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Fine Tuning of Fluorene-Based Dye Structures for High-Efficiency *p*-Type Dye-Sensitized Solar Cells

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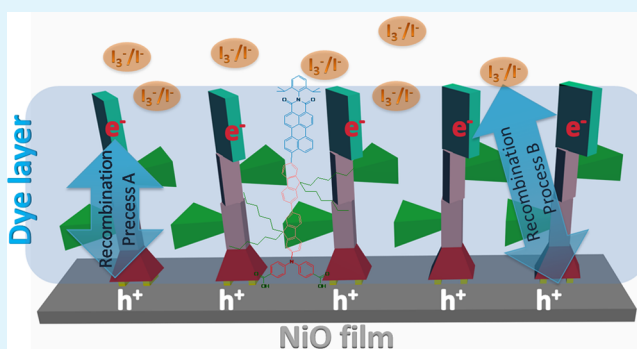
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S Supporting Information

ABSTRACT: We report on an experimental study of three organic push–pull dyes (coded as zxx-op1, zxx-op1–2, and zxx-op1–3) featuring one, two, and three fluorene units as spacers between donors and acceptors for *p*-type dye-sensitized solar cells (*p*-DSSC). The results show increasing the number of spacer units leads to obvious increases of the absorption intensity between 300 nm and 420 nm, a subtle increase in hole driving force, and almost the same hole injection rate from dyes to NiO nanoparticles. Under optimized conditions, the zxx-op1–2 dye with two fluorene spacer units outperforms other two dyes in *p*-DSSC. It exhibits an unprecedented photocurrent density of 7.57 mA cm^{−2} under full sun illumination (simulated AM 1.5G light illumination, 100 mW cm^{−2}) when the I[−]/I₃[−] redox couple and commercial NiO nanoparticles were used as an electrolyte and a semiconductor, respectively. The cells exhibited excellent long-term stability. Theoretical calculations, impedance spectroscopy, and transient photovoltage decay measurements reveal that the zxx-op1–2 exhibits lower photocurrent losses, longer hole lifetime, and higher photogenerated hole density than zxx-op1 and zxx-op1–3. A dye packing model was proposed to reveal the impact of dye aggregation on the overall photovoltaic performance. Our results suggest that the structural engineering of organic dyes is important to enhance the photovoltaic performance of *p*-DSSC.

KEYWORDS: *p*-type dye-sensitized solar cells, organic sensitizer, fluorene, charge recombination, NiO



INTRODUCTION

It is still challenging to convert solar energy to electricity in a cost-effective manner. Among many devices, dye-sensitized solar cells, especially *n*-type dye-sensitized solar cells (*n*-DSSC), have been extensively investigated.^{1–3} Recently, *p*-type dye-sensitized solar cells (*p*-DSSC) become attractive because of their potential to construct high-efficiency tandem cells with *n*-DSSC through molecular engineering of light-harvesting materials.^{4–8} However, low photocurrent densities from state-of-the-art *p*-DSSC limit output of current density of tandem cells; therefore, it has become a bottleneck for the development of high-efficiency tandem cells.^{4–7} To date, the highest photocurrent density for *p*-DSSC is 7.0 mA cm^{−2},⁹ which is much lower than the current density of its counterpart, which has a range of 15–20 mA cm^{−2}.^{2,3,10–13} The highest power conversion efficiency (PCE) of *p*-DSSC is 1.30%,¹⁴ which is also far below the record PCE of 13.0% in *n*-DSSC.^{10,13}

A key challenge in *p*-DSSC is how to reduce the charge recombination between the injected hole in the semiconductor and the reduced dyes or electrolytes.^{4,5,15–18} One strategy is to use rationally designed light-harvesting material to retard those processes. One method is to engineer push–pull dyes to enhance the hole injection capability.^{17,19–27} Sun et al. synthesized triphenylamine-based push–pull dyes and found that the photocurrent density increased significantly, to 5.48 mA cm^{−2}, on NiO films,^{19,28} which was much higher than its counterparts (coumarin dye, C343) without such a structure. Bäuerle et al. designed an oligothiophene-functionalized push–pull dye, coded as PMI-6T-TPA, and a photocurrent density of 5.35 mA cm^{−2} was obtained in similar cells.⁷ Recently, Bach et al. reported a record photocurrent density of 7.0 mA cm^{−2}

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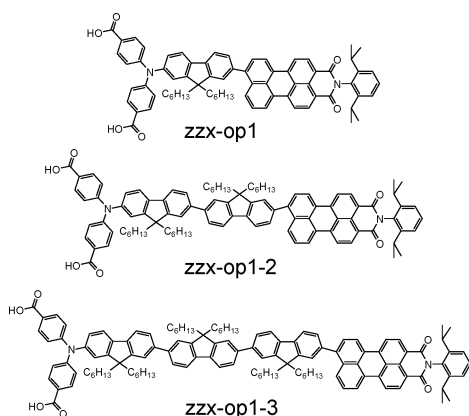
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when the same PMI-6T-TPA dye and the nanostructured NiO microballs were used.⁹ Previously, we designed three fluorene-based dyes—zzx-op1, zxx-op2, and zxx-op3—to modulate hole injection in *p*-DSSC.²⁹ Our study showed that ~ 0.80 eV of hole injection driving force was needed to achieve a high hole injection quantum yield when NiO and I^-/I_3^- were used in the devices. The dye zxx-op1 with a di(*p*-carboxy)phenylamine (DCPA) as a donor, a perylenemonoimide (PMID) as an electron acceptor, and one fluorene (FLU) as a spacer produced a photocurrent density of 4.38 mA cm^{-2} with a PCE of 0.184%, indicating the promising of this dye for *p*-DSSC. However, fast charge recombination between the injected hole and the reduced dye was observed in zxx-op1-sensitized *p*-DSSC. This comes primarily from a short distance between the PMID donor the surface of NiO nanoparticles, which results in a significant photocurrent losses.^{5,7}

