Hydrogenated ZnO Core–Shell Nanocables for Flexible Supercapacitors and Self-Powered Systems

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ABSTRACT Although MnO₂ is a promising material for supercapacitors (SCs) due to its excellent electrochemical performance and natural abundance, its wide application is limited by poor electrical conductivity. Inspired by our results that the electrochemical activity and electrical conductivity of ZnO nanowires were greatly improved after hydrogenation, we designed and fabricated hydrogenated single-crystal ZnO@amorphous ZnO-doped MnO₂ core–shell nanocables (HZM) on carbon cloth as SC electrodes, showing excellent performance such as areal capacitance of 138.7 mF/cm² and specific capacitance of 1260.9 F/g. Highly flexible all-solid-state SCs were subsequently assembled with these novel HZM electrodes using polyvinyl alcohol/LiCl electrolyte. The working devices achieved very high total areal capacitance of 26 mF/cm² and retained 87.5% of the original capacitance even after 10 000 charge/discharge cycles. An integrated power pack incorporating series-wound SCs and dye-sensitized solar cells was demonstrated for stand-alone self-powered systems.

KEYWORDS: supercapacitors · hydrogenation · core–shell · flexible · self-powered · ZnO

Specific energy storage devices possessing characteristics of flexibility, light weight, and even safety may meet the large proliferation of consumer electronics. Among various emerging energy storage technologies, supercapacitors (SCs), also named electrochemical capacitors (ECs), can be classified as two kinds: electrical double-layer capacitors (EDLCs), whose charges are adsorbed electrostatically at the electrode/electrolyte interface, typically are based on carbon materials; pseudocapacitors, which store energy by redox reactions on electrode materials, are usually based on conducting polymers and transition metal oxides. In fact, pseudocapacitors have higher specific capacitance than conventional EDLCs due to their fast and reversible redox reaction, so many efforts, including ours in this work, have been focused on further improving pseudocapacitors.

Compared to other transition metal oxides (RuO₂,⁹ CoO₆,¹⁰¹¹ NiO₆,¹² and FeO₆,¹³¹⁴), manganese dioxide (MnO₂) has significant superiorities such as low cost, low toxicity, abundant resource, and high theoretical specific capacitance (1370 F/g).¹⁵⁻¹⁸ Its poor electrical conductivity, however, limits the wide application of MnO₂. To overcome this problem, several strategies have been proposed, for instance, incorporating MnO₂ with carbon-based materials such as graphene,¹⁹⁻²¹ or graphene oxide and carbon nanotubes (CNTs),¹⁸,²³,²⁴ and core–shell structures (Co₉O₄@MnO₂,²⁵ SnO₂@MnO₂,²⁶ WO₃@Au@MnO₂,²⁷ Zn₅Sn₄O₆@MnO₂,²⁸ etc.). Recently, Jiang et al. reported MnO₂ nanowires/mesoporous carbon/MnO₂ nanoparticle hybrid nanowires, which significantly

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improved the conductivity of metal oxide materials.\textsuperscript{29} MnO$_2$ in this unique structure was efficiently utilized with the assistance of the highly conductive mesoporous carbon shell. Li et al. designed and synthesized novel MnO$_2$/Mn/MnO$_2$ sandwich-structured nanotube arrays with high supercapacitor performance.\textsuperscript{30} The special sandwich-structured nanotube architecture allowed highly efficient utilization of MnO$_2$ for charge storage with facilitated transport of ions and electrons, resulting in excellent electrochemical performance. Despite all of these achievements, enhancing the MnO$_2$ electrode conductivity still remains a challenge.

