


Cite this: *Mater. Adv.*, 2022,
3, 4831

Two-dimensional benzo[1,2-*b*:4,5-*b'*]difurans as donor building blocks for the formation of novel donor–acceptor copolymers†

Carmen L. Gott-Betts, Alfred A. Burney-Allen, David L. Wheeler and
Malika Jeffries-EL *

Four donor–acceptor copolymers combining the novel electron-donating moiety, 2,3,6,7-tetra(thiophen-2-yl)benzo[1,2-*b*:4,5-*b'*]difuran (BDF) with either 2,3,1-benzothiadiazole (BT) or 2-octyl-2*H*-benzo[*d*][1,2,3]triazole (BTz) as the electron-accepting unit were prepared via the Stille cross-coupling reaction. Two different alkyl chains were attached on the BDF monomers and the impact on the polymers' solubility and thin film morphology was evaluated. The resulting polymers had optical bandgaps between 1.7–1.9 eV and HOMO levels that ranged from –5.60 to –5.85 eV. In thin films, the BTz copolymer showed absorption bands in the range of 400–625 nm. The BT copolymer exhibited two absorption bands: one from 350–450 nm and a broad absorption in the range of 450–750 nm. When the polymers were used as donor materials with the electron-acceptor PC₇₁BM in bulk-heterojunction photovoltaic cells, power conversion efficiencies of up to 2.93% were obtained. These results indicate that BDF is a promising building block for the synthesis of donor materials for use in organic solar cells, due to the favorable energy levels. Although, additional modifications are needed to improve device performance.

Received 2nd February 2022,
Accepted 3rd May 2022

DOI: 10.1039/d2ma00116k

rsc.li/materials-advances

Introduction

The development of new conjugated polymers remains at the forefront of the field of materials chemistry due to their favourable optical and electronic properties which can be modulated through chemical synthesis.^{1–3} Additionally, these organic semiconductors can be fabricated into large area films using solution processing techniques, which would lower the cost of device fabrication.^{4,5} With a virtually infinite number of possible structural variants, synthetic chemists strive to achieve molecular perfection to optimize key figures of merit such as optical absorption, luminescence, frontier molecular orbital energy levels, charge transport and exciton dissociation.^{6,7} A common approach for tuning the optical and electronic properties of an organic semi-conductor is to alternate electron-donating and electron-accepting arenes within the pi-conjugated backbone, thus creating a “push–pull” system. The hybridized frontier molecular orbitals that result from this arrangement allow for facile tuning of the position of the lowest

unoccupied molecular orbital (LUMO), highest occupied molecular orbital (HOMO), and the optical gap.^{6,8,9} Using this approach, it is possible to develop materials with broad absorption profiles, low-lying HOMOs, and high charge carrier mobilities all of which are ideal for use in bulk-heterojunction (BHJ) organic photovoltaics (OPV)s.^{10–12}

The benzo[1,2-*b*:4,5-*b'*]dithiophene (BDT) moiety has been comprehensively investigated as the electron-donating species in donor–acceptor (D/A) conjugated polymers.^{13–16} BDT has a planar backbone that facilitates pi-stacking and enhances charge carrier mobilities. As a result, BDT-based organic semiconductors are among the highest performing materials in the field of OPVs with photocurrent efficiencies (PCE) of ~16.0%,^{17,18} and hole mobilities exceeding 0.1 cm² V^{–1} s^{–1}.¹⁹ Due to the success of BDT-based materials in these applications, our group has turned its focus towards the development its oxygen containing analog, benzo[1,2-*b*:4,5-*b'*]difuran (BDF). Furan is isoelectronic to thiophene, but oxygen has a smaller atomic radius than sulfur. Thus, the steric hindrance between adjacent heterocycles is reduced and the planarity between adjacent rings is increased.^{20,21} At the same time, BDF-based materials have been shown to exhibit improved solubility compared to their BDT analogs.^{20,22} Furthermore, the lower Dewar resonance energy of furan relative to thiophene makes the formation of quinoidal structures more favorable.²³

Department of Chemistry, Division of Materials Science and Engineering, Boston University, Boston, MA 02115, USA. E-mail: malikaj@bu.edu

† Electronic supplementary information (ESI) available: Synthesis of intermediates, ¹H NMR spectra, and CV traces. See DOI: <https://doi.org/10.1039/d2ma00116k>



Consequently, BDF containing materials have a narrower band-gap and more stabilized HOMO level in comparison to their BDT analogs.^{24–26}

Previously we published the synthesis of a new BDF-based building block, 5,5'-(3,7-didecylbenzo[1,2-*b*:4,5-*b'*]difuran-2,6-diyl)bis(4-alkylthiophene-5,2-diyl)bis(trimethyl-stannane) and utilized it in D–A copolymers with 6,6'-dibromo-*N,N'*-(2-octyldodecanyl)-isoindigo.²⁷ The resulting polymers exhibited two main absorption bands: a high energy band due to the π – π^* transition, and a low energy band due to intramolecular charge transfer between the donor and acceptor units. However, the poor solubility of the copolymer adversely impacted the film morphology. As a result, the OPVs fabricated from these polymers exhibited low PCEs of $\sim 1.0\%$. In another report, we detailed the copolymerization of BDF with the electron deficient 1,4-diketopyrrolo[3,4-*c*]pyrrole (DPP) moiety.²⁸ Although the absorption profile was strong and broad within the 600–800 nm range, like many DPP containing polymers, these materials also had poor solubility. As a result of the film morphology, modest device performances of $\sim 3.0\%$ were attained.