Two new oligo-fluorene-based dyes, zxx-op1-2 and zxx-op1-3, were designed to address this issue (Scheme 1). In

Scheme 1. Structure of Dyes zxx-op1, zxx-op1-2, and zxx-op1-3



these dyes, the DCPA was used as a donor, a PMID as an electron acceptor, two and three conjugated 9,9-dihexyl-9H-fluorene (FLU) units were used as spacers. We expect this design will tune the dye structure to retard the aforementioned recombinations. It was found that the insertion of FLU units only changed the hole driving force subtly. However, the FLU units extended the distance between PMID and the surface of NiO nanoparticles and retarded the charge recombination between the injected hole and the reduced dye, which reduced the photocurrent losses effectively.⁷ As a result, dyes zxx-op1-2 and zxx-op1-3 showed much better performance than zxx-op1. Under optimized conditions, an unprecedented photocurrent density of 7.56 mA cm^{-2} was obtained.

RESULTS AND DISCUSSION

The synthesis of dyes zxx-op1-2 and zxx-op1-3 is shown in Scheme 2. Brominated oligo-fluorene compounds **1** and **6** were synthesized according to the literature procedures,^{30,31} and reacted with diphenylamine under Buchwald–Hartwig reaction conditions to give diphenylamine–oligo-fluorene derivatives **3** and **7** in yields of 23% and 25%, respectively.²² Both then reacted with the borate–perylenemonoimide derivative (**4**) under Pd-catalyzed Suzuki cross-coupling reaction conditions to produce compounds **5** and **8** in yields of 88% and 83%, respectively.²² They were then hydrolyzed in the presence of TFA to give zxx-op1-2 and zxx-op1-3 in yields of 89% and

82%, respectively.²² The structures and purity of all intermediates and final compounds were characterized by NMR and high-resolution mass spectrometry (see details in the Electronic Supporting Information (ESI)).

The ultraviolet–visible (UV-vis) absorption spectra of all three dyes in THF are quite similar to each other. The details have been tabulated in Table 1. As shown in Figure 1, two major absorption bands in the range of 300–600 nm were observed for three dyes. The first band (~ 360 nm) comes from the FLU linker and the donor group. When the spectra were normalized at 520 nm, the intensity of the first band increased with the number of FLU repeating units. The second band (~ 520 nm) presented a small blue-shifted due to the π – π^* electron transition. This result is likely attributed to the aggregation of dyes in THF. When these dyes were stained on the NiO films, the absorption spectra became broader compared to those in solution (see Figure S1 in the ESI), and peak positions shifted to a shorter wavelength region. The broadened spectra may be ascribed to the aggregation on the surface of semiconductor, and the shifted peak positions may be ascribed to the deprotonation of the carboxylic acid or the different polarity of the medium and H-aggregation.^{25,32} The fluorescence spectra (see Figure S2 in the ESI) of three dyes were also quite similar to each other. The zero–zero transition energy ($E_{0-0}(S^*)$), determined from the crossing points of the normalized UV-vis absorption spectra and the fluorescence spectra of the dyes in THF (see Figure S2 in the ESI), were 2.21 eV for zxx-op1, 2.23 eV for zxx-op1-2, and 2.23 eV for zxx-op1-3, respectively. This indicated that increasing the number of FLU repeating units did not change the energy gap of the dyes significantly. The E_{ox} and E_{red} obtained from cyclic voltammograms (see Figure S3 in the ESI) for three dyes were almost identical to each other. As a result, the hole injection driving forces (ΔG_{inj}) for three dyes were evaluated to be -0.82 eV for zxx-op1, -0.82 eV for zxx-op1-2, and -0.83 eV for zxx-op1-3, respectively. Similar results for dye regeneration driving force were also observed (-0.66 eV for zxx-op1, -0.68 eV for zxx-op1-2, and -0.67 eV for zxx-op1-3, respectively). Those results indicate that the three dyes are energetically favorable for the operation of *p*-DSSC.

The photocurrent–voltage curves of the champion *p*-DSSC with the three dyes in different solvents are shown in Figure 2, and the averaged devices performance is shown in the ESI. The devices photovoltaic parameters are listed in Table 2. Acetonitrile has been widely used as a solvent for preparation of dye solution; however, we found that the three dyes exhibited low solubility in acetonitrile and the resulting NiO films were poorly stained. To overcome this problem, acetonitrile/THF solvent systems were used to prepare the dye solution. To our surprise, we found the solvent affected the overall cell performance significantly. When a dye solution of THF/acetonitrile (THF/AN, v/v = 2/1) was used, it was observed that the V_{OC} values of devices B (zxx-op1-2) and C (zxx-op1-3) were very similar to each other (~ 102 mV); both were higher than that of device A (zxx-op1, 96 mV). However, the short-circuit current density (J_{SC}) of devices varied in the following order:

$$\text{device A } (5.70 \text{ mA cm}^{-2}) < \text{device C } (6.53 \text{ mA cm}^{-2}) \\ < \text{device B } (7.03 \text{ mA cm}^{-2})$$

As a result, device B with zxx-op1-2 dye gave the highest PCE of 0.279%. This is consistent with its remarkably higher IPCE

