High Photo-Current in Solution Processed Organic Solar Cells Based on Porphyrin Core A- π -D- π -A as Electron Donor Material

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ABSTRACT

Two new conjugated acceptor-donor-acceptor (A- π -D- π -A) molecules with a porphyrin core linked by ethynylene bridges to two thiophene (**1a**) or thienylenevinylene thiophene (**1b**) units and both capped by N-ethylrhodanine have been synthesized. These compounds were used as the main electron donor moieties for bulk heterojunction small molecule organic solar cells (BHJ-SMOSC). The optimized devices, with PC71BM as the main electron acceptor molecule, show remarkable short circuit currents, up to 13.2 mA/cm², an open circuit voltage of around 0.85 V, and power conversion efficiencies up to 4.3% under 100 W/cm². The External Quantum Efficiency (EQE), Atomic Force Microscopy (AFM), hole mobility, Photo-Induced Charge Extraction (PICE) and Photo-Induced Transient Photo-Voltage (PIT-PV) were analyzed in devices based on **1a** and **1b** in order to account for differences in the final performance of the two molecules. The PIT-PV decays showed slower recombination kinetics for devices fabricated with **1b**. Moreover, the EQE was greater for **1b** and this is ascribed to the better nanomorphology, which allows better charge collection before carrier recombination takes place.

1. Introduction

Solution processed organic solar cells (OSCs) have improved significantly in recent decades,[1-4] due to the judicious design of push-pull low band-gap copolymers, optimization of the nanoscale morphologies of the photoactive layers and enhanced carrier mobility, all of which have led to power conversion efficiencies (PCEs) of 10% for single-junction polymer solar cells (PSCs)[5, 6] and up to 11% for tandem PSCs.[7] Moreover, small molecule organic solar cells (SMOSCs) have emerged as complementary systems to the polymeric materials and they have undergone rapid development in recent years as SMOSCs offer potential advantages such as the possibility of obtaining better-defined molecular structures, easier purification and better batch-to-batch reproducibility.[8, 9] Indeed, power conversion efficiencies for bulk heterojunction small molecule organic solar cells (BHJ-SMOSCs) have exceeded 10% and these are expected to be higher for tandem devices.[10-13]

Numerous different chemical architectures, especially as donor moieties, have been explored for SMOSCs [11, 14-19] and the most promising are the donor-acceptor (D-A) ‘push-pull’ molecules. Within this group, the use of porphyrin moieties (POR) as the main donor has recently been reported but this remains relatively unexplored.

Inspired by natural photosynthetic structures, such as chlorophylls, porphyrins and their derivatives, these excellent building blocks offer efficient light harvesting architectures, high stability, and their large and rigid planar structure allows fine tuning of their energetic and morphological properties.[20] For example, POR-based photosensitizers are among the best sensitizers in dye sensitized solar cells (DSSCs). In recent years, remarkable progress has been made on using POR as the main donor moieties[21-26] and efficiencies as high as 8% have been

reached in SMOSC [27] devices while in DSSC systems efficiencies close to 12% have been described.[28]

Recently, we described two similar conjugated acceptor-donor-acceptor (A- π -D- π -A) compounds that also contained a Zn-porphyrin but were capped by dicyanovinylene groups as acceptor units – these materials gave a photoconversion efficiency (PCE) of 3.21%.[29] With the aim of significantly improving the design and thus the performance of the final device, we report here: (a) the synthesis and characterization of two new conjugated POR-based acceptor-donor-acceptor (A- π -D- π -A) ‘push-pull’ molecules **1a** and **1b** (Fig. 1); (b) the device fabrication and performance of these materials in solution-processed SMOSCs using PC71BM as the main electron acceptor moiety and (c) analysis of the carrier recombination kinetics. Compounds **1a** and **1b** were used as electron donor molecules in the blend, with the Zn-porphyrin core acting as a donor linked by ethynylene groups to one or two units of thienylenevinylene and capped by *N*-ethylrhodanine as acceptor units. The strategy for this design is based on the use of the ethynylene units to make the system planar, the thienylenevinylene to extend the π -conjugation and the incorporation of hexyl chains on the thiophene units to enhance solubility. Finally, the use of rhodanine as the acceptor provides a stronger charge transfer character to the molecule and increases the light harvesting efficiency.