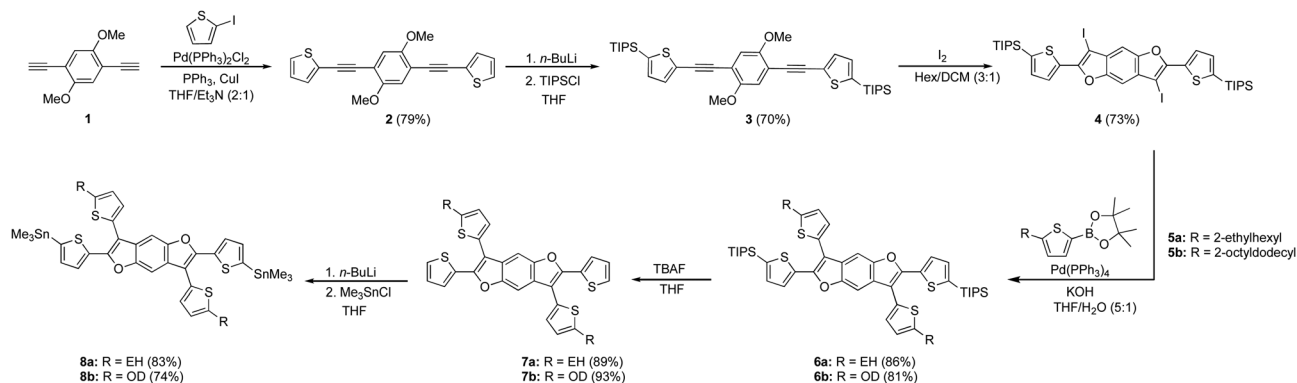
In this report, to further improve upon our BDF building block we introduced thienyl substituents onto the BDF moiety. Using aromatic substituents instead of alkyl or alkoxy-chains broadens absorption profiles and improves the intramolecular charge transfer (ICT) within the material.²⁴ Our functional BDF has reactive handles at the 3- and 7-positions. Thus, this molecule can be used to produce novel two-dimensional BDF monomers as most of the previous examples of two-dimensional BDT/BDF polymers have the aryl substituents at the 4- and 8-positions.^{19,22,29} Herein, We synthesized four donor–acceptor copolymers using a novel two-dimensional monomer 2,3,6,7-tetra(thiophen-2-yl)benzo[1,2-*b*:4,5-*b'*]difuran with either 2,3,1benzothiadiazole (BT) or 2-octyl-2*H*-benzo[*d*][1,2,3]triazole (BTz) as the comonomer. Here, BT and BTz were chosen for their facile synthesis and their demonstrated utility in high performing OPV materials.^{23,24} For the two-dimensional axillary thiophenes, alkyl side chains were attached to enhance solubility. The alkyl chain length was also varied to evaluate its impact on device performance.

Results and discussion

The synthesis of the benzodifuran monomers is shown in Scheme 1. Compound 2 was obtained from the Sonogashira cross-coupling reaction of 2-iodothiophene and 1,4-diethynyl-2,5-dimethoxybenzene.³⁰ The 5 and 5' positions of the thiophenes were protected with triisopropylsilyl (TIPS) groups to produce 3, which upon iodine mediated cyclization afforded the benzodifuran core 4. The subsequent Suzuki cross-coupling reaction of 4 with either 2-ethylhexyl (EH) 5a³¹ or 2-octyldodecyl (OD) 5b³¹ thiophenepinacolborane yielded 6a and 6b, respectively. Removal of the TIPS groups from 6a and 6b afforded 7a and 7b, respectively. The lithiation of 7a and 7b and subsequent quenching with trimethyltin chloride produced the target monomers 8a and 8b. A highlight of the monomer synthesis detailed here is that all intermediates and products were obtained in relatively high isolated yields ranging from 70%–93%.

The synthesis for the polymers is shown in Scheme 2. The polymers were all synthesized *via* Stille cross-coupling reaction of either 8a or 8b with the corresponding comonomers to yield polymers P1–P4. The yields ranged from 51–84% after purification *via* Soxhlet extraction. All polymers were soluble in chloroform at room temperature enabling characterization by proton nuclear magnetic resonance spectroscopy (¹H NMR).

The spectra obtained are consistent with the expected polymer structures. The molecular weights were assessed using gel permeation chromatography (GPC) at 25 °C using THF as the eluent. The resulting characterization data is summarized in Table 1. Comparing the molecular weights of polymers with the same alkyl chain (P1 vs. P2, P3 vs. P4), P2 has a slightly lower molecular weight (M_n and M_w) when compared to its BTz analog P1. Further, both polymers have low degrees of polymerization, indicating that the ethyl hexyl side chain is not adequate to impart solubility. Conversely, P3 has a lower molecular weight compared to P4. Assuming equal reactivity, this suggests that the additional alkyl chain on the BTz unit did not increase the solubility of the polymer. When comparing the polymers with the same acceptor units (P1 vs. P3, P2 vs. P4), the 2-ethylhexyl substituted polymers have lower molecular weights



Scheme 1 Synthesis of BDF monomers.



Scheme 2 Synthesis of D–A Copolymers **P1–P4**.

Table 1 Thermal and molecular weight data for polymers

Polymer	Yield ^a (%)	M_n^b (Da)	M_w^b (Da)	D	DP _n	T_d^c (°C)
P1	65	6155	9045	1.47	10	393
P2	84	4898	6246	1.28	7	363
P3	76	11 453	17 305	1.51	14	379
P4	51	20 054	29 558	1.47	26	389

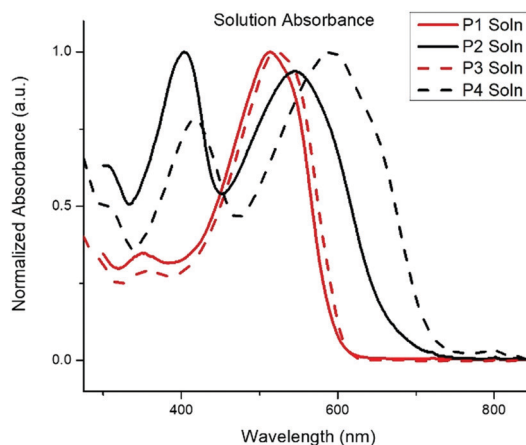
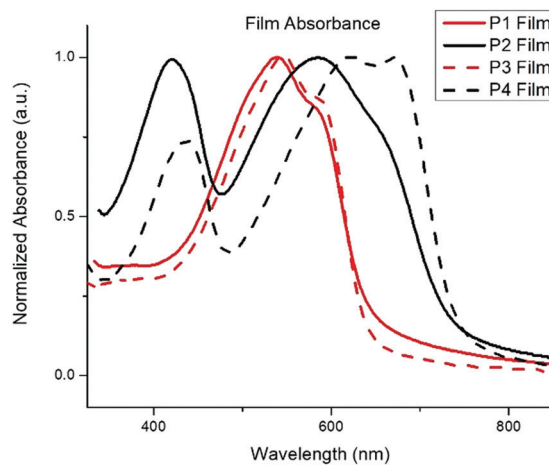
^a Isolated yield after Soxhlet extraction. ^b Molecular weight data was obtained by GPC (see ESI). ^c 5% weight loss temperature determined by TGA.

and degrees of polymerization than the 2-octyldodecyl variants. Assuming equal reactivity, this suggests that the longer branched alkyl substituents on the BDF core are better for improving solubility and increasing molecular weight.