Scheme 2. Synthesis Route of zzx-op1–2 and zzx-op1–3

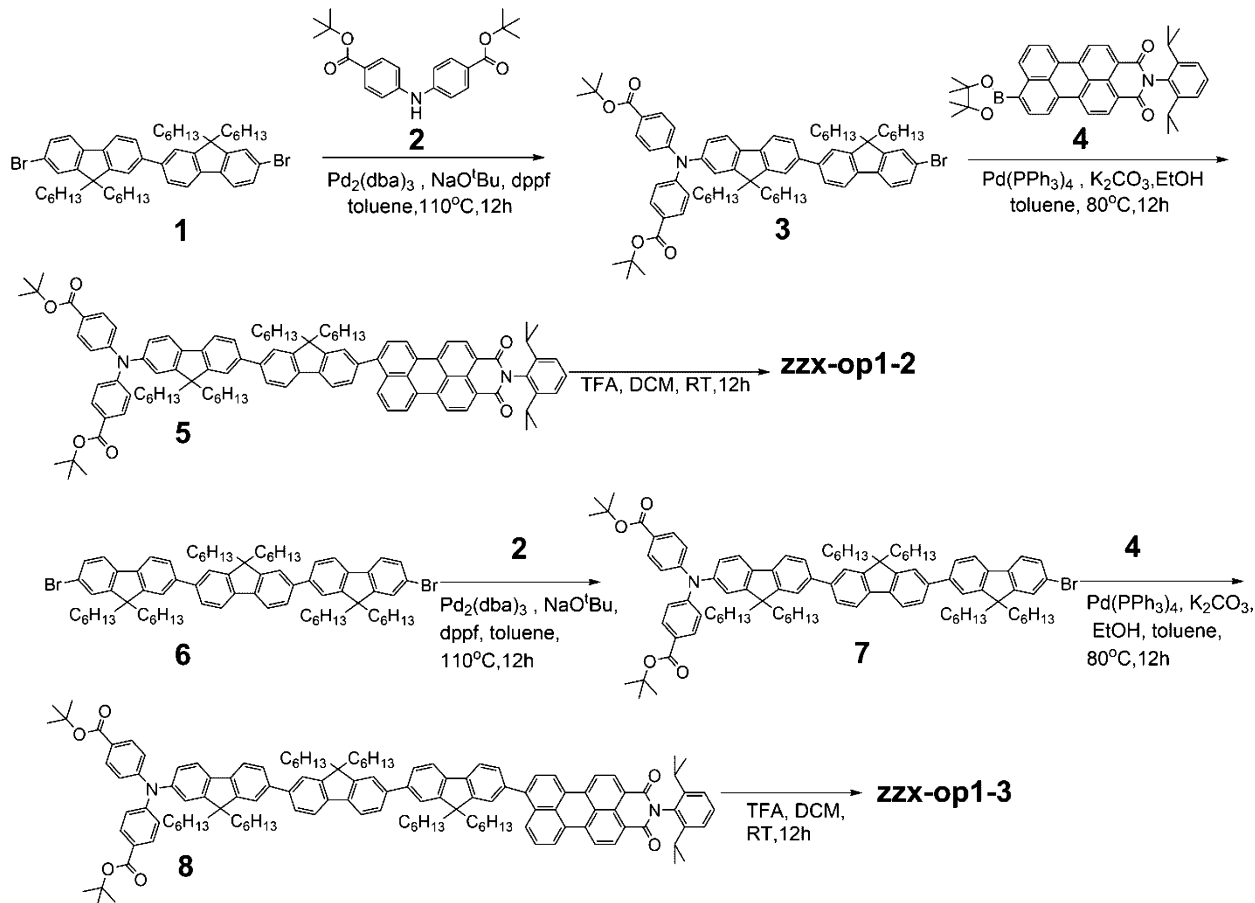


Table 1. Optical and Electrochemical Data of Dyes zzx-op1, zzx-op1–2, and zzx-op1–3

dye	λ_{\max}^a (nm)	E_{ox}^b (V, vs NHE)	E_{red}^b (V, vs NHE)	E_{0-0}^c (eV)	ΔG_{inj}^d (eV)	ΔG_{reg}^e (eV)
zzx-op1	354 518	1.28	−0.85	2.21	−0.82	−0.66
zzx-op1–2	362 517	1.24	−0.87	2.23	−0.82	−0.68
zzx-op1–3	369 517	1.24	−0.86	2.23	−0.83	−0.67

^aOptical properties of all dyes were measured in THF at room temperature. ^bElectrochemical properties of all dyes are measured in THF at room temperature. The supporting electrolyte was 0.1 M TBAPF₆. Potentials measured vs ferrocene/ferrocenium (Fc/Fc⁺) couple were converted to normal hydrogen electrode (NHE) by addition of +0.63 V. ^cThe zero-zero transition energy ($E_{0-0}(S^*)$) was estimated from the intersection of normalized absorption and emission curves. ^dCalculated according to the equation: $\Delta G_{\text{inj}} = e[E_{\text{VB}}(\text{NiO}) - (E_{0-0}(S^*) + E_{\text{red}}(S/S^-))]$, $E_{\text{VB}}(\text{NiO}) = 0.54$ V vs NHE. ^eCalculated according to the equation: $\Delta G_{\text{reg}} = e[E(M/M^-) - E_{\text{red}}(S/S^-)]$, $E(I_3^-/I_2^- \cdot) = -0.19$ V vs NHE.³³

with the maximum of 64.3% at 520 nm, compared to the 50.1% for device A with zzx-op1. When dye solutions of THF/AN (v/v = 1.5/1) and THF/AN (v/v = 1/1) were used for zzx-op1–2, both the photocurrent density and open-circuit voltage increased. Device E (with zzx-op1–2 dye) displayed a PCE of 0.353%, a V_{OC} of 117 mV, a fill factor (FF) of 0.40, and a J_{SC} of 7.57 mA cm^{−2}. To the best of our knowledge, this is the highest photocurrent density for *p*-DSSC. Its performance is consistent with the remarkably higher IPCE, with a maximum of 70.2% at 520 nm, as shown in Figure 3.

