

A non-fullerene electron acceptor modified by thiophene-2-carbonitrile for solution-processed organic solar cells



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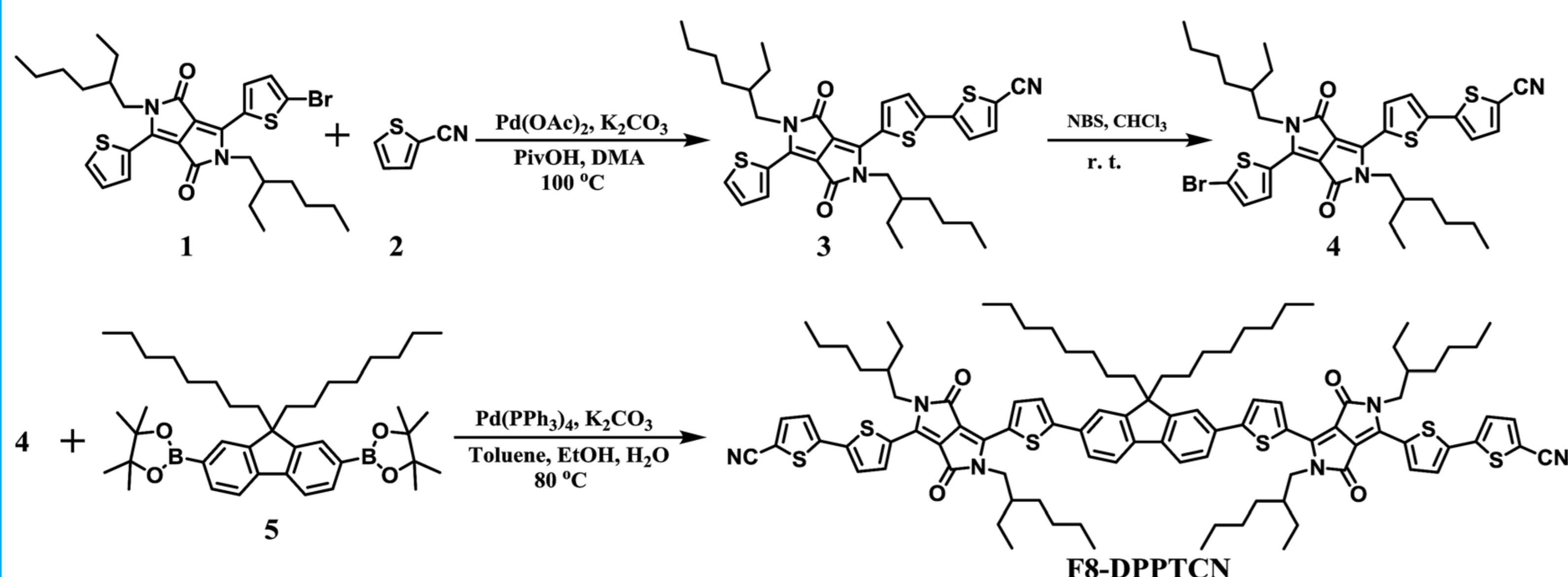


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Introduction

Effective electron acceptor materials usually have a deep lowest unoccupied molecular orbital (LUMO) energy level that can split excitons and generate current. A non-fullerene electron acceptor (F8-DPPTCN) was developed, using fluorene as the core with arms of diketopyrrolopyrrole (DPP) having thiophene-2-carbonitrile as the terminal units. The new molecule had a LUMO of -3.65 eV and a narrow bandgap (E_g) of 1.66 eV, owing to the electronegativity of the thiophene-2-carbonitrile group and its conjugation with DPP units. Organic solar cells (OSCs) with F8-DPPTCN as the acceptor and poly(3-hexylthiophene) (P3HT) as the donor were fabricated. A power conversion efficiency (PCE) of 2.37% was obtained with an open-circuit voltage (V_{oc}) of 0.97 V, a short-circuit current (J_{sc}) of 6.25 mA cm⁻², and a fill factor (FF) of 0.39.

Results and discussion



Scheme 1 Synthetic route of F8-DPPTCN.

