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High-quality conjugated polymers *via* one-pot Suzuki–Miyaura homopolymerization†

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This paper describes a one-pot Suzuki–Miyaura homopolymerization that involves *in situ* borylation/cross coupling of dibrominated donor–acceptor conjugated macromonomers, in contrast to the standard Stille copolymerization of thienosilole and isoindigo monomers. Reaction kinetics investigation reveals that bis(pinacolato)diboron promotes an efficient polymerization. The homopolymer showed blue-shifted absorption compared to the Stille copolymer, which is rationalized by quantum chemical calculations of a series of oligomers containing various donor–acceptor configurations. The calculations suggest that the homopolymerization of asymmetrical macromonomers likely introduced both acceptor–acceptor and donor–donor segments into the backbone. The acceptor–acceptor segment is found to contribute mostly to the blue-shift of maximum absorption wavelength. Furthermore, detailed analysis of MALDI-TOF (matrix-assisted laser-desorption ionization-time of flight) spectra of these two polymers indicated that while the homopolymer is well defined, the Stille copolymer is end-capped mostly with the thienosilole moieties and/or methyl groups, implicating that destannylation and methyl transfer are the most-likely chain-termination pathways that limit high molecular weight. This is in sharp contrast to the homopolymerization, where chain-terminators are required to control the molecular weight for obtaining soluble material. The photovoltaic performances of bulk-heterojunction solar cells based on these polymers are compared.

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Introduction

The Stille cross-coupling copolymerization¹ has been a method of choice for the development of donor–acceptor conjugated alternating copolymers for a host of applications such as polymer solar cells,^{2–4} polymer light-emitting diodes,⁵ thin-film transistors⁶ and even polymer-based sensors.⁷ Such popularity is largely due to its remarkable tolerance for many functional groups, which enables the design and preparation of a variety of functional polymers. Yet the convenience of Stille cross-coupled polymerization comes at a cost in terms of the unavoidable toxicity and the difficulty of purifying the organostannanes that are necessary to achieve high molecular weights, as well as possible synthetic complications due to tin–tin homocouplings when certain catalyst/organostannane combinations are used.⁸ These drawbacks may pose limitations on the molecular weights and structural integrity of the resulting polymers. In

this sense, a number of research groups have set out to prepare better copolymers from Stille copolymerization.^{9–12} Results suggest that high molecular weight, low polydispersity and better structural integrity can contribute to enhanced light absorption and improved morphology, which translate to better overall device performance.

Along with the popularity of Stille cross-coupled copolymerization, Suzuki cross-coupling has been widely used to synthesize high molecular weight alternating copolymers. However the monomer scope of the Suzuki copolymerization is primarily limited to mildly electron-rich building blocks such as carbazoles,^{13,14} and fluorenes,^{15,16} which is probably due to the tendency towards deborylation of the boronic acid/esters of such moieties. These challenges highlight the significance of developing one-pot reactions that eliminate the necessity to acquire purified boronic acids/esters. To this end, a handful of research groups have investigated the scope of one-pot synthesis of biaryls in terms of catalyst, base and aryl halides.^{17–22} Early discoveries have shown that bis(pinacolato)diboron could be used as a condensation reagent for the one-pot synthesis of low to moderate molecular weight poly(arylene)s,²³ and that certain bases such as K₂CO₃ may lead to oligomer formation in biaryl systems.²⁴ In 2009, Reynolds *et al.* revisited this synthetic pathway and specifically studied the effect of CsF on the homopolymerization of fluorenes.²⁵

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Furthermore, in one of our earlier contributions, it has been shown that with symmetrical model systems, *in situ* borylation/cross coupling demonstrates a general superiority over the Stille copolymerization in terms of end-group control.²⁶

In this work, we further develop the chemistry of one-pot borylation/cross-coupling polymerization. Instead of employing simple, head-to-tail symmetrical, relatively electron-neutral monomers, we constructed a head-to-tail asymmetrical donor-acceptor conjugated macromonomer that was subjected to the homopolymerization. Particularly, we compare the as-obtained homopolymer with a copolymerized counterpart synthesized from the standard Stille cross coupling, in terms of the optoelectronic properties and structural integrity.

Experimental section

General

All reagents and solvents were purchased from commercial sources and were used without further purification unless indicated otherwise. UV/vis measurements were carried out on a Perkin Elmer Lambda 9000 spectrometer in 1 cm quartz cuvettes with concentrations of 0.03–0.1 mg mL⁻¹ in CHCl₃. MALDI-TOF spectra were taken on a Biosystems Voyager apparatus. Samples were prepared by mixing the matrix (10 mg mL⁻¹ in THF) and polymer sample (1 mg mL⁻¹ in THF or CHCl₃) at room temperature in a 3 : 1 volume ratio. All measurements were performed in positive ion mode. NMR spectra were measured using a Varian AMX400 (400 MHz) instrument at 25 °C. FT-IR spectra were recorded on a Nicolet Nexus FT-IR fitted with a Thermo Scientific Smart iTR sampler. GPC measurements were done on a Spectra Physics AS 1000 series machine equipped with a Viskotek H-502 viscometer and a Shodex RI-71 refractive index detector. The columns (PLGel 5 m mixed-C) (Polymer Laboratories) were calibrated using narrow disperse polystyrene standards (Polymer Laboratories). Samples were made in CHCl₃ at a concentration of 2–3 mg mL⁻¹.

Synthesis

Compounds **1**,²⁷ **2** and **3** (ref. 28 and 29) were synthesized according to literature methods. The thienosilole precursors for **2** and **3** were synthesized with a modified procedure based on literature method.²⁹ The synthetic steps of the two polymers are shown in Scheme 1.