The thermal properties of the polymers were evaluated using differential scanning calorimetry (DSC) and thermal gravimetric analysis (TGA). DSC did not reveal any observable phase transitions for temperatures up to 250 °C. The TGA results are summarized in Table 1 and indicate that 5% weight loss onsets occurred between 363–393 °C. This thermal data supports that these polymer materials are thermally stable well above the operating temperature threshold for organic photovoltaic devices.³²

Optical and electrochemical properties

The normalized absorption spectra for **P1–P4** in dilute CHCl₃ solutions and thin films are shown in Fig. 1 and 2, respectively. The corresponding data is summarized in Table 2. The spectra for both BTz polymers **P1** and **P3** exhibit a weak high energy peak and a strong peak with a λ_{\max} at 539 and 546 nm respectively. Whereas the spectra for the BT polymers **P2** and **P4** have two distinct absorption bands as is typically seen in D/A copolymers: a low energy peak resulting from intramolecular charge transfer (ICT) and a higher energy peak associated with the π - π^* transition.³³ In the solid-state spectra, the small high energy bands for **P1** and **P3** have almost disappeared, and the lower energy peaks now display small shoulders, slightly broadened peak widths, and concurrent red-shifting of the λ_{\max} . The side chains did not have any impact on the solid-state

Fig. 1 Solution absorption profile of **P1–P4** with the concentration of 10^{-5} M in chloroform.Fig. 2 Thin film absorption spectra of **P1–P4**. Films spun from CHCl₃ solutions.Table 2 Optical and electronic data polymers **P1–P4**

Polymer	λ_{\max} (nm) Solution	λ_{\max} (nm) Film	$E_g^{\text{OPT } a}$ (eV)	HOMO ^b (eV)	LUMO ^c (eV)
P1	513	539	1.9	−5.85	−4.0
P2	403, 545	421, 585	1.7	−5.60	−3.9
P3	520	546	1.9	−5.71	−3.8
P4	417, 592	437, 622	1.7	−5.82	−4.1

^a Optical bandgap as measured by the optical onset ($E_g = 1240/\lambda_{\text{offset}}$). ^b HOMO levels measured by CV (see ESI). ^c Estimated LUMO = HOMO + E_g .

packing as the λ_{\max} for both **P1** and **P3** shifted by 26 nm from solution to film. Conversely, the spectra for the BT polymers showed a significant peak broadening between the solution and solid state along with a hypsochromic shift of all peaks. The extent of spectral broadening and shifting was dependent on the length of the side chains, with **P4** being the broadest.



To evaluate the electrochemical properties of the polymers, the redox profile was evaluated using cyclic voltammetry, see ESI.† All four polymers exhibit measurable and reproducible oxidation processes, but reduction curves were not observed. The HOMO levels were estimated from the onset of oxidation using the absolute energy level of ferrocene/ferrocenium (Fc/Fc⁺) as 5.1 eV under vacuum and are summarized in Table 2. The polymers had calculated HOMO values between -5.60 eV and -5.85 eV affording good stability in air, and open circuit voltage (V_{OC}), making them suitable for use in OPVs.³⁴ The LUMO values were estimated from the HOMO and the optical bandgap and ranged from -3.8 eV to -4.1 eV. As a result, the offset between the polymer LUMO and that of the fullerene (-4.1 eV) acceptor is small, which can negatively impact device performance.³⁵

Photovoltaic devices

The performance of all four polymers in OPV devices were evaluated using [6,6]-phenyl-C71-butyric acid methyl ester (PC₇₁BM) as the n-type electron acceptor with a device configuration of ITO/PEDOT:PSS/polymer:PC₇₁BM/Ca/Al. The active layers were deposited from 30 mg mL⁻¹ *o*-DCB with 0–2% of 1-chloronaphthalene (CN) as a high-boiling solvent additive to improve polymer/PCBM blend morphology. The current density–voltage curves are shown in the ESI.† The resulting short circuit current density (J_{SC}), open circuit voltage (V_{OC}), fill factor (FF), and power conversion efficiency (PCE) for the devices are shown in Table 3.

Among the devices fabricated without solvent additives, **P4** gave the highest PCE at 2.93% with the highest FF of 64.86. **P4** also demonstrated a moderate V_{OC} of 0.73 V and moderate J_{SC} of 6.18 mA cm⁻². **P1** and **P2** gave lower efficiencies at 1.81% and 1.63% respectively. The lowest performing polymer, **P2**, had a modest V_{OC} at 0.67 V but a lower FF of 35.07 and J_{SC} of 6.93 mA cm⁻². When CN was added, there was an increase in performance for all the polymers except for **P4**, due to increases in J_{SC} and/or V_{OC} . **P3** had the highest performing device at 2.67% under these conditions. The highest performing device for **P1** required a 2% CN additive compared to 1% for the other three polymers. The decrease in the **P4** device performance with the addition of CN additive is likely a result of its higher solubility than the other polymers.

A comparison of the alkyl chains shows that the ethylhexyl polymers **P1** and **P2** had significantly lower FF than their

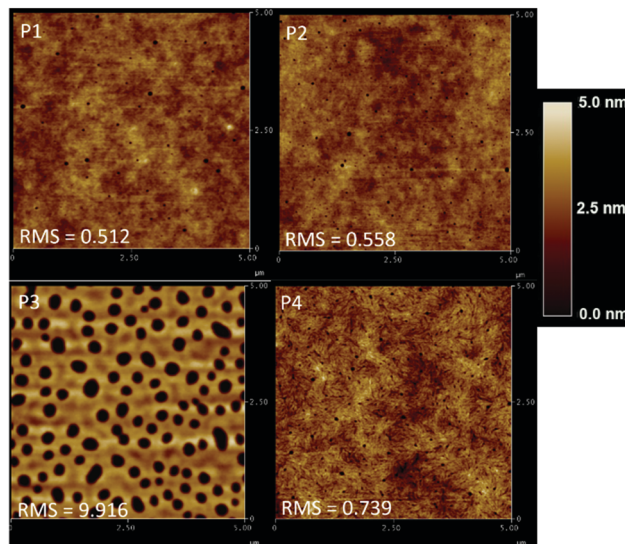


Fig. 3 AFM images of highest performing device active layers for **P1–P4**.

octyldodecyl variants, **P3** and **P4**. Conversely, **P1** and **P2** showed higher J_{SC} compared to the octyldodecyl polymers, **P3** and **P4**. Overall polymers with ethylhexyl side chains had poorer performance when compared to their octyldodecyl counterparts. Next, when comparing polymers with different electron withdrawing units, the nature of the electron deficient comonomer was not significant as **P1** performed better than **P2**, but **P4** performed better than **P3**.

