Dye-Sensitized Solar Cells Based on ZnO Films and Natural Dyes

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Abstract This work employs extracts from Walnuts, Rhubarb, and Pomegranate as natural dyes for fabrication of dye-sensitized solar cells (DSSCs). ZnO nanoparticles with crystallite mean value 12 nm as confirmed from XRD data have been synthesized at pH 12. SEM picture of the ZnO powder reveals homogeneous and well defined nanoparticles with size of about 15 nm. TEM micrograph shows that the powder has a porous agglomerate structure consisting mainly of spherical crystalline particles with about 15–20 nm diameter. ZnO films were deposited on Fluorinated Tin Oxide (FTO) coated glass sheets followed by sintering at 450°C. The samples were sensitized by soaking in the dye solution. A Graphite slab was used as a back electrode, and a redox was employed as an electrolyte. Each cell was illuminated with light intensities in the range 40,000–100,000 Lux to measure the photovoltaic parameters. The experimental results shows that the highest $I_{sc}$ value is obtained from the DSSC sensitized with Rhubarb extract while the highest $V_{oc}$ value is obtained from the DSSC sensitized with the Walnuts extract. The $P_{max}$ of the DSSC sensitized by the Walnuts extract is greater than those sensitized by Rhubarb and Pomegranate extracts. Moreover, the open circuit voltage $V_{oc}$ decay was found to closely follow a pseudo single exponential form.

Keywords Dye-Sensitized Solar Cell, Natural Dye, Walnuts, Rhubarb, Pomegranate

1. Introduction

Direct utilization of solar radiation to produce electricity is a steadily growing field and solar cell technology has received huge potential by physicists and engineers. Solar cells have many advantages over other energy techniques such as avoiding transmission losses, operating without noise, and requiring very little maintenance. Moreover, there are no toxic and greenhouse gas emissions in solar cell systems. Despite the considerable development in the last three decades, the high cost and the low efficiency of solar cells have been the main reasons behind the limited implementation of the technology.

Dye-sensitized solar cells (DSSCs) are promising, relatively low cost, and green energy photovoltaic devices[1-8]. A DSSC is usually comprised of a transparent conducting glass electrode coated with porous nanocrystalline wide-band gap semiconductor oxide film. The nanostructured oxide films are particularly attractive for DSSCs as they provide a large surface area for dye anchoring[9].

Dye molecules are attached to the surface of the nanostructured semiconductor oxide film. Photoexcitation takes place in the dye, and the photogenerated charges are separated at the dye-oxide interface. Moreover, a DSSC also comprises an electrolyte containing a reduction-oxidation couple such as $\Gamma^-/\Gamma_3^-$ and a catalyst coated counter-electrode. During illumination, the cell produces voltage over and current through an external load connected to the electrodes.

It is highly significant to reduce the charge traps in the semiconductor oxide to speed up the charge transport since the crystalline quality of the oxide film has a great effect on the charge transport. The amount of light entering the cell and the photocurrent extraction are determined by the transparent electrode. The selection of the proper semiconductor oxide and a corresponding transparent electrode is critical in the device design to achieve efficient light harvesting, charge separation, and current extraction.

The use of dye-sensitization in photovoltaics has received a great interest after the breakthrough achieved by Grätzel et al. in the early 1990’s. They developed a DSSC with energy conversion efficiency exceeding 7% in 1991[10] and 11.4% in 2001[11] by combining nanostructured electrodes to efficient charge injection dyes. Since then, TiO₂ nanoparticle films have been widely investigated for DSSCs. However, TiO₂ films also have some defects such as the lack of a large enough energy barriers between the interface of the films and electrolytes and the existence of plenty of elec-
tron-trapped surface states, which is the cause of the recombination\cite{12}. The photocatalytic activity of TiO2 is so high under the UV radiation of natural sun-light that organic materials in DSSCs may be decomposed during outdoor use, resulting in long-term reliability problems for the conversion efficiency. Hence, some investigations have been turned to ZnO films, which have a much lower photocatalytic activity and less electron-trapped surface states\cite{13}. However, the efficiency of ZnO films is still lower than that of TiO2.