Zinc oxide (ZnO) nanomaterials, owing to their special chemical and physical properties, and many technical advantages such as low cost, easy fabrication, and cleanroom compatibility, are widely studied in electronics and piezotronics devices,\textsuperscript{31,32} strain sensors,\textsuperscript{33,34} surface switches,\textsuperscript{35} and photocatalysis.\textsuperscript{36} Motivated by the idea that hydrogenation of metal oxides can greatly improve its electrochemical activity and electrical conductivity,\textsuperscript{37} here, we designed and fabricated a novel hybrid architecture by easily coating MnO$_2$ on hydrogenated zinc oxide nanowires (HZnO) grown on flexible carbon cloth. During the procedure, ZnO was found to be doped into the MnO$_2$ spontaneously. Since HZnO had improved free carrier concentration and better electrical stability, they could be a good scaffold for MnO$_2$ loading. Moreover, the conductive carbon cloth could serve as a lightweight and flexible current collector. Hydrogenated single-crystal ZnO@amorphous ZnO-doped MnO$_2$ core–shell nanocable (HZM) electrodes exhibited high electrochemical performance, and all-solid-state SCs were fabricated with these novel HZM electrodes using a polyvinyl alcohol (PVA)/LiCl neutral electrolyte via a simple and low-cost method. The fabricated HZM-based all-solid-state SCs exhibited many advantages in performance such as high capacitance, high flexibility, long lifetime, and low leakage current. In addition, the SCs integrated with dye-sensitized solar cells (DSSCs) could serve as a switchable power pack that is able to store energy as sunlight illuminates and light up commercial light-emitting diodes (LEDs) as light turns off, demonstrating the potential broad applications of SCs as a power pack for stand-alone self-powered systems.

RESULTS AND DISCUSSION

The general electrode fabrication procedure is schematically illustrated in Figure 1. Initially, aligned and ordered ZnO nanowires (NWs) were grown on carbon cloth using a wet chemical method. The obtained ZnO NWs were about 100–200 nm in diameter and 4.5–7 μm in length (Figure 2a,b). Hydrogenation of the ZnO NWs in a commercial tube furnace improved the electrical properties of ZnO. The uniform hydrogenated ZnO NWs were then soaked in a glucose aqueous solution and subsequently annealed in Ar gas. An amorphous carbon formed and served as a reducing agent for the deposition of MnO$_2$ via the following reaction at room temperature:\textsuperscript{38,39}

\[4\text{MnO}_4^{-} + 3\text{C} + \text{H}_2\text{O} = 4\text{MnO}_2 + \text{CO}_3^{2-} + 2\text{HCO}_3^{-}\]

(1)

As shown in Figure 2c,d, the ZnO NWs became slightly longer in length and larger in diameter after deposition of the carbon layer and a subsequent deposition of MnO$_2$ layer; however, the NW morphology was preserved uniformly.

Figure 3a and Figure S1a in Supporting Information show a typical scanning transmission electron microscopy (STEM) image of an individual HZM nanocable in which a thin shell (∼30 nm) uniformly covers the
surface of the HZnO nanowire. The corresponding selected area electron diffraction (SAED) pattern (inset of Figure 3a) suggested the single-crystal ZnO core and its [0001] growth direction. The amorphous shell was double confirmed by an electron diffraction pattern, in which no other diffraction peaks were observed besides those from ZnO (inset in Figure 3a) and random atom arrangement in high-resolution transmission electron microscopy (HRTEM) (Figure 3g). The enlarged HRTEM image of the core in the rectangular area of Figure 3g is shown in its inset, and the (0001) atomic plane spacing was measured to be 0.52 nm. Energy-dispersive X-ray spectrometry (EDS) mapping analysis of elements Mn, Zn, and O (shown in Figure 3d–f, respectively) from a select area of a single hybrid nanocable (Figure 3c) confirmed the HZM core–shell structure. EDS line scanning (indicated by a line in Figure 3a) of Zn, Mn, and O across the HZM nanowire could better illustrate the core–shell configuration (shown in Figure 3b); the profile of Zn showed a broad peak, which was located at the center of the profile, while the profile of Mn showed higher intensity on both sides. Analysis of the data indicated that the diameter of the HZnO core and the total outer diameter of the composite nanocable were about 150 and 225 nm, respectively, in accordance with the STEM image (Figure 3a).

