Dramatically different charge transport properties of bisthienyl diketopyrrolopyrrole-bithiazole copolymers synthesized via two direct (hetero)-arylation polymerization routes†

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Two bisthienyl diketopyrrolopyrrole (DPP)-bithiazole based copolymers were synthesised via two different direct (hetero)arylation polymerization (DHAP) routes. When a bisthienyl DPP and 5,5'-dibromo-2,2'-bithiazole were used as monomers, the resulting polymer PA-1 showed poor solubility, ill-organized chain ordering and low performance in organic thin film transistors (OTFTs). Surprisingly, the synthetic route using a dibrominated bisthienyl DPP and 2,2'-bithiazole as monomers produced a polymer PB-1 with improved solubility, much higher crystallinity and excellent charge transport performance in OTFTs. The electron and hole mobilities of up to 0.53 cm² V⁻¹ s⁻¹ and 0.06 cm² V⁻¹ s⁻¹, respectively, achieved for PB-1 are more than one order of magnitude higher than those of PA-1 and also better than the mobilities reported for a similar polymer synthesized via Stille coupling polymerization. The dramatically different carrier mobilities observed for these two polymers are accounted for by their different amounts of α-β coupling linkages and branched (and lightly cross-linked) structure defects formed in the respective synthetic routes. This work also demonstrated for the first time that 2,2'-bithiazole is a suitable monomer for the construction of conjugated polymers with good electron transport performance via DHAP.

Introduction

Polymer semiconductors have been widely studied as active materials for solution processable organic electronics such as organic thin film transistors (OTFTs),¹⁻⁵ organic photovoltaics (OPVs)⁶⁻⁹ and chemical sensors.¹⁰⁻¹¹ Compared to small molecules, polymers have better mechanical, thermal and thin-film forming properties, allowing for the fabrication of low-cost, large-area and flexible electronics using various printing technologies such as screen, inkjet or gravure printing.¹⁴,¹⁵ To enhance carrier mobility, electron-donating and electron-accepting building blocks can be combined in the repeat units to form donor–acceptor (D–A) polymers to shorten the π-stacking distance for more efficient charge hopping.¹¹,¹⁶,¹⁷ Diketopyrrolopyrrole (DPP) has been widely used as an electron-acceptor building block for the development of high-performance polymeric semiconductors for OTFTs and OPVs.⁴,¹⁷⁻²³ The majority of the DPP-based polymers reported in the literature were synthesized using either Suzuki or Stille coupling polymerization, which requires tedious steps to synthesize the organoboron or organotin monomers. Many organonit compounds are known to be highly toxic and are environmental hazards. Recently, a new alternative method, direct (hetero)arylation polymerization (DHAP),²⁴⁻²⁷ has been explored to construct conjugated polymers useful for organic electronics.²⁸⁻³¹ Since DHAP involves the C–C bond formation through a coupling reaction between a C–H bond in one conjugated monomer (an arene or heteroarene) and a C–X bond (X is usually a halide, Br or I) in another, this new method is much more environmentally friendly and cost effective than the Suzuki and Stille coupling methods.

Compared to thiophene, which is a commonly used electron donor building block, thiazole is a weak acceptor³⁴ and has been utilized to promote the electron transport performance of polymer semiconductors.³⁵,³⁶ Previously, DPP-quinethiophene based conjugating polymers (PDQT)¹⁹,³⁷,³⁸ were reported to show p-type hole transport performance with high mobility up to 6.9 cm² V⁻¹ s⁻¹.³⁷ If a bithiophene unit in PDQT is replaced by a 2,2'-bithiazole or 5,5'-bithiazole unit (Fig. 1), the resulting polymer PDBTz or PDBTz’ is expected to show electron transport behaviour. Very recently, a PDBTz polymer, PDBTz-27 (R = 5-decylheptadecyl) synthesized via Stille coupling polymerization was reported by Reichmanis...
and coworkers,\textsuperscript{39} which showed high electron mobility of up to 0.3 cm\(^2\) V\(^{-1}\) s\(^{-1}\) in OTFT devices. In this study, we used the DHAP method through two different routes to prepare two PDBTz polymers, PA-1 and PB-1 (R = 2-decyltetradecyl). The syntheses of these polymers reported here involved fewer steps than the Stille coupling method. It was found that PB-1 showed much higher electron mobility of up to 0.53 cm\(^2\) V\(^{-1}\) s\(^{-1}\) in OTFT devices due to its less structural defects formed during the polymerization compared to PA-1.