P-Stille

A 25 mL, flame-dried two neck flask, filled with dry N₂, was charged with **1** (100 mg, 0.155 mmol) and **2** (118 mg, 0.16 mmol), followed by the addition of Pd₂(dba)₃ (CHCl₃ adduct, recrystallized from acetone) (4 mg), P(*o*-tol)₃ (16 mg). Degassed dry toluene was then introduced into the reaction flask *via* a cannula. The reaction was then set at 100 °C for 24 h. Afterwards, the reaction mixture was allowed to cool to room temperature, and added dropwise into 500 mL of methanol. The precipitate was collected with a cellulose thimble, which was then subjected to Soxhlet extraction with methanol for 24 h, acetone for 7 h, hexane for 5 h, and CHCl₃ for 3 h, subsequently.

The chloroform fraction was obtained as a purple-blue solid (128 mg, 85%). ¹H NMR (400 MHz, CDCl₃) δ 9.08 (2H, br), 7.54 (1H, br), 7.43 (1H, br), 6.51 (1H, br), 3.74 (4H, br), 2.94 (1H, br), 1.22 (34H, br), 0.79 (30H, br). IR (cm⁻¹): 2954, 2920, 2853, 1689, 1608, 1453, 1376, 1352, 1170, 1102, 1073, 1013, 874, 817, 739.

Compound 4

A 50 mL, flame-dried three neck flask, filled with dry N₂, was charged with of **1** (500 mg, 0.776 mmol) and of **3** (480 mg, 0.83 mmol), followed by the addition of Pd(PPh₃)₄ (45 mg). Degassed dry toluene was then introduced into the reaction flask *via* a cannula. The reaction was then stirred at 100 °C overnight. After cooling to room temperature, most of the solvent of the resulting mixture was removed under reduced pressure. The resulting purple oil was subjected to silica gel chromatography with toluene/hexanes (v/v = 1/3) as the eluent. Compound **4** was obtained as a purple oil (450 mg, 60% yield). ¹H NMR (400 MHz, CDCl₃) δ 9.13 (d, *J* = 8 Hz, 1H), 9.02 (d, *J* = 8 Hz, 1H), 7.40 (s, 1H), 7.27 (dd, *J* = 8 Hz, 1H), 7.17 (dd, *J* = 8 Hz, 1H), 7.08 (d, *J* = 4 Hz, 1H), 6.92 (d, 1H), 6.86 (d, 1H), 3.62 (m, 4H), 1.84 (m, 2H), 1.35 (m, 20H), 1.18 (m, 10H), 0.93 (m, 20H), 0.80 (m, 12H). ¹³C NMR (101 MHz, CDCl₃) δ 168.59, 168.20, 149.84, 148.59, 145.94, 145.79, 144.51, 144.38, 143.31, 138.71, 133.19, 130.61, 130.55, 130.44, 130.01, 127.38, 126.12, 125.82, 124.85, 120.65, 120.35, 118.83, 111.30, 104.62, 44.30, 44.10, 37.77, 37.45, 35.92, 35.64, 30.81, 30.59, 28.87, 28.60, 24.28, 24.00, 22.97, 17.63, 14.17, 10.82.

Compound 5

A 50 mL round-bottom flask was charged with of **4** (320 mg, 0.33 mmol) and THF (20 mL). NBS (70.5 mg, 0.396 mmol) was added all at once. After 2 h, the product mixture was extracted with diethyl ether, and the organic phase was collected and dried under reduced pressure. The resulting purple oil was subjected to silica gel chromatography with toluene/hexanes (v/v = 1/3) as the eluent, yielding 245 mg of **5** (70% yield). ¹H NMR (400 MHz, CDCl₃) δ 9.14 (d, *J* = 8 Hz, 1H), 9.03 (d, *J* = 8 Hz, 1H), 7.37 (s, 1H), 7.26 (dd, *J* = 8 Hz, 1H), 7.16 (dd, *J* = 8 Hz, 1H), 7.03 (s, 1H), 6.93 (d, 1H), 6.89 (d, 1H), 3.64 (m, 4H), 1.85 (m, 2H), 1.37 (m, 24H), 1.17 (m, 10H), 0.96 (m, 16H), 0.79 (m, 12H).

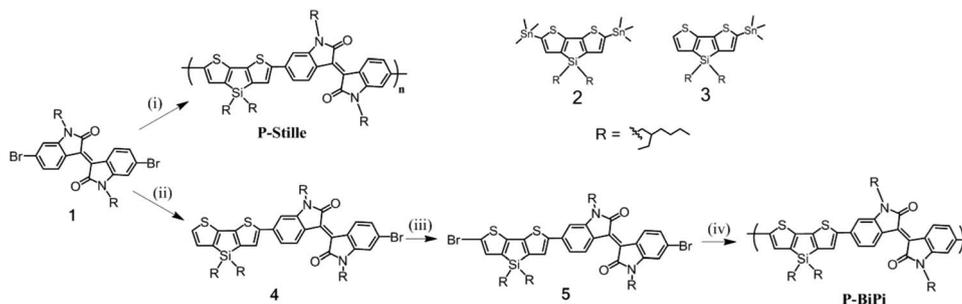
Reaction for P-BiPi kinetics

A 50 mL, flame-dried two neck flask, filled with dry N₂, was charged with **5** (80 mg, 0.0755 mmol) and then Pd(dppf)Cl₂ (CH₂Cl₂ adduct) (5 mg), K₃PO₄ (64 mg) and bis(pinacolato) diboron (19 mg, 0.0775 mmol). Degassed toluene/DMF (v/v = 4/1) (15 mL) was introduced into the reaction flask *via* a cannula. The reaction was set at 80 °C. A sample of 1 mL was taken from the reaction flask at given time spot for UV-vis and GPC measurements.

