroxide ion (OH<sup>-</sup>) and polysulfide ions (S<sub>x</sub><sup>2-</sup>).<sup>34</sup> Thus, the NiCo PBA nanocubes were easily etched by these species. As more defects were existed on the rough edges, the etching reaction preferentially took place on the edges of NiCo PBA nanocubes.<sup>29</sup> In this way,  $\gamma$ -NiOOH was easily formed due to the reaction between Ni<sup>2+</sup> and OH<sup>-</sup> ions. At the same time, the  $\gamma$ -NiOOH transition continuously converted into NiS<sub>x</sub> driven by the thermodynamic equilibrium due to the larger solubility product constant of hydroxides compared with sulfide (Fig. S4†). The EDX spectra showed the decrease of oxygen content and the increase of sulfur content as the reaction proceeds, evidencing the anion exchange between OH<sup>-</sup> and S<sup>2-</sup> (Fig. S5†). Similar phenomenon has been observed for many other transition metal hydroxides by matching the solubility product of their hydroxides and sulfides.<sup>35,36</sup>

Recently, many transition metal hydroxides have been utilized as precursors to prepare novel functional chalcogenides with satisfied performance. Herein, we converted the  $\gamma$ -NiOOH/NiS<sub>x</sub> microspheres to nickel sulfides *via* thermal decomposition and sulfuration reaction. As shown in Fig. 3a, all of the diffraction peaks were well indexed to cubic NiS<sub>2</sub> (JCPDS 65-3325, *Pa* $\bar{3}$ , *a* = *b* = *c* = 5.678 Å). No peaks from other crystalline

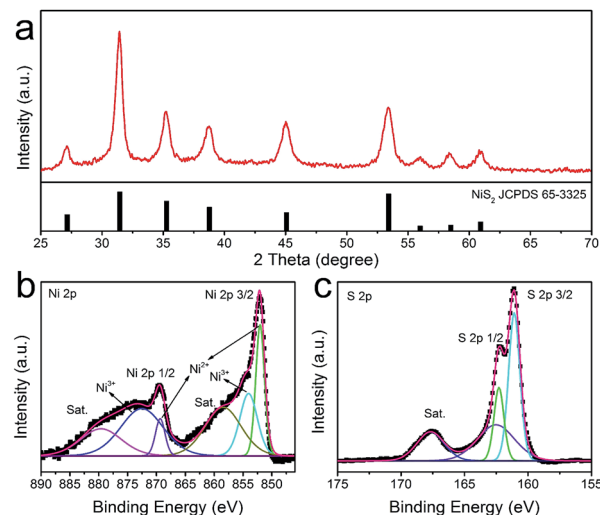


Fig. 3 XRD (a) and XPS spectrum of the as-prepared NiS<sub>2</sub> hierarchical microspheres: (b) Ni 2p and (c) S 2p.

byproducts could be identified, indicating the hydroxides phase was completely decomposed and converted into metal sulfide. X-ray photoelectron spectroscopy (XPS) was further used to analyze the valence state and the surface composition of the as-obtained product. By using a Gaussian fitting method, the Ni 2p spectrum can be deconvoluted into two spin orbit doublets and two shakeup satellites (identified as "Sat.") (Fig. 3b). It was found that the strong peaks at the binding energies of 851.9 and 869.4 eV could be attributed to 2p<sub>3/2</sub> and 2p<sub>1/2</sub> of Ni<sup>2+</sup> species, and the peaks at 854.1 and 872.5 eV could be assigned to 2p<sub>3/2</sub> and 2p<sub>1/2</sub> of Ni<sup>3+</sup> species, respectively.<sup>37</sup> The existence of Ni<sup>3+</sup> indicated the slight surface oxidation state of the sample, which was consistent with previous reports.<sup>38</sup> The XPS spectrum of S 2p was shown in Fig. 3c. The peaks at 161.1 and 162.3 eV were associated with S 2p<sub>3/2</sub> and S 2p<sub>1/2</sub>, respectively. Similarly, the minor peak near 162.5 eV suggested the presence of higher oxidation state sulfide species in the sample.<sup>39</sup> The XPS results were in agreement with XRD, confirming that the as-obtained sample was indeed composed of pure NiS<sub>2</sub>.

