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Bithienopyrroledione vs. thienopyrroledione based copolymers: dramatic increase of power conversion efficiency in bulk heterojunction solar cells†

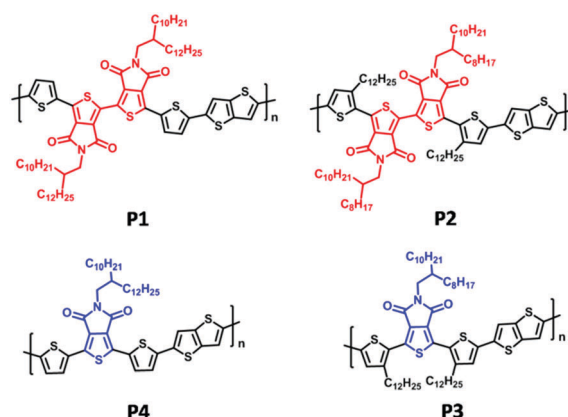
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Bithienopyrroledione (bi-TPD) based polymers P1 and P2 are designed and synthesized. Photovoltaic devices based on P1:PC₇₁BM and P2:PC₇₁BM blend films show power conversion efficiencies (PCEs) of 8.22% and 9.08%, respectively, whereas devices based on their thienopyrroledione (TPD)-based analogue P3 and PC₇₁BM blend films display a PCE of 5.10%.

Polymer solar cells (PSCs) have been intensively investigated in recent years due to their features of low cost, flexibility, and light weight.^{1–5} The development in active layer materials, interfacial materials, and device processing technologies has improved the power conversion efficiencies (PCEs) of PSCs greater than 10%.^{6–12} In view of the active layer materials, donor–acceptor (D–A) type polymers have attracted great attention due to their advantages in tuning the absorption spectra, HOMO/LUMO energy levels, intermolecular interactions and molecular orientations, and hence numerous D–A type semiconductor polymers have been synthesized and used as donors in bulk heterojunction (BHJ) structure solar cells.^{13,14} To date, D–A polymers with PCEs over 8% have rarely been reported and most of them employ benzo[1,2,5]thiadiazole and fluorinated thieno[3,4-*b*]thiophene derivatives as electron deficient units.^{15–24} Thus, the exploration of new electron-deficient units (acceptor units) is urgently required for the development of high performance

D–A polymers and the study of the relationships between polymer chemical structures and properties.

Thieno[3,4]pyrrole-4,6-dione (TPD) based polymers are widely used as electron donors in bulk heterojunction solar cells and a record PCE of 8.5% has been reported for TPD–benzodithiophene copolymerized polymers by using a classical device configuration.^{25–28} As a derivative of TPD, bithienopyrroledione (1,1'-bithieno[3,4-*c*]pyrrole-4,4',6,6'-tetraone, bi-TPD) has a good planar structure and stronger electron-withdrawing ability than TPD,^{29–31} making it a good candidate as an acceptor unit for D–A polymers. However, bi-TPD based D–A polymers have rarely been investigated in PSCs. This might be partially because of the commonly accepted view that the large internal dipole moment along the polymer chain facilitates charge separation in PSCs and thus results in high PCE, which were examined by Yu's and Tsige's groups on polymers PTB7 and PBB3.^{32,33} In this study, we designed and synthesized two bi-TPD based polymers **P1** and **P2** (Scheme 1) and investigated their application as donor materials in PSCs. The **P1** and **P2** based cells exhibit PCEs of 8.22% and 9.08% with a traditional device configuration of ITO/PEDOT:PSS/**P1** (**P2**):PC₇₁BM/Ca/Al, much higher than that of the cell of TPD based analogue **P3** (PCE: 5.10%) and the highest PCE value reported for bi-TPD



Scheme 1 Chemical structures of polymers **P1–P4**.

