Performance of natural-dye-sensitized solar cells by ZnO nanorod and nanowall enhanced photoelectrodes

Saif Saadaoui¹, Mohamed Aziz Ben Youssef¹, Moufida Ben Karoui¹,², Rached Gharbi*¹, Emanuele Smecca³, Vincenzina Strano⁴, Salvo Mirabella⁴, Alessandra Alberti³ and Rosaria A. Puglisi³

Abstract

In this work, two natural dyes extracted from henna and mallow plants with a maximum absorbance at 665 nm were studied and used as sensitizers in the fabrication of dye-sensitized solar cells (DSSCs). Fourier transform infrared (FTIR) spectra of the extract revealed the presence of anchoring groups and coloring constituents. Two different structures were prepared by chemical bath deposition (CBD) using zinc oxide (ZnO) layers to obtain ZnO nanowall (NW) or nanorod (NR) layers employed as a thin film at the photoanode side of the DSSC. The ZnO layers were annealed at different temperatures under various gas sources. Indeed, the forming gas (FG) (N₂/H₂ 95:5) was found to enhance the conductivity by a factor of 10³ compared to nitrogen (N₂) or oxygen (O₂) annealing gas. The NR width varied between 40 and 100 nm and the length from 500 to 1000 nm, depending on the growth time. The obtained NWs had a length of 850 nm. The properties of the developed ZnO NW and NR layers with different thicknesses and their effect on the photovoltaic parameters were studied. An internal coverage of the ZnO NWs was also applied by the deposition of a thin TiO₂ layer by reactive sputtering to improve the cell performance. The application of this layer increased the overall short circuit current $J_{sc}$ by seven times from $2.45 \times 10^{-3} \text{ mA/cm}^2$ to $1.70 \times 10^{-2} \text{ mA/cm}^2$. 

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Introduction

Energy demand has increased rapidly during the last forty years to reach a growth rate of 1.8% per year [1]. To satisfy this growing need, it is necessary to find new sources of renewable energy. For instance, photovoltaic (PV) technologies offer a promising green industry for the future power demand. Among these technological resources, dye-sensitized solar cells (DSSCs) have shown good performance since their first demonstration by O’Regan and Grätzel in 1991 [2]. Figure 1a shows the standard structure of DSSC: The first part of this structure represents the photoelectrode, composed of a wide band gap semiconductor thin layer coated on transparent conducting oxide (TCO) films. A mesoporous film layer is used as a semiconductor in the photoelectrode. Because of its stability and easy synthesis, titanium dioxide (TiO$_2$) is mostly used as the semiconductor in DSSCs [2-5]. Besides, the TiO$_2$ offers high electronic mobility for photogenerated electron collection, a suitable band gap, which adapts to the injection of the electrons of most studied dyes, and high surface area to enhance the dye loading by anchoring the dye [6,7].

Zinc oxide (ZnO) has been studied as a mesoporous wide band gap semiconductor for use in DSSCs. It presents itself in the form of different morphological nanostructures, such as nanorods, nanocrystals, nanowires, nanotubes and nanowalls that can be exploited to optimize the dye loading [6-9]. The main purpose of the photoelectrode is to collect the injected electrons from the excited dye absorbed by the semiconductor layer and to transport the electrons to the external load. The second main part of the structure is the counter electrode formed by a thin layer of platinum coated on TCO to accelerate the redox reaction with the electrolyte solution, which constitutes the third main part of the cell [3].

The internal process starts with the excitation of the sensitizer (S) through the absorption of a photon to obtain an excited sensitizer ($S^*$). The latter injects an electron into the conduction band of the semiconductor. Then, it stays in the oxidized state ($S^+$), as mentioned in Figure 1b. The injected electron reaches the external load through the semiconductor and the TCO layers. When arriving at the counter electrode, the electron reduces the redox solution and regenerates the sensitizer to complete the reaction [2,10]. Different high-ranked commercial dyes, such as N719 or N3, have been widely studied in the literature and their performance has been proven. Because these organometallic dyes contain environmental pollutants, such as heavy metals [6], recent studies have focused on finding a metal-free organic dye with high performance [8,11-13]. Indeed, several studies were carried out using natural dyes, such as spinach and red turnip as a sensitizer. However, their efficiency is lower than that of commercial organometallic dyes. In this work, we investigate two different natural dyes extracted from henna and mallow plants. We discuss also their application to different semiconductor structures.

