Structure and Photoelectrical Properties of Natural Photoactive Dyes for Solar Cells

Qian Liu, Nan Gao, Dejiang Liu, Jinglin Liu and Yuanzuo Li

Abstract: A series of natural photoactive dyes, named as D1–D6 were successfully extracted from six kinds of plant leaves for solar cells. The photoelectrical properties of dyes were measured via UV-Vis absorption spectra, cyclic voltammetry as well as photovoltaic measurement. To theoretically reveal the experimental phenomena, the chlorophyll was selected as the reference dye, where the ground and excited state properties of chlorophyll were calculated via density functional theory (DFT) and time-dependent density functional theory (TD-DFT). The experimental results show that the absorption peaks of those dyes are mainly distributed in the visible light regions of 400–420 nm and 650–700 nm, which are consistent with the absorption spectrum of chlorophyll. The photoelectrical conversion efficiencies of the solar cells sensitized by the six kinds of natural dyes are in the order of D1 > D4 > D2 > D5 > D6 > D3. The dye D1 performance exhibits the highest photoelectrical conversion efficiency of 1.08% among the investigated six natural dyes, with an open circuit voltage of 0.58 V, a short-circuit current density of 2.64 mA cm$^{-2}$ and a fill factor of 0.70.

Keywords: natural photo-active dyes; UV-Vis absorption spectra; density functional theory (DFT); Dye-sensitized solar cells (DSSCs)

1. Introduction

With the development and utilization of renewable energy, much works have been paid attention to the devices converting solar energy into electrical energy, among which the dye-sensitized solar cells (DSSCs) with its characteristics of low fabrication costs, environmentally friendly components and relatively high photoelectrical conversion efficiency (PCE) have received wide attention [1–5]. In 1991, Regan and Gratzel [6] created the DSSCs with overall PCE yield of 7.1–7.9% in simulated solar light from low- to medium-purity materials by low-cost process. DSSCs are mainly composed of a light absorbing photoanode, dyes, an electrolyte containing redox couple and a counter electrode [7]. It is remarkable that the dyes play a crucial role in DSSCs because firstly they absorb sunlight [8], followed by injection of the excited electron into semiconductor. In addition to the use of dyes containing metal elements [9–11], the metal-free organic dyes have been widely applied to the field of DSSCs [12–19]. However, the shortcomings of high cost, environmental pollution and also expensive chemical synthesis process are worth the attention of the researchers [20]. Fortunately, researchers have found that the application of natural dyes extracted from the plant flowers petals, leaves, roots in the DSSCs, also producing a considerable PCE [20–33], which has opened up a friend way for the conversion of solar energy into electricity. For example, Hosseinnezhad et al., used some natural dyes as photosensitizers (extracted from Celosia Cristata, Saffron, Cynoglossum, and eggplant peel) [21], in which PCE is found to be 1.38% (Celosia Cristata). Five different natural dyes with various anchoring
groups have been studied from various plants and used them as photo-sensitizers in DSSC [22], showing possess better absorption at 300–550 nm. Hamadanian et al. [23] used a series of natural dye extracted from plants for the DSSCs. Because Delphinidin was the main pigment of C. ajacis, interaction between the hydroxyl groups of the Delphinidin and the TiO$_2$ surface was very efficient, and this sensitizer owned the best photovoltaic performance among all the natural dyes. It was interesting that Maiaugree and co-workers [24] used carbonized mangosteen peel and mangosteen peel dye as a natural counter electrode and a natural photo-sensitizer for DSSC application, respectively; they found a distinctive mesoporous honeycomb-like carbon structure with a rough nanoscale surface in carbonized mangosteen peels, resulting in the highest PCE (2.63%) when using carbonized mangosteen peel and an organic disulfide/thiolate (T$_2$/T$^-$) electrolyte. Since the first report on Chlorophyll (Chl) and its derivatives as sensitisers in DSSCs [32], the greater sensitizer ability of Chl derivatives with respect to the pure untreated chlorophyll a was evidenced, which result can be explained considering the fact that the low attitude of this natural pigment should be efficiently absorbed on TiO$_2$ layer when dissolved in some solvent (ethanol, acetone, pyridine, etc.). Generally, the dye extracted from natural species are not pure chlorophyll but natural compounds derived by chlorophyll a and b. Recent study found the copper-Chle6 achieving the following parameters: FF of 0.65, $V_{OC}$ of 438 mV and PCE of 1.81% [33]. There are also several ways to improvement of solar cell performances by photoanodes modification, such as the photoanodes modification based on H-TiO$_2$ NPs scattering layer [34] and indium doped ZnO photoanode [35].