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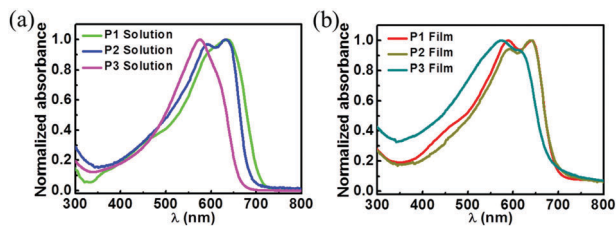


Fig. 1 UV-vis absorption spectra of **P1–P3** in dilute *o*-DCB solutions (a) and thin films (b).

contained polymers. These results demonstrate that bi-TPD based polymers have great potential for application in PSCs.

P1 and **P2** were synthesized through conventional Stille coupling of a stannyl reagent of thieno[3,2-*b*]thiophene with the corresponding 5,5'-bis[5-bromo-3-alkylthiophen]-1,1'-bithieno[3,4-*c*]pyrrole-4,4',6,6'-tetraone. For comparison, **P3**³⁴ and **P4**, the mono-TPD based analogues of **P1** and **P2**, were also synthesized using the Stille coupling reaction (Scheme S1 in the ESI†). Their number-average molecular weights (M_n) were measured using high temperature (150 °C) gel-permeation chromatography (GPC) in 1,2,4-trichlorobenzene using polystyrene as the standard. The M_n /PDI (polydispersity index) values were 82.9 kDa/2.88 for **P1**, 53.3 kDa/2.29 for **P2**, 154 kDa/4.99 for **P3** and 7.1 kDa/2.83 for **P4**. The low molecular weight of **P4** was due to its poor solubility, and solar cell devices based on **P4** were not fabricated and investigated because of its lower M_n .

The UV-vis absorption spectra of copolymers **P1–P3** were measured by a UV-vis spectrophotometer in *o*-dichlorobenzene (*o*-DCB) solution and in thin films spin-coated from *o*-DCB solution (Fig. 1). All polymers showed 0–0/0–1 peaks in solution and in thin films. For **P1**, the 0–1 peaks in solution and films were located at 588 nm, and the 0–0 peak was at 634 nm in solution and at 640 nm in films. For **P2**, similar to **P1**, the 0–1 peaks were at 592 nm in solution and films, and the 0–0 peak was shifted from 632 nm in solution to 640 nm in films. The 0–1 and 0–0 absorptions of **P3** were at 575 and 614 nm in both solution and thin films. The optical band gaps estimated from thin film absorptions were 1.80, 1.80, and 1.84 eV for **P1**, **P2**, and **P3**, respectively. The electrochemical properties of the **P1** and **P2** thin films were investigated through cyclic voltammetry (CV) (Fig. S1, ESI†). **P1** and **P2** displayed a quasi-reversible oxidation process with redox potentials located at 1.24 and 1.22 V vs. SCE, respectively, and the HOMO energy level was –5.61 eV for **P1** and –5.59 eV for **P2**. The LUMO energy levels calculated from the HOMO levels and optical band gaps were –3.81 eV for **P1** and 3.79 eV for **P2**. The HOMO/LUMO energy levels of **P1** and **P2** were much lower than that of **P3** (HOMO/LUMO: –5.07/–3.23 eV).³⁴ The energy levels of the **P1–P3** and PC₇₁BM (LUMO: –4.30 eV; HOMO: –6.00 eV) exhibited a suitable cascade stage for exciton dissociation, suggesting their potential application in polymer:PC₇₁BM blend solar cells.

The photovoltaic properties of polymers **P1–P3** were investigated using the conventional device structure of ITO/PEDOT:PSS/polymer:PC₇₁BM/Ca/Al. The current density–voltage (*J–V*) characteristics of the optimized devices are illustrated in Fig. 2a. The performance

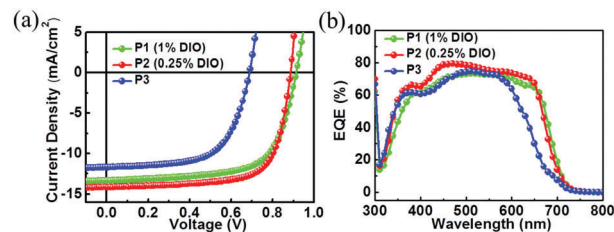


Fig. 2 *J–V* curves (a) and the corresponding EQE curves (b) of the polymer:PC₇₁BM solar cells with optimal performance.