It was worthy to point out that the Zn signal was observed from the shell structure, reflected from both EDS mapping and line scanning. The reason for this is explained in the following. During the annealing process, carbon could convert the ZnO in contact with itself to Zn metal, which was more volatile than the oxide, so the surface of ZnO nanowires was no longer as smooth as before annealing. Vaporized zinc atoms diffused into the remaining carbon shell, and sometimes it generated a tiny gap between the ZnO core and the carbon shell (Figure S1b). Subsequently, when the sample was removed from Ar atmosphere and placed in ambient conditions, the Zn atoms embedded in the outer carbon layer were unavoidably oxidized to ZnO again. As the outer carbon layer reacted with KMnO₄, it generated an amorphous ZnO-doped MnO₂ shell (Figure S1c). XPS data (Figures 4 and S2) showed that Mn 2p₃/2 and Mn 2p₁/2 peaks were located at ca. 642.2 and 653.8 eV, suggesting the element Mn in the sample was present in the chemical state of Mn⁴⁺. The Zn 2p₃/2 and Zn 2p₁/2 lines were found at the binding energies of about 1021.3 and 1044.4 eV, indicating the existence of ZnO. The electron energy loss spectroscopy (EELS) mapping image is shown in Figure 3h, indicating Mn (red color) in the shell (consistent with EDS data in Figure 3b,d). The element C (green color) was negligible in the HZM nanocable, implying that the carbon already fully reacted with KMnO₄. Green rectangular area with a protruding green dot on the HZM nanocable was carbon absorption induced by a previous STEM EDS mapping procedure.

Figure 3. (a) STEM image of HZM core/shell nanostructure. Inset is SAED pattern of HZM nanocable, showing only a set of diffraction patterns from ZnO and suggesting the single-crystal ZnO core and amorphous shell. (b) EDS line scan curves showing Zn, Mn, and O element profiles across the HZM core/shell nanocable indicated in (a). (c) STEM image of a select area. (d–f) EDS element mapping images from the same area as in (c). EDS mapping images and line scan curves both indicated two elements (Zn and O) were in the core, and three elements (Mn, Zn, and O) were in the shell. (g) HRTEM image of the HZM core/shell nanocable, confirming the single-crystal ZnO core and amorphous shell. Inset image is from an enlarged area in (g), showing the 0.52 nm distance between (0001) planes and [0001] growth direction of the ZnO core. (h) EELS mapping image showing Mn (red) and C (green) elements. The EELS mapping image indicated that Mn was in the shell (consistent with EDS mapping), and C was negligible in the HZM nanocable. Green rectangular area with a protruding green dot on the HZM nanocable was carbon absorption induced by a previous STEM EDS mapping procedure.
were located at energies of about 1021.3 and 1044.4 eV, which were consistent with the values reported for ZnO. The charge/discharge curves of different ZnO electrodes were further studied by the galvanostatic electrochemical performance of different ZnO electrodes, indicating a larger specific capacitance. The CV area of the HZnO sample was nearly rectangular shaped curves suggested that ZnO electrodes collected at the scan rate of 100 mV/s. The untreated ZnO, air-annealed ZnO (AZnO), and HZnO electrodes were compared to remove the ZnO core but preserve the ZnO that might work as an EDLC. The CV area of the HZnO sample was significantly higher than those for ZnO and AZnO electrodes. For instance, the areal capacitance of the HZnO electrode achieved 1.38 mF/cm² at a current density of 0.05 mA/cm², which was approximately a 3- and 5-fold enhancement compared with the AZnO (0.46 mF/cm²) and ZnO (0.27 mF/cm²) electrodes, respectively. This areal capacitance of HZnO was much improved than other ZnO-based (0.15 mF/cm² at 0.03 mA/cm²) SCs. Additionally, it showed a smaller IR drop (0.027 V), confirming the good electrical conductivity of the HZnO electrode. According to the electrochemical impedance spectroscopy (EIS) curves in Figure 5c, the equivalent series resistance (ESR) was only 2.22 Ω for HZnO samples of 1 cm² area, much lower than 3.83 Ω for AZnO and 3.77 Ω for ZnO, further proving the good electrical properties of the HZnO that resulted from the hydrogenation.