The increase of photocurrent density in zzx-op1–2 and zzx-op1–3 is consistent with their dye-loading densities on the NiO films. When the zzx-op1–2 was dissolved in THF/AN (v/v = 2/1), THF/AN (v/v = 1.5/1), and THF/AN (v/v = 1/1), the dye-loading density increased from 1.11×10^{-8} mol cm^{−2} μm^{-1} , to 1.24×10^{-8} mol cm^{−2} μm^{-1} , to 1.28×10^{-8} mol cm^{−2} μm^{-1} , because of the augmentation of adsorption–desorption equilibrium constant,³⁴ the photocurrent density increased

from 7.03 mA cm^{−2} to 7.44 mA cm^{−2} to 7.57 mA cm^{−2}. A similar trend was also observed when zzx-op1–3 was dissolved in THF/AN (v/v = 2/1, 0.98×10^{-8} mol cm^{−2} μm^{-1}) and chlorobenzene (1.11×10^{-8} mol cm^{−2} μm^{-1}). However, the discrepancy between dye-loading density and photocurrent density was also observed among three dyes. When the THF/AN (v/v = 2/1) was used as a solvent, the measured dye-loading densities of zzx-op1, zzx-op1–2, and zzx-op1–3 on NiO films were 1.20×10^{-8} , 1.11×10^{-8} , and 0.98×10^{-8} mol cm^{−2} μm^{-1} , respectively. The zzx-op1–2 with a moderate dye-loading density exhibited the highest photocurrent density. When zzx-op1–2 were dissolved in THF/AN (v/v = 2/1) and zzx-op1–3 in chlorobenzene, respectively, an identical dye-loading density (1.11×10^{-8} mol cm^{−2} μm^{-1}) was observed; zzx-op1–2 produced a higher photocurrent density than zzx-op1–3 (7.03 mA cm^{−2} vs 6.68 mA cm^{−2}). This discrepancy indicated the impact of structure difference on the overall cell performance.

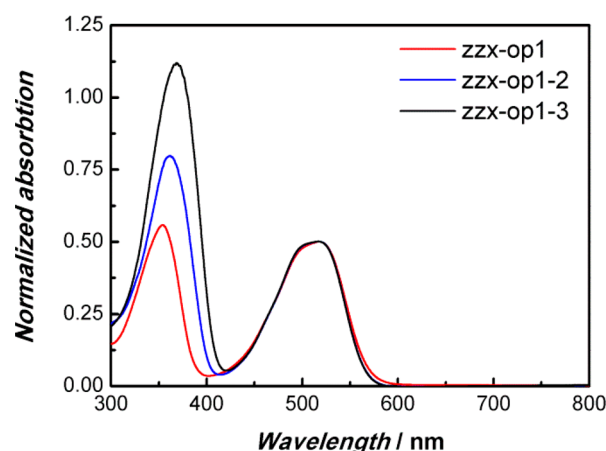


Figure 1. Absorption spectra of zzx-op1, zzx-op1-2, and zzx-op1-3 in THF.

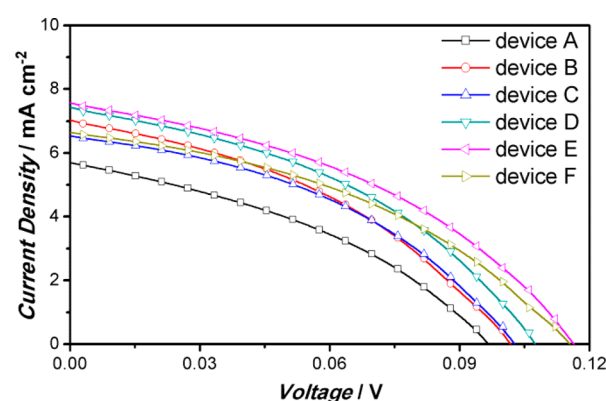


Figure 2. Current–voltage (J – V) characteristics of champion devices under simulated AM 1.5 (100 mW cm^{-2}) conditions. The NiO films for devices A–C were made with dyes zxx-op1, zxx-op1-2, and zxx-op1-3 in THF/AN ($v/v = 2/1$) solution, devices D and E with zxx-op1-2 in THF/AN ($v/v = 1.5/1$) and THF/AN ($v/v = 1/1$), and device F with dye zxx-op1-3 in chlorobenzene.

