

# NATURAL DYE PHOTOSENSITIZERS / SOLID STATE ELECTROLYTE JUNCTION FOR DSSC APPLICATION

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#### Abstract

Dye-sensitized solar cells (DSSCs) were constructed by using the Lawsonia inermis leaves, Sumac/Rhus fruits, and Curcuma longa roots as natural sensitizers of anatase-based nanostructure  $TiO_2$  thin film Paint-coated on ITO conducting glass. The orange-red Lawsone, red purple anthocyanin and yellow Curcumin are the main components in the natural dyes obtained from these natural products. A blend of 50 wt% chitosan and 50 wt% polyethylene oxide (PEO) was used as a solid state thin film electrolyte. The polymer blend was complexed with ammonium iodide (NH<sub>4</sub>I) and some iodine crystals were added to the polymer–NH<sub>4</sub>I solution to provide  $I^{\prime}I^{3}$  redox couple. The ionic conductivity of the polymer electrolyte is  $1.18 \times 10^{-5}$  S cm<sup>-1</sup> at room temperature. Structural and optical properties of the semiconductor thin films were characterized by X-ray diffractometer and UV-VIS spectrophotometer. The XRD shows nanocrystalline structures for  $TiO_2$  thin films (D=13nm). The photovoltaic properties of the cell have been studied and the best overall solar energy conversion efficiency of 1.75% was obtained, under AM 1.5 irradiation, with the red purple Sumac/Rhus extract, that showed a reasonable current density (Jsc = 0.93 mA/cm<sup>2</sup>).

The energy and fuel crisis is the prime concern worldwide. Fossils fuels are facing rapid resource depletion, but the

**Keywords:** natural dyes; DSSC; solid state electrolyte; photovoltaic; solar energy

# 1. Introduction

de mand for energy is growing day by day and many countries around the world have no alternative but to increase domestic oil process. So there is an urgent need of sustainable energy resources, such as the solar energy, which is considered as an environmentally friend, novel alternative and promising candidate to address this problem. However, solar energy has a limited application that directly related to its high cost of the per watt electricity generated. In present time, technology of solar cells based on crystalline silicon is facing a problem of silicon-based raw materials. So, low cost alternatives and hence new types of low cost solar cells is an urgent issue. Dyesensitized solar cells (DSSCs), a new type of solar cells, have attracted considerable attention due to their environmental friendliness and low cost of production. A DSSC is composed of a nanocrystalline porous semiconductor electrode-absorbed dye, a counter electrode, and an electrolyte containing iodide and triiodide ions. In DSCs, the dye as a sensitizer plays a key role in absorbing sunlight and transforming solar energy into electric energy. Numerous metal complexes and organic dyes have been synthesized and utilized as sensitizers. By-far, the highest efficiency of DSSCs sensitized by Ru-containing compounds absorbed on nanocrystalline TiO<sub>2</sub> reached 11– 12% [1,2]. However, noble metals limited in amount, and costly in production. On the other hand, organic dyes are not only cheaper but have also been reported to reach efficiency as high as 9.8% [3]. However, organic dyes have often presented problems as well, such as complicated synthetic routes and low yields. Nonetheless, the natural dyes found in flowers, leaves, and fruits can be extracted by simple procedures. Due to their cost efficiency, non-toxicity, and complete biodegradation, natural dyes have been a popular subject of research. Thus-far, several natural dyes have been utilized as sensitizers in DSSCs. Calogero and Marco reported that a conversion efficiency of 0.66% was obtained using red Sicilian orange juice dye as sensitizer [4]. Wongcharee et al. employed rosella as sensitizer in their DSSC, which achieved a conversion efficiency of 0.70% [5]. Roy et al. indicated that when using Rose Bengal dye as sensitizer, the Jsc and Voc of their DSSC reached 3.22mAcm-2 and 0.89 V, respectively, resulting in a 2.09% conversion efficiency [6]. Furthermore, Wang et al. carried out structural modification of coumarin and used the coumarin derivation dye as sensitizer in their DSSC, which provided an efficiency of 7.6% [7-10]. For ideal performance and excellent efficiency, electrolyte should have high ionic conductivity so that it can transfer

oxidized/reduced species to respective electrodes efficiently and should prevents back electrode reactions

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completely. Polyethylene oxide (PEO) has some exceptional properties of good mechanical strength, film forming properties and excellent ability to form complex with the ionic salts. Polymeric electrolyte is an ideal choice used in lithium ion batteries, supercapacitors, photoelectrochromic display devices and solar cells [11-13].