To further investigate the performance trends, the morphology of the highest performing polymer/PC₇₁BM devices was evaluated using atomic force microscopy (AFM) as shown in Fig. 3. The root mean-square (RMS) values are shown in Table 3. When first evaluating the effect of alkyl chain length on the BDF donor, it was observed that **P2** and **P4** formed relatively smooth films with RMS roughness values of 0.56 and 0.74. The OPV performance increased from 1.75% for **P2** to 2.93% with **P4** even though **P4** had a higher roughness value. Analogously, **P1** has a very smooth film compared to **P3** with RMS values of 0.51 and 9.9, respectively. Although **P3** has a very large RMS value, the overall device performance was like **P1** with PCE values of 2.67% and 2.33% respectively. While it has been reported that a higher RMS value may lead to better photovoltaic performance,³⁶ high RMS values indicate that

Table 3 Photovoltaic device performance for **P1–P4**

Polymer	Additive (%)	J_{SC} (mA cm ⁻²)	V_{OC} (V)	FF	PCE (%)	RMS
P1	None	8.47 ± 0.40	0.50 ± 0.01	34.77 ± 1.31	1.67 ± 0.10	—
	2% CN	8.80 ± 0.79	0.60 ± 0.01	37.19 ± 1.03	1.98 ± 0.21	0.512
P2	None	6.69 ± 0.33	0.64 ± 0.02	35.10 ± 0.79	1.51 ± 0.09	—
	1% CN	6.99 ± 0.52	0.64 ± 0.4	33.62 ± 1.15	1.53 ± 0.15	0.558
P3	None	4.60 ± 0.19	0.81 ± 0.00	58.89 ± 1.10	2.22 ± 0.08	—
	1% CN	5.21 ± 0.27	0.83 ± 0.00	56.15 ± 1.05	2.45 ± 0.15	9.916
P4	None	5.79 ± 0.34	0.72 ± 0.01	65.36 ± 0.90	2.75 ± 0.15	0.739
	1% CN	5.47 ± 0.40	0.73 ± 0.00	57.90 ± 2.35	2.33 ± 0.18	—

Average of 8 devices.



the device films made with **P3** have pinholes and/or completely void areas. In comparing the electron accepting unit within the D–A polymer it was observed that **P1** and **P2** have similar RMS values of 0.51 and 0.56 respectively. Thus, the heterocycles have less of an impact on film morphology than the side chains do. Whereas, for **P3** and **P4**, the RMS values were too different to determine if the difference in performance was due to the smoother film or to the intrinsic optical properties of the materials. However, it is impressive that **P3** has such a high PCE considering the severe film defects. These results suggest that a mid-range side chain may be beneficial to improve device properties. Further optimization of the **P3** device film would be necessary to produce truly optimized devices.

Conclusions

A series of donor–acceptor copolymers based on electron withdrawing 4,7-dibromo-2-octyl-2*H*-benzo[*d*][1,2,3]triazole or 4,7-dibromobenzo[*c*][1,2,5]thiadiazole units and electron donating 2,3,6,7-tetra(thiophen-2-yl)benzo[1,2-*b*:4,5-*b'*]difurans bearing different alkyl chains were synthesized. Polymer pairings with the same comonomer had similar absorption profiles and optical bandgaps, regardless of the side-chain length. Whereas the polymer sets with the same alkyl chain had different optical profiles due to the relative difference in the strength of the electron acceptor unit. It was hypothesized that the two-dimensional BDF monomer would lead to improvements in OPV performance. The resulting polymers had low-lying LUMO levels, and deep HOMO-levels both of which were favorable for a novel OPV donor materials. However, the band gaps ranging from 1.7–1.9 eV, which combined with the poor film forming properties lead to modest OPV performances. Despite these outcomes, the aforementioned synthetic ease and oxidative stability make the BDT moiety a promising building block for designing new materials.³⁷ In the future, we will investigate the BDF moiety with stronger electron-accepting comonomers. Doing this would both reduce the polymer bandgap and allow for the use of additional alkyl character to improve solubility and improve film formation.

Experimental methods

Materials

Air and moisture sensitive reactions were completed using standard Schlenk techniques. Solvents used for metal catalyzed reactions were deoxygenated prior to use by sparging with nitrogen while vigorously stirring for 30 minutes. SiliaFlash Irregular Silica Gels F60 was purchased from Silicycle for purification of materials using silica columns. All other chemical reagents were purchased from commercial sources. The molecules 4,7-dibromo-2-octyl-2*H*-benzo[*d*][1,2,3]triazole,³⁸ 4,7-dibromobenzo[*c*][1,2,5]thiadiazole,³⁹ 1,4-diethynyl-2,5-dimethoxybenzene (1),³⁰ and 4,4,5,5-tetramethyl-2-(5-(2-octyl-dodecyl)thiophen-2-yl)-1,3,2-dioxaborolane³¹ were prepared according to literature procedures. The synthesis of 2-[5-(2-

ethylhexyl)thien-2-yl]-4,4,5,5-tetramethyl-1,3,2-dioxaborolane is provided in the ESI.†

Characterization

Nuclear magnetic resonance (NMR) spectra were obtained using chloroform-*d* unless otherwise noted and recorded on an Agilent 500 MHz VNMRS spectrometer with a Varian ultra-shielded magnet or an Agilent 400 MHz VNMRS unity plus spectrometer with an Oxford Instruments superconducting magnet. The ¹H NMR spectra were internally referenced to the residual protonated solvent peak and all chemical shifts are given in ppm (δ) relative to the solvent, 7.26 for chloroform. The gel permeation chromatography (GPC) measurements were conducted at 25 °C using THF and a flow rate of 1.0 mL min⁻¹. Calibration was based on polystyrene standards. Thermogravimetric analysis (TGA) measurements were performed over an interval of 100–500 °C at a heating rate of 10 °C min⁻¹ under nitrogen atmosphere. Differential scanning calorimetry (DSC) was obtained using a scan heating rate of 10 °C min⁻¹ between -30 °C and 250 °C. Cyclic voltammetry (CV) was performed using an e-DAQ e-corder 410 potentiostat with a scanning rate of 100 mV s⁻¹. The polymer solutions (5 mg mL⁻¹ in chloroform) were drop-cast on a platinum electrode and Ag/Ag⁺ was used as the reference electrode and a platinum wire as the auxiliary electrode. The reported values are referenced to Fc/Fc⁺ (-5.1 eV *versus* vacuum).⁴⁰ All cyclic voltammetry experiments were performed in deoxygenated acetonitrile under an argon atmosphere using 0.1 M tetrabutylammonium hexafluorophosphate as the electrolyte. Absorption spectra were obtained on a photodiode-array Shimadzu UV-1800 ultraviolet spectrophotometer using polymer solutions in chloroform and thin films. The thin films were made by spin-coating 25 × 25 × 1 mm glass slides using solutions of polymer (5 mg mL⁻¹) in chloroform at a spin rate of 1200 rpm.