The photoelectrode is regarded as an important part in the DSSC where it represents the electron generator of the cell. Solar cell parameters, such as open-circuit voltage ($V_{oc}$), short-circuit current density ($J_{sc}$) and fill factor (FF), are the most significant parameters used to evaluate the enhancement of the power conversion efficiency (PCE) of solar cells [10]. The short circuit current density, $J_{sc}$, depends essentially on dye adsorption, the collection efficiency in the semiconductor thin layer film and the efficiency of the collected dye molecules [3,14,15]. The adsorption of the dye can be improved by various means, such as increasing the thickness and/or the porosity of the photoelectrode or by using organized structures, such as nanowalls or nanorods. The $V_{oc}$ can be improved by modifying the energy difference between the Fermi level ($E_F$) of the semiconductor potential and redox potential ($E_{redox}$) of the electrolyte [10].
Results and Discussion

Dye analysis

In order to understand the structure of natural dye molecules and to determine the main elements responsible for absorbing light, we used FTIR spectroscopy techniques and ultraviolet–visible spectrophotometry (UV–vis) to characterize dyes extracted from henna and mallow powder as shown in Figure 2.

The scopoletin molecule, shown in Figure 3b [15-17], is the major component in the mallow dye. From the FTIR spectrum of the mallow dye (Figure 3a), it is clear that mallow has a primary O–H bond associated with the peak at 3372 cm$^{-1}$. We notice also the existence of aliphatic groups, CH$_2$ and CH$_3$ (2904 cm$^{-1}$, 2949 cm$^{-1}$, 2990 cm$^{-1}$), an ether function ($\nu$C–O) at 1055 cm$^{-1}$, and a regular deformation band $\delta$C$_{sp2}$-H at 668 cm$^{-1}$.

2-Hydroxy-1,4-napthaquinone (C$_{10}$H$_6$O$_3$), frequently called lawsone [15,18], is one of the constituents of the natural dye henna in addition to other compounds, such as gallic acid, sterols, resin, tannin and coumarins [19]. Lawsone (Figure 4b), the main component of henna extract, has been used as a dye in the cosmetics industry. The most significant feature of the lawsone molecule is its ability to absorb visible light between 400 and 600 nm.

The FTIR spectrum of henna extract, presented in Figure 4a, shows the existence of three valence bands associated with C=C aromatics at 1338 cm$^{-1}$, 1421 cm$^{-1}$ and 1458 cm$^{-1}$, and also a stretching absorption band at 3368 cm$^{-1}$ corresponding to the vibration of the associated O–H bond.

The UV–vis measurements of the extracted dyes of henna and mallow powders diluted in ethanol are given in Figure 5. Both of the studied dyes show two remarkable peaks: the first around 470 nm (blue light), and the second at 665 nm (red light). Both peaks correspond to the mixture of chlorophyll [20]. In the region from 525 nm to 625 nm, the mallow dye spectrum reveals two small peaks at 536 nm and 608 nm. These peaks could serve to increase the charge-transfer reaction under sun illumination in the final DSSC [21].

Morphological results of the layers

In order to estimate the capacity of the dye loading, scanning electron microscopy (SEM) analysis was carried out to study the porosity and the morphology of the layers prepared by the CBD technique [8,22].

Figure 6a gives the SEM images illustrating the morphology of the ZnO NRs annealed in forming gas (FG) (N$_2$/H$_2$, 95:5). From this figure, we observe that nanorods cover the entire surface of the fluorine-doped tin oxide (FTO) conductive glass and
have a hexagonal shape with uniform size and length [23]. The width of the NRs varies between 40 and 100 nm, while their length ranges from 500 to 1000 nm, depending on growth time, as shown in Figure 6b and mentioned by Iwu et al. [8]. In Figure 6c and Figure 6d, we give the SEM images of the ZnO NR after being immersed in henna and mallow dye, respectively. The images show the formation of a thin layer between the NRs. This layering effect may be due to a reaction between the ZnO nanoparticles and the solvent in the dye solution. From these images, we notice that the number of individual hexagonal NRs is reduced and their shapes are no longer visible. This morphological change can be explained by aggregation of the dye and Zn\(^{2+}\) when the films were immersed in henna and mallow dye [24].