To investigate the underlying mechanism for the enhanced electrical properties of ZnO by hydrogenation, Mott–Schottky (MS) measurements were conducted. This method was based on the Schottky barrier formation between the semiconductor material and an electrolyte. The variation of the differential capacitance (Cₜ) of the space charge layer was measured as a function of the applied potential. As shown in Figure 5d, all ZnO samples exhibited a positive slope, indicating n-type semiconductor character. Carrier densities (Nₐ) of ZnO samples were calculated by the following equation:

\[
N_\text{d} = \frac{2}{\varepsilon \varepsilon_0 A^2} \left| \frac{dV}{dC} \right|^{-2}
\]

where \( \varepsilon \) is the relative dielectric constant of ZnO (usually 8.5), \( \varepsilon_0 \) is the permittivity of vacuum, \( V \) is the potential applied at the electrode, and \( A \) is the area in contact with the electrolyte. The carrier densities of the HZnO, AZnO, and ZnO NWs were calculated to be 2.65 \times 10^{20}, 1.86 \times 10^{18}, and 7.04 \times 10^{17} \text{ cm}^{-3}, respectively. Although the MS equation was derived from a planar model, these results are still qualitatively comparable as these electrodes have similar morphology.
and surface area. The significant enhancement in the carrier density, which is over 2 orders of magnitude, could be clearly attributed to the hydrogen insertion. The low-energy oxygen vacancies that might be formed in high concentrations would be neutral and electrically inactive in n-type ZnO; however, the substitutional hydrogen atoms for the oxygen vacancies acted as shallow donors.46 The improved electrical conductivity and stability indicated that hydrogen played an important role in n-type conduction as the donor. This result was also in accord with a previous report by Lu et al.37

The aforementioned experimental results confirmed that HZnO NWs had better electrochemical and electrical properties, which enabled them to serve as effective support for other capacitive materials to form composite structures. Here, HZM nanocables were synthesized on carbon cloth via a self-limiting process (see Methods).

To demonstrate the electrochemical performance of the composite structures, the electrochemical properties of nanocable samples were first characterized in three-electrode ECs with CV, galvanostatic charge/discharge, and EIS measurements in 0.5 M Na2SO4 aqueous solution. The CV curves of untreated ZnO/ZnO-doped MnO2 (ZM), air-annealed ZnO/ZnO-doped MnO2 (AZM), and HZM electrodes collected at the scan rate of 100 mV/s are shown in Figure 6a. The HZM electrode exhibited a quasi-rectangular shape and a much more enhanced capacitive current density. Figure 6b shows the galvanostatic charge/discharge curves of different nanocable electrodes collected at a current density of 1 mA/cm². The negligible IR drop (0.019 V) was even smaller than HZnO electrode discharged at 0.05 mA/cm², benefitting from the synergistic effect of improved electrical performance of hydrogenated ZnO core and ZnO-doped MnO2 shell. The areal capacitance \( C_a \) of the electrode was calculated by the following equation

\[
C_a = \frac{I \Delta t}{\Delta \overline{E} \cdot S}
\]  

where \( I \) is the discharge current, \( \Delta t \) is the discharge time, \( \Delta \overline{E} \) is the potential window during the discharge process (subtract the IR drop), and \( S \) is the effective electrode area. The areal capacitance of HZM electrodes were as high as 138.7 mF/cm² at a current density of 1 mA/cm², which was 1.6 and 2.8 times those of AZM (88.2 mF/cm²) and ZM (49.3 mF/cm²) electrodes (Figure 6c), respectively. Remarkably, this value is also higher than the values reported for other manganese oxide composites, such as TiN@MnO2 coaxial arrays (∼41 mF/cm² at 0.12 mA/cm²),47 PEDOT@MnO2 nanoparticles (∼62 mF/cm² at 5 mA/cm²),48 SnO2@MnO2 nanowires (∼64 mF/cm² at 0.08 mA/cm²),26 hydrogenated TiO2@MnO2 nanowires (∼70 mF/cm² at 2 mA/cm²),49 and WO3@Au@MnO2 nanowires (∼105 mF/cm² at 0.06 mA/cm²),27 and even comparable to graphene sheets/Mn3O4 composites (∼242 mF/cm² at 1 mA/cm²).50