The morphology details of the annealed sample was then investigated by SEM and TEM. As one can see in Fig. 4a, the hierarchical structure was basically maintained after calcination. However, the nanoflakes of the NiS<sub>2</sub> microspheres became porous with rough surfaces in contrast to the smooth surfaces of the hydroxides precursor (Fig. 4b). A typical TEM image in Fig. 4c indicated the nanoflake was composed of nanocrystals with size of several nanometers. The lattice fringes were clearly displayed in a high-resolution TEM image and the interplanar spacing of 2.83 Å and 2.53 Å could be ascribed to the (200) and (210) planes of cubic-phase NiS<sub>2</sub>, respectively (Fig. 4d). The selected area electron diffraction (SAED) confirmed the polycrystalline nature of the annealed sample (Fig. 4e). In virtue of the 3D hierarchal structure, the NiS<sub>2</sub> microspheres possessed a relatively high Brunauer–Emmett–Teller (BET) surface area of about 69.8 m<sup>2</sup> g<sup>-1</sup> (Fig. S6†). It's clear that this hierarchical structure is beneficial for mass transport and fast diffusion of the electrolyte during the electrocatalysis process.

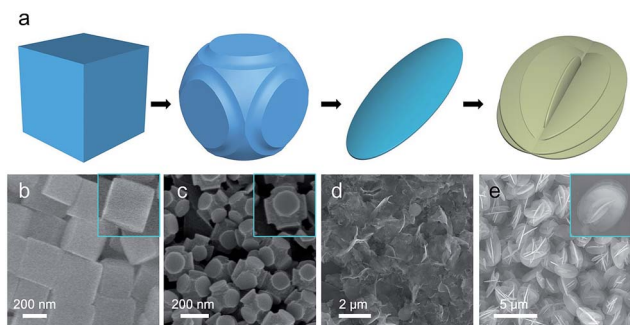


Fig. 2 Schematic illustration of the formation process of  $\gamma$ -NiOOH/NiS<sub>x</sub> hierarchical microspheres (a). FESEM images of the products obtained after reaction for (b) 0 hour, (c) 0.5 h, (d) 1.0 hour, and (e) 10 hour.



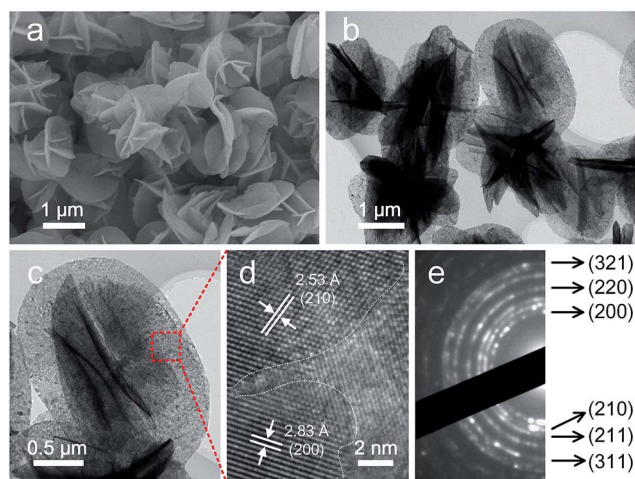


Fig. 4 SEM (a), TEM (b, c), HRTEM (d) images and SAED pattern (e) of the as-obtained NiS<sub>2</sub> microspheres after annealing at 350 °C for 2 hours under nitrogen atmosphere.