Recently, the theoretical method based on quantum chemistry calculation has provided the valuable reference for predicting the relationship between the structure and photoelectrical properties of molecules [36–40]. Qin et al. [41] reported a theoretical study of the betalain pigments using density functional theory (DFT) and time-dependent density functional theory (TD-DFT), and the results indicated that the lowest unoccupied molecular orbital (LUMO) had a close spatial proximity to bound TiO$_2$, assuming that betalain binding occured through the two carboxylic groups on the dihydropyridine ring. Khadtare et al. [42] analyzed the spectral features of lawsone (2-hydroxy-1,4-naphthoquinone), an active component of the natural dye henna in ethanol by using experimental and computational methods; the study showed that the 1,4-naphthoquinone classes of compounds can provide the basis for the design of novel compounds for DSSCs to enhance its efficiency. Charge transfer enhancement in DSSCs was reported by changing the electron donor of the famous porphyrin-based sensitizer YD2-o-C8 [43].

In this work, six natural dyes, named as D1, D2, D3, D4, D5 and D6, were successfully extracted from the leaves of Euodia meliaeifolia (Hance) Benth, Matteuccia struthiopteris (L.) Todaro, Corylus heterophylla Fisch, Filipendula intermedia, Pteridium aquilinum var. iatiusculum and Populus L (see Figure 1), respectively. The optical and electrical properties of the dyes applied in DSSCs were measured via UV-Vis absorption spectra and cyclic voltammetry. To compare with the experimental results, the chlorophyll was selected as the reference molecule, and the ground and excited state properties of chlorophyll were calculated via DFT and TD-DFT methods, respectively. The obtained results indicate the DSSC sensitized by the dye D1 exhibits the highest PCE of 1.08% among the investigated six natural dyes, with an open circuit voltage ($V_{OC}$) of 0.58 V, a short-circuit current density ($J_{SC}$) of 2.64 mA cm$^{-2}$ and a fill factor (FF) of 0.70.
The fresh leaves of the six plants were selected and placed into a vacuum drying oven maintaining 70 °C to remove moisture. The dried leaves were pulverized into powder in a mortar and weighed 1 g powder, and then mixed 20 mL ethanol with the power. After keeping the admixture away from the light in a brown jar for a week, the efficient natural dyes were obtained through a simple filtering process. The structure of prepared DSSC was mainly made of TiO<sub>2</sub> photo-anode, dye and electrolyte. The specific production process is as our previous works [16]: (a) the TiO<sub>2</sub> electrode was prepared through adding 10 mL isopropyl tianate into water, and keeping the hydrolysis for 3 h, and then adding nitric acid and acetate into the solution, and placing in an environment of 80 °C; the mixed solution was stirred until it became transparent blue; later, the hydrothermal reaction was executed at 200 °C for 12 h. After cooling, spin steaming, centrifuging, terpineol ethyl and cellulose were added into the ball grinder; the paste were prepared completely by ball mill, rotary steam and three roll. (b) the screen printing technology was adopted to print the TiO<sub>2</sub> paste to the clean surface of a conductive glass, and the active area of TiO<sub>2</sub> photo-anode was 0.16 cm<sup>2</sup>; after ethanol bathing and drying, the TiO<sub>2</sub> photo-anode was sintered, and then treated in TiCl<sub>4</sub> solvent. After this, the photo-anode was sintered again. In the later process, the photo-anode was immediately removed after the natural cooling to 80 °C, and the photo-anode was soaked in the natural dye without light for 24 h. (c) the photo-anode and the platinum plating counter electrode were fitted together into the cell, and in the middle of the two electrodes, the electrolyte solution (including 0.5 M LiI, 0.05 M I<sub>2</sub> TBP, GUSCN) was added. The SEM image of thin film of TiO<sub>2</sub> nanoparticles was shown in Figure 2, where the area of TiO<sub>2</sub> film was 16 mm<sup>2</sup> and thickness of the TiO<sub>2</sub> anode layer was about 16–18 μm; here, the characteristic of void and honeycomb of TiO<sub>2</sub> anode layer is benefit for enhancing the adsorption of dyes.