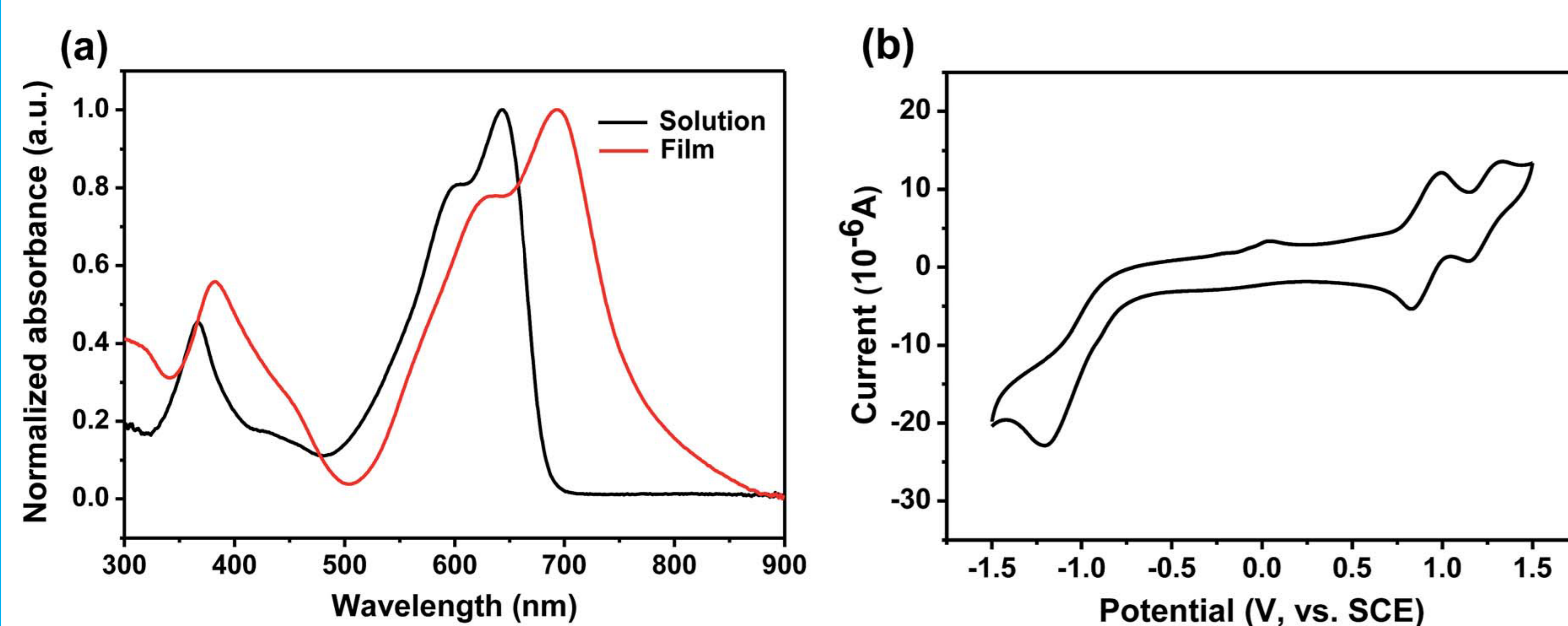


Fig. 1 (a) UV-vis absorption spectra of F8-DPPTCN in CHCl₃ solution and films; (b) cyclic voltammogram of F8-DPPTCN in CH₂Cl₂ solution.