Table 1 Photovoltaic parameters of PSCs based on polymer:PC₇₁BM

Polymers ^a	V_{oc} (V)	J_{sc} (mA cm ^{–2})	FF (%)	PCE ^b (%)
P1	0.91	13.45	67.18	8.22 (8.02 ± 0.20)
P2	0.89	14.15	72.13	9.08 (8.92 ± 0.16)
P3	0.69	11.70	63.15	5.10 (4.89 ± 0.21)

^a Blend ratios of **P1**:PC₇₁BM (w/w) were 1 : 2 with 1% DIO, **P2**:PC₇₁BM (w/w) were 1 : 2 with 0.25% DIO and **P3**:PC₇₁BM (w/w) was fixed at 1 : 1.5 without DIO. ^b In parentheses are average values based on more than 10 devices.

of the device was first optimized by using different polymer:PC₇₁BM weight ratios (Table S1, ESI†). The optimal weight ratio for bi-TPD based polymers **P1** and **P2** to PC₇₁BM was found to be 1 : 2, and that for **P3** to PC₇₁BM was 1 : 1.5. Then, the effect of the 1,8-diiodooctane (DIO) additive on the device performance was investigated (Fig. S2–S4 and Table S2, ESI†). The experimental results show that (Table 1) the cells of **P1**:PC₇₁BM with 1% DIO exhibited an optimal PCE of 8.22% with a V_{oc} of 0.91 V, a J_{sc} of 13.45 mA cm^{–2}, and a FF of 67.18%. For **P2**:PC₇₁BM based devices, a PCE up to 9.08% with a V_{oc} of 0.89 V, a J_{sc} of 14.15 mA cm^{–2}, and a FF of 72.13% was observed upon addition of 0.25% DIO. For **P3**:PC₇₁BM based devices, the optimal PCE was 5.10% with a V_{oc} of 0.69 V, a J_{sc} of 11.70 mA cm^{–2}, and a FF of 63.15%. This PCE value is comparable to the reported TPD and 2,5-di(thiophen-2-yl)thieno[3,2-*b*]thiophene (DT-TT) based polymers.³⁵ Apparently, the V_{oc} of **P1** and **P2** based solar cells was significantly larger than that of **P3** based devices. This is consistent well with the deeper HOMO energy levels of the bi-TPD based polymers. The PCEs of **P1** and **P2** based devices may be further improved through device engineering similarly to that of PTB7-Th:PC₇₁BM based solar cells.

The photo-response of the solar cells based on these polymers was investigated using their external quantum efficiencies (EQEs), and the EQE curves of the optimal devices are illustrated in Fig. 2b. The **P1**:PC₇₁BM and **P2**:PC₇₁BM devices showed a broader photo-response than the **P3**:PC₇₁BM device, consistent with their thin film absorptions. The **P2**:PC₇₁BM device exhibited the highest average EQE values among the three devices. The current density calculated from the EQE curves with a standard solar spectrum (AM 1.5G) was 13.25, 13.92, and 11.36 mA cm^{–2} for **P1–P3**, respectively, agreeing well with the ones obtained from *J–V* measurements.