Sandwich-structure devices were assembled to assess the potential application of NiS<sub>2</sub> hierarchical microspheres in DSSCs. The current density–voltage (*J*–*V*) curves were shown in Fig. 5a and the related photovoltaic parameters including short circuit current density (*J*<sub>sc</sub>), open circuit voltage (*V*<sub>oc</sub>), power conversion efficiency (PCE) and fill factor (FF), were summarized in Table 1 and S1.† The DSSC with NiS<sub>2</sub> microspheres (MSs) CE produced a *V*<sub>oc</sub> of 746 mV, a *J*<sub>sc</sub> of 16.23 mA cm<sup>−2</sup> and a FF of 0.70, thus yielding an overall PCE of 8.48%. This was an improved photovoltaic performance compared to that of the device employing Pt CE (*V*<sub>oc</sub> = 761 mV, *J*<sub>sc</sub> = 15.86 mA cm<sup>−2</sup>, FF = 0.65,  $\eta$  = 7.84%). Comparing these parameters, we can find that the enhancement of PCE was mainly contributed by the improvement of *J*<sub>sc</sub> and FF. The increased *J*<sub>sc</sub> can be attributed to the accelerated reduction of I<sub>3</sub><sup>−</sup> ions by NiS<sub>2</sub> microspheres, which subsequently resulted in the fast regeneration of N719 molecules.<sup>40</sup> The improvement of FF corresponded to the reduced charge-transfer resistance at the CE/electrolyte interface and diffusion resistance of I<sub>3</sub><sup>−</sup> in the electrolyte. Additionally, to demonstrate the structural advantage of 3D hierarchical structures in catalysis, the obtained irregular NiS<sub>2</sub> nanoparticles (NPs) were also used as CE catalyst for comparison. It was found that the performance of the device with NiS<sub>2</sub>

Table 1 Photovoltaic performance of DSSCs with various CEs under AM1.5G illumination (100 mW cm<sup>−2</sup>)

CEs	<i>V</i> <sub>oc</sub> (mV)	<i>J</i> <sub>sc</sub> (mA cm <sup>−2</sup> )	FF	PCE (%)
NiS <sub>2</sub> MSs	745 ± 3.77	16.28 ± 0.04	0.70 ± 0.01	8.46 ± 0.06
NiS <sub>2</sub> NPs	745 ± 2.16	15.27 ± 0.03	0.68 ± 0.00	7.76 ± 0.04
Pt	741 ± 3.87	15.66 ± 0.11	0.69 ± 0.01	8.04 ± 0.05

NPs CE was significantly inferior to those with NiS<sub>2</sub> MSs and Pt CEs. This observation demonstrated the higher catalytic activity of 3D nanostructure based CEs due to the numerous catalytic active sites exposed on the surface. These results were also supported by the IPCE measurement, in which the NiS<sub>2</sub> MSs based device exhibited a highest efficiency than the other two CEs (Fig. 5b). The IPCE followed the same trend with the difference of PCE and *J*<sub>sc</sub> values, confirming the as-prepared NiS<sub>2</sub> hierarchical microspheres CE could provide sufficient and fast regeneration of I<sub>3</sub><sup>−</sup> ions in the electrolyte.<sup>41</sup>

