A Novel Preparation of Nano-Copper Chalcogenide (Cu$_2$S)-based Flexible Counter Electrode

Enl Wu$^{1}$, Jingsha Jin$^{1}$, Shaowen Liu$^{1}$, Dan Li$^{1}$, Shufang Gao$^{1}$, Fei Deng$^{2}$, Xuemin Yan$^{2}$, Yan Xiong$^{1}$ & Haolin Tang$^{3}$

Copper nanowires (CuNWs) are used to prepare flexible, transparent conducting substrates due to their low cost and ease of fabrication on a large area. A CuNW/polymer composite substrate was prepared and vulcanized to create a novel flexible nano-Cu$_2$S/polymer composite substrate. The physical and photovoltaic properties of the substrate can be controlled by tuning the concentration of CuNW dispersion during the preparation of CuNWs and nano Cu$_2$S films. The nano-Cu$_2$S-based composite substrate was used as an effective flexible counter electrode of a quantum-dot-sensitized solar cell (QDSSC) and resulted in a maximum cell efficiency of 1.01%.

Dye-sensitized solar cells (DSSCs) have a photoelectric conversion efficiency of over 12%, making them a potential candidate for next-generation solar cells$^{12}$. Quantum-dot-sensitized solar cells (QDSSCs) use semiconductor quantum dots (QDs) in sensitizers, instead of organic dyes. As a result, QDSSCs exhibit the unique advantages of quantum size effect, multi-exciton effect, large absorption coefficient, and easy matching of energy levels between the electron donor and acceptor materials$^{13,14}$. The structure of a QDSSC is sandwich-like and mainly consists of a photoanode, electrolyte, and counter electrode. The photoanode is mainly composed of a conductive transparent substrate, such as fluorine-doped tin oxide (FTO) glass, with an overlying semiconductor oxide film such as TiO$_2$, ZnO that adsorbs a sensitizer, i.e. QDs. A polysulfide electrolyte works as a redox couple. The counter electrode, generally copper chalcogenide (Cu$_2$S or CuS), serves as a reduction catalyst$^{5,6}$. CdS/CdSe-cosensitized TiO$_2$ is widely studied as a classical co-sensitization system. The CdS QDs adsorbed on the TiO$_2$ films show a good effect on the deposition of CdSe QDs, finally forming a classical TiO$_2$/CdS/CdSe cascade structure$^{6,7}$.

Copper is the most frequently used metal material for industrial and commercial applications. Recent research has focused on the applications of copper nanowires (CuNWs)$^{8,9}$. A nanowire-based transparent conductive film has the advantages of excellent photoelectric performance, low preparation cost, and it could be used to prepare flexible devices. CuNWs are being used to prepare flexible transparent conducting substrates because of their low cost and ease of fabrication on a large area$^{10-12}$.

While DSSCs typically use Pt- or Au-coated conducting glass as the counter electrode, QDSSCs usually use a copper-chalcogenide-based counter electrode. This is because the sulfur-containing electrolyte absorbs preferentially and strongly on the Pt or Au surface, leading to surface passivation and decrease in the conductivity of electrodes$^{13}$. Bulk and nanostructured copper chalcogenides are used as the counter electrodes in QDSSCs. The bulk copper chalcogenides are mainly made using brass, and they have the best cell efficiency. However, the cells suffer from mechanical and chemical instability$^{13-15}$. The nanostructured copper-chalcogenide-based counter electrodes are usually fabricated by either synthesizing Cu$_{2-x}$S and then coating it on the conducting glass, or by assembling nano Cu$_2$S arrays on the rigid substrate$^{16}$.

In this study, a flexible CuNW-based composite substrate was prepared and then vulcanized to create a flexible nano-Cu$_2$S-polymer composite substrate. To the best of our knowledge, this is the first time that a nano Cu$_2$S film was fabricated as described, and a flexible nano-Cu$_2$S-based counter electrode on a polymer substrate was used in QDSSCs. The physical and photovoltaic properties of the substrate can be controlled by tuning the concentration of CuNW dispersion during the preparation of CuNWs and nano Cu$_2$S films. The novel composite substrate functioned well as the flexible counter electrode of a CdS/CdSe QD co-sensitized solar cell.