Dye-sensitized solar cell (DSSC) constructed using dye molecules, nanocrystalline metal oxides and organic liquid electrolytes have attractive features of high energy conversion efficiency and low production cost [14, 15]. However, presence of traditional organic liquid electrolytes in such cells has some problems such as a less long-term stability and a need for hermetic sealing. Moreover, the electrolyte loss caused by the leakage and volatility of the electrolyte solution has been pointed out to be one of the major problems, which stays the durability of the dye-sensitized solar cell low. Solid-state dye-sensitized solar cell did not need hermetic sealing, but energy conversion efficiency of them decreased in comparison to those of dye-sensitized solar cell with traditional organic liquid electrolytes. Various approaches to these problems have been tried so far. These approaches include employing a gel-type electrolyte to minimize the loss [16–20].

Since the preparation of synthetic dyes normally requires multistep procedures, organic solvents and, in most cases, time consuming chromatographic purification procedures, there is interest towards the possible use of natural dyes which can be easily extracted from fruits, vegetable and flowers with minimal chemical procedures [21–23]. The pigments are present in the different part of the plant including flowers petals, fruits, leaves, stems and roots. We report here the results of a series of experiments carried out on raw extracts of the following species: three types of natural dyes were extracted from leafs, flowers and roots, these are; Lawsonia inermis (Henna leafs), Sumac /sumach, (flowering plants) and Curcuma longa (turmeric) respectively. To the best of our best knowledge, natural dyes from Henna and Sumac are reported as sensitizers of DSSCs for the first time. Figure1shows a schematic chemical structure of lawsone pigment in henna leafs.

Figure 1. Schematic Chemical Structure of Lawsone pigment in Henna leafs

Henna (Lawsonia inermis, also called henna tree) [24] is a flowering plant used since antiquity to dye skin, hair, fingernails, leather and wool. The name is also used for dye preparations derived from the plant. Henna's coloring properties are due to lawsone, (2-hydroxy-1,4-naphthoquinone) [25], also known as hennotannic acid,  $C_{10}H_6O_3$ , a burgundy organic compound that has an affinity for bonding with protein. Lawsone is primarily concentrated in the leaves. Fresh henna leaves will not stain color until the lawsone molecules are made available (released) from the leaves and they are smashed with a mildly acidic liquid. The lawsone will gradually migrate from the henna paste/solution into the outer layer of the skin and bind to the proteins in it known as keratin, creating a fast stain. Lawsone is a skin protective since it strongly absorbs UV light.

Sumac is any one of approximately 250 species of flowering plants in the genus Rhus in the family Anacardiaceae. Sumacs grow in subtropical and temperate regions throughout the world, especially in Africa and North America [26].

Sumacs are shrubs and small trees that can reach a height of 1–10 meters (3.3–33 ft). The leaves are spirally arranged; they are usually pinnately compound, though some species have trifoliate or simple leaves. The flowers are in dense panicles or spikes 5–30 centimeters (2.0–12 in) long, each flower very small, greenish, creamy white or red, with five petals. The fruits form dense clusters of reddish drupes called sumac bobs. The dried drupes of some species are ground to produce a tangy purple spice [27]. Grounded dried fruits of sumac are usually purple–reddish in color, Tart and sour [28]. The pericarp owes its dark red color to anthocyanin pigments.

**Figure 2.** Anthocyanin chemical structure ( $R_1$  and  $R_3 = -H$ ) ( $R_2$ ,  $R_4$ ,  $R_5$ ,  $R_6$  and  $R_7 = -OH$ ) [29].

