Research Article

H₂O₂ Treatment of Electrochemically Deposited Cu₂O Thin Films for Enhancing Optical Absorption

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Cu₂O is considered to be promising as an absorber layer material of solar cells, but its band gap (about 2.1 eV) is larger than the optimum one (about 1.5 eV). CuO has a smaller band gap of about 1.35 eV. Therefore, we attempted to oxidize Cu₂O using H₂O₂ to increase oxygen ratio and decrease band gap. Cu₂O thin films were deposited on indium-tin-oxide-coated glass from an aqueous solution containing CuSO₄, lactic acid, and KOH by the galvanostatic electrochemical deposition at 40°C with current density of −1 mA/cm². Then, the as-prepared copper oxide thin film was dipped in H₂O₂ (30%) at fixed temperature to oxidize for some time. By the H₂O₂ treatment at room temperature, the oxygen content was increased, and the band gap was decreased.

1. Introduction

Solar cell devices can be divided into three major categories: silicon solar cells, inorganic compound semiconductor solar cells, and organic solar cells. At present, crystalline silicon solar cells have high photovoltaic conversion efficiency, but production cost is high. For amorphous silicon solar cells, the production cost can be significantly reduced, but the main problem is that the photovoltaic conversion efficiency is low. Organic solar cells in general have a severe problem of poor stability. As an alternative, compound semiconductor solar cells are considered to increase the conversion efficiency and effectively reduce production cost. CuInSe₂ (CIS) and CdTe solar cells are now in practical use as compound semiconductor solar cells. One of the advantages of CIS and CdTe solar cells compared to silicon-based solar cells is that their absorption coefficients are large so that the film thickness can be small. Moreover, a manufacturing process is not complicated, and hence production cost will be reduced. However, In and Te are rare-earth elements, and Se and Cd are toxic. In contrast, constituent elements of Cu₂O are abundant, inexpensive, and harmless to a human body. Rühle et al. reviewed recent developments of all-oxide PV systems, which until today were mostly based on Cu₂O as an absorber [1].

So far, the Cu₂O/ZnO heterojunction was fabricated by several methods, for example, sputtering technique [2–5], oxidation of metals [6, 7], photochemical deposition [8], and electrochemical deposition (ECD) [9–12]. ECD is a simple technique which can produce films over a large area and be easily scaled up to a commercial process at low cost. We also have been working on ECD of Cu₂O/ZnO heterostructures [13, 14]. However, one of the shortcomings of Cu₂O solar cells is that the Cu₂O band gap (about 2.1 eV) is larger than the optimal value (about 1.5 eV) of a solar cell. Cu₂O can absorb only the light with wavelength less than 0.6 μm. Thus, if the band gap is decreased, optical absorption of the film and the photocurrent can be increased. Since CuO has a smaller band gap (about 1.35 eV) [15–19], we may be able to reduce the band gap by increasing oxygen content of the film. Recently, it was reported that the performance of the Schottky contact of ZnO can be improved by the surface treatment with boiling hydrogen peroxide [20]. Through the H₂O₂ treatment, a ZnO₂ layer is formed on the ZnO surface. Thus one can expect that composition of Cu₂O can be made close to CuO by the H₂O₂ treatment. In this study, we attempt H₂O₂ treatment of ECD-Cu₂O films to reduce the band gap and increase the optical absorption.
2. Experimental Procedure

An aqueous solution used for the deposition of Cu$_2$O contained 0.2 mol/L CuSO$_4$ and 1.6 mol/L lactic acid. pH of the solution was adjusted to 12.5 by adding KOH. Cu$_2$O was deposited galvanostatically at 40°C with a current density of −1 mA/cm$^2$. The sample was deposited for 10 min. The indium-tin-oxide-(ITO-) coated glass sheets with sheet resistance about 8-9 Ω/square were used as substrates. Then, the as-prepared Cu$_2$O thin film is dipped in H$_2$O$_2$ (30%) for some time. After the treatment, the sample was washed with pure water.

The structural, compositional, and optical properties of the films were characterized by various techniques. Profile meter Accretech Surfcom-1400D was used to measure the thickness of the thin film. The optical transmission measurement was performed using a JASCO-570 spectrometer with the substrate as the reference. The photoconductivity of the film was examined by means of the photoelectrochemical (PEC) measurement [21, 22]. The PEC measurement was carried out using the three-electrode cell with saturated calomel electrode (SCE) as a reference electrode. The deposited film was used as a working electrode and 100 mM of Na$_2$SO$_4$ was used as an electrolyte. The incident light from an xenon lamp (about 100 mW/cm$^2$) was irradiated on the backside of the sample and turned off and on mechanically every five seconds. The current was measured under application of a ramp voltage with a scan rate of 5 mV/s, first in the cathodic bias range (from 0 to −0.5 V) and then in the anodic bias range (from 0 to +0.5 V). The Auger electron spectroscopy
2.3
2
1.7
1.4
1.1
0.8
0.5
0 50 100 150 200 250
Composition ratio ... i a l
orientation along (111). We did not find any peaks of other
phases such as CuO even after the H₂O₂ treatment. This

(AES) analysis and scanning electron microscopy (SEM)
observation were carried out using the model JEOL JAMP-
9500F at probe voltage 10 kV. Argon ion etching with an
acceleration voltage of 2 kV and a ion current of 2.6 μA was
used to sputter the film surface. The composition ratio was
calculated by using a standard CuO compound. Furthermore,
the X-ray diffraction (XRD) measurement was carried out
by the Rigaku SmatLab X-ray diffractometer using Cu Kα₁
radiation at 40 kV/30 mA.