P-BiPi

A 25 mL, flame-dried two neck flask, filled with dry N₂, was charged with **4** (11 mg, 0.0113 mmol), **5** (100 mg, 0.0943 mmol) and then Pd(dppf)Cl₂ (CH₂Cl₂ adduct) (4 mg), K₃PO₄ (80 mg)





Scheme 1 Synthesis of polymers. Reagents: (i) **2**, Pd₂(dba)₃ (CHCl₃ adduct, recrystallized from acetone), P(*o*-tol)₃, toluene; (ii) **3**, Pd₂(dba)₃ (CHCl₃ adduct), P(*o*-tol)₃, toluene; (iii) NBS, THF; (iv) bis(pinacolato)diboron, K₃PO₄, Pd(dppf)Cl₂ (CH₂Cl₂ adduct), toluene/DMF (v/v = 4/1). P-Stille refers to the polymer obtained from a standard Stille copolymerization, while P-BiPi is the polymer synthesized with the BiPi promoted polycondensation. BiPi means bis(pinacolato)diboron.

and bis(pinacolato)diboron (25 mg). Degassed toluene/DMF (v/v = 4/1) (10 mL) was introduced into the reaction flask *via* a cannula. The reaction was set at 80 °C overnight. Afterwards, the reaction mixture is cooled room temperature, and added dropwise into 500 mL of methanol. The precipitate was collected with a cellulose thimble, which was subjected to Soxhlet extraction with methanol for 24 h, acetone for 7 h, hexane for 5 h, and CHCl₃ for 5 h, respectively. Production of the chloroform fraction appears as a purple solid (70 mg, 70%). ¹H NMR (400 MHz, CDCl₃) δ 9.22 (1H, br), 9.19 (1H, br), 7.41 (1H, br), 7.31 (2H, br), 7.03 (1H, br), 7.00 (2H, br), 3.74 (4H, br), 1.93 (2H, br), 1.25 (34H, br), 0.83 (28H, br). IR (cm⁻¹): 2954, 2920, 2853, 1689, 1608, 1453, 1376, 1352, 1170, 1102, 1073, 874, 817, 739.

Density functional theory calculations

Geometries of dimer (DAAD, DADA and ADDA) and tetramers (DADADADA and ADDAADDA) were optimized using Density Functional Theory (DFT) (B3LYP/6-311G*) with ORCA³⁰ without any symmetry constraints. Subsequently, the vertical excitation energies were calculated using time-dependent DFT (TD-DFT) (B3LYP/6-311G*). The lowest 10 excited states were calculated for the tetramers.

Photoluminescence measurements

Photoluminescence measurements were carried out on solutions contained in quartz cuvettes (2 mm path length) in transmission mode. The excitation source was the second harmonic (approximately 400 nm) of a mode-locked Ti:Sapphire laser (Mira 900, Coherent) delivering 150 ps pulses at a repetition rate of 76 MHz. The laser power was adjusted using neutral density filters; and the excitation beam was spatially limited by an iris. The beam was focused with a 150 mm focal length lens onto a spot of approximately 50 μm. Steady state spectra were collected into a spectrometer with a grating of 50 lines per mm and recorded with an em-CCD array (Hamamatsu, ImagEM) sensitive between 400 nm and 900 μm. For time resolved measurements, the same pulsed excitation source was used. Spectra were collected on a Hamamatsu streak camera (Hamamatsu, Japan) working in Synchroscan mode (time

resolution 2 ps) with a cathode sensitive in the visible. All spectra were corrected for the spectral response of the setup using a calibrated lamp.

Solar cell device fabrications and measurements

Indium tin oxide (ITO) coated glass substrates were precleaned with deionized water, CMOS grade acetone, and then isopropanol, each for 15 min. Possible organic residues were removed by UV-ozone cleaning for 20 min. A layer of PEDOT:PSS (poly(3,4-ethylenedioxythiophene):poly(styrenesulfonate), VP A14083, H. C. Stark) (thickness 60 nm) was spin-cast on top at 1500 rpm for 50 s. After being baked at 140 °C for 10 min, the substrates were transferred into a nitrogen-filled glovebox (<0.1 ppm of O₂ and H₂O). Atop that, active layers of polymer:[60]PCBM (99.5%, Solenne B.V.) (D/A weight ratio = 1 : 4) were spin-coated in a N₂-filled glovebox from *o*-DCB solution with 3% (v/v) 1,8-diiodooctane (DIO, Sigma Aldrich) at 600 rpm for 5 s and 400 rpm for 120 s. The donor-to-acceptor total concentration was 30 mg mL⁻¹. The thickness of the photoactive layers was about 100 nm. Finally, LiF/Al (1 nm and 100 nm, respectively) were thermally evaporated at a pressure of <10⁻⁶ Torr on top of the organic layer to make the sandwiched structure ITO/PEDOT:PSS/active layer/LiF/Al. The active area of the device was 0.04 cm². The current-voltage characteristics of all solar cell devices were measured with a Keithley 2400 source meter and were conducted in a nitrogen-filled glovebox. An AM 1.5 G solar simulator (AAA grade, XES-70S1) was used as the light source. The light intensity of the solar simulator was calibrated to be 100 mW cm⁻² (at the position of sample) using a standard silicon reference solar cell (area 20 × 20 mm², the certification of the reference cell is accredited by NIST to the ISO-17025 standard). Hole-only devices were fabricated on bare glass substrates which were carefully cleaned by washing with detergent solution and ultrasonication in acetone and isopropyl alcohol, followed by UV-ozone treatment. A Cr(1 nm)/Au(30 nm) anode was then thermally evaporated onto the substrate, followed by the coating of a 60 nm thick film of PEDOT-PSS. The film was then baked at 120 °C for 10 min. Subsequently, polymer:fullerene blends were deposited by spin-coating from *ortho*-dichlorobenzene solutions under N₂ atmosphere. The hole-only



devices were finished by thermal evaporation of a Pd(20 nm)/Au(80 nm) top contact.