The specific capacitance was calculated as 1260.9 F/g since the loading of MnO2 was 0.11 mg/cm² determined by inductively coupled plasma atomic emission spectroscopy. Furthermore, HZM electrodes of 1 cm² area showed very low ESR values (2.74 Ω) in Figure 6d, suggesting that the HZM electrodes have very small resistance with good ion response at high-frequency ranges. The very low ESR value further confirmed that the HZnO NWs could indeed act as a good substrate for MnO2 loading.

An all-solid-state symmetric flexible SC was assembled with two HZM electrodes using a solid
PVA/LiCl electrolyte (Figure 7a). The CV curves in Figure 7b suggested that the SC could be bent and twisted without affecting the device performance much, showing good flexibility and stability. Galvanostatic charge/discharge tests were also performed with different current densities with a voltage window of 0–0.8 V (Figure 7c). The linear and symmetric curves revealed the good capacitive characteristic of the solid-state SC. The total areal capacitance of the device was nearly 26 mF/cm² (or 325 mF/cm³) at a current density of 0.5 mA/cm², which was higher than other solid-state devices reported previously, for instance, laser scribing graphene (∼1.84 mF/cm² at 0.82 mA/cm²)\(^5\) and bacterial nanocellulose/carbon nanotubes (∼20 mF/cm² at 0.4 mA/cm²),\(^6\) and even on the verge of carbon nanotubes papers (∼31 mF/cm² at 0.46 mA/cm²).\(^5\) Figure S3a showed the CV curves of the symmetric SC measured at scan rates ranging from 10 to 100 mV/s. These CV curves exhibited nearly rectangular shapes. The cycle life of the SC was tested through a cyclic charge/discharge process at a current density of 0.5 mA/cm², which is shown in Figure 7d. After 10 000 cycles, over 87.5% capacitance was maintained, demonstrating the excellent electrochemical performance of this HZM nanocable electrode material. The Coulombic efficiency (\(\eta\)) of the SC was characterized by a charge/discharge process according to

\[
\eta = \frac{\Delta t_d}{\Delta t_c}
\]  

where \(\Delta t_d\) and \(\Delta t_c\) represent the discharge and charge times, respectively. The Coulombic efficiency nearly exceeded 90% (Figure 7d), implying good charge/discharge reversibility for the solid-state SC. Potentiostatic EIS measurements were performed after first and 10 000th cycles to evaluate the internal resistance change during the cycling charge/discharge test (Figure S3b). An insignificant shift of ESR was observed from 5.9 to 9.0 Ω. The increase of charge transfer resistance might be caused by the distortion of charge transport pathway. Energy density (\(E\)) and power density (\(P\)) are two key factors for evaluating the application of SCs, which can be calculated as

\[
E = \frac{CV^2}{2U} \tag{5}
\]

\[
P = \frac{E}{t} \tag{6}
\]

where \(C\) is total capacitance of the device, \(V\) is the cell voltage, \(U\) is the device volume, and \(t\) is the discharge time. A power density of 2.44 mW/cm³ and an energy density of 0.04 mWh/cm³ were achieved. In particular, the energy density of our SC was higher than that of the other solid-state SCs (shown in Figure 7e), such as CNT-based SCs (0.008 mWh/cm³)\(^5\) and MnO₂-coated ZnO NWs (0.005 mWh/cm³).\(^4\)

As efficient energy storage devices, SCs should be able to not only drive electrical components but also store energy from various other energy sources, especially intermittent sources. Solar energy is probably the most clean and renewable energy in the world, but it is always limited by time, location, and weather. Integrating dye-sensitized solar cells (DSSCs) with energy storage devices such as SCs and batteries may offer a promising solution to build stand-alone self-powered systems,\(^5\) which potentially break the limitations of time, location, and weather. Specially, to demonstrate the potential applications of our newly fabricated all-solid-state SCs, we designed a stand-alone