The density functional theory (DFT) calculation were performed to study the structural conformation and electron density distribution of the dyes. As shown in Figure 4, the electron density distributions of the lowest unoccupied molecular orbitals (LUMOs) for the three dyes were quite similar to each other and was significantly localized on the PMID unit. The electron density distribution of the highest occupied molecular orbitals (HOMOs) was quite different. In zxx-op1, the electron density in the donor section was more pronounced and was extended significantly to acceptor moiety (PMID unit). In zxx-op1-2, the electron density in the donor reduced with concurrent increase on the first and second FLU

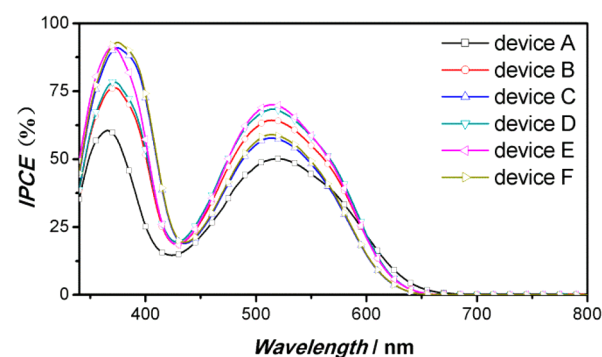


Figure 3. Photocurrent action spectra of devices A–F.

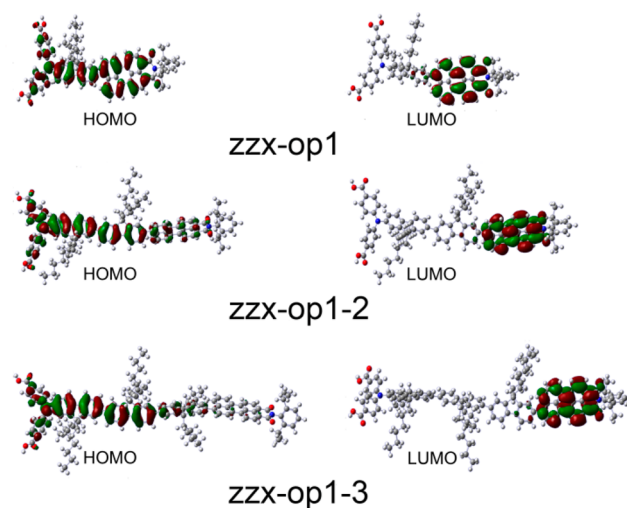


Figure 4. Geometry-optimized structures of zxx-op1, zxx-op1-2, zxx-op1-3 and the electron density distribution of frontier molecular orbitals from DFT calculations with the B3LYP/6-311G(d) model.

unit and extension to PMID unit. In zxx-op1-3, the electron density in the donor reduced with a concurrent increase on the first and second FLU unit and extension to the third FLU unit. In both cases, a disjoint character was observed, which will benefit the charge separation. The length of oligo-fluorene linkers was determined to be 6.90, 18.3, and 26.8 Å for zxx-op1, zxx-op1-2, and zxx-op1-3, respectively. The spatial distribution of the dyes is in an approximately straight line, and thus the molecule might be aligned predominantly perpendicular to the NiO surface. (The detailed torsion angle of π -linkers and PMID are shown in the Table S1 in the ESI.) The dyes with longer linkers (zxx-op1-2 and zxx-op1-3) presented better energy conversion efficiency, which was also observed in several other systems.^{7,35}

Table 2. Photovoltaic Data of zxx-op1, zxx-op1-2, and zxx-op1-3 Sensitized Champion Solar Cells under an Illumination of 100 mW cm^{-2} , AM1.5G Conditions

device	dye	solvent (v/v)	J_{SC} (mA cm^{-2})	V_{OC} (V)	fill factor, ff	η (%)
A	zxz-op1	THF/AN (2/1)	5.70	96	0.38	0.207
B	zxz-op1-2	THF/AN (2/1)	7.03	102	0.39	0.279
C	zxz-op1-3	THF/AN (2/1)	6.53	103	0.41	0.276
D	zxz-op1-2	THF/AN (1.5/1)	7.44	106	0.41	0.322
E	zxz-op1-2	THF/AN (1/1)	7.57	117	0.40	0.353
F	zxz-op1-3	chlorobenzene	6.68	115	0.40	0.308

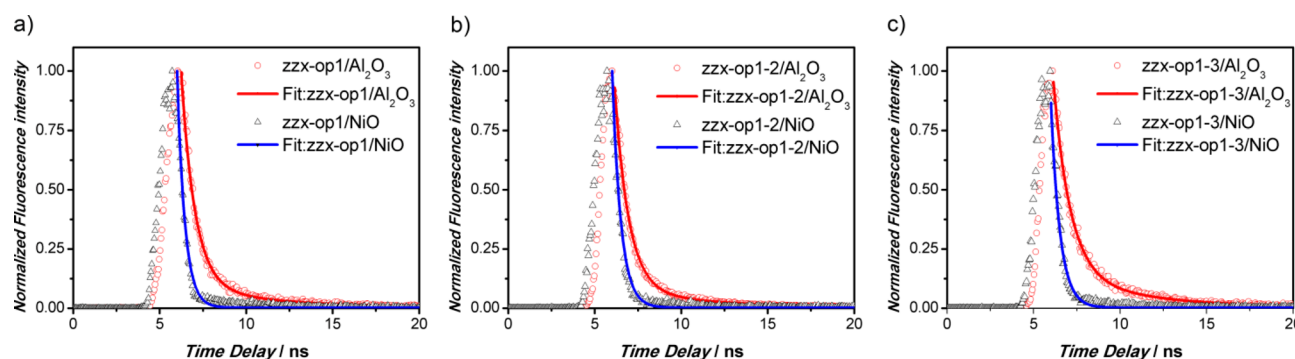
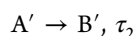
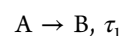


Figure 5. Time-resolved PL decay traces of dye-sensitized NiO films (black triangle, Δ), Al_2O_3 films (red circle, \circ), fitted line for dyes on NiO film (blue line), Al_2O_3 film (red lines): (a) zzx-op1, (b) zzx-op1-2, and (c) zzx-op1-3. Excitation wavelength = 445 nm.