**Results and discussion**

The synthetic routes to polymers PA-1 and PB-1 are outlined in Scheme 1. Stannylation of 2-bromothiazole resulted in compound 1, which was coupled with 2-bromothiazole to afford compound 2 in 78% yield. Compound 2 was then brominated at the 5,5'-positions using N-bromosuccinimide (NBS) to form compound 3 in 98% yield. The bithiazole compounds 2 and 3 are monomers for the following DHAP polymerization reactions. The DPP-containing monomers, compounds 4-1 and 5-1, were prepared according to the literature methods.\textsuperscript{37} The target polymers PA-1 and PB-1 could be synthesized via DHAP through two routes: (A) coupling between 3 and 4-1 to afford PA-1 and (B) coupling between 2 and 5-1 to afford PB-1.

The \(\alpha\) C–H bond (at the C2 or C5 position) of thiophene is known to have higher activity toward a C–X (Br or I) bond through DHAP than the \(\beta\) C–H bond (at the C3 or C4 position), allowing for synthesis of soluble polymers with minimal cross-linked structural defects under certain controlled conditions.\textsuperscript{31,33,40–43} Monomer 4-1 in Route A has two \(\alpha\) C–H bonds and four \(\beta\) C–H bonds in two thiophene units. If DHAP occurs predominantly at the desired \(\alpha\) C–H bonds, a linear polymer PA-1 could be obtained. The regioselectivity of the \(\alpha\) C–H bond (at C2 or C5) over the \(\beta\) C–H bond (at C4) in thiazole has also been reported to be very high in DHA reactions for small molecule syntheses.\textsuperscript{44–53} A bithiazole compound with the C4 positions substituted (blocked) with nonyl side chains was used for DHAP to make a soluble conjugated polymer without observable cross-linking side reactions.\textsuperscript{45} However, to the best of our knowledge, the use of a bithiazole compound containing both \(\alpha\) and \(\beta\) C–H bonds as a monomer for DHAP has not been demonstrated yet. Therefore, it is interesting to explore if the use of a \(\beta\) non-substituted bithiazole compound 2 as a monomer in Route B can form a soluble target polymer PB-1.

Two polymerization reactions were conducted under identical conditions, in the presence of a catalyst system containing \(\text{trans-bis(acetato)bis[O-di(di-tolylyphosphino)benzyl]dipalladium (n)}\) (Herrmann–Beller’s catalyst), tris(o-methoxyphenyl)phosphine, cesium carbonate and pivalic acid under reflux in toluene for 16 h, which were successfully used for the preparation of other conjugated polymers through DHAP.\textsuperscript{29–32,55} The crude polymers were purified by Soxhlet extraction using acetone, hexane, chloroform and 1,1,2,2-tetrachloroethane (TCE) sequentially. Negligible amounts of polymers were dissolved by acetone and hexane. For PA-1, 9% was dissolved by chloroform and 53% by TCE. The remaining (\(\sim\)38%) solid was insoluble in common known solvents. PB-1 showed much improved solubility with 66% dissolved in chloroform and 10% dissolved in TCE, but \(\sim\)24% of this polymer still remained insoluble (Scheme 1). High temperature gel permeation chromatography (HT-GPC) with 1,2,4-trichlorobenzene as an eluent and polystyrene as standards at a column temperature of 140 °C was used to determine the molecular weights of PA-1 and PB-1 due to the strong chain aggregation tendency at room temperature. The number-average molecular weight (\(M_n\))/polydispersity index (PDI) of PA-1 are 17 kDa/6.9 for the chloroform extracted fraction and 11 kDa/5.9 for the TCE extracted fraction. \(M_n/PDI\) of PB-1 are 18 kDa/3.8 for the chloroform extracted fraction and 23 kDa/3.9 for the TCE extracted fraction (Scheme 1).

As aforementioned, one immensely important issue associated with DHAP is that side reactions of C–X (X = Br or I) bonds with the undesirable (hetero)aromatic C–H bonds, such as via the strong chain aggregation tendency at room temperature.
as the $\beta$ C–H at the C3 or C4 position in thiophene,\textsuperscript{56} can occur, leading to formation of cross-linked, insoluble polymers and low molecular weight soluble fractions.\textsuperscript{32,40,54,57–59} Therefore, a possible reason for the presence of a significant amount of insoluble fractions for both polymers (PA-1: \~38%; PB-1: \~24%) is due to the formation of cross-linked structures via the side reactions between the $\alpha$ C–Br and the $\beta$ C–H bonds on thiophene and/or thiazole units. PB-1 showed much better solubility and narrower molecular weight distributions for its soluble fractions in chloroform and TCE compared with PA-1, indicating that less $\alpha$–$\beta$ coupling side reactions might have occurred in Route B.