Figure 1. Molecules **1a** and **1b**.

Synthesis and Characterization

Compounds **1a-b** were obtained in 82% and 45% yield, respectively, by Knoevenagel condensation of **2a-b**[28] with N-ethylrhodanine and piperidine as base (Scheme 1). Both new compounds were satisfactorily characterized by ^1H and ^{13}C NMR, FT-IR and MALDI-MS (see Supporting Information for synthetic details and full analytical and spectroscopic data). In the ^1H NMR spectra of **1a** and **1b** a new signal was observed at 8.00 ppm and 7.95 ppm, respectively, and this confirms the success of the condensation reaction. The mass spectrum of **1a** showed the molecular ion peak at m/z 1498.75 amu and **1b** gave a molecular ion peak at m/z 2051.69 amu.

Compounds **1a-b** showed excellent thermal stability up to 300 °C, with Td values of 377 °C and 355 °C for **1a** and **1b**, respectively (Figure S9 and S10); this thermal stability is suitable for photovoltaic applications.

Optical Properties

The optical properties of **1a** and **1b** in CH_2Cl_2 (DCM) solutions and in thin films are shown in Fig. 2 and the characteristics of semiconductors **1a** and **1b** are collected in Table 1.

The absorption spectra in DCM solution of both compounds exhibit a panchromatic absorption in the visible region. Compound **1a** shows the characteristic absorption pattern of a Zn-chelated porphyrin with an intense Soret band (B band) at 507 nm ($\log \varepsilon = 5.27$) and a broad intense intramolecular charge transfer (ICT) band with a maximum at 709 nm ($\log \varepsilon = 5.09$). Extension of the conjugation on the bridge by the introduction of a new thienylenevinylene unit had a significant impact on the absorption spectrum of **1b**. This change led to a broader absorption, which is bathochromically shifted, for the Soret band to 547 nm ($\log \varepsilon = 5.02$) and in the ICT band at 716 nm ($\log \varepsilon = 5.02$). Comparison of the corresponding absorptions in films shows that

the absorption bands became broader and red-shifted as a consequence of the closer intermolecular interactions in solution.

The fluorescence spectra of **1a-b**, measured in DCM (CH_2Cl_2) and with excitation at 481 and 482 nm, respectively, show emission bands at 727 and 746 nm, respectively (Figure S13 and S14).

Figure 2. (a) UV-Visible spectra of **1a** (black) and **1b** (red) in DCM solution. (b) Normalized UV-Vis spectra of **1a** (black) and **1b** (red) and in thin films (red line).

From absorbance and emission data, the E_{0-0} values were calculated. The HOMO-LUMO gaps are as narrow as 1.74 and 1.69 eV for **1a** and **1b**, respectively.

Table 1 UV-Vis, ^a Fluorescence Emission^a and OSWV^b data for compounds **1a-b**

	λ_{max} sol (nm)	log (ϵ)	λ_{max} film (nm)	λ_{em} (nm)	E_{0-0} ^c (eV)	E'_{ox} ^{b,d} (V)	E_{HOMO} ^e (eV)	E_{LUMO} (eV)
1a	481	5.16	435	727	1.74	0.26	-5.36	-3.62
	507	5.27	522					
	709	5.09	768					
1b	482	4.95	510	746	1.69	0.14	-5.24	-3.55
	547	5.02	540					
	716	5.02	762					

^a 10⁻⁵ M in dichloromethane (DCM); ^b 10⁻³ M in *o*-dichlorobenzene (ODCB)-acetonitrile (4:1) versus Fc/Fc⁺ (E_{ox} = 0.04 V) glassy carbon, Pt counter electrode, 20 °C, 0.1 M Bu₄NClO₄, scan rate = 100 mV s⁻¹; ^c estimated from the intersection between the normalized absorption and normalized emission spectra at λ_{max} ; ^d reversible processes; ^e calculated with respect to ferrocene, E_{HOMO} : -5.1 eV.