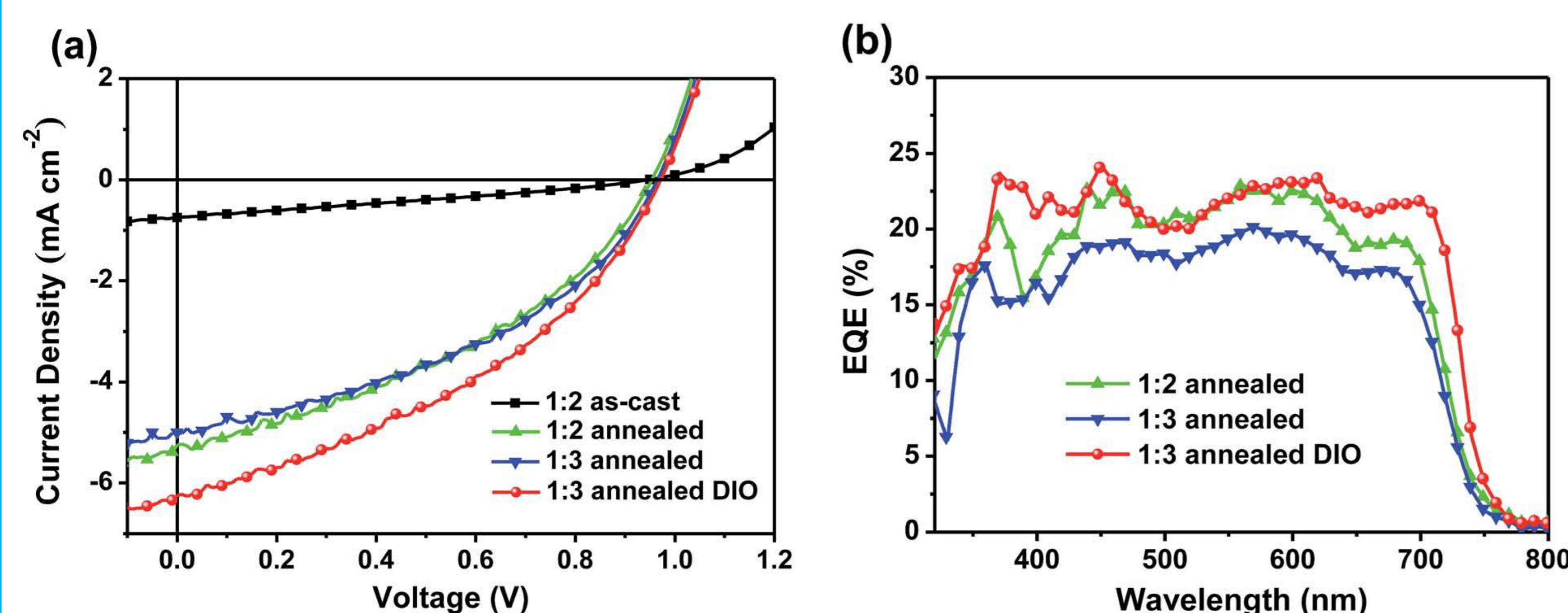


Fig. 2 (a) J - V curves and (b) EQE spectra of P3HT:F8-DPPTCN based OSCs.

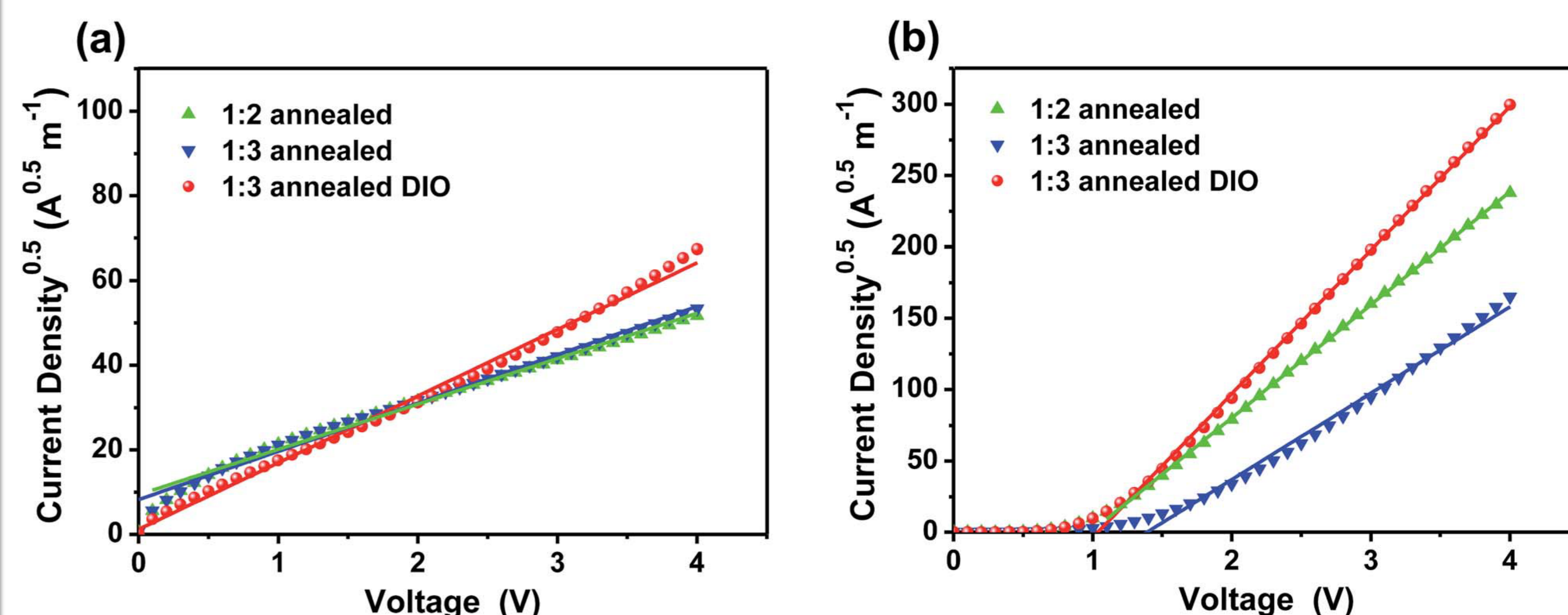


Fig. 3 (a) $J^{0.5}$ - V curves of (a) the hole-only and (b) electron-only devices based on P3HT:F8-DPPTCN blends.