$^1$H-NMR spectroscopy has been used to determine the presence of branched structures formed via $\alpha$–$\beta$ coupling side reactions of thiophene units.\textsuperscript{38,40,60,61} However, the $^1$H NMR spectra of these two polymers acquired at room temperature showed extremely poor resolutions due to the aggregation of polymer chains, which has been reported for some other D–A polymers.\textsuperscript{19,62} To improve the spectral resolution, the NMR measurements were conducted at an elevated temperature of 120 $^\circ$C. Three major peaks at 7.43, 8.09 and 8.82 ppm are clearly seen for both polymers, which can be assigned to $H_a$ and $H_c$ in the thiophene units and $H_b$ in the thiazole units (Fig. 2). However, several minor peaks, which appear at different positions for PA-1 and PB-1, are also observed. These may be originated from their different terminal monomer units and $\alpha$–$\beta$ coupled units. However, it is very difficult to assign the peaks that belong to the $\alpha$–$\beta$ coupling structures.

Another possible reason for the large amounts of insoluble products in PA-1 and PB-1 is due to the poor solubility of their high molecular weight fractions. To confirm if the formation of insoluble fractions was caused by cross-linking or the high molecular weights, two model polymers PA-2 and PB-2 with much larger tritriacontan-17-yl butyrate side chains\textsuperscript{63} were synthesized under the same reaction conditions used for PA-1 and PB-1 (Scheme 1). Most of PA-2 (81%) and PB-2 (89%) could be dissolved with hexane by Soxhlet extraction, indicating the greatly improved solubility of these two polymers. However, there were still some remaining products, which could not be dissolved by any solvent, for both polymers (PA-2: \~19%; PB-2: \~11%). Since the solubility of these soluble fractions is excellent (soluble in hexane at room temperature) due to their large side chains, the presence of insoluble products is considered to be caused by the cross-linking of the polymer chains by the side reactions rather than by their high molecular weights. The soluble fractions of these two polymers were subject to the GPC measurements. $M_n$/PDI values of 15 kDa/4.3 for PA-2 and 11 kDa/2.5 for PB-2 were obtained, which are comparable to the values for PA-1 and PB-1, even though the solubility of these two model polymers is much improved. This could be explained by the premature termination of the polymer chain growth during the polymerization. Since both polymers contain appreciable amounts of insoluble fractions, it is reasonable to consider that cross-linking occurred during the polymerization, which resulted in the formation of some insoluble polymer chains containing heavily cross-linked struc-

![Fig. 2](https://example.com/figure2.png)  
Fig. 2  The $^1$H-NMR spectra of polymers PA-1, PB-1, PA-2, PB-2, and monomers 2, 3, 4-1 and 5-1. The measurements were carried out at 120 $^\circ$C for the polymers and at room temperature for the monomers.
tures. Due to the excessive consumption of the C–Br groups by the cross-linked structures through α–β coupling, the remaining polymer chains did not have functional C–Br groups for further growth and consequently led to their smaller molecular weights. Because the solubility of PA-2 and PB-2 is much improved, their 1H-NMR spectra were measured in order to elucidate the defective structures formed by the side reactions. However, the 1H-NMR spectra of PA-2 and PB-2 are very similar to those of their corresponding polymers, PA-1 and PB-1 (Fig. 2), although the peaks of PA-2 and PB-2 are sharper as a result of their improved solubility. The major peaks, a, b, and c, are assigned as shown in Fig. 2 according to the NMR spectra of their corresponding monomers. The minor peaks at 7.3 ppm and 7.7 ppm for PA-1 and PA-2 are positioned similarly to those of the peaks of a’ and d’ of 4-1 and 4-2 (not shown), suggesting that 4-1 (or 4-2) is a terminal unit in PA-1 (or PA-2). On the other hand, there is only a very small peak at ~7.8 ppm that is close to the chemical shift of peak b’ of 3, indicating that the amount of 3 as a terminal unit in PA-1 and PA-2 is minimal. For PB-1 and PB-2, the peak at 7.9 ppm could be assigned to the β-H b″ of the terminal 2 unit, while the α-H of the terminal 2 unit might be hidden in peak a. On the contrary, there is no peak that corresponds to c′ of monomer 5-1, suggesting that 5-1 is unlikely a terminal unit in PB-1 and PB-2. The above observations indicate that 4-1 and 4-2 are the dominant terminal units for PA-1 and PA-2, respectively, while 2 is the dominant terminal unit for PB-1 and PB-2. The lack of the dibromo monomer terminal units (3, 5-1 or 5-2) substantiates that some of the C–Br bonds reacted with β-H’s. The α-H in an α–β coupling defect could further react with a C–Br group, consequently forming branched and cross-linked structures (see Scheme S1 in the ESI†). It should be noted that the major peaks, a, b, and c, are comprised of multiple peaks and/ or minor peaks, which might be originated from the sub-terminal units and the branched (or lightly cross-linked) structures formed by α–β coupling. Therefore, we may conclude that both polymerization routes (A and B) likely involved the α–β coupling reactions that led to the formation of cross-linked structures, whereas the polymers produced in Route A have more cross-linked structures than the polymers made by Route B based on the amounts of their insoluble fractions. Polymer chains in the soluble fractions may also contain structural defects such as branches and lightly cross-linked structures.