Electrochemical Properties

The electrochemical properties of **1a** and **1b** were investigated by Cyclic Voltammetry (CV) and Osteryoung Square Wave Voltammetry (OSWV) in *o*-dichlorobenzene (ODCB)-acetonitrile (4:1) (Table 1, Figures S15 and S16). On the cathodic side, both compounds showed a first reversible one-electron oxidation wave at 0.26 V for **1a** and 0.14 V for **1b** (vs Fc/Fc⁺ in all cases). This first oxidation potential is ascribed to the oxidation of the porphyrin core; it can be remarked that the extended conjugation in **1b** gives rise to a decrease in the E_{ox} value by 12 mV with respect to **1a**. A second reversible oxidation wave was observed at 0.65 V for **1a** and at 0.42 V for **1b**, and this is attributed to the thienylenevinylene units. Compound **1b** showed two more non-reversible oxidation waves at 0.63 and 0.88 V, which are attributed to oxidation of the thienylenevinylene moieties. The estimated E_{HOMO} values were determined to be -5.36 eV for **1a** and -5.24 eV for **1b**.

Low-lying HOMO levels should result in high open-circuit voltages (V_{oc}).^[30-32] The E_{LUMO} values of **1a-b** are higher than the E_{LUMO} of PC71BM (-4.0 eV), i.e., -3.62 and -3.55 eV for **1a**

and **1b**, respectively. Hence, the LUMO energy levels of these small-molecules (Fig. 3) clearly match the LUMO of PC71BM, thus enhancing charge transfer.

Figure 3. Schematic representation of the HOMO and LUMO energy levels for **1a**, **1b** and PC71BM.

Theoretical Calculations

In order to gain an insight into the structure and the electronic properties of the materials, theoretical studies were carried out by performing density functional theory (DFT) calculations at the B3LYP/6-31G* level *in vacuo* using Gaussian 03W.

The optimum geometries obtained for dyes **1a** and **1b** (Figure S17) show that the π -conjugated system, porphyrin macrocycle and the thienylenevinylene monomers are planar. The dihedral angles for this system are around 4°.

The calculated dihedral angle for the *N*-ethylrhodanine with the adjacent thiophene ring (θ) is around 1.5°. The planarity of the whole π -conjugated system is responsible for the extended conjugation and this situation is consistent with the calculated wavefunctions of the frontier molecular orbitals.

The HOMO and LUMO are spread over the whole porphyrin and the corresponding thienylenvinylene system. Thus, both orbitals are overlapped and this promotes the HOMO to LUMO electronic transitions (Figure S18).

The theoretical HOMO–LUMO gaps are similar for both dyes, with a slightly lower value for compound **1b** (1.72 eV) than for **1a** (1.85 eV). These values are of the same order of magnitude as those determined by electrochemical measurements.

The greater extent of the conjugation in dye **1b** results in an increase in the HOMO energy level and, as a consequence, a bathochromic shift in the maximum absorption wavelength, which in turn improves the light harvesting behavior.

The differences between the HOMO of the donor (**1a** or **1b**) and the LUMO of the acceptor (PC71BM)[24] (**1a**: 1.12 eV, **1b**: 0.86 eV) point to efficient Voc values. Furthermore, the difference between the LUMO of the dyes **1a** and **1b** and the LUMO of PC71BM (**1a**: 0.73 eV, **1b**: 0.85 eV) ensures efficient exciton dissociation at the D/A interface.

Device Characterization

The photovoltaic (PV) properties of **1a** and **1b** were investigated by fabricating solar cells using the conventional sandwich structure of ITO/poly(3,4-ethylenedioxythiophene): polystyrenesulfonate (PEDOT:PSS)/small-molecule:acceptor/Ca/Al. The fabrication method and other details are provided in the Supporting Information. The electrical characterization of all devices was performed under ambient conditions with AM 1.5 G simulated illumination at an intensity of 100 mW cm⁻². The current density-voltage characteristics and the external quantum efficiency (EQE) are shown in Fig. 4 and the performance parameters are summarized in Table 2 as a function of the donor:PC71BM weight ratios. Average values were taken from 12 devices.