To evaluate the electrocatalytic activities of the as-prepared CEs toward the reduction of I<sub>3</sub><sup>−</sup> ions, cyclic voltammetry (CV) measurements were carried out in a three-electrode system. As shown in Fig. 6a, all electrodes exhibited two typical pairs of oxidation–reduction peaks, suggesting that the NiS<sub>2</sub> CEs can exhibit Pt-like electrocatalytic behavior. The left peak in low potential range is assigned to the oxidation (Ox-1) and reduction (Red-1) of I<sup>−</sup>/I<sub>3</sub><sup>−</sup>, and the right one in high potential range corresponds to the oxidation (Ox-2) and reduction (Red-2) of I<sub>2</sub>/I<sub>3</sub><sup>−</sup>.<sup>42</sup> The peak current density of Red-A and peak-to-peak separation (*E*<sub>pp</sub>) between Red-1 and Ox-1 are the two important parameters for evaluating the catalytic activities of different CEs.<sup>43</sup> The NiS<sub>2</sub> MSs CE produced a higher peak current density (1.76 mA cm<sup>−2</sup>) than that of NiS<sub>2</sub> NPs (1.48 mA cm<sup>−2</sup>) and Pt CE (1.63 mA cm<sup>−2</sup>), indicating it had better electrocatalytic activity for the reduction of I<sub>3</sub><sup>−</sup> ions. Additionally, the *E*<sub>pp</sub> of NiS<sub>2</sub> MPs CE (330 mV) was also smaller than those of NiS<sub>2</sub> NPs (482 mV) and Pt CE (343 mV), which suggested a smaller overpotential for the reduction reaction on NiS<sub>2</sub> MSs CE.<sup>44</sup> The higher peak current density and lower *E*<sub>pp</sub> value indicated the NiS<sub>2</sub> MSs CE presented higher catalytic activity for the reduction of I<sub>3</sub><sup>−</sup> to I<sup>−</sup>. The excellent catalytic activity of NiS<sub>2</sub> MPs CE can be attributed to its large active surface area and abundant hierarchical pore structure, which provided more catalytic activity sites and the higher charge transfer rate. Additionally, the stacking CV plots of NiS<sub>2</sub> MPs CE were recorded at a scan rate from 10 to 150 mV s<sup>−1</sup> (Fig. S7†). The good linear relationships between the peak current density and the square root of corresponding scan rate implied the reduction process of I<sup>−</sup>/I<sub>3</sub><sup>−</sup> redox couples at NiS<sub>2</sub> CE obeyed a diffusion-controlled mechanism, and there was no specific interactions between NiS<sub>2</sub> CE and I<sup>−</sup>/I<sub>3</sub><sup>−</sup> redox couples.<sup>45</sup>

Electrochemical impedance spectroscopy (EIS) measurements were further used to investigate the internal resistance and charge transfer kinetics at the CE/electrolyte interface. The Nyquist plots for symmetric cells (CE/electrolyte/CE) were shown in Fig. 6b and the fitted results were listed in Table 2. The intercept on the real axis can be assigned to the ohmic series

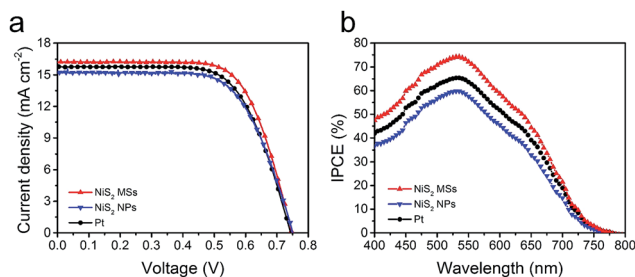


Fig. 5 *J*–*V* curves (a) and IPCE spectrum (b) of the DSSCs using NiS<sub>2</sub> MSs, NiS<sub>2</sub> NPs and Pt CEs, measured at AM1.5G illumination.