The branched and lightly cross-linked structures would cause twisting of the polymer backbone, while the α–β coupling linkages in the polymer backbone would disrupt the π-conjugation. As a result, the optical band gap of the polymer with more structural defects would be larger. To confirm this, UV-Vis absorption spectra of these polymers were measured in their dilute chloroform solutions and thin films. The wavelength of maximum absorbance ($\lambda_{\text{max}}$) of PA-1 was observed at 660 nm in solution and 746 nm in the thin film (Fig. 3a). PB-1 showed a longer $\lambda_{\text{max}}$ at 677 nm in solution. The as-cast film of PB-1 exhibited a $\lambda_{\text{max}}$ at 737 nm, which is shorter than that of the PA-1 film. However, the right side of the absorption spectrum of the PB-1 film extends farther into the near infrared region than the PA-1 film. As a result, the optical band gap of PB-1 (1.42 eV) is narrower than that of PA-1 (1.49 eV) calculated using the onset absorption wavelengths of their films. The larger optical band gap of PA-1 suggests its shorter effective conjugation length than that of PB-1. Because the $M_n$’s of PA-1 and PB-1 extracted with chloroform are very similar (17 kDa vs. 18 kDa) and the weight-average molecular weight ($M_w$) of PA-1 (117 kDa) is much larger than that of PB-1 (70 kDa), the effective conjugation length of PA-1 should be similar or larger than that of PB-1 if both polymers have linear main chain structures. Therefore, the blue-shift of the absorption spectrum of PA-1 seems to support the assumption that PA-1 contains more α–β coupling defects in the main chain and/or branched (and lightly cross-linked) structures, which disrupted the main chain conjugation. Due to the influence of the large side chains, the $\lambda_{\text{max}}$’s of PA-2 in solution (617 nm) and in the film (665 nm) (Fig. S8†) are both shorter than those of PA-1. The same trend is observed for PB-2, which showed shorter $\lambda_{\text{max}}$’s of 650 nm in solution and 685 nm in the film compared to PB-1. However, it is obvious that the spectra of PA-2 showed significant blue shifts in both solution and film with respect to those of PB-2, similar to the trend observed for PA-1 and PB-1. The optical band gap of PA-2 (1.54 eV) is thus wider than

![Image](https://example.com/image.png)
that of PB-2 (1.50 eV), indicating again the presence of more α-β coupling defects and/or branched (and lightly cross-linked) structures in PA-2 obtained by Route B.

Cyclic voltammetry (CV) was used to determine the frontier energy levels of PA-1 and PB-1 films (Fig. 3b). By using the onset oxidative/reductive potentials, the HOMO/LUMO levels are calculated to be −5.55 eV/−3.61 eV for PA-1 and −5.56 eV/−3.63 eV for PB-1, respectively. The energy levels of PA-1 and PB-1 are very similar and also close to those of the DPP-bithiazole copolymer PDQT-24 (−5.53 eV) due to the presence of the electron-deficient bithiazole units.34

The atomic force microscopy (AFM) images (Fig. S9†) of PA-1 and PB-1 showed that all the polymer thin films are very smooth and did not undergo dramatic morphological changes after annealed at temperatures ranging from 100 °C to 250 °C. The crystallinity of the polymer thin films was investigated by reflection X-ray diffraction measurement (XRD). The PA-1 film annealed at 100 °C showed no obvious diffraction peaks, indicating its very poor crystallinity (Fig. 4a). When the annealing temperature was increased to 150 °C, a weak primary (100) peak appeared at 2θ = 3.85°, which corresponds to a d-spacing distance of 2.29 nm. By increasing the annealing temperature to 200 °C, the crystallinity of the PA-1 film improved notably, as evidenced by the much intensiﬁed peak at 2θ = 3.94° (d-spacing = 2.24 nm). Further increasing the annealing temperature to 250 °C, the intensity of the primary peak and the d-spacing remained almost the same. As shown in Fig. 4b, the crystallinity of the PB-1 film improved notably, as evidenced by the much intensiﬁed peak at 2θ = 3.93°, which corresponds to a much shorter d-spacing distance of 2.25 nm. Upon annealing at higher temperatures of 200 °C and 250 °C, a continued increase in the intensity of the primary peak as well as a shift of this peak to 2θ = 3.99° (d-spacing = 2.21 nm) and 4.05° (d-spacing = 2.18 nm) was observed. A noticeable secondary peak also appeared. This manifests that the polymer chains were packed much more orderly and compactly with the aid of thermal annealing at high temperatures. The significantly higher crystallinity of the PB-1 films than that of the PA-1 films annealed at the same temperatures most likely resulted from the presence of less α-β coupling linkages and branched (and lightly cross-linked) structural defects in the former as discussed previously. Since no (010) peaks can be seen, the polymer chains of PA-1 and PB-1 in the spin-coated thin ﬁlm samples presumably adopted a layer-by-layer lamellar packing motif with an edge-on orientation, which has been observed for other crystalline conjugated polymers.64,65