Figure 7a shows the morphology of the ZnO NWs annealed in FG. The NWs intersect at different angles, creating hollow spaces, acting to increase the porosity of the layer, which may improve the dye loading as observed by Polkoo et al. [25]. By analyzing the cross-section of the ZnO NWs, we obtained a NW vertical length of around 850 nm, as shown in Figure 7b. In Figure 7c, we give the images of the ZnO NWs after being immersed in mallow dye. We notice that dye molecules cover the NWs without affecting their overall shape, as compared to NRs. The formed layer does not appear clearly in the surface of the NWs. However, there is a small difference in the intersection angles between walls. After the immersion, Figure 7c shows less acute angles of wall intersections compared to those presented in Figure 7b. This variation in the angle sharpness shows that a thin layer is formed in the different intersections and not in the walls.

Figure 8a illustrates the magnified image of the ZnO NWs over which a TiO\(_2\) layer was deposited by the sputtering technique. We observe that the nanowall vertical length is about 1000 nm. Besides, the shape is arranged in a more vertical orientation as compared to the nanowalls without the TiO\(_2\) blocking layer. The main idea is to cover the ZnO NWs with a thin layer of TiO\(_2\) (about 150 nm of thickness), which enhances the electron transport by exploiting the high mobility and diffusion rate of the ZnO. This covering layer also increases the efficient electron diffusion from the TiO\(_2\) layer to the ZnO NWs. Using this structure, the electrons were quickly transported from the sensitizer to the TiO\(_2\) to reach the FTO (Figure 8b). This solution created an energy barrier allowing the reduction of the charge recombination [26].

**Optical characterization**

Based on the SEM analysis, we conclude that the growth time can influence the layer thickness, which affects the light absorbance [9] and conductivity [27]. For these reasons, after the
Figure 6: SEM images of ZnO NRs (a) before immersion in dye, (b) cross-section view of ZnO NRs, (c) after immersion in henna dye and (d) after immersion in mallow dye.

Figure 7: SEM images of ZnO NWs (a) cross-section view of ZnO NWs (b) before immersion in dye and (c) after immersion in mallow dye.
growth of the ZnO NR layers on FTO, they were annealed in O₂, FG and N₂ gas at 300 °C.

As shown in Table 1, the sheet resistance of the ZnO NRs depends on the used annealing gas. By using O₂, we obtained the highest sheet resistance with a ratio of 10^3 compared to those prepared under N₂ or FG. The best sheet resistance was measured in the case of annealing in FG.

Figure 9 shows the ZnO NR layer absorbance in different conditions of the annealing gas and the growth time annealed at 200 °C. We see that the sample without annealing gas has the lowest absorbance, which implies the necessity of the annealing step. The sample, annealed in FG and grown for 45 minutes, has the highest absorbance in the region between 300 nm and 600 nm compared to those annealed in the same gas condition. However, the growth time influences the absorbance, which indicates that thin ZnO NR layers absorb more in this region.

### Table 1: Sheet resistance of ZnO layers for different ambient annealing gases and J–V measurement results of the assembled cells.