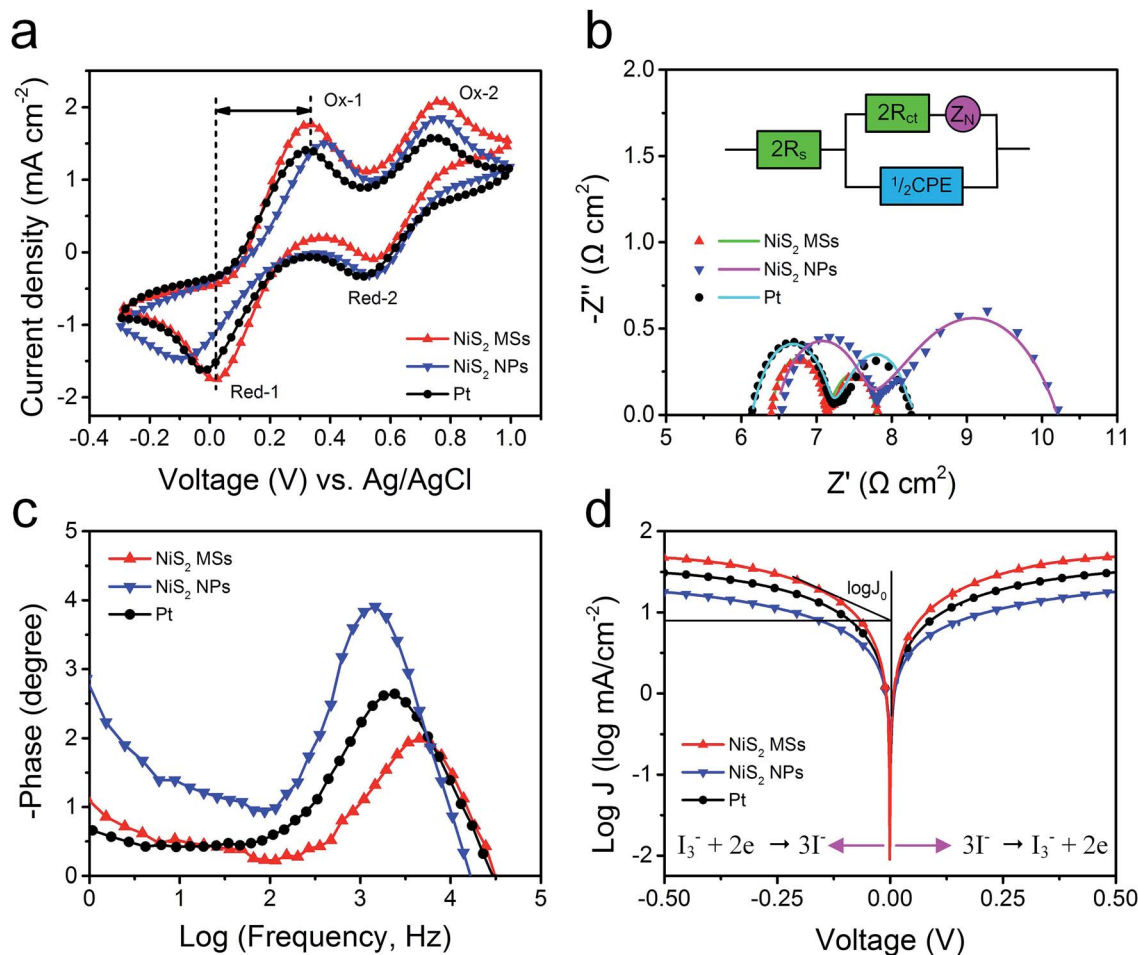


Fig. 6 (a) Cyclic voltammograms measured in an iodine-based electrolyte with a scanning rate of  $50 \text{ mV s}^{-1}$ . (b) Nyquist and (c) Bode EIS plots, and (d) Tafel polarization curves for the symmetric dummy cells fabricated by two identical CEs. The inset shows an equivalent circuit simulated by Z-view software.

Table 2 The electrochemical parameters for different CEs

CEs	$R_s$ ( $\Omega \text{ cm}^2$ )	$R_{ct}$ ( $\Omega \text{ cm}^2$ )	CPE-T ( $\mu\text{F cm}^{-2}$ )	CPE-P	$W_s - R$ ( $\Omega \text{ cm}^2$ )	$W_s - T$	$W_s - P$	$\tau$ ( $\mu\text{s}$ )	$E_{pp}$ (mV)
NiS <sub>2</sub> MSs	3.23	0.37	259.2	0.81	0.68	0.011	0.50	31.7	330
NiS <sub>2</sub> NPs	3.27	0.63	96.4	0.93	2.44	0.005	0.50	116.9	482
Pt	3.09	0.55	71.6	0.88	1.08	0.015	0.50	71.4	343