2. Methods

The fresh leaves of the six plants were selected and placed into a vacuum drying oven maintaining 70 °C to remove moisture. The dried leaves were pulverized into powder in a mortar and weighed 1 g powder, and then mixed 20 mL ethanol with the power. After keeping the admixture away from the light in a brown jar for a week, the efficient natural dyes were obtained through a simple filtering process. The structure of prepared DSSC was mainly made of TiO<sub>2</sub> photo-anode, dye and electrolyte. The specific production process is as our previous works [16]: (a) the TiO<sub>2</sub> electrode was prepared through adding 10 mL isopropyl tianate into water, and keeping the hydrolysis for 3 h, and then adding nitric acid and acetate into the solution, and placing in an environment of 80 °C; the mixed solution was stirred until it became transparent blue; later, the hydrothermal reaction was executed at 200 °C for 12 h. After cooling, spin steaming, centrifuging, terpineol ethyl and cellulose were added into the ball grinder; the paste were prepared completely by ball mill, rotary steam and three roll. (b) the screen printing technology was adopted to print the TiO<sub>2</sub> paste to the clean surface of a conductive glass, and the active area of TiO<sub>2</sub> photo-anode was 0.16 cm<sup>2</sup>; after ethanol bathing and drying, the TiO<sub>2</sub> photo-anode was sintered, and then treated in TiCl<sub>4</sub> solvent. After this, the photo-anode was sintered again. In the later process, the photo-anode was immediately removed after the natural cooling to 80 °C, and the photo-anode was soaked in the natural dye without light for 24 h. (c) the photo-anode and the platinum plating counter electrode were fitted together into the cell, and in the middle of the two electrodes, the electrolyte solution (including 0.5 M LiI, 0.05 M I<sub>2</sub> TBP, GUSCN) was added. The SEM image of thin film of TiO<sub>2</sub> nanoparticles was shown in Figure 2, where the area of TiO<sub>2</sub> film was 16 mm<sup>2</sup> and thickness of the TiO<sub>2</sub> anode layer was about 16–18 μm; here, the characteristic of void and honeycomb of TiO<sub>2</sub> anode layer is benefit for enhancing the adsorption of dyes.

Figure 1. Basic working principle of DSSC, which mainly contains four parts: photoanode, natural dyes, electrolyte containing redox couple and counter electrode, where photosensitive layer using the dyes (D1–D6) were extracted from the leaves.

Figure 2. SEM micrographs of the TiO<sub>2</sub> nanoparticles on the glass.
UV-Vis spectra of dye solutions and dye adsorbed on TiO$_2$ photo-anodes were measured with TU-1900 spectrometer (Beijing Purkinje General Instrument Co., Ltd., Beijing, China). Cyclic voltammetry properties were performed by using CH Instruments CHI832C Electrochemical Workstation (Shanghai Chenhua Instrument Co., Ltd., Shanghai, China). The redox potentials of the dyes were measured in ethanol solvent. The scan range was between $-1000$ mV and $+1000$ mV, and the initial scan potential was $-1000$ mV, at a scan rate of 50 mV/s. Here, a glassy carbon working electrode, platinum counter electrode and Ag/AgCl reference electrode were used to form electrode system, where 0.1 M KNO$_3$ was used as the supporting electrolyte [44]. Solar energy conversion efficiency measurements were completed with a solar simulation instrument (PEC-L15, Peccell Technologies, Inc., Yokohama, Japan), and light intensity was adjusted via a reference standard Si-solar solar cell at 1 sun light intensity of 100 mW cm$^{-2}$.