Figure 6. (a) CV curves for ZM, AZM, and HZM electrodes at a scan rate of 100 mV/s. (b) Galvanostatic charge/discharge curves of ZM, AZM, and HZM collected at a current density of 1 mA/cm². (c) Areal capacitances of the ZM, AZM, and HZM electrodes measured as a function of current density. (d) Nyquist plots for ZM, AZM, and HZM electrodes with an area of 1 cm².
self-powered system consisting of DSSCs, SCs, and LEDs. Simple schematic view of a self-powered system is illustrated in Figure 8a. The switchable power pack in this system consisted of an environmental energy-harvesting module (four series-wound homemade DSSCs) and an energy storage module (three series-wound SCs). The photovoltaic performance of a DSSC cell was measured under the illumination of AM 1.5 (100 mW/cm²) by recording the current density/voltage response (Figure S4). When sunlight was available, the switch was set at position “1”, so the DSSCs harvested solar energy and stored the energy in the SCs. After 120 s charging under the illumination, the voltage of three series-wound SCs reached approximately 2.3 V (inset in Figure 8b). When the solar source was turned off and the room became dark, the switch was then placed to position “2”, and the charged SCs could drive a commercial blue LED to illuminate (Figure S5b). A red LED in this stand-alone self-powered system could be lighted up for more than 30 min by three series-wound SCs, which were previously charged by four series-wound DSSCs for 2 min. Figure S5b and video S1 demonstrate that an LED was powered by three series-wound SCs. At the beginning, the LED twinkled very brightly, demonstrating the excellent power merit of SCs. Figure S5a showed the time courses of the open-circuit voltage of the series-wound SCs. It gently underwent a self-discharge course; furthermore, the SC devices showed an output of ∼1.5 V after 40 min. On the level of practical application, leakage current of the devices is critical. As shown in Figure 8b, the leakage current dropped significantly at the beginning (from 0.5 mA to 68.7 µA after 100 s and to 22.8 µA after 400 s) and then gradually became stable and small (finally to only 8.5 µA after 4500 s). This value was even smaller than that of the CNT/PANI supercapacitor, indicating few impurities in the electrode and electrolyte materials in the fabricated SCs. Such demonstration implied...
that the stand-alone self-powered systems had potential applications in combining mobile devices and driving small electrical compounds.

CONCLUSION

In summary, we have reported the design and synthesis of a novel nanocable material, HZM with a hydrogenated single-crystal ZnO core and amorphous ZnO-doped MnO2 shell, as well as the fabrication of a new kind of flexible all-solid-state SC configuration with two pieces of slightly separated carbon cloth grown with HZM nanocables as electrodes and PVA/LiCl gel as solid-state electrolyte. The SC devices exhibited many advantages in performance such as high capacitance, high flexibility, long lifetime, and low leakage current. The hydrogenated ZnO core with larger charge concentration and ZnO-doped MnO2 shell with better conductivity contributed to the high performance of SCs. Our HZM flexible SCs might greatly contribute to the fundamental research and technologies of flexible energy storage devices. A potential application of the flexible solid-state SCs as a switchable power source in stand-alone self-powered systems was demonstrated, which exhibited the potential broad applications of the SCs.

METHODS

Synthesis of HZM. First, ZnO NWs were grown on carbon cloth (WOS1002, 2.5 cm × 0.7 cm × 0.6 μm) by a wet chemical process. Briefly, carbon cloth was first soaked in 0.5 M KMnO4 for 30 min to form a seed layer. A precursor 100 mL solution was prepared with 0.015 M of zinc nitrate hexahydrate and 0.015 M of hexamethylenetetramine (HMTA) added with 4 mL of ammonia. The seeded carbon cloth substrates were dipped into the precursor solution then placed under 90 °C for 24 h. After the wet chemical process, ZnO NWs were grown on the carbon cloth. The HZnO NWs were obtained by annealing the ZnO NWs in hydrogen atmosphere at 350 °C for 3 h. ZnO NW samples were also annealed in air at 350 °C for 3 h to obtain AZnO as control samples for comparison. Second, the HZnO NWs were immersed into a 0.5 M aqueous glucose solution for 24 h. HZC samples were then annealed by carbonization at 450 °C in Ar gas for 2 h. Third, MnO2 was deposited on HZC via a self-limiting process.88 Simply, 0.5 M KMnO4 and 0.5 M Na2SO4 was mixed with equal volume. The HZC samples were immersed in the solution for 8 h at room temperature. Then the HZM was cleaned several times using deionized water. The loading mass of MnO2 was 0.11 mg/cm2.