The dye that is used as a photosensitizer plays an important role in the operation of DSSCs. The efficiency of the cell is critically dependent on the absorption spectrum of the dye and the anchorage of the dye to the surface of the semiconductor. Much work has been concentrated on organic dyes and organic metal complexes. On the other hand, natural dyes extracted from fruits and flowers have attracted the attention of many researchers\cite{14-16}, and many natural dyes have been proven to be efficient dyes as photosensitizers in DSSCs.

In this work, preparation of ZnO nanoparticles and characterization using XRD, SEM, and TEM is presented. Also, three natural dyes extracted from Walnuts, Rhubarb, and Pomegranate were investigated for dye sensitization. The absorption spectra of these extracts were studied. Besides, the performance of the DSSCs fabricated using ZnO nanoparticle films and extracts of these natural dyes were investigated.

2. Experimental

2.1. Synthesis and Characterization of ZnO Nanoparticles

ZnO nanoparticles used in this article have been synthesized using a method described by Rani et al.\cite{17} by dissolving 4.4 g of zinc acetate dihydrate, reagent Zn(CH3COO)2·2H2O (ACROS) in 100 mL of methanol to obtain a 0.2 M sol which was stirred overnight. A 3 M aqueous solution of sodium hydroxide (NaOH) was added dropwise to the solution under vigorous stirring. A white suspension was obtained and left under stirring for 12 h. A pH meter was used to adjust the pH of the solution was adjusted at pH=12 and then dried at 100 °C overnight for drying purposes. Then, an amount of 0.2 g of polyethylene glycol and 4 mL distilled water were added to two grams of ZnO nano-powder in order to form a paste. The resulted paste was dispersed mechanically using a home-made mill until a very soft uniform paste was obtained. Then, an amount of 5mL of ethanol was added to the paste. The ZnO samples were sensitized by soaking the nanoparticle films and extracts of these natural dyes for 30 min. then left overnight to cool down. The ZnO powder was coated on the FTO coated glass sheets. The samples were dried using a hot plate at 80 °C followed by sintering at 450 °C for 30 min. then left overnight to cool down. The ZnO samples were sensitized by soaking the samples in the dye solutions overnight at room temperature. Then, they were washed using methyl alcohol to remove the leftover dye. Finally, they were dried using a hot plate.

2.2. Natural Dye Extraction

Walnuts, Rhubarb, and Pomegranate were boiled in distilled water for many hours until solid extracts were obtained. The solid extracts were then left in an oven at 70 °C overnight for drying purposes. Then a small amount from each extract has been dissolved in ethylene glycol.

2.3. Preparation of ZnO Electrodes

Fluorinated Tin Oxide SnO2:F (FTO, K glass R =8Ω) coated glass sheets (1cm x 1cm), were first cleaned in a detergent solution using an ultrasonic bath for 15 min, rinsed with water and ethanol, and then dried. An amount of 0.2 g of polyethylene glycol and 4 mL distilled water were added to two grams of ZnO nano-powder in order to form a paste. The resulted paste was dispersed mechanically using a home-made mill until a very soft uniform paste was obtained. Then, an amount of 5mL of ethanol was added to the paste to obtain suitable viscosity for coating purposes. The ZnO paste was coated on the FTO coated glass sheets. The samples were dried using a hot plate at 80 °C followed by sintering at 450 °C for 30 min. then left overnight to cool down. The ZnO samples were sensitized by soaking the samples in the dye solutions overnight at room temperature. Then, they were washed using methyl alcohol to remove the leftover dye. Finally, they were dried using a hot plate.

![Figure 1](image.png)
3. Results and Discussions

3.1. Powder Structure

XRD pattern of the prepared ZnO powder is shown in Fig. 1. It is clear from the figure that the powder is highly crystalline and that its structure is in accordance with the typical wurtzite hexagonal structure (JCPDS No. 0036-1451, Wurtzite-0324982). The crystallite mean value is found to be 12 nm and the crystallite size calculated for the (101) and (002) peaks are found to be 12 nm and 11 nm, respectively. The lattice dimensions are found to be [a = b = 3.253(0) Å, and c = 5.213(4) Å].