$^1$School of Physics and Optoelectronic Engineering, Yangtze University, Jingzhou, 434023, P.R. China. $^2$College of Chemistry and Environmental Engineering, Yangtze University, Jingzhou, 434023, P.R. China. $^3$State Key Laboratory of Advanced Technology for Materials Synthesis and Processing, Wuhan University of Technology, Wuhan, 430070, P.R. China. Correspondence and requests for materials should be addressed to Y.X. (email: xiongyan1215@163.com) or H.T. (email: thln@whut.edu.cn)
Materials and Methods

Materials. Cadmium nitrate tetrahydrate (Cd(NO₃)₂·4H₂O ≥ 98.0%), sodium sulfide nonahydrate (Na₂S·9H₂O ≥ 98.0%), selenium (Se ≥ 99.5%), sodium sulfite (Na₂SO₃ ≥ 98.0%), cadmium sulfate hydrate (CdSO₄·8H₂O ≥ 99.0%), nitritolactic acid (C₆H₆NO₃ ≥ 99.0%), 2,2-Dimethoxy-2-phenyl-acetophenone (DMPA), Sulfur (S), potassium chloride (KCl ≥ 99.5%), nitritolactic acid and potassium hydroxide (KOH ≥ 98.0%) were purchased from Sigma-Aldrich. Isopropyl alcohol, methanol, ethanol, and acetone were obtained from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Conducting FTO glass and commercial TiO₂ nanoparticle (P25)-coated FTO were acquired from Yinkou OPV Tech New Energy Co. Ltd. (Yinkou, China). The TiO₂-coated FTO glass had an effective area of 0.16 cm². CuNWs and PVP ((C₆H₉NO)n) ethylene glycol dimethacrylate was purchased from Changxing Chemical Co., Ltd. (Zhuhai, China). All of the chemicals used in the preparation process were cleaned using detergent, deionized water, acetone, and isopropyl alcohol, and stored in the dark at room temperature.

Preparation of CdS/CdSe photoanode. For making CdS QDs, a TiO₂ film was first dipped into an ethanol solution containing 0.1 M Cd(NO₃)₂ for 1 min and rinsed with ethanol. Then, the film was dipped for another 1 min into a 0.1 M Na₂S methanol solution and rinsed with methanol. This two-step dipping procedure is regarded as one successive ionic layer adsorption and reaction (SILAR) cycle, and the incorporated amount of CdS can be increased by repeating the assembly cycles. A total of 12 SILAR cycles were performed, and then the glass was air dried. Next, CdSe was deposited on top of the CdS-coated glass by a chemical bath deposition (CBD) method. The CdSe deposition was achieved by using nitritolactic acid as a complex and selenosulfate as an Se source. First, for the Se source, Na₂SeSO₃ aqueous solution was freshly prepared by refluxing 0.2 M Se powder in an aqueous solution of 0.5 M Na₂SO₃ at 70 °C, for approximately 5 h. Nitritolactic acid and KOH were mixed to prepare K₃NTA solution. Then, a solution was prepared by mixing 80 mM CdSO₄, 160 mM K₃NTA, and 80 mM Na₂S in methanol. In the next step, the TiO₂ film was dipped into the prepared solution for 3–5 min to generate nano Cu₂S on the surface of the composite.

Preparation of CuNW-polymer composite substrate. Prior to the procedure, all of the glass substrates used in the preparation process were cleaned using detergent, deionized water, acetone, and isopropyl alcohol, under ultrasonication. CuNWs were dispersed in isopropanol with 1% (mass ratio) of PVP as agent. Here, PVP was used as a surfactant and stabilized reagent to prevent the nanowires from coalescing. The dispersion was treated under ultrasonication to obtain a uniformity.