Fabrication of photovoltaic devices

All organic photovoltaic devices were performed under inert atmosphere using solution-based spin casting. The indium tin oxide (ITO) substrates had a resistance of 40 Ohm. The device size was 25 mm. Conventional architecture was used to fabricate organic photovoltaic devices *via* (ITO)/Poly(3,4-ethylene-dioxythiophene): poly(styrene sulfonate) (PEDOT:PSS)/polymer:[6,6]-phenyl-C₇₁-butyric acid methyl ester (PC₇₁BM)/Ca/Al). ITO glass substrates were cleaned *via* sonication in (1) Mucosal, (2) deionized water, (3) acetone, and (4) isopropanol for 10 minutes each respectively. Slides were dried in an oven followed by UV-ozone treatment for 15 minutes. PEDOT:PSS layers (Clevios P VP Al 4083) were filtered (0.45 μ m) and spin coated onto ITO substrates at 3500 rpm for 2 minutes and annealed for 100 °C for 30 min in air. After cooling, substrates were transferred into a nitrogen filled glove box. All polymer:PC₇₁BM solutions were prepared in 1:2–3 and 1:1 blend ratio (20 mg mL⁻¹), respectively, with chlorobenzene as the processing solvent. The prepared solutions were allowed to stir for 24 h at 80 °C prior to spin coating onto the PEDOT:PSS layer at 1000 rpm for 2 minutes. Following spin coating, Calcium



(15 nm) and Aluminum (100 nm) electrodes underwent successful deposition *via* thermal evaporation. Films were also evaluated under post thermal annealing conditions at 100 °C for 10 minutes. Current-density (J - V) data were obtained *via* Keithley 2400 source meter and simulated AM 1.5 G illumination (100 mW cm⁻², Newport 91160) calibrated using a KG-5 filter Silicon reference cell.

Synthesis

2,2'-((2,5-dimethoxy-1,4-phenylene)bis(ethyne-2,1-diyl))dithiophene (2). A dry round bottom 250 mL flask with a stir bar was attached to a condenser under nitrogen environment and charged with **1** (5.44 g, 29.2 mmol), Pd(PPh₃)₂Cl₂ (0.41 g, 0.58 mmol), CuI (0.22 g, 1.17 mmol), and PPh₃ (0.31 g, 1.17 mmol). The flask was then purged and backfilled with N₂ three times before adding deoxygenated anhydrous THF/Et₃N (2:1) and 2-iodothiophene (7.10 mL, 64.27 mmol) then stirred overnight at room temperature. The reaction mixture was poured into ice and extracted three times with dichloromethane. The combined organic layers were washed with saturated aqueous NH₄Cl, water, and brine. The organic layer was dried over MgSO₄, evaporated, and purified using a silica plug (hex/DCM 4:1). The resulting solid was recrystallized from ethyl acetate/hexanes (2:1) to yield **2** as a yellow solid (8.09 g, 79% yield). ¹H NMR (500 MHz, CDCl₃) δ 7.31 (m, 4H), 7.02 (m, 4H), 3.90 (s, 6H). ¹³C NMR (125 MHz, CDCl₃) δ 153.87, 132.37, 127.75, 127.27, 123.28, 115.44, 113.28, 89.47, 88.40, 77.41, 77.16, 76.91, 56.58. LRMS (ESI) m/z : [M+H]⁺ calculated for C₂₀H₁₄O₂S₂: 350.04; found: 351.0513 m/z , error 1.0113 ppm.

((2,5-dimethoxy-1,4-phenylene)bis(ethyne-2,1-diyl))bis(thiophene-5,2-diyl)bis(triisopropylsilane) (3). Compound **2** (4.29 g, 12.24 mmol) was added to a dry 250 mL round bottom flask with a stir bar. The reaction vessel was then purged and backfilled with Nitrogen three times. Anhydrous THF was cannulated into the round bottom and the reaction vessel was cooled to -78 °C before *n*-BuLi (2.5 M in hexanes, 10.77 mL, 26.93 mmol) was added dropwise *via* syringe. The green reaction mixture was stirred at -78 °C for 1 hour before the dropwise, syringe addition of tri-isopropylsilyl chloride (5.76 mL, 26.93 mmol). The reaction was warmed to room temperature and stirred overnight. The reaction mixture was quenched with 20 mL of deionized water, stirred for 1 hours, and the solvent removed. The resulting solid was dissolved in CHCl₃ and washed with NH₄Cl, water, and brine. The organic layer was dried over MgSO₄, evaporated, and purified by recrystallization from Ethyl acetate/hexanes (3:1) (5.68 g, 70% yield). ¹H NMR (500 MHz, CDCl₃) δ 7.39 (d, J = 3.53 Hz, 2H), δ 7.15 (d, J = 3.5 Hz, 2H), δ 6.99 (s, 2H), δ 3.88 (s, 6H), δ 1.34 (m, 6H), 1.11 (d, J = 7.44 Hz, 36H). ¹³C NMR (125 MHz, CDCl₃) δ 153.57, 138.10, 135.28, 132.75, 127.74, 115.13, 113.13, 90.41, 88.22, 77.16, 76.91, 76.65, 56.33, 18.41, 11.65. HRMS (ESI) m/z : [M+H]⁺ calculated for C₃₈H₅₄O₂S₂Si₂: 663.3182; found: 663.3209 m/z , error 4.0704 ppm.

((3,7-diiodobenzo[1,2-*b*:4,5-*b'*]difuran-2,6-diyl)bis(thiophene-5,2-diyl)bis(triisopropylsilane) (4). A 500 mL Erlenmeyer flask was charged with **3** (5.30 g, 8 mmol) which was dissolved in CH₂Cl₂ and cooled to 0 °C. The solution was rapidly stirred as iodine (6.09 g, 23.98 mmol) in 200 mL CH₂Cl₂/hexanes (3:1) was added dropwise

over 10 minutes. Upon completion of the addition, the reaction mixture was stirred at 0 °C for 4 hours. The cold reaction mixture was then filtered to collect the crude product. The solid was stirred in hot hexanes, cooled, and filtered. The yellow solid was then recrystallized from toluene to afford yellow crystals (5.18 g, 73% yield). ¹H NMR (500 MHz, CDCl₃) δ 8.08 (d, J = 3.6 Hz, 2H), δ 7.45 (s, 2H), δ 7.33 (d, J = 3.6 Hz, 2H), δ 1.40 (m, 6H), 1.15 (d, J = 7.4 Hz, 36H). HRMS (ESI) m/z : [M+H]⁺ calculated for C₃₆H₄₈I₂O₂S₂Si₂: 886.07; found: 887.0829, error 3.0437 ppm.