Results and discussion

Synthesis of polymers

Scheme 1 presents the synthesis of the two polymers **P-Stille** and **P-BiPi**, where **P-Stille** is the copolymerization result of the dibrominated compound **1** and the distannyl thienosilole **2** and **P-BiPi** is the polymer obtained by the homopolymerization method. The obtainable molecular weight of the **P-Stille** product is maximized by gently refluxing the polymerization mixture for 24 h, giving a M_n and M_w of 11 and 16 kDa ($D = 1.45$), respectively. On the other hand, after a successful synthesis of the dibrominated macromonomer **5**, an initial attempt to homopolymerize this macromonomer **5** overnight at 100 °C led to a 97% yield of intractable black solid, which could be either be due to cross-linking or a very high molecular weight. The possibility of cross-linking under the given conditions was readily excluded with the high purity of **5** determined from its HPLC absorption trace (Fig. S1†), excluding significant over-bromination of compound **4**. Furthermore, infrared spectroscopy measurements of **5** and the insoluble material showed overlapping spectra (Fig. S2†), unambiguously indicating that the intractable material comprises ultra-high molecular weight polymer chains, based on macromonomer **5**. These observations underscore the capacity of the *in situ* borylation/cross coupling route to yield high molecular weight polymeric materials.

To better control the molecular weight, we further studied the kinetics of this specific homopolymerization by monitoring the molecular weight and light absorption changes over time, as shown in Fig. 1(a). It can be seen that the polymerization was notably fast, with the absorption saturating 4 h after the reaction was started at room temperature. The consecutively red-shifting absorptions suggest a continuous growth of the polymer chains. This observation is further correlated with the GPC

results shown in Fig. 1(b), which indicate that polymers with $M_n = 9.3$ kDa are obtained after a reaction period of 4.5 h. Further prolonging the reaction time quickly led to insoluble product. Moreover, Fig. 1(b) shows a strong attenuation in the growth of the molecular weight over time (and over the conversion of monomers). Additionally, the fact that **P-BiPi** becomes completely intractable in a small time window after the initial polymerization period hints to cross coupling of long oligomers, leading to long and insoluble polymer chains that precipitate. The kinetic study suggests that with the specific macromonomer, the one-pot SM polymerization yields molecular weights comparable to that of the standard Stille copolymerization. Since molecular weight is not our pursuit within the current chemical investigation, no further optimization on molecular weights was conducted.

Note that the above analysis does not provide direct insight into the polymerization mechanism (step-growth or chain-growth), but a highly efficient polymerization of dibrominated donor-acceptor macromonomers was indeed observed. In terms of controlling the molecular weight, an efficient way is to pre-mix chain-terminators with the starting materials at the very beginning, such that part of the bifunctional monomers will be partially deactivated towards further reaction. Indeed, when using compound **4** as the chain-terminator for reaction (iv), we were able to isolate CHCl_3 -soluble polymers with a yield of 70%. Compound **4** was chosen as the chain-terminator because it shows the same reactivity as the macromonomer, ensuring a proper and efficient chain termination. The molecular weight of the polymer is limited by the chain terminator, which is reflected by increased product solubility in our case. An initial attempt of adding 5% of chain terminator (molar percentage relative to macromonomer) did not allow a satisfactory isolation of CHCl_3 -soluble polymers, even when the reaction time was limited. Isolation of CHCl_3 -soluble polymers was possible when the chain terminator percentage was increased to 10%.

Optoelectronic properties

With the insights obtained from the kinetic study, we were able to synthesize soluble polymeric materials for characterization. We first measured the UV-vis absorption spectra of polymers **P-Stille** and **P-BiPi** in chloroform, as shown in Fig. 2. Interestingly,

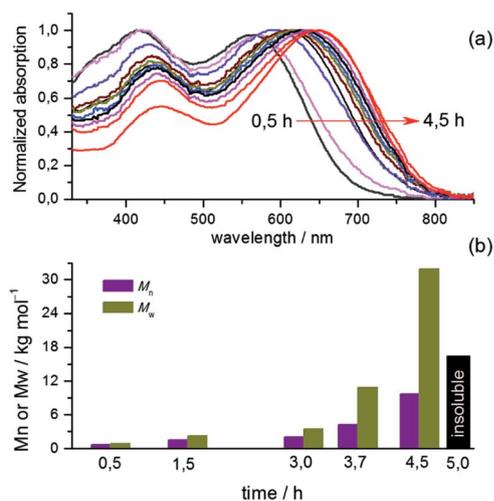


Fig. 1 (a) Normalized UV-vis absorptions and (b) molecular weight changes over time of the homopolymerization. The black bar in (b) means the reaction product has already become intractable.