Compared with experiment, the chlorophyll was selected as the reference dye in theory, and the ground state structure of chlorophyll a was optimized by using DFT [45], with B3LYP [46–48] functional at 6-31G(d) basis set. On the base of optimized ground state structure, the excited state properties of chlorophyll were calculated via TD-DFT [49] with different functionals, containing Cam-B3LYP, MPW1PW91 and PBEPBE [50–53] at 6-31G(d) basis set. By comparing the obtained theoretical and experimental results, the calculated absorption spectrum of chlorophyll by using the PBEPBE functional was well consistent with the experimental results. Therefore, we selected the theoretical results calculated by using PBEPBE functional to reveal the experimental results. The ethanol was selected as the solvent, and all calculations in solvent adopted the Conductor-like PCM (C-PCM) model [54]. DFT calculations was used to study photosensitizing features [55] and photophysical properties of expanded bacteriochlorins [56], where micromechanism of photosensitizing nature was revealed with DFT calculations. In addition, the three-dimensional (3D) real-space analysis method was used to describe the charge transfer process in the excited states [57–60]. All calculations were performed using Gaussian 09 package [61].

3. Results and Discussion

3.1. UV-Vis Absorption Spectra in Experiment and Theory

The UV-Vis absorption spectra of the dyes in ethanol and dye adsorbed on TiO$_2$ photo-anodes were measured, and the corresponding absorption spectra are presented in Figure 3a,b. The detected absorption peaks corresponding to the isolated dyes and dyes adsorbed on TiO$_2$ are listed in Table 1 in detail. As shown in Figure 3a, the absorption spectra of those six kinds of dyes have similar appearance, in which the absorption peaks are mainly distributed in the visible light regions of 400–420 nm and 650–700 nm. It is worth noting that all the absorption spectra of the investigated dyes have the characteristics of double peaks, which are advantageous to the absorption of dyes on the sun light. As listed in Table 1, the two absorption peaks of the six dyes are focused on about 412 nm and 664 nm. The absorption spectra of the dyes investigated in the present work are consistent with the absorption spectrum of chlorophyll [62,63]. In addition, the absorption spectra of dyes adsorbed on TiO$_2$ photo-anode are shown in Figure 3b, and the corresponding absorption peaks are listed in Table 1. As listed in Table 1, the absorption peaks in the region of 650–700 nm exhibit different extent of red-shifts compared with that of the isolated dyes, in which the red-shift values are 7.5 nm, 10 nm, 6.5 nm, 29.5 nm, 7.5 nm and 6.5 nm for D1, D2, D3, D4, D5 and D6, respectively.

<table>
<thead>
<tr>
<th>Dye</th>
<th>Absorption Peaks (nm)</th>
<th>Dye/TiO$_2$</th>
<th>Absorption Peaks (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>D1</td>
<td>416.0, 664.0</td>
<td>D1/TiO$_2$</td>
<td>416.5, 671.5</td>
</tr>
<tr>
<td>D2</td>
<td>419.5, 658.5</td>
<td>D2/TiO$_2$</td>
<td>421.5, 668.5</td>
</tr>
<tr>
<td>D3</td>
<td>412.5, 663.5</td>
<td>D3/TiO$_2$</td>
<td>416.5, 670.0</td>
</tr>
<tr>
<td>D4</td>
<td>411.5, 665.5</td>
<td>D4/TiO$_2$</td>
<td>695.0</td>
</tr>
<tr>
<td>D5</td>
<td>413.5, 664.0</td>
<td>D5/TiO$_2$</td>
<td>413.5, 671.5</td>
</tr>
<tr>
<td>D6</td>
<td>412.0, 665.0</td>
<td>D6/TiO$_2$</td>
<td>416.5, 671.5</td>
</tr>
</tbody>
</table>
To compare with the experimental results, the excited state properties of chlorophyll was investigated via TD-DFT method with different functionals at 6-31G(d) basis set based on the optimized ground state structure of chlorophyll, due to the fact that chlorophyll accounts for the largest proportion in the green plant leaves. The simulated absorption spectra of chlorophyll by using different functionals are presented in Figure 4, and the corresponding absorption peaks are listed in Table 2. Combined with the results of Figure 4 and Table 2, it can be seen that the simulated absorption spectrum of chlorophyll by using PBEPBE functional is more consistent with the experimental results; so we analyze the excited state properties of chlorophyll based on the results calculated by the PBEPBE functional. The calculated excitation energies, absorption peaks, electronic transition configurations and corresponding oscillator strengths of the first ten excited states are listed in Table 3. As listed in Table 3, the excited state S8 corresponds to the first maximum absorption peak 439.66 nm with the oscillator strength of 0.3846, which is mainly formed by the electrons transition from HOMO$^{-1}$ to LUMO$+1$. In addition, the excited state S1 corresponds to the second maximum absorption peak at 621.44 nm with the oscillator strength of 0.3042, which is mainly originated from the electrons transition from HOMO to LUMO. Moreover, the excited state S7 corresponds to the third maximum absorption peak at 456.17 nm (the oscillator strength of 0.2823), which mainly comes from the electrons transition from HOMO to LUMO$+1$.