Suzuki cross-coupling procedure for 6a and 6b. A round bottom flask was charged with **4** (1 equiv.), KOH (10 equiv.), alkyl boronic acid pinacol ester (3 equiv.), and Pd(PPh₃)₄ (0.05 equiv.) in THF/H₂O (5:1) under nitrogen. The reaction mixture was heated at 80 °C overnight. Next, the mixture was poured into 100 mL of water and extracted three times with dichloromethane. The combined organic layers were washed with deionized water, NH₄Cl, brine, and dried over MgSO₄. The solvent was removed *in vacuo* and the residue was sent through a silica plug using hexanes then used without further purification as a viscous, orange semi-solid. (**6a**, 86%): ¹H NMR (500 MHz, CDCl₃) δ 7.60 (s, 2H), δ 7.57 (d, J = 3.60 Hz, 2H), δ 7.17 (d, J = 3.61 Hz, 2H), δ 7.12 (d, J = 3.40 Hz, 2H), δ 6.86 (d, J = 3.39 Hz, 2H), δ 2.85 (d, J = 6.69 Hz, 4H), δ 1.67 (m, 2H), δ 1.30–1.45 (m, 22H), δ 1.11 (d, J = 7.4 Hz, 36H), δ 0.95 (m, 12H). HRMS (ESI) m/z : [M+H]⁺ calculated for C₆₀H₈₆O₂S₄Si₂: 1022.50; found: 1023.5139, error 1.0747 ppm. (**6b**, 81%): ¹H NMR (400 MHz, CDCl₃) δ 7.60 (s, 2H), 7.56 (d, J = 3.6 Hz, 2H), 7.17 (d, J = 3.7 Hz, 2H), 7.12 (d, J = 3.3 Hz, 2H), 6.85 (d, J = 3.3 Hz, 2H), 2.85 (d, J = 6.6 Hz, 4H), 1.71 (m, 2H), 1.36 (m, 70H), 1.11 (d, J = 7.3 Hz, 36H), 0.89 (m, 12H).

TBAF deprotection procedure for 7a and 7b. Compound **6** was added to a dry, one neck 250 mL round bottom flask. The flask was purged and refilled with Nitrogen three times before the addition of 100 mL of anhydrous THF. The flask was cooled to 0 °C and TBAF (4 equiv., 1M in THF) was added *via* syringe dropwise over 5 minutes. Next, the flask was warmed to room temperature and stirred overnight under nitrogen environment. The reaction mixture was then poured into NH₄Cl and extracted with chloroform. The organic layer was washed three times with water followed by brine and dried over MgSO₄. The solvent was then removed *in vacuo* and the crude product was purified by column chromatography using hexanes. The resulting solid was then recrystallized from ethanol to give orange/yellow crystals (**7a**, 89%): ¹H NMR (500 MHz, CDCl₃) δ 7.59 (s, 2H), δ 7.51 (d, J = 4.79 Hz, 2H), δ 7.12 (d, J = 3.42 Hz, 2H), δ 7.04 (m, 2H), δ 6.87 (d, J = 3.42 Hz, 2H), δ 2.86 (d, J = 6.35 Hz, 4H), δ 1.68 (m, 2H), δ 1.30–1.47 (m, 16H), δ 0.94 (m, 12H). ¹³C NMR (125 MHz, CDCl₃) δ 151.17, 148.65, 146.49, 132.58, 129.59, 129.34, 128.23, 127.54, 126.68, 126.21, 125.76, 110.80, 101.03, 77.41, 77.16, 76.91, 41.66, 34.48, 32.59, 29.07, 25.84, 23.20, 14.33, 14.29, 11.07. HRMS (ESI) m/z : [M+H]⁺ calculated for C₄₂H₄₆O₂S₄: 710.24; found: 711.2477, error 2.5308 ppm. (**7b**, 93%): ¹H NMR (500 MHz, CDCl₃) δ 7.58 (d, J = 0.6 Hz, 2H), 7.51 (d, J = 3.7 Hz, 2H), 7.30 (d, J = 6.1 Hz, 2H), 7.11 (d, J = 3.4 Hz, 2H), 7.07–7.00 (m, 2H), 6.86 (d, J = 3.4 Hz, 2H), 2.85 (d, J = 6.7 Hz, 4H), 1.74–1.69 (m, 2H), 1.36–1.21 (m, 64H), 0.88



(d, J = 5.1 Hz, 12H). ^{13}C NMR (125 MHz, CDCl_3) δ 151.20, 148.68, 146.51, 132.61, 129.59, 129.36, 128.24, 127.50, 126.64, 110.83, 40.25, 34.93, 33.51, 32.08, 31.75, 30.19, 29.87, 29.53, 26.84, 25.44, 22.85, 14.27. HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calculated for $\text{C}_{66}\text{H}_{94}\text{O}_2\text{S}_4$: 1046.61; found: 1047.6223, error 0.7636 ppm.

Stannylation procedure for 8a and 8b. Compound 7 was added to a dry 250 mL round bottom flask. The reaction vessel was charged with 100 mL of anhydrous THF and cooled to -78°C . Next, *n*-BuLi (2.10 equiv., 2.5 M in hexanes) was added dropwise to the flask *via* syringe. After stirring at -78°C for 1 hour, the reaction was quenched with trimethyltin chloride (2.5 equiv., 1 M in THF) added *via* syringe into the -78°C reaction mixture. The flask was then allowed to warm to room temperature, stirred overnight, poured into water, and extracted three times with dichloromethane. The organic layer was washed three times with water and brine. The solvent was removed *in vacuo* and the crude product used without further purification as an orange semi-solid. (**8a**, 83%): ^1H NMR (400 MHz, CDCl_3) δ 7.60–7.58 (m, 4H), 7.12 (m, 4H), 6.87 (m, 2H), 1.37 (m, 22H), 0.93 (m, 12H), 0.38 (s, 18H). (**8b**, 74%): ^1H NMR (400 MHz, CDCl_3) δ 7.59 (m, 4H), 7.10 (m, 4H), 6.86 (m, 2H), 1.26 (m, 70H), 0.89–0.85 (m, 12H), 0.38 (s, 18H).