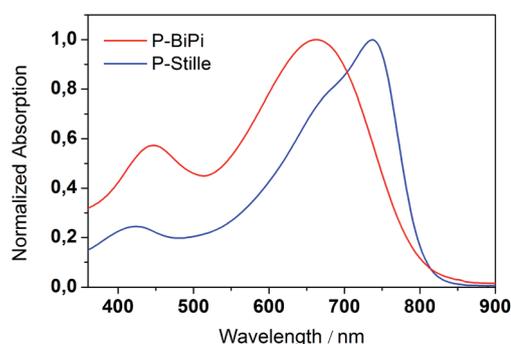


Fig. 2 UV-vis absorption spectra of the Stille copolymer **P-Stille** and Suzuki-Miyarau homopolymer **P-BiPi**.



while the cut-off wavelengths of the two absorptions are close, there is a 75 nm blue shift of the maximum absorption wavelength from **P-Stille** (737 nm) to **P-BiPi** (662 nm). **P-BiPi** also showed apparently enhanced absorption of high energy photons (an absorption band peaking around 410 nm) compared to **P-Stille**. These observations indicate a significant structural difference between **P-Stille** and **P-BiPi**. Considering the fact that both the Miyarau and Suzuki catalytic cycles involve oxidative insertion as the first step, which preferably takes place on the isoindigo end due to its electron deficiency, this structural difference is most likely related to the cross coupling of two isoindigo ends (forming an A–A segment). The resulting oligomers ending with donor moieties may then lead to donor–donor cross couplings (forming a D–D segment), in addition to the normal donor–acceptor cross coupling (forming a D–A segment).

To understand the structural origin of these absorption features, we considered a series of oligomers containing different conjugation segments, and calculated their absorption characteristics by time-dependent DFT (TD-DFT). We first calculated the absorption bands of DADA, DAAD and ADDA segments. These results are shown in Fig. S3.† It is clear that an ADDA fragment enables the absorption of low-energy photons, while a DAAD fragment absorbs the higher energy photons more amongst these three oligomers. We note that these three dimers were minimized in a coplanar conformation. In this sense, the calculated absorptions of these dimers are fairly qualitative. Nevertheless, the assessment between theory and measurement suggests that the existence of DAAD segments in the backbone will promote the absorption of high-energy photons, which is in accord with the observation shown in Fig. 1. This phenomenon is also present in the calculated absorption bands of DADADADA and ADDAADDADA tetramers, the latter of which contains both AA and DD segments (Fig. 2). On the basis of these observations, we further calculated the electronic transitions of the ADDAADDADA tetramer, for which an absorption spectrum was calculated as shown in Fig. 3(B). Despite the differences, it is clear that the calculated spectrum tends to replicate the absorption peaks of the experimental counterpart. This result provides indirect evidence that the as-presented *in situ* borylation/cross coupling polymerization has introduced DD, AA and DA segments into the backbone, probably in a random pattern.

These random donor–acceptor configurational patterns introduce structural disorder in the backbone of homopolymerized **P-BiPi**, which is further reflected in several optoelectronic properties of the material. For example, as shown in Fig. 4(a), while the Stille copolymer **P-Stille** shows a Stokes shift of 0.18 eV in chloroform solution, the Stokes shift of the homopolymer **P-BiPi** is notably increased to 0.35 eV. As noted by several authors, this might indicate that the structural disorder gives rise to a broader density of states, with emission occurring after relaxation towards low energy sites.³¹ Moreover, as seen in Fig. 4(b), while **P-Stille** clearly showed a mono-exponential photoluminescence decay with a fitted excitonic lifetime of 313 ps, the photoluminescence decay of **P-BiPi** is bimodal exponential, suggesting a multi-pathway decay with lifetimes of 41 ps and 5 ps.

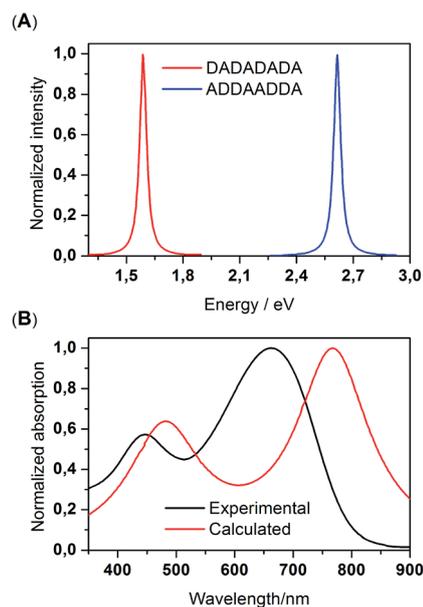


Fig. 3 (A) Calculated absorption bands of two planarized DA-tetramers, DADADADA and ADDAADDADA. (B) A comparison of the calculated absorption spectrum of an ADDAADDADA tetramer, and the measured absorption spectra of the **P-BiPi**.