The devices based on a **1a**:PC71BM blend film with a w/w ratio of 1:2 exhibited an average power conversion efficiency (PCE) above 3%, with an open-circuit voltage (Voc) of 0.85 V, high short-circuit current (Jsc) and poor FF of 30.8% (Fig. 4a). On the other hand, the photovoltaic device with **1b**:PC71BM at a blend ratio of 1:4 w/w showed a record PCE of 4.24%, with Voc 0.8 V, a Jsc of 13.2 mA/cm² and a FF of 40.1% (Fig. 4b). Moreover, the EQE values, which are displayed in Figures 4c and 4d, were measured under monochromatic light for **1a**:PC71BM and **1b**:PC71BM complete devices at different blend ratios. Both spectra exhibit a broad spectral response, ranging from 300 to 800 nm, whereas the mixture of **1b**:PC71BM at a 1:4 w/w ratio gave the highest incident photon-to-current efficiency (80%) at 423 nm. This finding is consistent with the devices with the highest Jsc and PCE. Furthermore, the EQE integration against the 1.5 AM G sun spectra results was in good agreement with the measured photocurrents. As can be seen, all the solar cells display fill factor values below 50% (**see Table 2**). This issue was independent of the donor/acceptor ratio although, for the best solar cells with a ratio of 1:4 we achieved the highest fill factor values. From these results, it is difficult and too speculative to infer the reasons for the low fill factor. Nevertheless, we believe that the low mobility measured for the “only hole devices” and the unbalanced charge transport between holes and electrons are, definitively important factors that effect a change on the fill factor

Table 2 Main performance parameters for **1a**:PC71BM and **1b**:PC71BM devices using different ratios. The values in parenthesis correspond to the highest efficiencies reached for each of the ratio conditions.

Active layer	V_{oc} (V)	J_{sc} (mA cm^{-2})	FF (%)	PCE (%)
1a :PC71BM (1:4)	0.79 ± 0.04	11.03 ± 0.31	31.4 ± 0.2	2.73 ± 0.2 (2.86)
1a :PC71BM (1:3)	0.87 ± 0.01	11.48 ± 0.25	30.2 ± 0.1	3.01 ± 0.1 (3.04)
1a :PC71BM (1:2)	0.85 ± 0.01	11.36 ± 0.21	30.8 ± 0.3	3.01 ± 0.1 (3.05)
1b :PC71BM (1:4)	0.80 ± 0.04	13.2 ± 0.21	40.1 ± 0.29	4.24 ± 0.1 (4.35)
1b :PC71BM (1:3)	0.82 ± 0.02	12.45 ± 0.24	38.5 ± 0.24	3.91 ± 0.1 (3.98)
1b :PC71BM (1:2)	0.84 ± 0.03	12.12 ± 0.37	34.7 ± 0.26	3.55 ± 0.1 (3.63)

Figure 4 (a) Current density-voltage (J-V) characteristics of **1a**:PC71BM devices; (b) **1b**:PC71BM devices, both at different ratios; (c) EQE of **1a**:PC71BM and (d) EQE of **1b**:PC71BM devices.

Figure 5. AFM images of (a) **1a**:PC71BM and (b) **1b**:PC71BM in the most efficient devices.

Once the solar cells had been evaluated, the morphologies of the active layer were characterized by atomic force microscopy (AFM) in tapping mode in order to gain an insight into the differences in device performance, using the best composition for each device, as shown in Figure 5. The surface topographies of the film of the **1b**:PC71BM (1:4) blend displays rather uniform nanometer-sized features with a root-mean-square roughness (rms) value of 0.53 nm. In contrast, the blend containing **1a** with PC71BM exhibited a coarser topography, with higher rms values of 0.62 nm. The surface of the former is smoother than that of the latter and it also has better-defined morphological features. This is often related to better phase intermixing, which in turn leads to better phase separation and is beneficial for efficient exciton diffusion and charge transport.

The effect that the blend nano-morphology has on the solar cell parameters was analyzed further by measuring the hole carrier mobility in complete devices with the appropriate selective contacts (see Supporting Information). The mobility was calculated using the Space Charge

Limited Current method (SCLC) as described in the Supporting Information (Figure S19). Values for **1a** and **1b** of $1.25 \pm 0.8 \times 10^{-5} \text{ Vcm}^{-2}\text{s}^{-1}$ and $1.53 \pm 0.3 \times 10^{-5} \text{ Vcm}^{-2}\text{s}^{-1}$, respectively, were obtained. It should be noted that, in contrast to previous publications,[33-40] the effect of the alkyl chains in these molecules on the hole carrier mobility is less critical. Moreover, it was determined that the hole carrier mobility is not responsible for the difference between the two porphyrin-based devices and, more importantly, this physical characteristic is not considered to be a determinant factor for the final device performance.