Table 2. Calculated absorption peaks and corresponding oscillator strengths (in bracket) of chlorophyll via TD-DFT method with different functionals at 6-31G(d).

<table>
<thead>
<tr>
<th>Peak</th>
<th>Cam-B3LYP</th>
<th>MPW1PW91</th>
<th>PBEPBE</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\lambda_{\text{max}1}$</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a 376.57 (0.9972)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>b 602.89 (0.3746)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>$\lambda_{\text{max}2}$</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a 375.28 (0.8763)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>b 586.59 (0.3816)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>439.66 (0.3846)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>621.44 (0.3042)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*a corresponds to the first absorption peak; b corresponds to the second absorption peak.
Table S1. Calculated absorption peaks is found to be 705.79 nm corresponding to the sixth excited state, which make red-shifted in comparison with the isolated dye in solvent. The closed strength is 0.3042 for external circuit.

Figure 4. Simulated absorption spectra of chlorophyll in ethanol via different functionals with 6-31G(d) basis set.

Table 3. Calculated excitation energies ($E$, eV), absorption peaks ($\lambda_{abs}$, nm), electronic transition configurations and corresponding oscillator strengths ($f$) of chlorophyll a in ethanol by using TD-DFT//PBEPBE/6-31G(d) method.

<table>
<thead>
<tr>
<th>State</th>
<th>$E$ (eV)</th>
<th>$\lambda_{abs}$ (nm)</th>
<th>Contribution MO</th>
<th>Strength $f$</th>
</tr>
</thead>
<tbody>
<tr>
<td>S1</td>
<td>1.9951</td>
<td>621.44</td>
<td>(0.65673)H–L</td>
<td>0.3042</td>
</tr>
<tr>
<td>S2</td>
<td>2.0275</td>
<td>611.51</td>
<td>(0.63801)H–1–L</td>
<td>0.0561</td>
</tr>
<tr>
<td>S3</td>
<td>2.3641</td>
<td>534.44</td>
<td>(0.68996)H–2–L</td>
<td>0.0027</td>
</tr>
<tr>
<td>S4</td>
<td>2.4662</td>
<td>502.73</td>
<td>(0.65897)H–4–L</td>
<td>0.0010</td>
</tr>
<tr>
<td>S5</td>
<td>2.4969</td>
<td>496.56</td>
<td>(0.65411)H–3–L</td>
<td>0.0086</td>
</tr>
<tr>
<td>S6</td>
<td>2.6161</td>
<td>473.93</td>
<td>(0.70631)H–5–L</td>
<td>0.0008</td>
</tr>
<tr>
<td>S7</td>
<td>2.7179</td>
<td>456.17</td>
<td>(0.46269)H–L+1</td>
<td>0.2823</td>
</tr>
<tr>
<td>S8</td>
<td>2.8200</td>
<td>439.66</td>
<td>(0.40417)H–1–L+1</td>
<td>0.3846</td>
</tr>
<tr>
<td>S9</td>
<td>2.9989</td>
<td>413.43</td>
<td>(0.61479)H–7–L</td>
<td>0.0555</td>
</tr>
<tr>
<td>S10</td>
<td>3.0548</td>
<td>405.87</td>
<td>(0.56616)H–6–L</td>
<td>0.0847</td>
</tr>
</tbody>
</table>

For Dye/TiO$_2$ system, calculated absorption information has been listed in supporting materials (Table S1). Calculated absorption peaks is found to be 705.79 nm corresponding to the sixth excited state, which make red-shifted in comparison with the isolated dye in solvent. The closed strength is located in 657.66 nm, which is in ninth excited state. Charge different density shows the distribution of charge during light absorption, which is shown in Figure 5. As shown, for the sixth excited state red electron is move into the semiconductor and hole is resided in the porphyrin ring. Hence this state is charge transfer (CT) state, and similar CT process can be found in ninth excited state, and more migration of electrons into semiconductor is benefit for electron transport for external circuit.