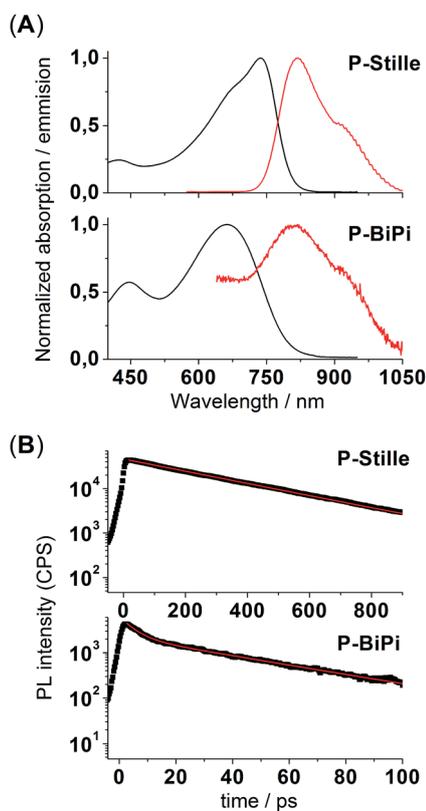


Fig. 4 (A) Normalized absorption (black lines) and emission (red lines) spectra of **P-Stille** and **P-BiPi** measured in chloroform. (B) Photoluminescence spectra of **P-Stille**, hexane and chloroform fractions of **P-BiPi**. The red lines are mono-exponential (**P-Stille**) or bimodal-exponential (**P-BiPi**) fits.



Although the detailed mechanism of this phenomenon is beyond the scope of this work, these observations seem to suggest that the existence of AA and/or DD segments in conjugated polymers adversely affects exciton lifetimes, and may also lead to significant fluorescence quenching as shown by the photoluminescence intensities in Fig. 4(B).

Mass spectrometry analysis

MALDI-TOF spectrometry has been widely applied to polymers, both with saturated backbones and with conjugated backbones, to infer structural information.^{32–34} Due to the soft-ionization nature of MALDI-TOF, the chemical composition of the polymers remains largely intact (*i.e.* absence of fragmentation). Therefore, given preliminary structural information and proper sample preparations, the analysis of MALDI-TOF spectra can be straightforward and reliable, although it does not give adequate quantitative information regarding the average molecular weight of the polymer sample. The MALDI-TOF spectra of **P-Stille** and **P-BiPi** are shown in Fig. 5. Note that due to the relatively poor solubility of **P-BiPi** in CHCl_3 , the hexane fraction instead of CHCl_3 fraction of the homopolymerization product was used to maintain a sufficient concentration at room temperature. As seen in Fig. 5(A), the *in situ* borylation/cross coupling polymerization yielded samples showing a clean repetition of peaks, with a repetition interval of 900 Da, in accord with the mass of a D–A repeating unit. Additionally, no obvious residual Br or boronic ester end groups were observed in the case of **P-BiPi**. This is most likely due to the fact that a H-capped chain terminator (compound **4** in Scheme 1) was used. There are some side-peaks which are of slightly higher m/z values following the major peaks, as shown in Fig. 5(B). One of them might be a polymer chain with hydroxyl groups transferred from the dithranol matrix. The structural origin of the other lower peaks is not clear.

In contrast to the clean and simple mass spectrum of **P-BiPi**, a very different scenario was recorded for the Stille-copolymerized **P-Stille**. Although well-defined repetitions are found for all major peaks, only some of them can be sensibly assigned to corresponding structures. As shown in Fig. 5(A), oligomeric molecules of the same chemical composition can be found for **P-Stille** and **P-BiPi**. Yet it is interesting to notice that compared to the dihydro-terminated oligomers based on the same number of donor and acceptor moieties (as defined by the **P-BiPi** spectrum of Fig. 5(A)), all peaks with the highest intensity in the **P-Stille** spectrum are 416 Da (mass of a repeating donor moiety) higher ($5 \rightarrow 6$, $7 \rightarrow 8$, and $9 \rightarrow 10$). This indicates two possibilities: (i) a majority of the resulted polymerized molecules are ended with a donor moiety; (ii) this extra donor moiety is incorporated into the backbone through tin–tin homocoupling. Since the spectrum is clearly free of polymer chains with trimethyltin end groups, the first possibility further implies that destannylation behaves as a major pathway for chain termination, impeding further chain growth. The origin of destannylation was not investigated specifically, but we hypothesize that destannylation is more likely to occur when electron-rich heteroarylstannanes are involved.³⁵ Destannylation as a side reaction is expected to imbalance the stoichiometry of the original monomer ratio, leading to polymer chains with

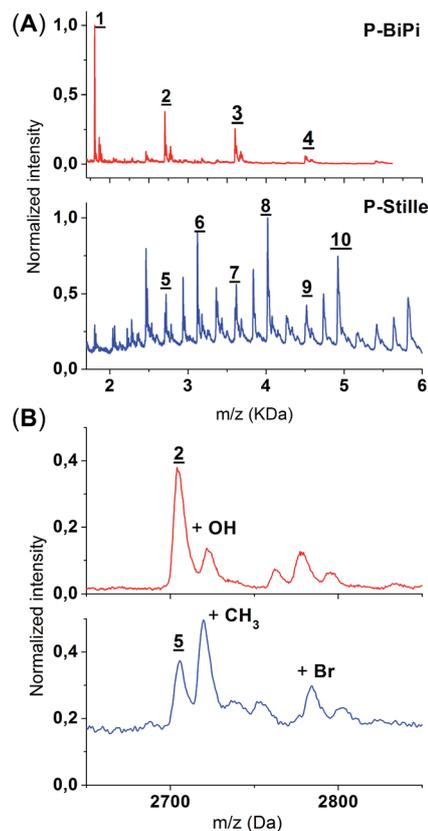


Fig. 5 (A) Comparison of the MALDI-TOF spectra of **P-Stille** and **P-BiPi**. Peaks of interest are marked with numbers. Peaks 1–4 (peak-to-peak interval: 900 Da) of polymer **P-BiPi** correspond to dimer to pentamer with hydrogen end groups as the homopolymerization result. Peaks 5, 7, 9 of polymer **P-Stille** correspond to peaks 2–4 of **P-BiPi**. The peak-to-peak interval of $5 \rightarrow 6$, $7 \rightarrow 8$ and $9 \rightarrow 10$ is 416, which indicates the presence of an extra donor moiety in the backbone. Other peaks are not convincingly assignable. Dithranol is used as the matrix for both spectra. (B) A zoom-in of peaks 2 and 5. The zoom-in reveals that in the case of the Stille copolymerization, there are peaks representing polymers + methyl group. This is mostly likely due to methyl transfer from the trimethyltin monomer. Residual bromo end groups are visible.