Time-resolved fluorescence decays measurements were performed to evaluate the hole injection kinetics. The measurements were performed on dye-coated NiO films and compared with those from Al_2O_3 films.^{36–39} The NiO films and Al_2O_3 films with the same thickness ($L \approx 3.0 \mu\text{m}$) were used. The result is shown in Figure 5. To resolve the hole injection kinetics, the time coefficients from an amplitude-averaged decay time model were used to determine the corresponding rate coefficients, $k_{\text{NiO}} = \tau_{\text{NiO}}^{-1}$ and $k_{\text{Al}_2\text{O}_3} = \tau_{\text{Al}_2\text{O}_3}^{-1}$, according to the report of Diau et al.^{36,40} In this model, the decay data were fitted with a parallel kinetic model:



$$f(t) = a \exp\left(-\frac{t}{\tau_1}\right) + b \exp\left(-\frac{t}{\tau_2}\right)$$

Components A and A' are described with a single-exponential function and decay coefficients τ_1 and τ_2 , respectively. The pre-exponential factors a and b represent the relative weights (or amplitudes) of components A and A', respectively. For Al_2O_3 films, two decay components are required to fit the fluorescence transients. This is because the PL decays of the dye-coated Al_2O_3 films reflect the radiative decay to the ground state, as well as intermolecular energy transfer, because of aggregation of the dye on the surface of Al_2O_3 film. These two processes were in the similar time scale. For NiO films, only one component is required to fit the data. This is because the hole injection is much faster than intermolecular energy transfer (IET). Because of the limitation of our instrumentation, the observed decay profiles only reflected the IET, which was of course affected by hole injection into the valence band of NiO.³⁶ Assuming the dye aggregation on the NiO and Al_2O_3 was similar when the absorption of the films are same, and the nonradiative relaxations are also similar to each other, the averaged fluorescence lifetime therefore can be used to reveal the hole injection kinetics. The average fluorescence lifetimes for zzx-op1, zzx-op1-2, and zzx-op1-3 sensitized Al_2O_3 films were evaluated to be 2.27, 2.26, 2.70 ns, respectively, while the strong quenching of the emission for those dyes were observed when they were stained on the NiO films. Considering the reconvolution of the instrument response function of ~ 89 ps, the IRF response and the data fitting results with parallel kinetics model indicated that the lifetime of the emission of the three dyes stained on the NiO films were 0.44, 0.50, and 0.53 ns, respectively. This result confirms a rapid hole injection from

the excited state of the dye into the valence band of NiO.³⁷ The hole injection quantum yield can be approximately described according to the equation $\Phi_{\text{inj}} = 1 - \tau_{\text{NiO}}/\tau_{\text{Al}_2\text{O}_3}$, which was 80% (zzx-op1), 78% (zzx-op1-2), and 80% (zzx-op1-3), respectively.³⁶ It indicates that the three dyes showed nearly identical hole injection process. It should be mentioned that the injection yield could be higher than those showed here due to deadsorbed dye inside the films.

The dark current and electrochemical impedance spectroscopy (EIS) studies revealed different charge recombination behavior of devices from three dyes.^{5,22,41–43} As shown in Figure 6, the zzx-op1 and zzx-op1-2 sensitized cells (devices A

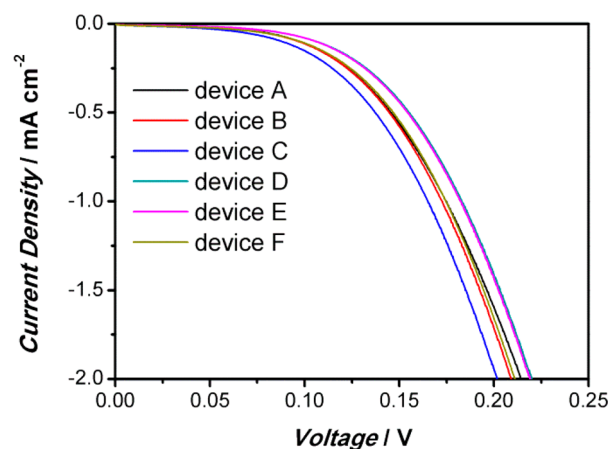


Figure 6. Dark current density as a function of applied bias obtained from dark current measurement of devices A–F.

and B) exhibited smaller dark current densities than that of zzx-op1-3 sensitized cell (device C) at the same bias, implying compact layers of zzx-op1 and zzx-op1-2 on the surface of the NiO particles were formed to prevent the charge recombination between the injected hole and the hole acceptor in electrolytes. This is consistent with higher recombination resistance observed for devices A and B in impedance results. When different solvent systems were used, smaller dark current densities were observed for devices D and E. In EIS study, the NiO/dye/electrolyte interfacial charge transfer resistance (i.e., charge recombination resistance, R_{rec}) was evaluated using a transmission line model.^{42–44} As shown in Figure 7, device A (zzx-op1) and device B (zzx-op1-2) showed higher R_{rec} than device C (zzx-op1-3) at a given bias. This result agrees with the observation on the dark current, indicating device C