To elucidate the in-plane chain packing motif, we measured the 250 °C-annealed PA-1 and PB-1 ﬂakes using transmission XRD (Fig. 4c). Both polymers showed a broad peak around 2θ = −20° (d-spacing = 0.44 nm), which represents the typical van der Waals distance for an amorphous polymer phase. The 250 °C-annealed PB-1 ﬂakes showed a distinct (010) peak at 24.6°, which corresponds to a π–π stacking distance of 0.36 nm. PA-1, on the other hand, exhibited a small hump at the similar position, suggesting that the polymer main chains are much less orderly packed along the π–π stacking direction in this polymer. Again, the presence of more α-β coupling linkages and branched (and lightly cross-linked) structural defects in PA-1 would sterically hinder the π–π stacking of the polymer main chains. It is noticed that the π–π stacking distance of these polymers is smaller than that of PDQT-24 (0.39 nm), indicating the stronger intermolecular interaction in the PDBTz polymers.
Based on the bithiazole units. In the n-channel operation mode, devices (Table 1), which is a result of the presence of electron-deficient more pronounced electron transport over hole transport 3.7 × 10\(^{-3}\) and PB-1

Annealing data. Polymer thin films annealed at a higher temperature of 300 °C were also tested, but the mobility values dropped. 100 3.3 × 10\(^{-3}\) (6.7 × 10\(^{-4}\)) 100 3.3 × 10\(^{-3}\) 100 1.6 × 10\(^{-2}\) (1.2 × 10\(^{-3}\))

PB-1 100 7.9 × 10\(^{-2}\) (1.1 × 10\(^{-2}\)) 250 7.9 × 10\(^{-2}\) (2.3 × 10\(^{-2}\)) 0.42 (0.11) 0.53

Table 1 Performance of OTFT devices using PA-1 and PB-1 annealed at different temperatures

To evaluate the charge transport performance of PA-1 and PB-1, we used these two polymers as channel semiconductors in top-gate, bottom-contact OTFT devices. The Au source and drain electrodes were deposited on a SiO\(_2\)/Si wafer substrate by a common photolithography method. A solution of PA-1 or PB-1 in chloroform (10 mg mL\(^{-1}\)) was spin-coated on the substrate to obtain a polymer thin film (∼40 nm), which was annealed at 100 °C, 150 °C, 200 °C or 250 °C for 15 min on a hotplate in nitrogen. Then the gate dielectric layer (∼570 nm) was formed by spin-coating a Cytop (a fluoropolymer) solution at 2000 rpm. After baking on a hotplate at 100 °C for 1 h in nitrogen, a ∼70 nm Al layer was deposited by thermal evaporation as the gate electrode. The devices were characterized in air in the absence of light. In contrast with PDQT that showed unipolar hole transport performance, 19,37,38 all devices of PA-1 and PB-1 exhibited ambipolar charge transport behaviour with more pronounced electron transport over hole transport (Table 1), which is a result of the presence of electron-deficient bithiazole units. In the n-channel operation mode, devices based on the PA-1 thin films showed electron mobility of up to 3.7 × 10\(^{-3}\) cm\(^2\) V\(^{-1}\) s\(^{-1}\) for the 100 °C-annealed films. The mobility improved as the annealing temperature increased. The highest electron mobility of 2.4 × 10\(^{-2}\) cm\(^2\) V\(^{-1}\) s\(^{-1}\) was achieved for a 250 °C-annealed film. On the other hand, PB-1 showed much superior electron transport performance. The average electron mobility increased from 1.6 × 10\(^{-2}\) cm\(^2\) V\(^{-1}\) s\(^{-1}\) for the 100 °C-annealed films to 0.42 cm\(^2\) V\(^{-1}\) s\(^{-1}\) for the 250 °C-annealed films. The maximum electron mobility of ∼0.53 cm\(^2\) V\(^{-1}\) s\(^{-1}\) was achieved for a 250 °C-annealed PB-1 film (Fig. 5). In the p-channel operation mode, both polymers showed hole transport characteristics, but their hole mobilities are about one order of magnitude lower than their respective electron mobilities at the same annealing temperatures. The best hole mobilities are 2.4 × 10\(^{-3}\) cm\(^2\) V\(^{-1}\) s\(^{-1}\) for PA-1 and 5.9 × 10\(^{-2}\) cm\(^2\) V\(^{-1}\) s\(^{-1}\) for PB-1, both achieved for the films annealed at 250 °C. The significantly lower charge transport performance shown by PA-1 is considered primarily due to the presence of a larger amount of irregular α-β coupling linkages and branched (and lightly cross-linked) structures, which reduced the main chain conjugation length and resulted in disordered chain ordering as revealed by the UV-Vis and XRD data. Polymer thin films annealed at a higher temperature of 300 °C were also tested, but the mobility values dropped.