| Cell N° | Dye     | Photoanode layer | Annealing gas | Annealing temp. (°C) | Sheet resistance (Ω/sq) | Growth time (min) | J_SC (mA/cm²) | V_OC (V) | FF (%) |
|---------|---------|------------------|---------------|----------------------|-------------------------|-------------------|--------------|----------|--------|
| 1       | henna   | ZnO nanorods     | FG            | 200                  | 5.3 × 10^3              | 90                | 1.26 × 10^-1 | 0.370    | 27     |
| 2       | mallow  | ZnO nanorods     | FG            | 90                   | 6.3 × 10^-1              | 90                | 1.55 × 10^-1 | 0.425    | 27     |
| 3       | FG      | FG               | FG            | 45                   | 1.49 × 10^-1              | 45                | 1.49 × 10^-1 | 0.428    | 29     |
| 4       | N₂      | as deposited     | 90 × 10^3     | 90                   | 5.35 × 10^-2              | 90                | 5.35 × 10^-2 | 0.136    | 30     |
| 5       | O₂      | as deposited     | 600 × 10^3    | 90                   | 4.24 × 10^-2              | 90                | 4.24 × 10^-2 | 0.379    | 27     |
| 6       | TiO₂ + ZnO nanowalls | FG | 200 | 1.4 × 10^-5 | – | – | – | – | – |
| 7       | TiO₂ + ZnO nanowalls | FG | 300 | 5.3 × 10^-3 | 5 | 8.02 × 10^-3 | 0.426 | 34 |
| 8       | as deposited | FG | 200 | 6.00 × 10^-3 | 1.23 × 10^-2 | 0.413 | 29 |
| 9       | as deposited | FG | 200 | 5.3 × 10^-3 | 2.45 × 10^-3 | 0.169 | 29 |
| 10      | ZnO nanowalls | FG | 200 | 5.3 × 10^-3 | 2.45 × 10^-3 | 0.169 | 29 |
The second UV–vis measurements were performed after the immersion of the ZnO NR layer in dye. Figure 10 demonstrates the absorbance spectra versus the variation in time and the concentration of mallow dye. The initial concentration is 3.25 g of mallow powder stirred in 40 mL of ethanol. By multiplying the concentration three times, we increase the loaded dye in the ZnO NR layer and we clearly observed the absorbance peak related to the collected dye in the photoanode side at 665 nm.

The prepared photoanode was assembled with a platinized counter electrode. The current–voltage (I–V) measurements were carried out using a Keithley 4200 device under AM1.5 conditions.

Figure 11a shows the I–V measurement response of the DSSC cells assembled with a ZnO NW layer on photoanodes annealed at different temperatures with two different structures formed by using only ZnO NWs in the photoanode and with applying a TiO₂ blocking layer deposited by sputtering (Table 1). The mallow dye was chosen because of its optimum absorbance in the visible region. The DSSC, based on NWs with a TiO₂ blocking layer annealed at 200 °C in FG, presents the highest $V_{OC}$ compared to the NW cell without a blocking layer prepared under the same thermal conditions, where the $V_{OC}$ increases from 0.169 to 0.380 V. This result verifies the main role of the blocking layer in decreasing the electron–hole recombination and the negative shift of the conduction band edge due to the increased electron density in TiO₂, as previously reported by Yao et al. [26].

We also noticed that the increase of the temperature treatment from 200 °C to 300 °C in FG maintains the same sheet resistance of the NW layer at 5.3 Ω/sq. However, the short circuit current density of the cell decreases from $1.70 \times 10^{-2}$ mA/cm² to $8.02 \times 10^{-3}$ mA/cm². This decrease can be explained by the fact that when increasing the annealing temperature, the stability of the ZnO nanoparticles was considerably affected, which resulted in higher reaction with the dye solution.

However, the DSSC assembled without an annealed photoanode side gave a slightly higher short circuit current and smaller variation of 5% in the FF as compared to the cell annealed at 300 °C using a TiO₂ blocking layer as shown in Figure 11. This increase can be explained by the high variation in the sheet resistance of the ZnO NWs in the photoanode side [28].

For the NR-based photoanode cells, we annealed using different gas sources. The forming gas provided the most conductive layers as compared to N₂ and O₂, as discussed above. More-
over, the sheet resistance increased from $5.3 \times 10^3 \ \Omega/\text{sq}$, when annealing in FG, to $90 \times 10^3 \ \Omega/\text{sq}$ in N$_2$. It also increased considerably in O$_2$ to reach $1.4 \times 10^3 \ \Omega/\text{sq}$ due to the ZnO ionization, which decreased the volume of Zn interstitials, resulting in a decrease of the carrier concentration [29].

Therefore, by comparing the two used sensitizers, under the same thermal conditions (200 °C) and at the same growth time (90 minutes), we noticed that the electrical performance was almost identical and they have the same FF value equal to 27%. By changing the growth time from 90 to 45 minutes, the FF value rises slightly from 27% to 29%. This increase implies that the reduction in the length of NRs, caused by the decrease of growth time, augments the internal shunt resistance of the cell, resulting in a small decrease of the short circuit current, $J_{sc}$, from 0.155 mA/cm$^2$ to 0.149 mA/cm$^2$ and a small increase of $V_{OC}$ from 0.425 to 0.428 V [30].