A CuNW film can be prepared by different methods, such as spin coating, Mayer rod coating, and spray coating. In this study, to fabricate a relatively thick film of CuNWs, a dispersion of CuNWs in isopropanol was drop cast on the glass substrate at room temperature. After the required thickness of CuNWs was drop cast, the glass substrate was air dried. Afterwards, the acrylate monomer with 1 wt% DMPA as a polymerization initiator was coated on the CuNW coating. The coating was then cured under an ultraviolet curing conveyor for approximately 90 s and peeled off as a transparent flexible composite.

CuNW vulcanization. In a water and methanol (1:1 volume) solution, 0.1 M Na₂S, 0.1 M S, and 0.2 M KCl were added to prepare a solution. The previously prepared CuNW-based flexible composite was placed in this solution for 3–5 min to generate nano Cu₂S on the surface of the composite.

QDSSC device fabrication and characterization. The sensitized TiO₂ film was used as a photoanode and the nano-Cu₂S-based composite substrate was used as the counter electrode. The electrolyte, which consisted of 0.1 M Na₂S, 0.1 M S and 0.2 M KCl in a water and methanol (1:1 volume) solution, was injected between the photoanode and counter electrode by siphonic action.

The photovoltaic performances [short circuit current (Jsc), open circuit voltage (Voc), fill factor (FF), and power conversion efficiency (η)] of the cell were measured. The current density-voltage (J-V) characteristics of the cells were measured using a Keithley 2450 source meter for a light intensity of 100 mW/cm² from a Xenon lamp (300 W; Nbet, HSX-F300). The microstructure of the CuNWs was analyzed with a field emission scanning electronic microscope (SEM; JEOL, JSM7100F). Electrochemical impedance spectroscopy (EIS) measurements were obtained using an electrochemical workstation (CorrTest, CS 350 H). Elemental analysis was performed with an energy dispersive spectrometer (EDS, Oxford X-MAX). X-ray diffraction (XRD) patterns were recorded by an X-ray diffractometer (Empyream, PANalytical).

Results and Discussion

Figure 1 shows the transmittance of CuNW-based flexible composite substrates prepared with dispersion of different CuNW concentrations. The transmittance of the film reflects the thickness and compactness of the film. This gives an indication of the quality of the film. For the substrates prepared with CuNW dispersion concentrations less than 5 mg/ml, the transmittance in the wavelength range of 350–800 nm is higher than 50%. As the CuNW concentration of the dispersion increases from 3 mg/ml to 8 mg/ml, the transmittance of the substrate gradually decreases. This decrease happens because the distribution density of the nanowires grows with the dispersion concentration.

Figure 2 shows a CuNW-based composite substrate (Fig. 2a) and a vulcanized substrate (Fig. 2b). Both substrates have good flexibility. After CuNW vulcanization, the color of the substrate turned from bronze to black, which is the typical color of Cu₂S. The average thicknesses of the CuNW layer and composite substrate were 4 μm and 44 μm, respectively.

SEM images of the surface of a CuNW-based composite substrate are shown in Fig. 3. The samples were prepared with dispersion of CuNW concentrations of 5 mg/ml, 7 mg/ml, and 8 mg/ml, respectively. CuNWs are
evenly distributed on the substrate. As the CuNW concentration is increased, the composite substrate is coated with denser CuNWs.

The CuNW-based composite substrates that were prepared with CuNW concentrations of 5 mg/ml and 7 mg/ml were sulfided to produce nano Cu₂S. The SEM images of the surface of these samples are shown in Fig. 4. The Cu₂S flakes are closely arranged. Compared to nano Cu₂S fabricated with lower CuNW dispersion concentration (5 mg/ml), the nano Cu₂S prepared with a higher CuNW dispersion concentration (7 mg/ml) are more densely distributed. The morphology of the nano Cu₂S is similar to the morphology found in literature13,21.