resistance ( $R_s$ ). The semicircles in frequency regions between 1 kHz and 100 kHz can be attributed to the resistance capacitance of the electrode/electrolyte interface, including the charge transfer resistance ( $R_{ct}$ ) and the corresponding double-layer capacitance ( $C_{\mu}$ ). The arc at low frequency regions can be assigned to the Warburg impedance ( $W_s$ ) in the electrolyte.<sup>46</sup> Typically,  $R_{ct}$  is regarded as an indicator to reveal the charge transfer process and thereby to evaluate the catalytic activity of catalysts. Constant phase element (CPE) is frequently used as a substitute for a capacitor in an equivalent circuit to fit the impedance behavior of the electrical double layer, which can be defined as:  $\text{CPE} = (\text{CPE-T})^{-1}(\omega)^{-(\text{CPE-P})}$ , where  $j^2 = -1$ ,  $\omega$  is the frequency and CPE-T, CPE-P are the frequency-independent

parameters of the CPE.<sup>47,48</sup> Generally, a larger CPE-T means an increase in the active surface area, and a decrease of CPE-P means an increase in the porosity. Additionally, the Warburg impedance equals:  $W_s = R \tanh([j\omega T]^P)(j\omega T)^{-P}$ , in which  $P = 0.5$ ,  $R$  is the diffusion impedance and  $T$  is the relationship of the effective diffusion thickness and the effective diffusion coefficient.<sup>49</sup> These electrochemical parameters can be obtained by fitting EIS spectra using a Z-view software.

The NiS<sub>2</sub> CEs yielded similar  $R_s$  value with Pt CE due to the addition of acetylene black, hence posing similar influence on the photovoltaic performances. The simulated  $R_{ct}$  value of NiS<sub>2</sub> MSs CE was  $0.37 \Omega \text{ cm}^2$ , smaller than those of NiS<sub>2</sub> NPs ( $0.63 \Omega \text{ cm}^2$ ) and Pt CE ( $0.55 \Omega \text{ cm}^2$ ), which suggested that the





NiS<sub>2</sub> MSs CE had better electrocatalytic activity for the reduction of I<sub>3</sub><sup>−</sup> ions. The simulated CPE-T values were 96.4 μF cm<sup>−2</sup> for Pt CE and 71.6 μF cm<sup>−2</sup> for NiS<sub>2</sub> NPs CE, both of which were much lower than the value of 259.2 μF cm<sup>−2</sup> obtained for the NiS<sub>2</sub> MSs CE. These results demonstrated the NiS<sub>2</sub> MSs CE had larger specific surface area and the higher electrolyte penetration in the hierarchical microstructure.<sup>50</sup> In addition, the diffusion impedance for the symmetric cells increased by a sequence of NiS<sub>2</sub> MSs (0.68 Ω cm<sup>2</sup>) < Pt (1.08 Ω cm<sup>2</sup>) < NiS<sub>2</sub> NPs (2.44 Ω cm<sup>2</sup>), revealing a faster diffusion velocity of the redox species in the electrolyte for NiS<sub>2</sub> MSs CE. The main role of CE is to reduce the I<sub>3</sub><sup>−</sup> to I<sup>−</sup>, thus, the actual lifetimes of electrons (τ) participating in the I<sub>3</sub><sup>−</sup> reduction reaction can also be used to evaluate the catalytic activity of different CEs, which can be calculated according to formula:<sup>51</sup>  $\tau = 1/2\pi f_p$ , in which  $f_p$  is the peak of the high-frequency region in the Bode spectra (Fig. 6c). The order of the calculated τ value is 31.7 μs (NiS<sub>2</sub> MSs) < 71.4 μs (Pt) < 116.9 μs (NiS<sub>2</sub> NPs). Apparently, the shorter lifetime in NiS<sub>2</sub> MSs CE implied it had higher electrocatalytic activity for the reduction of I<sub>3</sub><sup>−</sup> ions than that of Pt CE, which was consistent with the results derived from the CV data and photovoltaic parameters.