Because both PA-1 and PB-1 showed very good thermal stability with a 5% weight loss at 395 °C and 384 °C, respectively (see the thermogravimetric analysis (TGA) thermograms in Fig. S16†), the drop in mobility for the 300 °C-annealed polymer films is presumably due to the deteriorated semiconductor/dielectric or semiconductor/electrode interface. Devices based on the PA-2 and PB-2 thin films showed electron mobilities of up to 1.7 × 10\(^{-4}\) and 3.3 × 10\(^{-3}\) cm\(^2\) V\(^{-1}\) s\(^{-1}\), respectively, for the 250 °C-annealed films. Negligible hole transport performance was observed. The much lower mobili-
ties compared to PA-1 and PB-1 are due to the adverse effect of the large side chains on the chain packing. Nonetheless, the charge transport performance of PA-2 is still significantly lower than that of PB-2, since Route A produced more irregular coupling linkages.

Experimental

Materials and characterization

All chemicals were purchased from Sigma Aldrich and other used without further purification. Cytos (a perfluorinated amorphous resin) was purchased from AGC Chemicals. Compounds 4-1 and 5-1 were prepared according to the literature methods. NMR spectra were recorded on a Bruker DPX 300 MHz spectrometer with chemical shifts relative to tetramethylsilane (TMS, 0 ppm). UV-Vis spectra were collected by a Thermo Scientific GENESYS-20 spectrophotometer. Cyclic voltammetry (CV) data were obtained with a CHI660E electrochemical analyser using an Ag/AgCl reference electrode, a Pt wire counter electrode, and a Pt foil working electrode in 0.1 M tetrabutylammonium hexafluorophosphate in acetonitrile at a sweep rate of 50 mV s⁻¹. Ferrocene was used as a reference, which has a highest occupied molecular orbital (HOMO) of −4.8 eV. Atomic force microscopy (AFM) images of polymer thin films spin coated on SiO₂/Si substrates were recorded using a Dimension 3100 scanning probe microscope. After the AFM measurements, the reflection X-ray diffraction (XRD) patterns of the polymer samples were characterized using a Bruker D8 Advance powder diffractometer with a standard Bragg–Brentano geometry using Cu Kα radiation (λ = 1.5406 Å). Transmission XRD measurements were conducted on a Bruker Smart 6000 CCD 3-circle D8 diffractometer with a Cu RA (Rigaku) X-ray source (λ = 1.5406 Å) using polymer flakes stacked between two Mylar substrates. Gel-permeation chromatography (GPC) measurements of polymers were performed on a Malvern HT-GPC using 1,2,4-trichlorobenzene as an eluent and poly styrene as standards at 140 °C. The thermogravimetric analysis (TGA) was conducted on a TGA Q500 Thermogravimetric Analyzer (TA Instruments) at a heating rate of 10 °C min⁻¹ under nitrogen.

OTFTs fabrication and characterization

A top-gate, bottom-contact OTFT structure with heavily n-doped Si/SiO₂ wafer as the substrate was used to evaluate the charge transport performance of PA-1 and PB-1. The source/ drain electrode pairs were deposited using the conventional photolithography method to define the device dimensions with a channel length (L) of 30 μm and a channel width (W) of 1 mm. The substrate was cleaned using acetone and then isopropanol in an ultrasonic bath prior to use. A polymer film with a thickness of ~40 nm was deposited on the substrate by spin coating a polymer solution in chloroform (10 mg mL⁻¹) at 2000 rpm for 60 s and subsequently annealed at 100 °C, 150 °C, 200 °C or 250 °C for 15 min in a glove box. A Cytos layer (~570 nm) as the gate dielectric was then deposited by spin coating a Cytop solution, followed by drying at 100 °C on a hotplate for 1 h in a glove box before thermally depositing an Al gate electrode layer (~70 nm). All devices were characterized in air in the absence of light using an Agilent 4155C Semiconductor Analyser. Carrier mobility was calculated in the saturation regime according to the equation:

\[ I_D = \frac{WC}{2L} \mu (V_G - V_T)^2 \]

where \( I_D \) is the drain current, \( \mu \) is the charge carrier mobility, \( \mu \) is the capacitance per unit area of the insulator determined from a metal–insulator–metal structure (\( C_i \) is 3.2 nF cm⁻² for a 570 nm-thick Cytop film) and \( V_G \) is the gate voltage.