To further confirm the vulcanization of CuNWs, EDS analysis was conducted to investigate the elemental compositions of non-vulcanized and vulcanized CuNWs on the flexible substrates. Samples of CuNW-based composite substrates, which were fabricated with dispersion of different CuNW concentrations were analyzed. The details of the EDS analysis are listed in Table 1. The samples were primarily composed of C, O, and Cu. The Cu atomic percentage gradually increased as the CuNW dispersion concentration increases. The vulcanized samples were primarily composed of C, O, S, Cu, and Cu₂S. After vulcanization, the atomic percentage of Cu in the sample decreased. This decrease may be caused by the loss of CuNWs during the vulcanization in the solution. The ratio of S atoms to Cu atoms are similar for CuNW concentrations between 5 mg/ml to 8 mg/ml.

The vulcanization of CuNWs was also confirmed by the XRD patterns presented in Fig. 5. The diffraction peaks of the CuNW/polymer composite matched well with known Cu peaks (JCPDS 04-0836). After vulcanization, new peaks corresponding to Cu₂S (JCPDS 33-0490) were observed.

Figure 6 shows the J–V characteristics of the solar cells using a nano-Cu₂S-based composite substrate as counter electrode. Table 2 shows the key photovoltaic parameters (Jsc, Voc, FF, and maximum total energy conversion efficiency, η) of the devices. As the concentrations of CuNWs in the dispersion increases from 3 to 8 mg/ml, the cell efficiency of the corresponding QDSSCs first increases. A maximum efficiency of 1.01% is achieved when
the CuNW dispersion concentration is 8 mg/ml. After this, the efficiency decreases as the CuNW concentration is increased to 9 mg/ml and 10 mg/ml. The cell efficiency is 0.8% or more when the dispersion concentration of CuNWs is between 6 mg/ml and 9 mg/ml. This increased energy conversion efficiency can be mainly attributed to an increase in the photocurrent. The open circuit voltage of the QDSSCs with CuNW concentrations between 4 mg/ml and 9 mg/ml is in the range of 0.52 V to 0.6 V.

As the CuNW concentration in the dispersion increases, the as-prepared composite substrate and the corresponding vulcanized substrate obtains denser CuNWs or nano Cu2S on the substrate. Thus, the efficiency of the QDSSCs using nano Cu2S counter electrode increased until the saturation point is reached at about 7 mg/ml–8 mg/ml.

It should be noted that, during the fabrication of a CuNW/polymer composite, as the CuNW concentration was increased, the density of the drop-cast CuNW film on the glass substrate increased. It took less time for a CuNW film made with a higher CuNW concentration to air dry. In addition, CuNWs made with a higher CuNW concentration dispersion are expected to aggregate more easily. As the CuNW concentration was increased to more than 9 mg/ml, it was difficult to completely transfer CuNWs from the glass substrate to the acrylate polymer film; after the CuNW/polymer film was peeled off from the glass substrate, some CuNWs were still left on the glass substrate. These factors lead to a low efficiency of QDSSCs when the CuNW dispersion concentration is 10 mg/ml.

To evaluate the flexibility of nano-Cu2S-based counter electrodes, they were subjected to a bending test. The nano Cu2S counter electrode was bent along a curve of radius 0.7 cm, which corresponds to a curvature of $143 \text{ m}^{-1}$. After repeatedly bending the electrode 200 times, a tiny decrease in cell performance was observed. The cell efficiency was acceptable at 0.96%. The $J_{sc}$, $V_{oc}$ and $FF$ were 0.50 V, 7.08 mA/cm$^2$, and 0.27, respectively. Figure 7 shows the $J-V$ characteristics of the solar cells using a nano-Cu2S-based composite substrate that was bent 200 times. No fall-off or breakage was observed even after the photovoltaic test (Fig. 7, inset). While bulk copper chalcogenide electrodes are usually prepared by immersing a polished brass in a (poly)sulfide solution,14,15 when Cu2S coated brass is employed as the counter electrode of QDSSCs, the Cu2S on the surface exfoliates easily. Thus, the novel flexible nano-Cu2S-based counter electrode has better flexibility and mechanical stability than bulk copper chalcogenide electrodes.