3.2. Surface morphology and BET surface area analyses of ZnO powder

The key points for DSSC application are surface morphology and Brunauer–Emmett–Teller (BET) surface area. Thus it is significant to study the surface morphology and BET surface area of the ZnO powder.

Figure 2. SEM images of the nanostructured ZnO powder synthesized at pH 12. The scale bars correspond to 100 nm.

Figure 3. TEM image of the nanostructured ZnO powder synthesized at pH 12 and dried at 100°C for 12 h. The scale bar corresponds to 20 nm.

The BET surface area of the ZnO powder prepared at pH 12 and dried at 100°C was 42.3 m² g⁻¹. This value is higher than the 23.8 m² g⁻¹ reported by Aghababazadeh et al.[18] for powders obtained by mechanochemical processing and sintered at 400°C. This difference can be due to agglomeration of nanoparticles during sintering which accordingly reduce the surface area.

Figure 2 illustrates a SEM picture for the ZnO powder. The figure shows obviously that the nanoparticles are homogeneous and well defined with size of about 15 nm.

TEM micrograph of the powder is shown in Fig. 3. It is clear that the powder has a porous agglomerate structure consisting mainly of spherical crystalline particles with about 15–20 nm diameter.

3.3. Absorption spectra

The UV–vis absorption spectra of the three dyes were measured using a UV–vis spectrophotometer (thermo light Genesys 6). Figure 4 shows the representative UV–vis absorption spectra for the extracts of Walnuts, Rhubarb, and Pomegranate dissolved in polyethylene glycol.

Figure 4. Light absorption spectra of (A) Walnuts extract, (B) Rhubarb extract, and (C) Pomegranate extract dissolved in polyethylene glycol.

3.4. Electrical Measurements

A Graphite slab was used as a back electrode. A drop of iodine–iodide redox was inserted between the back electrode and the dyed ZnO. The I–V characteristic curves at different light intensities were conducted using National instruments data acquisition card (USP NI 6251). The power P was then calculated and plotted as a function of voltage V and the maximum power point was determined. The I–V curves of the fabricated DSSCs were obtained by applying an external reverse bias to each cell and measuring the generated photocurrent under different intensities of white light irradiation.

The incident light intensity in the range 40,000–100,000 Lux was measured and then converted to W/m² using the simplified relationship[19,20]:

\[
\frac{E_{\text{rad}}(\text{Lux})}{120,000} = \frac{E_{\text{rad}}(W/m^2)}{1,000}
\]

or
The $I-V$ characteristic curves of DSSCs sensitized with the extracts of Walnuts, Rhubarb, and Pomegranate are shown in Figs. 5, 6, and 7, respectively. From these figures the values of short-circuit current ($I_{sc}$) and open-circuit voltage ($V_{oc}$) of the cells can be obtained directly using the $I-V$ data corresponding to $10^5$ Lux luminance. These results are tabulated in Table I for the three devices. The cell output power has been calculated as $P = I \cdot V$ using the $I-V$ data corresponding to $10^5$ Lux intensity and plotted as a function of $V$ in Fig. 8. The maximum power ($P_{max}$) of the DSSCs for each cell is then obtained from Fig. 8. The current ($I_{mp}$) and the voltage ($V_{mp}$) corresponding to the maximum power point are then obtained. The values of the fill factor ($FF$) are calculated according to the following equation

$$FF = \frac{I_{mp} \cdot V_{mp}}{I_{sc} \cdot V_{oc}}$$  (3)

The efficiency $\eta$ is calculated as follows:

$$\eta = \frac{P_{max} \cdot V_{oc}}{E_{rad}}$$  (4)