3. Results and Discussion

3.1. Treatment at Room Temperature. Figures 1(a) and 1(b)
show the SEM images of the untreated Cu₂O thin film and the
film treated at room temperature for 7 hours, respectively, and
Figures 1(c) and 1(d) show those of the film treated at room
temperature for 24 hours (with different magnifications). The
thickness of the as-deposited film was about 0.6 μm, and
the thickness was not changed significantly by the room
temperature treatment. No significant change was observed
for the surface of the sample treated for 7 hours compared
with the untreated film. However, for the 24-hour-treated
film, we can see bumps on the surface, where adhesion to
the substrate was apparently weakened with narrow cracks
generated on the surface, as shown in Figure 1(d).

Figure 2 shows the change of Cu/O composition ratios
with the sputtering time. For the as-deposited sample, the
composition ratio of the films is 1.8. For the sample treated
for 24 hours, the Cu/O composition ratio near the surface is
close to unity, and the composition ratio was increased with
increasing sputtering thickness. Thus, the surface of the Cu₂O
thin film was converted to CuO by the H₂O₂ treatment while
the inner portion of the samples was not affected.

Figure 3 shows the XRD spectra of the as-deposited and
H₂O₂-treated samples. The Cu₂O films are polycrystalline
and have cubic-cuprite structure with weak preferential
orientation along (111). We did not find any peaks of other
phases such as CuO even after the H₂O₂ treatment. This
would be because the CuO layer formed near the surface by the H₂O₂ treatment is too thin or does not have crystallinity.

Figure 4 shows the comparative results of the optical transmission of the Cu₂O sample treated for 7, 16, and 24 hours at room temperature. From the figure, it was confirmed that by extending the treatment time, the optical absorption is increased further. The sample absorbed light of wavelength less than 560 nm completely; the absorption edge is around 1000 nm when treated for 24 hour. Figure 5 shows the plot of (αhν)² versus hν. The apparent band gap can be estimated from extrapolation of a straight line part of the curve to the photon energy axis. For the samples treated for 24 hours, the band gap seems to be about 1.6 eV. From these results, we can conclude that Cu₂O thin films can be oxidized to have a smaller band gap by the H₂O₂ treatment at room temperature.

Figure 6(a) shows the PEC measurement results of the as-deposited sample. In the PEC measurement, the current was measured under chopped illumination. Relative change in the carrier concentration due to the illumination is larger for the minority carriers than for the majority carriers, and thus the photocurrent due to photo-generated minority carriers is observed clearly. Thus negative photocurrent is generally significant for a p-type sample, and positive photocurrent for an n-type sample. For the as-deposited samples, the negative photocurrent (under cathodic bias) and the positive photocurrent (under anodic bias) were observed, and thus the conduction type is close to intrinsic. Figure 6(b) shows the PEC measurement results of the sample treated for 24 hours. The photocurrent is negative even under anodic bias. This indicates that the treated sample exhibited clearer p-type character.

3.2. Treatment at Higher Temperatures. As shown in the previous section, the H₂O₂ treatment at room temperature is effective to modify the composition and enhance the optical absorption, but the treatment time needs to be longer than several hours. For the practical application, shorter treatment time would be preferable. Thus we tried the H₂O₂ treatment at higher temperatures.

Figure 7 shows the optical transmission of the samples treated at 60°C for 30 minutes. From the figure, we can conclude that the optical absorption of the sample was not increased by the H₂O₂ treatment at 60°C for 30 min.

Figures 8(a) and 8(b) show the SEM images of the thin films treated at 60°C and at 90°C for 30 min, respectively. It can be seen that the surface of the sample was partly broken (holes were formed) when treated at 90°C. Similar damages were observed for the film treated at 70°C and 80°C. Thus, we can conclude that the H₂O₂ treatment at higher temperatures is not useful for the modification of the film composition.

4. Conclusions

Cu₂O thin films have been deposited electrochemically onto ITO-coated glass. Then, the thin film was dipped in H₂O₂ (30%) at fixed temperature for oxidization. By the H₂O₂ treatment at room temperature for 7, 16, and 24 hours, the oxygen content of surface was increased and the apparent band gap of the samples was decreased. Thus this treatment can be used to enhance optical absorption of the Cu₂O films.
Figure 8: SEM images of (a) the film treated at 60°C and (b) the film treated at 90°C for 30 min.

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References