Turmeric (Curcuma longa) is a rhizomatous herbaceous perennial plant of the ginger family, Zingiberaceae [30]. It is native to tropical South Asia and needs temperatures between 20 °C and 30 °C (68 °F and 86 °F) and a considerable amount of annual rainfall to thrive. When not used fresh, the rhizomes are boiled for several hours and then dried in hot ovens, after which they are ground into a deep orange-yellow powder commonly used as a spice in curries and other South Asian and Middle Eastern cuisine, for dyeing, and to impart color to mustard condiments. Its active ingredient is curcumin and it has a distinctly earthy, slightly bitter, slightly hot peppery flavor and a mustardy smell. Curcumin can be used to test the alkalinity or acidity of foods. It turns yellow in an acidic food, and it turns red in an alkaline food [31]. Turmeric contains up to 5% essential oils and up to 5% Curcumin, a polyphenol. Curcumin is the active substance of turmeric and Curcumin is known as C.I. 75300, or Natural Yellow 3. The systematic chemical name is (1E, 6E)-1, 7-bis (4-hydroxy-3-methoxyphenyl)-1,6-heptadiene-3,5-dione. It can exist at least in two tautomeric forms, keto and enol. The keto form is preferred in solid phase and the enol form in solution. Curcumin is a pH indicator. In acidic solutions (pH <7.4) it turns yellow, whereas in basic (pH > 8.6) solutions it turns bright red [32].

Figure3. Chemical Structures of Two Tautomeric Forms of Curcumin, keto and enol

As shown in Figure 1, 2, and 3, all dyes contain carboxylic functions which facilitate  $TiO_2$  surface binding [33]. These extracted dyes were characterized by UV–VIS absorption spectra and the semiconductor thin films were characterized by X-ray diffractometer and UV-VIS spectrophotometer. The preparation and characterization of the polymer matrix as polymer electrolyte is investigated in our previous work [13]. The photovoltaic properties of the DSSCs using these extracts as sensitizers and PEO as polymeric solid state electrolyte were investigated. The energy conversion efficiency ( $\eta$ ) of the solar cell device was calculated by the values of open-circuit voltage (Voc) and short-circuit current (Jsc), and fill factor (FF).

# 2. Experiment

We have prepared the solid state dye-sensitized solar cell devices, using natural dyes as photosensitizers, sandwiched with nanocrystalline semiconductor oxide of TiO<sub>2</sub> deposited and C-coated electrodes as working and counter electrodes respectively. The structure of the dye-sensitized solar cell device is shown in Figure 4. However the role of the polymer electrolyte is a channel for the redox couple.

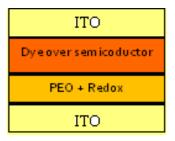


Figure 4. Schematic configuration of Natural dye photosensitizers / Solid state electrolyte junction

#### 2.1. Preparation of TiO<sub>2</sub>Working Electrode (Photoanode)

The photoanode is prepared by adsorbing a dye (s) on a porous  $TiO_2$  layer deposited on ITO conducting glass. By this approach, the dye extends the spectral sensitivity of the photoanode, enabling the collection of lower energy photons. The conductive glass plates (ITO glass, Indium-doped  $SnO_2$ , sheet resistance 15  $\Omega/cm^2$ ) and the titanium oxide ( $TiO_2$ ) nanopowder (13 nm) were purchased respectively from Nikko Materials, Japan and Aldrich. Solvents and chemicals were used as received. The semiconductor paste was prepared by blending 20 g of commercial  $TiO_2$  nanopowder, 3 ml of 0.1 M acetic acid, and 20ml of ethanol. The resulting suspension was stirred for 2 h and subsequently ultra-sonicated for additional 2 h; the resulting mesoscopic oxide film was around 7–20  $\mu$ m thick and opaque. Two edges of the ITO glass plate were covered with a layer of adhesive tape to control the thickness of the film and to mask electric contact strips. Successively the  $TiO_2$  paste was spread uniformly on the substrate by sliding a glass rod along the tape spacer. After heating up the ITO glass spreaded  $TiO_2$  nanoparticle to  $100^{\circ}$ C for about half an hour, the sintering process was completed and the  $TiO_2$  deposited- electrode was cooled down from  $100^{\circ}$ C to  $60^{\circ}$ C at cooling rate of  $3^{\circ}$ C/min to avoid cracking of the glass.