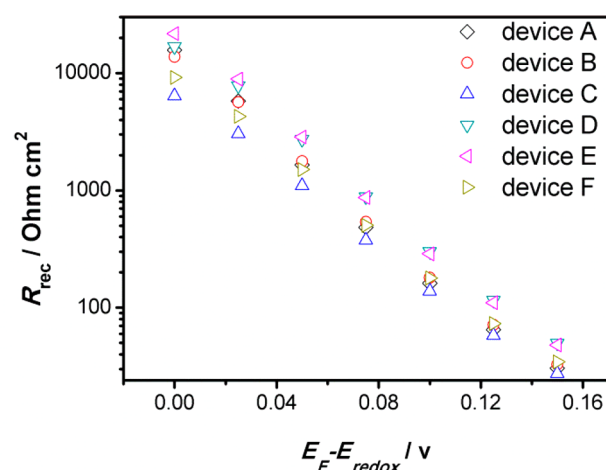


Figure 7. Interfacial charge recombination resistance (R_{rec}), as a function of applied bias obtained from electrochemical impedance measurement in darkness for devices A–F.

suffered from charge recombination between the injected hole and hole acceptor in electrolytes. This implies that zxx-op1 and zxx-op1–2 might form a more compact dye layers on the surface of NiO films than zxx-op1–3 when THF/AN ($v/v = 2/1$) was used. The situation can be changed with the solvent. For example, device F exhibited an increased R_{rec} value, which was due to the application of chlorobenzene.

Transient photovoltage/photocurrent decay measurements showed longer hole lifetime in zxx-op1–2 sensitized solar cells.^{45,46} Figure 8a presents plots of the hole density as a

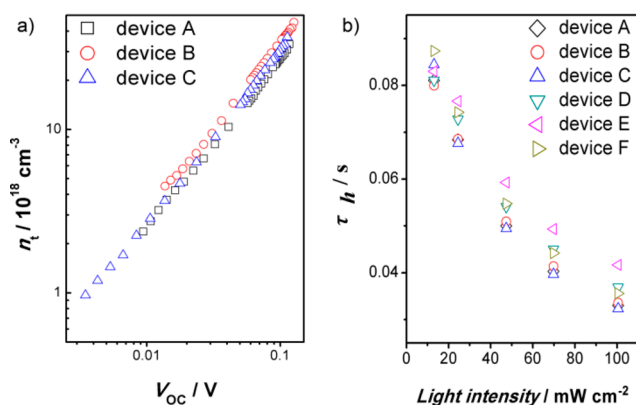


Figure 8. Transient photovoltage/photocurrent decay and charge-extraction measurements: (a) hole density as a function of bias voltage (V_{OC}) of devices A, B, and C; (b) hole lifetime as a function of light intensity of devices A–F.

function of the potential bias of device A (zxx-op1), device B (zxx-op1–2), and device C (zxx-op1–3). It perceived that devices B and C exhibited higher hole density than that of device A. As illustrated in Figure 8b, under the same light intensity, device B exhibited longer hole lifetime (τ_h) than that of devices A and C. These results are quite fitted with the results from dark current and impedance measurements. In the case of zxx-op1–2, as illustrated in Figure 8b, device E exhibited a longer lifetime than that of device B under the same light intensity. This was attributed to a more compact dye block layer for device E, efficiently retarding the charge recombina-

tion between the injected holes and the hole acceptor in electrolytes.

In *p*-DSSC, the overall performance linked tightly to the charge recombination of the injected hole in a semiconductor with the reduced dye (process A) and with hole acceptor in electrolyte (process B), as shown in Figure 9.^{4,5} It is critical to

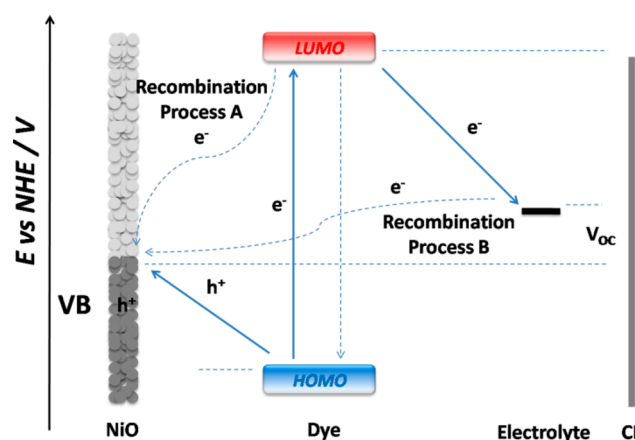


Figure 9. Schematic of charge transfer process in *p*-DSSC, arrows with solid lines denote desired transfer for energy conversion, and those with dashed lines denote undesired transfer. Recombination Process A represents the recombination between the injected hole and the reduced dye, and Recombination Process B represents the recombination between the injected hole in NiO film and the hole acceptor in electrolyte.

reduce such recombination in order to obtain better performance. This could be achieved by either applying a thin layer of metal oxide, depositing an insulation layer, or forming a more compact layer of organic dyes, providing a shield layer for the charge recombination.^{47–52} This can be used to explain the different photovoltaic performance of zxx-op1, zxx-op1–2, and zxx-op1–3. As illustrated in Figure 10, when the dyes adsorbed

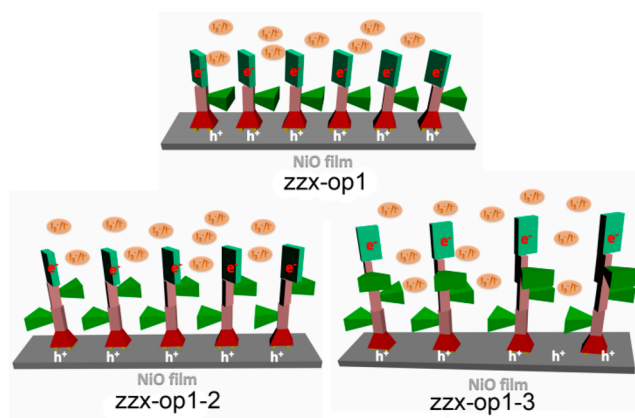


Figure 10. Pictorial representation of the self-assembled dye layer of zxx-op1, zxx-op1–2, and zxx-op1–3 on the surface of NiO.

on the NiO nanoparticles, they tend to anchored vertically on the surface of NiO nanoparticles due to the “Y”-shaped feature.⁷ The zxx-op1 with a more rigid structure formed a much compact dye layer, which retarded Recombination Process B. However, it will not provide sufficient shielding to retard Recombination Process A, because of the short length between the PMID unit and the surface of NiO nanoparticles.