Finally, the carrier losses were evaluated by measuring the interfacial carrier recombination kinetics as reported previously.[33, 34, 41-48]

The charge extraction of **1a-b** is depicted in Figure 6a and was determined by subtracting the contribution of the geometrical capacitance. The total extracted charges are shown in the Supporting Information (Figure S20), where the geometric contribution is indicated by a straight line and the calculated values are 80 and 64 nF cm⁻² for **1a** and **1b**, respectively. Hence, the charge values shown in Figure 6b correspond to the photo-induced charge, which can either be stored at the contacts, in the bulk of the device or both. As can be seen in Figure 6a, there is a shift in the exponential in devices made with 1a and this finding is in good agreement with the 5–10 mV increase in V_{oc} observed in the IV curves.

Moreover, on using **1b** the organic blend is capable of storing more charges at a given photo-induced voltage (the so-called light bias) and, as shown above, can extract these charges more efficiently when compared to **1a**. The Photo-Induced Transient Photo-Voltage (PIT-PV) technique was also applied to compare the charge lifetime of devices based on **1a** and **1b**. A clear difference can be seen between these two materials. Firstly, **1b** clearly has a slower carrier lifetime than **1a**. The slower carrier recombination dynamics, in addition to the observed shift in

the exponential charge density measurement, explain the higher Voc observed for **1b**. It is known that the Voc of a solar cell is closely related to the carrier recombination dynamics under illumination. The slower the carrier recombination the higher the Voc for devices with similar charge density.

Figure 6. (A) Charge density at different light intensities (light bias) of **1a** (red line) and **1b** (green line). The charges correspond to the extracted charges under different illumination conditions. (B) The carrier life-time of **1a** (red line) and **1b** (green line) at different charge values.

The results described above indicate that the molecular structure of **1b** favors better film nano-morphology and thus improved charge collection and slower carrier recombination dynamics.

Conclusions

Two conjugated molecules bearing a Zn-porphyrin, which act as an electron donor, linked by ethynylene units to one or two thienylenevinylene moieties and capped by ethylenethiol groups as acceptor units have been synthesized and characterized for solution processed small

molecule solar cells. The design strategy was based on the incorporation of electron-accepting ethylenethiophene moieties through the thiophene based bridges, which shift the absorption profiles bathochromically to the NIR due to a more intense intramolecular charge transfer band. Compounds **1a** and **1b** exhibit excellent light harvesting properties, good thermal stability and have low-lying HOMO levels of -5.36 and -5.24 eV, respectively. Complete SMOSCs were fabricated using **1a** and **1b** donors combined with PC71BM as the electron acceptor molecule. Compound **1b** gave a PCE of 4.3%. The EQE measurements are consistent with the J_{sc} values obtained, with an incident photon-to-current efficiency of 80% at 430 nm for the **1b**:PC71BM device with a 1:4 ratio. The hole carrier mobility was also measured and values of $1.25 \pm 0.8 \times 10^{-5}$ $\text{Vcm}^{-2}\text{s}^{-1}$ and $1.56 \pm 0.4 \times 10^{-5}$ $\text{Vcm}^{-2}\text{s}^{-1}$ were obtained, respectively. Finally, photo-induced charge extraction and photo-induced transient photo-voltage experiments were carried out in order to determine the origin of the difference in performance between devices based on **1a** and **1b**. The PICE showed a shift that mirrored the difference in V_{oc} and very similar distribution of charges versus applied light bias, although **1b** can store more charge at the same voltage. Moreover, the PIT-PV showed a marked difference between the two molecules; In the case of **1b** the carrier lifetime was considerably shorter than that of **1a**. Thus, for **1b** the presence of the thiophene groups not only improves the light absorption but also changes the nano-morphology and, hence, the carrier extraction and recombination. This is achieved without a substantial decrease in the device V_{oc} , as expected due to the reduced bandgap of **1b**.

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Graphical Abstract