Figure 5. Charge difference density (CDD) for chlorophyll/TiO$_2$. (a) S6; (b) S9.
3.2. Electrochemical Properties

The thermodynamic feasibility of electron injection from the excited dye molecule to the TiO$_2$ conduction band and dye regeneration utilizing electrolyte can be investigated through the research on the electrochemical properties of the dyes [64,65]. Herein, the electrochemical properties of the six kinds of natural dyes were studied via cyclic voltammetry measurement in the ethanol solvent, adopting 0.1 M KNO$_3$ as the supporting electrolyte. The measured cyclic voltammetry curves of the six dyes are exhibited in Figure 6, and the onset oxidation potentials of the dyes obtained from the curves are listed in Table 4. As listed in Table 4, the onset oxidation potentials of the six dyes are 0.34 V, 0.19 V, −0.07 V, 0.07 V, 0.11 V and 0.10 V for D1, D2, D3, D4, D5 and D6, respectively. From the previous report [66], it can be known that the HOMO energy can be obtained by the onset oxidation potential of dye, when an Ag/AgCl electrode is used as the reference electrode; the HOMO energy can be calculated via the following equation:

\[
\text{HOMO} = -e(E_{OX} + 4.40) \text{(eV)}
\]

where \(E_{OX}\) stands for the onset oxidation potential of dye. The calculated HOMO energies of the six dyes are listed in Table 4, which shows that the HOMO energies of the six dyes are −4.74 eV, −4.59 eV, −4.33 eV, −4.47 eV, −4.51 eV and −4.50 eV for D1, D2, D3, D4, D5 and D6, respectively. In terms of the work of Tian et al. [67], the dye with strong donating ability will exhibit the more negative HOMO level. Therefore, it can be concluded from the obtained HOMO energies of these dyes that the electron donating ability of these dyes in the order of D1 > D2 > D5 > D6 > D4 > D3. Their LUMOs are higher than that are higher than semiconductor band (−4.0 eV), meaning possible occurrence of electron injection for studied systems. Furthermore, the lowest HOMO can enhance the ability of dye regeneration because the larger driving force of dye regeneration between dye and electrolytic, which is contributed to the \(J_{SC}\).

Figure 6. Cyclic voltammograms of the six natural dyes. (a) dyes D1–D3; (b) dyes D4–D6.
3.3. Photoelectric Properties of DSSCs

The photoelectrical conversion efficiency ($\eta$) of the DSSC is defined via the following equation [69]:

$$\eta = \frac{J_{SC} \times V_{OC} \times FF}{P_{in}}$$

(1)

where $J_{SC}$ is the short-circuit current density, $V_{OC}$ represents the open circuit voltage, $FF$ is the fill factor, and $P_{in}$ is the intensity of the incident light, in which the $J_{SC}$, $V_{OC}$ and $FF$ are only obtained through the experiment.

The fill factor ($FF$) is defined as the ratio of the maximum power obtained from the DSSC and the theoretical maximum power, which can be understood by the following equation:

$$FF = \frac{I_m \times V_m}{J_{SC} \times V_{OC}}$$

(2)

where $I_m$ and $V_m$ represent the current and voltage related to the maximum power, respectively.

The measured current-voltage curves of the fabricated DSSCs sensitized by the six kinds of natural dyes are shown in Figure 7, and the photovoltaic parameters of the DSSCs, containing $V_{OC}$, $J_{SC}$, $FF$ and $\eta$ are listed in Table 5. It can be found from Table 5 that the PCE of the DSSCs sensitized by the six kinds of natural dyes are in the order of $D1 > D4 > D2 > D5 > D6 > D3$, in which the DSSC sensitized by the dye $D1$ exhibit the highest PCE of 1.08%, with $V_{OC}$ of 0.58 V, $J_{SC}$ of 2.64 mA cm$^{-2}$ and $FF$ of 0.70. It is worth noting that the DSSC sensitized by the dye $D1$ possesses the highest $J_{SC}$ among the investigated DSSCs, which can be promoted by the improved ability of dye regeneration can promote.