The highest $I_{sc}$ value is obtained from the DSSC sensitized with Rhubarb while the highest $V_{oc}$ value is obtained from the DSSC sensitized with the Walnuts. The $P_{max}$ of the DSSC sensitized by the natural dye extract of Walnuts (10.14 µW) is greater than those sensitized by Rhubarb (8.75 µW) and Pomegranate (3.85 µW). The lowest $P_{max}$ value come from the DSSC sensitized with Pomegranate which may be attributed to weak bonding between the dye molecule and ZnO particles. Good photo-to-electric conversion ability in a DSSC is strongly dependent on available bonds between the dye molecules and ZnO particles, through which electrons can transport from excited dye molecules to ZnO film. Obviously, as a photosensitizer, the interaction and bond between sensitizer (dye) and sensitized film (ZnO) is very important in enhancing the photoelectric conversion efficiency of a DSSC.
Table 1. Photoelectrical parameters corresponding to $E_{ad} = 100,000$ Lux of DSSCs sensitized by natural dyes

| Natural dye  | $V_{oc}$ (V) | $I_{sc}$ (μA) | $P_{max}$ (μW) | $V_{mp}$ (V) | $I_{mp}$ (μA) | FF | η (%) |
|--------------|--------------|--------------|----------------|--------------|--------------|----|------|
| Walnuts      | 0.304        | 73.3         | 10.14          | 0.198        | 49.1         | 0.39| 0.0104|
| Rhubarb      | 0.238        | 82.65        | 8.75           | 0.157        | 55           | 0.44| 0.0104|
| Pomegranate  | 0.159        | 63.2         | 3.85           | 0.096        | 37.4         | 0.36| 0.0043|

Table 2. The coefficients of the $V_{oc}$ decay fitting equation for cells with the three extracts

| Extract     | $V_0$           | $A_1$       | $t_1$       | $\tau_n$ | $k_{rec}$ |
|-------------|-----------------|-------------|-------------|----------|-----------|
| Walnuts     | 0.0872305       | -0.1703259  | -2.3029668  | 25.4855061 | 0.03924   |
| Rhubarb     | 0.0421363       | -0.3747801  | -6.8775522  | 8.1777440  | 0.12228   |
| Pomegranate | 0.04292611      | -0.1405300  | 0.3965477   | 3.1397002  | 0.31850   |

Figure 9 shows the open circuit voltage $V_{oc}$ decay due to turning the light source off after illuminating the sample for a few minutes. Normally, the photovoltage decay closely follows a pseudo single exponential form [21], thus the recombination rate constant, $k_{rec}$, can be extracted from the slope of the semi-logarithmic plot. Since the back reaction is usually taken as the pseudo-first order reaction, $k_{rec}$ is related to electron lifetime $\tau_n$ by the relation

$$k_{rec} = \frac{1}{\tau_n}$$  \hspace{1cm} (5)

The data is fitted with the following equation [21]

$$V = V_0 - A_1 \exp[-(t-t_1)/\tau_n]$$  \hspace{1cm} (6)

Values of the various coefficients of this fitting equation for the three cells are shown in table II.

4. Conclusions

Preparation and characterization of ZnO nanoparticles using XRD, SEM, and TEM are presented in this work. The SEM picture of the ZnO powder shows that the nanoparticles are homogeneous with size of about 15 nm. The TEM micrograph indicates that the powder has a porous agglomerate structure consisting mainly of spherical crystalline particles with about 15–20 nm diameter. Also, we have used three natural dye extracts as photosensitizers to assemble some DSSCs. A short circuit current from 63.2 μA to 82.65 μA, an open circuit voltage from 0.159 V to 0.304 V, a fill factor from 0.36 to 0.44, $P_{max}$ from 3.85 μW to 10.14 μW, and an efficiency from 0.00431% to 0.0104% were obtained from these cells. Based on our investigation, it was found that Walnuts possesses the best photosensitization effect among three extracts of natural dyes studied. Natural dyes as alternative sensitizers for DSSCs are expected to be promising because of many reasons such as the simple preparation technique and low cost.

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