The zzx-op1–3, having the longest molecule length, effectively retarded Recombination Process A,⁷ but not Recombination Process B. The less-compacted dye layer on the surface of NiO nanoparticles provides a large void for easy penetration of iodide electrolyte into dye assembly to increase the recombination probability of process B. The dye zzx-op1–2 with moderate length formed a compact dye layer, which could retard Recombination Processes A and B simultaneously. As a result, the dye zzx-op1–2 achieved better device performance than zzx-op1 and zzx-op1–3.

Finally, the long-term stability zzx-op1–2 sensitized solar cell (device E) was evaluated in darkness. Figure S7 in the Supporting Information shows the evolution of photovoltaic parameters of device E stored in darkness at ambient temperature (20 °C). During a month-long period of time, only a slight decrease of J_{SC} and V_{OC} was observed. Meanwhile, an averaged photocurrent density of 7.27 mA cm⁻² was observed. The PCE remained almost unchanged, with a slightly increased fill factor. This impressive result is indicative of the superior thermal stability of *p*-DSSC devices based on oligo-fluorene-based organic dyes.

CONCLUSIONS

In summary, we have synthesized three oligo-fluorene-based organic push–pull dyes for *p*-DSSC. The unprecedented photocurrent density of 7.57 mA cm⁻² with a power conversion efficiency (PCE) of 0.357% was achieved under AM 1.5G conditions. Time-resolved fluorescence decays measurement showed that three dyes were energetically favorable for efficient hole injection. The dark current and electrochemical impedance measurements revealed that the zzx-op1 and zzx-op1–2 sensitized cells (devices A and B) exhibited smaller dark current densities than that of the zzx-op1–3 sensitized cell. Transient photovoltage/photocurrent decay measurements showed longer electron lifetime in zzx-op1–2 sensitized solar cells. All results indicated that the dye zzx-op1–2 formed a much more compact dye block layer than zzx-op1 and zzx-op1–3, with regard to retarding the charge recombination between the injected hole in the NiO film and the hole acceptor in the electrolyte. The long-term stability study also showed the superior stability of *p*-DSSC devices based on oligo-fluorene-based organic dye.

EXPERIMENTAL SECTION

Devices Fabrication. NiO nanoparticle powder from Inframat (72.2 wt % Ni, Catalog No. 28N-0801, particle size \approx 20 nm, BET multipoint specific surface area, SSA > 50 m²/g) was used as received. NiO paste was prepared by mixing an ethyl cellulose solution in ethanol (44 wt %), terpineol (46 wt %), and NiO (10 wt %). The paste was screen-printed on FTO glass (NSG 15 Ω /□, 2.2 mm thick) by a commercial semiautomatic screen printer. The films were dried for 10 min at 120 °C. The thickness of films (3.1 μ m thick) was determined by a profile system (DEKTA, VEECCO, Bruker). The films were then sintered under an air flow at 500 °C for 30 min with a ramping time of 40 min from room temperature (\sim 25 °C) to 500 °C and then cooled to 80 °C in air. The films were then dipped into a dye solution either in a mixture of THF/AN or chlorobenzene at room temperature for 18 h. After washing with acetonitrile and drying with air flow, the sensitized NiO working electrode and the counter electrode were assembled together using a 25- μ m-thick Surlyn gasket. A pneumatic finger press was used to press the working electrode while heating the counter electrode. The counter electrodes were produced by applying one drop of H₂PtCl₆ (10 mM in ethanol) to FTO glass and thermally decomposing by firing at 400 °C for 15 min under a gentle flow of air. The electrolyte (1.0 M LiI, 0.1 M I₂ in

acetonitrile) was vacuum-backfilled in the cell cavity. The backfilled hole was sealed using an aluminum-Surlyn sheet, which was prepared by heating aluminum foil with a 25- μ m Surlyn sheet on a hot plate at 120 °C.

Devices Characterization. A 450 W xenon light source solar simulator (Oriel, model 9119) with an AM 1.5G filter (Oriel, model 91192) was used to give an irradiance of 100 mW cm⁻² at the surface of the solar cells. The current voltage characteristics of the devices under these conditions were obtained by applying external potential bias to the devices and measuring the generated photocurrent with a digital sourcemeter (Keithley, Model 2400, USA). A similar data acquisition system was used to control the incident photon-to-current conversion efficiency (IPCE) measurement. A white light bias (1% sunlight intensity) was applied onto the devices during the IPCE measurements with the AC model (10 Hz). The devices with the photoanode area of 0.40 cm \times 0.40 cm were tested with a metal mask: 0.30 cm \times 0.30 cm to prevent the scattering light.

ASSOCIATED CONTENT

Supporting Information

Synthetic procedures and characterization data of all new compounds, emission spectra, absorption spectra on NiO nanoparticles, cyclic voltammogram, PL decay, EIS and long-term stability data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

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