The interface characteristics of the QDSSCs were studied by the EIS method. Figure 8 shows the Nyquist plots of QDSSCs under illumination of 100 mWcm$^{-2}$. For CuNW dispersion concentrations of 3 mg/ml and 5 mg/ml, two semicircles are observed in the Nyquist curves for both CuNW concentrations. The smaller semicircle appears at high frequencies and represents the redox impedance of the counter electrode interface. The larger semicircle appears at middle frequencies and represents the impedance at the photoanode/dye (QDs)/electrolyte interface. The semicircles can be represented by an equivalent circuit, as shown in Fig. 8 (inset)22,23. The equivalent circuit is composed of a series resistance $R_s$, transfer resistances $R_{ct1}$ and $R_{ct2}$, and chemical capacitances $CPE_1$ and $CPE_2$. $R_{ct1}$ and $R_{ct2}$ represents the charge transfer resistance at the interface of the electrolyte and counter electrode. $R_{ct2}$ represents the charge transfer resistance at the interface of TiO$_2$/QD/electrolyte. $CPE_1$ and $CPE_2$ are constant phase elements of the capacitances corresponding to $R_{ct1}$ and $R_{ct2}$, respectively. Table 3 shows that a substrate prepared with a CuNW dispersion concentration of 5 mg/ml has a smaller $R_{ct1}$ value than that of a dispersion concentration of 3 mg/ml. This trend indicates that the carriers are easier to transfer at an electrode with denser nano Cu2S.

Figure 3. Scanning electronic microscope (SEM) images of the surface of CuNW-based flexible composite substrates, prepared with dispersion of CuNW concentrations of (a) 5 mg/ml, (b) 7 mg/ml, and (c) 8 mg/ml.
The Cu₂S-polymer composite used in this study is easy to fabricate. In addition, once CuNWs have been synthesized, they can be directly used to prepare a nano Cu₂S film. The density and morphology of the distributed nano Cu₂S on the polymer substrate is controllable by tuning the CuNW dispersion concentration. This convenient and simple method can also be generalized to the fabrication of similar flexible electrodes. The performance

![Figure 4. SEM images of the surface of nano-Cu₂S-based composite substrate prepared with dispersion of CuNW concentrations of 5 mg/ml (a–c) and 7 mg/ml (d–f).](image)

| Dispersion concentration (Atomic percentage) | 3 mg/ml | 4 mg/ml | 5 mg/ml | 7 mg/ml | 8 mg/ml |
|-----------------|---------|---------|---------|---------|---------|
| CuNW C-K        | 72.81   | 55.86   | 57.37   | 54.21   | 49.65   |
| CuNW O-K        | 13.43   | 19.56   | 16.52   | 9.89    | 8.25    |
| CuNW Cu-L       | 14.19   | 24.56   | 25.98   | 35.53   | 42.48   |
| Vulcanized CuNW C-K | --      | --      | 54.86   | 54.64   | 50.29   |
| Vulcanized CuNW O-K | --      | --      | 13.42   | 13.73   | 14.07   |
| Vulcanized CuNW S-K | --      | --      | 7.47    | 7.56    | 7.70    |
| Vulcanized CuNW Cu-L | --      | --      | 24.19   | 23.96   | 27.68   |

Table 1. Element distribution of CuNW-based composite substrate and vulcanized substrate, as seen using EDS analysis.
Figure 5. X-ray diffraction patterns of CuNW-based flexible composite substrate (a) and vulcanized substrate (b), prepared with dispersion of CuNW concentrations of 8 mg/ml.