EIS analysis of complete DSSCs were further studied under light illumination of 100 mW cm<sup>−2</sup> in the frequency range of 0.1 Hz to 100 kHz using an applied bias of V<sub>oc</sub> (Fig. S8†). The charge transport impedances ( $R_t$ ) at the CE/electrolyte interface for various CEs were higher than those measured with symmetric cells, but the value for NiS<sub>2</sub> MSs CE (0.84 Ω cm<sup>2</sup>) was still smaller than the Pt (1.12 Ω cm<sup>2</sup>) and NiS<sub>2</sub> NPs (1.39 Ω cm<sup>2</sup>) CEs. This result confirmed the NiS<sub>2</sub> MSs CE has the best electrocatalytic activity for the reduction of I<sub>3</sub><sup>−</sup>. Additionally, CPE magnitude of CE (CPE) increased in the order of NiS<sub>2</sub> NPs (52.15 mF cm<sup>−2</sup>) < Pt (144.42 mF cm<sup>−2</sup>) < NiS<sub>2</sub> MSs (278.80 mF cm<sup>−2</sup>), suggesting a same order of the active surface area. The conclusions for the catalytic activity derived from the EIS data with symmetric dummy cells and complete DSSCs were consistent.

To further examine the interfacial charge-transfer behaviors of the I<sub>3</sub><sup>−</sup>/I<sup>−</sup> couple on CE surface, Tafel polarization measurements were carried out on the symmetric cells used in the EIS experiments. Tafel curve can be divided into three zones depending on the overpotential value: the polarization zone at low overpotential ( $|V| < 120$  mV), Tafel zone at intermediate overpotential (with a sharp slope), and diffusion zone at high overpotential. In the Tafel zone, the intersection of the cathodic branch and the equilibrium potential line can be regarded as  $J_0$ , which can be obtained by extending the line to zero voltage and measuring the intercept on the  $J$ -axis ( $y$ -axis). As shown in Fig. 6d, the slope of a tangent for NiS<sub>2</sub> MSs CE was slightly higher than that of NiS<sub>2</sub> NPs and Pt CE, implying the NiS<sub>2</sub> MSs CE had larger exchange current densities and thereby better catalytic activity. The  $J_0$  value can also be calculated from the formula:<sup>52</sup>  $J_0 = RT/nFR_{ct}$ , where  $R$  is the gas constant,  $T$  is the absolute temperature,  $F$  is Faraday's constant and  $R_{ct}$  is charge transfer resistance obtained from EIS plots. The order of the calculated  $R_{ct}$  values was NiS<sub>2</sub> MSs < Pt < NiS<sub>2</sub> NPs, which was consistent with the results from the EIS analysis. In the diffusion zone, the intersection of the cathodic branch with the  $y$ -axis can be regarded as limited current density ( $J_{lim}$ ),

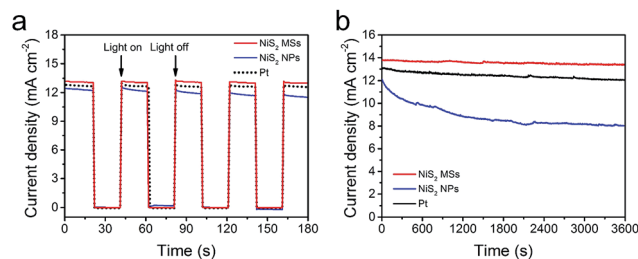


Fig. 7 Start-stop switches (a) and photocurrent stabilities (b) of the DSSCs using NiS<sub>2</sub> MSs and Pt CEs. The on-off plots are achieved by alternately irradiating (100 mW cm<sup>−2</sup>) and darkening (0 mW cm<sup>−2</sup>) the DSSC at 0 V, whereas the photocurrent stabilities are carried out under a sustained irradiation of 100 mW cm<sup>−2</sup>.