residual reactive Br end groups, which are clearly visible in the MALDI-TOF spectrum. Although homocoupling of organostannanes in conjugated polymer formation has been highlighted elsewhere,¹² we did not directly observe it in our case. We note that homocoupling of organostannanes should not be a noticeable problem in this case, primarily owing to the usage of a Pd(0) instead of a Pd(II) catalyst. Upon zooming in around peaks 2 and 5, methyl transfer from trimethyltin is evidenced with a clear peak,^{12,36} as shown in Fig. 5(B). This highlights the potential drawbacks of routine Stille copolymerizations that potentially hinder the achievement of high quality conjugated polymers. Future synthetic effort should be dedicated to diminish these structural flaws.

Photovoltaic characterization

The photovoltaic performance of these two polymers was compared by fabricating bulk heterojunction solar cells under



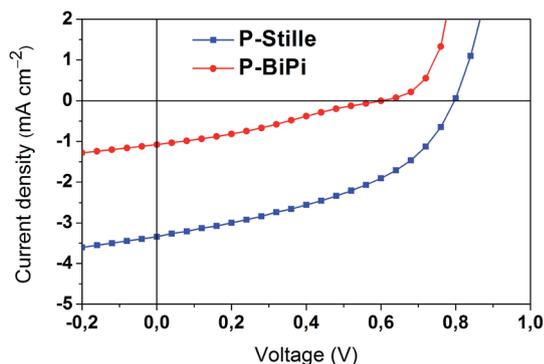


Fig. 6 Current–Voltage curves of the P-Stille and P-BiPi-based bulk heterojunction solar cells with DIO additive, under AM1.5 solar illuminations.

the same conditions, with a device structure of ITO/PEDOT:PSS/polymer:[60]PCBM/LiF/Al. AFM pictures (Fig. S4†) of the active layer indicate that both polymers can be mixed with [60]PCBM without noticeable phase separation, though **P-Stille** seems to be slightly aggregated in the blend. Despite the better chemical integrity of the homopolymerized **P-BiPi**, the random donor–acceptor configuration has notably reduced the absorption of the incoming light spectrum, which is indicated by the drastically lower molar extinction coefficient per repeating unit of **P-BiPi** than **P-Stille**, from 550 nm on (Fig. S5†). We believe this is largely responsible for its lower photocurrent of 1.08 mA cm^{-2} , compared to the 3.34 mA cm^{-2} of the **P-Stille** device, as shown in Fig. 6. The random donor–acceptor configurations in **P-BiPi** likely impair effective intermolecular packing, resulting in a lower hole mobility of $5 \times 10^{-8} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, compared to that of $6 \times 10^{-6} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ in the **P-Stille** case as measured by space charge limited current measurements (Fig. S6†). Note that the studied polymers were not optimized for high mobility, as it is well-known that a variety of structural variations may pose notable effect on this physical parameter. While the present research demonstrates the capacity of one-pot homopolymerization in producing polymers of enhanced chemical integrity, its translation into improved physical characteristics shall be achieved in conjunction with dedicated consideration on physical aspects. Overall, with a low photovoltage of 0.6 V and fill factor of 29%, the **P-BiPi**-based device showed a power conversion efficiency of 0.99%, while **P-Stille** generated a power conversion efficiency of 1.66% with a photovoltage of 0.79 V and a fill factor of 44%.

Conclusions

In conclusion, we have extended the concept of homopolymerization based on *in situ* borylation/cross coupling. This polymerization route is appealing in the sense that it not only completely eliminates the necessity to synthesize high-purity boron agents, but also can potentially produce high molecular weight polymeric materials through efficient polymerization, which is evidenced by our kinetic investigation. Yet when applied to head-to-tail asymmetrical donor–acceptor

macromonomers rather than simple, symmetrical monomers, such homopolymerizations tend to introduce donor–donor, acceptor–acceptor segments into the polymer backbone, apart from the expected alternating donor–acceptor structure. This type of structural complication was shown to occur in the homopolymerization by correlating the experimental UV-vis absorptions with calculated absorptions from TD-DFT. Nevertheless, MALDI-TOF measurements showed the capacity of this homopolymerization in producing polymers with improved structural integrity, compared to the standard Stille copolymerization which most likely suffers from destannylation and/or methyl transfer, affording moderate molecular weights even with prolonged reaction time. This work is to lead to new strategies for the synthesis of high-quality donor–acceptor conjugated polymers, which shall feature improved chemical integrity and high molecular weights.