To elucidate the observed performance of **P1–P3** based solar cells, we explored the morphological features of the optimal blend films using transmission electron microscopy (TEM) and



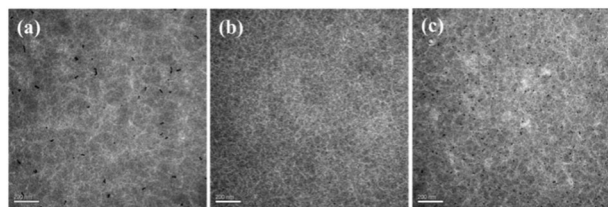


Fig. 3 TEM images of the polymer:PC₇₁BM films: (a) **P1**:PC₇₁BM; (b) **P2**:PC₇₁BM; and (c) **P3**:PC₇₁BM. The scale bars in the images are 200 nm.

atomic force microscopy (AFM). Fig. 3 illustrates the TEM images of the blend films. All the films exhibited obvious dark and light domains, corresponding to the aggregates of PC₇₁BM and the polymer, respectively. For **P1** (**P2**):PC₇₁BM films, nanofibrillar-like aggregates were formed and distributed throughout the blend films, resulting in an interpenetrating network. This kind of morphological feature is more favorable for efficient exciton separation and diffusion. **P3**:PC₇₁BM films showed similar morphological features to the other two blend films. However, some aggregated structures of **P3** (high brightness areas) were observed. AFM images revealed that **P1**:PC₇₁BM films were comprised of large size grains with an RMS value of 4.21 nm (Fig. S5, ESI[†]). Compared to **P1**:PC₇₁BM films, **P2**:PC₇₁BM films exhibited smaller size domains and smoother surfaces with an RMS value of 1.73 nm. The RMS values of **P3**:PC₇₁BM films were larger than 9 nm due to the formation of aggregated structures, which is consistent with the TEM results.

The crystallinity and molecular packing of the polymers in the active layers were probed using 2D grazing incidence X-ray diffraction (2D-GIXD) (Fig. 4). **P1–P3** were crystalline and adopted edge-on and face-on co-orientations in the blend films, as proved by the (100) diffraction peaks in both the q_{xy} -axis and the q_z -axis. Moreover, π - π stacking diffractions (010 peaks) were observed in the q_z -axis and the π - π stacking distances were calculated to be 3.52, 3.61, and 3.60 Å for **P1**, **P2** and **P3**, respectively. The crystalline nature and co-existence of the edge-on and face-on packing structures of the films facilitated charge carrier transport in polymer solar cell, leading to high photovoltaic performance.

The hole and electron transport properties of the optimal polymer:PC₇₁BM films in the direction perpendicular to the substrate were evaluated using space-charge-limited-current (SCLC) measurements. The J - V curves are shown in Fig. S6 (ESI[†]). For **P1**:PC₇₁BM films, the calculated electron (μ_e) and hole (μ_h) mobilities were $4.48 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $1.99 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ with a

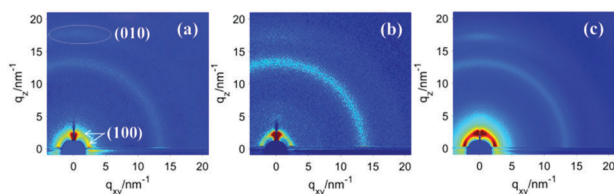


Fig. 4 2D-GIXD images of the polymer:PC₇₁BM films: (a) **P1**:PC₇₁BM; (b) **P2**:PC₇₁BM; and (c) **P3**:PC₇₁BM.

μ_e/μ_h ratio of 2.25. For **P2**:PC₇₁BM films, the calculated μ_e and μ_h were $3.86 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $2.49 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ with a μ_e/μ_h ratio of 1.55. For **P3**:PC₇₁BM films, the calculated μ_e and μ_h were $3.57 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $1.35 \times 10^{-4} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ with a μ_h/μ_e ratio of 2.64. The relatively balanced carrier mobility of **P2**:PC₇₁BM might be an important factor for its higher FF in solar cells.³⁶