Synthesis

2-(Trimethylstannyl)thiazole (1). A solution of 2-bromothiazole (1.64 g, 10 mmol) in anhydrous diethyl ether (25 mL) was slowly added into a stirred solution of n-butyllithium (4.4 mL, 2.5 M in hexane) in anhydrous diethyl ether (40 mL) at −78 °C. The mixture was stirred for 1 h, then a solution of trimethyltin chloride (11 mL, 1.0 M in hexane) was added dropwise over 15 min. After stirring for an additional hour at −78 °C, the reaction mixture was quenched with saturated aqueous sodium hydroxide carbonate solution (15 mL) and extracted with diethyl ether. The organic layer was dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The obtained yellow liquid was directly used in the next step without further purification. Yield: 2.51 g (~100%).

1H NMR (300 MHz, CDCl₃) δ 8.15 (d, J = 2.5 Hz, 1H), 7.56 (d, J = 2.7 Hz, 1H), 0.47 (s, 9H). Synthesis of compound 1 was reported previously using a different procedure.

2,2′-Bithiazole (2). A solution of 1 (0.694 g, 2.8 mmol), 2-bromothiazole (0.328 g, 2 mmol) and tetraakis(triphenylphosphine)palladium(0) (Pd(PPh₃)₄) (46 mg, 0.05 mmol) in toluene was refluxed for 24 h under a nitrogen atmosphere. The reaction mixture was allowed to cool down to room temperature before the solvent was removed under reduced pressure. The crude product was purified by column chromatography on silica gel with a mixture of hexane and ethyl acetate (v/v = 7/3) as an eluent to give compound 2 as white crystals.

Yield: 0.263 g (78.1%).

1H NMR (300 MHz, CDCl₃) δ 7.90 (d, J = 3.1 Hz, 2H), 7.44 (d, J = 3.2 Hz, 2H). Synthesis of compound 2 was reported previously using a different procedure.

5,5′-Dibromo-2,2′-bithiazole (3). N-Bromosuccinimide (NBS) (0.872 g, 4.9 mmol) was added in portions to a solution of 2 (0.206 g, 1.2 mmol) in N,N-dimethylformamide (DMF) (10 mL) at room temperature. The reaction mixture was then heated to 60 °C and stirred at this temperature for 5 h. The mixture was poured into water (200 mL) and extracted with dichloromethane. The organic phase was dried over anhydrous sodium sulphate. The crude product was purified by column chromatography on silica gel with a mixture of hexane and ethyl acetate (v/v = 3/1) as an eluent to give compound 3 as yellow crystals.

Yield: 0.390 g (98.0%).

1H NMR (300 MHz, CDCl₃) δ 7.75 (s, 2H).
Di(tritriacontan-17-yl)-4,4′(1,4-dioxo-3,6-di(thiophen-2-yl)pyrrole[3,4-c]pyrrole-2,5,1(1H,4H)-dipyridinatoe (4-2). 4-2 was synthesized following the similar procedure for the synthesis of 4-1 according to the literature methods.37 Yield: 0.67 g (61.0%). 1H NMR (300 MHz, CDCl3) δ 8.90 (d, J = 3.9 Hz, 2H), 7.64 (d, J = 5.0 Hz, 2H), 7.29 (t, J = 4.6 Hz, 2H), 4.86 (t, J = 6.2 Hz, 2H), 4.15 (t, J = 7.4 Hz, 4H), 2.44 (t, J = 7.6 Hz, 4H), 2.10 (q, J = 7.5 Hz, 4H), 0.87 (t, J = 6.5 Hz, 12H). 13C NMR (75 MHz, CDCl3) δ 172.57, 135.36, 131.01, 128.85, 74.87, 41.62, 34.21, 32.14, 32.09, 29.86, 29.82, 29.77, 29.72, 29.70, 29.52, 25.53, 25.48, 22.85, 14.28. HRMS (M + H)+ Calc. for C88H153O6N2S2: 1398.1109; found: 1398.1164.

Di(tritriacontan-17-yl)-4,4′(3,6-bis(5-bromothiophen-2-yl)-1,4-dioxopyrrolo[3,4-c]pyrrole-2,5,1(1H,4H)-dipyridinatoe (5-2). 5-2 was synthesized following the similar procedure for the synthesis of 5-1 according to the literature methods.37 Yield: 0.41 g (93.0%). 1H NMR (300 MHz, CDCl3) δ 8.63 (d, J = 4.2 Hz, 2H), 7.24 (d, J = 4.3 Hz, 2H), 4.87 (t, J = 6.1 Hz, 2H), 4.07 (t, J = 7.4 Hz, 4H), 2.43 (t, J = 7.4 Hz, 4H), 2.14–1.99 (m, 4H), 0.88 (t, J = 6.6 Hz, 12H). 13C NMR (75 MHz, CDCl3) δ 135.35, 131.92, 131.11, 77.36, 75.00, 41.68, 34.21, 32.09, 32.00, 29.87, 29.83, 29.78, 29.71, 29.53, 25.50, 22.86, 14.28. HRMS (M + H)+ Calc. for C88H155Br2O6N2S2: 1553.9302; found: 1553.9345.