a parameter depending on the diffusion coefficient ( $D_n$ ) of the redox couple in electrolyte. The larger intersection of the cathodic branch with the  $y$ -axis demonstrated the NiS<sub>2</sub> MSs CE had higher limited current density ( $J_{lim}$ ) and higher diffusion velocity.<sup>53</sup>

Fig. 7a showed the start/stop switching of the DSSCs with NiS<sub>2</sub> MSs and Pt CEs at alternatively on/off illumination. The abrupt increase in photocurrent density without delay in starting the cell indicated that both of NiS<sub>2</sub> MSs and Pt CEs were vigorous in catalyzing the reduction of I<sub>3</sub><sup>−</sup> ions.<sup>13</sup> However, only 67.5% and 92.1% of the initial photocurrent density was remained for the device with NiS<sub>2</sub> MSs and Pt CEs under persistent irradiation for 3600 s under illumination at 100 mW cm<sup>−2</sup>, in comparison to 97.2% for NiS<sub>2</sub> MSs based DSSC (Fig. 7b). This result indicated that NiS<sub>2</sub> MSs CE had better stability. In order to cross-check the stability, the two CEs were subjected to sequential scanning CVs for 100 cycles at a scanning rate of 50 mV s<sup>−1</sup> (Fig. S9†). The current densities and the  $E_{pp}$  value of NiS<sub>2</sub> MSs CE have nearly no change during 100 cycles, while the  $E_{pp}$  of Pt CE increases noticeably. Moreover, symmetrical dummy cells (CE/electrolyte/CE) were also subjected to sequential scanning of EIS for 10 cycles (Fig. S10†). There were no obvious changes in both  $R_s$  and  $Z_N$  for all CEs, suggesting that the potential cycling exerted negligible influence on the series resistance as well as the mass transport between the CEs and redox pairs. The  $R_{ct}$  for Pt CE increased from 0.92 Ω cm<sup>2</sup> in the first cycle to 1.31 Ω cm<sup>2</sup> in the final cycle. However, for the NiS<sub>2</sub> MSs CE, the  $R_{ct}$  kept nearly constant (increasing from 0.76 to 0.81 Ω cm<sup>2</sup>). These electrochemical characterization confirmed that the NiS<sub>2</sub> MSs CE had better electrochemical stability than that of Pt CE in iodine electrolyte.<sup>54</sup> This result also implied that the NiS<sub>2</sub> MSs CE had better corrosion resistance to the iodine-based electrolyte than Pt CE.

## 4. Conclusion

In summary, 3D hierarchical NiS<sub>2</sub> microspheres had been prepared through a facile chemical etching/anion exchange reaction followed by a controlled annealing treatment. The as-prepared NiS<sub>2</sub> hierarchical microspheres were constructed by 2D nanoflakes building blocks, and possessed large surface areas, high structural void porosity and accessible inner



surface. All of these features were beneficial for the mass diffusion and fast charge transport between electrolyte and CEs. A series of chemical characterization, including CV, EIS and Tafel-polarization, showed that the resultant NiS<sub>2</sub> hierarchical microspheres had excellent electrocatalytic activity and stability toward the reduction of I<sub>3</sub><sup>−</sup> ions. A typical device assembled with NiS<sub>2</sub> hierarchical microspheres achieved an impressive PCE of 8.46% under AM1.5G illumination, higher than that of pyrolysis Pt electrodes (8.05%). This strategy opened up a new way to synthesize a variety of hierarchical structures by selective etching of Prussian blue analogous.

## Conflicts of interest

There are no conflicts to declare.

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