General procedure for polymerization with 4,7-dibromobenzo[*c*]-[1,2,5]thiadiazole. The respective benzodifuran monomer (0.29 g, 0.21 mmol) was added to a dry, three neck round bottom 100 mL flask equipped with a stir bar and a condenser. 4,7-Dibromobenzo[*c*]-[1,2,5]thiadiazole (82 mg, 0.21 mmol) was added to the flask followed by $\text{Pd}_2(\text{dba})_3$ (3.9 mg, 0.004 mmol, 0.02 equiv.) and *P*(*o*-tol) $_3$ (5.1 mg, 0.016 mmol, 0.08 equiv.). The flask was purged and back filled three times with nitrogen. Anhydrous toluene was cannulated into the flask and the reaction heated to reflux for three days. On the third day, the polymer was end-capped by the addition of excess iodobenzene and trimethyl(phenyl)stannane. The reaction was heated at reflux overnight. After being allowed to cool to room temperature, the solvent was removed, the crude mixture dissolved in 10 mL of chloroform, and precipitated into 200 mL of methanol cooled to -78°C . The solid was collected in a Soxhlet thimble and the polymers purified *via* Soxhlet extraction using methanol, acetone, hexanes, and, lastly, chloroform. The chloroform fraction was evaporated until approximately 10 mL of chloroform remained and the solution was precipitated into -78°C methanol. The collected solid was again dissolved in 10 mL of chloroform and precipitated for a second time into methanol cooled to -78°C . The solid was collected once more and further purified *via* a chloroform silica plug. The chloroform was then removed, the product redissolved in 10 mL of Chloroform and precipitated a final time into -78°C methanol to yield **P1** (65%) and **P3** (76%). **P1**: ^1H NMR (500 MHz, CDCl_3) δ 8.04 (br, 2H), δ 7.82 (br, 1H), δ 7.45 (br, 4H), δ 7.12 (br, 2H), δ 6.88 (br, 2H), δ 6.78 (br, 1H), δ 4.79 (br, 2H), δ 2.74–2.90 (br, 4H), δ 2.20 (br, 2H), δ 1.28–1.71 (br, 28H), δ 0.87–0.98 (br, 15H). **P3**: ^1H NMR (500 MHz, CDCl_3 , δ) 8.08 (br, 2H), 7.84 (br, 1H), 7.58 (br, 2H), 7.43 (br, 2H), 7.12 (br, 2H), 6.92 (br, 2H), 6.77 (br, 1H), 4.80 (br, 2H), 2.89 (br, 4H), 2.21 (br, 2H), 1.24–1.39 (br, 76H), 0.86 (br, 15H).

General procedure for polymerization with 4,7-dibromo-2-octyl-2*H*-benzo[*d*][1,2,3]triazole. The respective benzodifuran monomer (0.31 g, 0.23 mmol) was added to a dry, three

neck 100 mL round bottom flask equipped with a stir bar and condenser. Next, 4,7-dibromo-2-octyl-2*H*-benzo[*d*][1,2,3]-triazole (89 mg, 0.23 mmol) was added to the reaction vessel followed by $\text{Pd}_2(\text{dba})_3$ (4.1 mg, 0.004 mmol, 0.02 equiv.) and *P*(*o*-tol) $_3$ (5.5 mg, 0.018 mmol, 0.08 equiv.). The flask was purged and back filled three times with nitrogen and then anhydrous toluene was cannulated into the flask and the reaction heated to reflux for three days. On the third day, the polymer was end-capped by the addition of excess iodobenzene and trimethyl(phenyl)stannane. The reaction was heated at reflux overnight. After being allowed to cool to room temperature, the solvent was removed, the crude mixture dissolved in 10 mL of chloroform, and precipitated into 200 mL of methanol cooled to -78°C . The solid was filtered into a Soxhlet thimble, and the polymers purified *via* Soxhlet extraction using methanol, acetone, hexanes, and, lastly, chloroform. The chloroform fraction was evaporated until approximately 10 mL of chloroform remained and the solution precipitated into -78°C methanol. The collected solid was again dissolved in 10 mL of chloroform and precipitated for a second time into methanol cooled to -78°C . The solid was collected once more and further purified *via* a chloroform silica plug. The chloroform was then removed, the product redissolved in 10 mL of Chloroform and precipitated a final time into -78°C methanol to yield **P2** (84%) and **P4** (51%). **P2**: ^1H NMR (500 MHz, CDCl_3) δ 8.03 (br, 1H), 7.76 (br, 2H), 7.49 (br, 3H), 7.10 (br, 3H), 6.92 (br, 2H), 6.77 (br, 1H), 2.75–2.90 (br, 6H), 1.26–1.73 (br, 16H), 0.89–1.06 (br, 12H). **P4**: ^1H NMR (500 MHz, CDCl_3 , δ) 8.08 (br, 1H), 7.81 (br, 2H), 7.55 (br, 3H), 7.15 (br, 3H), 6.93 (br, 1H), 2.92 (br, 4H), 2.74 (br, 2H), 1.26–1.57 (br, 63H), 0.86 (br, 12H).

Author contributions

Carmen Gott-Betts: conceptualization, methodology, investigation, writing – original draft, writing – review & editing, and visualization; Alfred Burney Allen: methodology, investigation, writing – original draft, and writing – review & editing; David Wheeler: methodology, investigation, writing – original draft, and writing – review & editing; Malika Jeffries-EL: conceptualization, methodology, investigation, writing – original draft, writing – review & editing, visualization supervision, project administration and funding acquisition.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors would like to thank Dr Norman Lee and Dr Paul Ralifo from the Chemical Instrumentation Facility at Boston University for analysis of our molecules and polymers. We also thank Dr Anlee Krupp for help with AFM analysis of polymer films. Next, the authors would like to thank the Center for Electronic Materials and Devices at the University of



Massachusetts-Amherst for device fabrication. We also thank Professor Mark Grinstaff for usage of thermal gravimetric analysis equipment and Aidan Murphy and Sara El-Arid for help with gel permeation chromatography analysis. Lastly, we thank the National Science Foundation (DMR-1410088/1640297) for and Boston University for financial support of this work.