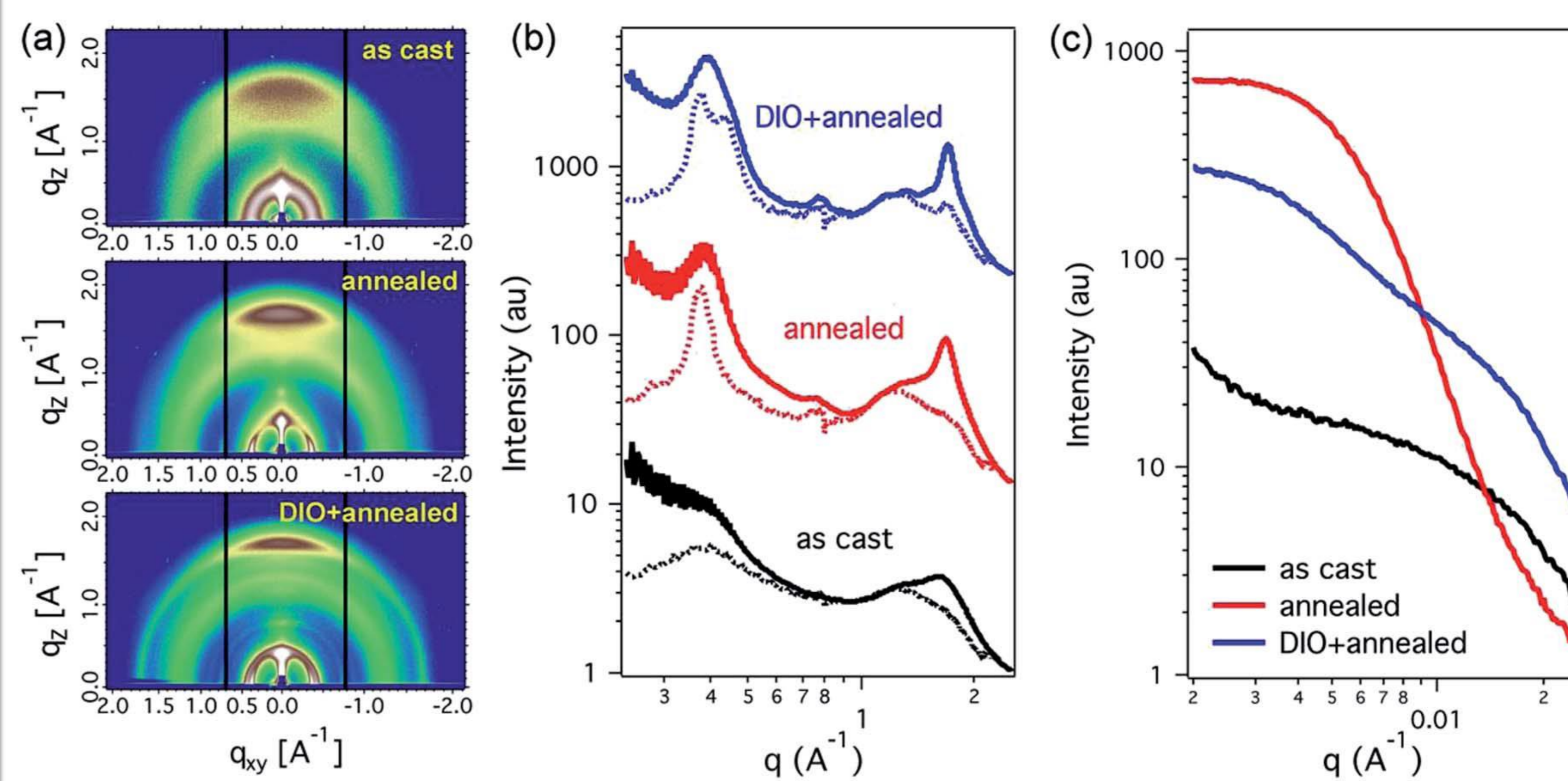


Fig. 4 (a) The 2D GIXD images of P3HT:F8-DPPTCN (1:3, by wt) blend films; (b) 1D out-of-plane (solid lines) and in-plane (dot lines) X-ray profiles extracted from GIXD; (c) RSoXS profiles of P3HT:F8-DPPTCN (1:3, by wt) blend films.

Conclusions

We have designed and synthesized an electron acceptor (F8-DPPTCN) with a fluorene core having DPP arms that are end-capped with thiophene-2-carbonitrile. The resulting material exhibits extended absorption to the near infrared region (up to 757 nm). F8-DPPTCN molecules exhibit good electron mobility (up to 10⁻³ cm²V⁻¹s⁻¹) in bulk heterojunction blends. The best performing solar cells are obtained by using an additive processing method, and the active layer shows multi-length scale morphology. In device, a PCE of 2.37%, and V_{oc} of 0.97 V, J_{sc} of 6.25 mA cm⁻², FF of 0.39 are obtained.

Notes and references

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