Assembly of the Solid-State SC. The SC was assembled by two pieces of HZM NW electrodes with a separator (NKK TF40, 0.7 cm × 1.5 cm) and LiCl/PVA gel as a solid electrolyte. LiCl/PVA gel was prepared by mixing LiCl (12.6 g) and PVA (6 g) in 60 mL of deionized water and heated at 85 °C for 30 min to form a seed layer. A precursor 100 mL solution was prepared with 0.015 M of zinc nitrate hexahydrate and 0.015 M of hexamethylenetetramine (HMTA) added with 4 mL of ammonia. The seeded carbon cloth substrates were dipped into the precursor solution then placed under 90 °C for 24 h. After the wet chemical process, ZnO NWs were grown on the carbon cloth. The HZnO NWs were obtained by annealing the ZnO NWs in hydrogen atmosphere at 350 °C for 3 h. ZnO NW samples were also annealed in air at 350 °C for 3 h to obtain AZnO as control samples for comparison. Second, the HZnO NWs were immersed into a 0.5 M aqueous glucose solution for 24 h. HZC samples were then annealed by carbonization at 450 °C in Ar gas for 2 h. Third, MnO2 was deposited on HZC via a self-limiting process.88 Simply, 0.5 M KMnO4 and 0.5 M Na2SO4 was mixed with equal volume. The HZC samples were immersed in the solution for 8 h at room temperature. Then the HZM was cleaned several times using deionized water. The loading mass of MnO2 was 0.11 mg/cm2.

Instrumentation. The structural properties of electrode materials were characterized by field-emission SEM (FEI Sirion 2000), TEM (FEI TECNAI F30) equipped with an energy-dispersive X-ray spectroscopy (EDS) detector, and electron energy loss spectroscopy (EELS) detector, X-ray photoelectron spectroscopy (XPS, Thermo K-alpha). The electrochemical properties of the products were investigated with cyclic voltammetry (CV) and chronopotentiometry measurements employing a CHI 660D electrochemical workstation (Chenhua, Shanghai) and VersaSTAT 3 (Princeton Applied Research), and the electrochemical impedance spectroscopy (EIS) was measured by an Autolab PGSTAT302N at a frequency ranging from 100 mHz to 10 kHz with a potential amplitude of 10 mV. The photoelectric conversions were characterized by recording voltage—current curves with a computerized digital multimeter (Keithley 2601A). An AM 1.5 solar simulator (Oriel 91197) equipped with an energy-dispersive X-ray spectroscopy (EDS) was used to record the photoelectric conversion under an AM 1.5 solar stimulator (Abet Technologies). The cycling stability was recorded by a battery test system (Neware BTS). For a single electrode test, a piece of electrode (∼0.7 cm × 1.5 cm, effective area ∼1 cm2) was dipped into 0.5 M Na2SO4 solution at room temperature. Ag/AgCl reference electrode and graphite rod counter electrode were used in the measurement.

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Supporting Information Available: Assembly of the DSSCs, TEM images for HZ and HZM, more XPS analysis data of HZM, self-discharge curve of the SC, more EIS data, schematic of the DSSCs, DSSC data, more electrical and electrochemical data for ZnO, AZnO, HZnO, 2M, AZM, and HZM electrodes, and a video of lighting up a red LED by three series-wound SCs. This material is available free of charge via the Internet at http://pubs.acs.org.

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