#### 2.2. Graphite Coated Counter Electrode

To prepare the counter (positive) electrodes, uncoated ITO plates were coated with carbon on the conducting side using a graphite rod or soft pencil to apply a light carbon film to the entire conductive side of the plate. Any loose graphite particles should be gently removed. This thin carbon layer serves as a catalyst for the triiodide-to-iodide regeneration reaction. For long-lasting the carbon-coated counter electrode was annealed at 450 °C for a few minutes and washed with ethanol and gently blotted dry before the device is assembled.

#### 2.3. Preparation of Solid State Polymer Electrolyte

The Polymer Electrolyte was prepared and characterized following the procedure reported in the literature of our previous work [13]. In these dye-sensitized solar cells, the polymer electrolyte, polyethylene oxide (PEO) was casted onto  $TiO_2$  electrode impregnated natural dye as photosensitizers.

## 2.4. Preparation of Natural Dye Sensitizers

The fresh Lawsonia inermis leaves and Curcuma longa roots were harvested in Kuala Lumpur Malaysia, while the red Sumac was taken from the Middle East. Lawsonia inermis leaves and Curcuma longa roots extracts were prepared by crushing the fresh leaves and roots respectively. 30 ml of ethanol was added to the pastes and bring to boil for 30 minutes. The as prepared extract solutions were filtered to remove solid fragments and stabilized at pH = 2.5, 3.0, and 3.5 by addition of aqueous (0.1 M) HCl. Extracts further purification was avoided to check whether an efficient sensitization could be achieved with minimal chemical procedures. If properly stored, protected from direct sunlight and refrigerated at about +4 °C, the acidic natural dye solutions (pH = 5.0) are usually stable, with a deactivation half-time of more than 12 months [34]. Grounded dried fruits of sumac (Rhus) are usually purple-reddish in color, Tart and sour, did not require HCl stabilization. The powder was soaked in ethanol and boiled for 30 minutes. TiO<sub>2</sub> coated electrodes were dipped in these solution extracts overnight (24 hours) for pigments stain.

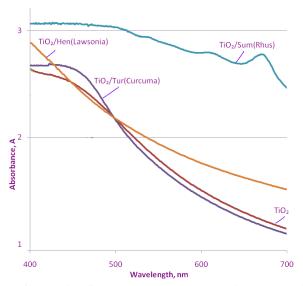
# 2.5. Assembly of DSSC or Grätzel Solar Cell

The dye-sensitized solar cells were assembled by fixing a  $TiO_2$  electrode casted with polymer electrolyte and a graphite counter electrode so that the dyed  $TiO_2$  plates facing down onto the coated graphite anode. They should be placed so that they are slightly offset to allow connections (for the crocodile clips). The two electrodes were pressed against the electrolyte and clamped firmly in a sandwich configuration. By illuminating the cells with a light source/sun light, the voltage across each individual cell can be measured.

#### 3. Results and Discussion

# 3.1. Absorption Spectra of pigment stained TiO<sub>2</sub>

Figure 5 shows the UV–VIS absorption spectra of pure  $TiO_2$  coated on ITO conducting glass and the adsorbed extracts on the  $TiO_2$  electrodes of;  $TiO_2$ /Lawsone extract (orange-red),  $TiO_2$ /Sumac extract (purple) and  $TiO_2$ /Curcuma longa extract (yellow) for two sets of experiments carried out at (a) different layers (thickness) and constant pH=3.0 and (b) different pH and constant thickness (1 layer). These absorptions originate from  $\pi$ – $\pi$ \* transitions have pointed out their essential charge transfer character. Upon adsorption on the  $TiO_2$  electrodes of all extracts the visible absorption band shifts to higher energy, showing a maximum absorption around 400–500 nm. The acidic environment was essential for obtaining sensitized photo-electrodes characterized by high optical densities, capable of wide absorption of visible photons in the 400–650 nm range. The reason is ostensibly related to protonation of carboxylic groups which are otherwise unable, in their anionic form to bind to the  $TiO_2$  surface.



**Figure5.** Absorption spectrum of pure TiO<sub>2</sub> film coated on ITO conducting glass and TiO<sub>2</sub> films adsorbed by three pigments; Lawsone, Sumac and Curcumin

# 3.2. Photovoltaic Properties

Figure 6 shows the Variation of current–voltage curve of lawsone based DSSCs. All experiments were carried out less than 1 sun illumination, ( $100 \text{ mW/cm}^2$ , and air mass 1.5) with solid state electrolyte and  $\text{TiO}_2/\text{Lawsone}$  extract (orange-red) photoanode.