In conclusion, new bi-TPD based polymers **P1** and **P2** were designed and synthesized. Their physicochemical properties were investigated. Compared to their TPD-based analogue **P3**, **P1** and **P2** have deeper HOMO energy levels and exhibit red-shifted thin film UV-vis absorption spectra. The photovoltaic characteristics showed that cells based on bi-TPD based polymers **P1** and **P2** exhibited much higher PCE than that of mono-TPD based polymer **P3**. The optimal PCE is 8.22% for **P1** cells, 9.08% for **P2** cells and 5.10% for **P3** cells. The PCE and FF values of **P2** cells are among the highest values reported for single junction solar cells. The blend films of **P1** (**P2**):PC₇₁BM exhibited high crystallinity with face-on and edge-on co-orientations and good phase separation. These results demonstrate that bi-TPD is an excellent electron-deficient unit that is rarely investigated for high PCE polymer semiconductors. Through further material optimization in the side chains and donor units as well as device engineering, we believe, bi-TPD based polymers with higher efficiencies can be envisioned.

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Notes and references

- 1 L. Dou, Y. Liu, Z. Hong, G. Li and Y. Yang, *Chem. Rev.*, 2015, **115**, 12633.
- 2 L. Lu, T. Zheng, Q. Wu, A. M. Schneider, D. Zhao and L. Yu, *Chem. Rev.*, 2015, **115**, 12666.
- 3 F. C. Krebs, N. Espinosa, M. Hösel, R. R. Søndergaard and M. Jørgensen, *Adv. Mater.*, 2014, **26**, 29.
- 4 P. Cheng and X. Zhan, *Chem. Soc. Rev.*, 2016, **45**, 2544.
- 5 Y. J. Cheng, S. H. Yang and C. S. Hsu, *Chem. Rev.*, 2009, **109**, 5868.
- 6 Y. Liu, J. Zhao, Z. Li, C. Mu, W. Ma, H. Hu, K. Jiang, H. Lin, H. Ade and H. Yan, *Nat. Commun.*, 2014, **5**, 5293.
- 7 W. Zhao, D. Qian, S. Zhang, S. Li, O. Inganäs, F. Gao and J. Hui, *Adv. Mater.*, 2016, **28**, 4734.
- 8 H. Bin, L. Gao, Z. G. Zhang, Y. Yang, Y. Zhang, C. Zhang, S. Chen, L. Xue, C. Yang, M. Xiao and Y. Li, *Nat. Commun.*, 2016, **7**, 13651.
- 9 J. Chen, C. Cui, Y. Q. Li, L. Zhou, Q. D. Ou, C. Li, Y. Li and J. Tang, *Adv. Mater.*, 2015, **27**, 1035.
- 10 X. H. Ouyang, R. X. Peng, L. Ai, X. Y. Zhang and Z. Y. Ge, *Nat. Photonics*, 2015, **9**, 520.
- 11 J. Zhang, Y. Zhang, J. Fang, K. Lu, Z. Wang, W. Ma and Z. Wei, *J. Am. Chem. Soc.*, 2015, **137**, 8176.
- 12 Y. Liu, Z. A. Page, T. P. Russell and T. Emrick, *Angew. Chem., Int. Ed.*, 2015, **54**, 11485.
- 13 Y. Li, *Acc. Chem. Res.*, 2012, **45**, 723.
- 14 J. W. Chen and Y. Cao, *Acc. Chem. Res.*, 2009, **42**, 1709.
- 15 M. Wang, D. Cai, Z. Yin, S. C. Chen, C. F. Du and Q. Zheng, *Adv. Mater.*, 2016, **28**, 3359.
- 16 K. Kawashima, T. Fukuhara, Y. Suda, Y. Suzuki, T. Koganezawa, H. Yoshida, H. Ohkita, I. Osaka and K. Takimiya, *J. Am. Chem. Soc.*, 2016, **138**, 10265.