Synthesis of PA-1

To a 25 mL Schlenk flask were added compound 3 (24.5 mg, 0.075 mmol), 4-1 (73.0 mg, 0.075 mmol), trans-bis(acetato)bis[o-(di-o-tolylphosphino)benzyl]dipalladium(u) (Herrmann-Beller’s catalyst) (1.4 mg, 1.5 μmol), tris(2-methoxyphenyl)phosphine (2.1 mg, 6.0 μmol), cesium carbonate (56.2 mg, 0.29 mmol), and pivalic acid (2.3 mg, 0.023 mmol). After degassing and refilling argon for 3 times, anhydrous toluene (3 mL) was added under argon. The mixture was refluxed for 16 h. After cooling down to room temperature, the reaction mixture was poured into methanol (100 ml). The precipitates were collected by filtration and subjected to Soxhlet extraction with acetone, hexane, chloroform and TCE. Yield: 7.7 mg (9.0%) from the chloroform-extracted fraction and 45.4 mg (53.2%) from the TCE-extracted fraction. The remaining solid (~38%) is insoluble in any solvent. HT-GPC data: Mn = 17 kDa and PDI = 6.9 (chloroform extracted fraction); Mn = 11 kDa and PDI = 5.9 (TCE extracted fraction).

Synthesis of PA-2

PA-2 was synthesized using compound 3 (24.5 mg, 0.075 mmol) and 4-2 (84.9 mg, 0.075 mmol), following the similar procedure for the synthesis of PA-1. Yield: 95.3 mg (81.3%) from the hexane-extracted fraction. The remaining solid (~19%) is insoluble in any solvent. HT-GPC data: Mn = 15 kDa and PDI = 4.3.

Synthesis of PB-1

PB-1 was synthesized using compound 2 (12.6 mg, 0.075 mmol) and 5-1 (84.9 mg, 0.075 mmol), following the similar procedure for the synthesis of PA-1. Yield: 56.5 mg (66.2%) from the chloroform-extracted fraction and 8.2 mg (9.6%) from the TCE-extracted fraction. The remaining solid (~24%) is insoluble in any solvent. HT-GPC data: Mn = 18 kDa and PDI = 3.8 (chloroform extracted fraction); Mn = 23 kDa and PDI = 3.9 (TCE extracted fraction).

Synthesis of PB-2

PB-2 was synthesized using compound 2 (12.6 mg, 0.075 mmol) and 5-2 (116.7 mg, 0.075 mmol), following the similar procedure for the synthesis of PA-1. Yield: 104.4 mg (89.1%) from the hexane-extracted fraction. The remaining solid (~11%) is insoluble in any solvent. HT-GPC data: Mn = 11 kDa and PDI = 2.5.

Conclusions

We synthesized two bithienyl DPP-bithiazole copolymers PA-1 and PB-1 using the DHAP method via two routes. PA-1 was synthesized by reacting 5,5′-dibromo-2,2′-bithiazole and the DPP monomer, while PB-1 was synthesized by reacting the dibrominated bithienyl DPP and 2,2′-bithiazole monomers. It was found that PA-1 showed a larger band gap, much poorer solubility and lower degree of crystallinity compared to PB-1, which were considered to be due to the presence of a larger amount of α-β coupling linkages and branched (and lightly cross-linked) structures in the former. PB-1, which has less structural defects, achieved very high ambipolar charge transport performance with electron and hole mobilities of up to 0.53 cm2 V−1 s−1 and 5.9 × 10−2 cm2 V−1 s−1, respectively, in OTFT devices. In contrast, PA-1 showed one order of magnitude lower mobilities. Our results demonstrated that placing the bromo groups on different monomers could significantly influence the occurrence of α-β coupling side reactions, which led to the formation of polymers with dramatically different properties. This work for the first time demonstrated that 2,2′-bithiazole is a suitable monomer with good regioselectivity for the construction of regular conjugated polymers with promoted electron transport performance via DHAP. The synthesis of PDBTz’ (Fig. 1), an isomer of PDBTz, by DHAP between 5,5′-bithiazole (or 2,2′-dibromo-5,5′-bithiazole) and the bithienyl DPP monomer 4-1 (or 5-1) is under way. Changing the positions of the nitrogen atoms in the bithiazole units may have an impact on the side reactions and the charge transport performance of the resulting polymers.

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Notes and references