The efficiency of the ZnO NR- and NW-based DSSCs is lower than that of conventional DSSCs. The main reason for this difference in performance is related to the nature of the photo-sensitizer used in the present investigation.

Furthermore, the small thickness of the ZnO NRs and NWs can be a major reason for this degradation in the efficiency. However, the chemical reaction of the ZnO nanostructures when immersed in dye solution was carried out at the surface of the layer, resulting in Zn–dye complex aggregates affecting the electron injection efficiency [31].

**Conclusion**

In this work, we reported the fabrication and characterization of natural DSSCs with two different ZnO layer structures. We analyzed two natural dye sensitizers extracted from henna and mallow plants in different concentrations and at various immersion times. Furthermore, we analyzed the properties of the different used dyes and we reported the most anchoring-dominant molecules. By studying different gases used in the annealing process, we noted that the FG gave the most conductive layers compared to N$_2$ or O$_2$, which was confirmed in the photovoltaic parameters of the final cells. We also highlight the fact that the use of a TiO$_2$ blocking layer in the structure ZnO NW/TiO$_2$/dye increases the photovoltaic performance of the realized DSSC by enhancing the morphological and physical properties of the photoanode.

**Experimental**

Both ZnO NRs and NWs were grown on FTO substrates ($1.5 \times 1.5$ cm$^2$) by CBD [8,22]. Before the ZnO NR preparation, a seed layer of ZnO crystallites was deposited by spin coating (1000 rpm, 60 s) using a solution of 5 mM zinc acetate dihydrate in ethanol, followed by 20 min annealing in air on a hot plate (at nominal 240 °C).

However, in the case of the ZnO NWs, the seed layer was replaced by an Al film (100 nm thick). The basic Al films were sonicated in soapy water, water, ethanol and acetone before being used. After the pre-layer deposition, we prepared the CBD bath [8,22] by mixing two solutions: a 100 mL solution of variable hexamethylenetetramine (HMTA) concentration in de-ionized (DI) water was well stirred and preheated at 90 °C. Then, it was added to a 100 mL solution of zinc nitrate hexahydrate in DI water.

The zinc nitrate concentration in the CBD bath was fixed at 25 mM both for ZnO NR and NW growth. The HMTA final concentration was set at 25 mM and 50 mM for three different growth times (45 min, 60 min and 90 min) in case of ZnO NRs. In the synthesis of NWs, the growth time was set to 5 min. The HMTA concentration was 25 mM. Moreover, 16 mM of ammonium hydroxide (NH$_4$F) was added to the synthetic solution since its presence enhanced the quality of the NWs [22].

The DSSC preparation started by covering an active area of $0.5 \times 0.5$ cm$^2$ TCO glass with a thin film ZnO layer (NRs, NWs, or TiO$_2$ on NWs). After the annealing step, the prepared film was submerged in a solution containing the natural dye. The used natural dyes were prepared using ethanol as the solvent. The electrode and the counter electrode were sealed with thermoplastic sealing film (Meltonix 1170-60PF from SOLARONIX), followed by injection of (Iodolyte AN-50) the electrolyte (SOLARONIX).

The used henna and mallow powders were prepared in-house by drying henna and mallow plants. Afterwards, the dried plants were milled and sieved to obtain the final powder.

Current–voltage characteristics and layer conductivity were measured using a computer-controlled Keithley 4200 instrument under standard conditions using a solar simulator based on a 150 W xenon lamp equipped with an ASTM-certified filter with a calibrated Si photodiode to produce the AM1.5 solar spectrum.

For the FTIR measurements, we used a Nicolet 560 FTIR spectrometer with a measurement region varying between 400 and 4000 cm$^{-1}$ with suitable scan resolution of 2 cm$^{-1}$. The UV–vis measurements were studied by means of a UV–vis Lambda 40 spectrophotometer with an integrating sphere. The study of the structural properties of ZnO NWs and NRs was performed using scanning electron microscopy (Gemini field emission SEM, Carl Zeiss SUPRATM 25).
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