- 17 H. Bin, Z. G. Zhang, L. Gao, S. Chen, L. Zhong, L. Xue, C. Yang and Y. Li, *J. Am. Chem. Soc.*, 2016, **138**, 4657.
- 18 R. S. Ashraf, I. Meager, M. Nikolka, M. Kirkus, M. Planells, B. C. Schroeder, S. Holliday, M. Hurhangee, C. B. Nielsen, H. Sirringhaus and I. McCulloch, *J. Am. Chem. Soc.*, 2015, **137**, 1314.
- 19 Y. Deng, J. Liu, J. Wang, L. Liu, W. Li, H. Tian, X. Zhang, Z. Xie, Y. Geng and F. Wang, *Adv. Mater.*, 2014, **26**, 471.
- 20 W. Chen, M. Xiao, L. Han, J. Zhang, H. Jiang, C. Gu, W. Shen and R. Yang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 19665.
- 21 M. Zhang, Y. Gu, X. Guo, F. Liu, S. Zhang, L. Huo, T. P. Russell and J. Hou, *Adv. Mater.*, 2013, **25**, 4944.
- 22 J. You, L. Dou, K. Yoshimura, T. Kato, K. Ohya, T. Moriarty, K. Emery, C. C. Chen, J. Gao, G. Li and Y. Yang, *Nat. Commun.*, 2013, **4**, 1446.
- 23 Z. He, B. Xiao, F. Liu, H. Wu, Y. Yang, S. Xiao, C. Wang, T. P. Russell and Y. Cao, *Nat. Photonics*, 2015, **9**, 174.
- 24 M. Zhang, X. Guo, S. Zhang and J. Hou, *Adv. Mater.*, 2014, **26**, 1118.
- 25 Y. Zou, A. Najari, P. Berrouard, S. Beaupré, B. R. Aich, Y. Tao and M. Leclerc, *J. Am. Chem. Soc.*, 2010, **132**, 5330.
- 26 E. Zhou, J. Cong, K. Tajima, C. Yang and K. Hashimoto, *Macromol. Chem. Phys.*, 2011, **212**, 305.
- 27 E. Zhou, J. Cong, K. Tajima, C. Yang and K. Hashimoto, *J. Phys. Chem. C*, 2012, **116**, 2608.
- 28 J. W. Jung, J. W. Jo, E. H. Jung and W. H. Jo, *Org. Electron.*, 2016, **31**, 149.
- 29 P. Berrouard, F. Grenier, J. R. Pouliot, E. Gagnon, C. Tessier and M. Leclerc, *Org. Lett.*, 2011, **13**, 38.
- 30 J. E. Donaghey, R. S. Ashraf, Y. Kim, Z. G. Huang, C. B. Nielsen, W. Zhang, B. Schroeder, C. R. G. Grenier, C. T. Brown, P. D'Angelo, J. Smith, S. Watkins, K. Song, T. D. Anthopoulos, J. R. Durrant, C. K. Williamsa and I. McCullocha, *J. Mater. Chem.*, 2011, **21**, 18744.
- 31 X. Qiao, Q. Wu, H. Wu, D. Wang and H. Li, *Polym. Chem.*, 2016, **7**, 807.
- 32 B. Carsten, J. M. Szarko, H. J. Son, W. Wang, L. Lu, F. He, B. S. Rolczynski, S. J. Lou, L. X. Chen and L. Yu, *J. Am. Chem. Soc.*, 2011, **133**, 20468.
- 33 R. S. Bhatta and M. Tsige, *ACS Appl. Mater. Interfaces*, 2014, **6**, 15889.
- 34 Q. Wu, M. Wang, X. Qiao, Y. Xiong, Y. Huang, X. Gao and H. Li, *Macromolecules*, 2013, **46**, 3887.
- 35 G. Y. Chen, Y. H. Cheng, Y. J. Chou, M. S. Su, C. M. Chen and K. H. Wei, *Chem. Commun.*, 2011, **47**, 5064.
- 36 G. Li, V. Shrotriya, J. S. Huang, Y. Yao, T. Moriarty, K. Emery and Y. Yang, *Nat. Mater.*, 2005, **4**, 864.