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References

- Z. Bao, W. K. Chan and L. Yu, *J. Am. Chem. Soc.*, 1995, **117**, 12426–12435.
- G. Yu, J. Gao, J. C. Hummelen, F. Wudl and A. J. Heeger, *Science*, 1995, **270**, 1789–1791.
- S. Günes, H. Neugebauer and N. S. Sariciftci, *Chem. Rev.*, 2007, **107**, 1324–1338.
- Y. Liang, Z. Xu, J. Xia, S.-T. Tsai, Y. Wu, G. Li, C. Ray and L. Yu, *Adv. Mater.*, 2010, **22**, E135–E138.
- J. Pei, W.-L. Yu, W. Huang and A. J. Heeger, *Macromolecules*, 2000, **33**, 2462–2741.
- J. Mei, D. H. Kim, A. L. Ayzner, M. F. Toney and Z. Bao, *J. Am. Chem. Soc.*, 2011, **133**, 20130–20133.
- D. T. McQuade, A. E. Pullen and T. M. Swager, *Chem. Rev.*, 2000, **100**, 2537–2574.
- V. Farina, V. Krishnamurthy and W. J. Scott, *Organic Reactions*, John Wiley & Sons, Inc., New York, 1997, vol. 50, p. 46.
- T. Qin, W. Zajaczkowski, W. Pisula, M. Baumgarten, M. Chen, M. Gao, G. Wilson, C. D. Easton, K. Müllen and S. E. Watkins, *J. Am. Chem. Soc.*, 2014, **136**, 6049–6055.
- H. Zhong, C.-Z. Li, J. Carpenter, H. Ade and A. K.-Y. Jen, *J. Am. Chem. Soc.*, 2015, **137**, 7616–7619.
- K. H. Hendriks, W. Li, G. H. L. Heintges, G. W. P. van Pruissen, M. M. Wienk and R. A. J. Janssen, *J. Am. Chem. Soc.*, 2014, **136**, 11128–11133.
- T. Vangerven, P. Verstappen, J. Drijkoningen, W. Dierckx, S. Himmelberger, A. Salleo, D. Vanderzande, W. Maes and J. V. Manca, *Chem. Mater.*, 2015, **27**, 3726–3732.



- 13 N. Blouin, A. Michaud, D. Gendron, S. Wakim, E. Blair, R. Neagu-Plesu, M. Belletête, G. Durocher, Y. Tao and M. Leclerc, *J. Am. Chem. Soc.*, 2008, **130**, 732–742.
- 14 N. Blouin, A. Michaud and M. Leclerc, *Adv. Mater.*, 2007, **19**, 2295–2300.
- 15 H. Fan, Z. Zhang, Y. Li and X. Zhan, *J. Polym. Sci., Part A: Polym. Chem.*, 2011, **49**, 1462–1470.
- 16 F. Parenti, P. Morvillo, E. Bobeico, R. Diana, M. Lanzi, C. Fontanesi, F. Tassinari, L. Schenetti and A. Mucci, *Eur. J. Org. Chem.*, 2011, **28**, 5659–5667.
- 17 A. Giroux, Y. Han and P. Prasit, *Tetrahedron Lett.*, 1997, **38**, 3841–3844.
- 18 L. Zhu, J. Duquette and M. Zhang, *J. Org. Chem.*, 2013, **68**, 3729–3732.
- 19 G. Cuny, M. Bois-Choussy and J. Zhu, *Angew. Chem., Int. Ed.*, 2003, **42**, 4774–4777.
- 20 C. F. Nising, U. K. Schmid, M. Nieger and S. Bräse, *J. Org. Chem.*, 2004, **69**, 6830–6833.
- 21 N. Ma, Z. Zhu and Y. Wu, *Tetrahedron*, 2007, **63**, 4625–4629.
- 22 P. B. Dzhevakov, M. A. Topchiy, D. A. Zharkova, O. S. Morozov, A. F. Asachenko and M. S. Nechaev, *Adv. Synth. Catal.*, 2016, **358**, 977–983.
- 23 A. Izumi, R. Nomura and T. Masuda, *Chem. Lett.*, 2000, **29**, 728–729.
- 24 T. Ishiyama, K. Ishida and N. Miyaura, *Tetrahedron*, 2001, **57**, 9813–9816.
- 25 R. M. Walczak, R. N. Brookins, A. M. Savage, E. M. van der Aa and J. R. Reynolds, *Macromolecules*, 2009, **42**, 1445–1447.
- 26 F. Brouwer, J. Alma, H. Valkenier, T. P. Voortman, J. Hillebrand, R. C. Chiechi and J. C. Hummelen, *J. Mater. Chem.*, 2011, **21**, 1582–1592.
- 27 J. Mei, K. R. Graham, R. Stalder and J. R. Reynolds, *Org. Lett.*, 2010, **12**, 660–663.
- 28 C. M. Amb, S. Chen, K. R. Graham, J. Subbiah, C. E. Small, F. So and J. R. Reynolds, *J. Am. Chem. Soc.*, 2001, **133**, 10062–10065.
- 29 J. Hou, H.-Y. Chen, S. Zhang, G. Li and Y. Yang, *J. Am. Chem. Soc.*, 2008, **130**, 16144–16145.
- 30 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 73–78.
- 31 J.-L. Brédas, J. Cornil and A. J. Heeger, *Adv. Mater.*, 1996, **8**, 447–452.
- 32 S. D. Hanton, *Chem. Rev.*, 2001, **101**, 527–570.
- 33 M. Jayakannan, J. L. J. van Dongen and R. A. J. Janssen, *Macromolecules*, 2001, **34**, 5386–5393.
- 34 S. M. Weidner and S. Trimpin, *Anal. Chem.*, 2008, **80**, 4349–4361.
- 35 V. Farina, V. Krishnamurthy and W. J. Scott, *Organic Reactions*, John Wiley & Sons, Inc., New York, 1997, vol. 50, p. 48.
- 36 V. Farina, V. Krishnamurthy and W. J. Scott, *Organic Reactions*, John Wiley & Sons, Inc., New York, 1997, vol. 50, p. 47.