References

- S. C. Rasmussen, *ChemPlusChem*, 2020, **85**, 1412–1429.
- R. M. Pankow and B. C. Thompson, *Polymer*, 2020, **207**, 122874.
- Z. Qiu, B. A.-G. Hammer and K. Müllen, *Prog. Polym. Sci.*, 2020, **100**, 101179.
- I. Burgues-Ceballos, M. Stella, P. Lacharmoise and E. Martinez-Ferrero, *J. Mater. Chem. A*, 2014, **2**, 17711–17722.
- G. Li, R. Zhu and Y. Yang, *Nat. Photonics*, 2012, **6**, 153–161.
- T. Mikie and I. Osaka, *J. Mater. Chem. C*, 2020, **8**, 14262–14288.
- I. Osaka, in *Organic Solar Cells: Energetic and Nanostructural Design*, ed. M. Hiramoto and S. Izawa, Springer, Singapore, 2021, pp. 89–121, DOI: [10.1007/978-981-15-9113-6_5](https://doi.org/10.1007/978-981-15-9113-6_5).
- E. E. Havinga, W. ten Hoeve and H. Wynberg, *Polym. Bull.*, 1992, **29**, 119–126.
- E. E. Havinga, W. ten Hoeve and H. Wynberg, *Synth. Met.*, 1993, **55**, 299–306.
- W. Chen, G. Huang, X. Li, Y. Li, H. Wang, H. Jiang, Z. Zhao, D. Yu, E. Wang and R. Yang, *ACS Appl. Mater. Interfaces*, 2019, **11**, 33173–33178.
- W. Zhao, S. Li, H. Yao, S. Zhang, Y. Zhang, B. Yang and J. Hou, *J. Am. Chem. Soc.*, 2017, **139**, 7148–7151.
- C. Lee, S. Lee, G.-U. Kim, W. Lee and B. J. Kim, *Chem. Rev.*, 2019, **119**, 8028–8086.
- J. G. Laquindanum, H. E. Katz, A. J. Lovinger and A. Dodabalapur, *Adv. Mater.*, 1997, **9**, 36–39.
- H. Yao, L. Ye, H. Zhang, S. Li, S. Zhang and J. Hou, *Chem. Rev.*, 2016, **116**, 7397–7457.
- Y. Liang and L. Yu, *Acc. Chem. Res.*, 2010, **43**, 1227–1236.
- Y. Che, Y. Zhang, Y. Yang, C.-H. Liu, R. Izquierdo, S. S. Xiao and D. F. Perepichka, *J. Org. Chem.*, 2020, **85**, 52–61.
- S. Liu, J. Yuan, W. Deng, M. Luo, Y. Xie, Q. Liang, Y. Zou, Z. He, H. Wu and Y. Cao, *Nat. Photonics*, 2020, **14**, 300–305.
- Y. Cui, H. Yao, J. Zhang, T. Zhang, Y. Wang, L. Hong, K. Xian, B. Xu, S. Zhang, J. Peng, Z. Wei, F. Gao and J. Hou, *Nat. Commun.*, 2019, **10**, 2515.
- Y.-W. Huang, Y.-C. Lin, Y.-S. Wu, Y.-T. Wong, M.-Y. Kuo, W.-C. Chen and C.-C. Chueh, *Ind. Eng. Chem. Res.*, 2020, **59**, 9105–9115.
- O. Gidron and M. Bendikov, *Angew. Chem., Int. Ed.*, 2014, **53**, 2546–2555.
- M. Jeffries-El, B. M. Kobilka and B. J. Hale, *Macromolecules*, 2014, **47**, 7253–7271.
- L. Liang, J.-T. Wang, X. Xiang, J. Ling, F.-G. Zhao and W.-S. Li, *J. Mater. Chem. A*, 2014, **2**, 15396–15405.
- R. S. Hosmane and J. F. Liebman, *Tetrahedron Lett.*, 1992, **33**, 2303–2306.
- L. Huo, T. Liu, B. Fan, Z. Zhao, X. Sun, D. Wei, M. Yu, Y. Liu and Y. Sun, *Adv. Mater.*, 2015, **27**, 6969–6975.
- J. Warnan, C. Cabanetos, A. E. Labban, M. R. Hansen, C. Tassone, M. F. Toney and P. M. Beaujuge, *Adv. Mater.*, 2014, 4357–4362, DOI: [10.1002/adma.201305344](https://doi.org/10.1002/adma.201305344).
- E. He, Y. Lu, Z. Zheng, F. Guo, S. Gao, L. Zhao and Y. Zhang, *J. Mater. Chem. C*, 2020, **8**, 139–146.
- B. M. Kobilka, A. V. Dubrovskiy, M. D. Ewan, A. L. Tomlinson, R. C. Larock, S. Chaudhary and M. Jeffries-EL, *Chem. Commun.*, 2012, **48**, 8919–8921.
- B. M. Kobilka, B. J. Hale, M. D. Ewan, A. V. Dubrovskiy and M. Jeffries-El, *Polym. Chem.*, 2013, **3**, 5329–5336.
- C. Gu, D. Liu, J. Wang, Q. Niu, C. Gu, B. Shahid, B. Yu, H. Cong and R. Yang, *J. Mater. Chem. A*, 2018, **6**, 2371–2378.
- J. L. Novotney and W. R. Dichtel, *ACS Macro Lett.*, 2013, **2**, 423–426.
- M. Scheuble, Y. M. Gross, D. Trefz, M. Brinkmann, J. T. López Navarrete, M. C. Ruiz Delgado and S. Ludwigs, *Macromolecules*, 2015, **48**, 7049–7059.
- G. Bardizza, E. Salis, C. Toledo and E. D. Dunlop, *Prog. Photovoltaics*, 2020, **28**, 593–600.
- B. C. Thompson, Y.-G. Kim and J. R. Reynolds, *Macromolecules*, 2005, **38**, 5359–5362.
- C. G. Tang, M. C.-Y. Ang, K.-K. Choo, V. Keerthi, J.-K. Tan, M. N. Syafiqah, T. Kugler, J. H. Burroughes, R.-Q. Png, L.-L. Chua and P. K.-H. Ho, *Nature*, 2016, **539**, 536–540.
- N. A. Ran, J. A. Love, M. C. Heiber, X. Jiao, M. P. Hughes, A. Karki, M. Wang, V. V. Brus, H. Wang, D. Neher, H. Ade, G. C. Bazan and T.-Q. Nguyen, *Adv. Energy Mater.*, 2018, **8**, 1701073.
- M. He, W. Han, J. Ge, Y. Yang, F. Qiu and Z. Lin, *Energy Environ. Sci.*, 2011, **4**, 2894–2902.
- X. Li, Y. Li, Y. Zhang and Y. Sun, *Small Sci.*, 2022, 2200006.
- A. Balan, G. Gunbas, A. Durmus and L. Toppare, *Chem. Mater.*, 2008, **20**, 7510–7513.
- B. Burkhart, P. P. Khlyabich, T. Cakir Canak, T. W. LaJoie and B. C. Thompson, *Macromolecules*, 2011, **44**, 1242–1246.
- C. M. Cardona, W. Li, A. E. Kaifer, D. Stockdale and G. C. Bazan, *Adv. Mater.*, 2011, **23**, 2367–2371.