Figure 6. J-V characterization of the solar cells, where J is the current density (mA/cm²), and V is the voltage.

| Dispersion concentration (mg/ml) | $V_{oc}$ (V) | $J_{sc}$ (mA/cm²) | $\eta_{max}$ (%) | FF  |
|----------------------------------|-------------|-------------------|-----------------|-----|
| 3                                | 0.34        | 1.88              | 0.19            | 0.29|
| 4                                | 0.54        | 3.23              | 0.43            | 0.25|
| 5                                | 0.60        | 3.42              | 0.53            | 0.26|
| 6                                | 0.54        | 5.50              | 0.80            | 0.26|
| 7                                | 0.57        | 7.79              | 0.99            | 0.22|
| 8                                | 0.57        | 6.89              | 1.01            | 0.25|
| 9                                | 0.52        | 5.66              | 0.90            | 0.30|
| 10                               | 0.48        | 4.92              | 0.59            | 0.24|

Table 2. Photovoltaic parameters of the solar cells.
of the as-prepared counter electrode depends a lot on the quality of the CuNWs and the corresponding nano Cu$_2$S film. The morphology and dispersity of CuNWs affect the electrical conductivity, uniformity, and vulcanization of the CuNW-polymer composite substrate, and consequently of the Cu$_2$S-polymer composite counter electrode. The efficiency of QDSSCs is expected to increase with the improvement of CuNWs, and we are exploring this.

Figure 7. $J$-$V$ characterization of the QDSSC using a nano-Cu$_2$S-based composite substrate that was bent 200 times, where $J$ is the current density (mA/cm$^2$), and $V$ is the voltage. Inset: nano-Cu$_2$S-based counter electrode after the photovoltaic test.

Figure 8. Nyquist plots of the QDSSCs under 100 mW/cm$^{-2}$ illumination and frequency from 0.1 Hz to 500 KHz at room temperature. Inset: equivalent circuit model of the QDSSCs.

| Dispersion concentration | $R_s$ ($\Omega$) | $R_{ct1}$ ($\Omega$) | $R_{ct2}$ ($\Omega$) |
|--------------------------|------------------|----------------------|---------------------|
| 3 mg/ml                  | 2.620            | 7.685                | 43.61               |
| 5 mg/ml                  | 3.608            | 5.455                | 86.48               |

Table 3. Photovoltaic properties obtained by fitting the impedance spectra of QDSSCs using the equivalent circuit shown in Fig. 8.
Conclusion

CuNW films were prepared on glass substrates by CuNW dispersion. A flexible CuNW-polymer composite substrate was then prepared by coating an acrylate monomer onto the CuNW coated glass and then curing under ultraviolet light. After vulcanization treatment, the CuNWs turned into nano Cu$_2$S and a novel flexible nano-Cu$_2$S-based composite substrate was obtained. The nano-Cu$_2$S based substrate was used as a counter electrode of QDSSCs. A maximum cell efficiency of 1.01% was observed. The effects of the concentration of CuNW dispersion on the physical and photovoltaic properties of the CuNW film, nano Cu$_2$S film, and QDSSCs were investigated by absorption spectroscopy, energy dispersive spectrometry, SEM, XRD, J-V characteristics, and EIS.

If the CuNW in the dispersion is thick enough, the as-prepared counter electrode would achieve a denser Cu$_2$S J-V characteristics, and EIS. 

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Acknowledgements

This work was supported by the Grants No. 11804032 and 51472034 from the National Natural Science Foundation of China, No. 201801023A from the Intellectual Property Office of Hubei Province of China, No. 2650-5006-0404 from Petro China Innovation Foundation.

Author Contributions

E.W. was responsible for the fabrication and characterizations of nano-Cu$_2$S-based composite substrate and QDSSCs; J.J., S.L. and D.L. contributed to this work by preparing the QDs and the photoanode. E.W. and F.D. contributed to the drafting and editing of the manuscript. S.G., F.D. and X.Y. carried out characterizations. The results obtained including UV-vis spectra, J-V curves, SEM images, EIS and EDS analysis were interpreted and analyzed, and the concepts and design for the experiment are planned and discussed by E.W., X.Y. and H.T. All authors gave final approval for publication.

Additional Information

Competing Interests: The authors declare no competing interests.

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