![Figure 7. Current-Voltage curves of DSSCs sensitized with the six natural dyes.](image-url)

Table 4. Electrochemical properties of the six natural dyes.

<table>
<thead>
<tr>
<th>Dye</th>
<th>$E_{OX}$ (V) $^a$</th>
<th>HOMO (eV)</th>
<th>LUMO $^b$ (eV)</th>
<th>Dye</th>
<th>$E_{OX}$ (V) $^a$</th>
<th>HOMO (eV)</th>
<th>LUMO (eV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>D1</td>
<td>0.34</td>
<td>−4.74</td>
<td>−2.873</td>
<td>D4</td>
<td>0.07</td>
<td>−4.47</td>
<td>−2.607</td>
</tr>
<tr>
<td>D2</td>
<td>0.19</td>
<td>−4.59</td>
<td>−2.707</td>
<td>D5</td>
<td>0.11</td>
<td>−4.51</td>
<td>−2.643</td>
</tr>
<tr>
<td>D3</td>
<td>−0.07</td>
<td>−4.33</td>
<td>−2.461</td>
<td>D6</td>
<td>0.10</td>
<td>−4.50</td>
<td>−2.636</td>
</tr>
</tbody>
</table>

$^a$ The onset oxidation potential of the dye; $^b$ estimation from HOMO and excitation energy [68].

Table 5. Measured photovoltaic parameters of the fabricated DSSCs.

<table>
<thead>
<tr>
<th>DSSCs</th>
<th>$V_{OC}$ (V)</th>
<th>$J_{SC}$ mA cm$^{-2}$</th>
<th>$FF$</th>
<th>$\eta$ (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>D1</td>
<td>0.58</td>
<td>2.64</td>
<td>0.70</td>
<td>1.08</td>
</tr>
<tr>
<td>D2</td>
<td>0.60</td>
<td>0.75</td>
<td>0.72</td>
<td>0.32</td>
</tr>
<tr>
<td>D3</td>
<td>0.56</td>
<td>0.68</td>
<td>0.69</td>
<td>0.26</td>
</tr>
<tr>
<td>D4</td>
<td>0.54</td>
<td>0.87</td>
<td>0.74</td>
<td>0.34</td>
</tr>
<tr>
<td>D5</td>
<td>0.56</td>
<td>0.74</td>
<td>0.73</td>
<td>0.30</td>
</tr>
<tr>
<td>D6</td>
<td>0.57</td>
<td>1.25</td>
<td>0.37</td>
<td>0.27</td>
</tr>
</tbody>
</table>
4. Conclusions

In this work, a series of natural dyes (D1, D2, D3, D4, D5 and D6) were successfully extracted from six kinds of plant leaves, and their optical and electrical properties were measured via UV-Vis absorption spectra and cyclic voltammetry. To reveal the experimental phenomena, the chlorophyll was selected as the reference molecule, and the DFT and TDDFT methods were adopted to calculate the ground and excited state properties of chlorophyll. The experimental results indicate that the absorption spectra of the six dyes exhibit two absorption bands in the visible light region of 400.0–420.0 nm and 650.0–700.0 nm, and the DSSC sensitized by the dye D1 exhibits the highest PCE of 1.08% among the investigated six natural dyes, with $V_{OC}$ of 0.58 V, $J_{SC}$ of 2.64 mA cm$^{-2}$ and $FF$ of 0.70. The high HOMO energy levels of D1 can be contributed to the highest $J_{SC}$ and further affect the photoelectric properties. In addition, the theoretical results imply that the excited states corresponding to the absorption peaks about 412 nm and 664 nm are mainly derived from electron transition of HOMO $-1$ to LUMO $+1$ and HOMO to LUMO, respectively. There exits charge transfer (CT) state, where electron is transferred into semiconductor during photoexcitation.


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