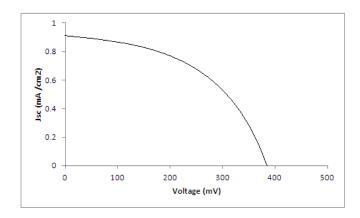


Figure6. Current–voltage curve for a sumac/Rhus extract sensitized solar cell.

Table 1 presents the performance of the DSSCs in terms of short-circuit photocurrent (Isc), open-circuit voltage (Voc) fill factor (FF) and energy conversion efficiency  $(\eta)$ .

<b>Table1.</b> Photovoltaic perform	nances with natural dyes from d	ifferent sources (area*= 0.5 mm <sup>2</sup> )
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Dye Source	Jsc (mA/cm <sup>2</sup> )	Voc (mV)	FF	η (%)
Lawsonia inermis	0.38	336	0.38	0.49
Sumac/Rhus	0.93	394	0.48	1.75
Curcuma longa	0.20	280	0.32	0.18

Sumac/Rhus extracts displayed promising photoelectrochemical performances showing Jsc = 0.93 mA/cm<sup>2</sup>, Voc = 394 m V, FF = 0.48 and  $\eta$  = 1.76% (to our knowledge among the highest efficiency so far reported with raw natural dyes), using a solid state electrolyte composed of PEO-chitosan blend, a non- volatile electrolyte, more suitable for practical applications. The performances of the Lawsonia inermis dyes were poorer to the Sumac/Rhus dyes, exhibiting a short circuit photocurrent close to 0.4 mA cm<sup>-2</sup>. On the other hand Curcuma longa were even poorer source of dyes as compared to the first two photosensitizers. Indeed cells fabricated with such raw extracts only achieved modest power conversion efficiencies, with maximum photocurrents slightly higher than 0.9 mA cm<sup>-2</sup> (see Table 1). In general, natural dyes suffer from low Voc. This can be due both to possible efficient electron/dye cation recombination pathways and to the acidic dye adsorption environment. In fact, it is well known that H+ are potential determining ions for TiO<sub>2</sub> and that proton adsorption causes a positive shift of the Fermi level of the TiO<sub>2</sub>, thus limiting the maximum photovoltage that could be delivered by the cells.

In dye-sensitized  $TiO_2$  solar cell the photoexcited dye transfers an electron to the semiconducting  $TiO_2$  layer via electron injection. The injected electron is then transported through the porous  $TiO_2$  layer and collected by the conductive ITO layer on the glass surface. Within the electrolyte, the mediator ( $\Gamma/I_3$ ) undergoes oxidation at the dye and regeneration at the catalyst-coated counter electrode as long as the cell is illuminated and current flows through the electrical load. The reactions found in the solar cell are:

$$Dye + light \rightarrow dye^* \tag{1}$$

Dye\* + TiO<sub>2</sub> 
$$\rightarrow$$
 e<sup>-</sup> (TiO<sub>2</sub>) + oxidized dye (2)

Oxidized dye + 
$$3/2 \Gamma \rightarrow \text{dye} + 1/2\Gamma_3$$
 (3)

$$1/2\Gamma_3 + e^{-1}$$
 (counter electrode)  $\rightarrow 3/2\Gamma$ 

# 4. Conclusion

In this work we have reported an investigation on three types of pigments as natural photosensitizers, describing and comparing their sensitization and Photoelectrochemical activities with respect to one another. The raw pigments simply extracted in acidic conditions from leaves, roots and fruits achieved solar energy conversion efficiency of 1.75%, which is the highest obtained among all sensitized cells. Natural dye based cells appear to be limited by low Voc and a large decrease in photocurrent, probably due to dye degradation. Finding different additives for improving Voc might result in larger conversion efficiencies. Although the efficiencies obtained with these natural dyes are still below the current requirements for large scale practical application, the results are encouraging and may boost additional studies oriented to the search of new natural sensitizers and to the optimization of solar cell components compatible with such dyes.

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